Circulating monocytes in atherosclerosis : local or systemic actors?

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Circulating Monocytes in Atherosclerosis: Local or Systemic Actors?

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Aan Stefanie and Finn,
Voor al hun liefde en ondersteuning

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Chapter 01

General Introduction & Thesis Outline

Cardiovascular diseases (CVD) are one of the leading causes of mortality and morbidity in the Western world. According to figures from the Netherlands Heart Foundation (NHF), approximately 40,000 people died from cardiac or vascular diseases in The Netherlands in 2010 ¹. Although, the advances in research have already contributed to a decreased number of cardiovascular deaths in developed countries, still millions of people suffer from these diseases worldwide. Next to the successful intervention in developed countries, the incidence of CVD is believed to increase in developing countries over the next years, but also in the USA the prevalence of CVD is expected to increase ². Despite extensive research and the implementation of novel interventions not all patients at risk of a cardiovascular event can be detected timely or treated successfully. The knowledge on the consequences of risk factors, including obesity ³, hypertension ⁴, diabetes ⁵, smoking ⁶, and hyperlipidemia ⁷, but also on disease detection and prediction need to be expanded even further, to obtain be able to truly develop personalized medicine strategies and resolve cardiovascular diseases on a global scale.

Atherosclerosis

Atherosclerosis is the key underlying cause of CVD, responsible for a significant number of cardiac and vascular events, such as stroke, myocardial infarction, and peripheral artery disease. Atherosclerosis is considered to be a chronic inflammatory disease of the medium and large arteries ⁸. The disease is characterized by two main hallmarks: (I) lipid/lipoprotein metabolism and (II) inflammation. It is the interplay between these two processes that defines the principle pathogenesis and distinguishes atherosclerosis from other chronic inflammatory diseases ⁹. Lipid and cholesterol accumulation in the subendothelial layer occurs at sites with disturbed flow, such as curvatures and bifurcations of the arteries ^{10,11}. The endothelial layer will hence become activated which is even further stimulated by accumulation of low density lipoprotein (LDL) cholesterol particles and the subsequent oxidation of these LDL particles ¹². This activated endothelium will produce inflammatory cytokines, but also increase the expression of adhesion receptors such as vascular cell adhesion molecule (VCAM) 1 and intercellular cell adhesion molecule (ICAM) 1 on the luminal membranes ¹³⁻¹⁵. The upregulation of adhesion molecules will result in an increased attraction and adhesion of leukocytes to this activated endothelium ¹⁶. Interestingly, the locations of the

vasculature which have higher incidence of plaque development, already have increased leukocyte counts and signs of low-grade inflammation before development of atherosclerosis¹⁷ (figure 1).

Initially, the main leukocyte population attracted to the early atherosclerotic lesion will be monocytes. Monocytes will roll along the vessel wall and adhere at sites with increased adhesion receptors (or integrins). Eventually, the monocytes will attach to the endothelial layer of the vessel and can transmigrate into the subendothelial layer^{16,18}. The inflammatory milieu of the intima will trigger the monocytes to differentiate into macrophages. The latter cell type is considered to be a tissue-resident, mature form of monocytes. Oxidized cholesterol present in the early lesion will be scavenged by these macrophages via class B scavenger receptors and CD36. However, as these cells are unable to metabolize these oxLDL molecules efficiently and cholesterol efflux is also inhibited, intra-cellular cholesterol will accumulate, giving rise to the formation of macrophage foam cells. These cells have a foamy appearance due to the presence of large amounts of cholesterol filled endosomes and cytoplasmic lipids¹⁹⁻²². The macrophages and foam cells present in the initial atherosclerotic lesions, together with other attracted leukocytes, such as T-lymphocytes, will produce pro-inflammatory cytokines and thereby stimulate further expansion of the lesion by attraction of additional leukocytes to the progressing atherosclerotic plaque.

Over time, advanced atherosclerotic lesions can develop, most often asymptomatic, and thus without any clinical symptoms for many years^{23,24}. However, some lesions can progress into more unstable lesions by thinning of the fibrous cap covering the a-cellular and necrotic material present in the core of the lesion. Thinning of the cap is caused by e.g. matrix metallo-proteases (MMP), secreted by macrophages/foam cells in the lesion. Matrix metallo-proteases have also been shown to be involved in proteolytic cleavage of some cytokines thereby stimulating the inflammatory response⁹. Eventual rupture of the cap will present the debris from the core of the lesion to the blood stream and trigger coagulation, giving rise to the formation of thrombi. These thrombi will in the end cause local or peripheral clinical symptoms and even death due to the occlusion of arteries and subsequent inhibition of blood flow

²⁴.

Lipid/lipoprotein Metabolism in Atherosclerosis

Cholesterol synthesis and transportation

Lipids and cholesterol are essential to most organisms as many processes depend on these compounds for normal function. Cholesterol is one of the main components of cell membranes and functions as precursor for steroid hormones²⁵ and vitamin D²⁶, while lipids in general are an important energy source for the body. As lipids and cholesterol are poorly soluble in aqueous solutions, they need to be encapsulated in order to be transported in the blood. Several different subsets of lipoproteins can be distinguished involved in cholesterol transport. The most important lipoprotein fractions are very low density lipoprotein (VLDL), low density lipoprotein (LDL), and high density lipoprotein (HDL). Generally, LDL is thought to be the 'bad' cholesterol^{27,28} and HDL the 'good' cholesterol^{29,30}.

The liver plays a central role in the turnover of lipids and is responsible for the conversion of free fatty acids and chylomicrons into ApoB100-bound VLDL particles which are secreted into the circulation. Upon lipolysis, governed by lipoprotein lipases, VLDL is converted into LDL which is subsequently sequestered by the liver, but can also be used as metabolites in other pathways throughout the body. HDL particles are bound to ApoA-I which is important in the sequestration of cholesterol from cells via dedicated receptors. ATP-binding cassette transporter (ABCA) 1 is one of these receptors and is present on macrophages, supporting (limited) efflux of cholesterol to HDL particles^{31,32}.

Lipoproteins in Atherosclerosis

Hypercholesterolemia is, already for many years now, considered one of the most important risk factors for the development of atherosclerosis³³. Disturbances in the lipid/lipoprotein metabolism can affect the levels of various lipoprotein subsets, as these processes are responsible for the transport of lipoproteins throughout the circulation³⁴. At sites with turbulent flow and thus decreased shear stress, LDL cholesterol can accumulate in the intima, where it is prone to oxidation by local radicals, resulting in the development of oxLDL. As mentioned, this will initiate atherogenesis by endothelial cell activation due to disturbed blood flow^{35,36}. The humoral response to the oxLDL accumulation in the intima will further aggravate this

immunological initiation of atherosclerosis³⁷. The presence of lipoprotein, and more in particular oxLDL, is one of the central hallmarks of atherosclerosis. Exposure of (free) lipids and cholesterol, as well as necrotic material entrapped in the lesion to the blood flow upon plaque rupture may give rise to the clinical manifestations of cardiovascular disease²⁴.

Next to free cholesterol, also cellular cholesterol contributes to atherogenesis. Oxidized LDL phagocytized by macrophages, stimulates the inflammatory response in the vessel wall by the release of pro-inflammatory cytokines from macrophage foam cells and the induction of necrosis of these cells. Intracellular cholesterol can be present either cytoplasmic or lysosomal. Further, cholesterol crystals can be observed in cells which have accumulated excessive amounts of lipids and cholesterol.

Figure 1 - Schematic overview of early atherosclerosis development

Atherosclerotic plaque development is initiated by low density lipoprotein (LDL) cholesterol accumulation in the subendothelial layer of the vessel wall. Free radicals present in the tissue will lead to the formation of oxidized LDL (oxLDL), resulting in activation of endothelial cells. Upon activation endothelial cells will release chemo-attractant cytokines (chemokines) and upregulate the expression of integrins, facilitating the adhesion, rolling, and transmigration of circulating monocytes. In the subendothelial tissue monocytes will differentiate into macrophages which will subsequently scavenge the cholesterol and stimulate the pro-inflammatory immune response.

Inflammation in Atherosclerosis

Inflammation is, next to the lipid metabolism, a second essential process in the pathogenesis of atherosclerosis⁹. The inflammatory response consists of a cellular and humoral component and includes many immune cells, but also interactions with local endothelial cells and smooth muscle cells. As described earlier, circulating monocytes and plaque macrophages are critical actors in initial lesion development, but also in lesion progression, and eventual destabilization of the atherosclerotic plaque¹⁹.

Monocyte subsets in human

Human monocytes can be divided in two subsets based on expression of the membrane markers CD14 and CD16³⁸. In humans, CD14 is a co-receptor for binding of lipopolysaccharide (LPS) in combination with toll-like receptor (TLR) 4 and LPS-binding protein (LBP)^{39,40}, while CD16 is a Fc receptor (FcγRIII) which bind the constant Fc region of antibodies⁴¹. The two monocyte subsets that are currently being described in literature are CD14^{high}/CD16⁻ and CD14^{low}/CD16⁺⁺. The most abundant monocyte population in the circulation is the 'classical' CD14^{high}/CD16⁻ monocytes, comprising over 90% of circulating monocytes⁴². The classical monocytes are also often defined as inflammatory monocytes⁴³. The second population comprises the 'non-classical' CD14^{low}/CD16⁺⁺ monocytes, which are also referred to as patrolling monocytes. There are however also indications of a third monocyte population which has intermediate expression for both CD14 and CD16 (CD14⁺/CD16⁺)⁴⁴.

The classical and non-classical monocyte populations show differential expression of chemokine receptors involved in chemotaxis towards inflammatory stimuli⁴⁵. In relation to atherosclerosis and other inflammatory diseases such as rheumatoid arthritis, both subsets of circulating monocytes are thought to be important, however recruitment of these cells towards the site of inflammation is regulated by different chemokine receptors^{42,46}. The CC-motif chemokine receptor 2 (CCR2) is expressed in high levels by 'classical' CD14^{high}/CD16⁻ monocytes, and the responsiveness towards monocyte chemo-attractant protein (MCP) 1 is considered to be a characteristic of this subset⁴⁷. The CX₃C-motif chemokine receptor 1 (CX₃CR1), or fractalkine receptor, is expressed by both monocyte subsets⁴⁸, however the expression is approximately two-fold higher in 'non-classical' monocytes⁴⁹. This receptor is involved not only in chemotactic responses, but also has a role in cell survival

^{50,51}. Other important chemokine receptors expressed by monocytes are CCR1 and CCR5 and to a lesser extent CXCR2, which also are important for the patrolling properties of monocytes together with CCR2 and CX₃CR1 ⁴⁷⁻⁴⁹. As monocyte migration into the atherosclerotic lesion is mainly studied in mice there is only limited information on the role of these human subsets and their migratory behavior in atherosclerosis.

