Tracing fast roads towards bone regeneration

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Tracing fast roads towards bone regeneration:
Strategies to augment the bioactivity of additive
manufactured scaffolds

María Cámara Torres

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Tracing fast roads towards bone regeneration: Strategies to augment the bioactivity of additive manufactured scaffolds

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Tracing fast roads towards bone regeneration: Strategies to augment the bioactivity of additive manufactured scaffolds

DISSERTATION

To obtain the degree of Doctor at the Maastricht University, on the authority of the Rector Magnificus, Prof.dr. Rianne M. Letschert in accordance with the decision of the Board of Deans, to be defended in public on Monday 13th September 2021, at 13:00 hours.

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Chapter 1

General introduction and aims

As one of the most frequently damaged tissues in our body, bone has always been in the spotlight of clinicians and scientists. In principle, bone tissue can self-repair. However, there are some cases, in which the injury is too large and severe, or the local environment is suboptimal for this process to happen on its own demanding external help to manage the injury. Unlike the axolotl, the human body does not have the unique ability to perfectly self-regenerate after complex damage.

For decades, the use of autografts, allografts or metallic prosthesis have been considered as the gold standard clinical practices to reconstruct large bone defects or non-unions. However, these approaches have never been considered optimum solutions, due to donor morbidity, immunogenicity, unavailability and/or structural failure. With the birth of the tissue engineering field, which aims at combining biomaterials, cells and/or biophysical and chemical signals to promote endogenous tissue regeneration, a new research perspective appeared to confront non-union bone injuries. Since then, a wide variety of novel approaches have been suggested towards a more effective clinical solution. Most of these revolve around the design of an ideal scaffold, which upon implantation can serve as a temporary structure mimicking bone extracellular matrix (ECM) to promote cell adhesion, proliferation, differentiation and new tissue formation. Several biomaterials and biofabrication techniques have been investigated trying to create a scaffold fulfilling the required biodegradability, bioactivity and mechanical properties. In this regard, natural and synthetic polymers, ceramics, metals, and their combinations have been processed through a variety of fabrication methods, including particulate leaching, gas foaming, thermally induced phase separation, freeze-drying, electrospinning and additive manufacturing (AM). Among all, AM, and melt extrusion AM (ME-AM) in particular, has stood out over the other biofabrication techniques, as it allows the reproducible control over scaffold size, shape, internal interconnected macroporosity, which are crucial factors for bone tissue engineered (BTE) scaffolds success. In spite of these benefits, ME-AM possess also its own drawbacks, associated with the inherent characteristics of synthetic polymers, which are the only processable materials through this technique and generally lack the required bioactivity to induce bone regeneration. In **Chapter 2** we summarize the research that has been carried out over the last decades on strategies to augmenting the physicochemical properties of such ME-AM scaffolds, in order to enhance their interaction with cells. These include modifications of the scaffold internal architecture and surface properties, the fabrication of hybrid scaffolds, combining the ME-AM framework with fibrous or hydrogel networks, and the use of polymer composites containing inorganic,

organic or metallic fillers. In spite of all these efforts, the search for optimal scaffolds that can boost bone regeneration continues, since an ideal ME-AM scaffold that shows an optimal *in vivo* tissue regeneration has not yet been found. To further contribute to the field, and by overcoming some of the identified challenges, this thesis aims at investigating different material-driven approaches to design and fabricate bioactive ME-AM scaffolds for BTE. These approaches are described in the different chapters of the thesis.

The first step towards the short and long-term functionality of a scaffold-supported engineered tissue is an efficient scaffold seeding. However, cell seeding on ME-AM scaffolds is challenging, due to the interplay of macroporosity, inert scaffold surface properties, and gravity. **Chapter 3** and **Chapter 4** propose two different alternatives to efficiently seed ME-AM scaffolds in terms of seeding efficiency and distribution. While some of the pre- or post-scaffold fabrication modification techniques presented in **Chapter 2** have been used on an attempt to circumvent these intrinsic cell seeding difficulties, there is still the need of a simple and reliable method that does not depend on the scaffold architecture or its material physicochemical properties. In this regard, **Chapter 3** describes a simple, unique and universal approach to statically seed ME-AM scaffolds in a precise and reproducible manner, regardless of the scaffold type. In this chapter, the independent addition to the cell seeding medium of two biocompatible macromolecules, dextran and Ficoll, is studied to adjust the medium viscosity and density, respectively, and tune cell settling velocity for better attachment during the seeding process. The influence of these seeding media on cell attachment and distribution along the scaffolds' cross section, as well as on the osteogenic differentiation capacity of human mesenchymal stromal cells (hMSCs), is evaluated.

In addition to the simple strategy presented in **Chapter 3**, the method described in **Chapter 4** relies on the use of a high-end technology for attaining enhanced cell attachment on ME-AM via plasma modification. This technology, which combines a ME printhead and an atmospheric pressure plasma jet (APPJ) module, not only allows scaffold production and treatment in a single-step, but also ensures a controlled and homogeneous surface modification, in contrast to conventional post-fabrication plasma approaches. Here, ME-AM polymeric scaffolds are plasma-polymerized with (3-Aminopropyl) trimethoxysilane and maleic anhydride-vinyltrimethoxysilane, which deposit rather stable amine- and carboxyl-based coatings on the scaffolds filaments. The effect of these uniquely performed

treatments on scaffolds' hydrophilicity, protein adhesion, cell adhesion mechanism and hMSCs osteogenic differentiation is evaluated.

After ensuring an efficient cell attachment, composite ME-AM scaffolds comprising the copolymer PEOT/PBT and either nanohydroxyapatite (nHA), reduced graphene oxide (rGO), or antibiotic-loaded lamellar fillers, are fabricated in **Chapters 5, 6 and 7**, respectively. Among all the strategies for augmenting the bioactivity of ME-AM polymeric scaffolds, the use of composites enable bulk scaffold properties modification (i.e. mechanical, physicochemical, electrical) without altering the scaffold architecture or the need of post-fabrication steps. To replicate the organic-inorganic composition of bone ECM, and taking advantage of the osteoinductive properties of calcium and phosphate, composite constructs comprising polymers and calcium phosphate materials, and in particular nHA, have been widely explored. Despite material advances and thorough studies on the printability of highly loaded nHA ME-AM composite scaffolds, the biological assessment of such scaffolds has been mostly overlooked and, in general, poorly characterized. In **Chapter 5**, we aimed to fill this important gap and provide an in depth *in vitro* biological characterization of highly loaded nHA scaffolds. Polymer-nHA composite scaffolds with up to 45 wt% nHA were fabricated, and the effect of nHA content on the osteogenic differentiation of hMSCs was intensively evaluated using several biochemical and molecular assays, such as q-PCR, immunofluorescence, ELISA and calcium staining. In addition, a mechanical analysis was also performed.

Towards the fabrication of ME-AM scaffolds with outstanding physicochemical and electroactive properties for bone formation stimulation, graphene based materials can be used as fillers. In spite of its thermal stability, electroconductivity and easy production, compared to other graphene derivative, rGO has been mostly overlooked as filler within ME-AM composite scaffolds due to its low bulk density compromising its safe handling during melt compounding. In order to overcome this limitation, rGO was densified in **Chapter 6**, and the effect of densification on rGO compaction degree and on the resulting melt compounded composites physicochemical properties and printability via ME-AM was assessed. In addition, the effect of rGO concentration on the materials' physicochemical properties and cell-material interactions in 2D and 3D, including osteogenic differentiation, was also investigated.

Bone infection is a common complication after trauma or implant surgery. On an attempt to circumvent the use of high doses of systemic antibiotic therapy, local antibiotic delivery

systems are desired. However, approaches to fabricate ME-AM scaffolds with antimicrobial properties are based either on post-fabrication and multi-step surface adsorption treatments, which lead to poor loading efficiencies and uncontrolled release kinetics, or on low temperature ME-AM (< 120 °C) to preserve the bioactive molecules functionality, which limits the polymer choice. In **Chapter 7**, the possibility of incorporating antibiotics within the polymer matrix of scaffolds fabricated by high temperature ME-AM (~190 °C), via antibiotic-loaded inorganic fillers, is discussed. In this chapter, two different antibiotic-nanofiller complexes are investigated, and the antibiotic release kinetics, drug stability after printing and scaffold antimicrobial properties against relevant orthopedic bacterial strains, evaluated. In addition, the osteogenic differentiation capability of hMSCs on the composite scaffolds is assessed to ensure their dual functionality.

In **Chapter 8**, the results described in the aforementioned chapters are discussed and place into the state of the art context, while providing future perspectives. To finalize, a reflection on the scientific and social impact of the research carried out in this thesis is presented in **Chapter 9**.

Chapter 2

Boosting bone regeneration using augmented melt-extruded additive manufactured scaffolds

Abstract

The bone tissue engineering (BTE) field is in active search of the ideal scaffold to give a clinical solution for bone regeneration in non-union fracture scenarios. During the last decades, the use of additive manufacturing (AM) technologies and, in particular, melt extrusion AM (ME-AM), has been thoroughly investigated towards this aim. ME-AM enables the cost-effective fabrication of personalized 3D scaffolds, with a controlled and highly interconnected porosity, through the solvent-free processing of biodegradable and mechanically robust thermoplastic polymers. In addition to these properties matching the requirements for BTE scaffolds, the synthetic polymers used to fabricate these constructs are also more amenable for further functionalization than metals or ceramics, to influence cell behavior, making thermoplastic materials a preferred choice for BTE. This review provides a comprehensive analysis of various ME-AM scaffolds developed for BTE, along with the approaches used to augment their bioactivity, which include architectural modifications, surface physical and chemical modifications, the incorporation of secondary fibrous or hydrogel networks within the scaffolds pores, and the use of filler-polymer composites for ME-AM scaffold fabrication. A detailed examination on how each of these strategies affect the final scaffold physicochemical and mechanical properties is presented. Special emphasis is placed on discussing the effect of these modifications on promoting cell osteogenic differentiation *in vitro* and bone tissue development *in vivo*.

Bone and its regenerative capacity

Bone is a complex hierarchically structured tissue that plays a critical role in human physiology including locomotion, organ protection, blood production, mineral storage and homeostasis, and source of mesenchymal stem cells in the marrow compartment.¹ The building blocks of the bone matrix consist of an inorganic phase of carbonated hydroxyapatite (HA), and an organic phase mainly composed of collagen type I (COL I), among other types of collagens, proteoglycans, fibronectin and specific bone proteins, such as osteopontin (OPN), bone sialoprotein (BSP) and osteocalcin (OCN), in minor percentages.² In the cortical bone, which extends along the shaft or diaphysis of long bones and becomes thinner towards the distal and proximal ends or epiphysis, mineralized collagen nanofibers organize parallel into concentric lamellae and surround the Haversian channel constituting the osteons.³ On the other hand, cancellous bone, which is present in long bones mostly in their epiphysis, is organized into irregular and convoluted lamellae, which create an intricate porous trabecular framework as housing for the soft and highly cellularized bone marrow.³ Due to their differences in porosity and architecture, cortical and cancellous bone possess an elastic modulus at the macroscale of 15 – 20 GPa and 0.1 – 2 GPa, and a compressive strength of 100 – 200 MPa and 2 – 20 MPa, respectively.⁴⁻⁵ These properties are highly dependent on age, health condition, anatomic site and loading direction.^{4, 6-8} At the tissue level, cortical bone possess slightly higher density and mineral content compared to trabecular bone, as it presents a lower rate of turnover.⁹ Due to this, microtensile measurements have also revealed differences in the elastic modulus of both tissues at the microscale (~18 GPa for cortical bone vs. ~ 11 GPa for cancellous bone). Yet, they are in the same order of magnitude, due to their similar lamellar nanostructure.⁸ These impressively high mechanical properties are owed to the mineralization of the collagen fibers, which provides stiffness and structural reinforcement to the viscoelastic and tough collagen.

Bone is constantly undergoing remodelling to adapt to mechanical stresses, resulting from physical exercise or loading, and to maintain the integrity of the skeleton, by repairing micro fractures caused by the bending and torsion of bones during locomotion.¹⁰ This is a dynamic process orchestrated by a variety of biochemical and mechanical factors that requires the interplay between different cell phenotypes: the bone-forming osteoblasts, the bone-degrading osteoclasts, and the bone lining cells and the osteocytes.¹¹ In line with its remodeling capacity, bone also has self-healing abilities after fracture.¹² In bone fractures,

not only the structural integrity of bone is disturbed, but also the vascular structures, limiting the nutrients flow at the injury site. Therefore, the healing process is much more complex than during spontaneous remodelling, and includes an inflammatory phase, a soft callous formation, and a remodelling phase, involving multiple cellular types, including mesenchymal stromal cells, and molecular activities that recapitulate the endochondral ossification happening during embryogenesis.¹³ However, improper stabilization, age and comorbidities of the patient, as well as large bone defects, caused by disease, traumatic injury, implant surgery, infection or tumor resection, can negatively influence the healing process, leading to non-union in up to 10% of the cases.¹⁴ Current strategies used in the clinics for the treatment of non-union or critical size bone defects mainly rely on: i) the distraction osteogenesis or Ilizarov method (**Figure 1A**),¹⁵ ii) the induced membrane or Masquelet technique,¹⁶⁻¹⁷ combined with autografts, such as intramedullary bone grafts (**Figure 1B-C**), iliac crest, or vascularized fibula graft (**Figure 1D**)¹⁸ iii) allografts, or iv) demineralized bone matrices.¹⁹⁻²⁰ Treatments based on the Ilizarov or Masquelet methods are often complex, requiring external fixation and multiple surgeries. Similarly, natural bone grafts (autografts or allografts) possess several disadvantages, including donor site morbidity, restricted availability, high costs, pain, prolonged rehabilitation, lack of structural properties, and can trigger inflammation, immunological rejections in the case of allografts, or infections.¹ Therefore, the management of non-union bone defects remains a major clinical challenge.

During the past decades, a large amount of efforts have been dedicated to investigate tissue-engineered strategies to regenerate bone in non-union scenarios, and avoid the aforementioned disadvantages associated with the currently used therapies.²¹ Particular attention has been given to additive manufacturing (AM) technologies, which enable to finely control the scaffold properties. In the following sections of this chapter, a brief overview on current AM techniques used to fabricate scaffolds for bone tissue engineering (BTE) applications will be provided, and special focus will be given to the advantages of melt extrusion AM (ME-AM) over other technologies. This includes the compatibility of the synthetic polymers used for ME-AM with various supplemental methodologies. While making this choice, we acknowledge the promising bone regeneration capacities of ceramic and biodegradable metal scaffolds, however they offer suboptimal biomechanical properties, being either too brittle or too stiff, respectively, compared to polymeric scaffolds. In this regard, current state-of-the-art strategies used to augment the bioactivity of polymeric ME-AM scaffolds towards an ideal BTE scaffold will be extensively discussed.

The related *in vitro* cell responses and *in vivo* bone regeneration produced by these strategies will be highlighted.

Figure 1 - Current gold standard techniques for the treatment of non-unions. (A) Schematic representation of the distraction osteogenesis technique. After the formation of a critical size defect, a bone segment is fractured and brought into close contact with the bone. The Ilizarov apparatus (external fixation) is used to gradually separate the bone segment, allowing new bone to form in the gap via conventional endochondral ossification during fracture healing. Adapted with permission from Ref. ²² (B) Clinical case in which the Masquelet technique was applied to treat a humeral non-union. (i) After debridement of the bone ends, a cement spacer was inserted for 2 months. (ii) After this period, a pearly membrane was induced around the cement, which was filled with a bone graft after spacer removal. Adapted with permission from Ref. ¹⁷ (C) Clinical case combining the Masquelet technique with an intramedullary bone graft to treat a tibial non-union. (i) After poly(methyl methacrylate) (PMMA) spacer removal and the induced membrane formation, the defect site was packed with autologous cancellous bone graft collected from the femur with the reamer-irrigator aspirator (RIA) technique. (ii) 7 months post-surgery X-rays. (D) Schematic representation of the vascularized fibula approach combined with a supporting construct comprising the following steps: resection of malignant tumor, construct placed around the vascularized fibular graft, insertion in the critical size defect and use of internal fixation. Adapted with permission from Ref. ²¹

Towards AM scaffolds for BTE applications

One of the pillars of tissue engineering (TE) are scaffolds. An ideal scaffold should provide a physico-chemically biomimetic environment to support and boost bone tissue ingrowth and regeneration. ²³ In this regard, the use of biocompatible materials to avoid the risks of local toxicity and undesired immune/inflammatory responses is fundamental. Moreover, the scaffold degradation and the bone formation rates should match, ensuring the stability of the defect post implantation. At the same time, scaffolds should provide mechanical support and stability, ideally matching the mechanical properties of neighboring bone

tissue to prevent stress shielding.²⁴ Surface properties such as chemistry, surface charge and topography are also critical, as they will dictate cell–material interactions for bone tissue ingrowth. While the aforementioned properties are of utmost importance, the architectural parameters of the scaffold i.e. pore interconnectivity, porosity, and pore size and shape, will ultimately determine the success of the scaffold in promoting tissue ingrowth leading to the regeneration of a fully functional tissue. Interconnected pores are necessary for efficient management of nutrients, oxygen, metabolic waste and material biodegradation products, as well as for continuous vascularization and tissue ingrowth within the scaffold. Moreover, scaffolds with pore sizes of at least 300 µm are needed to promote neovascularization and favor osteogenesis *in vivo*.²⁵⁻²⁷ Pore shape and curvature has also shown to influence bone formation by modulating cell response and rate of tissue regeneration.²⁸⁻²⁹ Since it is difficult to decouple porosity from mechanical properties and degradation rate, all these parameters need to be carefully balanced in order to optimize scaffolds for bone regeneration.^{25, 27}

It is already established that conventional scaffold fabrication techniques, such as particulate leaching, gas foaming, phase separation, solution casting or freeze drying, cannot ensure pore interconnectivity, nor the precise control over pore size, shape and spatial distribution.³⁰⁻³¹ On the contrary, biofabrication technologies derived from AM techniques for rapid prototyping, have demonstrated their potential to generate reproducibly complex-shaped tissue engineered constructs with a high degree of control on architectural parameters. Depending on the material type and its processing method, the most commonly used AM technologies in biofabrication of BTE constructs can be classified into:³²⁻³³

- i) powder based techniques, including 3D printing, selective laser sintering (SLS), selective laser melting (SLM) and electron beam melting (EBM),
- ii) vat photopolymerization techniques, including stereolithography (SLA), digital light processing (DLP) and two-photon polymerization (2PP),
- iii) ME techniques, including fused deposition modeling (FDM), 3D fiber deposition (3DF) and bioextrusion,
- iv) solution extrusion, including direct ink writing, low temperature deposition manufacturing (LDM), pressure-assisted microsyringe (PAM), and wet-spinning, and

- v) bioprinting, including bioplotting, ink-jet bioprinting and laser-assisted bioprinting.

Among all the aforementioned, ME-AM techniques are the most used AM techniques in BTE. Here, a thermoplastic polymer or composite in the form of a filament (FDM) or pellets (3DF and bioextrusion) is hot-extruded in molten form through a nozzle upon application of pressure via rollers pushing the filament (FDM), pressurized gas/piston (3DF) or a screw (bioextrusion), and deposited as solidified strut in a layer-by-layer manner.³⁴ Compared to other AM technologies, ME-AM does not require demanding hardware, its operation principle is straightforward, and the materials used do not require extensive development, in terms of chemistry. Moreover, ME-AM techniques offer one of the highest resolution-to-time of manufacturing (RTM) ratio among all AM techniques ($\sim 1 \times 10^{-3} \text{ m}^2/\text{minute}$). RTM is a parameter that defines the efficiency of the manufacturing process in terms of the ability to produce scaffolds with fine details in a short time, and is expressed as volume of material deposited per minute, divided by the minimum printed feature dimension.³³ An RTM ratio of $\sim 1 \times 10^{-3} \text{ m}^2/\text{minute}$ is also estimated for SLS, where a high power laser selectively sinters a powder bed layer-by-layer. In spite of producing scaffolds at a faster rate, SLS offers lower resolution than ME (400 μm vs. 200 μm).³³ Moreover, SLS and other powder-based methods such as 3D printing, which is based on the controlled deposition of a binder material on successive powder layers, require post-fabrication heat treatments to complete the binding reactions and increase the construct mechanical properties, as well as a depowdering step to remove excess material.³⁵ Similarly, scaffolds fabricated by solution extrusion methods, in which the material in a solvent is extruded at low temperature, are followed by post-lyophilization (i.e. LDM) or drying steps (i.e. PAM and wet-spinning) to remove the organic solvent, which also leads to scaffolds shrinkage and loss of mechanical properties.³⁶⁻³⁷ Besides, the long time required to fabricate thick constructs by solvent extrusion, due to their low minimal feature size ($\sim 20 \mu\text{m}$), contributes to a low RTM ratio ($\sim 0.5 \times 10^{-3} \text{ m}^2/\text{minute}$).³³ Generally, the use of binders and organic solvents is by itself a limitation of all the aforementioned processes. Such additives could potentially induce cytotoxic effects if not well removed prior to their contact with cells. Likewise, uncured resin and photoinitiator remnants in scaffolds produced by vat photopolymerization techniques, which work by selectively crosslinking layers of a photosensitive resin by laser (SLA and 2PP) or UV light (DLP), may be toxic.³⁸ Despite their accuracy, ability to fabricate complex shapes without additional support, and high speed (especially DLP),³³ the questionable biocompatibility of photopolymers and photoinitiators, together with the limited availability

of such systems and their demanding rheological specifications, has delegated vat photopolymerization techniques to a secondary plane within the BTE field.³⁸⁻³⁹

Post-processing steps involving solvent and binder removal, or post-curing, are not necessary in ME-AM, as here the raw polymeric material is only subjected to melting and subsequent solidification. Moreover, most of the commonly used biodegradable thermoplastics for bone scaffolds fabrication, such as poly(ϵ -caprolactone) (PCL), poly(lactic acid) (PLA), and poly(lactic-co-glycolic) acid (PLGA), have already been approved for multiple applications by the Food and Drug Administration (FDA). These materials have been used for decades in the clinic, processed by melt-based techniques. Some examples include melt spun PCL or PLGA resorbable sutures,⁴⁰⁻⁴¹ or injection molded PLA resorbable screws for anterior cruciate ligament reconstruction.⁴² The record of accomplishment of melt-based techniques in medical device manufacturing further emphasizes the enormous translational potential of ME-AM scaffolds to the clinic. Examples of this are the 3D implant Osteoplug™ and the thin interwoven mesh Osteomesh™, from Osteopore International, which are the first marketed 3D AM medical devices for bone regeneration applications. These implants are fabricated by FDM using PCL and used as Class II medical devices for craniofacial applications. These products have their origin in the research of Hutmacher et al., who first described the fabrication of FDM scaffolds for TE applications using PCL and PCL/tricalcium phosphate (TCP) in 2000.^{23, 43} Promising *in vitro* research led to *in vivo* studies in large animal models and to first-in-human clinical studies to evaluate the feasibility of the PCL scaffolds for skull bone regeneration.⁴⁴⁻⁴⁷ Subsequent clinical studies were performed to study their safety and efficacy for alveolar ridge preservation after tooth extraction and orbital floor fractures reconstruction.⁴⁸⁻⁴⁹

PCL, together with other commonly used thermoplastics, such as PLA, PLGA, poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT) and thermoplastic polyurethane (TPU), are biocompatible, biodegradable, possess relatively high mechanical properties, can be easily processed, and can support *in vitro* and *in vivo* tissue ingrowth which make them excellent candidates for bone scaffold materials.⁵⁰⁻⁵⁴ However, these synthetic polymers generally lack bioactivity to stimulate cell and protein adhesion, which are necessary for an efficient cell differentiation and tissue formation. This is due to such polymers being relatively hydrophobic, with static contact angles in the range of 75° - 90°, and lacking sufficient reactive side-chain groups for influencing cell

behavior.⁵⁵⁻⁵⁶ For this reason, research has been focused over the last two decades on augmenting the bioactivity of ME-AM scaffolds for BTE made of synthetic polymers. These include i) modifications of the scaffold architecture, ii) modification of scaffolds filaments' surface, in terms of roughness, microporosity, chemistry or bioactive coatings, iii) the fabrication of hybrid scaffolds, combining the ME-AM framework with fibrous or hydrogel networks, and iv) the use of polymer composites containing inorganic, organic or metallic fillers. (*note*: while the terms 'hybrid' and 'composite' are broad, permitting overlaps in the above defined categories, here 'hybrid' will be used for scaffolds prepared using a combination of material classes or fabrication techniques, and 'composite' will be used for cases where a filler is loaded into the thermoplastic polymer).

Scaffold architecture modification

A major strategy in BTE is the fabrication of scaffolds mimicking the mechanical properties of human bone. ME-AM offers the possibility of modulating the porosity and, therefore, the mechanical properties of the scaffolds, by simply tuning layer-by-layer the fiber deposition pattern, in terms of fiber distance (FD) and deposition angle.⁵⁷ Mechanical analysis of scaffolds fabricated from various polymers (PCL, poly(ester) based TPU (PEU) and PEOT/PBT) has proven that for a given deposition angle, increasing the FD in XY or the layer thickness in Z, leads to an increase in the scaffolds' porosity and a decrease in stiffness (**Figure 2A**).⁵⁸⁻⁶⁰ A larger amount of deposited polymer, and the subsequent decrease of porosity within the same scaffold volume, explained higher compression modulus upon FD reduction. Similarly, minimizing the angle of deposition from 90° towards 15° while keeping porosity, FD and layer thickness constant, has also shown to decrease the mechanical properties of scaffolds, due to reduced contact points between fibers increasing the local stress experienced by the structure (**Figure 2B**).⁶¹ In an attempt to maximize the scaffold stiffness while maintaining an optimal porosity (~ 60%), PCL scaffolds with triply periodic minimal surface (TPMS) geometries were recently manufactured by ME-AM (**Figure 2C**).⁶² Among other geometries, gyroid scaffolds had a compression modulus of 60-80 MPa, higher than previously reported PCL scaffolds with similar porosity but standard geometries.^{61, 63} While the mechanical properties that can be attained with scaffolds fabricated with commonly used polymers, such as PCL, lies in the range of cartilage and cancellous bone mechanical properties (up to 120 MPa), other materials, such as PLA, can provide scaffolds with much higher mechanical properties

(compression modulus up to 450 MPa), which are more suitable for load bearing applications.⁶⁴

Besides having an impact on the scaffold's mechanical properties, the scaffold pore architecture has also shown to influence cell seeding efficiency and distribution on scaffolds, which can directly affect the tissue formation rate and uniformity, and cell differentiation behavior.⁶⁵⁻⁶⁶ Generally, achieving a high seeding efficiency on ME-AM scaffolds is difficult, due to the open pore architectures and the poor cell-adhesion sites of the scaffold materials. In this regard, scaffolds displaying a gradient in pore size along the vertical direction have shown to improve cell seeding efficiency and distribution, without compromising porosity (**Figure 2D**).⁶⁷ Cell differentiation has also been shown to be modulated by scaffold architecture. In this regard, Di Luca et al. showed that human bone marrow-derived mesenchymal stromal cells (hBM-MSCs) preferentially differentiate towards the osteogenic lineage, over chondrogenic differentiation, in PEOT/PBT scaffolds with high FD (900 - 1100 μm), compared to smaller pore size scaffolds (500 μm), according to increased alkaline phosphatase (ALP) activity and ECM mineralization.⁶⁸ This was in agreement with the study of Woodfield et al., where a higher glycosaminoglycan (GAG) and Collagen II production by chondrocytes was observed on small pore size PEOT/PBT scaffolds.⁶⁹ In addition, higher ALP activity and the upregulation of osteogenic related genes has also been shown to be better supported on PCL scaffolds with rhomboidal pore shapes, given by a 0/15° lay down pattern, compared to other filament orientation angles.⁷⁰ In comparison, square shaped pores (0/90°) supported enhanced chondrogenic differentiation of hBM-MSCs. This pore shape limited the local oxygen availability, inducing the expression of hypoxia inducible factors and chondrogenesis, compared to rhomboidal pore scaffolds with same porosity.⁷⁰ In contrast to this *in vitro* result, PCL/TCP scaffolds with a 0/90° laydown pattern implanted on a rat calvarial critical bone defect for 12 weeks revealed the highest amount of bone formation compared to scaffolds with other angles of deposition.⁷¹ *In vivo*, the optimum porosity and FD have also not been clearly defined, as some studies have reported significantly higher amount of new bone formation on PCL scaffolds with low FD (130 μm), compared to scaffolds with high FD (500 - 700 μm),⁷² while others have not observed differences in bone formation volume among PCL scaffolds with pore sizes ranging from 350 μm to 800 μm after 8 weeks of implantation.⁷³ In short, pore sizes and shapes achievable using ME-AM provide a suitable way to tune tissue regeneration and implant mechanics, but attention needs to be paid to not modify

another interdependent property and reverse the beneficial effects of tailored pores, as was possibly the case in the conflicting results cited above.

Figure 2 - Modification of the scaffold architecture. (A) (i) SEM images of the top view and cross section of PEU scaffolds with different angles of deposition. (ii) Compressive Young's moduli of PEU scaffolds with different FD and angles of deposition. Adapted with permission from Ref. ⁶⁰ (B) μ CT images of PCL scaffolds with different lay-down patterns. Adapted with permission from Ref. ⁶¹ (C) (i) PCL gyroid scaffold. (ii) Representative confocal microscopy images of human adipose derived MSCs (hAD-MSCs) cultured for 30 days on a PCL (calcium phosphate coated) gyroid scaffold. (F-actin, red). Adapted with permission from Ref. ⁶² (D) (i) SEM images of the cross section of a homogeneous PCL scaffold (no gradient, pore size 750 μ m), and a vertical pore size gradient PCL scaffold (750 μ m $\rightarrow$ 200 μ m $\rightarrow$ 750 μ m). (ii) Fluorescent microscopy images of the cross section of the aforementioned homogeneous and gradient scaffolds seeded with SaOs-2 cells. Adapted with permission from Ref. ⁶⁷

Scaffold surface modification

Since cell-scaffold interactions mainly occur at the interface, numerous techniques have been used to modify the surface of chemically inert ME-AM scaffolds, without altering the bulk scaffold material and architectural properties. These include wet chemical modifications, plasma treatment, and coatings with proteins, growth factors, calcium phosphate (CaP) or other bioactive molecules. These can alter the scaffold surface roughness, surface topography, hydrophilicity, chemical composition and bioactivity,

towards enhanced protein adsorption, and improved cell adhesion, proliferation and differentiation.

Introduction of roughness and microporosity

Surface roughness and nano/microtopography have been shown to enhance protein attachment and influence cell adhesion, proliferation and differentiation.⁷⁴⁻⁷⁶ However, the surface of the filaments of ME-AM polymeric scaffolds is rather smooth, due to rounding of free surfaces in the melt state before solidification, and surface roughness can be introduced via scaffold immersion in an etching solution. For instance, Kumar et al. etched PCL scaffolds by immersion in a dichloromethane-acetone solution post-fabrication (**Figure 3A**).⁷⁷ Scaffolds filaments showed a 5-fold increase in average surface roughness, which induced hBM-MSCs osteogenic differentiation and matrix mineralization after 62 days of culture under basic media conditions. This effect was attributed to a change of cell morphology, as cells in etched scaffolds presented a lower spread area and smaller size, similar to differentiated cells in previous studies,⁷⁸ compared to the elongated and undifferentiated cells on non-etched scaffolds. Similarly, an alkaline treatment with NaOH has been shown to provide changes in the surface microtopography of PCL scaffolds, as well as to introduce functional carboxyl and hydroxyl groups to the filaments surface.⁷⁹ In this case, a combination of physical and chemical changes led to an increase of ALP activity and calcium deposition by MC3T3-E1 pre-osteoblasts, compared to treated scaffolds. In spite of requiring post-etching solvent removal and washing, post-fabrication wet chemical surface modifications are considered greener compared to other techniques, such as solvent extrusion AM, in which the organic solvent is used during the scaffold manufacturing process and cytotoxic traces can remain within the material's core. Nevertheless, some studies have reported significant reduction of the filament diameter after solvent etching, which negatively affected the mechanical properties of the scaffold for the final application.⁸⁰ This suggests that special care should be taken on the selection of the treatment parameters, in terms of solution concentration and immersion time.

Alternative to the introduction of surface topography by solvent etching, several organic solvent-free methodologies have been used to induce microporosity both in the surface and in the bulk of ME-AM scaffolds' filaments. Microporosity has previously been shown to increase the permeability of the scaffold and modify its degradation rate, as well as to enhance the integration between scaffold and surrounding tissue, essential for the

mechanical stability of the implant.⁸¹⁻⁸² In this regard, ME-AM and porogen leaching have been combined to fabricate scaffolds with surface and intra-filament microporosity. For instance, elongated pores were formed along the long axis of the extruded filaments on TPU/Poly(vinyl alcohol) (PVA) scaffolds by composite printing followed by PVA dissolution in water.⁸³ These scaffolds, in combination with a CaP coating enabled higher cell proliferation rates, as well as higher ALP activity and COL I deposition than smooth scaffolds.⁸³ Similarly, a combination of ME-AM with salt leaching was used to fabricate dual porosity scaffolds for drug delivery applications (**Figure 3B**).⁸⁴ Microporosity led to a significant increase in scaffold surface area, enabling a more efficient cefazolin (an antimicrobial molecule) loading upon soaking, and larger quantities released per timepoint. An additional Gelatin methacryloyl (GelMA) coating allowed to dampen the initial burst release and scaffolds showed antimicrobial activity over time.⁸⁴ An alternative technique to create microporosity in ME-AM scaffolds is gas foaming. PCL/PVA composite scaffolds were subjected to a supercritical CO₂ gas foaming step, followed by their immersion in water to leach the PVA and create open pores of around 2-10 µm within and on the surface of the filaments.⁸⁵ Similarly, the combined extrusion of PLA with a foaming agent, which decomposed at the printing temperature for gas foaming, led to dual macro/microporous PLA scaffolds (micropore size 10-60 µm) for significantly enhanced cell attachment and proliferation.⁸⁶ A major limitation of the gas foaming step is the formation of a skin layer on the filaments surface, which hinders the exposure of the inner filament pores. Castano et al. showed that this layer can be removed and create surface porosity using the breath figure approach, which in turn requires the use of organic solvents.⁸⁷ This technique consists on the short immersion of the scaffold in an organic solvent in a moist atmosphere. During the evaporation of the thin solvent layer deposited at the polymer surface, water vapor condensates as droplets on the surface of the scaffolds. After the complete evaporation of water and solvent, a honeycomb like porosity appears. Another drawback derived from the introduction of microporosity within the scaffolds filaments by all the aforementioned techniques is the loss of scaffold mechanical properties.

Plasma treatment

Plasma surface modification has the advantage of altering both the surface roughness and chemistry of ME-AM scaffolds, thus conferring hydrophilicity and positive or negative charge to the filaments, without the use of solvents and maintaining their bulk properties.⁸⁹ Surface charge does not only promote protein adsorption for cell attachment and to

trigger proliferation and differentiation pathways via integrin binding,⁹⁰⁻⁹¹ but it can also induce cell attachment through direct electrostatic interactions between the surface chemical groups and the cell surface.⁹²⁻⁹³ Oxygen, air, argon and helium non-thermal plasma (NTP) activation has been shown to increase the hydrophilicity of PCL or PLA ME-AM scaffolds.⁹⁴⁻⁹⁷ This was achieved due to plasma etching, which increased the filaments surface roughness, and due to the introduction via post oxidation steps of oxygen containing functional groups (mostly peroxide and hydroperoxide groups). Similarly, ethylene/nitrogen and ammonia plasma activation have shown to directly deposit nitrogen containing groups (amines) on PCL⁹⁸ and polystyrene (PS) scaffolds,⁹⁹ respectively, also leading to a significant increase on the scaffold wettability. Compared to untreated scaffolds, plasma activated ones displayed enhanced cell attachment and better cell distribution along scaffold cross section, regardless of the plasma gas or cell type used (MG63 osteosarcoma cell line, 7F2 mouse osteoblast, human Saos-2 osteoblast or hBM-MSCs). In addition, enhanced cell proliferation and early osteogenic differentiation in differentiation media cultures was also observed in some cases, when compared to untreated scaffolds.^{94-95, 99} Although in 3D scaffolds it has not been proven so far, cell differentiation on plasma modified 2D surfaces has been previously correlated to an enhanced integrin binding exposure of adhered proteins, as well as to high levels of recruitment of focal adhesion components and phosphorylation of focal adhesion kinases in the adhered cells, which are events required for osteoblast differentiation.^{90, 100}

To obtain a more unique and stable coating with higher density of specific functional groups, the plasma discharge can be fed with monomers, such as alkyl amine or acrylic acid, to coat the surface with a thin amorphous polymeric film containing primarily amine or carboxyl groups, respectively. Cools et al. have reported the use of such a plasma process, referred to as plasma polymerization, to treat ME-AM scaffolds for cartilage TE.¹⁰¹⁻¹⁰² Here, acrylic acid plasma polymerization successfully deposited carboxyl groups on the surface of PCL and PEOT/PBT scaffolds filaments, which enhanced cell adhesion of human adipose derived MSCs (hAD-MSCs), as well as proliferation and GAG production by chondrocytes, respectively, compared to untreated and helium activated scaffolds.¹⁰¹⁻

Figure 3 - Modification of the scaffold surface. (A) (i, ii) Introduction of surface roughness to PCL scaffolds filaments by post-fabrication solvent etching (dichloromethane:acetone). (iii) Alizarin red S (ARS) staining (calcium deposition) on the aforementioned unetched and etched scaffolds seeded with hBM-MSCs and cultured for 60 days in basic medium (without osteogenic factors). Insets correspond to the unseeded scaffolds cultured under the same media conditions. Adapted with permission from Ref. ⁷⁷ (B) (i, ii) Introduction of microporosity to the filaments of PCL scaffolds by combining ME-AM with salt leaching. SEM images depicting the scaffolds surface and cross section. (iii) μ CT images of solid and microporous scaffolds at representative cross sectional planes. Adapted with permission from Ref. ⁸⁴ (C) (i) Hybrid AM platform consisting of a ME printhead and an atmospheric plasma jet for whole scaffold treatment from top, layer-by-layer, or zonally in specific regions of interest within the scaffold volume. (ii) Selective maleic anhydride-Vinyltrimethoxysilane (MA-VTMOs) plasma polymerization treatment in a central spot of a PEOT/PBT scaffold using the hybrid AM platform. The patterned region was visualized using methylene blue staining. (iii) Specific hBM-MSCs attachment (F-actin, green) to an argon plasma-patterned scaffold. Adapted with permission from Ref. ⁸⁸

Despite promising results in enhancing ME-AM scaffolds surface bioactivity, NTP processes are usually carried out at sub-atmospheric pressures, and require costly vacuum and plasma chamber equipment. Most importantly, inhomogeneous functionalization along the scaffold volume, already observed for conventionally fabricated scaffolds, ¹⁰³⁻¹⁰⁵ has been also pointed out in some reports, where ME-AM scaffolds were plasma treated. For instance, lower oxygen atomic concentration has been observed on the filaments within the core of PCL scaffolds, with respect to the edges, after helium plasma activation and acrylic acid plasma polymerization. ^{95, 101} In addition, the average oxygen content over the scaffold cross section decreased with decreasing scaffold pore size. ¹⁰¹ Since plasma does not ignite inside the pores, the treatment of the core filaments

is limited to diffusion, which drops while entering the scaffold and with decreasing pore size. This leads to a prolonged residence time of the monomer in the active plasma discharge, which in turn leads to a more extensive fragmentation of the precursor, and thus a lower oxygen incorporation.¹⁰¹ The use of atmospheric pressure plasma jets (APPJ) and their assembly within a ME-AM printing platform eliminates the need of a reactor chamber and vacuum equipment, avoids the formation of unwanted gradients, and enables plasma treatment during the manufacturing step.¹⁰⁶ With such a hybrid setup, the flame exiting the jet can be directed into the scaffolds pores, and the scaffold can be treated as a whole from the top, layer-by-layer to reach deep pores in large anatomical scaffolds, or zonally in specific regions of interest within the scaffold volume. Two recent reports have explored these possibilities, employing oxygen plasma activation to treat PCL scaffolds,¹⁰⁷ and plasma polymerization of maleic anhydride-Vinyltrimethoxysilane (MA-VTMOS) on PEOT/PBT scaffolds, which deposits thin films containing carboxyl functional groups (**Figure 3C**).⁸⁸ In the latter study, selective plasma treatment in a central spot within the scaffold core enabled specific cell attachment to the treated region. The subsequent dispersion of a second cell population throughout the scaffold volume showed the cell-patterning potential of this hybrid technique, with applications in cell co-culture experiments.⁸⁸

Coatings with proteins, growth factors and other bioactive molecules

The coating of the scaffold's filament surface with proteins, peptides, growth factors or antimicrobial molecules can stimulate cell attachment and differentiation, or provide the scaffold with antimicrobial properties. The simplest method for bioactive molecules incorporation is surface adsorption upon scaffold soaking into a solution.¹⁰⁸⁻¹¹⁰ For instance, the adsorption of the antimicrobial peptide polylysine onto PCL/HA scaffolds rendered their surface antibacterial.¹¹¹ However, clear disadvantages of this technique are the requirement of potentially expensive highly concentrated solutions with the molecule of interest, the very low loading efficiency, and the loose binding to the polymer resulting in burst and uncontrolled desorption, which can lead to cytotoxicity effects or poor bioactivity in the long term. For instance, 50% of the total dexamethasone impregnated on PLA scaffolds by soaking was released within the first 6h upon immersion in phosphate buffer saline (PBS).¹⁰⁹ Similarly, complete VEGF released from PCL/HA scaffolds was observed after 2 days immersion in cell culture medium.¹¹⁰ Some studies have attempted to improve the binding efficiency of biomolecules by a prior protein coating. In this regard,

Sawyer et al. pre-coated a PCL/TCP scaffold with COL I prior to bone morphogenetic protein-2 (BMP-2) adsorption.¹¹² BMPs play a crucial role on bone formation, are essential signaling molecules involved in the recovery of bone fractures, mediate cell proliferation and osteogenic differentiation.¹¹³ While BMP-2 was still completely released within the first 6h of immersion in PBS at 37 °C, increased bone healing in a rat calvarial defect was observed compared to uncoated scaffolds.¹¹² Similarly, Yan et al. performed a layer-by-layer assembly of positively charged chitosan and negatively charged deferoxamine (an iron chelator with potential to promote vascularization and bone regeneration) on the filaments of aminolysed PCL scaffolds to increase the deferoxamine loading.¹¹⁴

Alternative to an adsorption mechanism, bioactive molecules have been covalently bonded to the surface of ME-AM scaffolds. For instance, copper based click chemistry has been used to attach BMP-2 to amino acid based-PEU scaffolds.¹¹⁵ This is a highly efficient method, however it involves multiple reaction steps and cytotoxicity studies should be performed to ensure that the residual copper absorbed to the scaffold filaments does not reach cytotoxic concentrations. Alternatively, crosslinkers such as glutaraldehyde or 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDC) have been used to mediate the conjugation of biomolecules to polymeric scaffolds. For instance, acetylated collagen was bonded to PLA scaffolds via glutaraldehyde and other linkers.¹¹⁶ Similarly, hydroxypropyltrimethyl ammonium chloride chitosan (HACC) was covalently bound to the surface of PLGA/HA scaffolds via EDC/ N-hydroxysuccinimide (NHS) chemistry.¹¹⁷⁻¹¹⁸ This latter functionalization provided antibacterial properties, as well as larger volumes of new bone formation *in vivo*, compared to control scaffolds.¹¹⁷⁻¹¹⁸ In spite of their versatility, crosslinker-based covalent immobilization methods require the introduction of amine or carboxyl functional groups to the scaffolds filaments surface prior to biomolecules binding, which is normally achieved by destructive aminolysis or hydrolysis steps. Avoiding these wet-chemical steps, reactive groups can be introduced through plasma activation. In most cases, plasma activation is followed by grafting of pre-synthesized polymer brushes, a process defined as plasma grafting. Polymer brushes are robust and versatile spacers, and offer a 3D distribution of functional groups, which can significantly increase the density of tethered biomolecules.¹¹⁹ Following this plasma grafting strategy, Declercq et al. activated PCL scaffolds with argon plasma, followed by 2-amino-ethylmethacrylate (AEMA) brushes UV grafting, which served for covalent attachment of gelatin, and subsequent physisorption of fibronectin.¹²⁰ Histological analysis of the scaffolds cross section showed a higher MC3T3-E1 cells colonization in the center of these scaffold, which

significantly helped osteogenic differentiation. In another study, Tardajos et al. coupled chitosan to the surface of PCL scaffolds through a similar surface functionalization procedure involving polymer brushes. First, scaffolds were plasma activated with argon, followed by methacrylic acid N-hydroxysuccinimide ester (NHSMA) brushes UV grafting, which enabled the coupling of chitosan.¹²¹ Such coating allowed cell growth while providing an antibacterial surface against *S. aureus* and *S. epidermidis*, which are bacterial strains commonly investigated when orthopedic implants are aimed. In addition to the aforementioned 'grafting to' approaches, 'grafting from' strategies, in which polymer brushes are grown into the surface via surface initiated polymerization, have also been used on ME-AM scaffolds for a better control of the functionality, density and thickness of the polymer brush. In this regard, poly[oligo (ethylene glycol) methacrylate] (POEGMA) brushes were uniformly grafted over the surface of PCL scaffolds, allowing the subsequent conjugation of fibronectin, for cell attachment, or of BMP-2 and transforming growth factor (TGF)- β 3, which triggered osteogenic and chondrogenic differentiation of hBM-MSCs, respectively (**Figure 4A**).¹²²⁻¹²³ Although immobilization of biomolecules through polymer brushes offers advantages in terms of high density of reactive functional groups, careful optimization of this technique is needed to avoid crowding of biomolecules, which can reduce their degree of conformational freedom and their resultant bioactivity.¹¹⁹

Polydopamine (PDA), which contains hydroxyl and amine functional groups, has also been investigated as primer coating for subsequent covalent conjugation of proteins onto ME-AM scaffolds. The dopamine molecule possesses a similar chemical structure to that of combined lysine and 3,4-dihydroxyphenyl-L-alanine (DOPA) aminoacids, which confer the adhesive properties to the mussel adhesive proteins (MAPs) secreted by marine mussels.¹²⁴ Inspired by mussel adhesiveness, a PDA coating can be deposited onto any surface via the oxidative self-polymerization of dopamine at a slightly basic pH. Following PDA coating, Teixeira et al. immobilized COL I on PLA scaffolds via EDC-NHS chemistry.¹²⁵ Here, the combined coating enhanced hBM-MSCs adhesion and ALP activity compared to untreated scaffolds. Besides, various reports have immobilized BMPs on PDA coated scaffolds via Michael addition reaction.¹²⁶⁻¹²⁹ For instance, Lee et al. grafted recombinant BMP-2 to PCL-PDA scaffolds, which was sustain released over time and upregulated osteogenic gene expression on preosteoblasts (MC3T3-E1) cultured on the scaffolds.¹²⁶ Similarly, bone forming peptide-1 (BFP-1, a short peptide derived from BMP-7) and SSVPT (an oligopeptide derived from BMP-2) were immobilized on PCL and PLA scaffolds, respectively, leading to enhanced cell adhesion and osteogenesis *in vitro* and increased

bone growth in critical size calvarial defects *in vivo* (**Figure 4B**).¹²⁸⁻¹²⁹ Due to the high adsorption of serum proteins to PDA, which maintain an active conformation exposing integrin binding domains,¹³⁰⁻¹³¹ a simple PDA coating onto ME-AM scaffolds has also been used to enhance cell adhesion and trigger differentiation. For instance, Kao et al. covered the filaments of a PLA scaffold with PDA.¹³² The PDA hydrophilic coating enhanced hAD-MSCs adhesion and proliferation, and supported osteogenesis by inducing the secretion of larger amounts of COL I and OCN, compared to bare PLA scaffolds. Interestingly, hAD-MSCs cultured on coated scaffolds also showed the upregulation of angiogenic genes. Due to its material-independent coating ability, simple deposition process, and high biocompatibility and stability *in vitro*, PDA constitutes a powerful tool for surface modification of ME-AM scaffolds. Moreover, preliminary *in vivo* studies have shown that PDA coatings can greatly reduce the inflammatory response of PLLA surfaces, evaluated during the first 14 days after implantation.¹³³ Nevertheless, its long term *in vivo* stability, degradation mechanism and biocompatibility should be still evaluated.

An alternative method to improve the bioactivity of scaffold consists of the coating with cell-laid ECM, containing several bioactive factors that can stimulate osteogenic differentiation *in vitro* and *in vivo*.¹³⁴ Compared to single protein coatings, which only provide fragmental components of the ECM that present specific functional receptors for cell attachment, such an approach better mimics the complexity of ECM. In this regard, Wu et al. cultured MG63 on PCL scaffolds for 5 days before a decellularization process with Triton, NH₄OH and DNase.¹³⁵ Human Wharton's Jelly-derived MSCs (hWJ-MSCs) seeded on these ECM-coated scaffolds exhibited enhanced cell adhesion, proliferation and the upregulation of ALP, COL I, BSP and OCN genes, as well as increased calcium deposition, compared to uncoated scaffolds. Bone formation on a rat calvarial defect filled with these scaffolds was also higher than on uncoated scaffolds.¹³⁵ In another study, the tissue formed by human nasal inferior turbinate tissue-derived MSCs (hNIT-MSCs) on PCL/PLGA/TCP scaffolds was decellularized through freeze-thaw cycles, which minimally affected the ECM bioactive components, compared to chemical methods.¹³⁶ Subsequently seeded hNIT-MSCs also showed osteogenic genes upregulation, and coated scaffolds induced greater ectopic bone formation *in vivo* than bare scaffolds (**Figure 4C**).¹³⁶ Similar to ECM, platelet rich plasma (PRP) can stimulate osteogenic differentiation and has been used to coat ME-AM scaffolds.¹³⁷ PRP is a source of growth factors that play an essential role in tissue healing, including platelet derived growth factor (PDGF), vascular endothelial growth factor (VEGF), TGF- β 1, basic fibroblast growth factor (bFGF), epidermal growth

factor (EGF), and insulin-like growth factor.¹³⁸⁻¹³⁹ PCL scaffolds coated with freeze dried PRP, which had previously been proven to preserve growth factors at high levels,¹⁴⁰ have shown to promote ALP activity and osteogenic gene upregulation on dental pulp stem cells, as well as higher levels of bone formation in a rat calvarial defect, compared to uncoated scaffolds.¹³⁷ Overall, coatings comprising a cocktail of biomolecules can exert a more profound effect on the phenotypic behavior of cells, compared to single proteins, peptides or growth factors. Moreover, they avoid the lack of reproducibility, availability, and tunability of bioactive decellularized ECM scaffolds while embracing their advantages. Nevertheless, further investigation on variables such as cell source, culture duration and media type, is necessary to optimize these unique and complex biologically engineered systems.

Figure 4 - Scaffolds surface modification via functional coatings. (A) (i) Methylene blue staining of hBM-MSCs adhered to PCL functionalized with POEGMA brushes. (ii) Homogeneous cell attachment on PCL-POEGMA scaffold grafted with fibronectin (FN). (iii) Vertical and (iv) radial gradient in coupled FN demonstrates gradients in cell attachment. The yellow arrows indicate the direction of decreasing FN concentration. Adapted with permission from Ref.¹²² (B) (i) SEM images of the surface morphology of PCL scaffolds coated with PDA and grafted with BFP1. Insets correspond to fluorescence images of BFP1 tethered on the scaffolds using FITC-labelled BFP1. (ii) Radiographic evaluation of bone formation around the scaffolds at 8 weeks post-implantation in a calvarial critical size defect in rabbits. Adapted with permission from Ref.¹²⁹ (C) (i) SEM images of uncoated and d-ECM-coated PCL/PLGA/TCP. (ii) Histological sections stained with H&E of critical size rat calvarial defects treated with the aforementioned scaffolds. Adapted with permission from Ref.¹³⁶ (D) (i) SEM images of uncoated and CaP-coated PLA scaffolds. (ii) Confocal

microscopy images showing cell attachment to CaP-coated scaffolds after 7 days of culture (F-actin, green, nuclei, red). (iii) ARS staining on scaffold seeded with hBM-MSCs and cultured for 14 days in mineralization media. Insets depict stained scaffolds without cells but cultured under the same media conditions. Adapted with permission from Ref. ¹⁴¹

CaP coatings

The presence of CaP deposits on the surface of ME-AM scaffolds can potentially stimulate cells' attachment and osteogenic differentiation through several cues, such as the release of calcium ions, the adsorption of serum proteins, or the nanoscale surface roughness. However, the direct growth of apatite layers on a polymeric surface is a challenging task due to the lack of nucleation sites. To overcome this, Jo et al. exploited the ability of PDA to nucleate apatite to create a bioinspired coating on PCL scaffolds, which allowed increased MC3T3-E1 preosteoblast infiltration and adherence within the scaffolds core. ¹⁴² Here, the catecholamine moieties in PDA acted as nucleation sites of calcium and phosphate from simulated body fluid (SBF, 1.5x), in which the scaffolds were incubated for 14 days. ¹⁴³ Following a different approach, Murab et al. activated the surface of PCL scaffolds with oxygen plasma, leading to the introduction of hydroxyl and carboxyl groups, which nucleated biomimetic HA crystals upon immersion in cell culture medium. ¹⁴⁴ Alternatively, Holmes et al. conjugated nano HA (nHA) particles to the surface of aminolysed PLA scaffolds through a glutaraldehyde crosslinker. ¹⁴⁵ This coating promoted osteogenic differentiation, observed in the form of COL I and calcium deposition, and vascularization upon co-culture of hBM-MSCs and human umbilical vein endothelial cells (HUVECs). Following a milder strategy, Jaidev et al. modified the surface of NaOH pre-treated PLA scaffolds by grafting polyethyleneimine and subsequently immobilizing citric acid. ¹⁴¹ Citric acid, which has been identified as a critical role player in precipitating the apatite phase during bone formation, ¹⁴⁶⁻¹⁴⁷ mediated CaP precipitation from 1.5x SBF onto the scaffold after 7 days incubation. HBM-MSCs cultured on these scaffolds showed BMP-2 gene upregulation and increased calcium deposition, compared to cells on bare PLA scaffolds (**Figure 4D**). ¹⁴¹ In an attempt to incorporate larger amounts of ceramics, Moroni and coworkers press fitted HA pillars into the pores of PEOT/PBT 3DF scaffolds. ¹⁴⁸⁻¹⁴⁹ Such approach enabled larger amounts of calcium and phosphate ions release near cells cultured on the scaffolds, which influence stem cell osteogenic differentiation leading to upregulation of OPN gene expression, compared to bare and PEOT/PBT-HA composite scaffolds.

In vivo studies assessing bone formation on CaP-coated ME-AM scaffolds have not always reported the beneficial effect of the CaP coating. For instance, Kim et al., who dip-coated PLGA/TCP scaffolds in a HA slurry, did not observe differences in new bone formation area among PLGA/TCP coated and uncoated scaffolds upon implantation in a rabbit femoral defect for 12 weeks.¹⁵⁰ Similarly, Chim et al. observed less vasculogenesis and mineralization on precalcified PCL scaffolds implanted subcutaneously in mice for 14 weeks, compared to uncoated PCL scaffolds.¹⁵¹ Here, the coating was prepared by priming the PCL scaffolds with a silicate gel prior to 4 weeks immersion in SBF (1 week in 1x SBF, followed by 3 weeks in 1.5x SBF). In both of the aforementioned studies it was hypothesized that flaking of the apatite coating might have impaired tissue ingrowth and osteogenesis, and that thinner layers might have yield more promising results. In accordance, Maia-Pinto, et al. reported increased bone formation on PLA scaffolds coated with a thinner CaP layer, only after 1 month of implantation in a rat calvarial defect, compared to bare PLA scaffolds.¹⁵² Here, the thin apatite layer was achieved through a shorter immersion time in SBF, 2 weeks (1 week in 1x SBF, followed by 1 week in 1.5x SBF), compared to the 4 weeks reported in the study of Chim et al. Taking into consideration that the aforementioned results derive from different *in vivo* models, we can still suggest that the optimization of the apatite layer, in terms of thickness, is necessary in order to obtain a functional coating for the final *in vivo* application. In this regard, ensuring a good interfacial bonding by selecting an appropriate scaffold priming methodology seems also crucial for the success of the coating.