Table 1 - Monocyte subsets in human and mouse (Adapted from: Hristov *et al* ⁵²)

Monocyte subset	Alternative name	Major markers
Human		
CD14 ^{high} CD16 ⁻	Classical/Inflammatory	CD14, CCR2, CD16, CD62L, SR-A, VEGFR1
CD14 ^{low} CD16 ⁺⁺	Non-classical/patrolling	CD16, HLA-DR, CD11c, CX ₃ CR1, CCR5
CD14 ⁺ CD16 ⁺	Intermediate	CD14, CD16, CCR2, Tie2, CD105, MHCII, HLA-DR
Mouse		
Ly6C ^{high} (Gr1 ⁺)	Classical	Ly6C/Gr1, CCR2, CD62L
Ly6C ^{-/low} (Gr1 ⁻)	Patrolling	CX ₃ CR1, CCR5, LFA1, CD11c

Monocyte subsets in mice

In mice different circulating monocyte populations can be distinguished within the (CD11b⁺ / Ly6G⁻) monocytes population. In contrast to the human situation where monocyte subsets are defined based on CD14 and CD16 expression, in mice the subsets are identified based on Ly6C (or Gr1; binds both Ly6C and Ly6G) expression in combination with the general marker for myeloid cells (CD11b) and exclusion of granulocyte (Ly6G). Classical or inflammatory monocytes express high levels of Ly6C (or Gr1; Ly6C^{high} monocytes), while patrolling or 'non-classical' monocytes express low levels of Ly6C (Ly6C^{low} monocytes) ^{48,53,54}. As in human monocytes, the classical (Ly6C^{high}) murine monocytes express high levels of CCR2, while patrolling (Ly6C^{low}) monocytes do not express CCR2, but do express CX₃CR1 and CCR5 ^{42,55}.

Studies in mice have shown that the different monocyte populations reflect different stages of development ⁵⁶⁻⁵⁸. The bone marrow produces Ly6C^{high} monocytes, which can differentiate into Ly6C^{low} monocytes in the periphery in the absence of inflammatory stimuli ^{55,59}. However, this progressive differentiation from Ly6C^{high} in the bone marrow to Ly6C^{low} in the periphery is still debated in literature ^{60,61}. Despite the similarities in chemokine receptor patterns for CCR2 and CX₃CR1 (table 1), it is still questionable to what extent monocyte subsets in mouse and man can be compared directly, as in

C57Bl/6 mice both subpopulations are roughly present in equal numbers⁶². Nevertheless, it has been shown that circulating monocytes are essential for atherogenesis, as inhibition or depletion of (subsets of) monocytes results in reduced plaque development^{57,58} (figure 2).

Monocyte/Macrophage Axis in Atherosclerosis

Monocytes are the first leukocytes to enter the sub-endothelial space in response to lipoprotein accumulation in the early stages of atherogenesis. However, also during plaque progression and eventual plaque destabilization resulting in clinical manifestation of the disease, monocytes (and macrophages) are pivotal^{63,64}. It has been shown that monocytes respond to tissue-derived signals by migrating into lymphoid and non-lymphoid tissues^{65,66}. Triggers for these tissue-derived signals can be tissue damage or infection. In atherosclerosis, it is however more complex as not only inflammatory signals, caused by modified cholesterol in the intima, but also disturbed laminar flow were seen to contribute to the recruitment of monocytes^{22,42}. This also explains why atherosclerotic lesions are almost exclusively detected in curvatures and bifurcations of arteries¹¹. Adhesion molecules on endothelial cells facilitate the adhesion of circulating monocytes to the arterial wall, and subsequently aid in the transmigration into the intima¹³⁻¹⁵. This process is supported by chemokines secreted by activated endothelial cells and inflammatory cells in the lesion.

For human disease the chronology of leukocyte plaque invasion is not completely understood, as tracking and fate-mapping studies are almost impossible in the human situation. In mice however, many studies have been performed in which monocytes have been tracked or depleted to get more insight in the processes involved in atherosclerosis development and the role of monocytes herein. Ly6C^{high} monocytes have been shown to expand under hypercholesterolemic conditions and promote the inflammatory processes during atherogenesis⁶⁷. The role of Ly6C^{low} monocytes is more debatable in the development of atherosclerosis, as these monocytes are often considered to be anti-inflammatory and are traditionally thought to give rise to anti-inflammatory M2 macrophages and dendritic cells. This dogma is being challenged nowadays with our growing understanding of monocyte development.

Figure 2 - Monocyte subsets in atherogenesis

Two important monocyte populations are described in mouse and humans related to atherosclerosis. On the one hand, the inflammatory $\text{Ly6C}^{\text{high}}$ (mouse) or $\text{CD14}^{\text{high}}/\text{CD16}^-$ (human) monocytes which mainly migrate towards inflammatory sites such as the atherosclerotic lesion via CCR2^+ and predominantly give rise to (tissue-resident) macrophages. On the other hand, the patrolling Ly6C^{low} (mouse) or $\text{CD14}^+/\text{CD16}^+$ (human) monocytes which mainly migrate via $\text{CX}_3\text{CR1}$ and give rise to both myeloid dendritic cells and macrophages.

Macrophages in Atherosclerosis

Macrophages are more comparable between mouse and man, as the subsets present in man are also observed in mice. Monocytes that infiltrate the tissue or inflammatory site (e.g. atherosclerotic lesion) differentiate into M0 macrophages. These macrophages are considered to be non-polarized and do not have specific pro- or anti-inflammatory properties. Polarization of these macrophages towards a pro- or anti-inflammatory phenotype is dependent on the local milieu⁶⁸⁻⁷⁰.

Macrophages, as part of the innate immune system, are responsible for the scavenging of foreign materials. In the atherosclerotic lesion these cells will ingest the oxidized LDL from the intima. However, this process results in cell activation and thereby propagating the already ongoing immune response. Cytokines and proteases secreted by macrophages have an important role in plaque development and destabilization of the fibrous cap covering the atheroma^{9,71,72}.

Local stimuli such as cytokines and cell-cell interactions can stimulate these tissue-resident macrophages to polarize into either classically activated, pro-inflammatory (M1) macrophages or alternatively activated, anti-inflammatory (M2) macrophages^{68,69}. Classically activated macrophages are considered to be the immune effector cells which are involved in the engulfment and digestion of microbes and other foreign particles. These cells are also important mediators of the inflammatory response and next to this have high proteolytic capacity. M1 macrophages are activated by lipopolysaccharide (LPS) and interferon (IFN) γ and produce high levels of interleukin (IL) 12, while no or low levels of IL10 are produced. Other cytokines produced by M1 macrophages are IL1 β , IL6, and tumor necrosis factor (TNF) α ⁷⁰. Alternatively activated macrophages (M2) macrophages can be divided in M2a and M2c macrophages based on their membrane marker expression and cytokine production. In contrast to 'inflammatory' M1 macrophages, M2 macrophages are considered to be anti-inflammatory or wound-healing cells^{70,73}. Polarization towards M2a or M2c phenotype can be induced by stimulation with IL4, or with IL10 and transforming growth factor (TGF) β , respectively⁷³. These M2 macrophages mainly produce IL10, stimulating anti-inflammatory pathways in e.g. T-lymphocytes^{9,70,73}.

In atherosclerosis the role of the different macrophage subsets has not yet been elucidated completely. Nevertheless, studies indicate that a heterogeneous set of macrophage subsets is present in the lesion^{68,69}. Besides the well-known pro-inflammatory M1, and anti-inflammatory M2a and M2c macrophages, also oxLDL polarized macrophages (Mox)⁷⁴ and CXCL4 polarized macrophages (M4)⁷⁵ can be found in atherosclerotic plaques^{70,76}. Due to the complexity of determining distinct macrophage subsets in the lesion in combination with potential phenotype switching of lesional macrophages^{77,78} the exact role of the different subsets in atherosclerosis development remains uncertain.

Hematopoietic Compartments

The development of atherosclerotic lesions is considered to be a local pathology, however plaque rupture can result in systemic effects and even death due to e.g. myocardial infarction or stroke. As the cells present in the lesion are attracted from the periphery via the circulation and also cytokines and chemokines are released into the circulation, there is also an important systemic contribution. Recently, several studies have shown that there is indeed an important role for the bone marrow and the spleen, which are both involved in myelopoiesis, and thus in the production of monocytes^{59,79-81}.

In mice it has been shown that atherosclerosis in combination with cardiovascular risk factors such as hyperlipidemia, result in systemic changes and adaptations of the hematopoietic process. Mainly, myeloid hematopoiesis (myelopoiesis) is affected, and leads to dynamic changes in bone marrow and spleen^{59,80}. Increased numbers of monocytes (monocytosis) and also granulocytes can be detected upon cholesterol rich diet feeding in both mouse and man^{82,83}. However, it is still disputable to what extent these changes are caused by hyperlipidemia, inflammation, or the combination of both factors. Despite the fact that still many questions need to be investigated, it is clear that hematopoiesis plays an important role in not only replenishing the pool of peripheral leukocytes and tissue-resident inflammatory cells, but also in boosting and regulating various inflammatory diseases.

Figure 3 - Myelopoiesis in pro-atherogenic mouse models

Inflammatory stimuli in combination with hypercholesterolemia trigger the release of myeloid progenitor cells as well as mature monocytes from the bone marrow into the circulation. Myeloid progenitor cells will preferentially migrate towards the spleen, giving rise to extramedullary hematopoiesis. In combination with accumulation of bone marrow-derived monocytes in the spleen, a splenic reservoir will be formed consisting of (mainly Ly6C^{high}) monocytes. Upon additional stimulus, these cells are quickly mobilized from the spleen and migrate towards e.g. atherosclerotic lesions to form plaque macrophages and dendritic cells.

In normal myelopoiesis we can distinguish several intermediate myeloid precursor subsets developing from multipotent hematopoietic stem cells in the bone marrow. Common myeloid progenitors (CMP) develop via granulocyte/macrophage progenitors (GMP) and eventually mature granulocytes and monocytes. This development is controlled by cytokines, such as granulocyte/macrophage colony stimulating factor (GM-CSF), granulocyte colony stimulating factor (G-CSF), and macrophage colony stimulating factor (M-CSF)⁸⁴⁻⁸⁷. During development of monocytes, and their differentiation into macrophages, different transcription factors are expressed. Initially, PU.1 and Kruppel-like factor (Klf) 4 are activated, while later in the differentiation process Early growth response protein (Egr) 1 and interferon

regulating factor (IRF) 8 are more important^{88,89}. The expression of these transcription factors is specific to cells of the myeloid lineage and more specifically monocytes and macrophages⁷⁹. IRF8 deficiency can also influence atherosclerosis development in mouse models indicating the importance of regulating and maintaining normal myelopoiesis⁹⁰.

Besides the bone marrow, the spleen is also an important compartment in atherosclerosis-associated myelopoiesis. According to studies performed in pro-atherogenic apolipoprotein E deficient (*ApoE*^{-/-}) mice, Ly6C^{high} monocytes migrate from the bone marrow into the circulation^{59,80,83}. Subsequently, these cells will migrate into the spleen and differentiate towards Ly6C^{low} monocytes⁹¹. The spleen thus serves as a reservoir for rapid mobilization of monocytes upon inflammatory stimuli. In a pro-inflammatory system, such as observed in atherosclerosis, not only mature cells are released from the bone marrow, but also progenitors will be mobilized. These hematopoietic progenitors will home to the spleen and locally start producing mature myeloid cells. This extra-medullar hematopoiesis is one of the mechanisms responsible for the monocytosis observed in hypercholesterolemic, but also hyperglycemic, pro-atherogenic mice^{92,93}. Overall, hematopoiesis contributes to atherosclerosis through enhanced production of myeloid cells and more specifically monocytes (figure 3).

From Cells to Biomarkers

Early detection and the selection of the optimal procedure for treatment are important aspects of cardiovascular biomarker research. Traditionally, different types of biomarkers can be distinguished and can be divided in diagnostic, prognostic, and predictive biomarkers. Diagnostic biomarkers are used to identify the presence of disease in a patient. These markers can be used in a clinical setting e.g. to distinguish between chest pain caused by myocardial infarction and chest pain due to non-cardiac related causes. Prognostic and predictive biomarkers give information on future risk of developing a specific event or disease. Prognostic biomarkers give information in untreated individuals, while predictive biomarkers can give more insight in the effectiveness of a treatment for individual patients. Both prognostic and predictive biomarkers can give indications for adapting the treatment.