Hybrid scaffolds

Fibrous networks

While the large pore size of ME-AM scaffolds (300-1000 μm) is advantageous for media permeation *in vitro*, as well as for tissue ingrowth and vascularization *in vivo*, such dimensions can turn out to be a limitation for cell seeding and scaffold coverage in an *in vitro* setting. Due to these large FDs, there is limited bridging between cells on different filaments, and pore closure happens slowly by cells forming concentric circles around the pore.²⁹ This can significantly slow the process of tissue maturation, for which cell-cell contact is essential.¹⁵³ Inspired by the fibrillary structure of ECM, some studies have embedded ME-AM scaffolds in a hydrogel solution and subsequent freeze-drying for the

creation of a fibrous network within the pores of the scaffolds. This strategy can create an environment to enhance cell retention during the seeding process, as well as cell-cell interactions and mobility on a mechanically stable and highly porous printed structure. For instance, Jensen et al. dip-coated a PCL scaffold with a mixture of hyaluronic acid and TCP, followed by freeze-drying to create a fibrous coating containing TCP particles around the scaffolds filaments.¹⁵⁴⁻¹⁵⁵ Human dental pulp stem cells (hDPSCs) and MSC derived from human induced-pluripotent stem cells (hiPS-MSCs) proliferated at a higher rate on these scaffolds, deposited higher amounts of calcium and showed an upregulation of osteogenic genes, compared to uncoated scaffolds. Similarly, Chen et al. freeze-dried PCL scaffolds immersed in hyaluronic acid to create an interconnected fibrous network. HMSCs were seeded in a methylated collagen and terpolymer hydroxylethyl methacrylate- methyl methacrylate- methacrylic acid (HEMA-MMA-MAA) solution.¹⁵⁶ This complex mimicked the constitution of the ECM, helped to achieve a more homogeneous cell distribution across the scaffold upon seeding and led to osteogenic genes upregulation, owed to the exposed integrin domains of the coated methylated collagen promoting focal adhesion assembly and activating osteogenesis pathways.

Inspired by the submicron key features in the ECM, some studies have reported the combination of ME-AM and electrospinning (ES), in order to introduce nanofibers to the microporous scaffolds. Some studies have reported the preparation of such hybrid scaffolds by alternating ES layers within ME layers,¹⁵⁷⁻¹⁵⁹ while others have opted for the surface addition of the ES fibers once the 3D scaffold was completely printed.¹⁶⁰⁻¹⁶¹ Interesting observations have been extracted from some of these studies. For instance, Park et al., who presented scaffolds consisting of PCL/collagen dense ES fibers intercalated within the layers of a PCL ME-AM scaffold, observed the proportional relation between number of intercalated ES layers and cell seeding efficiency and proliferation rate.¹⁵⁷ Moreover, the reports of Mota et al. and Vyas et al. described the alignment of nanofibers perpendicular to the direction of the printed filaments, which acted as a parallel plate collector, when depositing low fiber density electrospun layers (**Figure 5A**).¹⁵⁹⁻¹⁶⁰ These crossing nanofibers acted as bridges for cells to easily reach parallel printed filaments, otherwise not possible on conventional ME-AM scaffolds. Interestingly, Lara-Padilla et al. implemented a cooling system to allow the layer-by-layer intercalation of PCL nanofibers between PLA printed layers.¹⁵⁸ This was to avoid the PCL fibers to partially melt and lose their integrity upon ME of subsequent PLA layers, due to the higher melting point of PLA. However, the resulting hybrid scaffolds presented lower mechanical

properties, due to fast cooling reducing the bonding between printed layers. Going one step further, the combination of the ES and ME-AM technologies has given rise to gradient scaffolds for bone to ligament or bone to cartilage interfaces.¹⁶²⁻¹⁶³ In the latter case, the electrospun layer served as the tidemark of the osteochondral tissue and prevented blood vessel invasion towards the cartilage ME-AM region. In order to have a better control on the porosity of the secondary fibrillary network, Yang et al. used electrohydrodynamic direct writing (EHD-DW), instead of ES, in combination with ME-AM.¹⁶⁴ Here, layers of ME PCL filaments were alternated with layers of EHD-DW PCL threads (~100 μm , composed of microfibers of ~3 μm) precisely deposited with an offset with respect to the ME filaments. This created a secondary interwoven network with smaller dimensions and porosity, which allowed higher MG63 cell retention, ALP activity and calcium deposition than ME-AM or hybrid ES-ME-AM scaffolds (**Figure 5B**).

Overall, hierarchical scaffolds comprising ME filaments and a secondary fibrous network mimicking the micro/nanoscale details observed in tissues, such as bone, are desired to increase cell retention and promote tissue formation within the scaffold. On-going research on including such biomimetic features to ME-AM scaffolds has contributed to the development of combinatorial technologies, such as ES or EHD-DW, with ME-AM, which enable the automation and control over the whole scaffold fabrication process. While this requires currently complex experimental setups, more integrative technologies are expected to be designed in the future.

Hydrogels

Natural or synthetic hydrogels have also been used in combination with ME-AM scaffolds for cell encapsulation within the scaffold's pores. Such hybrid systems allow the retention of large cell concentrations compared to any other seeding methods, as the whole seeding volume containing the cells is maintained within the scaffolds pores after hydrogel crosslinking. For instance, sheep hBM-MSCs have been resuspended into a PRP solution and loaded on PCL/TCP scaffolds. The PRP solution formed a gel after crosslinking with thrombin, which enabled to retain the cells within the scaffold prior to their implantation on a sheep's tibial defect.¹⁶⁸⁻¹⁶⁹ With the same mechanism of action, fibrinogen has been used for seeding hBM-MSCs, which were embedded within the pores of a PCL scaffold after the addition of thrombin and fibrinogen polymerization into a fibrin gel.¹⁷⁰ After 2-3 days of culture, and in parallel to hydrogel degradation, cells started migrating towards the

PCL filaments. While in these systems the hydrogel was used as a degradable cell carrier, and the ME-AM scaffold was the main hybrid construct component for cells to grow on, other approaches have considered the ME-AM part solely as a structural support for the cell-laden hydrogel. For instance, Wang et al. took advantage of the round morphology of rabbit hBM-MSCs within PLGA-PEG-PLGA hydrogel carriers to stimulate chondrogenic differentiation, which was demonstrated to be better supported on the hybrid scaffolds, compared to bare PCL scaffolds.¹⁷¹ Moreover, Heo et al. used a PLA ME-AM scaffold to reinforce the surrounding hAD-MSCs-laden GelMA hydrogel, which in addition contained RGD conjugated gold nanoparticles for osteogenic gene upregulation and enhanced calcium deposition.¹⁷² Furthermore, hydrogels contained within ME-AM scaffolds have been used for the pre-vascularization of the constructs. For example, methacrylated hyaluronic acid/gelatin hydrogels carrying hAD-MSCs and HUVEC promoted the formation of capillary like networks within PCL/HA ME-AM scaffolds both *in vitro* and *in vivo* (**Figure 5C**).¹⁶⁵

Besides the incorporation of the cell-laden hydrogel component within ME-AM scaffolds by surface injection, the combination of bioprinting and ME-AM has been used to intercalate cell-laden hydrogel filaments within thermoplastic polymer filaments in a layer-by-layer manner. This hybrid multi-printhead deposition process enables i) the automation of the tissue construct fabrication, as it does not require post-scaffold fabrication cell seeding, ii) the precise control over the deposition of the hydrogel bioink at specific locations within the construct, and iii) the possibility of multimaterial and multicellular bioprinting within the same hybrid construct.^{166, 173-175} Yet, specific hydrogel rheological properties are required for bioprinting, and special attention should be taken to cell viability after shear during the extrusion process. Moreover, since the bioprinted filaments get in direct contact with the hot ME filament shortly after extrusion, low melting temperature thermoplastics, such as PCL, are normally used in these cases, to avoid heating of the cell-laden hydrogel by conduction. Working around these constraints, scaffolds with optimized microenvironment for the growth of 3D complex tissues have been fabricated. For instance, bioinks consisting of decellularized ECM of different tissues and hAD-MSCs have been bioprinted in combination with PCL ME-AM scaffolds to promote tissue specific differentiation.¹⁷⁶ Alternatively, Mekhileri et al. bioprinted human articular and nasal chondrocytes microtissues within the pores of PEOT/PBT scaffolds after the completion of the ME-AM scaffolds, in contrast to the aforementioned layer-by-layer process. The use of

microtissues instead of single cells enabled building complex tissues, such as the osteochondral interface, of clinically relevant sizes (**Figure 5D**).¹⁶⁶

Figure 5 Combination of ME-AM scaffolds with fibrous and hydrogel networks. (A) (i) SEM images of bare PCL scaffolds, and scaffolds fabricated combining ME-AM of PCL and ES of PLGA nanofibers. (ii) Fluorescent microscopy images of MC3T3 (F-actin, green) cultured for 7 days on the aforementioned scaffolds. Adapted with permission from Ref.¹⁶⁰ (B) (i) Optical and (ii) SEM images of bare PCL scaffolds, and hybrid scaffolds fabricated combining ME-AM of PCL and EHD-DJ of collagen struts. (iii) Confocal fluorescent microscopy images of MG63 (F-actin, red; nuclei, blue) cultured for 7 days on the aforementioned scaffolds. Adapted with permission from Ref.¹⁶⁴ (C) Immunohistochemistry fluorescent images of *in vivo* vascularization within PCL/HA scaffolds seeded with cell laden GelMA hydrogels containing either hAD-MSC, or hAD-MSC combined with HUVEC (CD31, red; α -smooth muscle actin, green; nuclei, blue). Scaffolds were implanted for 4 weeks subcutaneously. Asterisks indicate the PCL/HA scaffold filaments. Arrows indicate the stained cells/microvessels positive for human CD31. Adapted with permission from Ref.¹⁶⁵ (D) (i) Biphasic hemispherical construct with stained GelMA hydrogel micro-spheres representing the chondrogenic (red) and osteogenic (blue) phase of an osteochondral construct. (ii) This construct was fabricated by applying a bottom-up automated tissue bioassembly strategy, consisting on the automated insertion of micro tissues within the ME-AM scaffold pores. (iii) Fluorescence microscopy images of injected human articular cartilage micro tissues stained with Calcein (live cells, green) and Propidium Iodide (dead cells, red). Adapted with permission from Ref.¹⁶⁶ (E) (i) Representative X-rays and (ii) μ CT reconstructions of bone formation after hybrid scaffolds implantation on ovine tibial defects. Scaffolds consisted on a PCL ME-AM structure, loaded with a

platelet rich plasma hydrogel containing microparticles for the delivery of BMP-2 or a combination of BMP-2, VEGF and PDGF. (iii) Representative images from defect mid sections showing immunohistochemical staining for the endothelial marker Von Willebrand Factor. Adapted with permission from Ref. ¹⁶⁷

Hydrogels have also been used as cell-free delivery systems of growth factors within ME-AM scaffolds. They are an alternative strategy to surface adsorption, covalent immobilization or melt blending for the incorporation of bioactive molecules, which offer tunable and controlled release rates, as a function of hydrogel mesh size and degradation rate. PCL/PLGA and PCL/TCP scaffolds filled with a collagen hydrogel containing rhBMP-2 enabled the controlled release of the growth factor, which improved and accelerated bone formation *in vivo*. ^{168, 177} Similarly, decellularized ECM hydrogel containing human recombinant BMP-2 was dispensed within the filaments of a PCL/TCP ME-AM scaffold enabling a sustained release, which resulted in a larger volume of new bone formation, compared to PCL/TCP and PCL/TCP/decellularized ECM scaffolds. ¹⁷⁵ In another study, the central region of a PCL scaffold was loaded with a PRP hydrogel containing degradable PLGA microparticles, which carried VEGF, PDGF and BMP-2, or BMP-2 only. ¹⁶⁷ The polymeric microparticles were used to achieve a more sustained release, while the growth factors combination was chosen for obtaining a stable vasculature and induce bone formation in a sheep tibial segmental defect (**Figure 5E**).

Hydrogel laden ME-AM scaffolds bring together the ECM-like properties of hydrogels and the high structural support of ME-AM constructs, and the addition of cell, growth factors, drugs, or their combination to the equation adds up for extra functionality. The on-going research focused on the development of cell-laden hydrogels for bioprinting will allow in the future for further optimization of such hybrid systems towards advanced scaffolds for bone regeneration.

Composites

CaP based materials

In an attempt to mimic the organic-inorganic composition of bone ECM, CaP-polymer composites have been one of the most used combination of materials for the design of BTE scaffolds. CaP fillers can enhance the polymeric scaffolds mechanical properties, as well as offer osteoinductive and osteoconductive properties, which are dependent on their

specific chemical phase, structural properties (crystallinity, surface roughness, and porosity), mechanical properties and associated degradation behavior.¹⁷⁸⁻¹⁸⁰ Among all the CaP phases, HA and TCP, have been the most commonly used fillers for ME-AM scaffolds. Considering the average mineral content of bone tissue, which lies within the range of 50 to 74 wt%, depending on the body location,¹⁸¹⁻¹⁸² tremendous efforts have been conveyed to increase the CaP content within polymeric scaffolds to optimize their mechanical properties while retaining their printability. For instance, Esposito-Corcione et al. reported the fabrication of PLA/HA 50 wt% scaffolds, which presented a 2-fold increase in compression modulus, compared to bare PLA scaffolds.¹⁸³ Similarly, Jiang et al. and Ding et al. reported the fabrication of PCL/HA scaffolds with up to 40 wt% mineral content.¹⁸⁴⁻¹⁸⁶ Such scaffolds also presented a 2-fold increase in compression modulus and compressive strength compared to bare PCL scaffolds. Moreover, surface roughness and hydrophilicity were significantly increased, which enhanced cell attachment and proliferation. Interestingly, composites with higher HA concentration (60 wt%) were not printable in a continuous manner, due to particle aggregation leading to nozzle clogging and filament discontinuity.¹⁸⁶ In another report, Yu et al. fabricated PEU/HA scaffolds with up to 40 wt% mineral content, which were shown to support osteogenic differentiation in mineralization media cultures.¹⁸⁷ Here, MC3T3-E1 cells seeded on the highly loaded composite scaffolds presented higher proliferation rates, as well as increased ALP activity and BSP and OCN gene upregulation, compared to cells on scaffolds with lower HA content and bare PEU scaffolds. Enhanced osteogenesis was attributed to HA increasing protein adsorption and activating focal adhesion kinase (FAK) and mitogen-activated protein kinase/ extracellular signal-regulated kinase (MAPK/ERK) signaling pathways.¹⁸⁷ In order to further increase the osteogenic differentiation potential of polymer-HA composite scaffolds, Cho et al. eroded the surface of PCL/HA 40 wt% scaffolds with a NaOH surface treatment.¹⁸⁸⁻¹⁸⁹ This was done to increase the exposure of HA particles on the filaments' surface, and thus the possibility of Ca and P ions exchange with the medium, since Ca and P (re)precipitation from CaP materials is known to affect osteogenic pathways.¹⁹⁰⁻¹⁹² In this regard, NaOH erosion led to higher cell proliferation rates, calcium deposition and apatite layer formation *in vitro*, compared to other scaffold types.¹⁸⁸ *In vivo*, HA based scaffolds allowed for significantly higher bone formation compared to bare PCL scaffolds, and eroded PCL/HA scaffolds showed the best bone regeneration ability (**Figure 6A**).¹⁸⁹

While osteogenic differentiation has been supported *in vitro* on highly loaded HA scaffolds, enhanced osteogenesis has not been observed on scaffolds with low HA content, such as PEOT/PBT/HA or PCL/PLGA/HA with 15 % and 10% inorganic content, respectively.¹⁹³⁻
¹⁹⁴ However, *in vivo* experiments have revealed the osteoinductive/ osteoconductive properties of such low HA content scaffolds. For instance, new bone formation on the aforementioned PCL/PLGA/HA 10 wt% HA scaffolds was 100% higher than in bare polymeric scaffolds in a rat calvarial defect.¹⁹⁴ Besides, mechanical properties are still enhanced by low HA loadings, which makes these scaffolds still desirable for BTE applications.¹⁹³⁻¹⁹⁴ PCL/HA scaffolds with low HA content (10 wt% or 15 wt% HA) have also shown to promote bone formation *in vivo*.¹⁹⁵⁻¹⁹⁶

A couple of studies have assessed the effect of different CaP materials on the performance of composite ME-AM scaffolds. For instance, Nyberg et al. compared the *in vitro* osteoinductivity of PCL/HA, PCL/TCP, PCL/Bio-Oss® (bovine bone mineral matrix) and PCL/DBM (decellularized bone matrix) scaffolds with 30 wt% of inorganic content, post-treated with NaOH.¹⁹⁷ Overall, greater osteoinductivity was observed on hAD-MSCs cultured on PCL/Bio-Oss and PCL/DBM compared to PCL/HA and PCL/TCP scaffolds, in terms of calcium deposition and COL I and osteonectin gene upregulation in basic and mineralization media. Nevertheless, Runt-related transcription factor 2 (RUNX2) and OPN were also upregulated on PCL/HA scaffolds in basic media, compared to other scaffolds types. Here, TCP showed lower bioactivity compared to other CaP fillers. Similarly, Huang et al. also showed that despite PCL/TCP 20 wt% scaffolds displayed better compression modulus before implantation than PCL/HA 20 wt% scaffolds, they presented lower bioactivity, in terms of cell attachment and proliferation.¹⁹⁸ In spite of this, a large amount of reports have explored the use of TCP as filler for ME-AM scaffolds.⁴⁶ Several have attempted the addition of TCP up to 60 wt% to PCL composite scaffolds, which has shown to lead to an increase in compression modulus of up to 50%, although these high wt% diminished significantly the scaffolds compressive yield strength.¹⁹⁹⁻²⁰¹ For this reason, composite scaffolds with up to 20 wt% TCP have been mostly investigated, which have proven their bone regenerative capacity *in vivo*.^{46, 168, 202} In addition, promising *in vivo* results have led to clinical trials investigations, where the applicability of PCL/TCP scaffolds has been tested for craniomaxillofacial bone regeneration (NCT01119144, NCT03735199). The fast degradation of TCP compared to HA can confer the scaffold with more optimum ion exchange dynamics to stimulate bone formation through calcium activated pathways.^{190-191, 203} In order to accelerate the degradation dynamics of the

scaffold and of TCP, and to enhance the replacement of the scaffold material by newly formed bone, NaOH treatment²⁰⁴⁻²⁰⁷ and PLGA²⁰⁸⁻²¹⁰ have been further included into TCP based scaffolds. Yet, balancing the speed of degradation of TCP containing scaffolds is important, in order to keep the released calcium and phosphate concentration within the cell proliferation and differentiation stimulating range, without reaching apoptotic values.

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In order to further mimic the *in vivo* tissue structure and stimulate bone formation, polymer-inorganic composite scaffolds with gradients in HA or TCP concentration have been fabricated. For instance, Bittner et al. and Smith et al. have reported PCL/HA and PCL/TCP composite scaffolds, respectively, with vertical discrete gradients in inorganic content (from 30 or 20 to 0 wt% HA or TCP) to mimic the bone to cartilage interface (**Figure 6B**).²¹²⁻²¹³ In contrast to these multi-printhead and discrete gradient fabrication approaches, Sinha et al. have recently reported the fabrication of continuously graded HA-based ME-AM scaffolds using a single printhead approach (**Figure 6C**).⁸⁸ This printhead allowed the *in situ* mixing of two materials (PEOT/PBT only, and PEOT/PBT/HA 45 wt%) to obtain scaffolds with continuous HA composition gradients (among other inorganic fillers), which better mimic *in vivo* tissue interfaces and possess better mechanical properties than discrete gradient scaffolds.²¹⁴ Other studies have focused on the production of scaffolds with radial gradients in inorganic content and porosity have also been reported, to resemble the mechanical properties and bulk mineral content of the cortical to cancellous bone interface.²¹⁵⁻²¹⁶

CaP-polymer composite ME-AM scaffolds have extensively proven their mechanical and biological functionality for BTE, while benefiting from a single-step fabrication approach, compared to post-processing coating strategies. In spite of extensive material characterization and *in vitro* research, further *in vivo* investigations are still required to unravel the optimum scaffold composition, in terms of polymer type, CaP type and content, and degree of CaP exposure to the scaffold filament surface, which balance printability, mechanical properties and bone regeneration ability.

Silicates

Calcium silicate (CS) based materials, composed of CaO, SiO₂, and MgO or Al₂O₃, have been used as fillers in ME-AM scaffolds, due to their bioactive properties and charged surfaces being highly compatible with polymeric matrices.²¹⁷⁻²¹⁸ An apatite layer is

generally formed on CS when in contact with body fluid, which acts stronger interface and bonding with surrounding tissues upon implantation. Besides the known osteogenic capacity of Ca, the released Si ions also have the ability of upregulating MAPK/ERK and p38 signaling pathways, which are known to be involved in osteogenesis, and stimulating angiogenesis as well.²¹⁹ Lin et al. fabricated PCL/Mg-CS scaffolds, which released Mg and Si, and reprecipitated Ca and P when immersed in SBF, forming an apatite layer (**Figure 6D**).²²⁰ The highest hWJ-MSCs attachment, proliferation and osteogenic genes upregulation was observed for scaffolds with 50 wt% Mg-CS, which released larger amounts of the aforementioned ions. These scaffolds also showed the highest degradation rate, due to lower polymer content. Zhang et al. also observed the good performance of PCL/gliadin scaffolds containing 30 wt% Mg-CS.²²¹ Here, both Mg-CS and the natural polymer gliadin were added to the scaffold to enhance its bioactivity and degradability. In a rabbit femoral defect, scaffolds with 30 wt% Mg-CS allowed for the fastest degradation rate and largest new bone volume formation, compared to lower Mg-CS percentages and bare PCL/gliadin scaffolds.

A very widely used type of CS is bioactive glass (BG), which in addition to CaO and SiO₂, is composed of Na₂O and P₂O₃. *In vitro*, BG has been shown to outperform other CaP, like HA, in terms of biological response and osteogenic commitment when incorporated within a PLA scaffold at the same concentration, potentially due to enhanced protein adsorption.²²² Yet, *in vivo* studies have not reported differences among PCL/BG and PCL/CaP scaffolds within 16 weeks implantation time, in terms of bone formation.²²³ Strontium-substituted BG (SrBG) has also been investigated as filler for ME-AM scaffolds, as Sr is an important trace element in bone and can stimulate bone formation and reduce bone resorption.²²⁴ In this regard, Poh et al. prepared PCL/SrBG 10 wt%, which enhanced the formation of calcium deposits, compared to PCL/BG 10wt% scaffolds.²²⁵

CS based materials appear to be valid substitutes of CaP as osteoinductive materials for bone regeneration applications, due to their richer bioactive ion composition. Although these established advantages, insufficient efforts have been dedicated so far to investigate their use as fillers within ME-AM scaffolds. A reason for that could rely on their fast degradation rates compromising the scaffold mechanical properties already at early stages of the regeneration process.²²⁶ Nevertheless, further research is required to better understand the performance of CS in comparison with CaP fillers.

Metallic particles

Metals, such as titanium based alloys or cobalt chromium alloys, have been extensively used in the orthopedics field as permanent implants due to their good biocompatibility and high mechanical strength and wear properties.²²⁷⁻²²⁸ Others, such as iron- or magnesium (Mg)-based metals, are preferred for bone regeneration applications, since they are resorbable and can be degraded to be replaced by newly formed bone. In particular, Mg-based metals have been used as fillers for ME-AM scaffolds. Apart from improving the mechanical properties of composites, Mg ions are essential to bone metabolism, have been shown to be associated with the formation of biological apatite, and can stimulate osteogenic activity.²²⁹⁻²³¹ Since bare Mg corrodes in aqueous environment leading to the formation of gas cavities,²³² more stable compounds such as magnesium hydroxide (MgOH), magnesium oxide (MgO) or magnesium fluoride (MgF) have been used as fillers within ME-AM scaffolds. For instance, Abdal-hay et al. prepared PCL/MgOH scaffolds with up to 20 wt% of MgOH.²³³ Despite not enhancing the scaffolds' mechanical properties, potentially due to poor nanoparticle dispersion at high concentrations, Mg ions were released from the scaffolds, which led to increased cell proliferation and ALP activity. In another study, Shen et al. prepared PCL-PEG-PCL/MgO composite scaffolds.²³⁴ Here, silanization of the MgO particles allowed the addition of up to 4 wt% MgO without the formation of agglomerates, which resulted in a 2-fold increase in compression modulus compared to bare PCL scaffolds. Yet, the most optimum magnesium release rate was achieved with 1 wt% MgO, which favored cell adhesion, ALP activity and calcium deposition, as well as bone formation *in vivo*, compared to other MgO concentrations (**Figure 6E**).²³⁴ Hansske et al. also pre-coated MgF nanoparticles, in this case with a poly (ethylene glycol) (PEG)-based biomimetic coupling agent to improve their stability and compatibility with the polymer matrix, prior to loading within a PCL scaffold.²³⁵ In a subsequent study, these compatibilized MgF nanoparticles were used as fillers in a ternary composite scaffold consisting of PCL, nHA and MgF.²³⁶ The sole addition of 1 wt% MgF to the PCL/HA scaffolds enhanced the attachment and osteogenic differentiation of human fetal-derived MSCs (hF-MSCs), in terms of ALP activity, osteonectin gene expression and calcium deposition. In addition to the bioactivity of Mg, F is also known to play a key role in the development of mineralized tissues.²³⁷

Silver has also been used as metallic filler within ME-AM scaffolds. Besides enhancing mechanical properties, silver has shown to provide antimicrobial properties to the

scaffolds. For instance, Radhakrishnan et al. prepared PCL/silver 3 wt% scaffolds, which showed antibacterial activity against *E. coli* while maintaining osteoblast cell viability.²³⁸

Overall, metallic particles are an attractive filler option for ME-AM composite scaffolds, especially due to their ability to significantly enhance the scaffold's mechanical properties at very low concentrations. Nevertheless, compatibility and dispersion within polymers are the major issue associated to these materials, and the main reason why they have not been widely explored so far.

Figure 6 - Composite scaffolds. (A) (i) SEM images of the surface of PCL/HA 40 wt% composite scaffolds before and after etching with NaOH. Adapted with permission from Ref. ¹⁸⁸ (ii) μ-CT images of critical size rat calvarial defects depicting new bone formation after 8 weeks implantation of the aforementioned scaffolds. Increased bone formation is observed in the composite scaffolds, compared to the PCL and PCL etched scaffolds controls. (iii) Representative Masson's trichrome stain images of the calvarial defects 8 weeks post-implantation, depicting mineralized areas (red), which were more abundant in the HA containing scaffolds. Adapted with permission from Ref. ¹⁸⁹ (B) μCT images of PCL scaffolds with a discrete vertical gradient in HA concentration. Adapted with permission from Ref. ²¹² (C) μCT images of PEOT/PBT scaffolds with a continuous vertical gradient in HA concentration. Adapted with permission from Ref. ⁸⁸ (D) (i) SEM images of the

surface of PCL and PCL/CaS 50 wt% scaffolds, after immersion in SBF for 7 days, depicting the ability of PCL/CaS of nucleating HA. (ii) Fluorescent microscopy images of hWJ-MSCs (F-actin, green) cultured on the aforementioned scaffolds for 7 days, showing enhanced cell coverage on the composite scaffolds. (iii) ARS staining of calcium nodules deposited on the PCL and PCL/CaS scaffolds cultured with hWJ-MSC for 3 weeks. Adapted with permission from Ref. ²²⁰ (E) (i) Giemsa staining on histological sections of lateral epicondyle defects in rats 8 weeks post-implantation of PCL/PEG and PCL/PEG/MgO 1 wt% composite scaffolds (red, mineralized tissue). (ii) Fluorochrome-labeling analysis of new bone formation and mineralization 8 weeks post-implantation of the aforementioned scaffolds (calcein green, between 4 and 6 weeks; xylenol orange, red, between 6 and 8 weeks). 'OB' and 'NB' refer the location of original bone and newly formed bone, respectively. Adapted with permission from Ref. ²³⁴ (F) H&E stained sections of rat calvarial defects with implanted PCL and PCL/G (graphene) 0.78 wt% scaffolds, with or without electrical stimulation. Adapted with permission from Ref. ²³⁹ (G) (i) SEM images of the surface of PCL/PLGA and tobramycin-loaded PCL/PLGA scaffolds with various loading concentrations (2 wt% and 4 wt%). (ii) Results of disk diffusion test on day 1 (left) and day 7 (right), showing the aforementioned scaffolds' antimicrobial activity against with *S. aureus*. Adapted with permission from Ref. ²⁴⁰

Carbon-based nanomaterials

Polymer nanocomposites based on carbon-based materials have been used to improve the mechanical, electrical, chemical and osteogenic properties of polymers. The basic structure of carbon-based materials used for these applications is graphene, which is a one atom thick hexagonal lattice of sp² hybridized carbon atoms, a structure that confers outstanding mechanical properties (E modulus = 1 TPa, ultimate strength = 130 GPa), thermal conductivity (5000 W/ mK), electrical conductivity (up to 6000 S/m) and chemical properties. ²⁴¹ Depending on its dimensional arrangement, graphene is referred as 0-D fullerene or wrapped graphene, 2D- carbon nanotube (CNT) or rolled graphene, and 3D- graphite or stacked graphene. ²⁴² CNTs, generally prepared by chemical vapor deposition methods, can be single walled (SWCNTs), with a diameter of 1 nm, or with several nanotubes arranged concentrically into a multi-walled structure (MWCNTs), with diameters generally between 10 and 100 nm. ²⁴³ Despite their narrow diameter, they possess very high aspect ratios, which makes them ideal reinforcing fibers in composites. Moreover, their geometrical structure can be tuned to adjust their electronic properties, from metallic to semiconducting materials. ²⁴⁴ A few studies have reported the use of MWCNTs as fillers on ME-AM scaffolds for bone regeneration applications. For instance, Huang et al. loaded PCL scaffolds with up to 3 wt% MWCNTs, which were shown to be aligned in the extrusion direction. ²⁴⁵ This resulted in an increase in the compressive modulus from ~ 50 MPa to ~ 88 MPa. Moreover, hAD-MSCs showed increased proliferation with increasing MWCNTs content. Taking advantage of the alignment of the MWCNTs along the extrusion direction,

a subsequent study focused on the development of a ternary composite consisting of PCL, MWCNTs (0.75 wt%) and nHA (20 wt%), which aimed at mimicking the bone tissue structure at the nanoscale.²⁴⁶ The aligned MWCNTs induced the PCL chains in their vicinity to recrystallize along their walls, which surrounded by nHA particles, mimicked the highly aligned and mineralized collagen fibrils in bone. Such ternary scaffolds also possessed higher compression moduli and showed enhanced upregulation of ALP, COL I and OCN gene expression of hAD-MSCs, compared to bare PCL and PCL/MWCNT scaffolds.²⁴⁶

Wang et al. compared the effect of MWCNTs and graphene sheets on PCL scaffolds mechanical properties and cell-affinity.²⁴⁷ PCL/MWCNTs and PCL/graphene scaffolds with up to 3 wt% filler content were prepared. Interestingly, at a given filler concentration, PCL/graphene scaffolds presented higher compression modulus and strength, as well as higher protein adsorption and hAD-MSCs proliferation rates than PCL/MWCNT scaffolds. This was correlated to the better mechanical, physicochemical and conductive properties of graphene compared to MWCNTs. Indeed, graphene sheets prepared by liquid phase exfoliation of graphite, have been used as filler for PCL scaffolds in several studies from Bartolo's group.^{239, 248-250} In all of them, a maximum of 0.78 wt% filler was used to avoid potential hAD-MSCs cytotoxicity effects of graphene upon complete scaffold degradation, which have been previously shown to be dose and size dependent.²⁵¹ These reports demonstrated that the sole addition of 0.78 wt% graphene improved the scaffolds compression modulus by 50%. Moreover, PCL/graphene scaffolds showed enhanced hAD-MSCs proliferation compared to bare PCL scaffolds,²⁴⁸ which could be further improved by a chemical treatment with NaOH,²⁴⁹ or by the pre-adsorption of the P1-latex protein to the scaffolds filaments.²⁵⁰ Ultimately, untreated PCL/graphene scaffolds with 0.78 wt% filler were implanted in a rat calvarial defect, and electrical stimulation was applied, to assess their *in vivo* bone formation ability (**Figure 6F**).²³⁹ Histological analysis revealed that electrically stimulated PCL/graphene 0.78wt% scaffolds led to more tissue infiltration and mineralization at early timepoints (60 days), compared to other conditions. Yet, values were leveled at later timepoints (120 days).²³⁹

Due to the high synthesis costs and low scale production of graphene through the aforementioned CVD or direct exfoliation of graphite methods, currently the most commonly used method for large-scale production of graphene-like materials relies on graphene oxide (GO), the oxidized version of graphite, as starting material.²⁵² GO can be

easily mass produced by oxidation of graphite through well-established methods. Moreover, GO possesses oxygen-containing functional groups disrupting each carbon plane, including epoxy, carbonyl, ketone and hydroxyl groups, which offer the possibility of functionalization with biomolecules to tune its bioactivity.²⁵³ Unagolla et al. prepared PCL/GO 0.5 wt% and showed that while the addition of GO did not influence significantly the mechanical properties of the scaffolds, it led to a slight increase of matrix mineralization and osteogenic proteins expression at early timepoints, compared to bare PCL scaffolds.²⁵⁴ In contrast, Belaid et al. reported that the addition of 0.3 wt% GO to PLA scaffolds led to a 30% increase in the compression modulus. Moreover, MG63 proliferation was enhanced on these scaffolds, as well as calcium deposition when cultured in osteogenic medium.²⁵⁵ Chen et al. significantly augmented the filler loading, preparing TPU/PLA/GO scaffolds with up to 5 wt% GO, which led to a 125% increase in compression modulus, and good biocompatibility.²⁵⁶ Despite its facile production, GO is non-conductive and possesses poor thermal stability, which can lead to the formation of air pockets within the scaffold filaments when processed at high temperatures, like the ones required for PLA, PLGA or PEOT/PBT extrusion (~200° C).²⁵⁶ Electrical conductivity and thermal stability can be restored by chemical or thermal reduction of the GO, leading to exfoliated wrinkled reduced graphene oxide (rGO) sheets, similar to pristine graphene, but with some oxygen defects at the laminar edges, and holes on the carbon skeleton.²⁵⁷ Angulo-Pineda et al. fabricated PCL/rGO scaffolds with up to 10 wt% filler, which showed to be not only conductive, but also antimicrobial against *S. aureus*.²⁵⁸ Electrical stimulation on such scaffolds further enhanced the antimicrobial effect, while supporting higher hBM-MSCs proliferation rates than on non-stimulated scaffolds. In spite of rGO-polymer composite scaffolds possessing interesting bioactive properties for bone regeneration, very few reports have studied them. This might very well be due to the very low bulk density of rGO upon thermal reduction, which can lead to particle inhalation during handling and lack of free-flowability into twin-screw compounders, which is the main cost-effective route for composites production at large scale.²⁵⁹

The aforementioned research examples demonstrate the great potential of graphene-based materials as fillers for ME-AM scaffolds for BTE, since they can act as reinforcement agents, boost cell attachment and osteogenic differentiation, and provide antimicrobial activity. However, there are still concerns on the *in vivo* toxicity of these materials and their degradation products. Studies have reported that the cytotoxicity of graphene and derivatives is dependent on their production method and purity, physicochemical

properties (lateral dimensions, oxidation state, aggregation state, functionalization), concentration, exposure time and administration route.²⁶⁰ Yet, this information is scarce and contradictory results are often found, due to the various synthesis routes leading to diverse materials, which often are not properly characterized or described in literature.²⁶¹ In addition, biodistribution studies have mostly been performed upon single intravenous injection of dispersed graphene materials,²⁶² and the effect of local, sustained and long term injections, simulating the release of graphene from a polymeric scaffold, have not yet been considered. Overall, more thorough *in vitro* and *in vivo* biocompatibility studies paired to the growing interest in graphene-based scaffolds development are necessary. This would help to determine the maximum graphene-based filler loading within ME-AM scaffold, to ensure their functionality while retaining their biocompatibility in an *in vivo* scenario.

Bioactive molecules

The direct incorporation of drugs or antibiotics within the scaffolds filaments is desired in order to simplify the fabrication process, as well as to improve the loading capability and release profile. However, the main limiting factor for ME polymer-drug composites is the thermal stability of the incorporated molecules. This limits the bioactive molecule and polymer choice, as the large majority of the thermoplastics used for biomedical applications require high processing temperatures (~175 °C – 220 °C),^{60, 141} which are above the thermal stability limits of most commonly used antibiotics for orthopedic applications (~ 120 °C, depending on exposure time and antibiotic type).²⁶³ As an exception, PCL has low melting (~ 60 °C) and printing temperatures (up to 100 °C), thus several reports have investigated the incorporation of antibiotics or growth factors within PCL scaffolds. For instance, Lee et al. reported the fabrication of a PCL/rifampicin 2.5 wt% composite, which was melt blended and printed at 60 °C.²⁶⁴ During the 14 days evaluated, the antibiotic was sustain released and exerted antimicrobial activity against *E. coli* and *S. aureus*, while maintaining cell viability (**Figure 6G**). Shim et al. reported the preparation of PCL/PLGA/tobramycin scaffolds at 110 °C.²⁴⁰ Here, the drug was sustain released for a period of 50-60 days and when implanted on a rat femur model with osteomyelitis, was able to reduce inflammation and exert antimicrobial activity. In another study, Costa et al. printed PCL/poloxamine/dexamethasone scaffolds.²⁶⁵ The addition of the poloxamine copolymer in the blend led to filaments with lower surface porosity, which slowed down the release rate of dexamethasone. Similarly, the addition of HA to PLA/ nitrofurantoin

scaffolds allowed to tune the antibiotic release, since HA facilitated the infiltration of water within the polymer matrix, leading to faster drug release rates.²⁶⁶ The aforementioned studies also investigated the effect of the drug content (wt%) on the release profile, leading to the general observation that higher loadings led to higher drug amounts released over time.²⁶⁵⁻²⁶⁶ On the other hand, Kempin et al. investigated the effect of polymer type on drug release, using quinine as a model, concluding that for a given drug, desired release profiles might be obtained by choosing the appropriate polymer.²⁶⁷ For a given polymer type and drug concentration, drug type will also define the release profile, as observed by the comparison among two independent studies: PCL/gentamicin 5wt% scaffolds led to a burst release of gentamicin (60% release in 24h), while quinine was released in a sustained manner from PCL/quinine 5wt% scaffolds (60% release in 7 days).²⁶⁷⁻²⁶⁸

As presented, loading bioactive molecules within the polymer matrix of ME-AM ensures higher incorporation efficiencies, allows for larger loading amounts and avoids fast release rates, compared to other strategies. However, this simple and cost-effective technique is at the moment limited to low melting temperature polymers, such as PCL.

Other organic phase combinations

The addition of organic fillers within ME-AM scaffolds has been investigated to improve their mechanical or physicochemical properties. For instance, cellulose nanomaterials have been used to reinforce PLA filaments for FDM, as well as PCL scaffolds, due to their crystalline fibrillar structure.²⁶⁹⁻²⁷⁰ Composite PCL/microcrystalline cellulose scaffolds showed an increase in compression and elastic modulus from 25 MPa to 32 MPa and from 60 to 72 MPa, respectively, with the sole addition of 2 wt% filler.²⁷⁰ Besides its reinforcing capabilities, this biopolymer possesses high specific surface area and surface reactivity, which allows for the introduction of well controlled chemical, structural, and topographical features at a cell relevant scale, which is of great interest for the TE field, but have not yet been widely explored for ME-AM scaffolds.²⁷¹ Silk has also been used to improve the mechanical properties of ME-AM scaffolds. For instance, Aleman-Dominguez et al. demonstrated that the addition of up to 30 wt% silk microparticles within a PCL polymer matrix significantly increased the bulk and nano-mechanical properties of the scaffold, as measured by nanoindentation.²⁷² However, scaffold hydrophobicity was also significantly enhanced at this high concentration, which led to lower protein adsorption and subsequent poor cell affinity. This demonstrated that while a high filler content might be desired for

significantly augmenting the scaffold's mechanical properties, cellular interactions, which are of outmost importance for the ultimate scaffold application, might be compromised at the same time. In this regard, the concentration of polyaniline (PANI) particles within a PCL scaffold was reduced to a minimum to ensure its biocompatibility.²⁷³ While the addition of this conductive polymer increased both the conductivity and compressive mechanical properties of the scaffold, concentrations above 0.1 wt% were shown to be cytotoxic. Despite these limitations, PCL/PANI 0.1 wt% scaffolds possessed a conductivity of $2.46 \pm 0.85 \times 10^{-4}$ S/cm, which enabled their use as electroactive structures to modulate cellular response and enhance tissue regeneration.²⁷³ Camarero-Espinosa et al. also reported the fabrication of scaffolds with the ability of providing cell stimulation, consisted of PCL/PLA Janus filaments that can be nano-deflected via external application of ultrasounds.²⁷⁴ Such Janus scaffolds were spontaneously formed by phase separation of PCL/PLA blends during ME, and could perform as ultrasound transducers, enhancing cell proliferation, matrix deposition and osteogenic differentiation of hBM-MSCs, via the formation and activation of voltage-gated calcium ion channels.

Conclusions and future perspective

ME-AM enables the fabrication of complex-shaped personalized scaffolds for BTE applications, with an interconnected and highly controlled porosity and from a wide number of biodegradable and mechanically robust thermoplastic polymers. Compared to other AM technologies, ME-AM is a cost-effective and relatively simple technique, and does not require the use of potentially toxic binders, curing agents or solvents, which simplifies the clinical translation of the fabricated constructs. Furthermore, while synthetic thermoplastic polymers are often bioinert and lack cell-recognition sites to promote cell attachment, proliferation and osteogenic differentiation *in vitro* and *in vivo*, they are quite amenable to bioactive functionalization. A large amount of research has been carried out over the last decades on augmenting the bioactivity of ME-AM scaffolds. Modifying the scaffold architecture by adjusting the scaffold pore shape and size has been shown to influence cell attachment and differentiation. Similarly, the introduction of microporosity to the scaffold filaments' bulk and surface via ME-AM combined with salt leaching or gas foaming, has shown to increase the scaffold surface area for enhanced cell attachment. While these approaches have shown benefits, they can, in some cases, limit nutrient availability and diminish the scaffold's mechanical properties. On the contrary, surface

modification strategies aiming at the introduction of roughness, specific chemical functional groups, or bioactive coatings to the filaments surface, do not compromise the scaffold structural integrity, and thus have been more extensively studied. In particular, plasma surface modification has stood out as a multi-approach technique for enhancing cell attachment and steering cell differentiation towards a specific lineage. In this regard, surface roughness can be introduced via plasma activation, specific polar functional groups can be deposited via plasma polymerization, and scaffold filaments can be primed for the subsequent covalent binding of biomolecules via plasma grafting. In addition to the conjugation of biomolecules, biomimetic PDA, CaP and full ECM coatings have also been investigated to enhance the surface functionality of scaffolds towards enhanced cell attachment and differentiation.

Although surface modifications can successfully augment the bioactivity of ME-AM scaffolds, we have seen that these approaches frequently involve the use of wet chemistry, and most importantly, they often require complex and multiple post-fabrication steps. Further incrementing the scaffold complexity, fibrous or hydrogel networks have been combined with the ME-AM framework towards hierarchical and hybrid scaffolds, which have been shown to increase cell retention and provide a biomimetic platform for tissue formation. Polymer-filler composite scaffolds have shown to induce specific cell behavior with the sole addition of fillers to the polymer matrix, avoiding the need of post fabrication steps. Moreover, composites can enhance not only the scaffold physicochemical properties, but also their mechanical and degradation properties, towards a better scaffold performance. In addition, composite production via melt blending by twin screw compounding ensures a complete solvent-free scaffold fabrication process. For all these reasons, a large body of literature has been focused on the fabrication of composite ME-AM scaffolds using a variety of fillers, which include CaPs, silicates, metallic particles, carbon based materials, bioactive molecules and other organic materials, such as cellulose. Research has been focused on trying to identify the ideal filler type and concentration necessary to generate an optimum physicochemical and biological response, without compromising material printability, and while ensuring scaffold optimum mechanical properties and biocompatibility.

In spite of all these efforts, a ME-AM scaffold with optimum properties that can ensure optimum cell attachment and enhanced osteogenic differentiation towards a fully functional engineered *in vitro* and *in vivo* tissue is yet to be developed. There are still some

challenges to overcome to achieve this aim, as well as aspects that would need attention in future research.

Currently, newly developed scaffolds can be easily characterized, in terms of printability, surface physicochemical analysis, filler dispersion in a composite, and mechanical properties thanks to the fast pace with which material science behind scaffold fabrication is progressing. Yet, the *in vitro* biological evaluation methodologies of such scaffolds are often insufficient to determine their *in vivo* efficacy. As a common strategy, a short *in vitro* evaluation of cell attachment, proliferation, differentiation, and early stage mineralization is often discussed. While these parameters are relevant for an initial estimation of the scaffolds functionality, long-term evaluations, coupled to the long process of bone regeneration, would be necessary to accurately determine the actual scaffold performance. In order to obtain more reliable *in vitro* short and long term information, as well as to reduce the number of animal experiments towards clinical translation, scaffolds could be cultured in bioreactors, where they could potentially be exposed to relevant cell populations and pro-osteogenic external stimulus (chemical, mechanical, electrical, ultrasonic).²⁷⁵⁻²⁷⁶ The flexible (sometimes elastomeric) synthetic thermoplastic polymers used in AM are best suited for mechanical stimulation. Such dynamic systems, in combination with computational models, could help to better understand and predict how specific scaffold architectures and the subsequent forces transmitted to cells (mechanical strain, fluid shear stress) can affect cell fate.²⁷⁷⁻²⁷⁸ This information could be used to improve the scaffold design towards enhanced tissue formation. In addition, bioreactor systems would also allow to analyze the long term stability of the scaffold modifications, such as plasma treatments, protein coatings, or bioactive CaP layers. Moreover, polymer degradation profiles and fillers release rates from composite scaffolds could be monitored to evaluate their *in situ* cytocompatibility. This could lead to more complete and focused studies and help finding the ideal scaffold for bone regeneration.

Another overlooked matter on ME-AM scaffolds for BTE is the formation of a vascular network. Due to their macroporous structure and their relatively small size for *in vitro* cultures, ME-AM scaffolds can support cell ingrowth and waste-nutrients exchange *in vitro* without the need of vascularization. Yet, scaffolds aimed to be used in large bone defects will likely require the integration of a vascular network to support tissue formation. Therefore, it is of great significance to implement cues to promote vascularization within ME-AM scaffolds, in addition to cues for cell attachment and osteogenic differentiation. In

this regard, ME-AM scaffolds providing gradients in growth factors (i.e. VEGF) could induce blood vessels infiltration into the scaffold core.²⁷⁹ As presented in this review, current research is focused on the incorporation of bioactive cues to polymeric ME-AM scaffolds, while little efforts are being placed on developing new polymers, where the lack of functionality relies. Future research lines could relate to the design of new functional synthetic polymers, which can influence cellular behavior on their own, without the need of additional elements. Overall, this could lower the scaffold complexity, which would potentially simplify the production process, the *in vitro* and *in vivo* evaluation, the clinical translation path and, at the end of the chain, the final medical product price.

Despite current shortcomings, ME-AM scaffolds hold great promise in the future of bone regeneration. Appropriate scaffold design and architecture, specific bioactive cues for promoting cell attachment, differentiation and tissue formation, and thorough material and biological evaluations shall certainly be combined towards the construction of functional scaffolds for bone tissue regeneration.

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Chapter 3

**Improving cell distribution on 3D additive
manufactured scaffolds through engineered seeding
media density and viscosity**

Improving cell distribution on 3D additive manufactured scaffolds through engineered seeding media density and viscosity

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Abstract

In order to ensure the long-term *in vitro* and *in vivo* functionality of cell-seeded 3D scaffolds, an effective and reliable method to control cell seeding efficiency and distribution is crucial. Static seeding on 3D additive manufactured (AM) scaffolds made of synthetic polymers still remains challenging, as it often results in poor cell attachment, high cell sedimentation and non-uniform cell distribution, due to gravity and to the intrinsic macroporosity and surface chemical properties of the scaffolds. In this study, the biocompatible macromolecules dextran and Ficoll (Ficoll Paque) were used for the first time as temporary supplements to alter the viscosity and density of the seeding media, respectively, and improve the static seeding output. The addition of these macromolecules drastically reduced the cell sedimentation velocities, allowing for homogeneous cell attachment to the scaffold filaments. Both dextran- and Ficoll -based seeding methods supported human mesenchymal stromal cells viability and osteogenic differentiation post-seeding. Interestingly, the improved cell distribution led to increased matrix production and mineralization compared to scaffolds seeded by conventional static method. These results suggest a simple and universal method for an efficient seeding of 3D AM scaffolds, independent of their material and geometrical properties, and applicable for bone and various other tissue regeneration.

INTRODUCTION

In vivo, cells are surrounded by a complex 3D microenvironment that supports their growth and assembly into functional tissues. Upon tissue damage caused by trauma or disease, the structural integrity of this extracellular matrix (ECM) can be compromised leading to lack of essential cues for functional tissue performance, and the requirement of suitable replacements. By using scaffolds, tissue engineering and regenerative medicine (TERM) aim to provide artificial ECM-like structures, which try to mimic the native tissue in terms of architecture, chemistry or mechanical properties to guide cells into the first steps of the regeneration process.¹⁻² To allow cell infiltration, tissue ingrowth and vascularization, 3D scaffolds with a highly interconnected porous structure are required.³⁻⁴ In the past decade, additive manufacturing (AM) and, more specifically, melt extrusion (ME)-based techniques, have emerged as one of the most appealing technologies to produce scaffolds for TERM.⁵⁻⁶ Compared to other conventional methods such as gas foaming/particulate leaching,⁷⁻⁸ freeze-drying⁹⁻¹⁰ or phase separation,¹¹⁻¹² AM allows the reproducible fabrication of complex 3D structures with interconnected, high porosity, from various types of biocompatible and biodegradable polymers with optimum mechanical properties desired for tissue regeneration applications.

Efficient cell seeding of 3D scaffolds is a key step towards the development of *in vitro* tissues for TERM applications. Cell density and their spatial distribution in macroporous scaffolds are critical to ensure the functionality of the engineered tissue, as these parameters will dictate intercellular and cell-material interactions, thereby affecting different cellular behaviors such as migration, proliferation, differentiation and matrix production.¹³⁻¹⁴ To improve cell distribution within 3D scaffolds, different dynamic seeding methods have been applied. Compared to static seeding, dynamic seeding methods, which apply an external force to facilitate cell penetration, such as flow perfusion, centrifugation, orbital shaking, and spinner flask, have shown to result in better cell distribution.¹⁵⁻¹⁹ Moreover, culture of 3D scaffolds, fabricated with conventional techniques, in perfusion bioreactors after static seeding has also demonstrated to enhance cell distribution and tissue formation.²⁰ However the use of bioreactors require specific lab equipment and expertise and perfusion flow right after seeding results in significant decrease in cell number in 3D AM scaffolds.²¹ Overall, due to the intrinsic scaffold characteristics, i.e. large and interconnected pore architectures, most of the dynamic techniques show limited applicability for seeding or culture AM scaffolds. Furthermore, the

lack of specific cell adhesion sites in the majority of the polymers suitable for AM leads to poor cell–material interactions, which along with the gravitational force results in cell sedimentation during static seeding and, therefore, non-uniform cell distribution within the scaffold. Nevertheless, the simplicity of the static seeding over dynamic seeding has led to further developments of this method for seeding 3D AM scaffolds, starting by the modification of basic seeding parameters such as cell number, seeding volume, seeding time and seeding vessel optimization. For instance, seeding times up to 4h, seeding volumes comparable to the scaffold's free volume and not very large cell numbers have shown to give the highest cell seeding efficiency upon static seeding.²²⁻²⁴ On the other hand, different surface modification techniques capable of improving the chemical and biological performance of synthetic polymers have also shown to increase cell adhesion in 3D AM scaffolds. For example, plasma modification using oxygen²⁵ or nitrogen²⁶⁻²⁷ have shown to increase cell adhesion of osteoblastic cell lines in poly(ϵ -caprolactone) (PCL) scaffolds. Chondrocytes adhesion on poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT) 3D AM scaffolds has also been improved by the addition of carboxylic groups to the polymer surface via acrylic acid plasma polymerization.²⁸ Surface functionalization using adhesive polymers such as poly(dopamine) have also demonstrated to improve adhesion of different cell types on PCL and poly(lactic acid) (PLA) scaffolds.²⁹⁻³¹ As an alternative to chemical modification of polymer surfaces, better cell attachment along the scaffold cross section has been attained via decreasing the flow rate of the seeding media through the scaffold's pores³²⁻³³ By varying the strand distance or the angle of deposition in each layer, 3D AM scaffolds with pore size gradients along the vertical direction³² or with different lay-down patterns³³ were fabricated and resulted in enhanced cell attachment and distribution. This was, however, in exchange of varying the scaffolds mechanical properties and, more importantly, reducing the pore size, which hinders cell migration, nutrients and oxygen availability, waste management, and angiogenesis for pore sizes smaller than 300 μm .^{4, 34-36}

A large improvement in cell seeding efficiency and distribution was recently obtained by seeding 3D AM scaffolds with cell-laden hydrogels, which showed to retain almost the complete cell number mixed within the hydrogel precursor.³⁷⁻⁴¹ However, these hybrid systems are multi-step and require fast gelation times. In addition, cells are exposed to two different microenvironments: the hard scaffold filaments and the soft hydrogel carrier. As it is widely accepted that substrate stiffness can direct cell fate,⁴² having these types

of dual systems might mask the effect of the scaffold material and lead to unexpected results.

Although previous strategies have shown some improvement on cell seeding efficiency and distribution, they require modification of the scaffold's surface chemistry or architecture, influencing the overall microenvironment. Therefore, there is still lack of methodology and knowledge of an elegant cell seeding method applicable to all 3D AM scaffolds regardless of their characteristics. Here, we introduce a novel seeding technique based on controlling cell settling velocity upon static seeding that can be applied to all types of 3D AM scaffolds without requiring any chemical or architectural modification. Two biocompatible macromolecules (macroMs) were employed for the first time as supplements to independently tune i) the viscosity or ii) the density of the cell seeding media. Cells seeded with this novel technique were morphologically characterized using immunofluorescence and scanning electron microscopy. Furthermore, the osteogenic differentiation of human mesenchymal stromal cells (hMSCs) seeded with the macroM-based solutions on AM 3D scaffolds was investigated using various biochemical and molecular assays to evaluate the effect of macroMs on a specific cell differentiation.

MATERIALS AND METHODS

Scaffolds fabrication

Scaffolds were fabricated via a ME based AM technique (Bioscaffolder 3.1, SysENG, Germany). The copolymer PEOT/PBT, with a PEO molecular weight of 300 Da and PEOT/PBT blocks ratio 55/45 wt% (300PEOT55PBT45, PolyVation, The Netherlands) was used. Briefly, the cartridge was filled with PEOT/PBT pellets, heated at 195 °C and extruded by applying a pressure of 4 bar, an auger screw rotation of 40 rpm and a translation speed of 10 mm*s⁻¹. The scaffold architecture consisted of a 0-90 pattern, with a 250 µm fiber diameter, 200 µm layer thickness and 750 µm strand distance (center to center), giving an expected x-y porosity of approximately 500 µm. Cylindrical scaffolds of 4 mm diameter and 4 mm height were punched out from a 20x20x4 mm manufactured block using a biopsy punch and used for further experiments.

Density and viscosity measurements of Dextran and Ficoll based solutions

Ficoll-Paque™ Plus solution (GE Healthcare) (Ficoll-Pq) was diluted to 80, 60, 40 and 20 vol% with cell culture medium (CM), consisting of α MEM with Glutamax and no nucleosides (Gibco) supplemented with 10 vol% FBS (Sigma-Aldrich), penicillin (100 U/ml) and streptomycin (100 μ g/ml) (Fisher-Scientific). Similarly, 10, 5, and 2.5 wt% dextran (500 kDa, Pharmacosmos) solutions were prepared in CM. Densities were calculated by weighing known volumes of each of the solutions (accuracy ± 0.1 ml) in a high precision balance (accuracy ± 0.1 mg) and applying the density formula.. The relative viscosity (with respect to water) of the solutions was empirically determined by timing the flow of water and the same volume of each solution through the same fluidic circuit at the same constant pressure.

Determination of hMSCs density using density gradient separation

Density of hMSCs was measured using Ficoll-Pq density gradient centrifugation. The Ficoll-Pq solution was diluted to 80, 60, 40 and 20 vol% with appropriate volumes of CM containing phenol red, which helped to visualize the layers of the gradient. Starting from 100 vol% Ficoll-Pq solution, 2ml of each solution were carefully layered on top of each other in decreasing order in a 15 ml conical tube. One million cells in CM were layered on top of the 20% Ficoll-Pq layer. The tube was then centrifuged at 500 rcf for 20 min at room temperature (RT). Subsequently, the layers were carefully separated by pipetting and the cells in each layer were imaged via brightfield microscopy and quantified using a Neubauer chamber. As a comparative method, cell density was additionally determined through buoyancy force-driven separation, which does not require the application of mechanical forces. For the buoyancy separation, the various concentrations of Ficoll-Pq media were similarly layered in a 15 ml conical tube starting from 100 vol% Ficoll-Pq solution. Here, 1 million cells in medium were carefully pipetted at the bottom of the tube under the 100% Ficoll-Pq layer using a long syringe needle, without disturbing the layers. The tube was then incubated at RT for 3h to let cells be pushed by buoyancy forces to layers of similar density. After separation, the layers were collected separately and cells present in each solution were imaged and quantified using a Neubauer chamber.