Risk prediction for cardiovascular disease can be based on circulating, imaging, and genetic biomarkers⁹⁴. Traditionally, risk prediction is mainly

based on risk factors as hyperlipidemia, hypertension, diabetes mellitus, and smoking. These markers have been developed into predictive models such as the Framingham risk score⁹⁵. Research has shown that these models for assessing CVD risk at population level are limited in predicting the risk for individual patients, due to changes in biomarkers upon treatment⁹⁶ and lack of significant differences at the level of individual patients^{97,98}.

In recent years, many novel biomarkers have been presented which are developed in new (high-throughput) platforms⁹⁹. Based on the more traditional lipid and lipoprotein profiling additional biomarkers have been proposed. The ratio between apolipoprotein (Apo) B and ApoA-I, which are part of LDL and HDL, respectively, is a strong predictive marker of cardiovascular events^{100,101}. Another well-known biomarker for cardiovascular disease is C-reactive protein (CRP), which is a marker for systemic inflammation and is used in the clinic for many years¹⁰². The downside of CRP, and many more biomarkers in general, is the fact that different cohort studies result in different risk predictions¹⁰³⁻¹⁰⁵. Further, CRP is not specific for CVD as it is a more general marker for inflammation. Finally, biomarkers proposed so far have limited in their predictive value for an individual patient and often the biomarkers are inadequate with regard to specificity and sensitivity upon validation. Nevertheless, current and future biomarkers will contribute to personalized medicine extensively by improving diagnosis and prognosis for individual patients. By optimizing these strategies treatment and life style changes can be even more specified to the needs of a specific person, minimizing the risk for future cardiovascular morbidity and mortality.

Table 2 - Non-comprehensive list of CVD biomarkers

Biomarker		Pathology	Ref
CRP	<i>C reactive protein</i>	<i>Acute inflammation</i>	102,104
ApoB/ApoA-I	<i>Apolipoprotein B/A-I</i>	<i>Predictive marker CV events</i>	100,101
BNP	<i>B-type natriuretic peptide</i>	<i>Prognostic marker heart failure</i>	106,107
sCD40L	<i>Soluble CD40 ligand</i>	<i>Plaque instability</i>	108
CCL18	<i>CC-motif chemokine ligand 18</i>	<i>Unstable angina pectoris</i>	109
hsTNT	<i>High sensitive troponin T</i>	<i>Myocardial injury</i>	107,110,111
sPLA ₂	<i>Secreted phospholipases A2</i>	<i>Prognostic marker CV events</i>	112
MPO	<i>Myeloperoxidase</i>	<i>Risk for future CV events</i>	113,114
GDF15	<i>Growth differentiation factor 15</i>	<i>Heart failure</i>	107,115

Whole cells and also circulating monocytes are already being used as biomarker for disease. Both secreted and membrane-bound markers are related to different pathological conditions. Peripheral blood mononuclear cells (PBMCs) from patients with systemic sclerosis secrete more CD163, which might give information on prognosis ¹¹⁶. In Alzheimer's disease LPS induced IL-6 release by PBMCs functions as prognostic marker for disease severity and cognitive outcome ¹¹⁷. Besides these secreted proteins also membrane markers can be used as biomarker. Monocyte HLA-DR expression has a strong relationship with disease severity and prognosis in acute liver failure ¹¹⁸, while Sialic acid-binding Ig-like lectin 1 (Siglec-1) and Fcγreceptor I (CD64) expression function as biomarker in systemic lupus erythematosus ^{119,120}. For cardiovascular disease only one biomarker is described that is possibly derived from (plaque) monocytes, however the exact cell type responsible for the secretion of junction plakoglobin has not been determined yet ¹²¹. As monocytes play a critical role in the development of atherosclerosis, in depth research towards biomarker discovery based on these cells will be necessary, to be able to more efficiently diagnose and treat patients at risk for CVD.

Focus & Outline of the Thesis

Despite the fact that in some developed countries the morbidity and mortality of cardiovascular diseases is diminishing, there is still need for more research as the incidence of disease is increasing in developing countries, and current diagnostic tools as well as therapies appear to be insufficient. Furthermore, there is a need for earlier, predictive and less invasive methods to detect the risk for atherosclerosis associated disorders. It is believed that circulating cells are the new target to determine the state of disease and risk for future events in patients suspected to have progressed cardiovascular diseases. This thesis focuses on the role of circulating monocytes in relation to atherosclerotic plaque development on the one hand, and the possibility of using these cells as biomarker source for plaque phenotype, and possibly even the risk for a future cardiovascular event on the other hand.

Circulating monocytes in mouse models of atherosclerosis

In **Chapter 2**, the effect of depleting monocytes and macrophages from the circulation, but also the atherosclerotic lesion will be investigated, using a pro-atherogenic mouse model. As the monocyte/macrophage axis plays a central role in atherosclerosis, depleting these subsets should have profound impact on local lesion associated processes, but also shed new light on the systemic importance of circulating cells in atherosclerosis.

Subsequently, the role of monocyte chemotaxis towards the atherosclerotic lesion will be analyzed in **Chapter 3**. We aim to gain insight into the processes regulating mobilization of monocytes from the periphery upon chemokine release. This chapter addresses the role GRK2 plays in chemotaxis and the effects of hematopoietic and macrophage specific deficiency on atherosclerotic lesion development in LDL receptor knock-out mice.

As atherosclerosis development is also critically dependent on lipids, and more in particular cholesterol, **Chapter 4**, will address the role of hypercholesterolemia on circulating monocytes. The processes underlying foam cell formation in the lesion are already well described, however little is known on the peripheral processes and responses of leukocytes, and in particular monocytes, upon increased cholesterol levels. Through diet-induced hypercholesterolemia, we are able to unravel the systemic adaptations, as well as functional changes in circulating monocytes.

Circulating monocytes as biomarker in human atherosclerotic disease

The second part of this thesis focuses on the role of monocytes (and macrophages) in human cardiovascular disease.

First, in **Chapter 5**, the effects of cell activation, as observed in atherosclerosis, will be investigated on a protein level. Whole cell phage display is used to identify novel markers on human monocytes, which are specific for unstable angina pectoris and also advanced atherosclerotic lesions.

Subsequently, in **Chapter 6**, monocytes from stable and unstable angina pectoris patients will be studied on a mRNA level to determine whether there are baseline differences in monocyte activation or regulation due to the presence of unstable atherosclerotic plaques. Micro-array analysis, combined with computational analyses and flow cytometry are used to determine the profile of these monocyte pools, to verify whether monocytes from different patient groups do indeed have a different profile. We also include preliminary data on the differences in activation potential between monocytes from patients with an event during follow-up compared to controls.

Finally, a summary, general discussion of the results of this thesis, and future perspectives will be presented in **Chapter 7**.

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The background of the entire page is an abstract, artistic splatter of red paint or ink on a white surface. The splatters vary in size, from small dots to large, elongated streaks and blotches, creating a dynamic and textured visual field. The red color is a deep, slightly dark shade, reminiscent of blood or a specific pigment.

Chapter 02

Local but not systemic CD115⁺ myelocyte ablation aggravates atherogenesis

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Timo Rademakers, Erwin Wijnands, Leon Schurgers, Erik AL Biessen

In Preparation

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Abstract

Although macrophage apoptosis is deemed a critical process in plaque progression, macrophage ablation studies have so far yielded contradictory results on the actual impact of macrophage death on atherosclerosis. This may be partly due to ectopic effects of the ablation strategy on circulating monocytes as well as on other leukocyte subsets. To distinguish plaque macrophage cell death associated effects from that on peripheral monocytic cells, we here studied the impact of systemic versus plaque-targeted ablation of monocytic cells on plaque development. ApoE^{-/-} mice, bearing a Macrophage FAS Induced Apoptosis (MaFIA) suicide gene under control of the CD115 promotor were used to develop advanced atherosclerotic lesions by perivascular collar placement in response to Western type diet feeding. Rapalog dimerizer was administered either systemically via intravenous tail vein injections, or locally via catheterized osmotic minipumps connected to the peri-adventitial site of the carotid arteries. Systemic treatment did not result in any changes regarding plaque development, despite efficient depletion of intimal macrophages. However, while effecting a complete ablation of circulating monocytes and neutrophils at 1 week after administration, systemic ablation led to a profoundly dysbalanced myelopoiesis, with major expansion of mature and immature myeloid cells in the circulation after 2 weeks. In contrast, focal ablation of CD115+ macrophages led to increased macrophage apoptosis and necrotic core size and to plaque expansion without any systemic effects. Taken together, extramedullar myelopoiesis and increased mobilization of inflammatory spleen monocytes to the circulation, upon systemic induction of monocyte/macrophage apoptosis causes vulnerability of atherosclerotic lesions. Local induction of macrophage apoptosis induced lesion expansion but allowed macrophage turnover without lesion vulnerability.

Introduction

Macrophage accumulation in the atherosclerotic lesion is generally considered essential for plaque progression and destabilization ¹. Furthermore, macrophages exert a range of functions to control and clear infections ², regulate and modulate wound healing ³, and engulf (non-)foreign particles ⁴. In early atherosclerosis these myeloid cells are involved in the clearance of oxidized low density lipoproteins (oxLDL) from the vessel wall, whereas in later stages macrophages also mediate the clearance of apoptotic cell material from the lesion ^{1,4}. Hence, macrophage cell death will at later stages be associated with impaired efferocytosis, and deposition of macrophage contained lipid debris in the plaque, both processes deemed deleterious for plaque stability ⁵. Moreover, macrophage derived matrix metallo-proteases (MMP) contribute to the degradation of the plaque-stabilizing collagen filaments and thereby undermine the integrity of the atherosclerotic lesion ^{6,7}.

Although plaque macrophage build up may partly result from clonal expansion of resident vascular macrophages as well, the bulk was seen to originate from extravasated monocytes that had been attracted to the plaque ^{1,8}. The recruitment of monocytes is a continuous process throughout all stages of the atherosclerotic lesion development ⁹. Thus, by fostering macrophage accumulation, monocytes will contribute to the progression of lesions and more specifically to the perpetuation of the ongoing inflammatory state and the vulnerability of the advanced atherosclerotic lesion as well ^{10,11}. In support of this notion, human leukocytosis was demonstrated to be an independent risk factor and predictor of clinical cardiovascular events in atherosclerosis ¹²⁻¹⁵.

This recognition has inspired to a wealth of studies on the interaction between of the myeloid lineage and hyperlipidemia and atherosclerosis. Our insights into the role of monocytes and macrophages in atherosclerosis have been greatly aided by both pharmacological and genetic monocyte/macrophage specific ablation studies. Clodronate liposomes have been used widely to study the effects of macrophage ablation in atherosclerosis ¹⁶, but also other pathologies ^{17,18}. Calin *et al* reported that initially macrophage depletion in atherogenesis is beneficial, however increased lipid and collagen content ultimately results in plaque expansion. Furthermore, studies have applied diphtheria toxin dependent ablation models under control of CD11b ¹⁹. Induction of macrophage apoptosis via a CD11b-

diphtheria toxin model showed that depletion of monocytes/macrophages is influencing early lesions more pronouncedly than established lesion¹⁹. CD11b-hDTR and clodronate liposomes dependent myeloid cell depletion appear to indicate similar effects of monocyte/macrophage ablation on atherosclerosis development.