Cell seeding and culture

Cell expansion

HMSCs isolated from bone marrow were purchased from Texas A&M Health Science Center, College of Medicine, Institute for Regenerative Medicine (Donor d8011L, female, age 22). Cryopreserved vials at passage 3 were plated at a density of 1,000 cells*cm⁻² in tissue culture flasks and expanded until approximately 80% confluency in CM without Pen/Strep at 37 °C / 5% CO₂.

Cell seeding and culture in 3D scaffolds

Scaffolds were disinfected in 70% ethanol for 20 min, washed 3 times with phosphate buffered saline (PBS) and incubated overnight in CM for protein attachment. Before seeding, scaffolds were dried on top of a sterile filter paper and placed in the wells of a non-treated hydrophobic tissue culture well plate. Trypsinized passage 4 hMSCs suspension was centrifuged for 5 min at 500 rcf and then resuspended in CM, Dextran-based CM, or Ficoll-Fq -based CM at a density of 200,000 cells per 37 µl (approximate scaffold free volume). Dextran-based CM consisted of 10 wt% dextran in CM and Ficoll-Pq based CM consisted of 60 vol% Ficoll-Pq in a FBS adjusted cell culture medium (α-MEM supplemented with 25 vol% FBS and 1 vol% Penstrep, to obtain 10 vol% FBS in the final medium). A 37 µl droplet of each cell suspension each was placed on top of each scaffold, drawn into the scaffold's free volume by capillary forces and held inside the pores by the hydrophobic well under it. Scaffolds seeded with CM are referred to as conventional seeded (CS) scaffolds, while macroM seeded (MS) scaffolds are referred to as MS-Ficoll-Pq and MS-Dextran, when seeded with Ficoll-Pq -based CM and dextran-based CM, respectively. Seeded scaffolds were incubated for 4 h at 37 °C / 5% CO₂ to allow cell attachment. After this time, scaffolds were transferred to new wells containing 1.5 ml of basic media (BM) (CM supplemented with 200 µM L-Ascorbic acid 2-phosphate (Sigma-Aldrich)). BM was replaced after 24h and every two or three days from then on. To evaluate hMSCs osteogenic differentiation, scaffolds were cultured for another 21 days in BM or mineralization media (MM) after 7 days in BM. MM consisted of BM supplemented with 10 nM dexamethasone (Sigma-Aldrich) and 10 mM β-glycerophosphate (Sigma-Aldrich).

To visualize the potential entrapment of macroMs within the scaffold after seeding, cells were resuspended in 10 wt% dextran-FITC (Fluorescein Isothiocyanate-Dextran 500 kDA, Sigma-Aldrich) or 60 vol% Ficoll-Pq-FITC at the same cell density (200,000 cells in 37 µl).

The Ficoll-Pq-FITC based CM was prepared by first dissolving 5.7% (wt/vol) polysucrose-FITC (400 kDa, Sigma-Aldrich) and 9% (wt/vol) sodium diatrizoate hydrate (Sigma-Aldrich) in sterile water, to match Ficoll-Pq composition. 60 vol% of this solution was prepared in FBS adjusted cell CM as previously described. A 37 μ l droplet of each cell suspension was placed on top of each scaffold and incubated for 4 h at 37 °C/ 5% CO₂ to allow cells attachment. After this time, scaffolds were transferred to new wells containing 1.5 ml of BM and cultured for 24h.

To rule out macroM adhesion, as well as macroM-driven enhancement of serum proteins adsorption to the scaffold filaments, as the causes of improved cell seeding, scaffolds were pre-incubated for 4h with dextran- and Ficoll-Pq - based CM. Non-adsorbed macroMs were removed by a single scaffold wash with PBS and the scaffolds were then dried on top of a sterile filter paper. Subsequently, scaffolds were seeded with 37 μ l of CM containing 200,000 cells and incubated for 4 h at 37 °C / 5% CO₂ to allow cell attachment, after which scaffolds were analyzed.

Cell seeding in 2D for viability and osteogenic differentiation evaluation

Preliminary experiments in 2D were performed in order to assess the viability and osteogenic potential (mineralization) of cells pre-incubated in macroMs solutions. This was done in order to rule out the direct effect of macroM in cell death or in enhanced mineralization in a 3D setting. To assess the viability of cells at different timepoints when in a macroM solution, trypsinized cells were centrifuged and resuspended in CM, dextran-based CM or Ficoll-Pq -based CM at a density of 200,000 cells per 37 μ l. NucBlue® Live and NucGreen® Dead reagents (ReadyProbes® Cell Viability Imaging Kit, Invitrogen) were added as well, at concentrations defined by the manufacturer protocol. Cell suspensions were incubated at 37 °C/ 5% CO₂ in plates mimicking the 4h seeding in 3D scaffolds and imaged at t = 0 h and t = 4 h after seeding using an inverted fluorescent microscope (Eclipse, Ti2-e, NIKON). For optimal visualization in the reported images, live cells were false colored in green while dead cells were given a red color. In order to assess the osteogenic differentiation potential (mineralization) of cells pre-incubated with macroMs, cells pre-incubated for 4h in macroM based solutions were centrifuged and resuspended in fresh CM to mimic the post-seeding culture condition of cells in scaffolds. Cells were counted and seeded at a density of 5,000 cells*cm⁻² in tissue culture polystyrene well plates. Media was refreshed every two or three days. Scaffolds were cultured for 21 days in BM or MM after 7 days in BM.

Imaging and quantification of cell distribution within scaffold cross section

After seeding and 4h incubation for attachment, CS and MS scaffolds were washed with PBS and fixed with 4 wt% paraformaldehyde for 30 min, followed by washing steps with PBS. Cells were permeabilized using 0.1 vol% Triton-X for 30 min, washed twice with PBS and incubated with phalloidin (Alexa Fluor 568, 1:75 dilution in PBS) for 1h at RT. Finally, samples were washed with PBS. The bottom and cross section of scaffolds were imaged using an inverted fluorescent microscope. Cross section images were analyzed using ImageJ software. Briefly, images were converted to 8-bits format and the local contrast was enhanced (blocksize 50, maximum slope 3, no mask). Next, the background was subtracted and the images were converted to binary (rendering regions with cells in white and the rest in black). Each image was divided into four regions corresponding to heights of 0-1 mm, 1-2 mm, 2-3 mm, 3-4 mm starting from the bottom of the scaffolds and the total amount of white pixels in each region was quantified and normalized to the total number of pixels in the scaffold cross section. Values are reported as normalized cell coverage area in percentage.

Imaging of cell viability and macroMs entrapment in 3D scaffold

After 4h seeding and 1 and 7 days of culture, dead cells were stained for 20 minutes prior to fixation using the LIVE/DEAD™ Fixable Dead Cell Stain Kit (Thermo Fisher Scientific) at a concentration of 0.5 μ l stain in 500 μ l Hank's Balanced Salt Solution per scaffold. Subsequently, samples were washed and fixed with 4 wt% paraformaldehyde for 30 min, followed by three washing steps with PBS. Cells were stained with phalloidin as stated in section 2.5. Live/dead images of scaffolds at day 1 and 7 were acquired using an inverted fluorescent microscopy. Confocal laser scanning microscopy of macroMs entrapment/dead cells at 4h and day 1 was performed with a Tandem confocal system (Leica TCS SP8 STED), equipped with a white light laser (WLL). Samples were excited with the dye specific wavelengths using the WLL or a photodiode 405 in the case of DAPI. Emission was detected with PMT detectors (DAPI) or HyD detectors (phalloidin, dextran-FITC, Ficoll-FITC, dead cells).

Scanning electron microscopy (SEM) imaging of cells in scaffolds

After 4 h seeding, 1 day of culture in BM and 28 days of culture in MM, scaffolds were washed with PBS and fixed with 4 wt% paraformaldehyde for 30 min, followed by washing

steps with PBS. Sample water content was removed with a graded ethanol series (30, 50, 70, 80, 90, 96, 100 vol%), each 15 min at RT. Drying of samples was achieved by 15 min incubation in 100% ethanol- hexamethyldisilazane (HMDS) at a 1:1 ratio for 15 min, followed by 15 min incubation in HMDS. HMDS was removed and scaffolds allowed overnight drying at RT. Samples were gold sputter coated and examined using SEM (Phillips XL-30), with the detector operating at 5 kV.

Biochemical assays

Alkaline phosphatase (ALP) assay

ALP activity was evaluated after 7 and 21 days of culture in MM (time points day 14 and 28, respectively). 3D scaffolds were collected at every time point and washed with PBS, cut in half, stored at -80 °C and freeze-thawed 3 times. Samples were incubated for 1h at RT in a cell lysis buffer composed of 0.1 M KH_2PO_4 , 0.1 M K_2HPO_4 , and 0.1 vol% Triton X-100, at pH 7.8. 40 μl of the chemiluminescent substrate for ALP CPD star- ready to use reagent was added to 10 μl of cell lysate. After 15 min incubation, luminescence (emission= 470 nm) was measured using a spectrophotometer (Biodrop). Remaining cell lysates were used for DNA quantification. Values are reported normalized to DNA content.

DNA assay

DNA assay was performed on cells cultured on 3D scaffolds after 4h of seeding (time point 4h), after 7 days in BM (time point day 7), and after an extra 7 and 21 days of culture in BM or MM (time point day 14 and 28, respectively). In addition, DNA quantification was also performed on 2D samples using CyQUANT Cell Proliferation Assay Kit (Thermo Fisher Scientific). Samples (lysed samples from ALP activity assay or frozen samples after time point collection) were incubated overnight at 56 °C in Proteinase K solution (1mg*ml⁻¹ Proteinase K (Sigma-Aldrich) in Tris/EDTA) for matrix degradation and cell lysis. Subsequently, samples were freeze-thawed three times and incubated 1h at RT with lysis buffer (cell lysis buffer from the kit diluted 20x in dH₂O) containing RNase A (1:500) to degrade cellular RNA. Lysed samples were incubated with the fluorescent dye provided by the Cyquant kit (1:1) for 15 min and fluorescence was measured (emission/excitation = 520/480 nm) with a spectrophotometer. DNA concentrations were calculated from a DNA standard curve.

Alizarin red S (ARS) staining and quantification

Calcium mineralization was qualitatively determined by ARS staining after 7 and 21 days of culture in BM or MM (time point day 14 and 28, respectively). 3D scaffolds were washed with PBS and fixed with 4 wt% paraformaldehyde for 30 min, followed by three washing steps in distilled water. Subsequently, scaffolds were cut in half and each section was stained with ARS solution (60 mM, pH 4.1-4.3) for 20 min at RT, and washed several times with distilled water until no more stain was leaching out. Images were taken using a stereomicroscope (Nikon SMZ25).

After imaging, a protocol adapted from Gregory et al.⁴³ was used to quantify calcium deposition. Briefly, stained samples were incubated in Eppendorf's for 1h at RT with 30 vol% acetic acid while shaking, followed by 10 min incubation at 85 °C. Afterwards, scaffolds were removed and solutions were centrifuged at 20,000 rcf for 10 min. An appropriate volume of 5M ammonium hydroxide was added to the supernatants to readjust the pH to 4.1-4.3. The absorbance was measured at 405 nm using a spectrophotometer. Concentration of ARS was calculated from an ARS standard curve and the values were normalized to DNA content.

Gene expression

Gene expression analysis was performed at day 14 (7 days in MM) and day 28 (21 days in MM). RNA was extracted from cells in scaffolds using a Trizol isolation method at the selected time points. Briefly, samples were transferred to Eppendorf tubes and Trizol was added. ECM was precipitated at the bottom of the tube through a centrifugation step at 12,000 rcf for 5 min. The supernatant was transferred to a new tube and chloroform was added to isolate the RNA, present in the aqueous phase after phase separation via centrifugation at 12,000 rcf for 5 min. RNA was further purified using RNeasy mini kit column (Qiagen), according to the manufacturer's protocol. The purity and quantity of total RNA was evaluated using a spectrophotometer. Reverse transcription was performed using iScript™ (Bio-Rad) following suppliers' protocol. The obtained cDNA was used in combination with SYBRGreen master mix (Qiagen) and the selected primers (**Table S1**, Supplementary Information) to perform qPCR using CFX Connect™ Real-Time System (Bio-Rad) under the following conditions: cDNA was denaturated for 3 min at 95 °C, followed by 40 cycles consisting of 15 s at 95 °C and 30 s at 65 °C. Additionally, a melting curve was generated for each reaction in order to test primer dimer formation and non-

specific amplification. Gene transcription was normalized to the transcription of the housekeeping human B2M gene. The $2^{-\Delta\Delta Ct}$ method was used to calculate relative gene expression for each target gene. Normalization was done with respect to relative gene expression of cells in CS scaffolds at day 14.

Statistical analysis

All data is shown as average with error bars indicating the standard deviation of at least three replicates. Analysis of statistics was conducted with GraphPad Prism (version 8.0.1). A one-way or two-way analysis of variance (ANOVA) was performed followed by a Tukey's or Sidak's post-hoc multiple comparison test (as recommended by the software) to evaluate statistical significance and correct for multiple comparisons.

RESULTS

Optimization of the macroM seeding methods

If cells in suspension are considered as rigid spheres, upon which frictional (viscosity related), buoyant (density related) and gravitational forces are exerted, their settling velocity in medium can be derived from the Stokes law.⁴⁴ When the sum of the frictional and buoyancy forces exerted on the cell, due to the surrounding medium, balance the gravitational force, the constant settling velocity (v) can be defined by:

$$v = \frac{2(\rho_{\text{cell}} - \rho_{\text{medium}})}{9\mu_{\text{medium}}} gR^2$$

where ρ_{cell} refers to cell density, ρ_{medium} refers to density of the surrounding cell culture medium, μ_{medium} refers to the viscosity of the surrounding cell culture medium, g is the acceleration due to gravity, and R the cell hydrodynamic radius. According to this equation, to maintain cells in suspension the settling velocity can be decreased by increasing the media viscosity or reduced to zero by matching the density of the media and the cells. In this study, we investigated both approaches independently (**Figure 1**). The media viscosity was changed with the addition of dextran. As shown in **Figure S1A**, by the addition of dextran up to 10 wt%, the viscosity of CM increases exponentially up to around 25 times. As our second approach, the density of the seeding medium was increased by the addition of Ficoll-Paque™ Plus (Ficoll-Pq) to match the cell density and therefore obtain a theoretical settling velocity equal to zero ($v=0$). Ficoll-Pq is an aqueous solution that also

contains sodium diatrizoate, a compound that allows to have a high density solution while maintaining its viscosity relatively low (**Figure S1B**, Supplementary Information). Notably, the viscosity of this solution is only ~3 times higher than CM viscosity. A density gradient of Ficoll-Pq (**Figure S1D**) allowed for the experimental determination of the density of the hMSCs, which is known to be lower than $1.073 \text{ g}\cdot\text{ml}^{-1}$ ⁴⁵⁻⁴⁶. Both separation by gradient centrifugation and by buoyancy forces determined that the density of the hMSCs population lied in between 1.034 and $1.052 \text{ g}\cdot\text{ml}^{-1}$ corresponding to 40 vol% and 60 vol% Ficoll-Pq in CM (**Figure S2**). In order to encompass the majority of the hMSCs population (i.e. to maintain most of the cells in suspension) 60 vol% Ficoll-Pq solution (Ficoll-Pq - based CM) was chosen as the density-based media for seeding the MS-Ficoll-Pq, with a relatively ignorable influence on the viscosity of CM (only 1.65 fold increase) (**Figure S1B**). On the other hand, the viscosity is considered the only effective parameter on the cell settling velocity in a 10 wt% Dextran solution (Dextran-based CM), which was chosen as the viscosity-based media for seeding the MS-Dextran scaffolds. The density of 10 wt% dextran ($1.024 \text{ g}\cdot\text{ml}^{-1}$) is significantly lower than 60 vol% Ficoll-Pq density ($1.052 \text{ g}\cdot\text{ml}^{-1}$) as well as 40 vol% Ficoll-Pq density ($1.034 \text{ g}\cdot\text{ml}^{-1}$), and therefore, lower than hMSCs density. Specifically, its density is comparable to that of a 20 vol% Ficoll-Pq solution ($1.027 \text{ g}\cdot\text{ml}^{-1}$) (**Figure S1C, E**), where a minimum amount of cells were found during separation by Ficoll-Pq gradients (**Figure S2**).

Cell distribution using MS methods

To assess the cell distribution across highly porous PEOT/PBT scaffolds seeded with each of the seeding methods, cells in their cross sections and bottoms were imaged and the cross section images were divided into four horizontal regions of 1 mm each, from 0 to 4 mm height. Moreover, cell settling velocity and displacement towards the bottom of scaffolds during the seeding time was calculated and used to support the cell localization results. As demonstrated in **Figure 2A, B**, around 70% of the total cell number found in the scaffolds cross sections was concentrated in the lowest 1 mm region of the scaffold, as well as attached to the bottom when seeded with the CS method. Using this method, cells sedimented at a theoretical velocity of $1.8 \mu\text{m}\cdot\text{s}^{-1}$ and displaced 2.6 mm towards the bottom of the scaffold (~75% of the scaffold height) during the 4h seeding (**Figure S1F**). In contrast, theoretical cell settling velocity decreased ~50-fold for MS-Dextran when compared to the CS method, or remained zero when seeded with the MS-Ficoll-Pq

Figure 1. Schematic representation of the static seeding approaches discussed in this study: Conventional seeding (CS), Macromolecular seeding with 10 wt% dextran (MS-Dextran) and macromolecular seeding with 60 vol% Ficoll-Paque (MS-Ficoll-Pq). Influence of the properties of the cell seeding media (viscosity and density) on the settling velocity of cells and, therefore, on cell attachment along scaffold height is depicted

method, resulting in a theoretical displacement of $\sim 55 \mu\text{m}$ or $0 \mu\text{m}$ towards the bottom of the scaffold during the 4 h seeding, respectively (**Figure S1F**). This allowed the cells to remain in suspension and to attach to the filaments of the scaffolds in a more homogenous manner (**Figure 2A**). Around 20 to 30% of the total cell number in the cross section was found in each of the 1 mm subdivisions of the scaffold (**Figure 2C, D**), regardless of the use of the MS-Dextran or the MS-Ficoll-Pq method, suggesting that both approaches successfully improved cell distribution along scaffold's cross section. Furthermore, no cells were found at the bottom of scaffolds when seeded with the MS methods, contrary to what was observed in the case of CS scaffolds, which further demonstrates the enhanced cell attachment. Additionally, cells in suspension along the scaffold depth that did not attach to the filaments were washed away after immersing the scaffold in media.

As observed in **Figure 3A**, cells preferentially attached to the top curved surface of the scaffold filaments during the seeding and, due to cell clustering and a larger number of cells attached, cell projections were not clear in the MS methods when compared to the spindle-like cells resulted from the CS method after 4h seeding. Nevertheless, only after 24h post-seeding cells in the MS scaffolds showed a clear elongated morphology, characteristic of migrating cells, and covered the whole surface of the filaments in contrast to the less crowded filaments on CS scaffolds (**Figure 3B**).

Figure 2. Fluorescent staining (F-actin, red) showing that MS methods improved cell distribution along scaffold cross section. (A) Fluorescence microscopy images of hMSCs in the bottom sides (top) and cross sections (bottom) of scaffolds seeded with the CS, MS-Dextran and MS-Ficoll-Pq methods and (B-D) respective quantification of cell distribution across the scaffold height. Statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (* $p < 0.05$; ** $p < 0.01$; **** $p < 0.0001$). Scale bars 1mm.

Figure 3. Higher cell coverage and matrix production on scaffolds seeded with the MS-methods. SEM micrographs of hMSCs on scaffolds seeded with the CS, MS-Dextran and MS-Ficoll-Pq methods after 4h, 1 day and 28 days of culture (21 days in MM) showing cell-scaffold interactions, cell coverage of scaffold filaments and matrix production. Scale bars 100 μ m.

MacroMs in scaffolds

In order to evaluate the potential presence of macroMs in the scaffolds after seeding, scaffolds were seeded with solutions containing fluorescently labeled dextran or Ficoll. After 4 h seeding, macroMs were not found coating the scaffolds filaments. Besides, cells also sedimented in CS scaffolds, which were pre-incubated with dextran or Ficoll-Pq based CM (**Figure S3**). Collectively, both results suggest that macroMs did not adhere to the scaffold surface to act as cell adhesive molecules nor contributed to better cell adhesion by mediating adsorption of serum proteins to the scaffolds filaments. Interestingly, while for both MS-Dextran and MS-Ficoll-Pq, macroMs co-localized with the cells cytoplasm after 4h seeding (**Figure 4**), their fluorescence signals were drastically reduced after 24h and were found to co-localize only with dead cells, and (**Figure 4** and **Figure S4**). Notably, cell death was visualized within the first minutes of hMSCs incubation with both macroMs

based CM and CM (medium without macroMs) (**Figure S5**). Nevertheless, high cell viability was observed across the MS-Dextran and MS-Ficoll-Pq scaffolds over 7 days culture, as demonstrated in the **Figure 5**.

Figure 4. Initial retention (4h) and rapid elimination (24h) of macroMs in cells attached to the scaffolds. Representative confocal microscopy images of hMSCs (F-actin) and macroM (FITC-labeled) on top of scaffold filament 4h and 24h post-seeding with the MS-Dextran and MS-Ficoll-Pq methods. Dash lines delimitating the scaffold filament. Scale bars 100 μ m.

Osteogenic differentiation on MS scaffolds

To fully assess the functionality of our novel MS methods to seed AM 3D scaffolds for bone regeneration, hMSCs potency and their ability to undergo osteogenic differentiation after seeding was evaluated. Preliminary 2D experiments confirmed that proliferation and mineralization were not affected when cells were pre-incubated in macroMs solutions and seeded in 2D (**Figure S6**). These initial positive results allowed us to further study hMSCs response to macroMs seeding in 3D. Despite the significant effect of MS methods in cell distribution (**Figure 2**), no statistical differences were found in the total number of cells attached to the scaffolds when compared to the CS method (**Figure 6A**). Similarly, looking at cell proliferation (**Figure 6A**), constant cell numbers were observed in scaffolds seeded with all the three methods over time, with the exception of day 28 in MM. At this timepoint,

increased cell numbers were observed on MS scaffolds, compared with earlier time points and with CS scaffolds at day 28. SEM images in **Figure 3C** also demonstrated differences in ECM production at this timepoint among the seeding methods. In the case of MS scaffolds, cells were present both on the scaffold filaments and within pores' volume along with the formation of a dense ECM. On the contrary, ECM density was found to be much lower in CS scaffolds, which correlates to the low cell proliferation values over the whole culture period (**Figure 6A**).

Figure 5. Cell viability is preserved after seeding with MS methods. Representative fluorescent images depicting cell viability (F-actin, green; dead cells, red) 1 day and 7 days post-seeding with the CS, MS-Dextran and MS-Ficoll-Pq methods. Inserts represent the imaged area in scaffold cross-section. Scale bars 250 μm .

ALP activity was measured after 14 and 28 days of culture, corresponding to 7 and 21 days culture in MM (**Figure 6B**). No significant difference was observed in ALP activity at day 14 regardless of the seeding method. However, ALP activity was significantly lower at day 28 when comparing cells in scaffolds seeded with CS and with MS methods. Interestingly, ALP activity increased from day 14 to day 28 in cells seeded with CS method.

In contrast, ALP decreased from day 14 to day 28 in cells seeded with MS methods despite the macroM type.

Figure 6. MS method does not hinder osteogenic differentiation of hMSCs: Evaluation of the osteogenic differentiation potential of hMSCs seeded with the MS-based methods compared to CS. (A) Cell seeding efficiency and cell number progression over 28 days of culture in BM and MM on scaffolds seeded with the CS, MS-Dextran and MS-Ficoll-Pq methods. Statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (*\$ p < 0.05; **\$\$\$ p < 0.01; ****##### p < 0.0001; * for comparisons among seeding methods each time point; # for comparisons with D28 MM among MS-Dextran; \$ for comparisons with D28 MM among MS-Ficoll-Pq). (B) Comparison of ALP activity at day 14 and day 28 of culture (7 and 21 days in MM, respectively) of hMSCs when seeded with the three different methods. (C) Quantification of the ARS staining extracted from scaffolds after 14 and 28 days of culture (7 and 21 days in MM, respectively), normalized to cell number. Statistical significance performed using two-way ANOVA with Sidak's multiple comparison test (n.s. p > 0.05; *p < 0.05; ***p < 0.001; ****p < 0.0001, for comparisons among seeding methods each time point or among time points each seeding method) (D) Stereomicroscopy images of scaffold cross sections stained with ARS after 14 and 28 days of culture (7 and 21 days in MM, respectively). Scale bars 1 mm.

To gain more insight into the potential of osteogenic differentiation of hMSCs in scaffolds seeded with our novel method, matrix mineralization was evaluated using ARS staining to visualize calcium deposits after 7 and 21 days in MM (day 14 and day 28, respectively). Scaffolds cultured in BM were used as controls to ensure the inert behavior of the macroMs in the mineralization process (**Figure S7A**). Intriguingly, as it is shown in **Figure 6D**, calcium deposition on the cross section of MS-scaffolds was observed from day 14 (7 days in MM), and by day 28 these were distributed over the whole scaffold visualized by the dense red color homogeneously covering the cross section and outer surface of the scaffolds (**Figure S7B**). Quantification of the staining (**Figure 6C**) confirmed that calcium depositions increased ~10 to 15-fold within these two time points. Moreover, statistical differences among the amount of calcium deposited by cells seeded by the MS-dextran and the MS-Ficoll-Pq methods were evident when quantified. No ARS staining was observed in CS scaffolds at day 14, and only a small amount of calcium deposits was quantified after 28 days culture, which was comparable to the mineralization in MS scaffolds at day 14.

Ultimately, the expression of relevant osteogenic genes on cells seeded with the MS-methods was evaluated and compared to the CS method. Notably, no significant differences were found in the expression of runt-related transcription factor (RUNX2), collagen type I (COL I), osteocalcin (OCN) and osteonectin (ON) among the different groups at day 14 (7 days in MM), suggesting that the seeding with macroMs did not alter the early stage of hMSCs differentiation (**Figure 7**). Importantly, the gene encoding RUNX2 was upregulated on cells seeded with the MS-Dextran and MS-Ficoll-Pq at day 28 (21 days in MM) when compared to CS method (**Figure 7A**). Similarly, COL I expression was higher in MS-Ficoll-Pq samples at this time point (**Figure 7B**). In addition, OCN was more expressed in MS-Dextran with respect to CS (**Figure 7C**). On the other hand, no significant differences were found in the ON encoding gene expression among time points or seeding methods (**Figure 7D**).

DISCUSSION

When cells are seeded on a 2D substrate, gravity acts as one of the main parameters favoring cell attachment. On the contrary, gravity acts as a counterforce to cell attachment during static seeding of 3D scaffolds fabricated from synthetic polymers and with large and interconnected pores, where cell attachment is already dramatically hindered because of

lack of adhesion motifs, fast flow velocity and large scaffold's pore size. Rather than modifying the scaffold material properties or architecture, in this study we tackled the cell sedimentation problem from a different perspective, the cell seeding media physical properties (viscosity and density). This is particularly a beneficial approach as it can be applied to all AM 3D scaffolds independent of their material chemistry or geometries. In this study we tuned the viscosity and density of the seeding media by using two natural biocompatible macroMs, dextran and Ficoll. Dextran has previously shown to promote ECM deposition in vitro not only when used as macromolecular crowder mimicking the

Figure 7. Gene expression of hMSCs after 14 and 28 days of culture (7 and 21 days in MM, respectively) comparing the three different seeding methods. (A) RUNX2, (B) COL I, (C) OCN and (D) ON fold change expression values are relative to CS d14 MM. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Sidak's multiple comparison test (n.s. $p > 0.05$; * $p < 0.05$; ** $p < 0.01$; * for comparisons among seeding methods each time point; \$ for comparisons among time points each seeding method).

highly crowded/dense native ECM, ⁴⁷⁻⁴⁹ but also when employed as media thickener to increase fluid shear forces in perfusion culture. ⁵⁰ It has also been used to study erythrocyte aggregation and blood viscosity changes. ⁵¹⁻⁵³ On the other hand, Ficoll has been used for the isolation of cells from blood samples by centrifugation, ⁵⁴⁻⁵⁵ as well as macromolecular crowding agent. ^{47-49, 56} Despite being widely used in the biomedical field, to the best of our knowledge, these macroMs were employed here for the first time as media supplements to enhance scaffolds static seeding. Moreover, we demonstrated that MS-Dextran and MS-Ficoll-Pq are two independent approaches, since viscosity and density are the effective parameters on cell settling velocity of the chosen dextran-based (10% Dextran) and Ficoll-Pq -based (60 vol% Ficoll-Pq) seeding solutions, and therefore the two methods can be studied and used separately. We showed that both increasing the seeding media viscosity or matching its density to that of hMSCs appropriately reduced cell settling velocity, allowing cells to float longer in the suspension, thereby giving them more time to attach and hence promoting homogeneous cell attachment on the PEOT/PBT scaffold filaments. On the other hand, cells poorly attached to the scaffold cross section when using the CS method. PEOT/PBT was chosen as model synthetic material to produce 3D AM scaffolds. These polyether-ester multiblock copolymers are thermoplastic elastomers, whose surface energies, swellability, degradation and mechanical properties can be tuned for specific applications by controlling the PEOT/PBT ratio ^{2, 57} and whose *in vitro* and *in vivo* applicability in the orthopedic field has been extensively studied. ⁵⁸⁻⁵⁹ Due to their easy processability, they have been previously used to produce 3D AM scaffolds. ⁶⁰ However, they offer poor cell attachment upon conventional static seeding as shown here and in previous works. ^{24, 61}

Direct comparison of our cell distribution results with previously published seeding techniques on 3D AM scaffolds is very challenging, as most of these studies have not shown the full characterization of cell distribution and sedimentation, but rather addressed the seeding efficiency and its direct effect on the tissue regeneration process. In one exception, the amount of DAPI nuclear staining was quantified in each of the slices subdividing the cross section of scaffolds with vertical pore size gradients. ³² Despite the positive effect of gradients on cell attachment in each of the scaffold subdivisions, still large amounts of cells accumulated at the bottom of the scaffolds, leading to a non-uniform cell distribution over the scaffold cross section. Interestingly, the uniform cell distribution demonstrated by MS methods closely resembles a recently published report in which hMSCs were seeded in a polymeric scaffold by means of a cell-laden polyethylene glycol

dithiothreitol hydrogel.⁴⁰ In this hybrid system, the cell carrier hydrogel was hydrolytically degraded after 7 days, time that cells used to populate the scaffold filaments attaining a good distribution. Importantly, when using our novel MS methods, cells attached directly to the filaments without the means of hydrogel carriers. Moreover, we proved that the macroMs did not act as a cell adhesive coating in the scaffolds, but viscosity or density of the media, depending on the macroM type, were the only factors leading to cell attachment. Previous reports demonstrating reduced protein adsorption and cell attachment in dextran coated surfaces support these results: dextran and Ficoll should be functionalized to render them cell adhesive.^{62, 63}

The fact that after the 4h seeding the macroMs were found to colocalize with the cell cytoplasm and that the amount of macroMs was drastically reduced after 24h culture in CM, suggests that the macroMs were internalized by cells during the seeding period, followed by their exocytosis towards the cell culture media after 24 h. Fluorescently labeled dextran and Ficoll have been shown to enter different cell types, including hMSCs, by macropinocytosis⁶⁴⁻⁶⁵ and micropinocytosis,⁶⁶ respectively. Within the first 24 h after uptake, they have been seen to accumulate in lysosomal compartments, where they degrade into glucose and fructose moieties.⁶⁶⁻⁶⁸ Moreover, since the cell culture media was renewed after 24h culture and every 2 days from then on, the effect of the macroMs as macromolecular crowders influencing ECM deposition can be ruled out. The absence of macroM in the culture might be beneficial since no effect or even detrimental influence of macromolecular crowding has been shown on cartilaginous tissue formation in 3D.⁶⁹ This was hypothesized to be mainly due to the fact that in a 3D cell confluent system cells already reside in the crowded microenvironment consisting of their secreted matrices. Moreover additional crowding could lead to inefficient oxygen and waste management and activation of degradation enzymes. However, there is lack of literature addressing the effect of macromolecular crowding in 3D cell systems and thus, no definitive conclusions can be drawn.

Regarding cell death, it is hypothesized that it occurred before their incubation with macroMs as i) it is known that internalized macroMs remain in the lysosomes with no effect on cellular function before their exocytosis to the extracellular space and ii) cell death was also visualized within the first minutes of hMSCs incubation even with medium without macroMs. Hence, the presence of dead cells in the scaffold might be attributed to the low media-to-cell ratio during the 4h seeding and to the handling of cells prior to seeding

(centrifugation and resuspension in a small volume of medium), which are unavoidable issues of the seeding process. In this regard, the visualization of macroMs in the scaffold even after 24h culture is attributed to the ability of macroM to penetrate the permeable membrane of dead cells during the seeding and the subsequent inability of these cells to process the macroMs.⁷⁰

Consistent with a previously published report, a cell monolayer was formed at the bottom of scaffolds seeded with CS method.²⁴ Notably, this was not observed in the case of MS scaffolds. It is plausible that the high cell density at the bottom of CS scaffolds enabled cell-to-cell and cell-to-scaffold contact after their fast sedimentation, while the low cell number localized at the bottom filaments region of MS-scaffolds, due to no cell sedimentation during MS methods and unattached cells washing after seeding, prevented the formation of a cell monolayer. Importantly, the sedimented cells which attached to the bottom of CS scaffolds, might have significantly contributed to the relatively higher seeding efficiency values than expected (~55%) for the high amount of cell settling observed, making it comparable to the MS methods seeding efficiency (~80% and ~75%, for MS-Dextran and MS-Ficoll-Pq, respectively). In 2D, high cell seeding density (cells*cm⁻²) has been demonstrated to direct and control early and late stage osteogenic markers and the temporal expression of MSCs genes towards osteogenesis.⁷¹ However, cell density coupled to an optimum distribution of cells occupying the whole scaffold surface area are the parameters that will ultimately favor osteogenesis in a 3D environment. Therefore, cell number alone is not a primary factor and “efficiency” is defined by both cell density and cell distribution when culturing 3D scaffolds. In our case, due to the higher chance of cell attachment when seeded with the MS methods, cells were confluent occupying the scaffold filaments from the beginning of the culture, resulting in surface saturation and contact inhibition between adjacent cells,⁷² which explains the lack of proliferation on these scaffolds over 28 days in BM.⁷³ The increased matrix production in the MS scaffolds’ pores volume when cultured in MM, might have offered the possibility to cells to grow not only on top of the scaffolds’ filaments, but also on the ECM-filled scaffolds’ pores. This increase in surface area available for cell proliferation between day 14 and day 28 was translated into an increase in cell number at late timepoints (day 28 in MM). Overall, cell confluency over the MS scaffolds, coupled to the osteogenic factors included in MM, showed to have positive effect on the osteogenic differentiation of the seeded hMSCs. This effect was initially demonstrated by the evaluation of ALP activity over the culture time, which is an early stage enzyme expressed in bone development and its upregulation

precedes the upregulation of late osteogenic genes, such as OCN, and matrix mineralization.⁷⁴ Decreasing ALP activity over time, which enables the development of the bone matrix by preparing the extracellular matrix for the deposition of mineral,⁷⁴ has previously been observed in scaffolds with high cell seeding numbers, and therefore possibly better cell distribution.⁷⁵ Concordantly, a peak of ALP was seen at day 14 in MM, with values decreasing at day 28 in MM in cells seeded with MS methods despite the macroM type. On the contrary, ALP activity peaked after 28 days of culture in MM on scaffolds seeded with hMSCs via the CS method, as previously reported.^{61, 76} These dissimilarities are likely due to the aforementioned differences in cell number coupled to their distribution profile within the scaffolds. Concordantly with the ALP activity progression, extensive ECM was produced over time on MS scaffolds and ARS staining results showed enhanced and progressively increased mineralization on MS scaffolds over the culture in MM. However, no clear hypothesis has been drawn to explain the increased calcium deposition on MS-Ficoll-Pq scaffolds at day 28 when compared to MS-Dextran, since cell viability was preserved in both cases and cell numbers and ALP activity values at the time point are comparable between both scaffold types.

Analyzing the gene expression, we observed that the upregulation of relevant protein encoding genes involved in osteogenic differentiation at day 28 in MM in cells seeded with MS-methods complements the abovementioned ALP activity and calcium deposition results. This further confirms that either there is a delay in the progression of osteogenesis in cells seeded with the CS method, or an acceleration in cells seeded with the MS methods, likely due to the differences in homogeneity in cell distribution. In particular, the gene encoding RUNX2, which is an essential transcription factor regulator of hMSCs differentiation into the osteogenic lineage,⁷⁷⁻⁷⁸ was upregulated for cells on MS scaffolds. It is known to target specific osteogenic genes, such as COL I, one of the major bone ECM components, or OCN, one of the major bone non-collagenous proteins with the ability to bind bone hydroxyapatite.⁷⁹ Accordingly, COL I and OCN were upregulated in MS-Ficoll-Pq and MS-dextran scaffolds, respectively, when compared to CS scaffolds, suggesting matrix maturation. Although ON did not differentially express in the MS scaffolds, this calcium binding non-collagenous protein has been found to be expressed in other non-mineralizing tissues, and therefore, lacks specificity.⁸⁰ While longer culture periods of CS scaffolds in MM might possibly lead to comparable results among different seeding methods,⁸¹ the possibility of speeding the osteogenic process of hMSCs by MS-methods is desired for both the *in vitro* and *in vivo* applications of the scaffolds.

CONCLUSION

Cell seeding on 3D AM scaffolds made of synthetic polymers with interconnected macroporosity still remains a challenge. The aim of this study was to develop a simple and reliable seeding technique to improve cell distribution upon static seeding valid for all 3D AM scaffolds regardless of their material properties and architecture. This was achieved by controlling the settling velocity of cells by altering the viscosity and density of the seeding media via the addition of two different macroMs, Dextran and Ficoll (Ficoll-Pq). Compared to scaffolds seeded with conventional static techniques, a homogeneous cell distribution was attained with direct cell attachment to the scaffold filaments. Importantly, macroMs did not affect viability or osteogenic differentiation of hMSCs and their complete removal was demonstrated after the seeding period. Intriguingly, the improved cell distribution from the macroMs based seeding led to enhanced mineralization and bone matrix maturation compared to conventional seeding. This newly proposed seeding method is simple, reproducible, and has the potential to improve the long-term functionality of *in vitro* and *in vivo* cell-seeded constructs for tissue engineering applications.

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SUPPLEMENTARY FIGURES

Table S1. Primer sequences used for q-PCR. B2M: beta-2-microglobulin; RUNX2: runt-related transcription factor; COL1A1: collagen 1 alpha 1; OCN: osteocalcin; SPARC: osteonectin.

Gene	Forward primer	Reverse primer
B2M	ACAAAGTCACATGGTTCACA	GACTTGTCTTTCAGCAAGGA
RUNX2	TGGTTACTGTCATGGCGGGTA	TCTCAGATCGTTGAACCTTGCTA
COL1A1	GAGGGCCAAGACGAAGACATC	CAGATCACGTCATCGACAAC
OCN	TGAGAGCCCTCACACTCCTC	CGCCTGGGTCTTCTACTAC
SPARC	AGCACCCCATTGACGGGTA	GGTCACAGGTCTCGAAAAAGC

Figure S1. (A, B) Relative viscosity and (C, D) density of cell culture media containing different concentrations of dextran and Ficoll. (E) Comparison of density values of media containing specific dextran and Ficoll concentrations. (F) Cell settling velocity in CS, MS-Dextran and MS-Ficoll methods.

Figure S2. (A) Quantification of cell number in each layer of the Ficoll gradient used to determine hMSCs population density by separation by centrifugation or by separation by buoyancy forces. (B) Representative images of the cells found in each layer after each separation method. Scale bars 200 μ m.

Figure S3. Fluorescence microscopy images of hMSCs in the cross sections and bottom lids of scaffolds pre-incubated with dextran and Ficoll based solutions and seeded with the CS method. Scale bar 1 mm.

Figure S4. Representative confocal microscopy images of hMSCs (F-actin and nuclei), macroMs (FITC-labeled) and dead cells on top of scaffold fiber 24h post-seeding with the MS-Dextran and MS-Ficoll methods. Scale bars 100 μ m.

Figure S5. Representative fluorescent images depicting hMSCs viability (live, green; dead, red) right after their resuspension (0 h) and after 4h incubation in macroM based solutions, as compared to control (CS). Scale bars 250 μ m.

Figure S6. Cell culture and differentiation in 2D after hMSCs incubation in macroM based solutions during 4h mimicking the scaffold seeding conditions. (A) Fold increase in cell number after 7 days of culture in BM and pre-incubation in normal, dextran or Ficoll based CM. (B) Representative brightfield images of cells pre-incubated in normal, dextran or Ficoll based CM after 1 and 7 days of culture in BM. (C) Stereomicroscopy images of cell monolayers stained with ARS after 28 days of culture (21 days in MM) and pre-incubated in normal, dextran or Ficoll based CM. Scale bars 250 μ m.

Figure S7. Stereomicroscopy images of ARS stained scaffolds. (A) Cross- and outer sections after 28 days of culture in BM and (B) outer sections after 28 days of culture (21 days in MM). Scale bars 1 mm.

Chapter 4

Tuning cell behavior on 3D scaffolds fabricated by atmospheric plasma assisted additive manufacturing

Tuning cell behavior on 3D scaffolds fabricated by atmospheric plasma assisted additive manufacturing

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Abstract

3D scaffolds with optimum physico-chemical properties are able to elicit specific cellular behaviors and guide tissue formation. However, cell-material interactions are limited in scaffolds fabricated by melt extrusion additive manufacturing (ME-AM) of synthetic polymers, and plasma treatment can be used to render the surface of the scaffolds more cell-adhesive. In this study, a hybrid AM technology, which combines a ME-AM technique with an atmospheric pressure plasma jet, was employed to fabricate and plasma treat scaffolds in a single process. The organosilane monomer (3-Aminopropyl) trimethoxysilane (APTMS) and a mixture of maleic anhydride and vinyltrimethoxysilane (MA-VTMOS) were used for the first time to plasma treat 3D scaffolds. APTMS treatment deposited plasma polymerized films containing positively charged amine functional groups, while MA-VTMOS introduced negatively charged carboxyl groups on the 3D scaffolds' surface. Argon plasma activation was used as control. All plasma treatments increased the surface wettability and protein adsorption to the surface of the scaffolds, and improved cell distribution and proliferation. Notably, APTMS treated scaffolds also allowed cell attachment by electrostatic interactions in the absence of serum. Interestingly, cell attachment and proliferation were not significantly affected by plasma treatment induced aging. Also, while no significant differences were observed between plasma treatments in terms of gene expression, human mesenchymal stromal cells (hMSCs) could undergo osteogenic differentiation on aged scaffolds. This is probably because osteogenic differentiation is rather dependent on initial cell confluency and surface chemistry might play a secondary role.

INTRODUCTION

Over the past decades, additive manufacturing (AM), and in particular melt extrusion AM (ME-AM), has emerged as an advanced fabrication technique for the development of scaffolds for skeletal tissue engineering. This is due to the possibility of producing cost-effective, customizable, biocompatible and biodegradable 3D constructs, with interconnected macro-pores facilitating tissue ingrowth, while maintaining sufficient mechanical properties for load bearing applications.¹ In order to ensure tissue formation and to obtain reliable readouts when evaluating a scaffold functionality *in vitro*, an efficient cell attachment upon seeding, in terms of cell density and distribution, is particularly important.²⁻⁴ Moreover, cell colonization also plays a pivotal role to guarantee the *in vivo* success of both cell-laden and cell-free scaffolds. Despite their advantageous structural features, cell adhesion has shown to be challenging on ME-AM scaffolds, due to their large pores and lack of biological recognition sites to enable cell-material interactions, as they are mostly made of synthetic thermoplastic polymers, such as poly(ϵ -caprolactone) (PCL), poly(lactic acid) (PLA), poly(lactic-co-glycolic acid) (PLGA), poly(urethane) (TPU), or poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT). These factors, along with the gravity force, contribute to fast cell sedimentation towards the scaffold's bottom and mediocre attachment to the scaffold's filaments upon conventional static seeding, which is a simple and widely used seeding method that consists on placing a droplet of cell suspension on top of the scaffold surface, that gradually flows into the scaffold's pores.⁵⁻⁷

Different strategies have been considered to optimize cell attachment during static seeding of 3D ME-AM. For instance, loading cells into a natural or synthetic hydrogel solution, which crosslinks within the scaffolds pores, has shown to improve cell retention within the scaffold.⁸⁻⁹ Alternatively, we have recently proposed the use of macromolecules as temporary seeding media supplements to increase its viscosity or density, which has proven to reduce cell sedimentation velocity and result in homogeneous cell attachment along the scaffold's cross section.⁴ Besides the modification of the seeding solution, the alteration of the scaffold architecture, with regard to pore size, lay-down pattern, or the introduction of vertical gradients, has also been suggested to enhance cell attachment.^{6, 10} Furthermore, synthetic polymers have been blended with bioactive fillers, such as calcium phosphates,¹¹ bioactive glass¹² or graphene derivatives,¹³ to improve cell

attachment and cell-material interactions by increasing scaffolds' hydrophilicity and /or surface roughness.

Surface coatings have also been considered as an alternative to increase polymeric scaffolds' bioactivity, as some of the methods described above involve the modification of scaffold geometry or the material bulk properties and these can negatively affect the scaffolds mechanical properties. For example, NaOH etching or other wet chemical surface modifications can increase the hydrophilicity and surface roughness, or deposit specific monomers with functional groups on 3D scaffolds.¹⁴ Yet, the use of harsh chemicals can potentially compromise the integrity of the scaffolds' structure.¹⁵ Other examples exploit scaffold's filaments coatings with extracellular matrix (ECM) proteins or cell adhesive peptides to improve cell adhesion.¹⁶ Nevertheless, this approach has not been extensively explored due to proteins' limited half-lives and high costs.

Non-thermal plasma (NTP) surface modification has the advantage of altering the polymer surface chemistry and/or topography without the use of solvents, while maintaining their bulk properties.¹⁷ Various types of gases, such as oxygen, helium, argon, nitrogen, or their combination can be used to generate the plasma discharge and activate polymer surfaces by the incorporation of hydroxyl, carboxyl, aldehyde or amine groups. To obtain more stable coatings with a higher density of functional groups, the plasma discharge can be fed with monomers, such as alkyl amine or ammonia, or acrylic acid, to coat the surface with a thin polymeric film containing amine or carboxyl groups, respectively, in a process known as plasma polymerization. NTP is an established technique to treat 2D polymeric surfaces, where extensive research has revealed enhanced protein adsorption and cell attachment due to the presence of functional groups compared to methyl only containing surfaces.¹⁸⁻¹⁹ Moreover, carboxyl modified surfaces have shown to promote chondrogenesis, while amine groups have proved to direct cells towards the osteogenic pathway.²⁰⁻²²

Being an established technique to treat two-dimensional (2D) polymeric surfaces, NTP was used during the last decade to functionalize 3D ME-AM scaffolds as well. Oxygen plasma activation, as one of the primary focuses of reported studies, has shown to increase surface roughness, protein adsorption and cell attachment, as well as to promote alkaline phosphatase (ALP) expression and matrix mineralization on PCL scaffolds.²³⁻²⁴ Other studies have explored plasma polymerization techniques. Among those, nitrogen containing groups have been introduced on PCL or polystyrene scaffolds by allyl amine,

ammonia or ethylene/N₂, promoting Runt-related transcription factor 2 (RUNX2) and ALP upregulation in human mesenchymal stromal cells (hMSCs) compared to untreated scaffolds.²⁵⁻²⁶ Alternatively, in a study using PEOT/PBT scaffolds treated with acrylic acid plasma polymerization, enhanced proliferation and glycosaminoglycan production by chondrocytes was shown.²⁷ Despite promising results in enhancing ME-AM scaffolds surface bioactivity, NTP is still not a widely used technique for 3D scaffolds treatment. This is mainly due to the NTP process usually being carried out at sub-atmospheric pressures, where plasma has shown to not ignite inside the scaffold pores.²⁸⁻²⁹ This limits the treatment in the scaffold's core to diffusion, leading to inhomogeneous and uncontrolled gradient functionalization. Moreover, the process often requires costly vacuum equipment and its application demands multi-steps processes. Overcoming these limitations, atmospheric pressure plasma jets (APPJ) have been developed,³⁰ where the need of a reactor chamber and vacuum equipment is eliminated and the NTP exiting the jet can be directed into the scaffolds pores, or a region of interest. Moreover, APPJ enables its assembly to an AM platform, where the printing and plasma treatment processes can take place in a single hybrid system. In addition, scaffolds can be either plasma treated as a whole from the top, or in a layer-by-layer manner to reach deep pores in large anatomical scaffolds, or treat specific scaffolds' zones.³¹⁻³²

Taking these advantages, here we used a hybrid AM technique, consisting of a ME printhead and an APPJ module assembled on a 3-axis platform to fabricate plasma treated PEOT/PBT scaffolds in a single process.³² Despite its excellent processability and previous *in vitro* and *in vivo* applications for bone tissue engineering, PEOT/PBT demonstrates poor cell attachment upon cell seeding.^{4, 7} The organosilane monomers (3-Aminopropyl) trimethoxysilane (APTMS) and maleic anhydride-Vinyltrimethoxysilane (MA-VTMOS) were used to deposit polymer-like thin films containing amine and carboxyl functional groups, respectively, by plasma polymerization in argon on the PEOT/PBT scaffolds surface. Despite being already explored to modify 2D surfaces,³³⁻³⁴ these monomers were used here for the first time to treat 3D scaffolds. Compared to other precursors, organosilanes possess the advantage of providing a stable siloxane backbone to the coating, which is highly adherent and resistant to delamination in water conditions, ensuring good functional group retention under cell culture conditions.³⁵⁻³⁶ Pristine scaffolds, plasma polymerized scaffolds, and argon plasma activated scaffolds were seeded with hMSCs. Cell attachment and adhesion mechanism were evaluated.

Moreover, proliferation and osteogenic differentiation were assessed and compared among the different plasma conditions.

MATERIALS AND METHODS

Plasma treated scaffolds fabrication and characterization

Scaffolds were fabricated and plasma treated with a hybrid AM platform, consisting of a 3D axis stage (Bioscaffolder 3.0, Gesim) equipped with a custom made ME printhead and an APPJ (Plasma Stylus Noble, Nadir srl),³⁷ all enclosed in a polymethylmethacrylate box with fume extraction ventilation. In a first step, scaffolds were produced (**Figure 1A**). For that, PEOT/PBT pellets (300PEOT55PBT45, PEO molecular weight = 300 kDa, PEOT:PBT weight ratio = 55:45, intrinsic viscosity 0.51 dl/g, Polyvation, The Netherlands) were heated up to 195 °C in the cartridge of the printhead. Molten polymer was extruded through a 250 µm internal diameter needle (pressure 7 bar, translation speed of 16 mm/s), depositing a 0-90 pattern block (15x15x4 mm³) with 200 µm layer thickness and 750 µm strand distance (center to center) (**Figure 1A**).

Figure 1. Hybrid AM platform consisting of (A) a ME printhead and (B) an atmospheric pressure plasma jet (APPJ), mounted on a 3D axis stage.¹⁻² (C) Schematics of the APPJ, which is based on a dielectric barrier discharge scheme simultaneously powered by a high-voltage (HV) generator and a radio-frequency (RF) generator. Electrodes are positioned externally to an alumina duct where argon is fluxed and plasma is ignited. Previously mixed precursors in vapor phase are carried by argon and mixed with the ignited plasma right before the RF generator through an inner coaxial tube, which is connected to external bubblers with precursors in liquid (APTMS, VTMS) or solid (MA) phase. Nitrogen is carried through an outer coaxial duct for shielding and cooling at the exit of the torch.

Right after fabrication, scaffolds were plasma treated with the APPJ (**Figure 1B**). The device is based on a dielectric barrier discharge scheme simultaneously powered by a high-voltage generator (HV, 10 kV, 17 kHz) and a radio-frequency generator (RF, 27 MHz) (**Figure 1C**). Electrodes are positioned externally to an alumina duct where argon is fluxed (Ar 5.0 purity, 10 L/min) and plasma is ignited. An inner coaxial tube allows for the introduction of the precursors in vapor phase just before the RF electrodes. For APTMS plasma polymerization, APTMS (Sigma-Aldrich) was carried into the plasma zone by argon gas passing through a bubbler at room temperature (RT) (flow = 2 L/min). Similarly, for MA-VTMOS plasma polymerization, MA (Sigma-Aldrich) and VTMOs (Sigma-Aldrich) in two independent bubblers were carried into the plasma zone by argon at 1.764 L/min and 0.233 L/min, respectively. For argon activation, the RF generator was operated with a power output of 15 W in continuous mode. For APTMS and MA-VTMOS plasma polymerization, the RF generator was operated in pulsed mode, at a duty cycle of 5% (250 μ s ON, 5000 μ s OFF), with the power output for the ON duration set to 15W. An outer gas shell of nitrogen (15 L/min) was used during the process to prevent precursor oxidation by the environmental air during the deposition. The APPJ nozzle was positioned at 1 mm above the scaffold surface and moved in XY following the surface filaments' path at 1 mm/s. For further studies, plasma treated scaffolds were used within 2 days after plasma treatment ('fresh' samples) or after ~10 days being stored at RT in a sealed container ('aged' samples).

To assess the successful deposition of an APTMS plasma polymerized layer, fresh, aged and ethanol disinfected scaffolds, were incubated with the amine reactive fluorescent dye LIVE/DEAD™ Fixable dead Cell Stain Kit (Thermo Fisher Scientific) for 45 min at a concentration of 0.5 μ l / 500 μ l PBS. Untreated, argon, and MA-VTMOS treated scaffolds were also stained as controls. After PBS washes, scaffolds were cut and the cross sections were imaged using a fluorescent microscope (Eclipse, Ti2-e, NIKON). In order to confirm the deposition of a MA-VTMOS layer, fresh, aged and ethanol disinfected scaffolds were incubated for 20 s in a methylene blue solution (1 mg/ml). Untreated, argon and APTMS treated scaffolds were also stained as controls. After PBS washes, scaffolds were cut and imaged with a stereomicroscope (Nikon SMZ25). In order to evaluate the dynamic wettability of the scaffolds, a 35 μ l water droplet was carefully deposited onto the surface of the scaffolds, and the wetting behavior was captured with a digital camera at 1 frame per second (Krüss DSA25S).

Plasma treated 2D films fabrication and characterization

2D films were prepared from PEOT/PBT pellets. Briefly, 60 milligrams of pellets were molten at 190 °C and pressed with a coverslip against a Teflon sheet to obtain films with 14 mm diameter and ~300 µm thickness. Subsequently, PEOT/PBT films were plasma treated with argon, APTMS or MA-VTMOs using the APPJ, according to the parameters in the section “Plasma treated scaffolds fabrication and characterization”.

Static contact angle was measured in fresh and aged films using the sessile drop method. For that, a 4 µl water droplet was placed on top of the substrates by an automatic syringe dispenser (Krüss DSA25S). 20 seconds after droplet formation, the contact angle was calculated automatically by the device’s software using the Laplace–Young curve fitting.

Attenuated total reflectance-Fourier-transform-infrared spectroscopy (ATR-FTIR, Nicolet iS50, diamond ATR) was performed on untreated and fresh plasma treated films in order to further confirm the presence of the coating.

Zeta potential measurements were performed on fresh films with a size of 2x4 cm². Electrokinetic measurements were done at RT using a SurPASS™ system (Anton Paar GmbH, Germany), with a KCl solution (0.01M, pH 5-5.5) as electrolyte. The zeta potential was calculated from ten experimental points taken by the measurement of the streaming potential in a pressure range of 500 mbar (start) to 200 mbar (end) in the flow cell.

Cell seeding on 3D scaffolds

HMSCs isolated from bone marrow were purchased from Texas A&M Health Science Center, College of Medicine, Institute for Regenerative Medicine. Cryopreserved vials were plated at a density of 1,000 cells·cm⁻² in tissue culture flasks and expanded at 37 °C / 5% CO₂ in cell culture media (CM) consisting of αMEM with Glutamax and no nucleosides (Gibco) supplemented with 10% fetal bovine serum (FBS).

To investigate the cell attachment mechanism, fresh scaffolds were disinfected in 70% ethanol for 20 min and washed 3 times with Dulbecco’s PBS (5 min each). Scaffolds were incubated overnight in CM with FBS ((+)FBS) or CM without FBS ((-)FBS). Before seeding, scaffolds were dried on top of a sterile filter paper and placed on non-treated well plates. HMSCs were trypsinized and resuspended in (+)FBS or (-)FBS CM. A droplet of cell suspension (37 µl containing 200,000 cells) was placed on top of each scaffold, filling the pores within some seconds. Seeded scaffolds were incubated for 4 hours at 37 °C / 5%

CO₂ to allow for cell attachment. After this time, scaffolds were collected or transferred to new wells containing 1.5 ml of (+)FBS and cultured overnight before sample collection. Aged scaffolds incubated overnight in (+)FBS CM and seeded on (+)FBS CM were also cultured overnight before sample collection, to analyze the effect of aging on cell adhesion.

To evaluate the effect of cell seeding density on scaffold coverage and proliferation, fresh and aged scaffolds were incubated overnight in CM and seeded with hMSCs in CM at a density of 200,000 (200k) or 400,000 (400k) cells/scaffold, concentrated in a 37 μ l droplet. After 4h attachment, 200k and 400k scaffolds were transferred to new wells containing 1.5 or 3 ml BM (CM supplemented with 200 μ M L-Ascorbic acid 2-phosphate), respectively, and cultured for 7 days.

To evaluate hMSCs osteogenic differentiation, aged plasma treated scaffolds were incubated overnight in (+)FBS. Scaffolds seeded with 200,000 cells were further cultured for 7 days in BM and for another 47 days in mineralization media (MM) consisting of BM supplemented with dexamethasone (10 nM) (Sigma-Aldrich) and β -glycerophosphate (10 mM) (Sigma-Aldrich). Media was replaced every two or three days. As osteogenic differentiation 2D controls, cells were seeded in tissue culture polystyrene wellplates at a density of 5,000 cells·cm⁻², and cultured in the same media conditions as 3D scaffolds.

Imaging of cell attachment within scaffold cross sections

Scaffolds were fixed with 4 wt% paraformaldehyde for 30 min, permeabilized using 0.1 vol% Triton-X for 30 min, and incubated with phalloidin (Alexa Fluor 488, 1:75 dilution in PBS) for 1h at RT. The bottom and cross section of scaffolds were imaged using a fluorescent microscope. Background subtraction and contrast enhancement were performed on the images using the software Image J, in order to clarify their visualization. To assess cell coverage, cross section images were converted to binary (rendering regions with cells in white and the rest in black), and the total amount of white pixels in the scaffold area was quantified and normalized to the total number of pixels.

DNA quantification

Scaffolds collected at the desired timepoints were freeze-thawed 3x for cell lysis and incubated overnight at 56 °C in Proteinase K solution (1mg/ml Proteinase K (Sigma-Aldrich) in Tris/EDTA buffer) (1:1) for matrix degradation and cell lysis. Then, scaffolds were freeze-thawed 3x and incubated 1h at RT with a 20X diluted lysis buffer from the

CyQUANT cell proliferation assay kit (Thermo Fisher Scientific) containing RNase A (1:500), to degrade cellular RNA. Lastly, samples were incubated with the fluorescent dye provided by the kit (1:1) for 15 min and fluorescence was measured with a spectrophotometer (emission/excitation = 520/480 nm).

Protein adsorption quantification

Disinfected fresh and aged scaffolds were incubated overnight at 37 °C in (+)FBS or (-)FBS. After washing with PBS, scaffolds were blocked for 1 h at 37 °C in 1 wt/v % BSA/PBS and then incubated with specific bovine vitronectin (ab23444, Abcam) or fibronectin (ab2413, Abcam) primary antibodies, diluted 1:500 and 1:400, respectively, in 1 wt/v % BSA/PBS. After washing three times in PBS, scaffolds were incubated for 1h at 37 °C in horseradish peroxidase (HRP)-conjugated anti-mouse secondary antibody (abcam, 1:100000 dilution 1% BSA) for vitronectin detection, or HRP-conjugated anti-rabbit secondary antibody (abcam, 1:10000 dilution in 1% BSA) for fibronectin detection. After three washes in 1% BSA+ 0.05% tween 20, scaffolds were blotted in an adsorbent paper and incubated in 150 µl of 1-Step Ultra TMB ELISA substrate (Thermofisher Scientific). Color was allowed to develop for 10 min at RT and the reaction stopped with 50 µl of sulfuric acid (2M). Supernatants absorbance was measured at 450 nm with a spectrophotometer (CLARIOstar®, BMG Labtech). Scaffolds incubated in (-)FBS and scaffolds incubated in (+)FBS and reacted with only secondary antibodies were used as controls.

In order to analyze albumin adsorption, disinfected fresh and aged scaffolds were incubated overnight at 37 °C with 1mg/ml BSA-FITC (Sigma-Aldrich) solution. After washing three times with PBS, scaffolds were blotted in an adsorbent paper and incubated for 2h at RT in a 1% SDS solution. Supernatants fluorescence was measured at excitation/emission = 495 nm/519 nm.

Alizarin red S (ARS) staining

Scaffolds were collected at day 35 and 54 of culture, fixed, washed with distilled water, cut in half and stained with ARS (60 mM, pH 4.1 - 4.3) for 20 min at RT. Samples were thoroughly washed to remove staining residues. Scaffolds' cross sections were imaged using a stereomicroscope to visualize calcium deposition.

Immunostaining

After fixation, scaffolds were permeabilized for 30 min by incubating on Triton-X 100 (0.1 v%). Subsequently, scaffolds were blocked by 1h incubation in blocking buffer (BB, 3 % BSA + 0.01% Triton-X 100), cut in half, and incubated overnight at 4 °C with a primary antibody: either collagen I (COL I) rabbit polyclonal (ab34710, Abcam), or vinculin mouse monoclonal (sc73264, Santa Cruz Biotechnology), both diluted 1:200 in washing buffer (10x diluted blocking buffer). Washed samples were incubated for 1h at RT with the secondary antibody (1:200 Alexa Fluor 568 goat derived anti rabbit antibody, Thermo Fisher Scientific). Then, scaffolds were washed and stained for F-actin (488 Alexa Fluor Phalloidin, Thermo Fisher Scientific) for 1h at RT and imaged with a confocal laser scanning microscope (Leica TCS SP8 STED), equipped with a white light laser (WLL). Emission was detected with HyD detectors.

Gene expression

Gene expression was analyzed with qRT-PCR at 14 and 54 days of culture. RNA was extracted from cells by incubating the scaffolds in Trizol. Subsequently, samples were centrifuged at 12,000 rcf for 5 min to precipitate the scaffold and ECM at the bottom. In a last step, chloroform was added to the supernatant and centrifuged at 12,000 rcf for 5 min to isolate the RNA, present in the aqueous phase. RNA was further purified using RNeasy mini kit column (Qiagen), according to the manufacturer's protocol and quantified using a spectrophotometer. Reverse transcription was performed using iScript™ (Bio-Rad) following suppliers' protocol. qPCR was performed on the mix composed of cDNA, SYBRGreen master mix (Qiagen) and the selected primers (**Table S1**) using a CFX Connect™ Real-Time System (Bio-Rad) under the following conditions: cDNA denaturation for 3 min at 95 °C, 40 cycles of 15 s at 95 °C, and 30 s at 65 °C. Gene transcription was normalized to the transcription of the housekeeping gene B2M. The $2^{-\Delta\Delta Ct}$ method was used to calculate relative gene expression for each target gene.

Statistical analysis

All data is shown as average with error bars indicating the standard deviation of at least three replicates. Analysis of statistics was conducted with GraphPad Prism (version 8.0.1). A one-way or two-way analysis of variance (ANOVA) was performed followed by a Tukey's

post-hoc multiple comparison test to evaluate statistical significance and correct for multiple comparisons.

RESULTS AND DISCUSSION

Compared to conventional manufacturing techniques, the possibility of fabricating customizable scaffolds for bone tissue engineering in a reproducible manner by AM brings the produced constructs one step closer to clinical translation. The assembly of a surface modification module, such as an APPJ, into the printing platform further increases the value of the technology and fabricated scaffolds for their research and clinical applications. With such a hybrid AM platform developed within our groups,³² we were able to manufacture surface modified 3D PEOT/PBT scaffolds in a continuous process, where scaffolds are fabricated by ME-AM and subsequently functionalized by the APPJ module (**Figure 2**). To do this, the nozzle tip of the APPJ module was adjusted at 1 mm over the scaffold surface, ensuring the penetration of the plasma flame over the whole scaffold depth (4 mm), while moved in XY over the scaffolds filaments at the transitional speed of 1mm/s, covering the whole scaffold block area (15x15 mm).

Figure 2. Schematic representation of the hybrid platform for scaffolds fabrication combined with plasma functionalization. 3D scaffolds are fabricated via ME-AM and consequently plasma treated from the top using an APPJ, which moves in XY following the scaffolds filaments, while the plasma penetrates the scaffold depth. As a result, (a) untreated, (b) argon activated or (c) MA-VTMS and (d) APTMS plasma polymerized scaffolds can be obtained, each with a specific surface functionality.

Plasma scaffolds characterization

Plasma polymerization of APTMS and MA-VTMOS on the surface of the scaffolds was carried out in a pulsing mode with 5% duty cycle, since compared to continuous plasma, pulsed plasma polymerization at low duty cycles has shown to allow for better preservation of the monomers' functional groups integrity during radical formation, as well as for the deposition of smoother surfaces.^{33, 36, 38} A homogeneous negatively charged coating covering the filaments all along the scaffolds volume after MA-VTMOS plasma polymerization was confirmed by staining with the cationic dye methylene blue, as previously described (**Figure 3A, Figure S1**), as well as with zeta potential measurements (**Figure S2**).³⁹ FTIR analysis further identified the negatively charged functional groups as intact anhydride groups from the MA precursor (-C=O stretching at $1750\text{-}1800\text{ cm}^{-1}$), and carboxyl groups (-OH stretching at $3200\text{-}3650\text{ cm}^{-1}$) (**Figure 3B, Figure S3A**).^{38, 40} In addition, vinyl groups from the VTMOs molecules (C=C stretching at $1630\text{-}1660\text{ cm}^{-1}$) were also identified. To visualize the formation of a homogeneous positively charged (**Figure S2**) thin polymer-like film containing amine groups (-NH_2) after APTMS plasma functionalization, an amine-reactive fluorescent dye was used to stain the scaffolds (**Figure 3A, Figure S1**). Primary amines vibrations were also observed in FTIR analysis (N-H bending at $1560\text{-}1630\text{ cm}^{-1}$ and N-H stretching at $3000\text{-}3500\text{ cm}^{-1}$) (**Figure 3C, Figure S3B**). Stretching vibrations visible at $2980\text{-}2880\text{ cm}^{-1}$, confirmed the retention of some aminopropyl chains of the precursor.⁴¹ Nitrogen-containing groups were also visible as amides (-C=O stretching at $1630\text{-}1695\text{ cm}^{-1}$) and as oximes (C=N stretching at $1650\text{-}1680\text{ cm}^{-1}$ and O-H stretching at $3550\text{-}3600\text{ cm}^{-1}$).⁴²⁻⁴³ The presence of oximes and amides was probably due to precursor oxidation during the deposition process, despite the nitrogen being flushed in the outer duct of the APPJ, probably due to the jet movement and the flow being disrupted by the scaffold structure.⁴¹ In addition, the APTMS and MA-VTMOS plasma polymerized substrates spectra were dominated by absorption bands between 1000 and 1200 cm^{-1} , confirming the presence of a siloxane network. In contrast to plasma polymerization, argon plasma activation is a less specific treatment, where a variety of polar groups can be introduced. According to previous reports on argon plasma on PEOT/PBT substrates, it is hypothesized that hydroperoxides (-OOH) and peroxide (-OO-) groups were incorporated, formed after post oxidation in air of the free radicals generated by plasma, along with aldehyde or ketone groups (-CHO , -CO-).⁴⁴⁻⁴⁶ The presence of these negatively charged groups exposed in the surface of the scaffolds was visualized by a low intensity methylene blue staining and confirmed by zeta potential

measurements (**Figure 3A, Figure S2**). Yet, these changes were not detected by FTIR analysis, possibly due to the very subtle variation in surface chemistry (**Figure 3D, Figure S3C**). Moreover, it is plausible that argon plasma activation could have induced changes in the surface roughness of PEOT/PBT, as previously described.⁴⁶ This effect has been attributed to plasma UV irradiation etching, mainly on the PEO amorphous regions of the copolymer, which are more mobile and susceptible to plasma than the PBT regions. However, SEM image analysis did not show any changes in the microscale surface roughness of argon scaffolds, compared to untreated ones (**Figure S4**). Similarly, no changes in microroughness were noted on the plasma polymerized scaffolds (**Figure S4**). This might be due to the low contact time of the plasma jet with the polymer, compared to the long exposure-driven changes reported in literature (from 5 to 30 min),⁴⁴⁻⁴⁵ modifying only the roughness in the nanoscale, which was not appreciable with SEM.

Figure 3. Characterization of plasma functionalized 3D scaffolds and comparison among fresh and aged conditions. (A) Verification of the treated scaffolds surface functionalization by specific staining. MA-VTMOS and Argon scaffolds were stained with the cationic dye methylene blue (blue staining in light microscopy) and APTMS scaffolds with an amine specific dye (red staining in fluorescence microscopy). Inserts represent stained control untreated PEOT/PBT scaffolds (white in light microscopy and grey in fluorescent microscopy). Scale bars 1 mm. (B, C, D) FTIR spectra of MA-VTMOS, APTMS and Argon plasma treated 2D substrates (melt-pressed PEOT/PBT films).

To evaluate the effect of plasma treatment on the polymer wettability, water contact angle (WCA) measurements were performed on melt pressed PEOT/PBT films, where enhanced hydrophilicity was observed due the introduction of polar groups by all the plasma conditions (**Figure 4A**). WCA decreased from $\sim 80^\circ$ to $\sim 55^\circ$ in argon and APTMS treated scaffolds, and down to $\sim 35^\circ$ in MA-VTMOs scaffolds. Interestingly, similar values were previously reported for self-assembled monolayers coated with these specific functional groups.⁴⁷ In accordance to the WCA, the dynamic wettability of MA-VTMOs and argon treated scaffolds increased compared to untreated scaffolds. This was observed by a significant reduction in the time that a droplet of water placed on top of the scaffold took to fill the scaffolds pores (from 6 min on untreated scaffolds to 2 sec on treated scaffolds), mimicking the static seeding process (**Figure 4B**). Surprisingly, despite the low contact angle of the APTMS coatings, the wettability of these scaffolds decreased compared to untreated scaffolds, potentially due to the lower density of functional groups and higher density of hydrophobic silane/siloxane in the scaffolds compared to the 2D films, where the WCA was measured. Yet, this did not affect cell culture experiments, since scaffolds were always pre-wetted before cell seeding.

Figure 4. Hydrophilicity and wettability of plasma functionalized 3D scaffolds and comparison among fresh and aged conditions. A) Static contact angle measured on untreated and plasma treated 2D substrates. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (* $p < 0.05$, *** $p < 0.001$, **** $p < 0.0001$). (B) Dynamic wettability of plasma treated scaffolds. Scale bar 1 mm.

The effect of storage (~ 10 days at RT in a sealed container) on the preservation of the plasma activation or polymerized treatments was also evaluated. This is a relevant shelf life information due to the potential time-lapse between scaffold production and its use, both for *in vitro* and *in vivo* applications. It is well known that plasma treated polymeric

surfaces can undergo ageing over time, which accounts for surface restructuring and hydrophobicity recovery towards a more energetically stable state.⁴⁸ While bulk analysis in terms of staining intensity, contact angle values and wettability suggested stability of the coatings (**Figure 3A, Figure 4A-B**), FTIR analysis revealed slightly lower functional groups absorbance peaks on aged substrates compared to fresh ones (**Figure 3B-D**), as previously reported for such plasma treatments on 2D substrates.^{40, 49} Moreover, it is worth noticing that the carbonyl stretching absorbance in MA-VTMOS spectra (-C=O stretching at $1700\text{--}1725\text{ cm}^{-1}$) slightly increased in aged samples, which can be explained by the formation of carboxylic groups by the hydrolysis of the anhydrides through humidity.⁴⁰ In general, these results suggest small changes in surface chemistry given by a reduction in the functional groups density on aged plasma polymerized substrates, which could potentially influence interactions such as protein adsorption or cell attachment. It is likely that the semi-crystalline property of PEOT/PBT and PBT the particularly efficient but cold and homogeneous APPJ configuration might have contributed to the limited aging effect, since it has previously been shown that the hydrophobic recovery decreases with increasing polymer crystallinity and with decreasing the degree of cross-linking due to plasma jet surface interaction.⁵⁰⁻⁵²

Protein adsorption

The effect of plasma treatment type and aging on protein adsorption after scaffolds incubation in cell culture media containing serum ((+)FBS) was analyzed (**Figure 5**). It was observed that fibronectin (Fn) and vitronectin (Vn), two relevant cell adhesive proteins present in serum, adsorbed in greater amounts to fresh plasma treated scaffolds than to untreated scaffolds (**Figure 5A, B**). Upon (+)FBS incubation, albumin, which is present at 100-1000 times higher concentration than adhesion-promoting proteins, adsorbs in the early phase of the protein layer formation. It is hypothesized that the hydrophobic interactions of albumin with the untreated PEOT/PBT are strong and resistant to displacement by Fn and Vn, while these proteins can effectively displace the weakly adsorbed albumin on the hydrophilic and charged plasma treated surfaces.⁵³⁻⁵⁴ Notably, Fn and Vn adsorption was significantly higher on APTMS scaffolds, which can be attributed to the electrostatic interactions between the positively charged amine based scaffolds' surface and the negatively charged proteins in solution leading to high protein adsorption (**Figure 5A, B**). Yet, the theory of electrostatic interactions cannot explain the high adsorption of negatively charged Fn and Vn to the negative charge of the surface of MA-

VTMOS and Argon scaffolds, compared to untreated scaffolds. Previous studies have also shown that Fn can be adsorbed at similar rates and amounts to both hydrophilic positively charged and negatively charged surfaces with zeta potential values similar to those measured within this study.⁵⁵ This has been explained by charged micro domains of the proteins yielding short-range attractions, which can lead to hydrogen bonds formation once the Debye interactions overcome the macroscopic electrostatic repulsions. Alternatively, adsorption of Fn or Vn to negatively charged polymers has been explained by a first adsorbed layer of positively charged serum proteins, such as laminin, allowing subsequent electrostatic interactions with negatively charged proteins.⁵⁶ Interestingly, the reduction in functional groups density and, potentially, surface charge, caused by aging led to a significant reduction in Fn and Vn adsorption, and a significant increase in albumin adsorption (**Figure 5C**), to MA-VTMOS and APTMS aged scaffolds. This resulted in comparable Fn and Vn adsorption to both scaffolds types, regardless their surface charge. Notably, argon scaffolds were not affected in this regard by aging and presented the largest relative cell-adhesive protein adsorption values among aged plasma treatment types. This observation further strengthens the hypothesis that argon plasma activation mostly led to a surface roughness change, rather than to a chemical change, and that is the nanoroughness, which remains unaltered overtime, the feature responsible of cell-adhesive protein adsorption.⁵⁷

Figure 5. Protein adsorption to scaffolds upon incubation in serum containing medium ((+)FBS): effect of plasma functionalization and aging conditions. Relative amounts of adsorbed (A) fibronectin, (B) vitronectin, and (C) albumin to untreated scaffolds and fresh and aged MA-VTMOS, APTMS and Argon plasma treated scaffolds. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.0001$; * for comparisons among treatments within aged or fresh scaffolds; \$ for comparisons among fresh and aged for each plasma treatment).

Cell adhesion mechanism to plasma treated scaffolds

Enhanced cell attachment on 2D surfaces and 3D scaffolds containing polar functional groups has been attributed to surface wettability,⁵⁸ surface charge potential,^{55, 59} or amount of protein adsorption.¹⁹ However, most of these studies have been performed in serum containing media, or in protein-coated surfaces, masking the potential effect of direct cell interactions with the functional groups, and therefore not fully characterizing the mechanism of cell adhesion to a specific surface chemistry. Thus, in order to discern among attachment mediated by electrostatic interactions with the charged surfaces, and attachment via integrin binding to proteins adsorbed from serum, fresh scaffolds were pre-incubated for 24h in (+)FBS or serum-free medium ((-)FBS), and subsequently seeded with hMSCs for 4h in (+)FBS or (-)FBS. Representative fluorescent images of hMSCs on scaffolds' cross section after overnight culture suggested an increase on cell attachment on plasma activated and polymerized scaffolds, compared to untreated scaffolds, when these were pre-incubated in (+)FBS, regardless of the presence or absence of serum during the 4h seeding (**Figure 6A**). This can be attributed to the higher adsorption of cell adhesive proteins (Fn and Vn) to these scaffolds, as shown in **Figure 5**. Despite improving cell adhesion and distribution, cell sedimentation towards the bottom of the scaffold was observed both on untreated and treated scaffolds (**Figure S5A**). It is known that, regardless of the scaffolds surface properties, cells in the macropores, which are far from the scaffolds filaments during the seeding process, are unable to interact with the scaffold's surface, leading to sedimentation towards the bottom and a monolayer formation. Looking at the quantitative values on seeding efficiency presented in **Figure 6C**, no differences in total cell attachment among scaffold types were observed, when pre-incubated and seeded in (+)FBS. In turn, slightly higher cell attachment was noted on MA-VTMOS and Argon scaffolds compared to APTMS and untreated scaffolds, when pre-incubated in (+)FBS but seeded in (-)FBS. In the case of APTMS, since cell distribution images show homogenous cell attachment in the cross section comparable to other plasma conditions and to its (+) FBS seeding counterpart, it is plausible that lower cell adhesion on the bottom of the scaffold might have contributed to lower overall cell adhesion values. Importantly, the sedimented cells that attached to the bottom of the untreated scaffolds, might have significantly contributed to the relatively higher seeding efficiency values than expected for the low cell coverage on their cross section. Interestingly, despite being pre-incubated in (-)FBS, homogeneous cell attachment was observed along the cross section of plasma

Figure 6. Cell coverage and cell seeding efficiency on fresh plasma treated scaffolds under different pre-incubation and seeding conditions. Fluorescence microscopy images (F-actin, green) of hMSCs in the cross sections of scaffolds after 24h of culture, and (A) pre-incubated with (+)FBS or (B) (-) FBS and seeded with (+)FBS or (-)FBS. Scale bars 1mm. Quantification of cell seeding efficiency on scaffolds (C) pre-incubated with (+)FBS and (D) pre-incubated with (-)FBS. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (* $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$).

treated scaffolds seeded in (+)FBS (**Figure 6B**), but not in untreated scaffolds. Furthermore, cell seeding efficiency values were comparable to those of scaffolds pre-incubated and seeded in (+)FBS (**Figure S5C**). These observations suggested that pre-incubation in (+)FBS was not necessary, when the seeding process was performed in (+)FBS, regardless of the plasma treatment. This is likely due to protein adsorption reaching an equilibrium already in short incubation times. In fact, serum proteins have been shown to adsorb to surfaces containing polar functional groups within the first minutes of incubation.^{53, 55} On the other hand, seeding with (+)FBS was necessary for optimum cell attachment on MA-VTMOS and argon scaffolds, when pre-incubated in (-)FBS. In this case, seeding with (-)FBS on MA-VTMOS treated scaffolds led to poor, and comparable cell coverage in scaffold's cross section to untreated scaffolds (**Figure 6B**), which together with lack of sedimented cells layer formation on the bottom (cells remained on the well plate) contributed to significantly lower seeding efficiency (**Figure 6D** and **Figure S5B**). Similarly, lower cell coverage and absence of a cell monolayer at the bottom were observed in cross section of argon treated scaffolds pre-incubated and seeded in (-)FBS, compared to other pre-incubation and seeding conditions. In both cases, due to the lack of protein attachment sites, the monolayer preferentially attached to the bottom of the seeding well plate rather than to the scaffold. Surprisingly, APTMS scaffolds pre-incubated and seeded in (-)FBS demonstrated a homogeneous and confluent cell coverage, as well as seeding efficiency values comparable to other pre-incubation and seeding conditions (**Figure 6D** and **Figure S5D**). Overall, these results suggest that i) MA-VTMOS plasma polymerization support cell attachment to scaffolds mostly through cell-protein interactions, and ii) cell attachment on APTMS plasma polymerized scaffolds is driven by cell-proteins interactions, in the presence of serum, and by electrostatic interactions between the cell and the amine groups coating the scaffolds filaments, in the absence of serum. The latter conclusion is in agreement with previously published reports studying cell adhesion on 2D surfaces suggesting that in protein-free conditions, direct interactions between negatively charged chondroitin sulfate proteoglycans in the cell membrane and the positively charged amine groups on a surface are responsible of promoting cell attachment.^{58, 60} On the other hand, repulsive electrostatic interactions between the negatively charged groups on the MA-VTMOS scaffolds and the cells do not allow for cell attachment in serum free conditions. In the case of argon plasma treatment, we hypothesize that the nanoroughness itself⁶¹ or combined with the lower presence of

charged hydrophilic groups, might have favored a slightly higher cell attachment in these scaffolds in serum free conditions compared to MA-VTMOS treated scaffolds.

To further analyze cell attachment mechanism to the different plasma treated scaffolds, cell morphology and focal adhesion formation in the absence or presence of serum was studied (**Figure 7**). Pre-incubation and seeding in (+)FBS led to cell spreading and the formation of F-actin stress fibers on hMSCs adhered to both untreated and plasma treated scaffolds, only after the 4h seeding. Moreover, while hMSCs did not show well defined vinculin structures on untreated scaffolds, established focal adhesions at the end of the stress fibers in the periphery of the cells were visualized on the plasma treated scaffolds, which was correlated to the higher amount of cell-adhesive proteins adsorbed on these surfaces allowing for integrin binding (**Figure 7**). Larger vinculin expression, as well as increased focal adhesion kinase expression, which localizes to focal adhesions to activate migration, proliferation and differentiation pathways, were also previously reported on amine, carboxyl, hydroxyl and argon functionalized surfaces pre-incubated in Fn ²⁰ or with (+)FBS medium, ⁶²⁻⁶³ compared to untreated surfaces. On the contrary, the few attached hMSCs to the untreated, MA-VTMOS and argon scaffolds, showed a well-defined round morphology and lacked focal adhesion complexes when pre-incubated and seeded in (-)FBS, due to the repulsive electrostatic forces and the lack of cell adhesive proteins hindering cell attachment spreading (**Figure 7**). The fact that, when looking at the scaffolds cross section, these cells were only found on the top surface of the filaments, further suggested that cells passively laid on top of the filament due to gravity during the seeding process, and that not active attachment occurred. On the contrary, hMSCs on APTMS scaffolds pre-incubated and seeded in (-)FBS were observed occupying the whole filament surface area. Interestingly, these cells were able to form multiple protrusions and F-actin fibers, which were not colocalized with focal adhesions, suggesting cell adhesion to the positively charged scaffold surface without mediation of integrin binding sites (**Figure 7**). Similar microfilament bundle formation and lack of focal contacts formation was previously reported on fibroblast adhesion to 2D amine coated surfaces. ⁶⁰

Figure 7. hMSCs morphology and focal adhesion formation after 4h seeding on fresh plasma treated scaffolds pre-incubated and seeded in (+)FBS or (-)FBS. Representative confocal microscopy images of hMSCs (F-actin, green; nuclei, blue; vinculin, red) on top of scaffold filaments 4h post-seeding. Continuous arrows indicating focal adhesions. Dashed lines delimitating the scaffold filament, with dashed arrows indicating the filament's surface. Scale bars 50 μ m.

Cell proliferation

Further cell culture studies aiming to assess cell behavior on the different plasma treated scaffolds were performed on scaffolds pre-incubated and seeded in (+)FBS, in order to have comparable cell number and distribution among scaffolds, as these parameters can influence cell behavior by their own.⁶⁴ Initially, we confirmed that aging did not affect total cell attachment (**Figure 8A**) nor cell distribution (**Figure S6**), regardless of the plasma condition. Plasma effect on cell adhesion and distribution also showed not to be hMSCs donor dependent, suggesting the possibility of extrapolating our results to other hMSCs populations (**Figure S6B, C**). Interestingly, hMSCs did not proliferate significantly after 7 days of culture, neither on untreated nor on fresh or aged plasma scaffolds (**Figure 8B**). Despite this initial lack of proliferation, a significant increase in DNA content with respect to day 1 was observed on all fresh and aged plasma treated scaffolds after 14 days of culture (7 days in BM followed by 7 days in MM) (**Figure 8C**). It is plausible that an increase in available surface area, given by ECM production in the scaffold pores volume when cultured in MM, offered cells the possibility to further proliferate on plasma treated scaffolds, as previously suggested.⁴ Previous research also showed significant differences in proliferation among surface functionalized and untreated 2D substrates only after long culture periods (~10 days),²⁵ whereas the use of dynamic culture systems or growth factors was found necessary to increase ECM production and boost cell proliferation on 3D plasma treated scaffolds.^{25, 27} Notably, no differences in hMSCs proliferation rate were found among fresh scaffolds, nor among aged scaffolds. Yet, while hMSCs in MA-VTMOS scaffolds proliferated at the same rate in fresh and aged scaffolds, higher levels in hMSCs proliferation were found on fresh APTMS and argon treated scaffolds, compared to their aged counterparts. In the case of APTMS, this might be due to the lower density of functional groups and adsorbed proteins on aged scaffolds.

Osteogenic differentiation potential of hMSCs seeded on aged plasma treated scaffolds

The analysis of osteogenic differentiation was performed on aged scaffolds, due to their higher applicability, as previously mentioned. Moreover, since comparable total cell numbers among conditions were observed at any timepoint evaluated during the culture on these scaffolds (**Figure S7**), it was then possible to focus solely on the influence of

Figure 8. Effect of plasma treatment and aging conditions on cell seeding efficiency and proliferation. (A) Cell seeding efficiency, and fold increase in DNA content after (B) 7 days of culture (in BM), and (C) 14 days of culture (7 days in BM and 7 days in MM) with respect to day 1, which is marked with a dashed line. Scaffolds were pre-incubated and seeded with (+)FBS. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$, * $p < 0.05$, ** $p < 0.01$; * for comparisons among plasma treatments within aged or fresh scaffolds, and among fresh and aged scaffolds for each plasma treatment; γ for comparisons between the given timepoint and day 1 for each plasma treatment).

plasma activation and polymerization on differentiation, and exclude the cell number and distribution related effect on osteogenesis. Following this rationale, inefficiently seeded untreated control scaffolds were not included in the study. After seeding, scaffolds were cultured for a total of 54 days in MM (7 days in BM, followed by 47 days in MM). Immunofluorescence was used to evaluate collagen I (COL I) deposition at early and late timepoints of the differentiation process (day 14 and day 54). Representative fluorescent microscopy images in **Figure 9A** revealed that hMSCs were able to produce COL I, one of the main bone ECM proteins, from early timepoints in all plasma treated scaffold conditions. These images also revealed that ECM became denser and pores more filled in the course of the culture. Furthermore, the expression of relevant osteogenic genes was screened through PCR (**Figure 9B**) at day 14 and 54 of culture. ALP, an early osteogenic marker, indicated no significant differences among plasma conditions or timepoints. Similarly, no statistical differences in OCN expression, a protein that binds to hydroxyapatite, were found among plasma conditions and timepoints. RUNX2, a transcription factor modulating the expression of osteogenic proteins, was upregulated at day 54 in Argon treated scaffolds compared to other conditions. When analyzing osterix (OSX), bone sialoprotein (BSP) and bone morphogenetic protein-2 (BMP2) expression, proteins present in a matured bone ECM, no upregulation was observed unless in argon treated scaffolds at day 54 compared to day 14. Yet, no significant differences among

plasma conditions were found at this time point. Overall, no clear trends were observed within this study and no plasma treatment showed to provide a distinct osteogenic stimulation to hMSCs over the others. In 2D functionalized surfaces, an enhanced osteogenic effect of amine groups compared to other functional groups has been correlated to the enhanced exposure of the integrin domain $\alpha 5\beta 1$ in adsorbed Fn, and to high levels of recruitment of focal adhesion components and phosphorylation of focal adhesion kinases in the adhered cells, which are events required for osteoblast differentiation.^{20-22, 62, 65} In this regard, it is plausible that aging masked the osteogenic stimulation potential of the APTMS surfaces, and that some differences would have been evidenced under fresh conditions, since APTMS fresh scaffolds displayed the highest levels of protein adsorption and, potentially, of integrin domain $\alpha 5\beta 1$. It is also plausible that the effect of surface chemistry is only pronounced at an early stage, when cells are in direct contact with the scaffolds surface, and that upon ECM production by cells during longer-term culture, cell-substrate contact becomes limited and ineffective.⁶⁶ Nevertheless, due to the lack of published reports simultaneously comparing the effect of different surface chemistries on osteogenic differentiation in 3D scaffolds, comparing our results with previous literature becomes challenging. To the best of our knowledge, a single study as such has demonstrated that salt leached poly(carbonate-urea) urethane scaffold modified by allylamine plasma polymerization, significantly enhanced osteogenic differentiation of adipose derived stem cells, compared to a carboxylic acid modified scaffold.⁶⁷ Yet, no clear explanation on how the amine group affected the osteogenesis pathway was reported. Moreover, since those scaffolds were fabricated by a conventional manufacturing method, i.e. porogen leaching, the poor interconnected porosity might have induced a different cell behavior compared to what we observed in ME-AM scaffolds.

Besides, it is worth noticing that ECM mineralization was not observed in any scaffold type during the culture period evaluated (**Figure S8A**), despite the proven mineralization potential of these cells in 2D (**Figure S8B**). Also, no matrix mineralization was attained on fresh scaffolds (**Figure S9C**). Thus, it is believed that cells on the scaffolds did not reach yet a matured level of differentiation, and that the homogenous cell distribution after seeding given by a bioactive surface chemistry is not sufficient for a successful osteogenic differentiation and mineralization outcome. It is hypothesized that larger cell seeding numbers could have led to larger cell attachment, and therefore higher cell density and cell-cell contact for accelerated differentiation.⁶⁸⁻⁷⁰ In order to investigate this, preliminary experiments were conducted, in which scaffolds were seeded with a larger seeding density

(400,000 cells per scaffold (400k), instead of 200,000 (200k)). Despite higher cell seeding efficiency, as per DNA quantification (**Figure S9A**), cells in scaffolds seeded with 400k tended to form a denser monolayer in the scaffolds' bottom, while maintaining the scaffolds' filaments as populated as when seeded with 200k (**Figure S9B, C**, day 1 of culture). Yet, some pores were already filled on the plasma treated scaffolds seeded with 400k at day 7 (**Figure S9D**), a cell confluency level that was not observed with lower cell seeding density, and that could potentially derive in earlier osteogenesis. Future work will be aimed at understanding if an enhancement of cell confluency at earlier timepoints, at the expense of needing a larger cell stock, might lead to mineralization and tissue maturation. This will help us to unravel if a specific plasma induced surface chemistry can offer enhanced osteogenic differentiation or if, on the contrary, cell confluency regardless of the surface chemistry is the primary parameter effectively playing a role on osteogenic differentiation on 3D AM scaffolds.

Figure 9. Evaluation of the osteogenic differentiation potential of hMSCs seeded on aged plasma treated scaffolds. (A) Representative confocal microscope images of hMSCs (F-actin, green) on top of scaffolds filaments after 14 and 54 days of culture (7 and 47 days in MM, respectively) and stained for the relevant osteogenic marker COL1 (red). Scale bars 100 μ m. (B) Gene expression of hMSCs after 14 and 54 days of culture (7 and 47 days in MM, respectively) comparing the three different plasma treatments. ALP, RUNX2, OCN, OSX, BSP and BMP2 fold change expression values relative to MA-VTMS day 14. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$; * $p < 0.05$; ** $p < 0.01$).

CONCLUSION

ME-AM enables the reproducible fabrication of highly porous polymeric scaffolds, ideal for tissue regeneration. However, the lack of cell adhesion sites on synthetic polymers hinders an efficient cell adhesion to the scaffolds, the first step towards the development of a functional construct. The aim of this study was to fabricate 3D scaffolds using a hybrid AM technique that enabled scaffold fabrication by ME-AM and their subsequent atmospheric pressure plasma treatment, to enhance their bioactivity, within the same platform. While argon activation resulted in an unspecific surface plasma treatment, APTMS plasma polymerization enabled the deposition of a positively charged coating containing amine functional groups, and MA-VTMOs rendered the surface of the scaffold negatively charged, by depositing carboxyl and anhydride functional groups. All plasma treatments increased the surface wettability of the scaffolds, enhanced cell-adhesive proteins adsorption to their surface, and allowed for homogeneous cell attachment along the scaffold cross section. Interestingly, cell attachment was found to be driven by cell protein interactions in the presence of serum and by electrostatic interactions between the cell and the charged scaffold surface in serum free conditions. This latter feature allowed for cell attachment and scaffold population on APTMS treated scaffolds in the absence of serum, which possess relevant clinical applications. Scaffolds storage led to the aging of the surface treatment, in terms of a slight reduction of exposed functional groups. Yet, cell attachment and proliferation were not significantly affected. Notably, none of the plasma treatments stimulated osteogenic differentiation of hMSCs significantly more than the others, and it is hypothesized that initial cell confluency might play a major role, overruling effects of a specific surface chemistry on the osteogenic differentiation of hMSCs and ECM mineralization on 3D ME-AM scaffolds. Overall, this newly proposed method enabled an efficient workflow of scaffold production and surface treatment, and opens the door to future research on the effect of different plasma treatments on cell behavior on 3D ME-AM scaffolds.

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SUPPLEMENTARY FIGURES

Table S1. Primer sequences used for qRT-PCR.^a

gene	forward primer	reverse primer
B2M	ACAAAGTCACATGGTTCACA	GACTTGTCTTTCAGCAAGGA
ALP	ACAAGCACTCCCACTTCATC	TTCAGCTCGTACTGCATGTC
RUNX2	TCAACGATCTGAGATTTGTGGG	GGGGAGGATTTGTGAAGACGG
OCN	TGAGAGCCCTCACACTCCTC	CGCCTGGGTCTCTTCACTAC
SP7	CCTCTGCGGGACTCAACAAC	AGCCCATTAGTGCTTGTAAGG
BSP	CCCCACCTTTTGGGAAAACCA	TCCCCGTTCTCACTTTCATAGAT
BMP2	ACTACCAGAAACGAGTGGGAA	GCATCTGTTCTCGGAAAACCT

^a B2M: β -2-microglobulin, ALP: alkaline phosphatase, RUNX2: Runt-related transcription factor 2, OCN: osteocalcin, SP7: osterix, BSP: bone sialoprotein, and BMP2: bone morphogenetic protein-2

Figure S1. (A) Stereomicroscopy image of APTMS scaffold stained with the cationic dye methylene blue, and fluorescent microscopy images of MA-VTMOS and Argon scaffolds stained with amine specific dyes, to prove the specificity of the staining. (B) Stereomicroscopy image of MA-VTMOS scaffold and fluorescent microscopy image of APTMS scaffold stained with coating specific dyes after ethanol disinfection to verify the maintenance of the treatment before cell culture experiments. Scale bars 1mm.

Figure S2. Zeta potential measurements of untreated, and Argon, MA-VTMOS and APTMS treated 2D substrates (melt-pressed films). Data presented as average $\pm$ s.d. and statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (**** $p < 0.0001$).

Figure S3. Regions of interest of the FTIR spectra of (A) MA-VTMOS, (B) APTMS and (C) Argon plasma treated 2D substrates (melt-pressed PEOT/PBT films).

Figure S4. SEM micrographs the surface of untreated and MA-VTMOS, APTMS and Argon plasma treated scaffolds. Scale bars 50 μm.

Figure S5. Fluorescence microscopy images (F-actin, green) of hMSCs in the bottom lids of scaffolds after 24h of culture and pre-incubated with (A) (+)FBS or (B) (-) FBS and seeded with (+)FBS or (-)FBS. Scale bars 1 mm. Quantification of cell seeding efficiency on scaffolds (C) seeded with (+)FBS and (D) seeded (-)FBS. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$; **** $p < 0.0001$).

Figure S6. (A) Fluorescence microscopy images (F-actin, green) of hMSCs on the cross section of aged plasma treated scaffolds after 24h of culture, and pre-incubated and seeded with (+)FBS. Scale bars 1 mm. (B) Fluorescence microscopy images (F-actin, green) of hMSCs from a different donor (donor B) on the cross section of fresh and aged plasma treated scaffolds. Scaffolds were pre-incubated and seeded with (+)FBS. Scale bars 1 mm. (C) Seeding efficiency on plasma treated scaffolds comparing hMSCs from 2 different donors. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$).

Figure S7. DNA content on aged plasma treated scaffolds after 14 or 54 days of culture (7 days or 47 days in MM, respectively). Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$).

Figure S8. (A) Stereomicroscopy images of aged (plasma treatment) scaffold cross sections stained with ARS after 35 and 54 days of culture (28 and 47 days in MM, respectively) with hMSCs. (B) Stereomicroscopy images of hMSCs monolayers (osteogenic differentiation in 2D) stained with ARS after 35 days of culture (28 days in MM). (C) Stereomicroscopy images of fresh (plasma treatment) scaffold cross sections stained with ARS after 35 days of culture (28 days in MM) with hMSCs. Scale bars 1 mm.

Figure S9. (A) DNA content on plasma treated scaffolds after seeding with two different cell densities: 200,000 cells (200k) or 400,000 cells (400k) per scaffold. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (** $p < 0.01$; *** $p < 0.001$, for comparisons between 200k and 400k for each plasma treatment). (B) Quantification of cell coverage on plasma treated scaffolds seeded with 200k or 400k cells after 7 days of culture in BM. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$). (C) Comparison of cell coverage among plasma treated scaffolds seeded with 200k or 400k cells per scaffold: (C) Fluorescent microscopy images of hMSCs in a representative cross sectional area and in the bottom lids of scaffolds after 1 day of culture. (D) Fluorescent microscopy images of hMSCs in a representative cross sectional area of scaffolds after 7 day of culture in BM. White arrows signing scaffolds' pores coverage by cells Scale bars 500 μm .

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Chapter 5

Effect of Highly Loaded Nanohydroxyapatite Composite Scaffolds Prepared via Melt Extrusion Additive Manufacturing on the Osteogenic Differentiation of Human Mesenchymal Stromal Cells

Effect of Highly Loaded Nanohydroxyapatite Composite Scaffolds Prepared via Melt Extrusion Additive Manufacturing on the Osteogenic Differentiation of Human Mesenchymal Stromal Cells

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Abstract

The field of bone tissue engineering seeks to mimic the bone extracellular matrix composition, balancing the organic and inorganic components. In this regard, additive manufacturing (AM) of highly loaded polymer-calcium phosphate (CaP) composites holds great promise towards the design of bioactive scaffolds. Yet, the biological performance of such scaffolds is still poorly characterized. In this study, melt extrusion AM (ME-AM) was used to fabricate poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT)-nanohydroxyapatite (nHA) scaffolds with up to 45 wt% nHA, which presented significantly enhanced compressive mechanical properties, to evaluate their *in vitro* osteogenic potential as a function of nHA content. While osteogenic gene upregulation and matrix mineralization were observed on all scaffold types when cultured in osteogenic media, human mesenchymal stromal cells did not present an explicitly clear osteogenic phenotype, within the evaluated timeframe, in basic media cultures (i.e. without osteogenic factors). Yet, due to the adsorption of calcium and inorganic phosphate ions from cell culture media and simulated body fluid, the formation of a CaP layer was observed on PEOT/PBT-nHA 45 wt% scaffolds, which is hypothesized to account for their osteoinductivity in the long term *in vitro*, and osteoconductivity *in vivo*.

INTRODUCTION

Bone is one of the most recurrently damaged tissues in our body in form of fractures or defects caused by bone disease, traumatic injury, implant surgery, infection or tumor resection. ¹ In case of long bones, bone loss in length can lead to critical-size defects, which will not heal on their own. Current strategies used in the clinics for the treatment of critical size bone defects mainly rely on i) distraction osteogenesis and induced membrane techniques, ²⁻³ or ii) on the use of graft materials, such as autografts, allografts, or demineralized bone matrices, that fill and / or help to regenerate the defect. ⁴⁻⁵ Treatment methods based on the first option are often complex, demanding for the patient, and require external fixation and multiple surgeries. Similarly, natural bone grafts, despite being osteoinductive and / or osteoconductive, possess several disadvantages, including donor site morbidity, restricted availability, high costs, pain, prolonged rehabilitation, lack of structural properties, and can trigger inflammation, immunological rejections or infections. ⁶ Therefore, the management of critical-size bone defects still remains a major clinical challenge, and consequently, a large amount of efforts are dedicated to investigate alternative strategies, such as tissue-engineered constructs to heal and regenerate bone in such scenarios. ⁷

Mimicking the physical and chemical properties of bone has been the main approach towards the design of the optimal scaffold for bone regeneration. The bone matrix mainly consists of an organic phase of collagen type I (COL I), among other proteins, and an inorganic phase of carbonated hydroxyapatite. ⁸ Accordingly, bioceramics, and in particular calcium phosphates (CaP), have been extensively applied for hard tissue repair. Widely applied CaP phases include hydroxyapatite (HA), tricalcium phosphate (TCP), biphasic and triphasic CaP, and amorphous CaP, ⁹ and their osteoinductive and osteoconductive performance is determined by various properties, such as chemical phase, structural properties (crystallinity, surface roughness, macro-, micro- and nanoporosity), mechanical properties and associated degradation behavior. ¹⁰ Yet, the role of the individual properties and their combination on the biological response to CaP-based bone graft substitutes has not yet been fully elucidated, with inconsistency in results found in the literature, which hampers the selection or design of the optimal CaP material for a specific application. ¹¹⁻¹² Several reasons for this are that studies lack sufficient material and/or biological characterization, or use different cell types *in vitro*, unavoidable inter- and intra-species variations in *in vivo* experiments, convoluted material properties that cannot

be easily tuned independently, and different fabrication techniques used for scaffolds manufacturing.¹¹⁻¹²

CaP processing into 3D macroporous scaffolds favors cell migration, tissue ingrowth, and vascularization. Moreover, the increase in surface area and body fluid accessibility provided by macroporosity can accelerate the degradation rate of the CaP material and boost the cell-material interactions towards an enhanced biological response, in terms of adhesion, proliferation and differentiation.¹³ Compared to conventional 3D scaffold fabrication techniques for polymers, such as gas foaming, particulate leaching, freeze-drying and solvent casting, additive manufacturing (AM) has emerged as a promising macroporous scaffold fabrication technique. AM enables the fabrication of patient specific constructs with a fully interconnected network of pores, whose size and shape can be tuned to control cell response and the scaffold mechanical properties, of special importance for bone tissue engineering.¹⁴ Nevertheless, pure CaP AM scaffolds, processed by powder bed and binder jetting techniques, require post-sintering steps, the removal of binders, and present poor mechanical properties, due to the inherent material brittleness and low fracture toughness, which overall limit their in vivo applications.¹⁵⁻¹⁶ Therefore, CaP have been processed by solution extrusion AM in combination with carrier liquids or solvents¹⁷⁻¹⁹, or by melt extrusion AM (ME-AM) in combination with synthetic biodegradable thermoplastic polymers. Among these, ME-AM of polymer-CaP composite materials has been widely explored, as it mimics the organic-inorganic composition of bone tissue, displays a high processability window, and allows the fabrication of scaffolds with tunable degradation rates and mechanical properties.²⁰⁻²¹ In particular, the combination of polymers with HA has gained interest, as HA closely mimics the bone mineral phase. Considering the average mineral content of bone tissue, which lies within the range of 50 wt% to 74 wt%, depending on the body location,²²⁻²³ tremendous efforts have been conveyed to increase the HA content in the scaffold, while retaining their printability and optimizing the mechanical properties. A few studies demonstrated feasibility of production of highly HA-loaded scaffolds processed by ME-AM, including poly(ϵ -caprolactone) (PCL)-HA composite with 40 wt% ceramic phase,²⁴⁻²⁵ poly(lactic acid) (PLA)-HA composite with 50 wt% ceramic,²⁶ and poly(ether urethane) (PEU)-HA composite with 40 wt% ceramic.²⁷ Despite material related advances, most of previously published reports focus only on the material synthesis and printability, and the structural characterization of the scaffold,^{24-26, 28-30} while their biological properties are poorly characterized.³¹⁻³²

Here, we studied the *in vitro* osteogenic differentiation potential of highly loaded poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT) – nano HA (nHA) composite scaffolds, fabricated by ME-AM. The biodegradable thermoplastic polyether-ester multiblock copolymer PEOT/PBT was selected due to its processability, mechanical properties and proven *in vitro* and *in vivo* applicability in the orthopedic field.³³⁻³⁷ While in previous studies this polymer has been blended with HA to a maximum of 25 wt% micro-HA,³⁸⁻³⁹ for the first time, PEOT/PBT-HA composite scaffolds with up to 45 wt% nHA were fabricated using ME-AM in the present report. To study their functionality in bone regeneration, human mesenchymal stromal cells (hMSCs) were chosen, due to the suitability of these stem cells over already committed cell lines to study the ability of a material to support osteogenic differentiation,⁴⁰ as well as for future clinical translation. We evaluated the effect of the scaffolds' nHA content on the osteogenic differentiation of hMSCs by analyzing cell attachment, proliferation, matrix production and mineralization, and the expression of a relevant set of osteogenic genes and proteins.

MATERIALS AND METHODS

Polymer-nHA composite production by solvent blending and characterization

The production of PEOT/PBT-HA composites with 20 wt% and 45 wt% nHA (20-nHA and 45-nHA, respectively) was carried out by a solvent blending procedure. PEOT/PBT pellets (PEO molecular weight = 300 kDa, PEOT:PBT weight ratio= 55:45, intrinsic viscosity 0.51 dl/g, Polyvation, The Netherlands) were dissolved in chloroform (Scharlab) under magnetic stirring for 45 min. An amount of 20 or 45 wt% nHA powder (particle size <200 nm, Sigma Aldrich) was added to the polymer solution and mechanically stirred at 750-1000 rpm for 15 min (Dissolver Dispermat CV3). After stirring, the dispersion was kept for 2h at 250 rpm, time during which part of the chloroform was evaporated and the viscosity of the blend increased. Subsequently, the polymer composite was precipitated in a mixture of diethylether and chloroform (Scharlab) (diethylether:chloroform = 10:1), and the supernatant removed. The composite was dried at RT, followed by a vacuum drying step at 90 °C for 2h to ensure the removal of solvent traces (diethylether <0.01 ppm, chloroform <1 ppm,). The foam-like composite formed during the precipitation step was compression molded at 150-175 °C and cut into pellets. The dispersion of nHA within the polymer was

examined using backscattered scanning electron microscopy (BSEM, FEI's Quanta 200 environmental scanning microscope, beam voltage 30 kV, spot size 5).

Scaffolds fabrication and characterization

45-nHA, 20-nHA and PEOT/PBT (0-nHA) scaffolds were fabricated via ME-AM. A commercial platform (Bioscaffolder 3.0, Gesim) was equipped with a custom-made printhead with two separate heating sources for the cartridge and extrusion screw.⁴¹ The printing parameters applied to each material are briefly described in **Table 1**. A needle with 250 μm internal diameter was used to print 0-90 layout scaffolds, with 200 μm layer thickness and 750 μm strand distance (center to center). For cell culture experiments and scaffolds morphological evaluation, 4x15x15 mm³ (HxWxD) blocks were manufactured and cylindrical scaffolds of 4 mm diameter and 4 mm height were cored out using a biopsy punch. For mechanical testing, 4 mm diameter, 15 mm high cylindrical scaffolds were printed and directly used for analysis. These scaffold dimensions resemble the type of scaffold needed for a rabbit critical segmental bone defect study, as a real case example which will be performed in future experiments. The scaffolds' architecture consisted also on 250 μm fiber diameter, 200 μm layer height, and 750 μm strand distance, and the scaffold layers had a 0-90 filament layout in the center, while at the edges, the filaments were deposited along the circular boundary. These scaffolds presented sufficient lateral porosity (**Figure S4A-B**).

Table 1. Printing parameters applied to obtained 3D scaffolds with each of the polymer composites

Scaffold	T cartridge (° C)	T screw (° C)	Motor rotation (rpm)	Cartridge Pressure (bar)	Printing speed (mm/s)
0-nHA	195	195	60	7	16
20-nHA	205	210	45	8	18
45-nHA	210	220	30	8	15

Scaffold morphology and porosity was assessed using a stereomicroscope (Nikon SMZ25) and scanning electron microscopy (SEM, XL-30, beam voltage 10 kV, spot size 3). Presence and distribution of fillers within gold sputter-coated scaffolds' filaments surface and cross-section was examined using BSEM (XL-30, beam voltage 20 kV, spot size 5). SEM operating at 25 kV coupled with energy dispersive X-ray spectroscopy (EDS) was used to quantify the chemical composition of the samples. Particle size distribution on

scaffolds was assessed on the acquired BSEM images and using the image analysis software tool XT Pro v3.2 (Soft Imaging System, GmbH).

Mechanical tests (compression at a rate of 0.1 mm/min., using a 100 N load cell) were carried out using an Instron Universal Mechanical test machine. Three scaffolds each, made from 0-nHA or 45-nHA were tested.

Cell seeding and culture

Cell expansion

HMSCs isolated from bone marrow were purchased from Texas A&M Health Science Center, College of Medicine, Institute for Regenerative Medicine (Donor d8011L, female, age 22). Cryopreserved vials at passage 3 were plated at a density of 1,000 cells cm⁻² in tissue culture flasks and expanded until approximately 80 % confluency in culture media (CM), consisting of α MEM with Glutamax, no nucleosides (Gibco), supplemented with 10% fetal bovine serum (FBS), without penicillin-streptomycin (PenStrep) at 37 °C and 5% CO₂.

Cell seeding in 3D scaffolds

Scaffolds were disinfected in 70% ethanol for 20 min, washed 3 times with Dulbecco's PBS and incubated overnight in CM for protein attachment. Before seeding, scaffolds were dried on top of a sterile filter paper and placed in the wells of non-treated plates. Passage 4 hMSCs were trypsinized and resuspended in a dextran solution (10 wt/wt % dextran in CM, 500 kDa, Farmacosmos) to slow down cell settling and guarantee homogeneous cell attachment throughout the scaffold. Dextran is a temporary supplement and will be washed away during the first 24h.⁴² A droplet of cell suspension (37 μ l containing 200,000 cells) was placed on top of each scaffold before slowly filling the scaffolds' pores. Seeded scaffolds were incubated for 4 hours at 37 °C and 5% CO₂ to allow cells to attach. After this time, scaffolds were transferred to new wells containing 1.5 ml of basic media (BM), consisting of CM supplemented with 200 μ M L-Ascorbic acid 2-phosphate and 1% PenStrep. Scaffolds were cultured for 7 days in BM followed by another 28 days in BM or mineralization media (MM), consisting of BM supplemented with dexamethasone (10 nM) and β -glycerophosphate (10 mM) (Sigma-Aldrich). Media was replaced after 24h and every two or three days during the culture period.

Biochemical assays

Alkaline phosphatase (ALP) activity and DNA assay

ALP activity and DNA content were evaluated over the culture period. For that, scaffolds were collected at every timepoint and washed with PBS, stored at -80 °C and freeze-thawed 3 times, to improve lysis efficiency. Samples were incubated for 1h at RT in a cell lysis phosphate buffer containing Triton X-100 (0.1 v %), at pH 7.8. After the addition of the chemiluminescent substrate for ALP CDP star® ready to use reagent (Roche) at a 1:4 ratio, luminescence was measured using a spectrophotometer (CLARIOstar®, BMG Labtech). Reported ALP values were normalized to DNA content, which was quantified using the CyQUANT cell proliferation assay kit (Thermo Fisher Scientific) on the remaining cell lysates. Samples for DNA quantification were incubated overnight at 56 °C in Proteinase K solution (1mg/ml Proteinase K (Sigma-Aldrich) in Tris/EDTA buffer) (1:1) for further cell lysis and matrix degradation. Subsequently, samples were freeze-thawed 3 times and incubated 1h at RT with a 20X diluted lysis buffer from the kit containing RNase A (1:500) to degrade cellular RNA. Samples were incubated with the fluorescent dye provided by the kit (1:1) for 15 min and fluorescence was measured (emission/excitation = 520/480 nm) with a spectrophotometer.

Extracellular matrix (ECM) mineralization assay

Calcium (Ca) deposits were qualitatively determined by alizarin red S (ARS) staining on composite scaffolds after 35 days of culture in BM or MM. Freshly fabricated scaffolds, and non-cell seeded scaffolds cultured in BM or MM and treated as cell seeded scaffolds in terms of media refreshment, were used as controls. Briefly, every timepoint scaffolds were fixed with 4% paraformaldehyde and washed with distilled water. Subsequently, scaffolds were stained with ARS (60 mM, pH 4.1 - 4.3) for 20 min at RT, and washed with distilled water. Scaffolds' cross sections were imaged using a stereomicroscope. After imaging, ARS was extracted from the scaffolds by 1h incubation at RT with 1 ml 30 v% acetic acid while shaking, followed by 10 min at 85 °C. Afterwards, scaffolds were removed and solutions were centrifuged at 20,000 rcf for 10 min. An appropriate volume of ammonium hydroxide (5 M) was added to the supernatants to bring the pH to 4.2. The absorbance was measured at 405 nm using a spectrophotometer. Concentration of ARS was calculated from an ARS standard curve and the values were normalized to DNA content. The signal from control scaffolds was subtracted.