The key role of monocytes and macrophages in atherosclerosis development¹, but also other inflammatory diseases such as rheumatoid arthritis²⁰, has already been demonstrated. However, the aforementioned models for macrophage depletion are often lack specificity, and directly or indirectly impacted leukocyte subsets other than macrophages, thwarting data interpretation and resulting in unwanted side effects^{19,21-23}. CD11b is expressed by not only monocytes/macrophages, but also by neutrophils and dendritic cells, in contrast, CD115 is more constricted to the monocyte/macrophage axis, and thus CD115 will give a more specific insight in the role of this axis in atherosclerosis development.

In this study we employed a loss-of-function approach using a MaFIA transgenic mouse model that carries an inducible FAS construct under control of the CD115 (also known as macrophage-colony stimulating factor receptor [M-CSFr] or colony stimulating factor 1 receptor [CSF-1R]) promotor²⁴. In mouse and man, the monocytic lineage depends entirely on M-CSF for their ontogenesis²⁵⁻²⁷, and they produce and express high levels of this cytokine as well as of the cognate receptor CD115. Together with other cytokines such as granulocyte/macrophage colony stimulating factor (GM-CSF) and interferon gamma (IFN- γ)²⁸, M-CSF is essential for myeloid lineage commitment in the bone marrow²⁹. The key role of the M-CSF receptor in the myelopoiesis leads to the concept of ablating macrophages based on this receptor to study the role of macrophages in atherosclerosis and myelopoiesis³⁰. This model allows us to deplete CD115⁺ monocytes and macrophages in a time-controlled manner, by administering a dimerizer. Lymphocytes, and also granulocytes and dendritic cells, are considered to be M-CSF receptor low or negative and predominantly develop in an M-CSF independent manner³¹.

We provide evidence that depletion of M-CSF receptor (CD115) positive myeloid cells in a FAS-dependent manner, only contributes to atherosclerotic lesion formation in mice on a pro-atherogenic Apo lipoprotein E deficient (*ApoE*^{-/-}) background when depleted locally in the atherosclerotic lesion. Systemic ablation of M-CSF receptor positive monocytes and macrophages does not influence atherosclerosis development, however it does result in deregulation of erythropoiesis and myelopoiesis.

Materials & Methods

Animals

C57BL/6J-Tg(Csf1r-EGFP-NGFR/FKBP1A/TNFRSF6)2Bck/J MaFIA^{+/+} male mice obtained from the Jackson Laboratories³², were backcrossed to *ApoE*^{-/-} mice for more than 12 generations to obtain homozygous *ApoE*^{-/-} // *MaFIA*^{+/+} mice. The MaFIA transgene was inherited according to Mendelian laws as demonstrated by qPCR analysis of gDNA (PureLink Genomic DNA Purification Kit, Invitrogen). Primer sequences were: 5'-CCACATGAAGCAGCAGGACTT-3' for the forward primer and 5'-GGTGCCTCCTGGACGTA-3' for the reverse primer.

Male mice expressing the Macrophage EAS Induced Apoptosis transgene (*MaFIA*^{+/+}; coexpressing green fluorescent protein (GFP) and the dimerizer inducible suicide receptor under control of the endogenous CD115 promotor) on a pro-atherogenic Apo lipoprotein E deficient (*ApoE*^{-/-}) background were obtained from the animal breeding facility at the Maastricht University. Dimerizer treated *ApoE*^{-/-} mice or solvent-treated *ApoE*^{-/-} // *MaFIA*^{+/+} mice served as controls for the *in vivo* studies. Mice were fed regular chow diet (RM3; Special Diet Services [SDS], Essex, U.K.) or Western type diet (WTD; 0.25% cholesterol; SDS). Drinking water supplemented with antibiotics (60,000 Units/L Polymixin B Sulfate [Sigma-Aldrich, St. Louis, MO, USA] and 100 mg/L Neomycin [Sigma-Aldrich]) was introduced two weeks before the introduction of Western type diet. During experiments drinking water and food were provided *ad libitum*. During all experiments animals were housed in filter top cages. All *in vivo* studies were approved by the Maastricht University Ethics committee for animal experiments and performed according to the Dutch government guidelines for animal experiments.

Dimerizer

Lyophilized AP20187 was dissolved in ice-cold pure (100%) ethanol to obtain a 0.455 mM (6.25 mg/mL) stock solution, which was stored at -20° C until further use. For *in vitro* experiments the stock solution was diluted in medium to a final concentration of 1 – 5 μ M in the culture medium with an end concentration of ethanol below 0.5%. For intra-venous (iv) injections, the stock solution was diluted 25-fold in water containing 4% ethanol, 10% poly(ethylene) glycol (PEG) 400, and 1.7% Tween, resulting in an end concentration of 0.25 mg/mL AP20187. All intra-venous injections were administered within 30 minutes after preparation of the injection solution. The applied dose was 1 mg/kg body weight (150 μ L/mouse on average). *ApoE*^{-/-} control animals received solvent containing AP20187 at the same dose as the MaFIA mice, *ApoE*^{-/-} // *MaFIA*^{+/+} controls received solvent-only injections. For the *in vivo* experiments in which osmotic minipumps (Alzet, Cupertino, CA, USA) were used, stock solution was diluted as before, however the administered dose dimerizer was 100-fold lower (0.01 mg/kg).

In vivo experiments

IV injection atherosclerosis study. Ten week old *ApoE*^{-/-} // *MaFIA*^{+/+} animals (*n*=14) and *ApoE*^{-/-} animals (*n*=13) were placed on a Western type diet. Two weeks after introducing the WTD animals received flow restricting collars around the right and left carotid arteries to induce atherosclerotic plaque formation, as described before³³. Three weeks later, mice received an intravenous AP20187 tail vein injection every 72 hours for two weeks (final dose of 1 mg/kg body weight). Bodyweight was measured to adjust the dose according to the individual weight of each mouse and monitor the effect of the treatment. Mice were sacrificed 18 hours after the last injection with the dimerizer.

Osmotic minipump atherosclerosis study. Ten week old *ApoE*^{-/-} // *MaFIA*^{+/+} animals (*n* = 8) and *ApoE*^{-/-} animals (*n* = 8) were placed on a Western type diet. Two weeks after introducing WTD, semi-constrictive collars were placed bilaterally, at the common carotid arteries, to induce atherosclerotic plaque formation; simultaneously an osmotic minipump containing AP20187 with catheters attached to the perivascular site of the carotid artery was placed. Mice were sacrificed two weeks after implantation of the osmotic minipumps.

Upon sacrifice the integrity and localization of the catheters was verified and animals with (suspected) malfunctions in the catheters or minipumps were excluded from the study.

Dynamics study. Ten week old *ApoE*^{-/-} // *MaFIA*^{+/+} animals (*n* = 16) and *ApoE*^{-/-} animals (*n* = 8) were placed on a Western type diet for five weeks. Subsequently, *MaFIA* mice received an intravenous tail vein injection containing AP20187 (1 mg/kg body weight) every 72 hours for one or two weeks (*n* = 8 / *timepoint*). Control *ApoE*^{-/-} mice were processed similarly. Bodyweight was measured to adjust the dose according to the individual weight of each mouse. Mice were sacrificed 18 hours after the last injection with the dimerizer.

In vitro experiments

Macrophage culture. Bone marrow cells were isolated from *ApoE*^{-/-} // *MaFIA*^{+/+} and *ApoE*^{-/-} mice (*n* = 4 / *genotype*). Bone marrow cells were isolated from one femur and one tibia per mouse and pooled samples were cultured in RPMI-1640 (Invitrogen) supplemented with 10% fetal calf serum and L929 conditioned medium (LCM) to generate bone marrow-derived macrophages (BMDM) as previously described³⁴. Medium was replaced every three days and differentiated BMDM were used for *in vitro* assays after seven days.

Macrophage polarization. BMDM were incubated for 24 hours with LPS (100ng/mL), IFN- γ (100 U/mL), TNF- α (40 ng/mL), IL-10 (10 ng/mL), or IL-4 (20 ng/mL). Subsequently, mRNA was isolated using the Trizol (Invitrogen) method. 500 ng total RNA was reverse transcribed using the iScript cDNA Synthesis Kit (BioRad, Veenendaal, the Netherlands). Quantitative PCR was performed using 10 ng cDNA, 100 nM of each primer, and SYBR Green (Invitrogen) in a total volume of 15 μ L. Gene expression levels were corrected for glyceraldehyde 3-phosphate dehydrogenase (GAPDH) as housekeeping gene. Primer sequences are available upon request.

Apoptosis assay. Peritoneal macrophages isolated from *ApoE*^{-/-} // *MaFIA*^{+/+} and *ApoE*^{-/-} mice (*n* = 5 / *genotype*) were cultured for 24 hours in RPMI-1640 (Invitrogen, Paisley, UK) supplemented with 10% fetal calf serum (FCS). Subsequently, AP20187 was added to the medium to a final concentration of

4.55 μM and cells were incubated for 18 hours. Apoptosis was measured by immunohistochemically staining for cleaved caspase-3 and counterstaining with DAPI. Pictures were made (Leica DMIL fluorescence microscope) and overlays were analyzed with ImageJ Software.

Histology

Carotid arteries from *ApoE*^{-/-} // *MaFIA*^{+/+} and *ApoE*^{-/-} mice were collected, formalin-fixed and paraffin embedded. Subsequently, 4 μm thick sections were stained with haematoxylin and eosin (H&E) and lesion size and necrotic core area were determined in five sections per animal. Representative sections on separate slides were stained immunohistochemically with a macrophage-specific antibody (MAC-3; Becton & Dickinson, NJ, USA) or with a smooth muscle cell-specific antibody (α -smooth muscle cell actin antibody; Sigma-Aldrich; St. Louis, MO, USA). Apoptosis was visualized using a terminal deoxynucleotidyl transferase dUTP nick-end labeling (TUNEL) kit (Roche; Woerden, Netherlands). Apoptotic cell content was determined by assessment of the TUNEL positive area per section. Collagen was visualized by Sirius-Red staining of representative slides.

Spleen and tibias were analyzed for the effects of the treatment on the hematopoiesis. After fixation for 24 hours in 1% paraformaldehyde spleen tissue was paraffin embedded and 4 μm thick sections were cut for histological analysis. Tibias were fixated for 24 hours and subsequently decalcified for 2 weeks in decalcification buffer (PBS with 0.05 mM EDTA). Buffer was replaced every 3 days to obtain sufficient level of decalcification. Subsequently, tibias were paraffin embedded and 4 μm thick sections were cut for RUNX2 and TRAP histological analysis. All histological analyses were performed, blinded, by an independent operator using Quantimet (Leica) with Qwin3 quantification software (Leica).

M-CSF and cholesterol levels

Blood samples were collected in tubes containing 5 μL 0.5M EDTA and centrifugated for 10 minutes at 2100 rpm. Subsequently, plasma was isolated and used for M-CSF and total cholesterol measurement. M-CSF (ligand for CD115) was determined using the mouse M-CSF Quantikine ELISA kit (R&D Systems). Total plasma cholesterol levels were measured using a colorimetric assay (CHOD-PAP, Roche, Mannheim, Germany).

Flow cytometry

Upon sacrifice blood, spleen, bone marrow, and peritoneal leukocytes were collected. In blood absolute counts were obtained using BD Trucount tubes, according to the protocol supplied by the manufacturer (BD Biosciences). In short, Fc-receptor blocking antibody was added to the Trucount tubes. Subsequently, 50 μ L anti-coagulated whole blood was added and the tube was gently vortexed. After incubation for ten minutes, the antibody cocktail was added and incubated at room temperature in the dark for an additional 20 minutes. Finally, hypotonic lysis buffer was added and samples were measured on a FACS Canto II flow cytometer (BD Biosciences). Blood parameters were measured using flow cytometry for the leukocyte fraction and a Coulter counter for determining other variables. Red blood cell and platelet numbers were determined as well as hemoglobin and hematocrit levels.

Myeloid progenitors were measured in spleen and bone marrow. First, cells from spleen and bone marrow were stained with biotin-conjugated antibodies against lineage markers (CD5, CD45RA [B220], CD11b, Gr-1 [Ly6C/G], 7-4, and Ter-119; Miltenyi Biotec). Next, lineage positive cells were depleted using streptavidin conjugated magnetic beads (Miltenyi Biotec) and LS columns (Miltenyi Biotec). Lineage negative fractions were subsequently stained with antibodies for Sca-1, c-Kit (CD117), CD34, CD16/32, CD115, and lineage markers (CD5, CD45RA [B220], CD11b, Ly6C, Ly6G, Ter-119) and measured on a FACS Canto II (BD Biosciences).

Single cell suspensions were made from spleen by crushing the tissue over a 70 μ m cell strainer (BD Biosciences, NJ, USA). To measure mature leukocytes, erythrocytes in spleen and bone marrow were removed by incubation with hypotonic lysis buffer (8.4 g NH_4Cl and 0.84 g NaHCO_3 per liter distilled water). Non-specific Fc-receptor binding was blocked by the addition of anti-CD16/CD32 antibody (eBioscience, San Diego, USA). Antibodies used for these flow cytometry experiments are shown in supplemental table 1. Samples and buffers were kept on ice throughout the experiment unless indicated otherwise. All measurements were performed on a FACS Canto II (BD Biosciences) and analysis of acquired data was performed using FACS Diva software (BD Biosciences).

Colony forming unit assay

Bone marrow cells were isolated from one tibia and femur per mouse. Cells were counted twice using a Countess automated cell counter (Sigma-Aldrich) and the concentration was calculated for each sample. Per well 10,000 bone marrow cells were added to 2 mL methylcellulose medium with recombinant cytokines (MethoCult Medium, StemCell Technologies, Grenoble, France). For spleen 250,000 cells were added to 2 mL methylcellulose medium with recombinant cytokines (Stemcell Technologies). After incubation for 7 days (37°C; 5% CO₂) the total number of colonies was quantified by an independent operator, and GM-CFU, E-CFU, G-CFU, and M-CFU colonies were specified based on morphology.

Statistics

Data are expressed as mean $\pm$ SEM. To compare individual groups, 2-tailed Students t-test was used; non-parametric data were analyzed using Mann-Whitney U test. All statistical analyses were performed using GraphPad Prism 5 software (GraphPad Software Inc, La Jolla, CA, USA) and p-values below 0.05 were considered statistically significant.

Results

Systemic treatment ablates plaque macrophages, but only affects lesion composition partially

As previously described, the MaFIA mice show weight loss due to the treatment with the dimerizer (data not shown)³², the weight loss is not present in solvent treated MaFIA mice or dimerizer treated *ApoE*^{-/-} controls. Pilot experiments showed the ablation of both monocytes and granulocytes after systemic treatment for 7 days using a dose of 1 mg/kg (figure 1A+B). Effectiveness of the treatment on ablation of macrophages was verified *in vitro* using peritoneal macrophages isolated from *ApoE*^{-/-} // *MaFIA*^{+/+} and *ApoE*^{-/-} mice. In the MaFIA peritoneal macrophages an almost complete induction of dimerizer-dependent cell death (>80%) could be observed, while there was no apoptosis measurable in the control macrophages (figure 1C). For bone marrow derived macrophages *in vitro* no apoptosis could be induced. In depth analysis revealed that *in vitro* differentiation of BMDM resulted in downregulation of CD115 and thus diminished expression of the MaFIA suicide

gene on these cells (supplemental figure 1A+B). This process was shown to be dependent on the M-CSF in the culture medium, indicating a negative feedback of M-CSF on the expression of the M-CSF receptor (CD115).

Figure 1 - *In vitro* and *in vivo* depletion of myeloid cells is functional

In vivo and *in vitro* depletion of monocytes, granulocytes, and macrophages using AP20187 is efficient. **(A, B)** Monocytes **(A)** and granulocytes **(B)** can be efficiently depleted *in vivo* after 7 days of treatment with 1 mg/kg intravenous injections of AP20187 (filled bar), compared to solvent-treated controls (open bar). **(C)** *In vitro* assays show highly efficient induction of apoptosis in peritoneal macrophages from ApoE^{-/-} // MaFIA^{+/+} animals. Blue is DAPI (nuclei) and red is polycaspase (apoptosis). *** $p < 0.001$

To study the effect of ablating plaque resident macrophages on atherosclerosis development, we examined *ApoE*^{-/-} // *MaFIA*^{+/+} (MaFIA) mice treated with 1 mg/kg AP20187 via intra venous injections in the tail vein. Treatment with AP20187 did not influence plasma cholesterol levels (supplemental figure 2A+B), nor in plasma M-CSF levels (supplemental figure 2D). At baseline, we observed no differences in collar-induced plaque characteristics between *ApoE*^{-/-} mice and (untreated) *ApoE*^{-/-} // *MaFIA*^{+/+} mice (supplemental figure 3). After two weeks of treatment with AP20187, MaFIA mice did not have any changes in total plaque area (figure 2A) nor in necrotic core size (figure 2B) compared to AP20187 treated *ApoE*^{-/-} mice. Plaque area did increase over the course of the two week treatment period compared to baseline (dashed line; figure 2A+B). Collagen was decreased significantly in *ApoE*^{-/-} // *MaFIA*^{+/+} mice compared to *ApoE*^{-/-} controls (figure 2C). As expected an ablation of plaque macrophages was observed in the dimerizer-treated *ApoE*^{-/-} // *MaFIA*^{+/+} mice compared to *ApoE*^{-/-} controls (figure 2D). However, the increased apoptosis of macrophages did not result in any effects on necrotic core, indicating that this reduction could possibly be caused by decreased monocyte migration towards the atherosclerotic lesion. Surprisingly, we also noted a significant reduction in smooth muscle cells (SMC) in lesions from *ApoE*^{-/-} // *MaFIA*^{+/+} mice compared to control *ApoE*^{-/-} mice (figure 2E). To verify whether this was related to increased apoptosis of SMCs we performed a TUNEL staining. An almost 4-fold increase in SMC apoptosis could be observed indicating that also smooth muscle cells are directly or indirectly affected by the systemic treatment with AP20187 *in vivo* (figure 2F). As there were no effects observed on smooth muscle cells in the control animals, it was not solvent-induced apoptosis of SMCs, but a specific reaction to the dimerizer treatment. Next, it could be noted that lesions from MaFIA mice had less pronounced lipid cores compared to wild type animals. However, total lipid core was unchanged between *ApoE*^{-/-} // *MaFIA*^{+/+} and *ApoE*^{-/-} mice (data not shown).

Figure 2 - Systemic treatment with AP20187 reduces plaque macrophage content

(A – C) *In vivo* treatment with 1 mg/kg AP20187 via intravenous tail vein injections does not affect total plaque area **(A)** and necrotic core area **(B)**, while collagen content decreases **(C)** in *ApoE*^{-/-} // *MaFIA*^{+/+} animals. **(D, E)** Intimal macrophages content **(D)** and smooth muscle cells **(E)** are reduced in *ApoE*^{-/-} // *MaFIA*^{+/+} animals upon AP20187 treatment. **(F)** This decreased number of SMCs is accompanied by an increased SMC apoptosis in *ApoE*^{-/-} // *MaFIA*^{+/+} animals. **(G)** Overview of the experimental setup for the *in vivo* treatment with AP20187. **(H)** Representative pictures of macrophage (Mac3) and alpha smooth muscle cell actin (αSMA) staining for *ApoE*^{-/-} and *ApoE*^{-/-} // *MaFIA*^{+/+} animals. White bars are *ApoE*^{-/-} controls and black bars are *ApoE*^{-/-} // *MaFIA*^{+/+} animals. * *p* < 0.05

Myeloid cells are depleted systemically; however recover after sustained treatment

Systemic treatment (1mg/kg) resulted in decreased numbers of CD45 positive leukocytes in the circulation of AP20187 treated *ApoE*^{-/-} // *MaFIA*^{+/+} mice compared to *ApoE*^{-/-} mice after one week of treatment. This was mainly contributable to the decreased number of myeloid cells in *MaFIA* mice compared to controls. Contrary to our expectations, total leukocyte numbers recovered after two weeks of continuous treatment with AP20187 to levels equal, or even slightly increased, compared to *ApoE*^{-/-} control mice. However, leukocyte composition was clearly affected, lymphocytes were affected only

partially by the AP20187 treatment; this was shown by unaltered numbers of CD3 positive T lymphocytes (data not shown). B lymphocytes (B220+ cells) were decreased in MaFIA mice after already 1 week of treatment and did not recover after two weeks of treatment (data not shown).

Total myeloid cells were affected more drastically and showed an increase after prolonged treatment of two weeks. Monocytes (CD11b+/Ly6G-) and surprisingly also granulocytes (CD11b+/Ly6G+) were efficiently depleted in the circulation of the *ApoE*^{-/-} // *MaFIA*^{+/+} compared to *ApoE*^{-/-} mice after the first week of treatment with AP20187 (figure 3A+B). In contrast to our expectations, monocytes and granulocytes in MaFIA mice recovered to baseline values after continuous treatment for two weeks compared to short (one week) treatment (figure 3A+B). We also noted the expansion of a CD11b⁺/Ly6G^{mid} myeloid subset, not being (mature) monocytes or granulocytes, after one and two weeks of treatment with AP20187 (figure 3C). The amount of Ly6C^{high} monocytes recovered faster than the Ly6C^{low} monocytes (figure 3D) upon the depletion after 1 week of dimerizer treatment. This could be indicative of expanded production or release of Ly6C^{high} monocytes from bone marrow and spleen in the AP20187 treated MaFIA mice.

As the massive depletion of granulocytes was unexpected we analyzed the expression of both the GFP reporter and also the M-CSF receptor (CD115) on these cells. Monocytes were CD115^{high} as expected, however also granulocytes showed low to intermediate levels of the M-CSF receptor. The activity of the CD115 promotor in granulocytes was confirmed by the GFP signal, which was comparable to the GFP level in monocytes (figure 3F). Based on the GFP levels we could also determine that a subset of B-cells has low expression of the construct, explaining the limited effects observed on circulating B-cells (data not shown). Treatment with AP20187 resulted in depletion of CD115 positive cells, however GFP expression was still measurable in CD115 negative cells (figure 3F), which could be indicative of post-translational regulation of the M-CSF receptor expression.