Osteocalcin (OCN) secretion

OCN secretion in media was analyzed after 14, 21 and 35 days of culture in BM or MM. Scaffolds' culture media was refreshed on each timepoint by adding 1.5 ml of the corresponding media without FBS, in order to avoid background OCN levels from the FBS. Upon collection (48 h later), media was centrifuged at 2,000 rcf for 10 minutes to remove debris and stored at - 80 °C until analysis. OCN content in medium was detected using a quantitative sandwich ELISA kit (ab195214) according to the manufacturer's protocol.

Gene expression

Gene expression analysis was performed after 7, 14, 21 and 35 days of culture. RNA was extracted from cells in scaffolds using a TRIzol isolation method. Briefly, TRIzol was added to each sample in an Eppendorf tube and centrifuged at 12,000 rcf for 5 min to precipitate the scaffold and ECM at the bottom. Chloroform was added to the supernatant and centrifuged at 12,000 rcf for 5 min to isolate the RNA, present in the aqueous phase. RNA was further purified using RNeasy mini kit column (Qiagen), according to the manufacturer's protocol. The purity and quantity of total RNA was evaluated using a spectrophotometer. Reverse transcription was performed using iScript™ (Bio-Rad) following suppliers' protocol. qPCR was performed on the mix composed of cDNA, SYBRGreen master mix (Qiagen) and the selected primers (**Table 2**) using a CFX Connect™ Real-Time System (Bio-Rad) under the following conditions: cDNA denaturation for 3 min at 95 °C, 40 cycles of 15 s at 95 °C, and 30 s at 65 °C. Additionally, a melting curve was generated for each reaction in order to test primer dimer formation and non-specific amplification. Gene transcription was normalized to the transcription of the housekeeping gene B2M. The $2^{-\Delta\Delta C_t}$ method was used to calculate relative gene expression for each target gene. Normalization was done with respect to relative gene expression of cells in 0-nHA scaffolds at day 7 in BM.

Table 2. Primer sequences used for q-PCR. B2M: beta-2-microglobulin, RUNX2: runt-related transcription factor, COL I: collagen 1 alpha 1, OPN: osteopontin, OCN: osteocalcin, BSP: bone sialoprotein, SP7: osterix.

Gene	Forward primer	Reverse primer
B2M	ACAAAGTCACATGGTTCACA	GACTTGTCTTTCAGCAAGGA
RUNX2	TCAACGATCTGAGATTTGTGGG	GGGGAGGATTTGTGAAGACGG
COL I	GAGGGCCAAGACGAAGACATC	CAGATCACGTCATCGCACAAAC
OPN	CTCCATTGACTCGAACGACTC	CAGGTCTGCGAACTTCTTAGAT

OCN	TGAGAGCCCTCACACTCCTC	CGCCTGGGTCTCTTCACTAC
BSP	CCCCACCTTTTGGGAAAACCA	TCCCGTTCTCACTTTCATAGAT
SP7	CCTCTGCGGGACTCAACAAC	AGCCCATTAGTGCTTGTAAGG

Immunofluorescence

Cells in scaffolds were fixed in 4% paraformaldehyde after 35 days of culture in BM or MM, and permeabilized by 30 min incubation in Triton-X 100 (0.1 v%). Subsequently, the scaffolds were cut in half and washed 3 times with PBS. For assessing cell content and distribution on scaffolds, some scaffolds were directly stained for F-actin (488 Alexa Fluor Phalloidin, Thermo Fisher Scientific) for 1 h. Scaffolds cross-sections were imaged with a fluorescent microscope (Eclipse, Ti2-e, NIKON). Background subtraction and brightness adjustments were performed on the images using the software Image J, in order to improve their visualization. For antibody staining, scaffolds were further blocked for 1 h in blocking buffer (3 % BSA + 0.01% Triton-X 100). Afterwards, primary antibodies diluted 1:200 in washing buffer (10x diluted blocking buffer) were added (COL I rabbit polyclonal (ab34710, Abcam), osteopontin (OPN) rabbit polyclonal (ab8448, Abcam), OCN rabbit polyclonal (ab93876), each to a different sample half, and incubated overnight at 4 °C. Washed samples were incubated for 1h at RT with the secondary antibody (1:200 Alexa Fluor 647 goat derived anti-rabbit antibody, Thermo Fisher Scientific). Then, scaffolds were washed and stained for F-actin (488 Alexa Fluor Phalloidin, Thermo Fisher Scientific) for 1 h, followed by three washing steps with PBS. Confocal laser scanning microscopy was performed with a tandem confocal system (Leica TCS SP8 STED), equipped with a white light laser (WLL). Samples were excited with the dye specific wavelengths and emission was detected with HyD detectors. For optimal visualization in the reported images, F-actin was colored in green, COL I in red, OPN in blue and OCN in magenta.

Imaging of cultured scaffolds

After 35 days of culture, cell seeded and non-cell seeded scaffolds in BM or MM were fixed with 4 wt% paraformaldehyde for 30 min. Scaffolds were dehydrated by a graded ethanol series (30, 50, 70, 80, 90, 96, 100 v%), during 15 min each at RT, followed by 15 min incubation in 100% ethanol- hexamethyldisilazane (HMDS) at a 1:1 ratio, and 15 min incubation in HMDS. HMDS was removed and scaffolds were allowed further drying overnight at RT in the fume hood. Samples were gold sputter coated. Cells, ECM and scaffolds surface were visualized on scaffolds cross-section using scanning electron

microscopy (SEM, beam voltage 10 kV, spot size 3). The presence of mineral deposits on cell seeded scaffolds was examined by backscattered SEM (BSEM, beam voltage 20 kV, spot size 5). SEM operating at 25 kV coupled with Energy-dispersive X-ray spectroscopy (EDS) was used to identify Ca, phosphorous (P), and other elements on the ECM.

Bioactivity test in Simulated Body Fluid (SBF)

An SBF solution was prepared as previously described ⁴³. Briefly, chemicals were dissolved in distilled water in the following order: NaCl, NaHCO₃, KCl, K₂HPO₄, MgCl₂·6H₂O, CaCl₂·2H₂O and Na₂SO₄. The solution was then buffered to pH 7.4 at 36.5° C using Tris and HCl (1M). The final solution had an ion concentration (mM) as follows: Na⁺, 142.0; K⁺, 5.0; Mg²⁺, 1.5; Ca²⁺, 2.5; Cl⁻, 147.8; (CO₃)⁻, 4.2; (PO₄)²⁻, 1.0; (SO₄)²⁻, 0.5. Scaffolds were immersed in 1ml SBF and incubated at 37 °C. SBF was refreshed every 3-4 days. After 4 and 14 days, scaffolds were carefully rinsed in distilled water, blotted on an adsorbent paper and dried at RT. Then they were cut in half, gold sputtered and observed with SEM at 15 kV, spot size 3.

Ion release analysis

Before media refreshment at timepoints day 7, 14, 21 and 35 of the cell culture experiment, medium was collected from each scaffold and stored at -30 °C until analysis. In a separated experiment, scaffolds were incubated in 1ml simulated (0.8% NaCl, 50 mM HEPES, pH 7.3) for 28 days at 37 °C. This was done to analyze Ca and P release in a non Ca and P saturated solution. Every 3-4 days, an aliquot of the supernatant was collected, stored at -30 °C until analysis, and the same volume of SPS was refreshed. Concentrations of Ca and P in cell culture medium and SPS were quantified by Inductively coupled plasma mass spectrometry (ICP-MS, iCaP Q, Thermo Scientific). Upon analysis, all samples, including fresh cell culture medium or fresh SPS, were thawed, vortexed for 30 s, centrifuged at 20,000 rcf for 15 min to remove debris, and diluted 1:50 in 1 v% HNO₃ with the addition of 20 ppb scandium as internal standard. A standard curve of 20, 10, 5, 2.5 and 1.25 ppm Ca and P in 1 v% HNO₃ and 20 ppb scandium was prepared. Quantification was performed in standard mode with He as collision gas.

Statistical analysis

All data is shown as average with error bars indicating the standard deviation of at least three replicates. Analysis of statistics was conducted with GraphPad Prism (version 8.0.1). A one-way or two-way analysis of variance (ANOVA) was performed followed by a Tukey's post-hoc multiple comparison test to evaluate statistical significance and correct for multiple comparisons.

RESULTS

Scaffolds fabrication, nHA distribution and mechanical properties.

Composite pellets prepared by solvent blending of PEOT/PBT and 20 or 45 wt% nHA (**Figure S1**) were used to fabricate PEOT/PBT-nHA composite scaffolds via ME-AM (**Figure 1**). SEM micrographs in **Figure 1A, B** showed that both 20-nHA and 45-nHA scaffolds maintained an interconnected macroporosity, comparable to the control 0-nHA scaffolds. BSEM images in **Figure 1C** and EDS analysis in **Figure S2A** confirmed an increasing amount of nHA, and therefore Ca and P, on the scaffold filaments with increasing nHA content in the composite blend. The presence of nHA on the surface of the 20-nHA and 45-nHA scaffolds was further confirmed by ARS, which binds to the Ca of nHA (**Figure S2B, C**). An increase in the pink-red stain intensity on the surface of scaffolds was observed with increasing nHA content. Despite the nanometer size of the raw nHA (<200 nm), particle aggregation was already noticed in the solvent blending process prior to ME, possibly occurring during the precipitation step of the polymer composite solution in the non-solvent (**Figure S1B**). In the 45-nHA scaffolds, nHA micro aggregates occupied around 24% of the filaments' surface area. Quantification on their size distribution revealed that ~75% of the nHA micro aggregates possessed a diameter in the range of 2-10 μm , while ~5% of the particles diameter lied in the range of 30 to 90 μm (**Figure S3**). The other 76% of the 45-nHA scaffolds filaments surface area was occupied by the polymer and nanometer size nHA particles. The presence of these nHA micro aggregates provided a higher surface roughness to the composite scaffolds when compared to the smooth 0-nHA scaffolds (**Figure 1B**). The addition of nHA up to 45 wt% led to a 2-fold increase in the scaffolds' compressive modulus, compared to copolymer only scaffolds (**Figure S4C**). Compressive strength at yield was also slightly increased on the 45-nHA

scaffolds (**Figure S4D**). Yet, no significant differences were observed compared to 0-nHA scaffolds.

Figure 1. PEOT/PBT-nHA composite scaffolds fabrication and nHA distribution in scaffold filaments. (A) SEM micrographs of scaffolds cross-section showing their macroporosity. Scale bars 1 mm. (B) Representative BSEM micrographs of scaffolds filaments depicting nHA distribution and increasing surface roughness with increasing nHA content. Scale bars 100 μm . (C) Magnified images of the regions marked with dashed lines in (B). Scale bars 20 μm .

hMSCs seeding efficiency, proliferation and osteogenic differentiation on PEOT/PBT-nHA composite scaffolds

Scaffolds were seeded using a viscous seeding solution to ensure homogeneous hMSCs attachment throughout the scaffold. Consequently, all scaffolds qualitatively showed uniform cell distributions from the earliest timepoints. Moreover, high seeding efficiency values in the order of ~80 % were maintained among different scaffold types without statistical differences in terms of cell attachment (**Figure 2A**). To understand the effect of nHA on the osteogenic differentiation of hMSCs, scaffolds were cultured up to 35 days in

BM or MM. To support cell proliferation before the addition of osteogenic factors, scaffolds were cultured initially for 7 days in BM. Yet, DNA content analysis demonstrated no cell proliferation during this period (**Figure 2A**). DNA content was further monitored over the culture time (**Figure 2B**). Although no statistical differences were observed in terms of cell number among scaffold types and time points, a decrease in cell number from day 14 to day 35 of culture in BM was observed for all scaffold types. Overall, **Figure 2B** also demonstrated lower cell numbers in scaffolds cultured in BM when compared to MM at late timepoints (day 21 and day 35).

Figure 2. Cell seeding and osteogenic differentiation on PEOT/PBT-nHA composite scaffolds. (A) Seeding efficiency and proliferation 7 days after seeding. (B) DNA content progression on 0-nHA, 20-nHA, 45-nHA scaffolds over 35 days of culture in BM or MM. (D) ALP activity of hMSCs progression over 35 days when seeded on the different scaffold types in BM or MM. (E) OCN secretion by hMSCs in media progression over 35 days when seeded on the different scaffold types in BM or MM. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test. For (A,B): n.s. $p > 0.05$. For (C): *\$ $p < 0.05$; **\$\$ $p < 0.001$; ***\$\$\$ $p < 0.001$; ****\$\$\$\$ $p < 0.0001$; * for comparisons among timepoints each scaffold type; \$ for comparisons to 0-nHA each timepoint; comparisons among BM and MM samples performed separately. For (D): n.s. $p > 0.05$; *\$ $p < 0.05$; **\$\$ $p < 0.001$; ***\$\$\$ $p < 0.001$; ****\$\$\$\$ $p < 0.0001$; * for comparisons among timepoints each scaffold type; \$ for comparisons to 45-nHA each timepoint; comparisons among BM and MM samples performed separately.

ALP activity, an early osteogenic marker, was evaluated at day 14, 21 and 35 (**Figure 2C**). ALP values were higher when cells were cultured in MM compared to BM. A peak in ALP activity was shown at day 14, with a progressive and significant decrease on day 21 and day 35, both in BM and MM. While in MM no statistical differences among scaffold types were observed at any time point, ALP values in BM were found to be significantly higher at day 14 and 21 in 0-nHA scaffolds compared to 20-nHA and 45-nHA scaffolds. When analyzing OCN secretion in media, higher values were observed in MM than in BM (**Figure 2D**). Since OCN is a late marker for osteogenic differentiation, a peak was observed at day 35, in contrast to ALP. As noted with ALP, no significant differences were observed in OCN secretion among scaffold types in MM. Nevertheless, a significantly higher OCN signal was recorded on 45-nHA scaffolds already at an early time point (day 14) when cultured in BM, compared to 20-nHA and 0-nHA scaffolds in BM. Notably, other scaffold types cultured in BM reached this levels of OCN secretion at day 21 or 35 at the earliest.

Gene expression

The expression of bone related transcription factors and protein encoding genes was assessed at day 14, 21 and 35, under BM or MM conditions. **Figure 3** shows the relative gene expression change with respect to 0-nHA day 7 BM condition (**Figure S5**). In general, RUNX2 gene expression was found to be higher in MM than in BM for all scaffold types over the whole culture period (**Figure 3A**). At day 14 in MM, significantly higher expression on 0-nHA scaffolds was observed. However, no significant differences among scaffold types were observed at later time points in MM. Similarly, despite higher COL I gene expression on 0-nHA scaffolds at day 14 in MM compared to the nHA composite scaffolds, comparable COL I gene expression levels were maintained at later culture time points among all scaffold types (**Figure 3B**). Interestingly, the culture media appeared not to have any effect on COL I gene expression and no significant differences were found between BM and MM at a given time point and scaffold type over the culture period. In the case of OPN gene expression, higher levels were noticed when cultured in BM compared to MM for all scaffold types at days 14 and 21 (**Figure 3C**). Yet, at day 35 the gene expression level of OPN was comparable among all scaffold types in both BM and MM media. Independent of the culture media, a low gene expression of OCN was maintained in all scaffold types until a significant upregulation, slightly higher in 0-nHA scaffolds, occurred at day 35 in MM (**Figure 3D**). A significant upregulation in the BSP and Osterix gene expression was observed for all scaffold types cultured in MM from day 14 to day 21, and

to day 35 in the case of Osterix (**Figure 3E, F**). At day 35, gene expression levels in 0-nHA and 45-nHA remained comparable and significantly higher than in BM.

Figure 3. Gene expression of hMSCs cultured on scaffolds with different nHA concentrations for 14, 21 and 35 days in BM or MM. (A) RUNX2, (B) COL 1, (C) OPN, (D) OCN, (E) BSP, and (F) Osterix fold change expression values relative to 0-nHA d7 BM. Data presented as average $\pm$ s.d., and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (*\$ $p < 0.05$; **\$\$ $p < 0.01$; ***\$\$\$ $p < 0.001$; ****\$\$\$\$ $p < 0.0001$; * for comparisons among timepoints each scaffold type; \$ for comparisons to 0-nHA (control) each timepoint; comparisons among BM and MM samples performed separately).

Protein expression

Complementary to gene expression, protein expression at day 35 was evaluated by immunofluorescence on all scaffold types and culture conditions. Extracellular COL I was found covering filaments on all scaffold types and culture media conditions (**Figure 4**). Supporting the gene expression profiles, OPN protein expression was found slightly more abundant on scaffolds cultured in BM compared to MM, except in 45-nHA scaffolds in MM, where OPN protein expression was higher than in 20-nHA or 0-nHA scaffolds. Following an opposite trend, OCN protein expression was higher in scaffolds cultured in MM compared to BM (**Figure 4**). Interestingly, OCN protein expression was also recorded on nHA containing scaffolds (20-nHA and 45-nHA) cultured in BM.

Figure 4. Influence of nHA content on the hMSCs expression of different osteogenic proteins. Representative confocal microscopy images of hMSCs (F-actin, green) on top of the filaments of 0-nHA, 20-nHA and 45-nHA scaffolds after 35 days of culture in BM or MM and stained for the relevant osteogenic markers COL I (red), OPN (blue) and OCN (magenta). Scale bar for all images: 200 μ m.

Extracellular matrix (ECM) production and mineralization

ECM mineralization was examined at day 35 of culture in BM and MM. An increasing amount of Ca deposition with increasing nHA content was observed, both quantitatively (**Figure 5**) and qualitatively, in scaffolds cultured in MM (**Figure 6A, Figure S6**). Although

at lower levels than in MM, ARS quantification also revealed Ca deposits on 20-nHA and 45-nHA scaffolds when cultured in BM (**Figure 5**), after subtraction of the signal given by non-cell seeded stained scaffolds. In BM Ca deposition was mostly appreciable on 45-nHA scaffolds, where a more intense red color, compared to the non-seeded controls, was observed covering most of the surface area of the 45n-HA scaffolds filaments, as well as the scaffold filaments' exposed cross sections on the scaffolds outer surface (**Figure 7A**, **Figure S6**).

Figure 5. Quantification of the ARS staining extracted from scaffolds after 35 days of culture in BM or MM normalized to cell number. Data presented as average $\pm$ s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test. *\$ $p < 0.05$; ***\$\$\$ $p < 0.001$; ****\$\$\$\$ $p < 0.0001$; * for comparisons among BM and MM each scaffold type; \$ for comparisons among scaffold types each culture media type.

Next, cell and ECM coverage at day 35 of culture was investigated. When cultured in MM, cells in all scaffold types were confluent occupying both the scaffolds' filaments and pores along the whole scaffold cross-section (**Figure 6B** and **Figure S7**). Moreover, a dense and fibrillar ECM network produced by cells was observed for all scaffold types (**Figure 6C** and **Figure S8**). BSEM images in Figure 5D revealed bright points intercalated within the ECM in scaffolds pores, which were discriminated via EDS as Ca and P deposits, as well as a minimal amount of Na salts from the culture medium (**Figure S9A**). In addition, a peak of N, corresponding to the ECM proteins, was also observed. In contrast to the dense ECM in scaffolds cultured in MM, a thin layer of cells and ECM was visualized covering the scaffolds' filaments when cultured in BM (**Figure 7B-C**, **Figure S7**, **Figure S8**). Notably, BSEM images of the ECM of cells cultured in BM did not depict bright points corresponding to mineral deposits (**Figure 7D**) and EDS analysis demonstrated significantly lower Ca and P signals compared to the analysis on scaffolds cultured on MM (**Figure S9B**). To be

able to better discern among the EDS signal coming from the Ca and P already contained in the scaffold material, and the newly deposited minerals, not only the cell and ECM directly deposited on top of the scaffolds filaments was analyzed, but also the cell and ECM layer detached from the filament or present in the pores. Ca and P content in the ECM on the pores of MM scaffolds was found to be significantly higher than that on BM scaffolds (**Figure S9**).

Figure 6. Influence of scaffolds' nHA content on matrix mineralization when cultured for 35 days in MM (7 days in BM followed by 28 days in MM). (A) Representative stereomicroscope images of scaffolds cross sections stained with ARS. Inserts represent the corresponding control scaffolds

without cells incubated in MM and stained with ARS. Scale bars main images and insets 1 mm. (B) Representative fluorescent microscopy images of hMSCs (F-actin, green) on scaffolds' cross sections. Scale bars 200 μ m. (C) Representative SEM micrographs of scaffolds' cross sections depicting cell and ECM coverage. Scale bars 200 μ m. (D) BSEM images of the regions marked with dash lines (ECM in scaffolds' pores) in (C) for examining the mineral deposits. Scale bars 20 μ m.

Figure 7. Influence of scaffolds' nHA content on matrix mineralization when cultured for 35 days in BM. (A) Representative stereomicroscope images of scaffolds cross sections stained with ARS. Inserts represent the corresponding control scaffolds without cells incubated in BM and stained with ARS. Scale bars main images and insets 1 mm. (B) Representative fluorescent microscopy images of hMSCs (F-actin, green) on scaffolds' cross sections. Scale bars: 200 μ m. (C) Representative SEM micrographs of scaffolds' cross sections depicting ECM coverage. Scale bars: 200 μ m. (D) BSEM images of the regions marked with dash lines (ECM in scaffolds' pores) in (C) for examining mineral deposits. Scale bars 20 μ m.

Ca and P ion exchange dynamics with the medium

To investigate the degradation of the nHA contained in the scaffolds, Ca release into SPS, which does not contain these ions, was quantified by ICP-MS. nHA-containing scaffolds released Ca into SPS continuously over the period evaluated (26 days) (**Figure S10**). In agreement with the nHA content on the scaffolds, 45-nHA scaffolds released larger amounts and at a higher rate (50 - 200 μM every 3-4 days) than 20-nHA scaffolds (10 - 50 μM every 3-4 days).

To investigate the Ca and P content of the medium when scaffolds are in a cell culture setting, ICP-MS analysis was carried out on media collected at different time points during culture. This enabled to elucidate the ion exchange dynamics of Ca and P between the scaffolds and the media. The lower concentrations of Ca in media in contact with scaffolds compared to fresh media concentrations demonstrated that all scaffolds, cell seeded and non-cell seeded, depleted the media from Ca, especially when cultured in MM (**Figure S11**). Similarly, scaffolds cultured in MM depleted the media from P, while P levels in BM remained similar to fresh media ones. Interestingly, in general no significant differences in Ca and P depletion from media were found among cell seeded and non-cell seeded control scaffolds, nor among scaffolds with different nHA concentrations (**Figure S11**). To complement these results, the surface of the filaments of cell seeded (areas without cells) and non-cell seeded scaffolds cultured in BM and MM were analyzed by SEM and EDS (**Figure S12** and **Figure S13**). Interestingly, while according to ICP-MS 0-nHA scaffolds depleted the media from Ca in BM, and from Ca and P in MM, no Ca and P was observed on any of the 0-nHA scaffolds surface. In contrast, the surface of 20-nHA and 45-nHA scaffolds in MM was found to be covered by CaP spheres in non-cell seeded scaffolds and by a compact CaP layer in cell seeded scaffolds. In BM, these scaffolds showed a higher amount of white particles in their surface, compared to as-prepared scaffolds, potentially corresponding to the deposition of a Ca based mineral phase. While not easy to verify by either imaging or EDS, the surface of 45-nHA scaffolds in BM also looked more populated with bright spots, potentially corresponding to Ca deposits.

The scaffolds were also immersed in SBF, a solution that closely mimics human blood plasma and is used to test the ability of a material to deposit mineral on its surface. SEM showed a more pronounced deposition of mineral on 45-nHA scaffolds compared to 20-nHA scaffolds (**Figure 8**). As expected, no mineral deposition occurred on the 0-nHA

scaffolds even after 14 days of immersion, when a thick CaP layer was already covering the 45-nHA scaffolds.

Figure 8. Representative SEM micrographs of scaffolds' filaments surface mineralization after immersion in SBF for 4 and 14 days. As-prepared scaffolds (day 0) are presented as controls. Scale bars: 20 μm . Inserts represent magnified regions. Scale bars 5 μm .

DISCUSSION

During the last decades, AM has rapidly grown into the field of tissue engineering offering patient specific 3D scaffolds from a wide range of biocompatible materials, including polymer-CaP composites. CaP fillers are highly desired when aiming to mimic the composition of bone and to provide scaffolds with high strength, tunable degradation and favorable bioactivity. While solvent extrusion has permitted the fabrication of AM scaffolds with up to 90 wt% HA, ⁴⁴⁻⁴⁵ the need of low volatility organic solvents, i.e. plasticizers, has shown to lead to solvent remnants within the scaffold, as well as to elastic constructs with poor applicability in load bearing applications. The advantage of thermoplastic ME-AM for bone tissue engineering over other AM techniques, such as stereolithography, mainly lies

in the possibility of processing viscous composite materials including fillers, such as CaP particles, in a cost effective manner, and without requiring the use of organic solvents. Since the firstly reported PCL-CaP ME-AM scaffolds, containing up to 25 wt% CaP,⁴⁶⁻⁴⁹ the field has largely evolved seeking to fabricate scaffolds with an inorganic content closer to native bone ECM, and with new polymeric materials. Here, we were able to fabricate composite 3D scaffolds based on PEOT/PBT, which have a great potential in the orthopedics field, and with up to 45 wt% nHA, closer to native bone mineral content than the majority of the until now reported scaffolds. Despite the high nHA loading, composites were printable and scaffolds' filaments maintained a good shape fidelity and macroporosity (pore size 500 μm), which is expected to favor tissue infiltration and vascularization *in vivo*.⁵⁰ When looking at the filaments' surface and cross-sections of the composite scaffolds, nHA aggregates with a size in the order of tens of microns were observed, in accordance with previous reports, where melt extruded filaments with HA concentrations above 10 wt% demonstrated HA agglomeration.⁵¹ In spite of these microaggregates, nanosized particles were still present in a well-dispersed manner within the polymer matrix of the scaffolds and on their surface, which were expected to elicit a higher degree of biological response compared to micro-sized counterparts, due to their high surface-to volume-ratio and to their small size, allowing uptake by cells.⁵²⁻⁵³ Most importantly, the high nHA content significantly enhanced the mechanical properties of the scaffolds, with 45-nHA scaffolds having a compressive modulus of 92 ± 25 MPa and compressive strength of 4.4 ± 0.9 MPa, both in the range of cortical bone mechanical properties.⁵⁴⁻⁵⁵ Moreover, these scaffolds are stiffer than previously reported melt extruded AM scaffolds prepared with PCL-HA composites with similar inorganic loadings. This suggests the suitability of 45-nHA scaffolds for their application in bone regeneration scenarios, compared to bare PEOT/PBT scaffolds, as well as highly loaded PCL scaffolds, which is a commonly used polymer for bone scaffolds.²⁴⁻²⁵

In vivo, osteoinductive materials have the ability of recruiting stem cells and differentiating them towards an osteogenic phenotype in a heterotopic location.^{12, 56} Due to the complexity of the *in vivo* microenvironment, *in vitro* models do not always reliably predict osteoinductivity.⁵⁷ Yet, culturing stem cells on biomaterial scaffolds and analyzing cells' ability to differentiate can give some initial insights into a material osteogenic potential and aid to understand the osteoinduction mechanism. Thus, we cultured hMSCs on scaffolds in BM, without osteogenic factors, i.e. dexamethasone and β -glycerophosphate, and assessed their phenotype as a function of nHA content. In addition, we evaluated the

synergistic effect of the material and these soluble factors that stimulate osteogenic differentiation/mineralization, by culturing in MM. Previously, it has been shown that cell distribution and density upon seeding can significantly influence hMSCs osteogenic differentiation in 3D scaffolds.^{42, 58-59} Accordingly, poor cell attachment and lack of confluency within the scaffold cross-section have been shown to result in poor differentiation and lack of matrix mineralization *in vitro*. Besides, some studies reported enhanced cell attachment on polymer-CaP composites, likely due to the hydrophilicity and protein adsorption capacity of the CaP.⁶⁰⁻⁶¹ In the present study, in order to decouple the osteogenic potential of the PEOT/PBT-nHA composite scaffolds from the effect of attachment efficiency, a viscous solution was used as seeding media, thereby ensuring comparable cell attachment and homogeneity among all scaffolds regardless of the nHA content, as previously reported.⁴² Due to scaffold surface saturation with cells upon seeding, no cell proliferation was observed in the first 7 days of culture, nor during the subsequent 28 days in BM on any scaffold type, as ECM production was limited to the scaffolds filaments surface. On the other hand, ECM was also produced within the pores when cultured in MM, which increased the growth surface area enabling cell migration and a slight, yet not statistically significantly different, increase of cell number. An important general conclusion that can be drawn from the DNA quantification, and cell and ECM imaging on the scaffolds, is that 3D ME-AM PEOT/PBT scaffolds with up to 45 wt% nHA are able to support cell growth, both in BM and MM.

Osteogenic differentiation was initially evaluated by ALP activity, an early marker for osteogenesis,⁶² and OCN secretion, one of the major bone non-collagenous proteins with the ability to bind to the bone HA.⁶³ Moreover, the gene expression of RUNX2, an essential transcription factor to boost the expression of bone ECM proteins⁶⁴⁻⁶⁵, as well as other bone markers for osteogenesis at mRNA level (i.e. COL I, OPN, OCN, osterix and BSP), were analyzed during culture. According to the progression of bone markers during osteogenic differentiation suggested by literature, ALP activity was at its highest at early time points, and RUNX2 gene expression was upregulated from early time points, while OCN, osterix and BSP genes were highly expressed at later time points when cultured in MM.^{62, 66-67} COL I and OPN expression was maintained close to basal levels in BM, and slightly downregulated, with respect to day 7, in MM. Interestingly, no significant effect of nHA was observed in MM, suggesting that the soluble osteogenic factors might play a more pronounced role than HA in the stimulation of osteogenesis, and the absence of a synergistic effect. In the case of culture in BM, all gene expression levels were much lower

compared to MM cultures, maintaining gene expression at basal levels, with a slightly upregulation at day 35. Similar to MM cultures, no major differences in the gene expression levels were observed among scaffold types when cultured in BM. Yet, immunofluorescence revealed OCN and OPN protein expression at day 35 in BM only on 45-nHA scaffolds, while the ELISA analysis showed that OCN was also secreted in BM at early timepoints only in 45-nHA scaffolds. Despite these results suggesting the potential induction of osteogenesis by 45-nHA scaffolds at late time points, this could not be concluded with a high degree of confidence, since the qualitative immunofluorescence observations were not supported by the aforementioned quantitative gene expression profiles. Moreover, OCN secretion in 45-nHA scaffolds measured by ELISA did not increase at later timepoints and levelled to other scaffold types. Previous studies have already suggested the lack of osteogenic response of cells to HA in composite scaffolds produced by ME-AM. This has been shown for polymer-HA composite scaffolds with different HA amounts. For example, PCL/PLGA-HA scaffolds with 10 wt% ceramic content in MM, as well as PEOT/PBT-HA scaffolds with 15wt% HA in BM or MM, did not enhance osteogenic differentiation of rabbit MSCs and hMSCs, respectively, when compared to pure polymer scaffolds.^{39, 60} Similarly, PLA-HA scaffolds with 50 wt% HA seeded with hMSCs did not show any differences in gene expression compared to bare polymeric scaffolds when cultured in BM.⁶⁸ Consistently, our results also demonstrated that increasing the nHA-to-copolymer ratio did not enhance the osteogenic potential of the scaffolds. We hypothesize that this is due to the suboptimal ion exchange dynamics between the scaffold and the medium. On one hand, we observed that upon incubation in SPS, scaffolds were able to release Ca and P from the nHA scaffolds, at the rate of 10 to 200 μM every 3-4 days. On the other hand, ICP-MS measurements revealed overall Ca and P depletion from media, as previously reported for HA or HA-polymer composite scaffolds incubated in cell culture medium.⁶⁸⁻⁷⁰ Therefore, it is plausible that the slow release dynamics, compared to the fast (re)precipitation events, led to insufficient Ca concentration in the medium so to affect Ca signaling pathways involved in osteogenesis. Previous reports have shown that Ca and P salts added to culture media⁷¹⁻⁷³ or released from polymeric scaffolds,⁷⁴⁻⁷⁵ at concentrations in the mM range, affect Ca and P signaling pathways involved in osteogenesis, leading to gene upregulation and matrix mineralization. Accordingly, scaffolds containing more soluble CaPs (e.g. biphasic CaP (BCP) or TCP) and, therefore, more optimum ion exchange dynamics, have shown to outperform HA based scaffolds, both in *in vitro* and *in vivo* scenarios.^{13, 69, 76-77}

Ca and P depletion from media and SEM images of scaffolds surface after incubation in cell culture media and in SBF, confirmed CaP (re)precipitation on the scaffolds filaments surface. This was due to the presence of nHA particles on the filaments surface, negatively charged at physiological pH, acting as nucleation sites and triggering electrostatic interactions with the Ca^{2+} cations in the media, which can accumulate on the surface of scaffold forming a positively charged Ca-rich layer and stimulate phosphate anions accumulation.⁷⁸ This sequential process ultimately leads to the formation of a crystal phase of apatite, which *in vivo* has shown to promote the co-precipitation of endogenous proteins (e.g. BMPs), ultimately triggering undifferentiated cells to commit to the osteogenic lineage.^{13, 40} This information, together with the lack of osteogenic differentiation in BM, suggests that the ARS staining in 20-nHA and 45 n-HA scaffolds after 35 days of culture in BM was not an hMSCs induced effect, but rather the consequence of a mineral phase (Ca carbonate or CaP) precipitating from media onto the scaffolds' nHA. Notably, since both 45-nHA cell seeded and non-cell seeded scaffolds depleted BM equally of Ca, and MM of Ca and P, and yet non-cell seeded scaffolds presented lower ARS staining and thinner ions layer adsorption, it is hypothesized that the ECM produced by cells on the nHA scaffolds helped to further stabilize the mineral layers, respectively. While 0-nHA scaffolds demonstrated depletion of Ca from media over the culture period, they did not show any ARS staining in BM, or CaP layer formation in SBF. This is likely due to the lack of nucleation sites for CaP precipitation, and the weaker interaction of Ca with PEOT/PBT in 0-nHA scaffolds, leading to an unstable adsorption, in contrast to the strong Ca-nHA interactions on 45-nHA scaffolds. This hypothesis is supported by previous research on PEOT/PBT copolymers calcification, showing that efficient Ca-PEO complexations only occurred in copolymers with higher PEO molecular weight and higher PEOT-PBT ratios, compared to the one used in this study.^{33, 79} Consequently, it is plausible that CaP precipitated in the cell culture media.

To better elucidate the osteogenic property of the nHA containing scaffolds, longer *in vitro* culture times could be considered. Alternatively, enhancing the exposure of HA to the surface of the scaffold could potentially improve the ion exchange dynamics between scaffolds and medium and, therefore, improve the osteogenic potential of the nHA scaffolds. To do this, ME-AM scaffolds surface erosion using NaOH,^{77, 80-81} as well as bare polymeric scaffolds coating with HA by ultrasonication,⁸² or pre-calcification by immersion in an SBF solution,⁸³ have been proposed. While these studies commonly suggest the correlation of osteogenic genes upregulation or increased matrix mineralization to the

enhanced HA exposure, we believe that careful investigations are still required, as using such surface optimization methods have also shown to dramatically change the surface roughness of the scaffolds, making it hard to decouple the effect of HA bioactivity and surface roughness, as the latter is known to affect osteogenic differentiation significantly.⁸⁴

CONCLUSION

Due to a combination of interconnected macroporosity, tunable biodegradability, optimal mechanical properties and bioactivity, 3D polymer-HA composite scaffolds prepared by ME-AM are considered important candidates, when intending to regenerate critical sized bone defects. The aim of this study was the fabrication of highly loaded PEOT/PBT-nHA composite scaffolds with up to 45 wt% nHA using ME-AM and the assessment of their mechanical and in depth biological performance as a function of the nHA content. 45-nHA scaffolds presented significantly enhanced compressive modulus, compared to bare PEOT/PBT copolymer scaffolds, which lied in the range of cancellous bone mechanical properties. In terms of cell behavior, hMSCs were able to differentiate into the osteogenic lineage in all scaffolds regardless of the HA content in MM. Although no differences were observed in osteogenic differentiation at the gene and protein level, increased matrix mineralization was observed on 45-nHA scaffolds compared to 0-nHA and 20-nHA scaffolds. While no osteogenic differentiation of cells was observed in BM, the observed ARS signal in 45-nHA scaffolds in BM suggested the precipitation of a CaP layer from Ca and P ions present in the cell culture media. Since such a CaP layer was also formed upon immersion in SBF, 45-nHA scaffolds are thought to potentially serve as osteoconductive substrates in an *in vivo* situation, favoring osteoblasts adhesion and proliferation. Overall, our results suggest the enhanced mechanical properties of the presented highly loaded composite scaffolds (45-nHA) and their ability to reprecipitate a CaP layer, thus supporting their *in vivo* applicability. Yet, future research and further optimization of polymer-HA composite scaffolds prepared by ME-AM for stimulating bone regeneration, as well as on the validation of their performance through *in vivo* studies are needed.

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SUPPLEMENTARY FIGURES

Figure S1. (A) Image of a 45-nHA pellet obtained after the solvent blending of nHA and PEOT/PBT process. The top side of the pellets is marked as side A and the cross section of the pellet as side B. Scale bar: 1 mm. (B) BSEM micrographs of the side B of 20-nHA pellets, and the side A and B of 45-nHA pellets at two different magnifications (1000x and 5000x). Scale bars top row 50 μ m, lower row 10 μ m.

Figure S2. (A) Elemental analysis spectrum obtained with EDS and corresponding elements weight and atomic % of a $\sim 80 \times 80 \mu\text{m}^2$ area from a 0-nHA and 45-nHA surface filament and 45-nHA cross section. Stereomicroscopy images of a- prepared 0-nHA, 20-nHA and 45-nHA scaffolds (A) before ARS staining and (B) after ARS staining.

Figure S3. nHA aggregates size distribution on 45-nHA scaffolds.

Figure S4. (A) Cylindrical 0-nHA and (B) 45-nHA scaffolds with 15 mm height and 4 mm diameter. Scale bars 1 mm. (C, D) Mechanical properties in compression of 0-nHA and 45-nHA scaffolds.

Figure S5. Gene expression fold change of hMSCs cultured on scaffolds with different nHA concentrations for 7 days in BM. Normalized to 0-nHA d7 BM. Data presented as average $\pm$ s.d., and statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$; * $p < 0.05$; ** $p < 0.01$)

Figure S6. Stereomicroscopy images of scaffolds stained with ARS after 35 days of culture in BM or MM (7 days in BM followed by 28 days in MM). (A) High magnification images of scaffolds cross sections. Scale bar 500 μ m. (B) Images of scaffolds outer surface of the scaffolds. Scale bar 1 mm.

Figure S7. Fluorescent images of hMSCs (F-actin, green) on scaffolds cross sections after 35 days of culture in BM or MM. Scale bar 1 mm.

Figure S8. SEM micrographs of scaffolds cross sections after 35 days of culture in BM or MM. Scale bars 1 mm.

Figure S9. Elemental analysis spectrum obtained with EDS and corresponding elements weight and atomic % of a $\sim 80 \times 80 \mu\text{m}^2$ area of the ECM formed in 0-nHA and 45-nHA scaffolds after 35 days in culture in BM or MM.

Figure S10. Ca ions release over time from nHA composite scaffolds upon immersion in SPS, measured by ICP-MS.

Figure S11. ICP-MS measurements of the calcium (Ca) and phosphorous (P) concentrations in cell culture medium after 7, 14, 21, and 35 days of culture on 0-nHA, 20-nHA and 45-nHA scaffolds with and without cells. The dash lines represent fresh medium concentrations. Data presented as average $\pm$ s.d.

Figure S12. Representative SEM micrographs of 0-nHA, 20-nHA and 45-nHA non-cell seeded and cell seeded scaffolds incubated for 35 days in BM or MM. Scale bars 20 μ m.

Figure S13. Elemental analysis spectrum obtained with EDS and corresponding elements weight and atomic % of a $\sim 80 \times 80 \mu\text{m}^2$ area of the surface of filaments in 0-nHA non-cell seeded scaffolds after 35 days in culture in BM or MM.

Chapter 6

Effect of reduced graphene oxide (rGO) compaction degree and concentration on rGO-polymer composites printability and cell interactions

Effect of reduced graphene oxide (rGO) compaction degree and concentration on rGO-polymer composites printability and cell interactions

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Abstract

Graphene derivatives combined with polymers have attracted enormous attention for bone tissue engineering applications. Among others, reduced graphene oxide (rGO) is one of the preferred graphene-based fillers for the preparation of composites via melt compounding, and their further processing into 3D scaffolds, due to its established large-scale production method, thermal stability, and electrical conductivity. In this study, rGO (low bulk density 10g/L) was compacted by densification using a solvent (either acetone or water) prior to melt compounding, to simplify its handling and dosing into a twin-screw extrusion system. The effects of rGO bulk density (medium and high), densification solvent, and rGO concentration (3, 10 and 15% in weight) on rGO dispersion within the composite, electrical conductivity, printability and cell-material interactions were studied. High bulk density rGO (90 g/L) occupied a low volume fraction within polymer composites, offering poor electrical properties but a reproducible printability up to 15 wt% rGO. On the other hand, the volume fraction within the composites of medium bulk density rGO (50 g/L) was higher for a given concentration, enhancing rGO particle interactions and leading to enhanced electrical conductivity, but compromising the printability window. For a given bulk density (50 g/L), rGO densified in water was more compacted and offered poorer dispersability within the polymer than rGO densified in acetone, and resulted in scaffolds

with poor layer bonding or even lack of printability at high rGO percentages. A balance in printability and electrical properties was obtained for composites with medium bulk density rGO densified in acetone. Here, increasing rGO concentration led to more hydrophilic composites with a noticeable increase in protein adsorption. Moreover, scaffolds prepared with such composites presented antimicrobial properties even at low rGO contents (3 wt%). In addition, the viability and proliferation of human mesenchymal stromal cells (hMSCs) was maintained on scaffolds with up to 15% rGO and with enhanced osteogenic differentiation on 3% rGO scaffolds.

INTRODUCTION

Since the isolation of graphene, graphene based materials have been thoroughly exploited for various biomedical applications, and in particular for bone tissue engineering, due to their excellent mechanical, chemical and electrical properties, as well as to their unique ability of directing stem cell differentiation.¹⁻³ Pristine graphene is a one atom thick, hexagonal lattice structure of sp² hybridized carbon atoms, which confer all the aforementioned properties.⁴⁻⁶ Due to high synthesis costs and small scale production of graphene through bottom-up processes, such as chemical vapour deposition,⁷ or by direct exfoliation of graphite, such as micromechanical cleavage or direct sonication,⁸ currently the most promising method for large scale production of graphene-like materials relies on graphene oxide (GO) as starting material.⁹ GO can be easily mass-produced by oxidation of graphite through well-established methods, and used as-prepared or after exfoliation into graphene-like sheets.⁹ In addition, GO possesses oxygen containing functional groups disrupting each carbon plane, including epoxy, carbonyl, ketone and hydroxyl groups, which allows for good water and polymers dispersability and offers the possibility of functionalization with biomolecules to tune its bioactivity, unlike graphene sheets directly obtained from graphite.¹⁰ However, GO is electrically insulating and thermally unstable, requiring at least partial reduction to restore these properties when required for the final application. GO can be reduced by different chemical and thermal procedures, leading to exfoliated wrinkled reduced graphene oxide (rGO) sheets, similar to pristine graphene, but with some oxygen defects and holes on the carbon skeleton.¹¹

In recent years, graphene, GO and rGO have been used as fillers in both natural and synthetic polymer nanocomposites to improve their physicochemical properties for bone tissue engineering applications. Among different types of such composites, including electrospun fibers,¹²⁻¹³ hydrogels,¹⁴⁻¹⁵ or additive manufactured (AM) scaffolds,¹⁶⁻¹⁹ the latter are superior when aiming towards load bearing applications, as they provide higher mechanical properties. Compared to other AM methods, melt extrusion AM (ME-AM) is considered a cost effective and established technique within the tissue engineering field, which enables the processing of a wide range of biocompatible and biodegradable thermoplastic materials, and provides full control on the internal pore architecture of the scaffolds. Accordingly, three main routes have been explored for the production of graphene derivatives-polymer composites to be used by ME-AM: solution blending, melt blending, and *in situ* polymerization.²⁰ While solvent blending is a simple path to obtain a

good filler dispersion, it generally requires the use of expensive and non-environmentally friendly organic solvents, which can potentially stay as residues within the composite matrix.²¹ *In situ* polymerization has shown to offer even a higher level of dispersion due to monomer intercalation between filler layers, followed by polymerization. However, this process can also require the use of solvents and is limited to specific polymer types.²² In spite of not providing the same level of filler dispersion as the aforementioned techniques, melt blending by twin screw compounding holds the most promise for large-scale composite fabrication, due to its lower cost, green production and industrial applicability.²³⁻²⁴ However, some reports have described the formation of air pockets within extruded polymer composite filaments containing GO at high concentrations, due to its thermal instability and *in situ* thermal reduction at the extruding temperature.²⁵ rGO has also offered some challenges when used as a filler, despite its thermal stability. This is due to its very low bulk density and volatility after volume expansion upon thermal reduction, which impedes its free-flowability into melt compounders and can lead to nanoparticle intake by inhalation during handling.²⁶ To overcome this, rGO compaction or densification by dispersion in a solvent and drying,²⁴ pre-coating with polymer particles in solution,²⁷ or the preparation of a highly concentrated masterbatch by solvent blending, prior to melt blending, have been considered.²⁸ Yet, the attention from rGO as filler for ME-AM scaffolds for bone tissue engineering applications has been deviated and, to the best of our knowledge, the vast majority of previous studies have focused only on graphene and GO fillers for this application, despite pristine graphene's low yield production and GO's poor thermal stability and lack of electrical conductivity.^{17, 29-31}

Here, we prepared densified rGO and studied, for the first time, the effect of rGO densification on its dispersion within melt blended rGO- poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT) composites, at various rGO concentrations. While most of previously reported graphene based ME-AM scaffolds contain only up to 3 wt% filler content, and very rarely up to 10 wt%, in this study we investigated the feasibility of preparing composites with up to 15 wt% rGO, and evaluated their conductivity and printability as a function of rGO compaction and concentration. Moreover, the effect of rGO concentration on the material physicochemical properties, in terms of hydrophilicity, protein adsorption, and antimicrobial properties was assessed. To evaluate its application for bone tissue engineering, we further assessed human mesenchymal stromal cells (hMSCs) adhesion, proliferation and osteogenic differentiation on the prepared 3D ME-AM scaffolds.

MATERIALS AND METHODS

rGO synthesis and characterization

GO was synthesized by Abalonix AS according to a proprietary in-house GO manufacturing method, and thermally reduced to obtain partially reduced rGO. During the reduction process, GO was introduced for few seconds into a tubular oven at 600 °C. However, due to cold air flowing through the oven chamber, the real temperature that the material experiences was estimated to be ~ 300 °C. Three different rGO batches were produced following the same protocol. Densified / compacted (the two terms will be used interchangeably) forms of these rGOs (hereafter referred to as d-rGO) were produced by dispersion into a solvent and subsequent drying. rGO was dispersed in acetone at 50 mg/ml and dried at room temperature to obtain d-rGO-B, with a bulk density of ~ 50 g/L. To obtain d-rGO-A, rGO was dispersed in water at 50 mg/ml and dried at 100 °C. The water dispersion and drying process was carried out two times to obtain d-rGO-C with a bulk density of ~ 90g/L. Overall, three different d-rGO materials were obtained: d-rGO-A (densified in water, bulk density 50 g/L), d-rGO-B (densified in acetone, bulk density 50 g/L) and d-rGO-C (densified in water, bulk density 90 g/L).

In order to evaluate the atomic composition of each d-rGO material, X-ray photoelectron spectroscopy (XPS, K-Alpha – Thermo Scientific, US) was used. Moreover, X-ray diffraction (XRD, D8 Advanced, Brukers, Germany) was employed to investigate the different rGO batches crystallinity and their layered structure. In addition, SEM was carried out to characterize the microstructure of the different d-rGO materials.

Composite production and characterization

d-rGO-polymer composites were obtained by compounding individually the various d-rGO materials (d-rGO-A, d-rGO-B and d-rGO-C) with PEOT/PBT (300PEOT55PBT45, pellets form, PEO Mw 300 kDa, PEOT:PBT weight ratio= 55:45, intrinsic viscosity 0.51 dl/g, Polyvation, The Netherlands). Production was carried out in a lab scale co-rotating twin-screw extruder installed in Nadir S.r.l., consisting of a screw profile with 8 zones, 3 interposed kneading sections, and a screw dimensions of 11 mm diameter and 40 length-to-diameter ratio. PEOT/PBT pellets and d-rGO powder were fed at different concentrations (3, 10, 15 wt%) into the main hopper with a volumetric feeder. The screw rotation speed was fixed at 80 rpm, while the barrel temperature was set at 140°C for the

first zone and 145-150°C for the following zones. Resulting composite wires were taken at the die exit, solidified in air and pelletized in a pelletizer machine. These will be referred to as “X% d-rGO-Y” pellets or composites, where “X” refers to the rGO concentration in wt% (3, 10, 15) and “Y” refers to the d-rGO batch (A, B, or C).

TGA measurements of the d-rGO batches and composites were carried out in a vertical thermo-balance, TGA Discovery 55 analyser (Water-TA Instruments). About 11 mg samples were conducted under nitrogen atmosphere (99,999% N₂) with a flow rate of 90 ml/min and using a temperature heating rate of 10 °C/min up to 900 °C. Residual weight percentages at 550 °C extracted from the TGA curves were used to calculate the experimental rGO loading of each composite. The 550 °C temperature was selected from the separate rGO and polymer only TGA's, as the temperature where most of the polymer was burnt off and most of the rGO was still left.

To analyze potential aggregation of d-rGO after the compounding process, pellets of each material were dissolved in chloroform. After mechanical stirring, a sample of each solution was dispensed on a scanning electron microscopy (SEM) sample holder, dried at room temperature, and imaged using SEM (XL-30) operating at a beam voltage of 15 kV and a spot size of 3. Particle size distribution was assessed using the image analysis software ImageJ.

Scaffolds fabrication and characterization

Scaffolds were fabricated via ME-AM. The platform consisted of a custom-made printhead, with separate heating sources for the cartridge (polymer reservoir) and extrusion screw, mounted on a three- axis stage (Bioscaffolder, Gesim).³² Briefly, the cartridge was filled with the pellets of the composite material to be printed, and scaffolds with a 0-90 architecture, 250 µm fiber diameter, 200 µm layer thickness and 750 µm strand distance (center to center) were fabricated according to parameters in **Table 1**. Cylindrical scaffolds of 4 mm diameter and 4 mm height were punched out from 15x15x4 mm³ manufactured blocks using a biopsy punch and used for further experiments.

Scaffolds morphology and porosity were assessed using a stereomicroscope (Nikon SMZ25). Scaffolds surface roughness was examined using SEM, operating at 10 kV, and a spot size of 3. The electrical conductivity of extruded filaments (length 6 cm, diameter 340 µm and 800 µm) of each material was measured using a digital multimeter (maximum measurable resistance: 2000 MΩ). To do this, filaments with a known diameter and length

were connected to the multimeter electrodes with conductive silver paste (Chemtronics Silver Conductive Adhesive Epoxy) to ensure good contact. The volume electrical conductivity was calculated by **Equation 1**.

Cell studies and antimicrobial analysis was performed only on the d-rGO-B scaffolds due to their superior combined properties for the final application, in terms of processability and electrical conductivity, compared to the d-rGO-A and d-rGO-C composites.

Table 1. Fabrication parameters of d-rGO-polymer composite scaffolds

Scaffold	T reservoir (° C)	T screw (° C)	Pressure cartridge (bar)	Screw rotation speed (rpm)	Translation speed (mm/s)
3% d-rGO-A	195	200	8	60	15
10% d-rGO-A	220	220	8	60	15-20
15% d-rGO-A	not extrudable				
3% d-rGO-B	195	200	8	60	15
10% d-rGO-B	200	210	8	40	15-20
15% d-rGO-B	200	210	8	60	15-20
3% d-rGO-C	200	205	8	60	15
10% d-rGO-C	200	205	8	60	15
15% d-rGO-C	200	210	8	60	15-20

Equation 1. Volume electrical conductivity (σ) formula, where l is the filament length (6 cm), R is the electrical resistance and A is the filament cross sectional area (diameter 340 μm and 850 μm).

$$\sigma = \frac{l}{R \cdot A}$$

Films preparation and characterization

2D films were prepared from around 60 milligrams of (3, 10, 15)% rGO-B pellets and PEOT/PBT pellets, which were molten at 190-210 °C and pressed with a coverslip against a Teflon sheet to obtain 14 mm diameter, ~300 μm thickness films. Static contact angle was measured on these films using the sessile drop method. To do this, a 4 μl water droplet was placed on top of the films by an automatic syringe dispenser (Krüss DSA25S). 20 seconds after droplet formation, the contact angle was calculated automatically by the device's software using the Laplace–Young curve fitting.

Protein adsorption to the films was analysed by incubation in a bovine serum albumin-FITC protein solution (1mg/ml, Sigma Aldrich) overnight at 37 °C. After washing in PBS, films were blotted in an adsorbent paper and incubated for 2h at RT in a 1% SDS solution. Supernatants fluorescence was measured at excitation/emission = 495 nm/519 nm. Cell studies and antimicrobial analysis was performed only on the d-rGO-B films, due to their superior combined properties for the final application, in terms of processability and electrical conductivity, compared to the d-rGO-A and d-rGO-C composites.

Antibacterial activity

The antibacterial activity of (3, 10, 15)% d-rGO-B scaffolds and films against *P. aeruginosa* (CECT 116) and *S. epidermidis* (CECT 231) was evaluated by means of the flask shake test method. Briefly, scaffolds were immersed in concentrated bacterial suspensions (10^6 CFU·L⁻¹) in nutrient broth (1:500) for 24 h at 37 °C. After incubation, the number of viable bacteria present in the suspensions was measured by placing aliquots of the suspensions in sterile petri dishes with molten nutrient agar per triplicate and swirled gently. The petri dishes were incubated at 37 °C for 24 h, after which the colonies present on the plates were counted. Values are reported as the log base 10 reduction (R), calculated as the difference in the log base 10 of the viable cell counts found on a suspension that has not been in contact with the scaffolds and the counts on a suspension that has been in contact with the scaffolds.

In addition, the antibacterial activity of (3, 10, 15)% d-rGO-B scaffolds was evaluated by means of the Agar disk-diffusion method. For that, scaffolds were incubated for 3 days in 1 ml PBS. Mueller- Hinton agar plates were spread with a standardized inoculum of *P. aeruginosa* and *S. epidermidis*. At each timepoint (24h, 48h, 72h), 20 µl of each of the scaffolds' supernatant was collected and used to impregnate commercial paper discs, which were subsequently placed on the agar surface. The agar plates were incubated under 37 °C during 18-24 hours. After incubation, zones of growth inhibition (ZOI) around each of the discs (including disc diameter) were measured to the nearest millimeter. For reporting, the disc diameter was subtracted.

Cell seeding and culture

HMSCs isolated from bone marrow were purchased from Texas A&M Health Science Center, College of Medicine, Institute for Regenerative Medicine (Donor d8011L, female,

age 22). Cryopreserved vials at passage 3 or 4 were plated at a density of 1000 cells $\times$ cm⁻² in tissue culture flasks and expanded until approximately 80 % confluency in complete media (CM). CM was composed of α MEM with Glutamax and no nucleosides (Gibco) supplemented with 10% fetal bovine serum (FBS), without penicillin-streptomycin (PenStrep) at 37 °C and 5% CO₂.

Cell seeding on 2D films

PEOT/PBT and (3, 10, 15)% d-rGO-B films were disinfected in 70% ethanol for 20 min, washed 3 times with PBS, and fixed on the bottom of well-plates with the help of biocompatible O-rings (Eriks, 10023241). Prior to cell seeding, films were incubated in CM for 2h at 37 °C and 5% CO₂ for protein adhesion. Trypsinized hMSCs were re-suspended in fresh media and seeded at 2,500 cells per film. HMSCs were cultured on films for 7 days, and media was replaced every 2 or 3 days.

Cell seeding on scaffolds

For cell attachment experiments, PEOT/PBT and (3, 10, 15)% d-rGO-B scaffolds were disinfected in 70% ethanol and incubated in CM for 2h at 37 °C and 5% CO₂ to allow protein attachment. Subsequently, scaffolds were blotted on top of a sterile filter paper and placed in the wells of a non-treated wellplate. Trypsinized hMSCs were resuspended in a dextran solution (500 kDa, Farmacosmos) (10 wt% dextran in CM), to achieve uniform cell distribution,³³ and were seeded at a density of 2 $\times$ 10⁵ cells with 37 μ l of CM per scaffold. After 4h incubation for cell attachment, scaffolds were transferred to new wells containing 1.5 ml of basic media (BM) (CM supplemented with 200 μ M L-Ascorbic acid 2-phosphate). BM was replaced after 24h and every 2 or 3 days from then on. Scaffolds were cultured for 7 days. For osteogenic differentiation experiments, PEOT/PBT and 3% d-rGO-B scaffolds were seeded as mentioned before, and cultured for 7 days in BM, after which scaffolds were further cultured either in BM or in mineralization media (MM), consisting of BM supplemented with dexamethasone (10 nM) (Sigma-Aldrich) and β -glycerophosphate (10 mM) (Sigma-Aldrich) for another 28 days.