Figure 3 - Sustained treatment with AP20187 results in disturbed leukocyte pattern

(A, B) Treatment with AP20187 results in initial depletion of monocytes (A) and granulocytes (B) after 1 week, but apparent recovery after two weeks of treatment. (C) Leukocyte patterns are disturbed upon AP20187 treatment and a novel CD11b⁺/Ly6G^{mid} population increases over time. (D) Monocyte subset (Ly6C^{high}/Ly6C⁻) ratio show a pattern that is similar to total monocytes and granulocytes, with initial depletion and subsequent recovery. (E) Representative dot plots of monocytes and granulocytes (top panels) and monocyte subsets (bottom panels). (F) Representative histograms showing CD115 (M-CSF receptor) and GFP (promotor activity) levels in monocytes, granulocytes, and control cells. White bars (controls, light purple bars (1 week treatment), and dark purple bars (2 weeks treatment). * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

Unexpectedly, analysis of splenic leukocytes revealed a significant and persistent increase in monocytes (figure 4A) and granulocytes (figure 4B) after 1 and 2 weeks of treatment. Also CD11b⁺/Ly6G^{mid} cells were massively expanded in the spleen (figure 4C). Flow cytometry also revealed that the monocyte expansion was CD115 independent, as CD115⁺ monocytes were almost completely ablated (figure 4D), and thus all remaining monocytes were CD115 negative.

Figure 4 - Spleen composition and function is perturbed by CD115⁺ cell depletion

(A – C) Treatment of MaFIA mice with 1 mg/kg AP20187 results in increased monocytes (A), granulocytes (B), and CD11b⁺/Ly6G^{mid} myeloid cells (C) in the spleen. (D) These splenic monocytes are however predominantly CD115 negative. (E) Total spleen cellularity is significantly increased after sustained AP20187 treatment in MaFIA mice compared to controls. (F, G) Flow cytometry analysis of the spleen indicated that Lineage⁻/Sca-1⁻/c-Kit⁺ (LK) cells were reduced upon treatment (F), however within this subset granulocyte/macrophage progenitors (GMP) (G) were increased. (H, I) Colony forming unit (CFU) assay showed that total number of colonies and macrophage colony forming units (M-CFU) were increased after 1 and 2 weeks of treatment in MaFIA mice, as could also be observed in representative pictures of CFU assay (I). * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

Myelopoiesis shifts to spleen after AP20187 treatment

As there was an unexpected increase in myeloid cells in the circulation (figure 3) and spleen (figure 4A-D), hematopoiesis was analyzed more in depth. As the spleen was enlarged (data not shown) and splenic cell content was increased (figure 4E), myelopoiesis was analyzed in the spleen to determine whether extramedullar hematopoiesis did occur in these treated MaFIA mice. In spleen, no effects on the LK fraction could be observed after one week of treatment. Two weeks treatment did result in a significant decrease of this progenitor subset (figure 4F). More in depth analysis did show that the decrease in LK cells was accompanied by an increase in GMP cells after both one and two weeks of treatment (figure 4G). Indeed, splenic myelopoiesis was dramatically increased after 1 and 2 weeks of treatment with AP20187. Total CFU was more than 3-fold increased, which was almost completely attributable to increased numbers of macrophage colony forming units (M-CFU), while GM-CFU and G-CFU were not significantly increased (figure 4H).

Macroscopically, it could be noted that there was a clear distinction between the bones (femur and tibia) from *ApoE*^{-/-} mice and *ApoE*^{-/-} MaFIA^{+/+} mice treated with AP20187. MaFIA mice had paler bone marrow which already indicated effects on bone marrow composition (figure 5A), which was confirmed by the decreased number of cells in the bone marrow (figure 5B). When myelopoiesis in bone marrow was analyzed by colony forming unit (CFU) assays it could be observed that there was a decrease in total CFU after 1 week of treatment, while after two weeks of treatment with the dimerizer levels normalized (figure 5C). These effects were similar to the effects observed in blood regarding monocyte and granulocyte numbers (figure 3). There were no specific subsets that were responsible for the decreased number of CFUs in the bone marrow compartment, as all subsets showed equal responsiveness to the treatment after one and two weeks of treatment (figure 5C). To confirm these results from the CFU assays, flow cytometry was performed to determine progenitors in bone marrow compartment. It could be observed that Lineage⁻/Sca-1⁻/c-Kit⁺ (LK) cells were significantly diminished in bone marrow after one week of treatment (figure 5D), which is in line with the effects observed in the CFU assays. Within this LK fraction the relative percentage of granulocyte/macrophage progenitors (GMP) was increased, indicative of a relatively small decrease or even expansion of this progenitor subset in the bone marrow (figure 5E). Overall, it could be shown that in spite of treatment

myelopoiesis did expand. Most likely this is a compensatory effect for the depleted mature myeloid cells in the periphery. Especially, the splenic hematopoiesis is thought to be responsible for the effects observed in blood and spleen regarding the recovery of myeloid cells after prolonged treatment.

Systemic treatment results in bone marrow osteoclast depletion

As we observed massive extramedullar hematopoiesis in MaFIA mice treated systemically (i.v. injections), we sought out to investigate which mechanism is responsible for the peripheral phenotype observed. Since we observed a shift in hematopoiesis from the bone marrow to the splenic compartment, we wanted to analyze the effect of the dimerizer (AP20187) treatment on cells involved in maintaining normal bone marrow niches. Staining for osteoclasts in decalcified femur sections showed that osteoclast levels were mildly reduced in MaFIA mice (supplemental figure 4).

Figure 5 - Stromal hematopoiesis is decreased in treated MaFIA mice

(A) Femurs from AP20187 treated MaFIA mice were visibly paler than femur from control animals. (B) Total cells per bone were decreased in MaFIA mice. (C) Colony forming unit (CFU) assay showed that total number of colonies (and subsets) were decreased after 1 week of treatment, but recovered after two weeks of sustained treatment in MaFIA mice. (D, E) Flow cytometry analysis of the bone marrow indicated that Lineage⁻/Sca-1⁺/c-Kit⁺ (LK) cells were reduced upon treatment (D), however within this subset granulocyte/macrophage progenitors (GMP) (E) were increased. (F) Representative dot plots indicating relevant gating strategy. ** $p < 0.01$, *** $p < 0.001$

Local treatment with AP20187 accelerates atherogenesis

To exclude any systemic effects on atherosclerosis development, we performed a second atherosclerosis study in which animals received a collar around the carotid arteries combined with an osmotic minipump with catheters attached to the perivascular side of the carotid arteries near the collar. The dimerizer AP20187 was constantly delivered for two weeks at a dose of 0.01 mg/kg body weight, which did not affect plasma cholesterol (supplemental figure 2C). Analysis of the carotid arteries after two weeks of treatment showed that total plaque area was over two-fold increased when the animals were treated continuously and locally compared to treated *ApoE*^{-/-} controls (figure 6A). Necrotic core (figure 6B) and collagen content (figure 6C) were equally expanded in the treated MaFIA mice. As expected, macrophages were not increased in the plaques from MaFIA mice compared to the controls despite the massively expanded plaque size (figure 6D). Intra-plaque smooth muscle cells were in contrast to the systemic study not affected by the local treatment with the dimerizer (figure 6E+F). Analysis of circulating leukocytes did not reveal any significant differences between *ApoE*^{-/-} MaFIA^{+/+} mice and *ApoE*^{-/-} controls (supplemental figure 5).

Figure 6 - Local treatment to the atherosclerotic lesion increases lesion formation

(A) Local treatment with AP20187 results in aggravated atherosclerosis formation in MaFIA mice. (B, C) Local treatment does not affect necrotic core (B), and plaque collagen content (C) in MaFIA mice compared to controls. (D) Intimal macrophage content is reduced upon treatment. (E, F) However, smooth muscle cell (SMC) content (E), and SMC apoptosis (F) are unaffected by local treatment. (G) Representative pictures of haematoxylin and eosin (H&E) stained lesions from control and MaFIA mice. * $p < 0.05$

Discussion

The MaFIA model for induction of CD115⁺ cell apoptosis in *ApoE*^{-/-} pro-atherosclerotic mice was used for this study. Cells of the myeloid lineage express an inducible suicide gene under transcriptional control of the CD115-promoter³². Apoptosis is prompted by administration of AP20187 which results in dimerization of extracellular FK domains and subsequently their cytoplasmic FAS domains triggering the extrinsic pathway of cell apoptosis³².

Macrophages are essential in atherosclerotic lesion development and progression, and contribute to clinical outcomes by destabilizing the atherosclerotic lesion^{35,36}. Macrophages have an important role throughout atherogenesis, from early intimal thickening until advanced rupture-prone atheromas³⁷. However, it has been difficult to unravel the role of plaque macrophages, next to circulating myeloid cells, in relation to the progression of the lesions. Here, we show that specific depletion of these cells, in a FAS dependent manner, exposed both local and systemic effects which contribute differentially to atherosclerosis, but also to the conservation of normal hematopoiesis.

Lesion composition in general was not affected by systemic ablation of macrophages, despite a substantial reduction of plaque resident macrophages. Vascular smooth muscle cells (VSMC) were partially reduced by the whole body treatment. However, it has been shown that VSMC which are in contact with an atherosclerotic lesion can express the M-CSF receptor^{38,39}, explaining the depletion of these cells *in vivo*. These smooth muscle cells show also other characteristics of the monocyte/macrophage lineage such as phagocytic capacity and even the transdifferentiation into foam cells³⁸. Nevertheless, the increased apoptosis of VSMC and macrophages did not contribute to any effects on atherosclerotic lesion phenotype.

Peripheral leukocyte analysis revealed a dramatic decrease in total leukocytes after seven days of AP20187 treatment. This reduction was almost completely contributable to ablation of myeloid cells, both monocytes/macrophages and granulocytes, with minor effects on B-lymphocytes. The effects on granulocytes and B-cells were in part unexpected, however could be explained by the role of PU.1 in controlling M-CSF receptor expression^{40,41}. PU.1 was already shown to be involved in myeloid cell development and B-cell development in a PU.1 knock-out model⁴². The ablation of granulocytes, but also B-cells, can be related to the PU.1 dependent

transcription of the M-CSF receptor and thus also transcription of the MaFIA transgene. The transcription, and even expression, of the M-CSF receptor on both B-cells and granulocytes has only been scarcely described so far, but could have profound impact on the current knowledge on myelopoiesis and cell maturation.

Systemic ablation of M-CSF receptor (CD115) positive macrophages and monocytes was shown to result in disturbed myelopoiesis with increased extra-medullar hematopoiesis and the release of immature myeloid cells into the circulation. The production and release of progenitors in the bone marrow is amongst others controlled by bone marrow monocytes, macrophages, and osteoclasts, which are considered to be related to the myeloid lineage⁴³. Osteoclasts respond to similar cytokines as macrophages, and also express comparable surface markers, including the M-CSF receptor⁴⁴. We were able to show that *in vivo* treatment with AP20187 results in decreased numbers osteoclasts. As macrophages as well as osteoclasts are ablated upon AP20187 treatment, the relevant contribution of each these cell types to the observed neutrophilia cannot be directly derived from these data⁴⁴⁻⁴⁶. However, in a G-CSF mobilization study Winkler *et al* argued that bone marrow macrophages are the most important cell type in regulating stromal release⁴⁵. Furthermore, osteoclast activation is associated with decreased stromal release⁴⁷, yet osteoclast activation is increased under inflammatory conditions⁴⁸. Osteoclast depletion may also be responsible for the observed effects on B-lymphocytes as the development of mature B-cells also is regulated by these cells in the bone marrow⁴⁹. Overall, systemic AP20187 treatment was shown to have profound effects on myelopoiesis, which could, at least in part, be explained by depletion of cells responsible for maintaining normal bone marrow niches.