Cells imaging on films and scaffolds

Films and scaffolds seeded and cultured with cells were fixed in 4% paraformaldehyde and incubated for 30 min in Triton-X 100 (0.1 v%). Then, cell cytoskeleton and nuclei were

stained with 488 Alexa Fluor Phalloidin (Thermo Fisher Scientific, 1:75 dilution in PBS, 1h at RT) and DAPI (0.1 µg/ml in PBS, 15 min), respectively.

Biochemical assays

Alkaline Phosphatase (ALP) activity

To measure ALP activity, scaffolds were collected after 14 and 35 days of culture, freeze-thawed 3 times and incubated for 1h at RT in a cell lysis buffer composed of KH_2PO_4 (0.1 M), K_2HPO_4 (0.1 M), and Triton X-100 (0.1 v%), at pH 7.8. The chemiluminescent substrate for ALP CDP star® ready to use reagent (Roche) was added to the cell lysate at a 1:4 ratio, and luminescence was measured using a spectrophotometer. Remaining cell lysates were kept for DNA quantification. ALP values were reported normalized to DNA content.

DNA assay

DNA assay was performed on cells cultured on films and scaffolds using the CyQUANT cell proliferation assay kit (Thermo Fisher Scientific). Lysed scaffolds after ALP assay and frozen films after collection were incubated overnight at 56 °C in Proteinase K solution (1 mg×ml⁻¹ Proteinase K (Sigma-Aldrich) in Tris/EDTA buffer) for matrix degradation and cell lysis. After three freeze-thawing cycles, samples were incubated for 1h at RT with a 20X diluted lysis buffer from the kit containing RNase A (1:500) to degrade cellular RNA. Subsequently, the fluorescent dye provided by the kit (1:1) was added and fluorescence was measured after 15 min incubation (emission/excitation = 520/480 nm). DNA concentrations were calculated from a DNA standard curve.

Alizarin red S (ARS) staining and quantification

Calcium deposits on scaffolds cultured for 35 days in BM and MM were stained by ARS (60 mM, pH 4.1 - 4.3, 20 min incubation at RT) and visualized using a stereomicroscope (Nikon SMZ25). After imaging, ARS was extracted and quantified. To do this, scaffolds were incubated for 1h at RT with 30 v% acetic acid, followed by 10 min incubation at 85 °C. After a centrifugation step at 20,000 rcf for 10 min, ammonium hydroxide (5 M) was added to the supernatants to bring the pH to 4.2. The absorbance was measured at 405 nm using a spectrophotometer. Concentration of ARS was calculated from an ARS standard curve and the values were normalized to DNA content.

Statistical analysis

All data is shown as average with error bars indicating the standard deviation of at least three replicates. Analysis of statistics was conducted with GraphPad Prism (version 8.0.1). A t-test analysis, or a one-way or two-way analysis of variance (ANOVA) followed by a Tukey's post-hoc multiple comparison test correcting for multiple comparisons, was performed to evaluate statistical significance.

RESULTS AND DISCUSSION

Effect of densification on resulting d-rGO properties

One of the most attractive methods for dispersing fillers into polymers, in terms of costs, scalability and green production, is melt compounding.³⁴ Due to the low scale separation of exfoliated graphene sheets from bulk graphite, and the low thermal stability of GO, the use of rGO within melt blended polymer-graphene derivatives composites is preferred.³⁵ However, the low bulk density of rGO makes its feeding into compounders very difficult. This low bulk density originates from the exfoliation and volume expansion of GO caused by the CO₂ formed upon the decomposition of oxygen-containing functional groups during the high temperature thermal reduction process.^{24, 26, 28} For this reason, prior to compounding with PEOT/PBT, the as-produced very low bulk density rGO (10 g/L) was densified (d-rGO) to ease its handling and dosing into the twin-screw compounding system, as well as to reduce the nanoparticles' inhalation hazard. In order to investigate the effect of different densification parameters (i.e. densification solvent and final rGO bulk density) on the resulting d-rGO properties, as well as on the compounding and printing processes, two different densification solvents, acetone and water, were used. Previous literature suggests the applicability of these solvents as densification medium.^{24, 36-37} Here, three d-rGO materials were obtained after dispersion in either of these two solvents (**Figure 1**): i) d-rGO-A: densified in water, bulk density 50 g/L (ρ_{medium}), ii) d-rGO-B: densified in acetone, bulk density 50 g/L (ρ_{medium}), and iii) d-rGO-C: densified in water, bulk density 90 g/L (ρ_{high}).

Since each of the d-rGO came from a different rGO source, their elemental composition was analysed to confirm equal C/O ratios and to allow for subsequent comparisons. According to the XPS analysis in **Table S1**, all d-rGO materials were partially reduced, due to the presence of oxygen in their composition. The C/O ratios were maintained between

Figure 1. Schematic of the polymer-rGO composite scaffold fabrication route. The low bulk density as-prepared rGO was densified (d-rGO), and three different d-rGO types were obtained: d-rGO-A (densified in acetone, medium bulk density), d-rGO-B (densified in water, medium bulk density) and d-rGO-C (densified in water, high bulk density). Each of the d-rGO was melt blended via twin screw extrusion with PEOT/PBT to obtain composites at three different d-rGO concentrations (3, 10 and 15 wt%). Each of the composites was used to fabricate scaffolds via melt extrusion AM.

5 and 6 in all rGO batches, verifying a consistent reduction degree among batches. Moreover, all batches presented traces of N, Al, Si, S, Cl or Fe lower than 1 atomic%. Impurities of sulfur and nitrogen were likely to be due to the covalently bonded sulfates and nitrates during the Hummers method to produce GO.³⁸⁻³⁹ Metallic contaminations are thought to derive from the starting graphitic material itself or the synthesis process.⁴⁰⁻⁴¹ The XRD patterns also showed no differences in the crystalline structure among the different d-rGO's (**Figure S1**). All diffractograms showed a (002) diffraction peak at 24 degrees, corresponding to rGO materials with an interlayer distance of 0.37 nm, and a (100) peak at ~ 42 degrees.^{38, 42} Upon calculation of the full width at half maximum (FWHM) of the (002) peaks, the same crystallite size among the different d-rGO's was confirmed (**Figure S1B**). The low intensity and broadness this peak revealed the amorphous structure and the short-range order of the d-rGO's.⁴³⁻⁴⁴ Other peaks at ~28-33 degrees, which correspond to crystalline mineral impurities, were also appreciated in some the diffractograms.⁴⁵ In addition to these, some spectra showed low intensity sharp

peaks at ~ 26.5 degrees, which correspond to traces of unoxidized graphite. Overall, all d-rGO display similar XRD diffractograms, suggesting that the densification step did not have an impact on the reduction degree, nanocrystalline structure or crystallite size of the samples. However, it is believed that the high porosity of as-prepared rGO created due to volume expansion and CO₂ formation during the thermal reduction process, is reduced during densification, leading to a more compacted and aggregated material. rGO aggregation has been previously reported during the complete removal of water after an aqueous GO reduction process, or after drying rGO in an acetone solution.^{24, 46} Furthermore, previous studies have revealed that re-dispersion of rGO in water or ethanol followed by a drying step, lead to agglomeration of the powder and a shrinkage of their macroporous and mesoporous structure.³⁷ For these reasons, we hypothesize that d-rGO-C, with the highest bulk density, was the highest compacted material after solvent evaporation. This was confirmed by SEM, where both d-rGO-B and d-rGO-C displayed particles aggregating together, but d-rGO-C clearly revealed a less porous and more compacted structure at higher magnifications (**Figure S2**). Although possessing the same bulk density, differences among d-rGO-A and d-rGO-B were also expected, since they were densified in different solvents. Previous literature suggests that the shrinkage of rGO pore size after drying aqueous solutions is more pronounced compared to that of rGO dried in solvents with lower polarity, surface tension, and wettability, such as ethanol or, in our case, acetone.³⁷ This is due to the better interaction of water with rGO, which exerts a higher capillary force upon evaporation, causing a significant decrease in the macro- and mesopore volume. For this reason, d-rGO-A densified in water is hypothesized to be a more compacted material than d-rGO-B. Therefore, the three different d-rGO could be classified according to their compaction degree, being d-rGO-C the most compacted (densified in water, bulk density 90 g/L), followed by the medium compacted d-rGO-A (densified in water, bulk density 50 g/L), and the low compacted d-rGO-B (densified in acetone, bulk density 50 g/L).

Effect of d-rGO densification on melt compounding

After densification, each of the d-rGO materials was melt compounded with PEOT/PBT to form PEOT/PBT-d-rGO composites (**Figure 1**). TGA measurements of the experimental d-rGO loading on each of the composites suggested the reproducibility of the melt compounding process regardless the d-rGO used, for all d-rGO concentrations (**Table 2**, **Figure S3**). Yet, it is worth noting that when d-rGO-B was blended at high concentrations

(15 wt%), the experimental values deviated slightly from the theoretical ones, and the composites contained lower d-rGO amounts than expected. This can be likely due to an inhomogeneous distribution of d-rGO within the pellets and the low amount of sample (a few pellets at most) used for TGA measurements.

Table 2. Experimental loadings (wt%) of each of the composites calculated by TGA.

	3 wt%	10 wt%	15 wt%
d-rGO-A	2.91 ± 0.34	11.22 ± 0.45	17.73 ± 0.28
d-rGO-B	2.18 ± 0.84	11.68 ± 1.03	11.73 ± 1.29
d-rGO-C	4.64 ± 0.28	9.47 ± 6.09	15.01 ± 1.03

The differences in compaction degree mentioned above correlated well with the filler dispersion within the PEOT/PBT-d-rGO composite after compounding. Filler aggregation as an indirect measure of d-rGO dispersion was evaluated by dissolving the composites in chloroform, and visualizing and quantifying the size of d-rGO particles. **Figure 2A** displays representative SEM images of d-rGO within each of the composites. Small particles up to 50 μm were abundant for all conditions (relative frequency $\sim 80\%$) (**Figure 2A, B**). In addition, big d-rGO aggregates were also observed mainly in d-rGO-A composites (ρ_{medium}), at all d-rGO concentrations, and on d-rGO-B composites (ρ_{medium}) at high concentrations (10% and 15%). Interestingly, d-rGO-C composites (ρ_{high}) displayed much lower d-rGO aggregation, as revealed by the SEM images and the lower number of $> 200\ \mu\text{m}$ aggregates compared to d-rGO-A and d-rGO-B composites (**Figure 2B**). Among all, the d-rGO-A composites displayed the largest number of $> 200\ \mu\text{m}$ aggregates, especially at 3% and 15% d-rGO. It is believed that the high bulk density d-rGO-C occupied a smaller volume fraction within the composite at a given wt% compared to d-rGO-A and d-rGO-B, even for high d-rGO concentrations. This avoided particle interaction within the composites and reduced the number of visible aggregates. On the other hand, due to the lower bulk density of d-rGO-A and d-rGO-B, these were able to reach a critical volume occupancy within the composites at much lower concentrations (wt%). This increased the probability of d-rGO particles interaction, and the formation of particles overlaps into bigger agglomerates, by hydrophobic interactions.^{10, 47-48} In addition, due to the lower porosity and higher compaction degree of d-rGO-A compared to d-rGO-B, filler exfoliation and dispersion within the d-rGO-A composites was more difficult, leading to the visualization

of a larger number of aggregates. Compared to melt compounding, other composite production methods such as solvent blending or *in situ* polymerization, have shown to ensure more uniform d-rGO dispersions.^{24, 49} Moreover, several reports have suggested the preparation by solvent blending of rGO-polymer masterbatches, to be used as starting material to prepare other rGO composites by melt compounding.^{28, 50-51} This approach has shown to lead to better rGO dispersion, as it skips the densification or compaction step. However, it involves the use of organic solvents in the process, which is not desirable for biological applications, as the solvents can remain as residues within the composite matrix.

Figure 2. (A) Representative SEM micrographs of d-rGO particles and aggregates after dissolving each composite in chloroform. Scale bars 200 μm . Inserts correspond to high magnification images. Scale bars 50 μm . (B) Size distribution of d-rGO particles and aggregates measured from SEM micrographs in (A).

Notably, d-rGO volume occupancy and dispersion within the composite also correlated well with the electrical properties of extruded composite filaments, as presented in **Table 3**. d-rGO-C extruded filaments' high resistance (out of the device range) impeded the report of their conductivity values at any d-rGO concentration. In case of the d-rGO-A and d-rGO-B filaments, their electrical conductivity was measurable from 10% d-rGO, and they were more conductive with increasing d-rGO content, as previously reported.²⁸ Interestingly, the conductivity of 10% d-rGO-A filaments was 35-fold and 3-fold higher compared to 10% d-rGO-B and 15% d-rGO-B, respectively, suggesting the lower percolation threshold of the d-rGO-A composites. As 15% d-rGO-A was not extrudable, no filament was obtained for this measurement. It is evident that the high volume fraction and poor dispersability of d-rGO-A within the composites helped to form a more connected d-rGO network leading to conductive composites at lower loadings than the less compacted and better dispersed d-rGO-B. On the other hand, the lower volume fraction of d-rGO-C within the composites could not create such conductive pathways and electrical conductivity was not measurable.^{24, 46, 52-53}

Table 3. Conductivity values of extruded filaments from each composite.

	Conductivity 3% (mS/m)	Conductivity 10% (mS/m)	Conductivity 15% (mS/m)
d-rGO-A	< 0.32	13.8 ± 1.1	n.a.
d-rGO-B	< 0.32	0.44 ± 0.1	5.4 ± 1.2
d-rGO-C	< 0.32	< 0.32	< 0.32

Effect of d-rGO compaction on composite printability

The d-rGO volume occupancy, compaction degree and dispersion also influenced composites printability by ME-AM (**Figure 3**). Interestingly, due to the large volume fraction and high frequency of large d-rGO aggregates in the 15% d-rGO-A composites, as shown in **Figure 2**, this material remained a paste-like material rather than a melt, even at 220 °C, and was not printable within the operational limits of the AM device (in terms of printing temperature and cartridge pressure). Reducing the filler loading down to 3 wt% made it possible to produce scaffolds with a fiber diameter of 250 µm (**Figure 3** and **Figure S4**). The fabrication of 10% d-rGO-A scaffolds was challenging, since lack of bonding between layers prevented their stability and handling, in spite of printing at the highest operational

temperature (220 °C) and increasing the fiber diameter (up to 340 μm) for promoting a bigger area of contact in between layers (**Figure 3** and **Figure S5**). On the other hand, scaffolds with a fiber diameter of 250 μm were obtained with all d-rGO-B composites, including 10% and 15% d-rGO-B (**Figure 3**, **Figure S4**). The printability and good interlayer bonding of these materials, compared to d-rGO-A, was possible due to the increased filler dispersion within the d-rGO-B composites and the lower number of aggregates compared to d-rGO-A (**Figure 2**). These parameters might have led to a lower viscosity of the 10% d-rGO-B melt compared to the 10% d-rGO-A melt at a given temperature, allowing an enhanced interlayer contact and bonding.⁵⁴ Printing temperature was adjusted for each d-rGO-A composite, ranging between 195 °C and 210 °C, since viscosity of the melt was found to increase with increasing d-rGO concentration for all composites.⁵⁵ Notably, while 3 and 10 % d-rGO-B scaffolds fabrication was reproducible in terms of resulting filament morphology and z-porosity, 15% d-rGO-B scaffolds lateral porosity did not always remain constant across the scaffold height (**Figure 3**). This was possibly due to sudden viscosity changes during the extrusion process, probably caused by the discontinuous presence of d-rGO-B aggregates changing the flow of the molten composite and, therefore, causing filaments sagging at some points during the printing process. Interestingly, d-rGO-C scaffolds printing was very stable, and scaffolds with optimum lateral porosity were obtained for all d-rGO concentrations (**Figure 3**). Looking at d-rGO-C scaffolds filament cross section by SEM (**Figure S6**), it can be observed that PEOT/PBT (showing a flowy appearance) occupies much larger volume than d-rGO particles (with a grainy and spiky appearance), which are present as dispersed islands within a polymer matrix for all d-rGO concentrations, making these materials relatively easy to print. On the other hand, d-rGO dominantly occupied the d-rGO-A and d-rGO-B scaffolds filaments volume, especially on 10% and 15% d-rGO scaffolds, leading to difficulties when printing. Overall, these results suggest that d-rGO bulk density, and therefore volume fraction, as well as d-rGO densification solvent, which dictates compaction and dispersability degree, plays key roles in the maximum filler loading to process composites into ME-AM scaffolds. While previous studies have reported ME-AM scaffolds production only up to 10 wt% pristine graphene or rGO,^{19, 31} we were able to produce scaffolds with up to 15 wt% d-rGO. Despite the apparent lower rGO content in these studies, equivalent volume percentages or graphene densities would be required to make fair comparisons with the composites studied herein. Nevertheless, to the best of our knowledge, solvent blending and direct ink writing (solvent-based printing) are the only

described manufacturing methods for scaffolds with higher graphene derivatives concentrations (50-75 wt% graphene), which both suffer from requiring the application of organic solvents.^{18, 53}

Figure 3. Stereomicroscopy images of 3D ME-AM scaffolds cross sections and top view (insets) obtained using each of the d-rGO composites, depicting scaffolds morphology and interconnected porosity. Scale bars 1mm.

The ME-AM scaffolds microscale surface roughness was analysed by SEM (**Figure 4**). As a general observation, the qualitative scaffold surface roughness increased with increasing d-rGO content for all scaffolds types, given by the irregular protuberances formed by the underlying rGO particles. Interestingly, the surface of 3% d-rGO scaffolds remained rather

smooth, regardless of the d-rGO type (**Figure 4**). This is in agreement with previous reports, in which the surface of ME-AM scaffolds containing less than 5 wt% graphene (or derivatives) did not present changes in their surface microroughness compared with bare polymeric scaffolds.⁵⁶ In the case of 10% and 15% d-rGO scaffolds, the degree of roughness was influenced by the type of d-rGO, i.e. its bulk density, compaction degree and volume fraction within the filament, for a given d-rGO concentration. In this regard, 10% d-rGO-A presented higher roughness than 10% d-rGO-B, and both higher than 10% d-rGO-C. Similarly, 15% d-rGO-B scaffold surface was much rougher than that of 15% d-rGO-C scaffolds, where only few wrinkles raised on the surface. This is likely due to the much lower volume fraction occupied by the highly compacted d-rGO-C particles decreasing their probability of populating the filaments surface.

Figure 4. Representative SEM micrographs of composite scaffolds depicting the scaffold's surface roughness. Scale bars 50 μm .

Effect of d-rGO concentration on wettability and protein adsorption

Prior to assessing the interaction of cells with the ME-AM scaffolds, different material properties, such as wettability and protein adsorption, as well as cell-material interactions in 2D were evaluated. For this and further studies, only the d-rGO-B composites were taken into consideration as they demonstrated a wider printability window (up to 15 wt%) compared to d-rGO-A composites, and superior electrical conductivity compared to the d-rGO-C composites, properties that are considered relevant for their application in bone tissue regeneration. The static contact angle measurements on composite films (**Figure 5A**), suggested that the material remained rather hydrophobic (contact angle $\sim 80^\circ$) up to 10% d-rGO-B. 15% d-rGO films contact angle was reduced down to $\sim 50^\circ$, denoting the composite hydrophilicity. While the increased hydrophilicity of polymers upon d-rGO-B addition is in agreement with some previous reports,^{19, 57} others have suggested the opposite trend.⁵⁸⁻⁵⁹ The increase in hydrophilicity of 15% d-rGO-B observed within our study might be explained by the d-rGO-B being partially reduced and containing remnant hydrophilic oxygen functional groups, as well as by the presence of larger amounts of d-rGO on the surface of the films with increasing d-rGO-B concentrations. Protein adsorption was assessed by incubating the films in a BSA solution, as a model protein for studying protein-substrate affinity (**Figure 5B**). While 3% d-rGO-B films adsorbed as much protein as PEOT/PBT films, d-rGO-B content over 3% led to a significant increase in protein adsorption to the films. It is believed that rGO interacts with proteins mainly through hydrophobic van der Waals and π - π stacking interactions, due to the carbon structure.⁶⁰⁻⁶¹ Yet, electrostatic interactions and hydrogen bonding are also possible due to the partial presence of oxygen groups on rGO.⁶⁰⁻⁶¹ As **Figure 5B** demonstrates, increasing the d-rGO-B concentration up to 15% led to a 4-fold increase in protein adsorption in 15% d-rGO-B films, likely due to a larger amount of interaction points with proteins, compared to lower d-rGO-B content films.

Surface physicochemical properties, i.e. roughness, hydrophilicity and protein adsorption ability, can regulate cellular behaviour. To evaluate this effect, hMSCs were cultured on films pre-incubated in cell culture medium. Due to a potential partial loss of conformation (i.e. hiding of cell-binding sites) of the adsorbed proteins when adhered to the rGO-based surfaces, previously described in the literature,⁶² as well as gravity acting as the main factor favouring cell attachment on 2D substrates, cell attachment to films did not significantly increase with increasing d-rGO-B content. This was observed upon DNA

quantification at day 1 (**Figure 5C**). Yet, compared to the spread morphology of cells on PEOT/PBT films, hMSCs cultured on d-rGO-B based films, especially at high d-rGO-B content (10 and 15 wt%), showed a spindle-shape, elongated and branched morphology at both day 1 and 7, suggesting higher affinity of hMSCs towards d-rGO-B containing materials, probably due to the aforementioned enhanced protein adsorption (**Figure 5D**). It is also plausible that the ridges and groves on the films rough surface, given by the underlying d-rGO-B, provided contact guidance cues for cells to acquire an elongated morphology.⁶³⁻⁶⁴

Figure 5. Effect of rGO concentration on physicochemical properties and cell-material interactions in 2D. (A) Static contact angle measured on PEOT/PBT and d-rGO-B composite films. (B) Protein adsorption to d-rGO-B films upon incubation in a BSA solution, normalized to protein adsorption to PEOT/PBT films. (C) Quantification of hMSCs attachment to d-rGO-B films after 1 day of culture. (D) Fluorescent microscopy images of hMSCs (F-actin in green, nuclei in blue) cultured on films for 1 and 7 days. Scale bars 100 μm. Data presented as average ± s.d. and statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (n.s. p > 0.05, * p < 0.05, **** p < 0.0001).

Effect of d-rGO concentration on scaffolds antimicrobial activity

The antimicrobial activity of scaffolds was evaluated against relevant Gram - and Gram + bacterial strains in the orthopedic field: *P. aeruginosa* and *S. epidermidis* respectively. Preliminary experiments demonstrated the biocidal activity of d-rGO-B at different concentrations and of composite films when in direct contact with bacteria (**Table S2** and **Table S3**). Importantly, the d-rGO-B ME-AM scaffolds also demonstrated higher antibacterial activity with increasing d-rGO-B concentration within the scaffolds (**Figure 6A**). Interestingly, antibacterial effects were only evident when bacteria were placed in direct contact with the scaffolds, and no biocidal effect of the supernatant, in which scaffolds were incubated up to 3 days (**Figure S7**), was observed. This suggests that reactive oxygen species (ROS)-dependent oxidative stress might have not been the dominant antibacterial mechanism, and that membrane mechanical stress induced by sharp edges or corners of d-rGO-B exposed to the surface of the scaffolds, which act as nano-knives or nano-needles and disrupt bacteria membranes, could have been the prevailing mode of antibacterial action.^{19, 59, 65} Besides, rGO has been shown to produce lower oxidative stress compared to GO, because of the lower amount of oxygen functional groups.⁶⁶ Mechanical stress causing bacterial damage would also explain the increasing antimicrobial activity with increasing d-rGO-B concentration, potentially due to more d-rGO-B exposed to the filaments surface (**Figure 6A**). Furthermore, this mechanism would support scaffolds being more potent against *P. aeruginosa* (Gram -) than *S. epidermidis* (Gram +) (**Figure 6A**). As previously described, this could be attributed to the elongated and curved shape of *P. aeruginosa* being more vulnerable to rGO mechanical damage compared to *S. epidermidis*, with less surface area and a spherical shape.⁶⁷

Due to the conductivity of rGO, charge-transfer oxidative stress can also be considered a main antimicrobial mechanism when bacteria get in contact with the surface of the scaffolds. Here, rGO can act as a conductive bridge over the bacteria lipid bilayer to mediate electron transfer from bacterial intracellular components to the external environment, interrupting the bacteria membrane respiratory chain, and leading to bacteria death.⁶⁵⁻⁶⁶ Overall, this and all the aforementioned antibacterial mechanisms will be a matter of investigation in future studies.

Cell seeding on scaffolds and osteogenic differentiation

In spite of scaffolds displaying antibacterial activity, hMSCs viability was not compromised when seeded on the d-rGO-B composite scaffolds, at any d-rGO-B concentration. As fluorescent microscopy images of cells (F-actin) attached to the scaffolds' filaments in **Figure 6C** depict, cells populated the scaffolds from the beginning of the culture, and showed viable cells characteristics, i.e. a spread or elongated morphology. DNA analysis further confirmed the biocompatibility of the scaffolds up to 15% d-rGO-B, as cell number did not decrease after 7 days of culture, but was maintained rather constant (**Figure 6B**). While various reports have addressed the cyto- or genotoxicity of graphene and GO in contact with different cell lines, ⁶⁸⁻⁷⁰ only a few have tried to understand the biocompatibility of rGO, which greatly differs from the other derivatives in terms crucial factors for biointeractions, such as size (number of layers, available surface area), functional groups (reduction degree) and protein interactions. It has been demonstrated that the cytotoxic effect of GO and partially reduced rGO can be attenuated by proteins in serum binding to their surface, which can reduce the GO/rGO ability to penetrate or physically damage the cell membrane, both in *in vitro* cell cultures and *in vivo*. ⁷¹ In addition, rGO has been found to be, in general, more biocompatible than GO and to provide a reduced inflammatory response upon *in vivo* implantation, ⁷² but still its dose- and size-dependent cytotoxicity towards hMSCs has been shown. ⁷³ Due to the d-rGO-B encapsulated within the polymer matrix and the polymer slow degradation rate such a cytotoxic effect was not evidenced during the culture period addressed here (35 days). ⁷⁴⁻⁷⁵ Future studies on scaffolds degradation kinetics and d-rGO-B release rate will be essential, as polymer degradation will eventually occur over time in an *in vivo* situation, releasing potentially toxic amounts of d-rGO-B.

Considering the aforementioned potential long-term toxicity of highly loaded rGO scaffolds, preliminary osteogenic differentiation studies were performed with hMSCs seeded on 3% d-rGO-B scaffolds, a low concentration that has been commonly studied in previous reports. ^{56, 76-78} hMSCs were able to proliferate both in PEOT/PBT and 3% d-rGO-B scaffolds over the culture period (35 days) especially in MM, likely due to more ECM formation within the scaffolds pores acting as cell growth support, compared to BM (**Figure S8**). As an early osteogenic marker, **Figure 6D** demonstrates the ALP activity of hMSCs scaffolds. For both bare PEOT/PBT and 3 % d-rGO-B containing scaffolds, a peak in ALP activity was shown at day 14, with a significant decrease on day 35, both in BM and MM,

which is in agreement with the osteogenic differentiation progression.⁷⁹ However, no statistical differences on ALP activity were found among the d-rGO-B containing scaffolds and the PEOT/PBT control.

Figure 6. Cell-material interactions in d-rGO-B scaffolds: effect of d-rGO-B concentration. (A) Antibacterial activity of scaffolds against *P. aeruginosa* and *S. epidermidis*. (B) DNA content on scaffolds seeded with hMSCs after 1 and 7 days of culture. (C) Fluorescent microscopy images of hMSCs attachment (F-actin, green) to composite scaffolds cross section. Scale bars 250 μm. (D) ALP activity of hMSCs seeded on PEOT/PBT and 3% d-rGO-B scaffolds after 14 and 35 days cultured in BM or MM. Data presented as average ± s.d. and statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (* p < 0.05, ** p < 0.01). (E) Stereomicroscopy images of PEOT/PBT and 3% d-rGO-B scaffold cross sections stained with ARS after 35 days of culture in MM. Scale bars 1 mm. (F) Quantification of the ARS staining extracted from scaffolds in (E), normalized to cell number. Data presented as average ± s.d. and statistical significance performed using T-test (n.s. p > 0.05).

ECM mineralization was examined at day 35 of culture in BM and MM with ARS, as a late osteogenic marker. Interestingly, both PEOT/PBT and 3% d-rGO-B scaffolds revealed calcium deposition on the ECM matrix produced by cells inside the pores of scaffolds cultured in MM, as depicted by the ARS in **Figure 6E**. In spite of a slight increase in the calcium deposition on 3% d-rGO-B scaffolds, no statistical differences were observed among these and the PEOT/PBT scaffolds (**Figure 6F**). These results imply that 3% d-rGO-B scaffolds support osteogenic differentiation, observed by both early and late stage markers. Yet, these scaffolds seem to not enhance differentiation compared to bare polymeric scaffolds. While this is in disagreement with some previous reports, other studies only point out the enhanced cell adhesion and proliferation ability on composite scaffolds with such low graphene derivatives concentrations (up to 1 wt%).^{16, 29, 76} As previously mentioned, this might very well be correlated to differences in rGO bulk density affecting the final volume fraction of rGO within the scaffolds. It is plausible that the low d-rGO-B concentration used here (3 wt%) did not allow for sufficient exposure of d-rGO-B to the scaffolds filament surface, preventing it to act as a pre-concentration platform for osteogenic inducers (e.g. dexamethasone and β -glycerolphosphate), which is believed to be the origin of the enhanced stem cell differentiation exerted by graphene derivatives.⁸⁰⁻⁸² In this regard, we hypothesize that the increased hydrophilicity and protein adsorption on 10% and 15% d-rGO-B scaffolds, together with their surface microroughness, would induce accelerated osteogenic differentiation, compared to 3% d-rGO-B and PEOT/PBT scaffolds. Nevertheless, future studies will be dedicated to validate this hypothesis, as well as to strike a balance between in vivo biocompatibility and bone formation ability.

CONCLUSION

Due to its very low bulk density, rGO has to be densified prior to melt compounding with a polymer. The aim of this study was to understand the effect of rGO densification parameters (densification solvent and rGO bulk density after densification) on rGO compaction degree and on the resulting melt compounded composites physicochemical properties and printability via ME-AM. The effect of rGO concentration on the materials' physicochemical properties and cell-material interactions in 2D and 3D was also investigated. It was observed that high d-rGO bulk density (90 g/L) correlated with higher d-rGO compaction, which was translated into smaller d-rGO volume fraction for a given d-rGO concentration within the polymer composite, and printability up to 15 wt% d-rGO. On

the other hand, medium bulk density d-rGO (50 g/L) occupied a greater volume within the melt compounded composites, which presented some challenges upon ME-AM at high d-rGO contents (10 and 15 wt%), in terms of loss of filament fidelity, poor layer bonding, or even lack of printability. At a given bulk density (50 g/L), when comparing acetone and water as densification solvent, it was observed that densification in water led to a more compacted d-rGO than d-rGO densified in acetone, and d-rGO was poorly dispersed within the polymer matrix. This led to bigger aggregates formation within the composites, whose connections were able to create conductive pathways within the composites, making these materials the most conductive among all d-rGO types. Composites prepared medium bulk density d-rGO-B densified in acetone were chosen for further characterization, due to a balance of printability and electrical properties. It was observed that increasing d-rGO content led to increasing material hydrophilicity and protein adsorption, as well as to increasing surface roughness due to higher rGO exposure to the surface. Scaffolds prepared with 3, 10 and 15% d-rGO were found to possess increasing antibacterial properties with increased d-rGO content, without affecting hMSCs viability. Notably, 3% d-rGO scaffolds were able to support hMSCs proliferation and osteogenic differentiation. Overall, this study demonstrated that rGO compaction degree and concentration greatly affects composites printability and scaffolds physicochemical and electrical properties. In this regard, a careful selection of the rGO densification parameters has to be made in order to ensure the most adequate properties of the final scaffolds required for bone tissue engineering applications.

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SUPPLEMENTARY FIGURES

Figure S1. (A) XRD diffraction patterns of the three different d-rGO used in this study, each coming from different rGO batches. (B) FWHM of (002) diffraction peaks.

Figure S2. Representative SEM images of d-rGO-B and d-rGO-C at different magnifications, displaying different degrees of compaction. Scale bars 20 μm (top row), and 2 μm (middle and bottom row).

Figure S3. TGA curves of (A) d-rGO batches, (B) PEOT/PBT-d-rGO-A composites, (C) PEOT/PBT-d-rGO-B composites and (D) PEOT/PBT-d-rGO-C composites. For each concentration, two curves, measured from two different samples are represented.

Figure S4. SEM micrographs of 3D ME-AM scaffolds cross sections obtained using each of the d-rGO composites, depicting scaffolds morphology and interconnected porosity. Scale bars 1mm.

Figure S5. Demonstration of poor layer bonding of 10% d-rGO-A upon punching of scaffold using a biopsy puncher, compared to other scaffold types, such as the 15% d-rGO-C scaffold. Scale bars 2 mm.

Figure S6. SEM micrographs of 3D ME-AM scaffolds filaments cross sections cut with a razor blade. Yellow arrows indicate rGO particles within the polymer matrix. Scale bars 50 μm .

Figure S7. (A) Disk diffusion agar test diagram. (B) Antimicrobial activity of d-rGO-B scaffolds against *P. aeruginosa* and *S. epidermidis*, measured through the disk diffusion agar test and reported as ZOI values. (C) Images of representative disk diffusion test plates depicting ZOIs around a disk impregnated with an aliquot of the scaffold supernatant after the initial 24h of incubation.

Figure S8. DNA content progression on PEOT/PBT and 3% d-rGO-B scaffolds over 35 days of culture in BM or MM.

Table S1. Atomic compositions (%) of each of the d-rGO measured by XPS.

	C	O	C/O ratio	N	Al	Si	S	Cl	Fe
d-rGO-A	85.3	14.4	5.8	0.0	0.0	0.2	0.0	0.1	0.0
d-rGO-B	83	16	5.2	0.2	0.0	0.6	0.2	0.2	0.0
d-rGO-C	82	16.5	5.0	0.2	0.2	0.6	0.1	0.4	0.1

Table S2. d-rGO-B antimicrobial activity at different concentrations in contact with *P. aeruginosa* and *S. epidermidis*.

Samples	<i>P. aeruginosa</i> (R)	<i>S. epidermidis</i> (R)
3mg/ml d-rGO-B	> 5.8	> 4.8
10 mg/ml d-rGO-B	> 5.8	> 4.8
15 mg/ml d-rGO-B	> 5.8	> 4.8

Table S3. Antimicrobial activity against *P. aeruginosa* and *S. epidermidis* of PEOT/PBT- d-rGO-B films (0.1 g/mL) containing different d-rGO concentrations.

Samples	<i>P. aeruginosa</i> (R)	<i>S. epidermidis</i> (R)
3% d-rGO-B film	1.3	2.0
10% d-rGO-B film	5.8	4.7
15% d-rGO-B film	6.2	4.7

Chapter 7

3D Additive Manufactured Composite Scaffolds with Antibiotic-loaded Lamellar Fillers for Bone Infection Prevention and Tissue Regeneration

3D Additive Manufactured Composite Scaffolds with Antibiotic-loaded Lamellar Fillers for Bone Infection Prevention and Tissue Regeneration

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Abstract

Bone infections following open bone fracture or implant surgery remain a challenge in the orthopedics field. In order to avoid high doses of systemic drug administration, optimized local antibiotic release from scaffolds is required. 3D additive manufactured (AM) scaffolds made with biodegradable polymers are ideal to support bone healing in non-union scenarios and can be given antimicrobial properties by the incorporation of antibiotics. In this study, ciprofloxacin and gentamicin intercalated in the interlamellar spaces of magnesium aluminum layered double hydroxides (MgAl) and α -zirconium phosphates (ZrP), respectively, are dispersed within a thermoplastic polymer by melt compounding and subsequently processed via high temperature melt extrusion AM (~190 °C) into 3D scaffolds. The inorganic fillers enable a sustained antibiotics release through the polymer matrix, controlled by antibiotics counterions exchange or pH conditions. Importantly, both antibiotics retain their functionality after the manufacturing process at high temperatures, as verified by their activity against both Gram + and Gram - bacterial strains. Moreover, scaffolds loaded with filler-antibiotic do not impair human mesenchymal stromal cells osteogenic differentiation, allowing matrix mineralization and the expression of relevant osteogenic markers. Overall, these results suggest the possibility of fabricating dual functionality 3D scaffolds via high temperature melt extrusion for bone regeneration and infection prevention.

INTRODUCTION

In the orthopedics field, surgical procedures involving fracture stabilizations or implants can develop infections in up to 5% of the cases, while this rate rises up to 50% in open bone fracture scenarios.¹⁻² Bacterial infections can trigger osteomyelitis, an inflammatory reaction in bone that can ultimately lead to bone destruction or osteolysis. Moreover, bacteria can invade and survive within osteoblasts slowing down or preventing new bone formation, or and create biofilms, which present a challenge to the antibiotic therapy, due to reduced antimicrobial susceptibility leading to prolonged infection.³⁻⁵ The standard medical strategy to overcome osteomyelitis involves the debridement of infected bone and soft tissue and the removal of the implant or fixation device, accompanied by the systemic administration of antibiotics from 6 to 16 weeks depending on the severity of the infection.⁶⁻⁷ Typically, high antibiotic concentrations are administered systemically to achieve effective concentrations at the infected site, which can lead to the development of resistant bacterial strains, as well as cell toxicity and adverse effects in the bone regeneration process.⁸ Since the removal of necrotic bone tissue commonly leaves a critical size defects, antibiotic-laden poly(methyl methacrylate) (PMMA) cements or beads were designed to act both as bone spacer and as local antibiotic delivery systems, maximizing target tissue concentrations and minimizing toxicity risks.⁹⁻¹⁰ Despite being one of the gold standard approaches, PMMA cements possess a series of disadvantages including lack of biodegradability, requirement of a second surgery to replace it by other material that allows bone regeneration. Furthermore, PMMA cements present a suboptimal drug elution kinetics, which is associated to a non-reproducible antibiotic-polymer mixing procedure, and to the exothermic polymerization of PMMA. Moreover, they are restricted to heat-stable antibiotics. antibiotics can be incorporated.¹¹⁻¹² With the recent development in the field of biomaterials and bone tissue engineering, a number of alternative natural (i.e. collagen or calcium phosphates) and synthetic (i.e. poly(lactic acid) (PLA) and poly(lactides-co-glycolides) (PLGA)) carriers have been designed to allow biodegradability and the capability of delivering locally a wider variety of antimicrobial agents.¹³⁻¹⁴ These carriers are mainly collagen, chitosan, calcium sulfate, calcium phosphates, demineralized bone, as well as synthetic polymers, such as polyurethanes, polyanhydrides, PLA and PLGA.¹³⁻¹⁴ However, processing these materials into 3D scaffolds using conventional fabrication techniques, such as gas foaming/particulate leaching, freeze-drying or phase separation, can result in constructs that lack reproducibility or the structural integrity and

mechanical properties needed to stabilize a non-union defect, making them also not fully functional candidates for facilitating bone healing following infection management.

In the past decade, additive manufacturing (AM) and, more specifically, melt extrusion (ME)-based techniques (ME-AM), have emerged as one of the most appealing technologies to produce scaffolds for bone tissue engineering.¹⁵ This method allows for the reproducible fabrication of patient-personalized 3D scaffolds, from a variety of biocompatible and biodegradable thermoplastic polymers, with an interconnected pore network, high porosity, and optimum mechanical properties for bone regeneration. Osteoinductive and antibacterial scaffolds have recently been developed by combining AM and drug delivery. Antimicrobial peptides,¹⁶⁻¹⁷ or other antimicrobial compounds, such as silver,¹⁸ bioactive glass,¹⁹ or quaternized chitosan,²⁰⁻²¹ have been incorporated onto the surface of 3D AM scaffolds. These alternative compounds have been used to avoid the use of antibiotics due to their potential cell toxicity and bacterial resistance risks. Yet, the *in vivo* efficacy of these substitutes compared to conventional antibiotic therapies is still unproven. Therefore, as a common strategy, antibiotics have been adsorbed on the surface of AM scaffolds. However, surface adsorption via the immersion of the scaffold in an antibiotic solution has shown a limit retention of the loaded antibiotic, amount that correlates to the scaffold surface area, resulting in poor loading efficiency and leading to the burst release of the antibiotic.²² In order to increase the loading efficiency and allow for a more sustained drug release, different strategies have been explored, such as porogen leaching to create microporosity in the scaffold's filaments, hydrogel coatings, or loading of antibiotics within degradable polymeric microparticles.²²⁻²⁴ The combination of several of these loading methods in the same scaffold has also been considered to allow the incorporation of different antibiotics and to obtain a sequential release profile for a more potent therapy.²⁵ However, in spite of enhancing the scaffold antimicrobial properties, these multi-step post-modification approaches are complex and laborious. Thus, the direct incorporation of the antibiotic within the scaffold material would be desired to simplify the fabrication process, as well as to improve the loading capability and release profile. With the exception of ME of polymers at low temperatures, such as polycaprolactone (PCL),²⁶⁻²⁷ the incorporation of antibiotics in the scaffold material is not frequently investigated due to the sensitivity of bioactive molecules to temperature. This in turn limits the polymer choice, as a large majority of the thermoplastic polymers for biomedical applications require long exposure to high processing temperatures (~175 – 220 °C),²⁸⁻²⁹ which are above the thermal stability limits of most commonly used antibiotics (~ 120 °C, depending

on exposure time and antibiotic type).³⁰ Moreover, most of current delivery systems lack the optimum release kinetics of the cargo. A desirable sustained drug release should ensure that an adequate antibiotic concentrations is delivered on site, above the minimal inhibitory concentration (MIC) values and over a sufficient time to cover the critical window period post-surgery, followed by a sustained release at an effective level for inhibiting the occurrence of a latent infection.³¹⁻³²

Here, we introduce a novel approach that allows the direct fabrication of drug loaded scaffolds via high temperature ME-AM. The composite scaffolds consist of the biocompatible and biodegradable copolymer poly(ethyleneoxideterephthalate)/poly(butylene terephthalate) (PEOT/PBT) and the inorganic layered compounds magnesium aluminum layered double hydroxide (MgAl) or α -Zirconium phosphate (ZrP), which contain antibiotics intercalated within their lamellar structures. Ciprofloxacin (CFX) and gentamicin (GTM), two commonly used antibiotics to treat orthopedic infections, are used as model anionic and cationic molecules to be intercalated within the positively charged MgAl and negatively charged ZrP fillers, respectively. We evaluate the dual functionality of these scaffolds as i) local antibiotic delivery systems against *Staphylococcus epidermidis* (Gram +) and *Pseudomonas aeruginosa* (Gram -), identified as two of the major responsible for implant associated infections,³³ and ii) their potential to support bone tissue formation.

MATERIALS AND METHODS

A list of materials described in this study and the abbreviations used to refer to them can be found in **Table 1**.

Table 1. List of materials

Nomenclature	Description	Nomenclature	Description
CFX	Ciprofloxacin	5, 10, 20 % ZrP-GTM _p	Pellets composed of 5, 10, 20 wt% ZrP-GTM dispersed in PEOT/PBT
GTM	Gentamicin	5, 10, 20 % ZrP-GTM _s	Scaffold composed of 5, 10, 20 wt% ZrP-GTM dispersed in PEOT/PBT

<i>MgAl</i>	Magnesium aluminum layer double hydroxide	5, 10, 20 % <i>ZrP-GTM_f</i>	Film composed of 5, 10, 20 wt% ZrP-GTM dispersed in PEOT/PBT
<i>ZrP</i>	α -Zirconium phosphate lamellar compound	10 % <i>CFX_p</i>	Film composed of 10 wt% CFX dispersed in PEOT/PBT
<i>MgAl-CFX</i>	MgAl filler loaded with CFX	10 % <i>CFX_f</i>	Film composed of 10 wt% CFX dispersed in PEOT/PBT
<i>ZrP-GTM</i>	ZrP filler loaded with GTM	10 % <i>GTM_p</i>	Film composed of 10 wt% GTM dispersed in PEOT/PBT
<i>PEOT/PBT_p</i>	PEOT/PBT pellets	10 % <i>GTM_f</i>	Film composed of 10 wt% GTM dispersed in PEOT/PBT
<i>PEOT/PBT_s</i>	PEOT/PBT scaffolds	10 % <i>MgAl_p</i>	Film composed of 10 wt% MgAl-CO ₃ dispersed in PEOT/PBT
5, 10, 20 % <i>MgAl-CFX_p</i>	Pellets composed of 5, 10, 20 wt% MgAl-CFX dispersed in PEOT/PBT	10 % <i>MgAl_s</i>	Scaffold composed of 10 wt% MgAl-CO ₃ dispersed in PEOT/PBT
5, 10, 20 % <i>MgAl-CFX_s</i>	Scaffold composed of 5, 10, 20 wt% MgAl-CFX dispersed in PEOT/PBT	10 % <i>ZrP_p</i>	Film composed of 10 wt% ZrP-H dispersed in PEOT/PBT
5, 10, 20 % <i>MgAl-CFX_f</i>	Film composed of 5, 10, 20 wt% MgAl-CFX dispersed in PEOT/PBT	10 % <i>ZrP_s</i>	Scaffold composed of 10 wt% ZrP-H dispersed in PEOT/PBT

MgAl synthesis and CFX intercalation

MgAl in nitrate form (MgAl) was obtained adapting the urea method from Costantino et al.³⁴ Solid urea was added to 0.5 M magnesium nitrate hexahydrate (M II) - aluminum nitrate nonahydrate (M III), having molar fraction $M(III)/(M(III) + M(II))$ equal to 0.33, until the molar ratio urea/M (III) reached the value of 6. The mixture was heated at 100 °C for 48 h. The final product was filtered, washed with water and dried in oven at 60 °C. MgAl-CFX was prepared via anionic intercalation. A carbon dioxide free water solution of NaOH 1 M (36 ml) was added to a suspension of CFX (12.2 g) in a hydroalcoholic solution (370 ml, water: EtOH 1:1) in order to obtain a sodium salt solution. Then MgAl (10 g) was added (MgAl:CFX 1:1 molar ratio). The suspension was kept under N₂ and stirred for one day at

RT. The reaction mixture was centrifuged, washed twice with a water-ethanol solution and dried in oven at 45°C.

ZrP-synthesis and GTM intercalation

ZrP synthesis and GTM intercalation was carried out as previously described.³⁵ Briefly, crystalline zirconium phosphate $\text{Zr}(\text{HPO}_4)_2 \cdot \text{H}_2\text{O}$ (ZrP) was obtained by refluxing zirconyl chloride ($\text{ZrOCl}_2 \cdot 8\text{H}_2\text{O}$) in a 10 M phosphoric acid solution for 48 h. The residual solid was centrifuged, washed 3 times with water and dried in oven at 60 °C. A pre-intercalated phase of ZrP with an expanded interlayer distance was obtained via ion exchange with propylamine, followed by HCl mixing to regenerate the acid form. ZrP-GTM was prepared by adding of GTM (18.9 g) to the gel. The mixture was stirred for 24 h at RT until complete exchange of the protons with GTM. The reaction mixture was centrifuged, washed twice with deionized water and dried in the oven at 40 °C.

Composite production

The production of 5, 10, 20 % MgAl-CFX_p, 5, 10, 20 % ZrP-GTM_p, 10 % MgAl_p and 10 % ZrP_p composites was carried out in a lab scale co-rotating twin screw extruder installed in Nadir S.r.l., with a screw diameter of 11 mm and a length-to-diameter ratio (L/D) of 40. The screw profile was composed of 8 zones with three interposed kneading sections. PEOT/PBT pellets (PEO molecular weight = 300 kDa, PEOT:PBT weight ratio= 55:45, intrinsic viscosity 0.51 dl/g, Polyvation, The Netherlands, PEOT/PBT_p) were fed in the main hopper with a volumetric feeder. Using a double inlet, MgAl-CFX or ZrP-GTM powder were fed in main hopper at 5, 10, 20 wt%. The screw rotation speed was fixed at 80 rpm while the barrel temperature was set at 140°C for the first zone and temperatures from 145 to 150°C for following zones. The nanocomposite wire was taken at the die exit, solidified in air and pelletized in a pelletizer machine. The production of 10 % CFX_p and 10 % GTM_p was carried out in co-rotating twin screw extruder by pre-mixing grinded PEOT/PBT_p and CFX or GTM powder (9:1 by weight). The screw rotation speed was fixed at 80 while the barrel temperature was set at 150°C. The nanocomposite wire was taken at the die exit, solidified in air and manually pelletized.

Composite materials characterization

TGA measurements of CFX, GTM, MgAl-CFX, ZrP-GTM, MgAl-CFX_p and ZrP-GTM_p were carried out with an STD Q600 thermal analyser (TA Instruments, USA) in air flow with a heating rate of 10 °C/min up to 800 °C. Inorganic filler content within MgAl-CFX and ZrP-GTM was calculated to be 54 wt% MgAl (46 wt% CFX) and 46 wt% ZrP (54 wt% GTM), respectively. These results in combination with the MgAl-CFX_p and ZrP-GTM_p TGA curves were used to calculate the experimental filler-antibiotic loading of the each polymer composite. Results are presented in **Table S6**.

Particle size distribution was assessed using the image analysis software tool XT Pro v3.2 (Soft Imaging System GmbH).

For high-performance liquid chromatography (HPLC) analysis, a precise amount of sample was weighted. When a polymer composite was analysed, sample was dissolved in chloroform by sonication for 20 min at RT, followed by 10 min centrifugation at 4600 rpm. The supernatant was aspirated and the residual chloroform was allowed to evaporate under N₂. The pelleted MgAl-CFX or ZrP-GTM was treated with a suitable volume of HCl (1 M) or in HCl/KCl (3 M), respectively, for antibiotic extraction. Then, the solution was filtered (pore size 0,2 µm) or centrifuged for filler separation and appropriately diluted. The amount of CFX was assessed by means of a HPLC column (Zorbax SB C18 4,6 x 250 mm, 5 µm) with a flow rate of 0.8 ml min⁻¹, and the eluent consisting of water and 2% acetic acid/acetonitrile at a ratio 84/16 v/v%. In order to assess the uncertainty involved in the samples dilutions, all samples were diluted and analyzed three times. A five point calibration curve was used to quantitatively determine the CFX content.

The amount of GTM was assessed by means of a LC-MS Bruker qTOF Compact. The chromatographic separation was conducted using an Agilent Zorbax SB-C18 column (4.6 mm x 150 mm, 3.5µm) with a flow rate of 0.4 ml/min and the eluent consisting of trifluoroacetic acid (TFA) water (0.2 M) and methanol at a ratio 92/8 v/v%. The mass spectrometer was operated in full scan mode, and for the mass calibration, sodium formate clusters were used. According to literature, the four main isoform components of GTM were separated and quantified by extracting the high resolution mass chromatogram of the selected compounds with an uncertainty of 0.01 a.u.. For the quantification of the total amount of antibiotic loaded into the composites, the sum of all four species was taken into account. In order to assess the uncertainty involved in the samples dilutions, all samples were diluted and analyzed twice.

Scaffolds fabrication and characterization

Scaffolds were fabricated via ME-AM. The platform (Bioscaffolder, Gesim) was equipped with a custom-made print head with two separate heating sources for the cartridge and screw. Briefly, the cartridge was filled with MgAl-CFX_p or ZrP-GTM_p, heated at 185 °C and extruded at 190-195 °C by applying a pressure of 8 bar, an auger screw rotation of 50-60 rpm and a translation speed of 15-20 mm/s. Parameters were adjusted slightly around these values for each material type and composition. 10% MgAl_p and 10% ZrP_p were also used to prepare only filler control scaffolds with the same aforementioned parameters. Similarly, PEOT/PBT_p were heated and extruded at 195 °C by applying a pressure of 4 bar, an auger screw rotation of 30 rpm and a translational speed of 15 mm/s. The scaffolds architecture consisted of a 0-90 pattern, with a 340 µm fiber diameter, 250 µm layer thickness and 850 µm strand distance (center to center), giving an expected x-y porosity of 510 µm and an expected z porosity of 90 µm. Cylindrical scaffolds of 4 mm diameter and 4 mm height were punched out from 15x15x4 mm³ manufactured blocks using a biopsy punch and used for further experiments.

Scaffold morphology and porosity was assessed using a stereomicroscope (Nikon SMZ25). Stereomicroscopy images were used to measure the experimental values of fiber diameter and x-y-z porosity, and these values were inserted in the Landers equation (**Equation 1**) to calculate the average scaffold porosity.³⁶ Presence and distribution of fillers within gold sputter-coated scaffolds' filaments surface and cross section was examined using backscattered scanning electron microscopy (BSEM, XL-30 beam voltage 20 kV, spot size 5). Scanning electron microscopy (SEM) operating at 25 kV coupled with energy dispersive X-ray spectroscopy (EDS) was used to observe the chemical composition of the samples. Particle size distribution on scaffolds was assessed using the image analysis software tool XT Pro v3.2 (Soft Imaging System GmbH).

Equation 1. Landers formula for scaffold porosity calculation, where d1 is the experimental fiber diameter, d2 is the experimental strand distance and d3 is the experimental layer thickness.

$$Porosity (\%) = \left(1 - \frac{\pi}{4} \cdot \frac{1}{\left(\frac{d2}{d1}\right)} \cdot \frac{1}{\left(\frac{d3}{d1}\right)} \right) \cdot 100$$

Antibiotic release kinetics from films and 3D scaffolds

To assess the antibiotic release from each of the composite 3D scaffolds, 5, 10, 20 % MgAl-CFX_s and 5, 10, 20% ZrP-GTM_s were disinfected (20 min, 70% ethanol), to mimic

the disinfection performed to the samples used for *in vitro* culture, and subsequently incubated for 4 weeks in 1 ml Dulbecco's phosphate buffered saline (dPBS) at 37 °C at static conditions.

2D films were prepared from 10 % CFX_p; 10 % GTM_p; 5, 10 and 20 % MgAl-CFX_p; and 5, 10, and 20 % ZrP-GTM_p. Briefly, 60 milligrams of pellets were molten at 190 °C and pressed with a coverslip against a Teflon sheet to obtain 14 mm diameter, ~300 µm thickness films. To evaluate the effect of the filler on antibiotic release, films were incubated after disinfection (20 min, 70 % ethanol) for 3 weeks in 1 ml dPBS at 37 °C at static conditions, with both sides being exposed to the solution. To assess the effect of pH on antibiotic release, films were incubated in buffer solutions of acidic, neutral and basic pH: pH 4 (potassium hydrogen phthalate based buffer), pH 7.5 (dPBS) and pH 10.5 (sodium tetraborate based buffer). To evaluate the effect of the eluent's ionic concentration on antibiotic release, 20 % MgAl-CFX_f were incubated for 3 weeks in 1ml of dPBS (containing 10 mM Na₂HPO₄, or 6.7 mM PO₄³⁻) or modified dPBS (with 2mM or 50 mM Na₂HPO₄, equivalent to 1.3 mM or 33.5 mM PO₄³⁻, respectively). Similarly, 20 % ZrP-GTM_f were incubated for 3 weeks in 1ml of dPBS (containing 135 mM NaCl, or 53 mM Na⁺) or modified dPBS (with 27 mM or 685 mM NaCl, equivalent to 11 mM or 265 mM Na⁺, respectively).

At each timepoint, the supernatant was collected and films or scaffolds placed on 1 ml of fresh corresponding solution. CFX in the supernatants was detected by direct measurement of CFX absorbance at 275 nm using a nanodrop UV-Vis spectrophotometer (Biodrop). CFX concentrations were calculated from a CFX standard curve. GTM was detected by mixing equal volumes of supernatant, isopropanol and o-phthaldialdehyde reagent (Sigma-Aldrich) and adding 4 µl ml⁻¹ 2-mecaptoethanol (Sigma Aldrich). ³⁷ After 30 min incubation at RT, the fluorescent complex formed was detected using a spectrophotometer (CLARIOstar®, BMG Labtech) at excitation/emission= 340/455 nm. GTM concentrations were calculated from a GTM standard curve.

To investigate the drug release mechanism, 20 % MgAl-CFX_s and 20% ZrP-GTM_s release curves (% antibiotic release from the total loading vs. time) were fitted to 7 commonly used drug kinetic models: zero order, first order, Ritger-Peppas, Higuchi, Bhaskar, modified Freundlich and parabolic diffusion.

Antibacterial activity

Susceptibility of bacterial strains to CFX and GTM

In order to analyze the susceptibility of bacterial strains to CFX and GTM the Agar disk-diffusion method was applied. Mueller- Hinton agar plates were spread with a standardized inoculum of *P. aeruginosa* (CECT 116) and *S. epidermidis* (CECT 231). Then, commercial filter paper discs impregnated with 20 μl of antibiotics at different concentrations (500, 250, 100, 75, 50, 25, 15, 10, 5 and 0 $\mu\text{g ml}^{-1}$) were placed on the agar surface. The Petri dishes were incubated under 37 °C during 18-24 hours. After incubation, zones of growth inhibition (ZOI) around each of the discs (including disc diameter) were measured to the nearest millimeter.

Antibacterial activity of filler-antibiotic complexes and films

Antibacterial activity of fillers-antibiotic complexes (MgAl-CFX and ZrP-GTM) at different concentrations (5, 10 and 20 mg/ml), and films (5, 10, 20% MgAl-CFX_f and 5, 10, 20% ZrP-GTM_f) was evaluated by shaking the samples in a concentrated bacterial suspension (106 CFU ml^{-1}) in nutrient broth (1:500) for 24 h at 37 °C. After incubation, the number of viable bacteria present in the suspension was measured by placing aliquots of the suspensions and their dilutions (in triplicate) in sterile petri dishes with molten nutrient agar and swirled gently. The petri dishes were incubated at 37 °C for 24 h, after which the colonies present on the plates were counted. Values are reported as the $\log_{10}$ reduction (R), calculated as the difference in the log base 10 of the viable cell counts found on a suspension that has not been in contact with the sample and a suspension that has been in contact with the sample.

Antibacterial activity of scaffolds

The antibacterial activity of the antibiotics released from 5, 10, 20 % MgAl-CFX_s and 5, 10, 20% ZrP-GTM_s scaffolds incubated in dPBS solution was evaluated using the Agar disk-diffusion method, as described in previous section.

Cell seeding and culture

HMSCs)isolated from bone marrow were purchased from Texas A&M Health Science Center, College of Medicine, Institute for Regenerative Medicine (Donor d8011L, female, age 22). Cryopreserved vials at passage 3 were plated at a density of 1000 cells cm^{-2} in tissue culture flasks and expanded until approximately 80 % confluency in complete media

(CM) consisting of α MEM with Glutamax and no nucleosides (Gibco) supplemented with 10% fetal bovine serum (FBS), without penicillin-streptomycin (PenStrep) at 37 °C / 5% CO₂.

Metabolic activity assay to determine antibiotic toxicity

The cytotoxicity of CFX and GTM was evaluated via PrestoBlue™ assay. Initially, trypsinized hMSCs at passage 4 were seeded at a density of 5000 cell cm⁻² in the wells of a 48 wellplate, with CM without PenStrep, and allowed to attach for 24 hours at 37 °C / 5% CO₂. The medium was then replaced with a fresh medium containing CFX or GTM at different concentrations, starting from 1 mg ml⁻¹ or 2.5 mg ml⁻¹, respectively. CM without antibiotics was used as control. After 24h, cells were washed with PBS and 250 μ l of Presto Blue solution (10% PrestoBlue in CM) was added to each well and incubated for 1.5 h at 37 °C / 5% CO₂. After incubation, the supernatants were collected and the fluorescence was measured (emission/excitation = 590/560 nm) using a spectrophotometer.

Cell seeding on 3D scaffolds

Scaffolds were disinfected in 70% ethanol for 20 min, washed 3 times with dPBS and incubated for 1.5 h in CM for initial burst release of the antibiotic. Scaffolds that were not pre-incubated in CM were used as controls. Before seeding, scaffolds were dried on top of a sterile filter paper and placed in the wells of a non-treated wellplate. Passage 4 hMSCs were trypsinized, centrifuged 5 min at 500 rcf and resuspended in a dextran solution (500 kDa, Farmacosmos) (10 wt/wt % dextran in CM), to achieve uniform cell seeding, at a density of 200,000 cells per 35 μ l.³⁸ The cell suspension (35 μ l) was placed on top of each scaffold as a droplet, which was drawn into the scaffolds' pores and retained inside due to the hydrophobicity of the wellplate. Seeded scaffolds were incubated for 4 hours at 37 °C / 5% CO₂ to allow cells to attach. After this time, scaffolds were transferred to new wells containing 1.5 ml of basic media (BM) (CM supplemented with 200 μ M L-Ascorbic acid 2-phosphate). BM was replaced after 24h and every two or three days from then on. To evaluate hMSCs osteogenic differentiation, after 7 days in BM, scaffolds were cultured in mineralization media (MM) consisting of BM supplemented with dexamethasone (10 nM) (Sigma-Aldrich) and β -glycerophosphate (10 mM) (Sigma-Aldrich) for another 28 days or until mineral nodules were observed (49 days in the case of MgAl-CFX_s, or 35 days in the case of ZrP-GTM_s).

Imaging of cell viability in 3D scaffolds

After 1h seeding and after 7 days of culture, dead cells were stained for 20 minutes prior to fixation using the Live/Dead™ fixable far-red dead cell stain kit (Thermo Fisher Scientific) at a concentration of 0.5 µl stain in 500 µl Hank's Balanced Salt Solution per scaffold. Subsequently, samples were washed, fixed with 4% paraformaldehyde for 30 min and permeabilized using 0.1% Triton X-100 for 30 min. Samples were incubated with phalloidin (Alexa Fluor 488, 1:75 in PBS) for 1h at RT as counterstain. Finally, samples were washed 3 times with PBS. Scaffolds cross sections were imaged with a fluorescent microscope (Eclipse, Ti2-e, NIKON). Background subtraction and brightness adjustments were performed on the images using the software Image J, in order to clarify their visualization. No image quantification was performed on the processed images.

Biochemical assays

Alkaline phosphatase (ALP) assay

ALP activity was evaluated on MgAl-CFX_s or ZrP-GTM_s after 47 days and 35 days of culture in MM, respectively (timepoints day 56 and day 42, respectively). 3D scaffolds were collected at every timepoint and washed with PBS, stored at -80 °C and freeze-thawed 3 times, to improve lysis efficiency. Samples were incubated for 1h at RT in a cell lysis buffer composed of KH₂PO (0.1 M), K₂HPO₄ (0.1 M), and Triton X-100 (0.1 v%), at pH 7.8. The chemiluminescent substrate for ALP CDP star® ready to use reagent (Roche) was added to the cell lysate at a 1:4 ratio. Luminescence was measured using a spectrophotometer. Remaining cell lysates were kept for DNA quantification. ALP values were reported normalized to DNA content.

DNA assay

DNA assay was performed on cells cultured on all 3D scaffolds after 1 and 7 days in BM (timepoints day 1 and day 7). Additionally it was performed on PEOT/PBT_s, 10% MgAl_s and 10% ZrP_s after 28 days in MM (timepoint day 35), on MgAl-CFX_s after 28 and 49 days in MM (timepoints day 35 and day 56), and on ZrP-GTM_s after 28 and 35 days in MM (timepoint day 35 and day 42). CyQUANT cell proliferation assay kit (Thermo Fisher Scientific) was used. Lysed samples from ALP assay or frozen samples after collection, were incubated overnight at 56 °C in Proteinase K solution (1mg/ml Proteinase K (Sigma-Aldrich) in Tris/EDTA buffer) for matrix degradation and cell lysis. Subsequently, samples

were freeze-thawed three times and incubated 1h at RT with a 20X diluted lysis buffer from the kit containing RNase A (1:500) to degrade cellular RNA. Lysed samples were incubated with the fluorescent dye provided by the kit (1:1) for 15 min and fluorescence was measured (emission/excitation = 520/480 nm) with a spectrophotometer. DNA concentrations were calculated from a DNA standard curve.

Alizarin red S (ARS) staining and quantification

Calcium deposition was qualitatively determined by ARS staining on MgAl-CFX_s or ZrP-GTM_s after 28 and 49 days (timepoints day 35 and day 56, respectively), or 28 and 35 days (timepoints day 35 and day 42, respectively) of culture in MM, respectively. PEOT/PBT_s, 10% MgAl_s and 10% ZrP_s scaffolds were stained after 28 days of culture in MM (timepoint day 35). Briefly, 3D scaffolds were fixed with 4% paraformaldehyde and washed with distilled water. Subsequently, scaffolds were stained with ARS (60 mM, pH 4.1 - 4.3) for 20 min at RT, and washed with distilled water. Images of scaffolds' cross sections were taken using a stereomicroscope (Nikon SMZ25). After imaging, stained samples (whole scaffold) were incubated for 1h at RT while shaking, followed by 10 min incubation at 85 °C, with 30 v% acetic acid. Afterwards, scaffolds were removed and solutions were centrifuged at 20,000 rcf for 10 min. An appropriate volume of ammonium hydroxide (5 M) was added to the supernatants to bring the pH to 4.2. The absorbance was measured at 405 nm using a spectrophotometer. Concentration of ARS was calculated from an ARS standard curve and the values were normalized to DNA content.

Immunostaining

Cells were fixed in 4% paraformaldehyde after 35 days (PEOT/PBT_s and ZrP-GTM_s) or 47 days (MgAl-CFX_s) of culture in MM (timepoints day 42 and day 56, respectively). Triton-X 100 (0.1 v%) was added and incubated for 30 min. Samples were washed 3 times with PBS and then blocked for 1 h with blocking buffer (3 % BSA + 0.01% Triton-X 100). Afterwards, primary antibodies were added (1:200 Collagen I (COL I) polyclonal antibody rabbit-derived (ab34710, Abcam), or 1:200 Osteopontin (OPN) polyclonal antibody rabbit-derived (ab8448, Abcam), in washing buffer (10x diluted blocking buffer)) and incubated overnight. Samples were washed 3 times with washing buffer prior to secondary antibody incubation (1:200 Alexa Fluor 488 goat derived anti rabbit antibody (Thermo Fisher Scientific), for PEOT/PBT and MgAl-CFX scaffolds, or 1:200 Alexa Fluor 647 goat derived anti rabbit (Thermo Fisher Scientific), for ZrP-GTM based scaffolds, in washing buffer).

After 1 h incubation, scaffolds were washed as before and stained for F-actin (647 Alexa Fluor Phalloidin (Thermo Fisher Scientific) for PEOT/PBT_s and MgAl-CFX_s, and 568 Alexa Fluor Phalloidin (Thermo Fisher Scientific) for ZrP-GTM_s (1:75 in PBS)), incubated for 1 h and washed 3 times with PBS. Confocal laser scanning microscopy was performed with a tandem confocal system (Leica TCS SP8 STED), equipped with a white light laser (WLL). Samples were excited with the dye specific wavelengths using the WLL. Emission was detected with HyD detectors (F-actin, OPN, COL I). For optimal visualization in the reported images, F-actin was colored in green, COL I in red and OPN in blue for all sample types.

Statistical analysis

All data is shown as average with error bars indicating the standard deviation of at least three replicates. Analysis of statistics was conducted with GraphPad Prism (version 8.0.1). A one-way or two-way analysis of variance (ANOVA) was performed followed by a Tukey's post-hoc multiple comparison test to evaluate statistical significance and correct for multiple comparisons.

RESULTS AND DISCUSSION

Scaffolds fabrication and characterization

Although the sustained release of drugs from MgAl³⁹⁻⁴⁰ and ZrP⁴¹⁻⁴³ has been previously reported, the combination of antibiotic-loaded fillers with polymers has not been carefully examined in terms of release kinetics, antimicrobial functionality or assurance of no negative effect on the bone regeneration response. Most importantly, no previous studies have reported their processing into 3D AM scaffolds, but only as implant coatings⁴⁴ or incorporated on electrospun scaffolds.⁴⁵ Here, 3D polymer composite scaffolds with filler-antibiotic concentrations up to 20 wt% (MgAl-CFX_s or ZrP-GTM_s, **Table 1**) were successfully fabricated for the first time via ME-AM of previously prepared melt-blended composite pellets (MgAl-CFX_p or ZrP-GTM_p) (**Figure 1** and **Figure S1**). The biodegradable polyether-ester multiblock copolymer PEOT/PBT was selected as model thermoplastic polymer to produce the scaffolds due to its processability at high temperatures (≥ 190 °C), and its proven *in vitro* and *in vivo* applicability in the orthopedic field.⁴⁶⁻⁵⁰ All composite scaffolds exhibited interconnected macropores, and experimental

filament diameters, as well as X-Y and Z pores sizes consistently matched the theoretical values (**Figure S2**). Scaffolds presenting pores of $>300\ \mu\text{m}$ and porosity values in the range of $\sim 50\%$ (**Figure S2B**) suggest that these constructs will support bone regeneration.⁵¹

Figure 1. Schematic representation of the antibacterial composite scaffolds (PEOT/PBT-filler-antibiotic) fabricated by melt-extrusion AM. These polymeric composite scaffolds contain either MgAl intercalated with CFX (MgAl-CFX), or ZrP intercalated with GTM (ZrP-GTM). When submerged in an eluent solution, anions can exchange with CFX, which diffuses out of the filler and polymer matrix into the eluent. On the other hand, cations can exchange with GTM in the ZrP-GTM based system, allowing the diffusion of GTM out of the filler and polymer matrix. The release of antibiotics confer antimicrobial activity to the scaffolds.

As it is observed in **Figure 2**, BSEM images confirm an increasing amount of filler (visible as brighter points) with increasing filler-antibiotic concentrations on the surface and cross section of the scaffolds filaments. Using EDS the fillers were discriminated as MgAl and ZrP containing particles and the distinctive atomic elements of the CFX (F and N) and GTM (S and N) molecules were found to co-localize with their respective lamellar particles (**Figure 2C-D**). This strongly indicates that the filler and antibiotic were not separated in the filaments, but showed a preserved filler structure after the compounding and ME-AM process. Moreover, particle size was maintained for both types of fillers within the after-synthesis values during the processing steps. A slight size reduction was observed during the ME-AM process, likely due to the higher shear forces in the print-head screw (**Figure S3**). Despite the presence of some filler aggregates close to the filaments' surface, the composite scaffolds surface roughness was not significantly altered in the microscale, presenting a smooth surface comparable to the PEOT/PBT scaffolds (**Figure 2** and **Figure**

S4). The sub-surface aggregates do form topographical projections, but given their low numbers and large separations (on the order of 100 μm), the cells are not likely to interact with multiple projections simultaneously.

Figure 2. Filler distribution and antibiotics localization on composite scaffolds. BSEM micrographs of scaffolds cross sections and filaments surface depicting filler distribution (as white points) on (A) MgAl-CFX_s and (B) ZrP-GTM_s at different filler-antibiotic concentrations. Scale bars 250 μm . EDAX elemental mapping of representative fiber cross section of (C) 20 % MgAl-CFX_s and (D) 20 % ZrP-GTM_s. Scale bars 50 μm .

In vitro antibiotic release

In order to evaluate the kinetics of antibiotic release from MgAl-CFX_s and ZrP-GTM_s, composite scaffolds were incubated in dPBS, and CFX and GTM released amounts were monitored over time. As demonstrated in **Figure 3A**, by increasing the MgAl-CFX concentration within MgAl-CFX_s, larger amounts of CFX were released over time. A similar trend was also observed in the release profile of GTM from ZrP-GTM_s (**Figure 3B**). This tendency demonstrates the possibility of designing a customized scaffold, which releases the desired amount of antibiotic over time, by tuning the filler-antibiotic

concentration mixed with the polymer matrix. Although both systems demonstrated a filler-antibiotic concentration-dependent release profile, significantly different release kinetics were observed on MgAl-CFX_s and ZrP-GTM_s. While the release of CFX from MgAl-CFX_s was sustained over the course of the evaluated time (1 month), there was a GTM burst release from ZrP-GTM_s during the first 24h followed by a slower release up to the completion of the study (**Figure 3A-B**). When active molecules are fixed into the inorganic lamellae of layered double hydroxides (LDH), such as MgAl, their de-intercalation mechanism occurs mainly by anion exchange, a process in which the intercalated active molecule and free anions in the elution solution can be exchanged via a concentration gradient.⁵² On the other hand, the release of active molecules from metal hydrogen phosphates, such as ZrP, can occur not only by ion exchange (in this case, of cations), but also via the disruption of acid-base interactions between the active molecule and the filler, which rely on a pH switch in the environment about the acid dissociation constant (pKa) of the intercalated molecule (**Figure 1**).⁵² A series of factors are involved in the release kinetics of antibiotics when the lamellar particles are dispersed within a polymeric matrix: i) the permeation and diffusion of water molecules carrying counter ions through the polymer matrix; ii) the intra-particle diffusion of water, ions and active molecules, which depends on guest-host interactions and determines the rate of de-intercalation, and iii) the diffusion rate of active molecules (antibiotics in this study) from the polymer matrix.⁵³ In order to determine the release rate-limiting step, the release profiles of the 20 wt% filler-antibiotic scaffolds, taken as a representative cases, were fitted to various release kinetic models. 7 relevant models were used and the corresponding correlation coefficients are reported in **Figure S5**. Bhaskar model was found to best describe the release profile of CFX from 20% MgAl-CFX_s. This model has previously been used to explain the release of active molecules from LDH systems and indicates that the release of CFX is an intra-particle diffusion controlled process, and therefore, ion exchange rate is the limiting step.⁵⁴⁻⁵⁷ Accordingly, the decreased release rate of CFX over time (**Figure 3A**) is likely due to the formation of a phase boundary between internal zones of the lamellar filler containing intercalated CFX and the external zones, in which anions from dPBS have already replaced the antibiotic, which progressively declines the antibiotic release process.⁵⁸

To gain more insight into the release mechanism of 20% ZrP-GTM_s, the total progress was separated into two stages: a rapid release stage I (<24h) followed by a slow release stage II (>24h). The modified Freundlich kinetic model, which describes heterogeneous diffusion from flat surfaces of clays via ion exchange, was found to explain the release in

Stage I (**Figure S5**). This implies that GTM diffuses into the medium solution via anion exchange during the first 24h, at a fast speed due to the antibiotic high solubility in water-based neutral pH eluents.⁵⁹⁻⁶¹ Since after the studied period the GTM released did not reach 100% (**Figure S5A**), it is hypothesized that GTM released in stage I came from the ZrP particles located on the surface of scaffold filaments, after which the release of GTM from internal ZrP particles was hindered by the polymer network, resulting in the plateau-like profile observed in Stage II (**Figure 3B**). This type of behavior was already reported and is likely due to the low diffusion rate of water molecules through the polymer network, given by the poor water uptake/swellability capability of PEOT/PBT (4% water uptake) at the specific PEO molecular weight (300 kDa) and PEOT:PBT segments ratio (55:45).^{48, 53} In this case, the GTM still incorporated into the system could be released upon polymer degradation, a process that could be potentially speeded up by increasing the molecular weight of the PEO segments of the copolymer or by using a different and faster degrading material. Following the same reasoning, the fact that a plateau was not reached in the CFX release curve from MgAl-CFX_s, and that only ~15-20% of the total CFX was released (**Figure S5A**) in the analyzed time, suggests that the CFX released during the evaluated period was also originated from the MgAl particles localized on the surface of the filaments. It is plausible that due to the low solubility of CFX at neutral pH, and possibly the stronger filler-antibiotic interactions, the fast release seen in Stage I in 20% ZrP-GTM_s did not happen in 20% MgAl-CFX_s.

Effect of filler, pH and ionic concentration

As mentioned above, the de-intercalation of molecules from fillers has been shown to be driven by ion exchange and pH changes in the eluent media. To further investigate these effects on our composite systems, easy to manufacture films were incubated in solutions with different pH values and ionic concentrations, and the release profiles were compared relative to each other without considering the absolute release amounts. In addition, the effect of filler was explored by preparing composite films (20 % MgAl-CFX_f) and films made of the same amount of CFX directly dispersed in the copolymer matrix (10% CFX_f) was compared (**Figure 3C**). The presence of the filler enabled the maintenance of a lower antibiotic release rate over time. The effect of filler on the sustained release of CFX was further confirmed upon incubation of 20% MgAl-CFX_f in acidic pH (pH = 4) (**Figure 3C**). Here, the acidic attack dissolved the layered MgAl, allowing for a faster release of CFX, whose release was not governed by ion exchange anymore and became comparable to

Figure 3. Antibiotic release profiles from MgAl-CFX and ZrP-GTM copolymer composites under different eluent conditions. Cumulative release of (A) CFX from MgAl-CFX_s and (B) GTM from ZrP-GTM_s immersed in dPBS during 4 weeks at 37 °C, proving the filler-antibiotic concentration dependent release. Influence of filler presence and pH of the eluent solution on the cumulative release percentage of (C) CFX and (D) GTM from filler-antibiotic composite films (20% MgAl-CFX_f and 20% ZrP-GTM_f, respectively) and films composed of comparable amounts of antibiotic directly dispersed in the copolymer matrix (10% CFX_f and 10% GTM_f), immersed in buffers at acidic, neutral and basic pH during 3 weeks at 37 °C. Influence of the ionic concentration of the eluent solution on the cumulative release percentage of (E) CFX and (F) GTM from 20% MgAl-CFX_f and 20% ZrP-GTM_f, respectively, during 3 weeks a 37 °C.

the release of CFX from 10% CFX_f at pH = 4.^{56, 62} Interestingly, at pH ≥ 7 , 10% CFX_f release profile remained unchanged and lower release rates than at pH = 4 were observed, due to lower CFX solubility at this pH range. Similarly, 20% MgAl-CFX_f release profile did not vary at pH ≥ 7 , as MgAl remained stable and intact (**Figure 3C**). The role of ZrP on maintaining a more sustained release, especially in the stage I, was also demonstrated for 20% ZrP-GTM_f, when comparing to 10% GTM_f (**Figure 3D**). However, while MgAl in MgAl-CFX_f was affected by acidic pH, the release of GTM from 20% ZrP-GTM_f was affected by alkaline conditions (pH = 10.5), due to the different nature of the system (**Figure 3D**). Considering the isoelectric point of GTM (pKa = 8),⁶³ the hydroxyl groups of GTM were deprotonated at pH = 10.5, disrupting the hydrogen bonds with the co-

intercalated water molecules as well as producing a negatively charged GTM, which was repelled from the ZrP filler due to the disrupted acid-base electrostatic interactions between the GTM amine groups and the phosphate groups of the ZrP layers.⁴¹ It is hypothesized that oxidative degradation of PEOT/PBT, happening at alkaline conditions,^{48, 64} allowed for further increase in the diffusion of GTM from the polymer matrix, which is visible as a slight increase in the release rate of GTM in Stage II on both 20% ZrP-GTM_f and 10% GTM_f release curves at pH =10 (**Figure 3D**).

The aforementioned results indicate that the GTM release from ZrP-GTM_f in stage II can be controlled by varying the pH (acid base-interactions), without filler degradation. This mechanism of release was found to be more effective than ion exchange. As depicted in **Figure 3F**, the release of GTM does not vary upon increasing or decreasing the concentration of cations (sodium, Na⁺) in the eluent, within the evaluated concentration range. On the contrary, CFX release is clearly affected by the concentration of anions (phosphate, PO₄³⁻) in the elution buffer, as depicted in **Figure 3E**, further demonstrating the ion-exchange related release of the system. Compared to other inorganic anions present in dPBS, phosphate ions present the highest affinity with LDH fillers, due to their higher charge.⁴⁰ By a 5-fold decrease in the phosphate concentration, CFX release decreased by 52% after 3 weeks. Interestingly, a 5-fold increase over the dPBS phosphate base concentration did not affect the system, suggesting the existence of a concentration threshold.

Overall, by varying the filler and antibiotic type incorporated within the scaffold, two antibiotic delivery systems were fabricated, whose sustained release can be controlled not only upon changing the filler-antibiotic concentration, but also by varying both the eluent ionic concentration or pH (in the case of MgAl-CFX), or pH (in the case of ZrP-GTM). The ionic concentration (*i.e.* phosphate, calcium, sodium, potassium) is a parameter that cannot be predicted during the bone regeneration process. In this regard, *in vivo* release experiments would be necessary to understand the *in situ* antibiotic release from the scaffolds. On the other hand, the local pH in a fracture during bone healing is known to be maintained in the range of pH 6 to pH 7.5 and, in principle, our system is not tunable in this range.⁶⁵⁻⁶⁶ However, the property of some biocompatible and biodegradable polymers, such as the aliphatic polyesters PLA, PGA or PLGA, to dramatically decrease the local pH upon degradation (down to pH=4),⁶⁷ could be use advantageously to further customize the *in vivo* antibiotic release from filler-antibiotic composite scaffolds.

The polymer degradation profile of the scaffold and its effect on the antibiotic release kinetics would also be a matter of evaluation in future studies. Previous literature suggests the slow degradation of the PEOT/PBT used in this study (PEO molecular weight = 300 kDa, PEOT:PBT weight ratio = 55:45), which is mainly due to the low molecular weight of the PEO hydrophilic segment preventing the swelling of the material and lowering the possibility of hydrolysis. Accordingly, the intrinsic viscosity and mechanical properties of PEOT/PBT with similar characteristics has been shown to remain constant after 24 weeks immersion in PBS at 37 °C.⁶⁸ Similar results were obtained after 24 weeks implantation *in vivo*, where the mass loss was only up to 3% and minimum cracks were observed in the surface by SEM.⁶⁹ It is plausible that in our case, the fillers increased the degradation time of the polymer, as previously shown on PEOT/PBT – nanosilicate scaffolds.⁷⁰ Yet, long term *in vitro* and *in vivo* studies are required to further evaluate this.

Thermal stability of antibiotics and scaffolds antimicrobial activity

One of the main concerns involved in the AM of 3D scaffolds via high-temperature ME of polymers containing bioactive molecules is the preservation of their bioactivity after the process' thermal stresses. It has been previously demonstrated that the incorporation of some bioactive molecules into lamellar inorganic fillers increases their thermal stability.^{56, 71-72} However, CFX decomposes already at high temperatures due to its stable aromatic structure, what might have neutralized the potential shielding effect of the MgAl filler in our system, which in turn maintained CFX stability, as demonstrated by TGA (**Figure S6A**), as well as in previously studies.⁷² While in our TGA analysis CFX does not show any degradation bellow 255 °C, according to literature, MgAl-CFX weight loss bellow 255 °C is attributed to the loss of MgAl interlamellar water, and CFX decarboxylation occurs at ~146 °C.⁷³⁻⁷⁴ This inconsistency might be due to experimental conditions differences. Yet, HPLC confirmed molecular decomposition of CFX after the melt-blending process at around 150 °C (~ 73% degradation), potentially attributed to the aforementioned decarboxylation of CFX at 146 °C (**Figure S6B, C**).⁷⁴ Further decomposition occurred during the scaffolds production via ME at 190-195 °C (~ 45% degradation of the remaining GTM after melt blending, equivalent to ~12.5 % degradation of the total initial GTM loading).

Similarly, GTM decomposition starts at 220 °C following some water loss, and the ZrP filler did not show to increase nor decrease its decomposition temperature, as shown in **Figure S7A**.⁶³ The loss of interlamellar water of ZrP-GTM occurs in the range 20–220 °C, while

the step above 220 °C is attributed to the condensation of the PO_4^{3-} groups and the decomposition of GTM.³⁵ Accordingly, HPLC-MS analysis of GTM probed the stability of the molecule after each of the processing steps, with ~ 1% and ~ 8% degradation after melt-blending and ME-AM, respectively (**Figure S7B, D**). Interestingly, this analysis also proved the preferential intercalation of the C1a GTM isoform during the ZrP-GTM synthesis process, due to its less steric bulky form, with respect to the other isoforms (**Figure S7C**).

Preliminary experiments demonstrated that the antimicrobial activity of the antibiotics was preserved after their intercalation within the filler, as well as in MgAl-CFX_f. Similarly, ZrP-GTM and ZrP-GTM_f presented antibiotic activity (**Table S1** and **Table S2**). Most importantly, the antimicrobial activity of both MgAl-CFX_s and ZrP-GTM_s against *S. epidermidis* and *P. aeruginosa*, was preserved on ME-AM scaffold (**Figure 4**), despite the aforementioned degradation. The antimicrobial activity of CFX consists of inhibiting the action of two essential enzymes that are involved in the modulation of the chromosomal supercoiling required for bacterial DNA synthesis, transcription and cell division.⁷⁵ Despite its degradation, important active sites for CFX interaction with the target enzymes remained intact at the processing temperatures during melt-blending and posterior ME into scaffolds, thus preserving its antimicrobial activity.⁷⁵ Importantly, the antimicrobial activity of CFX contained in MgAl-CFX_s was preserved at the same levels as unprocessed CFX, observed when comparing the experimental ZOI and the expected ZOI (*i.e.* ZOI given by corresponding amounts of pure unprocessed CFX) (**Table S3** and **Table S4**). On the contrary, previous studies demonstrate a decrease in CFX efficiency against certain bacterial strains upon exposure to high temperatures (120 °C for sterilization).³⁰ However, these tests were performed under oxygen atmosphere, unlike the nitrogen conditions in which the ME-AM takes place, which in combination with MgAl might have acted as a protective shield in our system. In accordance with the thermal stability results of GTM and supported by previous GTM heat stability studies,³⁰ the efficiency of this antibiotic released from ZrP-GTM_s was also preserved at the same levels as unprocessed GTM (**Table S3** and **Table S5**).

Figure 4. Antimicrobial activity of polymeric composite 3D AM scaffolds with different filler-antibiotic concentrations over 4 weeks. Graphs depicting ZOI of (A, B) MgAl-CFX_s and (C, D) ZrP-GTM_s in dPBS at 37 °C against *S. epidermidis* and *P. aeruginosa*. Images of representative disk diffusion test plates depicting ZOIs around a disk impregnated with an aliquot of the scaffold release supernatant after the initial 24h of release. Scale bars 5 mm.

CFX MIC against both *S. epidermidis* and *P. aeruginosa* and is 1 µg/ml and 0.5 µg/ml, respectively.⁷⁶ Since the non-cumulative release of CFX from MgAl-CFX_s per timepoint was significantly above the MIC thresholds at all MgAl-CFX concentrations (**Table S4** and **Figure S8**), the antimicrobial activity was preserved over the period evaluated (30 days) showing an increasing activity with increasing MgAl-CFX concentration (**Figure 4A, B**). On the other hand, the required GTM concentration to inhibit bacterial growth is higher (MIC against *S. epidermidis*: 1 µg/ml; MIC against *P. aeruginosa* 4 µg/ml).⁷⁶ Therefore, in accordance with the amount of GTM released per timepoint (**Table S5** and **Figure S8**), the antibacterial activity of ZrP-GTM_s against *S. epidermidis* lasted for 3 weeks, with a gradual decrease after a peak of activity in the first 48h (**Figure 4C**), while only 24h against *P. aeruginosa* (**Figure 4D**). The concentration dependent activity was also observed in the case of ZrP-GTM_s as it is shown in **Figure 4C, D**.

While *in vivo* experiments would be necessary to fully understand the *in situ* antimicrobial function of the scaffolds, which is linked to the aforementioned *in vivo* conditions for antibiotic release (ionic concentration and pH), this results already suggest the antimicrobial potential of the presented systems.

Cell viability

To further assess the cytotoxicity of the antibacterial systems, hMSCs viability was evaluated on MgAl-CFX_s and ZrP-GTM_s at the three different filler-antibiotic concentrations. Preliminary experiments demonstrated a significant reduction in cell number after 7 days of culture in 20% MgAl-CFX_s and 20% ZrP-GTM_s scaffolds (**Figure S9A, B**). Despite their well-known biocompatibility,^{43, 77} some reports describe the dose, time and size dependent cytotoxic effects of layered lamellar compounds, such as MgAl and ZrP when uptaken by cells.⁷⁸⁻⁷⁹ While this is not likely to occur during our *in vitro* experiments period as the fillers remain trapped within the copolymer matrix, fillers could potentially play a role in cell toxicity upon polymer degradation. Yet, we hypothesize that the sustained release of fillers into the surroundings, due to the slow polymer degradation process, would not lead to toxic amount of fillers as these are always under the toxicity limits previously reported in literature (~200-500 µl/ml).^{43, 80-82} Thus, it is plausible that the high amount of antibiotic released during the 4h seeding time concentrated in a small volume, negatively affected hMSCs attachment and viability. hMSCs cultured in tissue culture polystyrene showed a dose-dependence sensitivity to CFX (**Figure S10A**), which has shown to influence hMSCs proliferation upon continuous exposure.⁸³ Similarly, although GTM has an impact on hMSCs viability only at high concentrations (>1000 µg/ml) (**Figure S10B**), short exposure (20 min) to high levels of GTM was found to influence osteoblasts proliferation rate up to 3 days post-exposure.⁸⁴

To prevent a decrease in cell viability, surface extraction by pre-incubating the scaffolds in cell culture media prior to seeding (for 1.5h) was considered, as previously described.⁸⁵ This method was used to reduce the initial drug burst release prior to cell contact and showed to significantly improve cell viability within our scaffolds (**Figure 5D, E**). Accordingly, no meaningful differences in cell number were found among filler-antibiotic concentrations and comparable levels of viability were maintained for 7 days. Representative live/dead images of the scaffolds cross sections further confirmed high hMSCs viability regardless of the filler-antibiotic concentration over 7 days for both MgAl-

CFX_s and ZrP-GTM_s (**Figure 5A-C**). Interestingly, cells in the composite scaffolds showed a spread morphology comparable to the control samples (*i.e.* polymer-only scaffolds) after 1 day of culture, which changed to the characteristic elongated morphology of migrating cells after 7 days.

Figure 5. HMSCs viability is preserved after seeding MgAl-CFX_s and ZrP-CFX_s pre-incubated in media for 1.5 h. Representative fluorescent images depicting cell viability (F-actin, green; dead cells, red) 1 day and 7 days post-seeding (a) MgAl-CFX_s, (b) ZrP-GTM_s and (c) PEOT/PBT_s. Quantification of DNA content 1 day and 7 days post seeding (d) MgAl-CFX_s and (e) ZrP-GTM_s, compared to PEOT/PBT_s control. Scale bars 250 μm. Statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$).

Osteogenic differentiation

In order to gain more insight into the potential of MgAl-CFX_s and ZrP-GTM_s to promote new bone formation, the osteogenic differentiation capacity of hMSCs seeded on the different scaffolds was analyzed. Preliminary results suggested that the osteogenic potential of hMSCs was influenced by the presence of the filler-antibiotic complexes. As

depicted in **Figure S11**, after 28 days in mineralization media (MM), no mineral deposition was visualized or quantified on MgAl-CFX_s for all concentrations, nor on 20% ZrP-GTM_s, while a ~10-fold reduction in calcium deposition was observed in 5 and 10 % ZrP-GTM_s with respect to the control samples (**Figure S11**). Yet, calcium deposits were observed in the polymer- and filler-only control scaffolds after 28 days in MM at levels reported in previous studies.³⁸ This indicates that the fillers were not affecting hMSCs osteogenic behavior. This is further supported by a recently published report demonstrating the significant upregulation of several osteogenic genes including ALP, bone sialoprotein, Runt-related transcription factor 2 (RUNX2), OPN, and osteocalcin in osteoblasts when adding MgAl particles to the cell culture media.⁸⁰ In order to investigate if the antibiotics were impairing or just delaying the progression of hMSCs differentiation towards the creation of a mineralized matrix, scaffolds were cultured in MM for longer periods. After a total of 35 days in MM, all ZrP-GTM_s showed calcium deposition, which were inversely proportional to the ZrP-GTM concentration (**Figure 6B**). This suggests a GTM dose-dependent effect and its role in delaying mineralization. The amount of calcium in 5, 10, and 20 % ZrP-GTM_s increased ~3, 2 and 25-fold with respect to the previous timepoint, respectively. After a total of 49 days in MM, both 5 and 10 % MgAl-CFX_s showed matrix mineralization, which was not observed at earlier timepoints, yet at levels lower than the control conditions.

Interestingly, 20% MgAl-CFX_s did not mineralize even after 49 days in MM, suggesting that the continuous release of higher CFX amounts had a starker effect on the hMSCs differentiation process. This result was further supported by the lower DNA and higher values of ALP activity observed on 20% MgAl-CFX_s compared to lower MgAl-CFX concentrations and the control (**Figure S12**), as the ALP activity is known to decrease during the mineralization process under optimum conditions.⁸⁶ A strong inhibition of mineralization has been reported when confluent cultures of osteoblasts were treated with CFX. The effect was apparent with 2.5-5 µg/ml CFX and progressively increased to more than a 90% decline in the calcium/protein ratio at 20-40 µg/ml CFX.⁸⁷ These values are in the range of the release of CFX from 5-10% MgAl-CFX_s and 20 % MgAl-CFX_s, respectively, therefore supporting our results. While *in vitro* experiments agree in the dose dependent interference of CFX in the bone healing process, controversial results have been reported for *in vivo* studies. Abnormalities in cartilage morphology of fracture calluses were observed after treatment in rat models with ciprofloxacin (2.4 µg/ml in serum),⁸⁸ while higher CFX release (5 µg/ml per day) from a bone defect filler in rabbits allowed bone

formation within 3 months in another study.⁸⁹ This suggests that *in vivo* studies are necessary to gather a better insight over the effect of our scaffolds on bone formation. Compared to CFX, consistent *in vitro* and *in vivo* results have been previously published on the dose-dependency effect of GTM. For instance, mineralization in osteoblast and hMSCs cultures was not impaired after short exposure of up to 16 $\mu\text{g/ml}$ GTM,⁸⁴ or continuous exposure of 200 $\mu\text{g/ml}$.⁹⁰ Similarly, *in vivo* GTM serum concentration of 4.5 $\mu\text{g/mL}$,⁹¹ or a continuous release from a bone filler with a peak of 58 μg in the first 4h,⁹² did not show reduced healing effect in rat models. However, a burst release from a bone cement (10 mg GTM released in 3 days) resulted in significantly less mineralized tissue than the control in a sheep model.⁹³

Figure 6. Influence of filler-antibiotic concentration on matrix mineralization. Representative stereomicroscopy images of scaffolds cross sections stained with ARS: (a) MgAl-CFX_s after 56 days of culture (49 days in MM) and (b) ZrP-GTM_s and (c) PEOT/PBT_s after 42 days of culture (35 days in MM). Inserts represent the corresponding control scaffolds without cells incubated in MM and stained with ARS. Scale bars 1 mm. (d) Quantification of the ARS staining extracted from scaffolds normalized to cell number. Statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (**p < 0.01; ***p < 0.001).

Along with the study in matrix mineralization, the progression in the osteogenic differentiation of hMSCs was further evaluated by immunostaining of two relevant osteogenic makers, COL I and OPN. COL I constitutes nearly 90% of bone organic ECM and serves as minerals nucleation sites.⁹⁴ On the other hand, OPN is one of the most

abundant non-collagenous proteins in bone. It reaches its peak during the mineralization period supporting its progression and the prevention of premature precipitation of calcium phosphate is suggested as one of its key functions.^{86, 95} Interestingly, a dense layer of COL I fibrils was formed by hMSCs on top of scaffold filaments regardless of the scaffold composition and no significant differences were observed compared to the PEOT/PBT control (**Figure 7A** and **Figure S13**). This implies that COL I fibers formation was not affected by antibiotics. However, OPN expression was reduced in MgAl-CFX_s when compared to PEOT/PBT_s and ZrP-GTM_s, qualitatively observed when comparing among scaffold types the ratio of OPN-stained hMSCs over the total cell number (**Figure 7B** and **Figure S13**). Intriguingly, this result lays in agreement with the delay in matrix mineralization observed on these scaffolds. Particularly, in 20 % MgAl-CFX_s, where OPN expression is more clearly downregulated, the pronounced delay in the osteogenic differentiation of hMSCs cultured on these scaffolds is confirmed once more. However, the trend of the current results indicates that longer culture periods might lead to comparable osteogenesis and matrix mineralization results among different scaffolds types.

Figure 7. Influence of filler-antibiotic concentration on the osteogenic differentiation of hMSCs. Representative confocal microscopy images of hMSCs (F-actin, green) on top of the filaments of 20 % MgAl-CFX_s after 54 days of culture (47 days in MM) and 20% ZrP-GTM_s and PEOT/PBT_s after 42 days of culture (35 days in MM) and stained for the relevant osteogenic markers COL I (red) and OPN (blue). Scale bars 100 μm.

Overall, the aforementioned results suggest that bone formation is supported *in vitro* on the filler-antibiotic based scaffolds presented in this study. Further studies should aim at evaluating the bone bonding and heterotopic bone formation capabilities of the scaffolds by *in vitro* biomineralization experiments. Yet, conclusions should be drawn carefully after these, since it has been previously shown that the formation of an apatite layer on the material surface *in vitro* is not always correlated with bone formation *in vivo*.⁹⁶ Ultimately, *in vivo* experiments will be necessary to fully characterize the scaffolds bone regeneration performance.

CONCLUSION

Local delivery of antibiotics is desired to prevent infections after bone fracture of implant-related surgery. The aim of this study was to develop antibiotic loaded 3D scaffolds fabricated via high temperature ME-AM. In order to protect antibiotics from the fabrication process thermal stresses and obtain a sustained release, the antibiotics CFX and GTM were intercalated within the lamellar structure of the inorganic layered fillers MgAl and ZrP, respectively, prior to their dispersion within the polymeric matrix. Compared to no-filler systems, a more sustained release was obtained in the filler-based scaffolds. The antibiotic was released via diffusion through the polymer matrix and by a mechanism governed by ion exchange or pH changes. The functionality of the antibiotics after high temperature AM was confirmed against Gram + and Gram - bacterial strains demonstrating same antibacterial levels as unprocessed antibiotics. Importantly, the incorporation of fillers and antibiotics into the scaffolds neither affected hMSCs viability nor prevented their osteogenic differentiation, as verified by matrix mineralization and COL I and OPN expression. These newly proposed method enables the fabrication of dual functional scaffolds via direct incorporation of antibiotics in scaffolds fabricated at high temperatures, and has the potential to improve infection management while allowing bone regeneration. Moreover, this approach opens the door to future research on the use of other relevant antibiotics and biodegradable thermoplastic polymers to further optimize the system towards clinical applications.

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SUPPLEMENTARY FIGURES

Figure S1. (A) Schematic representation of the melt-compounding process of the different copolymer-filler-antibiotic composites. Initially, CFX was intercalated within MgAl to obtain MgAl-CFX, while GTM was intercalated within ZrP to obtain ZrP-GTM. Each of the two filler-antibiotic complexes separately were mixed at 5, 10 and 20 wt% with PEOT/PBT using a twin screw extruder. Pellets of the 6 different materials (MgAl-CFX_p and ZrP-GTM_p) were obtained to be subsequently used for the fabrication of 3D scaffolds via ME-AM. (B) SEM micrographs of the cross section of MgAl-CFX_p and ZrP-GTM_p with different filler-antibiotic concentrations. Scale bars 200 μm .

Figure S2. (A) Stereomicroscopy images of PEOT/PBT_s and composite scaffolds cross section and top view (inserts) depicting the scaffolds' interconnected porosity in XYZ. Scale bars 1mm. (B) Experimental fiber diameter (d1), strand distance (d2), layer thickness (d3), and porosity calculated for each scaffold,

Figure S3. (A) MgAl-CFX particle size (D90) after synthesis, melt blending and ME-AM. (B) MgAl-CFX size distribution on MgAl-CFX_p and (C) MgAl-CFX_s. (D) ZrP-GTM particle size (D90) after synthesis, melt-blending and ME-AM. (E) ZrP-GTM size distribution on ZrP-GTM_p and (F) ZrP-GTM_s.

Figure S4. SEM micrographs of PEOT/PBT_s and composite scaffolds depicting the scaffolds' filaments surface roughness. Insert micrographs represent the scaffolds cross section. Scale bars 100 μm. insert corresponding to low magnification images. Sale bars 1 mm.

Figure S5. (A) Cumulative release (%) of CFX and GTM from MgAl-CFX_s and ZrP-GTM_s, respectively. (B) Kinetic models used to fit the antibiotic release profiles from 20% MgAl-CFX_s and 20% ZrP-GTM_s (stage I only) and corresponding correlation coefficients (R²). *t* refers to time, *M_t* refers to amount of drug released in time *t*, *M₀* refers to the total amount of drug loaded in the system, and *k*, *a*, and *n* are model-specific constants.

Figure S6. Thermal stability of CFX. (A) TGA curves of pure CFX and MgAl-CFX. (B) HPLC of pure CFX and CFX extracted from MgAl-CFX, 20% MgAl-CFX_p and 20% MgAl-CFX_s. (C) Percentage of complete CFX molecule recovery after each processing step.

Figure S7. Thermal stability of GTM. (A) TGA curves of pure GTM and ZrP-GTM. (B) HPLC-MS of pure GTM and GTM extracted from ZrP-GTM, 20% ZrP-GTM_p and 20% ZrP-GTM_s. (C) Relative amount of each GTM isoform intercalated in ZrP after each processing step. (D) Percentage of complete GTM molecule recovery after each processing step.

Figure S8. Non cumulative release of (A) CFX and (B) GTM over time from MgAl-CFX_s and ZrP-GTM_s, respectively.

Figure S9. DNA content after 1 and 7 days of culture in BM in (A) MgAl-CFX_s and (B) ZrP-GTM_s, directly seeded without pre-incubation in media. Statistical significance performed using two-way ANOVA with Tukey's multiple comparison test (* $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$; **** $p < 0.0001$).

Figure S10. Dose dependence of toxicity of (A) CFX and (B) GTM towards hMSCs, in their pure form (not exposed to T) or previously exposed to 200 °C for 30 min under a N₂ atmosphere (exposed to T).

Figure S11. Stereomicroscopy images of (A) PEOT/PBT_s and MgAl-CFX_s and (B) ZrP-GTM_s cross sections stained with ARS after 35 days of culture (28 days in MM). Inserts represent the

corresponding control scaffolds without cells incubated in MM and stained with ARS. Scale bars 1 mm. (C, D) Quantification of the ARS extracted from scaffolds after 35 days of culture (28 days in MM), normalized to total cell number. Statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (** $p < 0.01$; *** $p < 0.001$; **** $p < 0.0001$).

Figure S12. (A) DNA content and (B) ALP activity on MgAl-CFX_s after 56 days of culture (49 days in MM). (C) DNA content and (D) ALP activity on ZrP-GTM_s after 42 days of culture (35 days in MM). Statistical significance performed using one-way ANOVA with Tukey's multiple comparison test (n.s. $p > 0.05$, ** $p < 0.01$).

Figure S13. Representative confocal microscopy images of hMSCs (F-actin, green) on top of the filaments of (A) 5 and 10 % MgAl-CFX_s after 56 days of culture (49 days in MM) and (B) 5 and 10 % ZrP-GTM_s after 42 days of culture (35 days in MM) stained for relevant osteogenic markers COL1 (red) and OPN (blue). Scale bars 100 μ m.

Table S1. Antimicrobial activity of MgAl-CFX and ZrP-GTM at different concentrations against *S. epidermidis* and *P. aeruginosa*.

Samples (0.1 g/ml)	<i>S. Epidermidis</i> R (log)	<i>P. Aeruginosa</i> R (log)
ZrP-GTM 5mg/ml	>4.7	>6.7
ZrP-GTM 10/ml	>4.7	>6.7
ZrP-GTM 20m g/ml	>4.7	>6.7
MgAl-CFX 5mg/ml	>4.8	>5.8
MgAl-CFX 10 mg/ml	>4.8	>5.8
MgAl-CFX 20 mg/ml	>4.8	>5.8

Table S2. Antimicrobial activity of MgAL-CFX_f and ZrP-GTM_f against *S. epidermidis* and *P. aeruginosa*.

Samples (0.1 g/ml)	<i>S. Epidermidis</i> R (log)	<i>P. Aeruginosa</i> R (log)
5% MgAl-CFX_f	1.08	>6.85
10% MgAl-CFX_f	1.51	>6.85
20% MgAl-CFX_f	5.51	>6.85
5% ZrP-GTM_f	>4.94	1.95
10% ZrP-GTM_f	>4.94	3.5
20% ZrP-GTM_f	>4.94	4.89

Table S3. Susceptibility of *S. epidermidis* and *P. aeruginosa* and strains to pure GTM and CFX evaluated by the disk diffusion agar test.

Concentration ($\mu\text{g/ml}$)	In disc (μg)	S.Epidermidis				P. Aeruginosa			
		GTM		CFX		GTM		CFX	
		ZOI (mm)		ZOI (mm)		ZOI (mm)		ZOI (mm)	
		Av.	s.d.	Av.	s.d.	Av.	s.d.	Av.	s.d.
500	10	16	1.2	33	2.0	9	0.0	25	0.6
250	5	16	1.2	30	0.6	8	0.5	25	1.0
100	2	12	0.0	25	0.6	6	0.0	23	1.0
75	1.5	11	0.6	21	0.6	6	0.0	22	1.5
50	1	10	0.6	18	0.0	6	0.0	18	0.6
25	0.5	10	1.5	16	0.6	6	0.0	16	0.6
15	0.3	7	1.2	12	0.6	6	0.0	14	0.6
10	0.2	6	0.6	6	0.0	6	0.0	13	0.0
5	0.1	6	0.0	6	0.0	6	0.0	6	0.0
0	0	6	0.0	6	0.0	6	0.0	6	0.0

Table S4. Non cumulative release of CFX over time from 20 wt% MgAl-CFX_s and the corresponding experimental and expected ZOI against *S. epidermidis* and *P. aeruginosa*. Expected ZOI were estimated from the susceptibility test in Table S3.

20% MgAl-CFX_s								
time (days)	CFX release (ug)		S. Epidermidis			P. Aeruginosa		
			ZOI experimental (mm)		ZOI expected (mm)	ZOI experimental (mm)		ZOI expected (mm)
	Av.	s.d.	Av.	s.d.		Av.	s.d.	
1	62.02	3.89	21.3	0.6	18-21	22.0	1.0	18-22
2	31.37	0.75	20.0	0.0	16-18	15.3	0.6	16-18
3	36.38	1.40	21.3	0.6	16-18	17.7	0.6	16-18
7	63.67	3.23	24.0	1.0	18-22	22.0	1.0	18-22
10	52.46	1.98	20.7	1.2	18	20.0	1.0	18
14	51.06	2.50	28.3	0.6	18	18.3	0.6	18
17	43.99	2.73	24.0	1.0	16-18	19.7	0.6	16-18
21	42.93	2.27	22.0	1.0	16-18	20.3	0.6	16-18
24	32.20	1.25	23.3	1.2	16-18	22.3	0.6	16-18
29	37.34	2.86	23.3	0.6	16-18	18.0	1.0	16-18

Table S5. Non cumulative release of GTM over time from 20 wt% ZrP-GTM_s and the corresponding experimental and expected ZOI against *S. epidermidis* and *P. aeruginosa*. Expected ZOI were estimated from the susceptibility test in Table S3.

20% ZrP-GTM_s								
Time (days)	GTM release (ug)		S. Epidermidis			P. Aeruginosa		
			ZOI experimental (mm)		ZOI expected (mm)	ZOI experimental (mm)		ZOI expected (mm)
	Av.	s.d.	Av.	s.d.		Av.	s.d.	
1	491.63	18.00	16.7	0.6	16	11.0	0.0	9
2	28.50	1.18	11.3	0.6	10	6.0	0.0	6
3	14.34	1.92	8.3	0.6	7	6.0	0.0	6
7	25.50	7.93	8.3	0.6	10	6.0	0.0	6
10	10.69	0.44	7.0	0.0	6	6.0	0.0	6
14	9.50	1.03	7.0	0.0	6	6.0	0.0	6
17	5.24	0.77	6.7	0.6	6	6.0	0.0	6
21	9.93	5.64	7.3	0.6	6	6.0	0.0	6
24	4.57	0.75	6.0	0.0	6	6.0	0.0	6
29	5.04	1.19	6.0	0.0	6	6.0	0.0	6

Table S6. Theoretical and experimental (determined by TGA) MgAl-CFX and ZrP-GTM loadings of MgAl-CFX_p and ZrP-GTM_p at different filler-antibiotic concentrations.

	Theoretical loading (%)	Experimental loading (%)
5% MgAl-CFX_p	5	5.4
10% MgAl-CFX_p	10	7.1
20% MgAl-CFX_p	20	20.9
5% ZrP-GTM_p	5	8.0
10% ZrP-GTM_p	10	11.3
20% ZrP-GTM_p	20	18.6

Chapter 8

General Discussion

In this Chapter, the most important findings of the thesis will be discussed and contextualized within the overall aim of the thesis, which is the design and fabrication of bioactive melt extrusion additive manufactured (ME-AM) scaffolds for bone tissue regeneration. We highlight the reasons for material and cell source choice, the importance of cell confluency in osteogenic differentiation, and the unique functionality provided by the investigated scaffold modifications.

Why PEOT/PBT?

As noticed in the literature review within **Chapter 2**, poly(ϵ -caprolactone) (PCL) is the most used biodegradable thermoplastic polymer to fabricate ME-AM scaffolds for bone tissue engineering (BTE) applications. The main motivation for researchers to invest on this polymer is its approval by FDA or CE mark certification for drug delivery devices, tissue adhesion barriers, and 3D printed implants for bone regeneration, which consequently smoothens the clinical and marketing path of any subsequent newly designed PCL based scaffolds. However, most of the studies have been performed using research-grade PCL, which has been shown to differ from medical-grade PCL in terms of molecular weight, impurities content, physicochemical properties (mechanical properties, degradation) and processing parameters, often leading to results being unsuitable for translation.¹⁻² Other important reasons for PCL monopolizing ME-AM BTE research are its high compression strength; long degradation time; and its low printing temperature, which diminishes the printer hardware demands.² Despite these appealing characteristics, PCL also possess some drawbacks, including its poor cell adhesion and limited cell spreading,³ which are non-optimal properties for applications in which strong cell adhesion and spreading are required, such as bone formation.⁴ For these reasons, the exploration of other synthetic polymers towards the design of an optimum scaffold for BTE is still desired. One of these polymers is the block copolymer poly(ethylene oxide terephthalate)/poly(butylene terephthalate) (PEOT/PBT), which is commercially available in medical-grade, as well as approved by regulatory authorities in the composition 1000PEOT70PBT30 (1000 kDa PEO segment and a PEOT:PBT ratio of 70:30) for its use as drug delivery system, cement restrictor and 3D printed scaffold for cartilage regeneration. PEOT/PBT copolymers are thermoplastic, biocompatible and biodegradable, and their mechanical, physicochemical and swelling properties can be tuned within a wide range by varying the PEO segment molecular weight, as well as the PEOT:PBT ratio, to meet the requirements of a specific regenerative application.⁵⁻⁸ Taking this into consideration, PEOT/PBT and scaffolds made

from these materials have proven their *in vitro* and *in vivo* applicability in skeletal tissue regeneration, including bone⁹⁻¹⁶, cartilage¹⁷⁻²⁰ and osteochondral tissue engineering,²¹⁻²⁴ as well as in skin,²⁵ and nerve regeneration,²⁶ or islets of Langerhans delivery.²⁷⁻²⁹ Among other formulations, the PEOT/PBT used within this thesis, 300PEOT55PBT45 (300 kDa PEO segment and a PEOT:PBT ratio of 55:45), constitutes a valid alternative to PCL for the fabrication of ME-AM scaffolds for BTE. This is owed to its long degradation time (3% mass loss after 24 weeks implantation *in vivo*), optimum mechanical properties (tensile modulus ~ 50 MPa, compression modulus ~ 120 MPa), and good cell attachment and spreading *in vitro* without further functionalization.^{5-8, 10} Furthermore, it has a relatively low melting temperature (~160 °C) and low viscosity, which enables its easy processing through standard ME-AM platforms at 165 -195 °C.³⁰⁻³¹

Why human mesenchymal stromal cells (hMSCs)?

Because of their ability to differentiate into osteoblasts and their role in tissue regeneration, hMSCs are commonly used in tissue engineering.³² hMSCs have the ability to migrate chemotactically to injured tissues, where they differentiate into tissue-specific cells, and help in the regeneration process by the secretion of factors, which can modulate the inflammatory response, promote vascularization, and induce proliferation and differentiation of other endogenous cell types.³³⁻³⁴ In an attempt to support the endogenous homing of hMSCs, BTE approaches have considered the direct injection to the injured site of these cells or, most interestingly, the implantation of hMSC-laden scaffolds.³⁵⁻³⁶ When combined with hMSCs, the scaffold would not only mechanically support the fracture and promote a material-induced bone growth response, but also ideally boost the implanted hMSCs and, subsequently, other endogenous cell types, towards an accelerated bone regeneration response. hMSCs can be extracted and isolated from the patient's own bone marrow without leading to donor site morbidity, unlike other autografts, and can be expanded in culture to generate sufficient numbers while maintaining their multipotency.³⁵

Prior to implantation, and regardless of the final scaffold application being cell-free or cell-laden, the functionality of a newly designed scaffold is usually tested by seeding it with hMSCs and evaluating these cells ability to undergo osteogenic differentiation on the scaffold.³⁷⁻³⁸ Several reports have already dedicated initial efforts to evaluate this way the *in vitro* biological performance of PEOT/PBT (300PEOT55PBT45) ME-AM scaffolds for

BTE.^{13-16, 21-24} This common *in vitro* practice gives hints on the osteoinductive and osteoconductive properties of the scaffold, i.e. the scaffold's ability to recruit bone forming cells or simply support bone formation on scaffold surface, respectively. Accordingly, hMSCs were used in this thesis to test the developed ME-AM scaffold's ability to support or induce bone formation *in vitro*.