As the unanticipated effects observed on the myelomonocytic cells after systemic dimerizer administration could well have masked local effects of macrophage ablation, we analyzed plaque-specific effects in a separate study. The dimerizer was administered locally to the carotid arteries via catheterized osmotic minipumps. Site-restricted administration of the dimerizer was shown not to result in any significant systemic effects. In the atherosclerotic lesion we did observe a completely inhibited expansion of the number of intimal macrophages. In contrast to systemic administration of the dimerizer, plaque size was increased more than two-fold in animals treated locally. As macrophage content was similar between the treated MaFIA mice and

controls, the plaque and necrotic core expansion were most likely caused by the immediately induced cell death of newly infiltrating monocytes. As monocytes are not ablated peripherally in mice treated locally, more monocytes will migrate towards the inflammatory locus⁵⁰. Hereby, more cells will enter the lesion, where AP20187-dependent apoptosis is induced. Due to an inhibited expansion of mature macrophages, and thus decreased efferocytosis, these apoptotic monocytes will be cleared less efficiently giving rise to secondary necrosis and progressive development of the necrotic core and total plaque. Taken together, our data confirm the current paradigm that influx of monocytes might be more important than reduction of plaque macrophages via efflux of apoptosis for controlling atherosclerotic plaque development⁵⁰⁻⁵³. Another explanation may lie in the mildly shifted Ly6C^{high} to Ly6C^{low} monocyte balance which may have favored the differentiation into classically activated pro-inflammatory macrophages⁵⁴⁻⁵⁶.

Overall, we can conclude that macrophages, and other M-CSF receptor positive cells, are critical controllers of many processes essential to atherogenesis, including hematopoiesis. Depletion of macrophages will result in disturbed physiology indicative of the essential role of myeloid cells in maintaining health and reducing plaque development.

Acknowledgements

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Supplemental figures

Supplemental figure 1 - M-CSF reduces CD115 expression on bone marrow derived macrophages

(A, B) M-CSF containing medium (LCM) results in down regulation of CD115 (M-CSF receptor) within 24 hours compared to M-CSF free cultured bone marrow derived macrophages.

Supplemental figure 2 - Plasma cholesterol is not affected by AP20187 treatment

(A, B, C) Plasma cholesterol is not affected at baseline (A), after intravenous treatment with AP20187 (B), or after local treatment (minipump) with AP20187 (C) of MaFIA mice compared to controls. (D) Plasma M-CSF levels after 1 and 2 weeks intravenous AP20187 administration.

Supplemental figure 3 - At baseline no differences in plaque size and morphology could be detected

At baseline no differences in total plaque area (A), necrotic core (B), collagen content (C), intimal macrophage content (D), smooth muscle cell content (E), and apoptosis (F) could be detected between control animals and AP20187 treated MaFIA mice.

Supplemental figure 4 - Osteoclasts are depleted from the bone upon AP20187 treatment

(A) Osteoclasts were stained and quantified in the bones of solvent-treated (control) animals (white bar) and dimerizer (AP) treated animals (black bar). A clear reduction in the number of osteoclasts could be shown in the AP-treated animals. (B) Representative pictures of bones from control and AP-treated animals. Osteoclasts are stained in purple, arrowheads indicate the positive cells. Note that in the AP-treated animals the staining is less pronounced than in the control animals. * $p < 0.05$

Supplemental figure 5 - Local treatment does not induce systemic disturbances in leukocyte composition

After local treatment with AP20187, no differences could be detected in circulating lymphocytes **(A)**, granulocytes **(B)**, and monocytes **(C)** between control animals and AP20187 treated MaFIA mice.

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The background of the entire slide is a white surface covered with numerous red splatters and droplets of varying sizes, resembling blood. Some splatters are large and circular, while others are thin, elongated streaks or fine mist-like particles. The distribution is dense, with some areas having more large droplets and others having more fine spray.

Chapter 03

Hematopoietic G-protein-coupled receptor kinase 2 deficiency decreases atherosclerotic lesion formation in LDL receptor- knockout mice

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Abstract

Leukocyte chemotaxis is deemed instrumental in initiation and progression of atherosclerosis. It is mediated by G-protein-coupled receptors (*e.g.*, CCR2 and CCR5), the activity of which is controlled by G-protein-coupled receptor kinases (GRKs). In this study, we analyzed the effect of hematopoietic deficiency of a potent regulator kinase of chemotaxis (GRK2) on atherogenesis. LDL receptor-deficient (*LDLr*^{-/-}) mice with heterozygous hematopoietic GRK2 deficiency, generated by bone marrow transplantation (*n*=15), displayed a dramatic attenuation of plaque development, with 79% reduction in necrotic core and increased macrophage content. Circulating monocytes decreased and granulocytes increased in *GRK2*^{-/-} chimeras, which could be attributed to diminished granulocyte colony-forming units in bone marrow. Collectively, these data pointed to myeloid cells as major mediators of the impaired atherogenic response in *GRK2*^{-/-} chimeras. *LDLr*^{-/-} mice with macrophage/granulocyte-specific GRK2 deficiency (*LysM-Cre GRK2*^{flox/flox}; *n*=8) failed to mimic the aforementioned phenotype, acquitting these cells as major responsible subsets for GRK2 deficiency-associated atheroprotection. To conclude, even partial hematopoietic GRK2 deficiency prevents atherosclerotic lesion progression beyond the fatty streak stage, identifying hematopoietic GRK2 as a potential target for intervention in atherosclerosis.

Introduction

Leukocyte migration toward atherosclerotic lesions is deemed instrumental in atherosclerosis initiation and progression ¹. By releasing cytokines ² and proteolytic enzymes, infiltrated leukocytes may also promote plaque destabilization, a process generally preceding thrombotic rupture and subsequent acute cardiovascular events. Because leukocyte migration may be a potentially attractive target for the prevention or control of plaque destabilization and ischemic disease, this research focuses on the pivotal role of leukocyte migration in atherosclerosis.

Chemokines, a family of cytokines with strong chemotactic capacity ^{3,4}, are key regulators of leukocyte transmigration into the vessel wall during atherogenesis ^{5,6}. Chemokines are inducible and control cellular recruitment, especially to inflammatory sites ⁷. In response to inflammatory stimuli, chemokines can be released by various cell types relevant to atherosclerosis, including endothelial cells, platelets, and leukocytes ⁸⁻¹¹. They act by binding to dedicated receptors of the G-protein-coupled receptor (GPCR) family. GPCRs are coupled to heterotrimeric G proteins and induce, after ligand binding, activation of downstream signaling cascades ^{12,13}.

The expression and activity of GPCRs is regulated not only at the mRNA or protein level but also functionally. One such mechanism is receptor desensitization, which dampens the response to prolonged or repeated stimuli ^{14,15}. Dedicated GPCR kinases (GRKs) can induce receptor desensitization by phosphorylation of the ligand-occupied receptor, thereby enhancing its affinity for arrestin proteins. Binding of arrestins to the phosphorylated receptor will result in uncoupling and internalization of the receptor ^{16,17}. The GRK family comprises 7 serine/threonine kinases ^{18,19}. They regulate a range of processes, including cell migration ^{20,21}, neuronal pathways ²², and cell survival ²³. Dysfunction of GRKs, particularly GRK2, has been implicated in various human pathologies, such as multiple sclerosis ²⁴, rheumatoid arthritis ²⁵, chronic pain ²⁶, and heart failure ²⁷, whereas GRK2 overexpression induces hypertension ^{28,29}. Recently, GRK5 was shown to be involved in atherogenesis in mice through smooth muscle cells and endothelial cells. However, monocyte migration and function were also affected by GRK5 deficiency ³⁰. Because key chemokine receptors in atherosclerosis [CC-motif chemokine receptor (CCR) 2 and CCR5] are targeted by GRK2 ^{ref 31}, decreased levels of this receptor kinase conceivably result in excessive migration of inflammatory cells toward

atherosclerotic lesions. Therefore, the aim of this study was to assess the role of leukocyte specific GRK2 deficiency on the development of atherosclerosis in LDL receptor (LDLr)-deficient mice.

Materials & Methods

Animals

Female *LDLr*^{-/-} mice³² were obtained from the animal breeding facility at Maastricht University (Maastricht, The Netherlands). Mice were fed a regular chow diet (RM3; Special Diet Services, Essex, UK) or a Western-type diet (WTD; 0.25% cholesterol; Special Diet Services). Drinking water and food were provided *ad libitum*. Experiments were performed at the Maastricht University animal facility. Male heterozygous GRK2 (*GRK2*^{-/-}) mice³³, *LysM-Cre GRK2*^{flox/flox} mice^{34,35}, and littermate controls were used as donors for bone marrow transplantations and *in vitro* assays. Donor mice were maintained at the Utrecht University Medical Center (Utrecht, The Netherlands) animal facilities. All experimental protocols were approved by the Maastricht University ethics committee for animal experiments.

Bone Marrow Transplantation

Female *LDLr*^{-/-} mice were lethally irradiated with a single dose (9 Gy, 0.5 Gy/min, MU 15 F/225 kV; Philips Healthcare, Best, The Netherlands) total-body irradiation 1 day before transplantation. On the next day, irradiated recipients received an intravenous tail vein injection of 5 x 10⁶ bone marrow cells. Bone marrow was isolated from *GRK2*^{-/-} and littermate controls or *LysM-Cre GRK2*^{flox/flox} and littermate controls by flushing femurs and tibiae. Drinking water supplemented with antibiotics [60 U/mL polymyxin B sulfate (Sigma-Aldrich, St. Louis, MO, USA) and 100 g/mL neomycin (Sigma-Aldrich)] was provided starting 1 week before the bone marrow transplantation. After 6 weeks of recovery, mice were fed a WTD for 12 weeks and subsequently euthanized.

In vivo Migration and Proliferation

Female wild-type (WT) controls and *GRK2*^{-/-} chimeras (n=4/5) were intraperitoneally injected 24 hours before euthanasia with PBS or CC-motif chemokine ligand (CCL) 5 [RANTES; 200 ng/mouse; Peprotech, London, UK] to

stimulate cell migration. At euthanasia, leukocytes were isolated by peritoneal lavage (sterile PBS), and lavages were subsequently analyzed for leukocyte content on an automated differential cell counter (XT-2000i; Sysmex, Norderstedt, Germany).

Splenocytes were isolated from spleen, after the tissue was gently homogenized over a 70 μm cell strainer (BD Biosciences, San Diego, CA, USA). To remove erythrocytes, cell suspensions were incubated in ice-cold erythrocyte lysis buffer (155 mM NH_4Cl in 10 mM Tris/HCl, pH 7.2) for 3 min. Splenocytes ($n=6/\text{group}$) were then resuspended in RPMI 1640 medium (Invitrogen, Paisley, UK) supplemented with l-glutamine, 100 U/mL streptomycin/penicillin, and 10% FCS and cultured for 48 h in quadruplicate in a 96-well round-bottom plate (2×10^5 cells/well; Corning, Lowell, MA, USA). Cells were stimulated with CCL5 (RANTES; 100 ng/mL). Concanavalin A (ConA; 8 $\mu\text{g}/\text{mL}$; Sigma-Aldrich) was used as a positive control. Proliferation was determined after addition of [^3H]thymidine (0.5 Ci/well; Amersham Biosciences, Roosendaal, The Netherlands), incubation of the cells for the last 16 h, and measurement of cell-associated (incorporated) [^3H]thymidine by a liquid scintillation analyzer (Tri-Carb 2900R; PerkinElmer, Waltham, MA, USA).