Addressing the cell seeding problem on ME-AM scaffolds

A challenge encountered when seeding hMSCs on PEOT/PBT ME-AM scaffolds was the poor cell attachment along the scaffold cross section, with the majority of cells sedimenting and forming a monolayer at the scaffold's bottom. This is a persistent issue associated with the macroporosity and synthetic nature of polymeric ME-AM scaffolds, as well as to gravity,^{22, 39} which up to now has been managed by architectural or surface scaffold modifications, or by complicated seeding methods. **Chapter 3** of this thesis has demonstrated that by simply altering the physical properties of the seeding media (viscosity or density), a significant enhancement on cell distribution along the scaffold cross section can be attained upon static seeding. This was possible by including the macromolecules (macroM) dextran or Ficoll to the seeding media at specific concentrations, which dramatically reduced hMSCs settling velocity, allowing cells to float longer in the suspension and giving them more time to attach to the scaffold filaments. This approach differs from previous efforts in terms of: i) its simplicity, as it does not require of additional equipment, such as a rocking bed for dynamic seeding, a vacuum chamber for plasma treatment, or complicated chemistry for a hydrogel carrier formation; ii) its universality, as it can be used for any scaffold type, regardless of its material and architecture; iii) its efficiency, as it avoids the use of large cell stocks; and iv) its temporary activity, as none of the supplements attach to the scaffold filaments and can be quickly expelled by cells when endocytosed.

Alternatively, enhanced cell attachment and distribution was achieved by fabricating PEOT/PBT scaffolds via atmospheric pressure plasma assisted AM in **Chapter 4**. In this case, cell attachment was induced by the deposition of hydrophilic and protein adhesive coatings on the scaffolds filaments, generated by plasma-polymerization, prior to conventional static seeding. Plasma had been previously used to treat 3D ME-AM scaffolds,⁴⁰⁻⁴² but with our approach, we have tackled several issues that had been previously identified when this surface modification technique was used to treat scaffolds.

First, the use of an atmospheric pressure plasma jet (APPJ) allowed to direct the plasma flame into the scaffold pores, thus avoiding the inhomogeneous and uncontrolled gradient functionalization along the scaffold volume resulting from conventional vacuum plasma.⁴³ Furthermore, we investigated the use of plasma polymerization, which offers a more specific surface functionalization via the deposition of specific functional groups, in contrast to plasma activation with inert gases. Specifically, we employed organosilane precursors ((3-Aminopropyl) trimethoxysilane (APTMS) and a mixture of maleic anhydride and vinyltrimethoxysilane (MA-VTMOS)), which provide stable coatings resistant to delamination in aqueous conditions compared to other precursors.⁴⁴⁻⁴⁵ In this way, we were able to precisely analyze, for the first time, cell attachment to 3D ME-AM scaffolds as a function of specific surface charge and functional groups, in the presence or absence of serum. In particular, we observed that all plasma treatments increased the surface wettability of the scaffolds, enhanced cell-adhesive proteins adsorption to their surface, and allowed for homogeneous cell attachment along the scaffold cross section. Interestingly, APTMS treated scaffolds allowed cell attachment in the absence of serum, driven by electrostatic interactions between the cell and the charged scaffold surface, which possess relevant clinical applications.

Homogeneous cell distribution and cell confluency in 3D ME-AM scaffolds are necessary for MSCs osteogenic differentiation and mineralization

Although both macroM seeding and plasma treatment improved homogenous cell coverage on the PEOT/PBT scaffolds and led to high seeding efficiencies of 70-80%, macroM seeding performed better at retaining cells along the scaffold cross section. While a large portion of the seeded cells adhered to the plasma treated scaffolds' bottom in form of a monolayer, reducing the cell number attached to the filaments throughout the scaffold's height, no monolayer was formed in the macroM seeded scaffolds. This was translated into a larger cell density and cell-cell contact points on the filaments of macroM seeded scaffolds, where cell agglomerates were already observed after seeding, and cells confluent covered the filaments after 1 day of culture. At the same timepoint, cell-cell contact was less abundant on the significantly less crowded plasma-treated scaffolds' filaments. Cell density and cell-cell contact in 2D cultures have been previously observed to greatly influence the osteogenic potential of hMSCs. According to the results in **Chapter 3** and **Chapter 4**, we hypothesized that these parameters also played a key role in the differentiation of hMSCs on our ME-AM 3D scaffolds. Here, calcium deposition, a late

marker for osteogenic differentiation, was observed only on scaffolds with high cell confluency on their filaments (macroM seeding), while osteogenic differentiation was delayed and calcium deposition was not observed on scaffolds with less cell density (plasma-assisted seeding), even after longer culture periods (54 days in MM). In 2D, studies have reported increased alkaline phosphatase (ALP) activity with increasing number of cell-cell contacts.⁴⁶⁻⁴⁷ Similarly, hMSCs plated at high density and cultured in osteogenic media have shown to lead to enhanced calcium deposition, compared to hMSCs plated at lower density.⁴⁸⁻⁵⁰ In particular, 100% confluent hMSCs cultures have resulted into higher calcium deposition and osteocalcin protein expression, compared to lower confluency values,⁵¹ and over-crowding (> 100%) has been shown to maintain the osteogenic differentiation potential of hMSCs.⁵²⁻⁵³ The upregulation of osteogenic markers through cell density variations in 2D has been explained by the activation of cell contact-dependent signaling, such as gap junctions,⁴⁶ and the activation of integrin⁵⁰⁻⁵¹ and cadherin⁵⁴ signaling pathways, which are all involved in the regulation of osteogenic differentiation.⁵⁵⁻⁵⁷

While cell studies in 2D have clearly demonstrated the positive correlation between cell confluency and enhanced osteogenic differentiation and calcium deposition, such effects have not been clearly observed so far in 3D. Some researchers have reported the detrimental effect of high seeding density on the ALP activity in osteoblasts or hMSCs, when cultured on 3D scaffolds,⁵⁸⁻⁵⁹ while others have shown a positive impact of high cell seeding density on hMSC osteogenic differentiation.⁶⁰⁻⁶³ Overall, we hypothesize that these contradictory results in 3D are mainly owed to a lack of homogeneous cell distribution along the scaffold volume, which results in an inhomogeneous cell population and an erroneous averaged analysis. In this regard, we believe that the homogenous cell distribution attained with the macroM seeding method in **Chapter 3** and with the plasma-assisted conventional seeding method in **Chapter 4**, allow us to draw more accurate conclusions on the effect of cell density, confluency and cell-cell contact on hMSCs osteogenic differentiation in 3D scaffolds, in particular in 3D ME-AM scaffolds. Taking this into account, it can be hypothesized that cell confluency is necessary for hMSCs osteogenic differentiation and calcium deposition on 3D AM-scaffolds, and this effect is potentially driven through the same mechanisms as in 2D.

Cell confluency vs. other pro-osteogenic cues: in the race to osteogenic differentiation

It cannot be dismissed that plasma-treated seeded scaffolds come along with an extra variable that has previously been shown to affect osteogenic differentiation on its own, i.e. surface chemistry. Functionalized 2D surfaces with polar functional groups have been shown to promote osteogenic differentiation, through an enhanced exposure of the integrin binding domains of adsorbed cell-adhesion proteins, which can lead to high levels of focal adhesion components and phosphorylated focal adhesion kinase (p-FAK) in the adhered cells, required for osteoblast differentiation.⁶⁴⁻⁶⁶ While in **Chapter 4** we hypothesized that the lower protein adhesion to aged scaffolds (where osteogenic differentiation was evaluated) might have hampered MSC osteogenic stimulation and calcium deposition, it is also possible that the surface chemistry's pro-osteogenic effect is not as strong as the effect of cell confluency. Some studies in 2D have pointed out that substrates with increasing amount of polar functional groups or surface roughness give rise not only to enhanced calcium deposition, but also to increased cell attachment and density.⁶⁷⁻⁶⁸ This makes it difficult to identify the relevant osteogenic cues, among surface properties and cell confluency. Besides these, other cues that have been found to influence hMSCs osteogenic differentiation are, for example, substrate stiffness, substrate roughness, cell shape, and cell size. In some studies, where cells were exposed to competing pro-differentiation signals, cell-cell contact has been found to be a powerful cue, overruling the effect of other signals. For instance, through alterations in cytoskeletal tension, ALP and RUNX2 gene expression have been proven to be upregulated on cells with a large spread area, large perimeter, or with an aspect ratio around 2, compared to other shapes or sizes.⁶⁹⁻⁷¹ Yet, hMSCs cultured at high density also displayed the upregulation of osteogenic markers, together with calcium deposition, in spite of the physical constraints imposed by an increase density leading to "non-osteogenic morphologies" (i.e. small cell spread areas).⁷⁰ Similarly, stiff substrates have been previously shown to direct hMSCs towards the osteogenic lineage, according to the upregulation of osteogenic markers.⁷² However, hMSCs cultured on stiff substrates were capable of depositing high amounts of calcium only when cultured at high density.⁴⁹

The fact that cell confluency can override other pro-osteogenic factors in ME-AM scaffolds was further supported by the results observed in **Chapter 5** and **Chapter 6**. In **Chapter 5**, hMSCs were seeded on PEOT/PBT-nanohydroxyapatite (nHA) composite scaffolds using

the macroM seeding method and osteogenic differentiation was analyzed as a function of the HA content. Similarly, in **Chapter 6** PEOT/PBT-reduced graphene oxide (rGO) composite scaffolds were seeded with hMSCs by the macroM seeding method, and osteogenic differentiation was assessed. While some previous studies had shown that the addition of HA and graphene based fillers to ME-AM composite scaffolds lead to the upregulation of osteogenic markers in hMSCs or other cell types *in vitro*,⁷³⁻⁷⁶ we did not observe significant differences in the osteogenic profile of these cells cultured on the composite scaffolds, compared to bare PEOT/PBT scaffolds. Indeed, hMSCs could undergo osteogenic differentiation and deposit a mineralized extracellular matrix (ECM) at the same levels on every scaffold type, regardless of the presence or absence of the fillers. One reason might be the polymer matrix being a barrier for fillers release in nHA containing scaffolds due to the poor exposure of the nHA particles to the surface, as suggested in **Chapter 5** and by previous reports.^{74, 77} This, together with the slow degradation time of nHA, possibly led to suboptimal ion exchange dynamics between the scaffold and the medium and a calcium concentration in the medium, which was insufficient to affect calcium signaling pathways involved in osteogenesis. Nevertheless, calcium phosphate (CaP) re-precipitation after incubation in simulated body fluid was an indicator of the potential of these highly loaded HA scaffolds *in vivo*, in terms of tissue integration and osteoconduction. Another possible reason why differences were not observed in the osteogenic outcome between bare polymeric and filler-containing scaffolds could be correlated to the variable of seeding density being ruled out by seeding all the scaffold types with the macroM seeding method. In addition to accelerated osteogenic differentiation, previous studies have also observed enhanced cell attachment on such composite scaffolds, due to the fillers increasing the scaffold hydrophilicity and surface physical properties.⁷⁸⁻⁸⁰ Therefore, it is plausible that the observed accelerated cell differentiation on these composite scaffolds was induced by the higher cell density and confluency indirectly promoted by the fillers. Currently ongoing animal studies, in which such scaffolds were implanted in critical size ulnar defects in rabbits, will allow us to draw better conclusions on the bone regeneration performance of HA based composite scaffolds, compared to bare polymeric scaffolds.

These results raise an important question: if cell confluency is the main pro-osteogenic factor in ME-AM scaffolds, and it can be attained by simple strategies, such as macroM seeding on bare polymeric scaffolds, would it be necessary to invest on further scaffold modification strategies to regenerate bone, such as plasma treatments or the use of

composite materials? Before answering this question, it is worth to mention that osteogenic differentiation was only attained in macroM seeded scaffolds when hMSCs were provided with osteogenic medium (**Chapter 3** and **Chapter 5**), and it is not known whether endogenous factors would be sufficient to trigger bone formation in confluent scaffolds in an *in vivo* scenario. Previous *in vivo* studies evaluating the bone regeneration ability of cell-laden polymeric ME-AM scaffolds (seeded with hydrogel carriers or following protein coatings) have shown contradictory results, and it is not clear yet whether the presence of cells along with the scaffold has overall beneficial effects.⁸¹⁻⁸⁴ Nevertheless, these studies have never assessed the state of cell confluency prior to scaffold implantation, and there is still no consensus on the optimum cell number, state of cell differentiation (the use of pre-differentiated or undifferentiated cells), and amount of viable cells after transplantation.⁸⁵ This highlights the necessity of future research on more controlled and systematic studies to investigate all these parameters. We believe that the macroM seeding method could serve as a reliable platform to study the bone regeneration capabilities of cell-laden scaffolds and to unravel the importance of cell density and confluency *in vitro* and *in vivo*. Given that the effect on *in vivo* bone regeneration of cell confluency in 3D scaffolds is still unknown, and that the use of the macroM seeding would be only applicable for cell-laden scaffolds strategies, the answer to the aforementioned question would be: yes, the investigation of other modification strategies such as plasma or the use of composites is still required.

Unique functionalities of plasma-modified and composite ME-AM scaffolds

Plasma for better tissue integration

As presented in **Chapter 4**, plasma activation and polymerization enhanced the surface energy and wettability of polymeric scaffolds, and led to protein adsorption and enhanced cell adhesion *in vitro*. *In vivo*, this could be translated into better scaffold integration with the existing bone defect ends and surrounding tissue (also called osseointegration),⁸⁶⁻⁸⁸ which is critical for implant stability and is considered a prerequisite for implant loading and long-term clinical success. Moreover, these plasma modifications can promote endogenous cell adhesion, as a first step towards osteogenesis and bone formation. Notably, the aged scaffolds still showed to provide enhanced cell adhesion compared to untreated scaffolds. This experiment simulated the clinical scenario of scaffold production, where the storage after surface modification may occur for several days prior the

implantation into the patient's defect. The automated production of scaffolds within an integrated ME-AM and APPJ platform further validates the translational potential of such plasma-modified scaffolds. Although not presented in this study, this integrated platform also allows to treat the scaffold layer by layer, to reach deep pores in large anatomical scaffolds, or zonally, to facilitate cell attachment in specific scaffold regions.⁸⁹

Composites for mechanical properties

Among all the ME-AM scaffold modification strategies presented in **Chapter 2**, the incorporation of bioactive fillers within the scaffolds' polymer matrix towards composite scaffolds is the only method that enables increasing the scaffold mechanical properties without compromising its architecture and porosity. This is of great importance since, regardless of the polymer type used, polymeric ME-AM scaffolds still possess very low mechanical properties compared to bone, whereas scaffolds with high mechanical strength are fundamental for load bearing scenarios. In **Chapter 5**, we observed how the addition of HA led to a significant increase in the scaffold compression modulus and strength at yield, reaching the range of cortical bone mechanical properties. Although not discussed within this thesis, the addition of rGO in **Chapter 6** and of lamellar inorganic fillers in **Chapter 7** can also improve the mechanical properties of the PEOT/PBT scaffolds, in terms of compression modulus and yield compression strength.⁹⁰

Electrically conductive scaffolds

In addition to improving their mechanical properties, the incorporation of fillers can provide the scaffold with other functional properties, such as electrical conductivity. In **Chapter 6**, polymeric scaffolds were rendered conductive with the addition of rGO fillers. Bone is a piezoelectric material and can generate electric potentials in response to mechanical stress, promoting bone resorption or formation.⁹¹ Moreover, stem cell attachment, migration, proliferation, and differentiation have been shown to be influenced by electrical stimulation through the modification of the electrochemical environment, resulting in the activation of ion transporting channels, and membrane receptor complexes.⁹² For all these reasons, electrical stimulation has been considered as a tool in the clinics to stimulate bone healing, and its combination with electrically conductive scaffolds opened the door to a more localized therapy.⁹³ During the last decades, scaffolds prepared through conventional fabrication methods and coated with conductive polymers, such as poly(3,4-ethylenedioxythiophene) PEDOT, polypyrrole (PPy), and polyaniline (PANI) have been explored as conductive substrates for bone regeneration.⁹⁴ As a result of the

establishment of AM as preferred scaffold fabrication technology, together with the rise of carbon-based materials as desired materials for biomedical applications, the field has recently been focused on the investigation of electrically conductive carbon based composite ME-AM scaffolds.⁹⁵⁻⁹⁶ The incorporation of the conductive material within the polymer matrix, rather than as a coating, provides mechanical reinforcement to the scaffold, enables to reduce its exposed amount ensuring higher biocompatibility, and avoids coating delamination due to weak interactions with the underlying substrate. Most recent studies have explored the *in vivo* bone formation capabilities of PCL/graphene and PCL/multiwall carbon nanotubes (MWCNT) ME-AM scaffolds combined with electrical stimulation.⁹⁷⁻⁹⁸ The presented results confirm the promising character of this line of research, to which we have contributed within this thesis mostly from a materials science point of view. Specifically, within **Chapter 6** we evaluated how the densification of rGO into a more dosable and less inhalation-hazardous material affects the resulting composite scaffold printability, conductivity and physicochemical properties. Among different graphene based fillers, rGO is preferred due to its electrical conductivity, thermal stability, and availability at large scale through established synthesis methods. Our study reveals that while increasing the rGO bulk density leads to a more dosable material and to composites with a wider printability window, it also results in scaffolds with lower electrical conductivity, at a given wt%. The solvent used for densification also showed to play a role in rGO compaction and subsequent dispersability within the composites. In this regard, for a given bulk density, densification in water resulted in a more compacted and aggregated rGO material than when densified in acetone, which reduced its dispersion within the polymer matrix and printability window. Overall, our study establishes relevant knowledge for potential future studies using rGO as a filler, and provides insight on the relevance of the rGO bulk density and volume occupancy for attaining ME-AM scaffolds with specific properties.

Antimicrobial scaffolds for managing infections

Beyond providing cues to promote bone formation, composite scaffolds can be used as drug carriers. In the clinics, up to 20% of open bone fractures, surgeries for fracture internal fixation, or implants revision surgeries develop into an infection.⁹⁹ The gold standard clinical strategy to manage infections consists of implant removal, tissue debridement and defect filling with a bone cement (poly(methyl methacrylate) (PMMA)) impregnated with antibiotics, which allows the reduction of systemic antibiotic administration.¹⁰⁰⁻¹⁰¹ With the

advances in biodegradable calcium-based bone grafts in the last decade, some of these products have included antibiotics in their composition to treat osteomyelitis, while allowing bone regeneration in the defect for avoiding a revision surgery.¹⁰¹ Clinical studies have proven the ability of such systems (based on calcium sulfate, alone or combined with CaPs) in eradicating infections in 90-100% of the cases. Yet, bone healing and defect filling in non-unions was not always achieved, showing success rates ranging from 34 to 100%.¹⁰²⁻¹⁰⁴ The fast degradation time of these bone void fillers (3-6 months), together with their low mechanical properties, have been identified as responsible parameters leading to failure in the bone regeneration process. For this reason, slower degrading and stiffer ME-AM scaffolds with antibiotic-eluting properties that can support the defect during the bone formation process are considered as more optimum alternatives. Generally, ME-AM processes to obtain such scaffolds are limited to a few low melting temperature polymers, mainly PCL, in order to maintain the stability of the dispersed heat-sensitive drugs. As an alternative to developing new polymers with lower melting temperatures, or adding biocompatible plasticizers to the composites for tackling thermal stability issues,¹⁰⁵⁻¹⁰⁶ we have investigated the use of lamellar protectors. In this regard, composite PEOT/PBT ME-AM scaffolds with lamellar inorganic fillers (magnesium aluminum layered double hydroxide (MgAl) and α -Zirconium phosphate (ZrP)) intercalated with antibiotics (ciprofloxacin (CFX) and gentamicin (GTM), respectively) were presented in **Chapter 7**. These fillers had been previously shown to increase the thermal stability of sensitive drugs.¹⁰⁷⁻¹⁰⁸ We also showed that the antibiotics maintained their antimicrobial activity after high temperature ME-AM, which was proven against some of the most common bacteria responsible for osteomyelitis (*S. epidermidis* and *P. aeruginosa*). In addition to this, we have shown that the fillers allowed for a more sustained antibiotics release, especially on scaffolds containing MgAl-CFX. The release profile was further tunable by adjusting the weight percentage of filler-antibiotic complexes within the scaffolds, as well as by the ionic concentration and pH changes in the surrounding environment. Such controlled antibiotic eluting systems are ideal, as they avoid potentially cytotoxic burst releases and prolong antimicrobial activity over time. Yet, such strategies require further development since standard antibiotic therapy for osteomyelitis lasts for 4-12 weeks,^{99, 109} a period that is much shorter than the time PEOT/PBT scaffolds' take to degrade, which is required for complete antibiotic release. In vivo, the PEOT/PBT used in this study (PEO molecular weight = 300 kDa, PEOT:PBT weight ratio= 55:45) showed only a 3% mass loss after 24 weeks of subcutaneous implantation in rats (film of thickness 400-600 μ m).¹¹⁰ It is possible

that our scaffolds porous structure, with increased surface exposed to surrounding body fluids, lower filament diameter (max 350 μm), and increased hydrophilicity given by the fillers, display shorter degradation times. In this regard, a careful evaluation of the scaffold degradation and antibiotic release in vivo is necessary, since a slow polymer degradation time can lead to a prolonged antibiotic treatment with local concentrations lower than the minimal inhibitory concentration (MIC), which can lead subsequently to the creation of antibiotic-resistant bacteria.¹¹¹ A couple of approaches could be considered to accelerate the scaffold degradation time, and therefore the translation potential of such scaffolds. Inspired by other scaffold modification strategies, microporosity could be introduced in the composite scaffolds' filaments to enhance water permeability. Alternatively, a faster degrading thermoplastic material could be selected, such as PEOT/PBT with higher PEO molecular weight, or poly(lactic-co-glycolic) acid (PLGA) copolymers.^{110, 112} Furthermore, the filler-antibiotic particles could be coated on the scaffold filaments, rather than being dispersed within the polymer matrix. Here, the fillers would still control the antibiotic release and prevent burst release, but the overall release would be completed faster due to the absence of the polymer matrix barrier. While the first suggested options would probably decrease the scaffold mechanical properties, the latter would not allow to control the amount of loaded antibiotic, which would mainly depend on the scaffold surface available area. Overall, it is evident that due to the control needed on multiple interdependent properties, tradeoffs are needed to be made to introduce antimicrobial properties to the scaffolds. Nevertheless, we encourage further investigations on the use of lamellar inorganic fillers for the incorporation of antibiotics within ME-AM scaffolds towards the development of an optimum system, as they are a reliable strategy to introduce bioactive molecules within high temperature melting polymers.

In the future

Multifunctional scaffolds

Looking to the future, a combination of the aforementioned scaffold modification strategies into a single scaffold is envisioned towards further enhancing their augmented bioactivity. For instance, a composite scaffold combining nHA and rGO, would give rise to a multifunctional scaffold with electrical and antimicrobial properties (given by rGO), and mechanical and osteogenic properties, given by both nHA and rGO. The synergistic effect of nHA and rGO added to the cell culture media in osteogenic differentiation in 2D has

been previously shown,¹¹³ and such a binary combination has been already presented in 3D solvent based AM scaffolds.¹¹⁴ The combination of antimicrobial agents with nHA has also been investigated in solvent based AM scaffolds. In this regard, PLGA/HA scaffolds coated with quaternized chitosan proved their bone regeneration capabilities and infection management in an infected *in vivo* bone model.¹¹⁵⁻¹¹⁶ This suggests the potential of combining our filler-antibiotic and HA into one composite scaffold for osteogenic and antimicrobial properties. Such binary composite scaffolds could be easily prepared by loading the single-filler composites used within this thesis into the double reservoir print-head developed by our group.⁸⁹ This print-head would allow the *in situ* mixing of two materials at desired ratios towards a binary scaffold, simple or graded, avoiding the pre-printing fabrication of binary composites by compounding. Ultimately, plasma treatment could further enhance the bioactivity of any of the suggested single or binary composite scaffolds.

Other CaP fillers

Due to the slow degradation time of nHA compared to other CaP fillers, such as tricalcium phosphate (TCP) or biphasic calcium phosphate (BCP), HA-based ME-AM scaffolds might maintain over a longer period of time their initial structural and mechanical properties, which are beneficial to support the fracture site while the new bone formation volume is still minimal. However, results in **Chapter 5** have also suggested that the slow dissolution of HA might be hampering the osteoinductive potential of HA-based ME-AM scaffolds. Very few studies have compared side by side the osteogenic differentiation or bone regeneration capacity of ME-AM polymeric scaffolds containing different CaP fillers, and no definitive conclusions can be obtained from here.¹¹⁷ However, reports comparing the biological performance of different porous CaP scaffolds have shown that BCP or TCP scaffolds, with more optimum calcium exchange dynamics, outperform HA scaffolds in their *in vitro* and *in vivo* bone formation capacity.¹¹⁸⁻¹²⁰ While the use of TCP as filler for ME-AM scaffolds has been studied in various reports, BCP, which is a combination of TCP and HA has not been so far a common filler of choice. In future studies, an exhaustive *in vitro* biological evaluation of ME-AM scaffolds containing BCP with different TCP/HA ratios would be of interest. By adjusting the TCP/HA ratio, a balance in mechanical properties, scaffold degradation time and ion exchange dynamics could be struck, towards an optimum ME-AM scaffold for BTE. The supplementation of the CaP filler with bioinorganics, such as strontium, silicon or magnesium, which are found in trace

amounts in bone ECM and have shown to improve the bone regeneration capacity of CaPs, ¹²¹ has also not been investigated in ME-AM scaffolds and could also be considered in the future.

Other relevant antibiotics

Future studies could be dedicated to investigate the loading of other antibiotic types within ME-AM scaffolds following the strategy presented in **Chapter 7**, which is based on the prior intercalation of antibiotics within lamellar inorganic fillers. Apart from the already tested CFX and GTM, MgAl and ZrP fillers can intercalate many other positively or negatively charged molecules, respectively, within their lamellar structure. This could enable the loading of a wide variety of antibiotics within ME-AM scaffolds, matching or widening the spectra of already existing local antibiotic delivery cements. Some of the antibiotics that could be considered are tobramycin, vancomycin, clindamycin, erythromycin or colistin. Although the first commercially available PMMA bone cements contained GTM, ¹²² subsequent generations have broaden the spectra and included the aforementioned antibiotics, alone or combined, to cover a wider variety of pathogens involved in orthopedic infections. ^{101, 109} Similarly, vancomycin or tobramycin have also been included in the formulation of commercial degradable cements based on calcium sulfates, due to their broad antimicrobial coverage. ^{101, 123-124} Rifampin administration, combined with levofloxacin or ciprofloxacin, is a common systemic antibiotic therapy against *Staphylococci*, which is the most common bacterial strain in implant associated infections. ⁹⁹ While this antibiotic has not been included so far within commercialized gold standard cements, due to its interference with the polymerization mechanism of PMMA, ¹²⁵ our polymer- intercalated filler approach could offer a suitable platform for its investigation within local delivery systems.

Despite the versatility of this system, the antibiotic size could be a limiting factor hampering its incorporation within the lamellar structure of the fillers. The access of a drug to the filler interlayer spaces can be restricted by its large molecular weight, or hydrodynamic radius, leading to adsorption to only the external filler surface. Among the aforementioned antibiotics, tobramycin, clindamycin, erythromycin and rifampin possess a molecular weight similar to that of CFX and GTM (300-800 Da), which have been already proven to fit within the interlayer spaces of lamellar fillers. ¹²⁶ On the other hand, vancomycin has a molecular weight of ~1,500 Da, and previous studies have already demonstrated their uptake only on the outer surface of such fillers. ¹²⁷ This is an aspect to take into account,

as it could influence the release kinetics, as well as impeded the molecule heat protection. For this reason, a thorough investigation of each antibiotic loading efficiency and mode, release kinetics, and antimicrobial activity maintenance after scaffold processing is encouraged.

Revisiting the polymer and cell choice

As previously mentioned, PEOT/PBT was initially chosen as the base material within the composite scaffolds, since previous studies had reported its potential for BTE applications, in terms of optimal mechanical, physicochemical (degradation time, melting temperature), and biological properties (biocompatibility, cell adhesion). While our results demonstrate that PEOT/PBT did not interfere in the functional properties of PEOT/PBT-rGO composite scaffolds (electrical conductivity, antimicrobial properties) and PEOT/PBT plasma functionalized scaffolds (cell adhesion), it proved to be one of the players diminishing the overall functionality of PEOT/PBT-nHA and PEOT/PBT-filler-antibiotic composite scaffolds. Here, the interaction of the fillers with the surrounding environment was necessary to exert their bioactivity, and the slow degrading PEOT/PBT acted as a barrier, leading to suboptimal CaP exchange dynamics and antibiotic release durations, respectively, and challenging its choice as scaffold material. In these cases, we argued that the use of a polymer with faster degradation times and increased water uptake capabilities would have alleviated these issues, yet compromising other equally important properties, such as mechanical stability. We certainly believe that further research on biocompatible and printable polymers with optimum physicochemical properties is required to widen the polymer library and be able to strike a balance in these situations.

Re-evaluating the polymer choice also guides us to revisit other initial selections, such as the cell type chosen to assess the functionality of the newly developed scaffolds. Among different cell sources for osteoblastic cells, hMSCs are preferred by the scientific community for their use in BTE strategies over primary cells, such as osteoblasts derived from bone or periosteum.¹²⁸ This is due to the possibility of isolating hMSCs from the bone marrow or adipose tissue in large numbers through minimal donor morbidity methods, in addition to their multi-lineage differentiation capabilities and participation in tissue regeneration processes. However, various reports have revealed that tissue source, isolation procedure, and donor characteristics (age, gender, and history) influence the phenotype, yield, growth rate, *in vitro* response to osteogenic signals, and *in vivo* bone formation capability of hMSCs.¹²⁹⁻¹³⁴ Such efficacy challenges are mostly overlooked in

BTE research. In the present study, although ensuring hMSCs osteogenic potential in 2D conditions parallel to each experimental procedure in 3D, such a variability was also not taken into consideration, and the extent of its effect on the reliability of the results and hypothesis presented within this thesis is not known. For this reason, the replication of the experiments with different hMSCs donors would be recommended to further proof all the hypothesis made.

Overall, we have proposed within this thesis significant scientific and technological advances applicable to the fabrication of more optimum ME-AM scaffolds for BTE. We have highlighted the advantages of each of them, as well as pointed out their limitations, while suggesting paths that future researchers could potentially take to further optimize our proposed systems.

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Chapter 9

Impact

Scientific impact

Although bone has been studied since the birth of the tissue engineering (TE) field, scientists are still searching for the optimal strategy for its regeneration (**Figure 1A**). Bone is a complex tissue, in terms of mechanical properties, structure, and composition, and many molecular factors and cell types are involved in its formation and regeneration. For this reason, clinical problems, such as non-union fractures or craniofacial reconstructions, are also complex to approach, and not easy to resolve. Among the three components of the classical tissue engineering paradigm (cells, scaffolds and growth factors), scaffolds have been the main focus of study in the bone tissue engineering (BTE) field. Besides providing a bioactive environment for new tissue formation, scaffolds in bone regeneration have the additional significant role of providing mechanical support, mimicking the high strength of the native extracellular matrix. Aiming towards size and shape personalized structures with high mechanical properties, but at the same time porous for cell ingrowth, additive manufacturing (AM) has stood out within the last 15 years as one of the preferred biofabrication technologies for the fabrication of scaffolds for BTE applications. Over the years, large technological advancements have been made around AM to facilitate an efficient scaffold design, i.e. multiprinthead melt extrusion AM (ME-AM) platforms have been developed for the fabrication of multimaterial scaffolds, sophisticated light-based or bioprinting AM platforms have emerged, new printable polymer materials or bioinks have been designed. Yet, an AM scaffold with the optimal set of properties for efficient bone regeneration has not been fully attained, suggesting that further investigations and advancements are still required.

Within this thesis, courageous choices were made to generate large scientific and technological strides providing solutions to existing issues within the field. We believe that the diverse strategies presented, consisting on optimized or new composite materials, new surface modification techniques, and novel cell seeding methodologies, contribute to the better understanding of AM scaffolds and their use for bone regeneration applications and provides a stepping stone for future studies on the application of functional composites in BTE. We also consider that this research has a scientific impact beyond the BTE field and can inspire future research, in the biomedical, biomaterial and overall tissue engineering scientific community.

Figure 1. Scientific publications on new bone scaffolds since 1970. Pubmed search, April 2021.

Within the first chapters, we found extremely relevant to revisit the recurrent cell seeding efficiency problem on macroporous AM scaffolds. On one hand, in **Chapter 3** we proposed a simple, reproducible and universal cell seeding approach that could be easily implemented by other research groups working with 3D AM for different applications. We are certain of the high scientific impact of this method, since it helps minimizing the required cell stock for *in vitro* experimentation and maximize cell seeding efficiency on scaffolds fabricated with any existing or newly developed synthetic material. It encourages researchers to redirect their efforts towards investigating more relevant issues, once the basic cell seeding problem is tackled. In **Chapter 3**, we went one step further and, while addressing the cell seeding issue, we explored a portion of the possibilities that a hybrid platform consisting of a melt extrusion printhead and plasma jet could bring to the design of surface-functionalized scaffolds. Our results open the door to future research on the effect of different plasma treatments on cell behavior on 3D ME-AM scaffolds. Furthermore, since this technique allows for a single-step workflow of scaffold production and surface homogeneous treatment, we envision its implementation within the scaffold fabrication pipeline of many laboratories in the future, as well as its impact in a smoother translation of ME-AM scaffolds into the clinic.

In subsequent chapters, several optimized or novel composite materials were investigated and complex methodologies from several scientific fields, transiting from material engineering, chemistry, and biology were used to analyze their suitability for BTE applications and their impact in the field. In particular, **Chapter 4** showed that blending high concentrations of nanohydroxyapatite within a polymer, as a common focus of the community trying to mimic the bone composition, does not necessarily result in a highly osteoinductive material. Thus, here we raised important questions to the scientific community related to the appropriate calcium phosphate material and their incorporation

method in ME-AM composite scaffolds for the stimulation of bone regeneration. On the other hand, the incorporation of graphene based materials into ME-AM scaffolds is not yet a well-established line of research in the field. For this reason, we believe that our study in **Chapter 5** about the effect of rGO density and concentration on composite scaffolds production and cell-material interaction constitutes a stepping stone for scientists entering this field in the future. Finally, we believe that the approach presented in **Chapter 6** to fabricate antibiotic-releasing AM scaffolds could open the door to future research on the use of other relevant inorganic nanofillers, antibiotics and biodegradable polymers to further optimize the system towards clinical applications.

Social impact

Bone is one of the most commonly injured organs in our body. Bone fractures can result from trauma, such as car accidents, or sport injuries, as well as from osteoporosis, a disease that weakens the bones and increases the risk of fragility fractures. Inappropriate initial treatments and co-morbidities, such as diabetes, genetic factors and poor life style, can increase the risk of delayed bone healing or non-unions. Resection of bone tumors, infections or prosthetic revisions in long bones can also result in non-unions. Around 4 million fractures are estimated to occur in Europe annually, from which approximately half a million require hospitalization and bone repair intervention.¹ Non-unions, which occur in 5-10% of the cases, are the most difficult to treat and have a high financial impact, with a total cost of € 8,000 to € 91,000 per case.²⁻³ In European healthcare systems, around 82.8–93.3% of this cost has been estimated to derive from indirect costs, such as productivity losses, due to long healing times leading to patients being unable to resume daily life activities and work within a short time.⁴ The remaining expenditures derive from care and intervention costs.

Current available clinical solutions for bone repair are based on natural and synthetic bone grafts. The natural grafts are divided into autografts and allografts, the latter ones including demineralized bone matrices (DBMs). The synthetic materials are based on ceramic, collagen-ceramic composites, polymers and bone morphogenetic proteins. In 2020, the global bone graft market size, based on allografts and synthetic substitutes, was estimated at € 2.36 billion (Grand View Research, market analysis report 2021, ID: GVR-1-68038-154-2). Due to the aging population and the increased incidence of osteoporosis and bone fractures, this market is expected to grow at an annual rate of 5.8% to reach € 3.7 billion

by 2028. Although the market is currently dominated by allograft materials (~ 60% share in 2020, mainly DBMs), the synthetic segment is anticipated to show significant growth over the years, due to the investment in product development, arise of new technologies and acceptance growth among patients. Bone grafts and substitutes are mostly used for spinal fusion, dental, and joint cemented prosthesis applications, while only 5% of the products are dedicated to long bone repair. This is partly because current marketed solutions, such as DBMs, or resorbable calcium phosphate cements have been shown to be inefficient in managing non-union scenarios in long bones, mainly due to their poor mechanical properties, unable to withstand the load required in these locations. Similar limitations are offered by collagen sponges infused with bone morphogenetic proteins (BMPs), which additionally have shown to induce undesired uncontrolled ectopic bone formation in clinical trials.

Here, AM scaffolds could bring many benefits for non-union defects. Such scaffolds can be fabricated to fit the specific patient complex defect using CT scan images of the patient's anatomy. This avoids the need of manipulation or modification of the scaffold intraoperatively, reducing the surgery time and ensuring good implant fixation within the defect. Moreover, AM scaffolds can be fabricated with specific structural parameters and mechanical properties from polymeric biodegradable materials or composites, to allow new bone formation and avoid a revision surgery. Despite their advantages, the implementation of biodegradable AM scaffolds in the clinics is still slow, as it is a rather new field and new medical devices require long regulatory paths, which subsequently leads to the current absence of companies marketing such products. Accordingly, many key players in the business of bone grafts, such as *DePuy Synthes*, *Medtronic*, *Wright Medical*, *AlloSource*, *Stryker*, *Johnson and Johnson* and *Zimmer Biomet*, only offer products based on DBM, ceramics or BMPs, and so far do not possess 3D AM scaffolds within their product portfolio. On the other hand, emerging small companies, such as *Mimetis* and *Cerhum*, have entered the sector by jumping in the 3D printing wave, yet developing AM ceramic scaffolds based on tricalcium phosphate (TCP) and hydroxyapatite (HA), whose performance in load bearing scenarios is arguable and still to be proven. Up to now, only one company (*Osteopore International*) has been able to commercialize 3D ME-AM polymeric scaffolds, for craniomaxillofacial defects, and currently is performing first-in-human trials for their future application in long bone non-unions. *Osteopore's* poly(caprolactone) (PCL) scaffold was recently implanted in a patient with a non-union tibial defect at Maastricht University Hospital. The success of this clinical case, and many

others to follow, will certainly open in the future the clinics' door to other novel ME-AM scaffolds. We believe that the research presented within this thesis will have a direct impact in shaping the future of such novel products, as we propose several strategies to improving their functionality, which could further accelerate bone healing times and reduce the socioeconomic burden associated with non-unions. Joining the success of the aforementioned basic polymeric ME-AM scaffolds with the extensive road of ceramics in the clinics, the clinical success of the PEOT/PBT-nHA scaffolds presented in **Chapter 5** can be seen as a natural next step to come. Such scaffolds would combine i) the bioactivity and osteoconductivity provided by the ceramic component, ii) the mechanical support brought by the polymer, the reinforcing ceramic, and the scaffold structure itself, and iii) the concept of personalized medicine. Overall, a blend of properties that current commercial products lack. Although not directly investigated within this thesis, the further addition to these scaffolds of the antibiotic loaded lamellar fillers presented in **Chapter 7**, would certainly further boost their functionality and transform them into a very unique product. As mentioned in previous chapters, further optimization would be required before obtaining a "ready to use" scaffold, but we believe that we are on a good track, which will surely guide us to implement a novel, more reliable, probably cheaper and, importantly, functional therapy for bone non-union regeneration, compared to current gold standards.

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Appendix

Summary

Current clinical strategies to treat critical bone size defects, mainly based on autografts, are inefficient and have limitations. For this reason, the orthopedic field is seeking innovative and effective solutions to regenerate bone in such scenarios, with the help of tissue engineering. Melt extrusion additive manufacturing (ME-AM) has stood out as an ideal biofabrication technique to generate patient personalized, highly porous, biodegradable and mechanically robust scaffolds that can provide support for new bone tissue formation in non-unions. However, these scaffolds are made out of synthetic polymers lacking bioactivity, which generally require further functionalization to improve their biological performance. In this regard, several strategies have been investigated over the last decades to augment the bioactivity of ME-AM scaffolds, which are carefully reviewed in **Chapter 2** of this thesis. Strategies such as architectural modifications, surface physical and chemical modifications, the incorporation of secondary fibrous or hydrogel networks within the scaffolds pores, and the use of filler-polymer composites are reviewed. As the search for optimal scaffolds that can boost bone regeneration continues, the aim of this thesis is to further contribute to the field by investigating different material-driven approaches to design and fabricate bioactive ME-AM scaffolds for bone tissue regeneration. To start with, **Chapter 3** and **Chapter 4** provide two elegant solutions to the recurrent cell seeding efficiency issue associated to macroporous 3D scaffolds. In **Chapter 3**, a simple and reliable static seeding method was developed to improve the cell seeding efficiency and distribution in ME-AM scaffolds. This method consisted of the supplementation of Dextran or Ficoll to the cell seeding media, which modified the medium viscosity and density, respectively, and reduced the settling velocity of cells, allowing for better attachment and distribution during the seeding process, compared to conventional static seeding. The macromolecules were completely removed after the seeding period and cell distribution along the scaffold cross section was dramatically enhanced. In addition, osteogenic differentiation of human mesenchymal stromal cells (hMSCs) was accelerated, in terms of enhanced mineralization and bone matrix maturation, compared to conventional seeding. In **Chapter 4**, the cell seeding issue was tackled from another perspective. In this case, ME-AM scaffolds were fabricated using a hybrid AM technique that combined ME-AM and an atmospheric pressure plasma jet, which was used to plasma-functionalize their surface. While argon activation resulted in an unspecific treatment, (3-Aminopropyl) trimethoxysilane (APTMS) and maleic anhydride-

Vinyltrimethoxysilane (MA-VTMOS) plasma polymerization deposited positively and negatively charged coatings, respectively. This enabled to investigate the cell attachment mechanism to the scaffolds as a function of surface charge in the presence or absence of serum. Furthermore, the coatings increased the surface wettability of the scaffolds, enhanced cell-adhesive proteins adsorption, and allowed for homogeneous cell attachment along the scaffold cross section. Cell proliferation was enhanced even on aged scaffolds, and hMSCs were able to express osteogenic genes in all scaffold types.

After ensuring an efficient cell attachment, composite scaffolds comprising a polymer matrix mixed with bioactive fillers possessing distinct properties were fabricated in **Chapters 5, 6 and 7**. In **Chapter 5**, highly loaded PEOT/PBT-nanohydroxyapatite (nHA) composite scaffolds with up to 45 wt% nHA were fabricated, mimicking the organic-inorganic bone composition. Here, the effect of nHA content on the osteogenic differentiation of hMSCs was evaluated in depth, and results suggested that in spite of no observed differences among scaffold types at the gene and protein level, increased matrix mineralization was observed on 45 wt% nHA scaffolds when cultured in mineralization and basic media. Moreover, a denser calcium phosphate layer reprecipitated on these scaffolds upon immersion in simulated body fluid. These results in combination with the enhanced compression modulus measured on 45 wt% nHA scaffolds, could be predicting their enhanced performance *in vivo*. In **Chapter 6**, ME-AM composite scaffolds containing reduced graphene oxide (rGO) were fabricated to provide electrical conductivity, protein adhesion and osteogenic properties. Initially, the low-density rGO was densified by solvent evaporation, in order to ease handling and ensure safety during melt compounding. It was observed that high density led to more compacted rGO which facilitated the printing of these composites up to 15 wt% rGO formulations. On the other hand, composites prepared with medium density rGO possessed a lower printability window, but presented higher electrical conductivity. In addition, increasing rGO content led to increasing material hydrophilicity, protein adsorption, and antibacterial properties, without affecting hMSCs viability. Furthermore, 3% rGO scaffolds were able to support hMSCs osteogenic differentiation. In **Chapter 7**, antibiotic-eluting scaffolds were fabricated via high temperature ME-AM. The antibiotics ciprofloxacin and gentamicin were intercalated within the lamellar inorganic fillers magnesium aluminium layer double hydroxide (MgAl) and zirconium phosphate (ZrP), respectively, prior to their dispersion within the polymeric matrix, in order to protect them from the fabrication process thermal stresses and obtain a sustained release. Compared to no-filler systems, a more sustained release was obtained

in the filler-based scaffolds, which was governed by diffusion through the polymer matrix and by ion exchange or pH changes. After printing, scaffolds demonstrated antimicrobial activity against Gram + and Gram - bacterial strains, at the same levels as unprocessed antibiotics. In addition, fillers and antibiotics did not affect hMSCs viability and osteogenic differentiation as shown by matrix mineralization and osteogenic proteins expression.

All the aforementioned results are discussed and placed into the state of the art context, in **Chapter 8**, while providing future perspectives. To finalize, a reflection on the scientific and societal impact of the research carried out in this thesis is presented in **Chapter 9**.

Samenvatting

De huidige klinische strategieën om kritieke botafwijkingen te behandelen, voornamelijk gebaseerd op autografts, zijn inefficiënt en hebben beperkingen. Om deze reden is de orthopedische sector op zoek naar innovatieve en effectieve oplossingen om bot te regenereren in dergelijke scenario's, met behulp van tissue engineering. Melt extrusion additive manufacturing (ME-AM) heeft zich onderscheiden als een ideale biofabricagetechniek om op de patiënt afgestemde, zeer poreuze, biologisch afbreekbare en mechanisch robuuste scaffolds te genereren die ondersteuning kunnen bieden voor de vorming van nieuw botweefsel in niet-unionen. Deze steigers zijn echter gemaakt van synthetische polymeren met een gebrek aan bioactiviteit, die over het algemeen verder gefunctionaliseerd moeten worden om hun biologische prestaties te verbeteren. In dit opzicht zijn er de laatste decennia verschillende strategieën onderzocht om de bioactiviteit van ME-AM steigers te vergroten, die in **hoofdstuk 2** van dit proefschrift zorgvuldig worden besproken. Strategieën zoals architectonische modificaties, oppervlakte fysische en chemische modificaties, de incorporatie van secundaire vezelachtige of hydrogel netwerken binnen de scaffolds poriën, en het gebruik van vulstof-polymeer composieten worden besproken. Aangezien de zoektocht naar optimale steigers die botregeneratie kunnen stimuleren doorgaat, is het doel van dit proefschrift om verder bij te dragen aan het veld door verschillende materiaal-gedreven benaderingen te onderzoeken om bioactieve ME-AM scaffolds te ontwerpen en te fabriceren voor botweefsel engineering. Om te beginnen, **hoofdstuk 3** en **hoofdstuk 4** bieden twee elegante oplossingen voor de terugkerende cel seeding efficiency probleem geassocieerd met macroporeuze 3D scaffolds. In **hoofdstuk 3**, werd een eenvoudige en betrouwbare statische zaaien

methode ontwikkeld om de cel zaaien efficiëntie en distributie in ME-AM scaffolds te verbeteren. Deze methode bestond uit de toevoeging van Dextran of Ficoll aan de cel zaaien media, die het medium viscositeit en dichtheid, respectievelijk gewijzigd, en verminderde de afwikkeling snelheid van de cellen, waardoor voor een betere hechting en distributie tijdens het zaaien proces, in vergelijking met conventionele statische zaaien. De macromoleculen werden volledig verwijderd na de zaaiperiode en de celdistributie langs de scaffoldsdoorsnede werd drastisch verbeterd. Bovendien werd de osteogene differentiatie van hMSCs versneld, in termen van verbeterde mineralisatie en botmatrix maturatie, vergeleken met conventionele seeding. In **hoofdstuk 4** werd het probleem van de celzaaiing vanuit een ander perspectief benaderd. In dit geval werden ME-AM scaffolds vervaardigd met behulp van een hybride AM techniek die ME-AM combineerde met een atmosferische druk plasmastraal, die werd gebruikt om het oppervlak te plasma-functionaliseren. Terwijl argon activering resulteerde in een niet-specifieke behandeling, (3-Aminopropyl) trimethoxysilaan (APTMS) en maleïnezuuranhydride-Vinyltrimethoxysilaan (MA-VTMOS) plasma polymerisatie afgezet positief en negatief geladen coatings, respectievelijk. Dit maakte het mogelijk het aanhechtingsmechanisme van de cellen aan de scaffolds te onderzoeken als functie van de oppervlaktelading in aanwezigheid of afwezigheid van serum. Bovendien verhoogden de coatings de bevochtigbaarheid van het oppervlak van de scaffolds, verbeterden ze de adsorptie van celadhesieve eiwitten en maakten ze een homogene celhechting langs de doorsnede van de scaffolds mogelijk. De celproliferatie was verbeterd, zelfs op verouderde scaffolds, en hMSCs waren in staat om osteogene genen tot expressie te brengen in alle scaffold types.

Na het verzekeren van een efficiënte celaanhechting, werden composiet scaffolds bestaande uit een polymeermatrix gemengd met bioactieve vulstoffen met verschillende eigenschappen gefabriceerd in de **hoofdstukken 5, 6 en 7**. In **hoofdstuk 5** werden hoogbelaste PEOT/PBT-nHA composiet scaffolds met tot 45 wt% nHA gemaakt, die de organisch-inorganische botsamenstelling nabootsten. Hier werd het effect van HA gehalte op de osteogene differentiatie van hMSCs diepgaand geëvalueerd, en de resultaten gaven aan dat ondanks geen waargenomen verschillen tussen scaffoldtypes op gen- en eiwitniveau, verhoogde matrixmineralisatie werd waargenomen op 45 wt% nHA scaffolds wanneer gekweekt in MM en BM. Bovendien reprecipiteerde een dichtere CaP-laag op deze steigers na onderdompeling in SBF. Deze resultaten in combinatie met de verhoogde compressie modulus gemeten op 45 wt% nHA scaffolds, zouden hun verbeterde prestaties in vivo kunnen voorspellen. In **hoofdstuk 6**, werden ME-AM composiet scaffolds met rGO

gemaakt om elektrische geleiding, eiwit adhesie en osteogene eigenschappen te bieden. In eerste instantie werd de lage dichtheid rGO verdicht door oplosmiddel verdamping, om het hanteren te vergemakkelijken en de veiligheid te garanderen tijdens het smelten. Er werd vastgesteld dat een hoge dichtheid leidde tot meer verdichte rGO, wat het drukken van deze composieten tot 15 wt% rGO formuleringen vergemakkelijkte. Anderzijds hadden composieten bereid met rGO met een gemiddelde dichtheid een lagere bedrukbaarheid, maar een hoger elektrisch geleidingsvermogen. Ook leidde het verhogen van het rGO-gehalte tot een verhoging van de materiaalhydrofiliëteit, eiwitadsorptie en antibacteriële eigenschappen, zonder de levensvatbaarheid van hMSC's aan te tasten. Bovendien waren 3% rGO scaffolds in staat om hMSCs osteogene differentiatie te ondersteunen. In **hoofdstuk 7**, werden antibiotica-eluting scaffolds gefabriceerd via hoge temperatuur ME-AM. De antibiotica CFX en GTM werden geïntercaleerd in de lamellaire anorganische vulstoffen MgAl en ZrP, respectievelijk, voordat ze werden gedispergeerd in de polymere matrix, om ze te beschermen tegen de thermische spanningen in het fabricageproces en om een duurzame afgifte te verkrijgen. Vergeleken met systemen zonder vulstof werd een duurzamere afgifte verkregen in de op vulstof gebaseerde scaffolds, die werd bepaald door diffusie door de polymere matrix en door ionenuitwisseling of pH-veranderingen. Na het printen toonden de steigers antimicrobiële activiteit tegen Gram + en Gram - bacteriestammen, op dezelfde niveaus als onbewerkte antibiotica. Bovendien hadden vulstoffen en antibiotica geen invloed op de levensvatbaarheid en osteogene differentiatie van hMSC's, zoals aangetoond door matrixmineralisatie en expressie van osteogene eiwitten.

Alle bovengenoemde resultaten worden in **hoofdstuk 8** besproken en in de context van de huidige stand van de wetenschap geplaatst, waarbij ook toekomstperspectieven worden gegeven. Tot slot wordt in **hoofdstuk 9** een beschouwing gegeven over de wetenschappelijke en maatschappelijke impact van het onderzoek dat in dit proefschrift is uitgevoerd.

List of Publications

Publications related to this thesis

M. Cámara-Torres, R. Sinha, C. Mota, L. Moroni, *Improving cell distribution on 3D additive manufactured scaffolds through engineered seeding media density and viscosity*, Acta Biomaterialia 2020, 101, 183-195, DOI: 10.1016/j.actbio.2019.11.020.

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M. Cámara-Torres, R. Sinha, A. Sanchez, P. Habibovic, A. Patelli, C. Mota, L. Moroni, *Effect of highly loaded nano hydroxyapatite composite scaffolds prepared via melt extrusion additive manufacturing on the osteogenic differentiation of human mesenchymal stromal cells*. (Submitted).

M. Cámara-Torres, R. Sinha, C. Mota, L. Moroni, *Boosting bone regeneration using augmented additive manufactured scaffolds*. (In preparation)

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R. Sinha, M. Cámara-Torres, P. Scopece, E. Verga Falzacappa, A. Patelli, L. Moroni, C. Mota, *A Hybrid Additive Manufacturing Platform to Create Bulk and Surface Composition Gradients on Scaffolds for Tissue Regeneration*. Nature Communications 2021, 12 (1), 500, DOI: 10.1038/s41467-020-20865-y.

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Scientific communications

Oral presentations:

3D additive manufactured composite scaffolds mimicking the bone composition and the bone-to-soft tissue interface: effect of high loadings of hydroxyapatite on the osteogenic differentiation of hMSCs, 11th World Biomaterials Congress, Online event, 2020.

High loading Reduced Graphene Oxide 3D Scaffolds for Bone Tissue Engineering, 11th World Biomaterials Congress, Online event, 2020.

Boosting bone regeneration: Additive manufacturing of continuous gradient scaffolds using functional composite materials. 3D Medical Printing Conference & Expo, Maastricht, The Netherlands, 2020.

3D additive manufactured scaffolds with antimicrobial activity for bone regeneration. eCM XIX: Orthopaedic Infection, Davos, Switzerland, 2019.

Continuous gradient hydroxyapatite-based 3D additive manufactured scaffolds for bone interface tissue engineering, TERMIS European Chapter Meeting, Rhodes, Greece, 2019.

3D additive manufactured scaffolds with antimicrobial activity for bone regeneration, 3rd MERLN PhD Symposium, Maastricht University, The Netherlands, 2018.

3D additive manufactured scaffolds with antimicrobial activity for bone regeneration, 29th Annual Conference of the European Society of Biomaterials, Maastricht, The Netherlands, 2018.

Poster presentations:

3D additive manufactured composite scaffolds for bone and bone-to-soft tissue interface: Effect of high loadings of hydroxyapatite on the osteogenic differentiation of hMSCs. 4th MERLN PhD Symposium, Maastricht, The Netherlands, 2019.

Atmospheric plasma assisted additive manufacturing of 3D scaffolds for tissue regeneration applications, 29th Annual Conference of the European Society of Biomaterials, Maastricht, The Netherlands, 2018.

Continuously graded reduced graphene oxide 3d scaffolds for bone regeneration, TERMIS World Congress, Kyoto, Japan, 2018.

Novel static cell seeding method to improve cell distribution in 3D additive manufactured scaffolds, TERMIS World Congress, Kyoto, Japan, 2018.

Novel Static Cell Seeding Method for Improving Cell Distribution on 3D Additive Manufactured Scaffolds, 2nd MERLN PhD Symposium, Maastricht, The Netherlands, 2017.

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Biography

María was born on the 16th August 1991 in Madrid, Spain. Pursuing her interest for engineering and biomedical sciences, she completed her BSc. in Biomedical Engineering at TECNUN School of Engineering- University of Navarra, Spain, in 2013. Aiming to gain international experience, she enrolled in a MSc. in Biomedical Engineering at FH-Aachen University of Applied Science, Germany.

She performed her master thesis at the DWI-Leibniz Institute for Interactive Materials in Aachen, Germany, where she developed injectable magnetic microgels and macroporous hydrogels to obtain tissue regeneration structures *in situ*. With this experience, her passion for biomaterials and tissue engineering grew, and in 2016, she joined MERLN institute for Technology-inspired Regenerative Medicine, Maastricht University, as a PhD candidate. There, she worked on a multinational and interdisciplinary European project entitled FAST, which aimed at developing novel 3D-additive manufactured composite scaffolds for bone regeneration in critical-size defects. In her research, she explored the effect of newly developed functional composites and surface modifications on the differentiation of stem cells towards bone cells. Keeping in mind the ultimate goal of tissue engineering and regenerative medicine, in the future she aims at bringing new biomedical products to patients.