Histological Analysis

For the *GRK2*^{-/-} study, aortic roots were isolated, and 10- μm cryostat sections were stained with Oil-Red-O. Total plaque area and necrotic core size were determined in 7 sections/animal. Corresponding sections on separate slides were stained immunohistochemically with a macrophage-specific antibody (MOMA-2; Serotec, Oxford, UK). For the *LysM-Cre GRK2*^{flox/flox} study, aortic root sections were collected, formalin-fixed, and paraffin-embedded. Subsequently, 4 μm -thick sections were stained with hematoxylin and eosin. Lesion size and necrotic core area were determined in 3-7 sections/animal. Corresponding sections on separate slides were stained immunohistochemically with a macrophage-specific antibody (MAC-3; BD, Franklin Lakes, NJ, USA). Movat's pentachrome stain was used to elastin and to obtain insight into plaque progression stage and stability. Plaque collagen content was quantified based on Sirius Red staining and subsequent analysis of the plaques under polarized light. Vascular smooth muscle cells were analyzed more in depth by α -smooth muscle cell-specific actin (αSMA ; Sigma-Aldrich) staining. T-lymphocyte infiltration was quantified based on staining with a CD3 ϵ -specific antibody

(CD3ε; Dako, Glostrup, Denmark). Apoptosis was visualized using a TUNEL kit (Roche, Woerden, The Netherlands). Apoptotic cell content was determined by assessment of the TUNEL-positive area per section. Histological analyses were performed in a blinded manner by an independent operator using Quantimet (Leica, Wetzlar, Germany) with QWin3 quantification software (Leica).

***In vitro* Macrophage Culture**

Bone marrow cells were isolated from WT and *GRK2*^{-/-} mice, and cells were pooled and cultured in RPMI 1640 medium supplemented with fetal calf serum and L929 conditioned medium to generate bone marrow-derived macrophages (BMDMs) as described previously³⁶. Medium was replaced every 3 days, and differentiated BMDMs were used for *in vitro* assays after 7 days.

Macrophage Foam Cell Formation Assay

BMDMs were cultured and plated in 24-well cell cluster plates (Corning). Macrophages were incubated for 24 hours with VLDL (25 µg/mL) to induce foam cell formation. Subsequently, Oil Red O staining was performed to visualize the cellular lipid accumulation. Pictures were taken using a DM IL LED fluorescence microscope (Leica) with QWin3 software, and foam cell formation was scored in a blinded manner by an independent operator, calculating the number of positive Oil Red O-stained cells, as well as the staining intensity at a per cell basis in >3 fields/condition (x100 view).

Macrophage Apoptosis and Phagocytosis Assay

BMDMs were cultured and plated in 48-well cell cluster plates (Corning). Macrophages were incubated for 18 hours in plain medium or medium supplemented with camptothecin (CPT; 500 µM) or oxidized LDL (oxLDL; 50 µg/mL). oxLDL was produced as described previously³⁷. Cells were labeled with annexin V-Alexa 488 (Invitrogen) and stained with propidium iodide to detect apoptosis and necrosis, respectively. Pictures were taken using a DM IL LED fluorescence microscope with Qwin3 software. Overlays were analyzed with ImageJ software (U.S. National Institutes of Health, Bethesda, MD, USA) to quantify the number of apoptotic and necrotic cells per field.

To investigate phagocytic capacity, cultured BMDMs were labeled with Cell Tracker Red (Invitrogen). Jurkat T cells labeled with Cell Tracker Green (Invitrogen) were exposed to a UV light source (UVS-26, 6-W bulb, 0.02 J/s/cm²; UVP, Upland, CA, USA) for 20 minutes and incubated (37°C with 5%

CO₂) for 45 minutes to induce apoptosis. Apoptotic Jurkat cells were added to the BMDMs at a ratio of 5:1 and incubated for 50 minutes. Macrophages were fixed with 1% paraformaldehyde and counterstained with DAPI (Invitrogen). Pictures were taken (DM IL LED fluorescence microscope), and overlays were analyzed with ImageJ.

Flow Cytometry

At euthanasia, blood, spleen, and peritoneal leukocytes were collected. Blood and peritoneal leukocytes were analyzed for cellular composition on an automated differential cell counter (XT-2000i). Single-cell suspensions were made from spleen by crushing the tissue over a 70- μ m cell strainer (BD) after treatment with DNase and Liberase (Roche). Erythrocytes in blood and spleen were removed by incubation with hypotonic lysis buffer (8.4 g of NH₄Cl and 0.84 g of NaHCO₃ per liter of distilled water). Nonspecific Fc receptor binding was blocked by the addition of anti-CD16/CD32 antibody (eBioscience, San Diego, CA, USA). To determine lymphocyte subsets, cells were labeled with CD3 ϵ -FITC, CD8 α -eFluor450, B220-PE-Cy7, CD25-APC (all eBioscience), and CD4-PerCP (BD). To detect monocytes and granulocytes, the cells were labeled with CD11b-PE-Cy7, Ly6G-PE (BD), and Ly6C-FITC (Miltenyi Biotec, Bergisch Gladbach, Germany). Dendritic cells were detected by labeling with CD3 ϵ -PerCPCy5.5, CD19-PerCP-Cy5.5, MHC II-FITC, CD8 α -eFluor450 (all eBioscience), CD11c-PE-Cy7, and CD4-APC-H7 (both BD) for resident dendritic cells. Chemokine receptor expression was determined by labeling peripheral blood leukocytes with CCR5-PE, CCR7-PerCP-Cy5.5, CD3 ϵ -eFluor450, NK1.1-APC (all eBioscience), CD11b-PE-Cy7, Ly6G-APC-Cy7, B220-V500 (all BD), and Ly6C-FITC (Miltenyi). After labeling, the samples were washed and analyzed on a FACSCanto II flow cytometer (BD). Samples and buffers were kept on ice throughout the experiment. Gating strategies used for the different flow cytometry stainings are shown in the supplemental data.

Colony-Forming Unit (CFU) Assay

Bone marrow cells were isolated from one tibia and one femur per mouse. *WT*, *GRK2*^{-/-} and *LysM-Cre GRK2*^{flox/flox} chimeras were used for the CFU assays. Cells were counted twice using a count chamber, and the concentration was calculated for each sample. A total of 10,000 bone marrow cells/well were added to 2 mL of methylcellulose medium with recombinant cytokines (MethoCult Medium; StemCell Technologies, Grenoble, France). After incubation for 7 days (37°C with 5% CO₂), the total number of colonies was quantified by an independent operator, and granulocyte-macrophage CFU (GM-CFU), granulocyte CFU (G-CFU), and macrophage CFU (M-CFU) colonies were specified based on morphology of the individual cells and the colony as a whole.

Statistical Analysis

Data are expressed as mean $\pm$ sem. To compare individual groups, a 2-tailed Student's *t* test was used; nonparametric data were analyzed using a Mann-Whitney *U* test. All analyses were performed using GraphPad Prism 5 software (GraphPad Software Inc., LA Jolla, CA, USA), and values of *P* < 0.05 were considered statistically significant.

Results

Partial Leukocyte GRK2 Deficiency Attenuates Atherosclerosis

Partial hematopoietic GRK2 deficiency did not affect body weight and total cholesterol levels (Supplemental Fig. S1). Hematopoietic *GRK2*^{-/-} chimeras in *LDLr*^{-/-} mice (*n*=15) showed significantly decreased atherosclerotic lesion development in the aortic root compared with that of controls ($403.0 \pm 43.8 \times 10^3$ vs. $585.0 \pm 56.4 \times 10^3 \mu\text{m}^2$, *P*=0.017; Fig. 1A). The necrotic core size was 4-fold smaller in heterozygous GRK2 than in WT chimeras (9.5 ± 2.3 vs. $44.6 \pm 6.1\%$, *P*<0.0001; Fig. 1B). The TUNEL-positive apoptotic cell burden (Fig. 1C) did not differ between GRK2 and WT chimeras. Further differentiation for TUNEL-positive cell density in the cap vs. the atheroma revealed that in both WT and *GRK*^{-/-} chimeras 95% of TUNEL-positive cells were located in the atheroma (data not shown). Moreover, because we did not observe a shift in localization over these two compartments and because the atheroma mainly consists of macrophages (next to cell debris), we may infer that partial GRK2 deficiency

has not affected macrophage apoptosis. Next, collagen content was not affected in *GRK2*^{-/-} chimeras compared with WT chimeras (Fig. 1D).

Surprisingly, intimal MOMA-2-positive macrophage content in heterozygous *GRK2* chimeras was 2-fold higher than that of controls (32.3 ± 2.9 vs. $15.3 \pm 2.4\%$, $P=0.00015$; Fig. 1E). Analysis of the lesions did not reveal significant differences in intimal (CD3ε+) lymphocyte numbers (data not shown). Intimal smooth muscle cell content, as assessed by smooth muscle cell-specific actin staining, was increased by heterozygous *GRK2* deficiency (Fig. 1F). General assessment of plaque morphology, from Movat's pentachrome staining, showed a significant reduction of overall necrosis in lesions from *GRK2*^{-/-} chimeras (Supplemental Fig. S2A). In line with the results obtained based on MOMA-2 staining, foam cell macrophage content was significantly increased in *GRK2*^{-/-} lesions (Supplemental Fig. S2B). For neutrophil infiltration, it was observed that the number of granulocytes per lesion were very low, which was confirmed by the virtual absence of positive Ly6G staining (data not shown), whereas intraplaque/adventitial neutrophil numbers tended to be decreased in heterozygous *GRK2* animals, based on morphological scoring of the lesions (Supplemental Fig. S2C). We did, however, observe a significant decrease in the amount of cells adhering to the luminal side of the atherosclerotic lesions in the *GRK2*^{-/-} chimeras (Supplemental Fig. S2D), which could not be further attributed to a specific cell type. Overall, atherosclerotic lesions in *GRK2*^{-/-} chimera were considered less advanced and more stable than those of controls, despite unchanged collagen levels (Supplemental Fig. S2E), based on plaque morphology and composition.

Figure 1 - Atherosclerotic plaque development is attenuated in GRK2^{+/-} chimeras.

(A) Plaque area and (B) necrotic core size were determined based on Oil-Red-O staining. (C) Relative number of apoptotic cells was determined by TUNEL staining. (D) Collagen content was measured by analysis of Sirius Red staining under polarized light. (E) Macrophage content was measured in sections stained for the monocyte/macrophage marker MoMa-2. (F) Smooth muscle cell content was measured by an staining against alpha-smooth muscle cell actin (αSMA). (G) Representative pictures of Oil-Red-O, Sirius Red (polarized light; insert bright field image), and MoMa-2 staining are shown for wild type and GRK2^{+/-} chimeras. The top panels represent WT and the bottom panels GRK2 chimeras. Wild type controls (WT) are shown in white and GRK2^{+/-} chimeras are shown in black; scale bars are 100 μm. * $p < 0.05$ and *** $p < 0.001$

Figure 2 – Phagocytosis, apoptosis, and foam cell formation of bone marrow derived macrophages is not affected in vitro.

(A) The phagocytic capacity of BMDM for apoptotic Jurkat T-cells was determined *in vitro*. Phagocytosis was determined by scoring yellow staining in overlays from macrophages (cell tracker red), apoptotic Jurkat T-cells (cell tracker green) and nuclei (DAPI; blue). **(B)** *In vitro* apoptosis and **(C)** necrosis of bone marrow-derived macrophages was measured at baseline (-) and upon camptothecin (CPT), or oxidized LDL (oxLDL) exposure. Cell death was quantified based on Annexin V-Alexa488 (apoptosis; green) and propidium iodide staining (necrosis; red/orange). **(D)** *In vitro* foam cell formation was measured upon VLDL exposure for 24 hours using Oil-Red-O staining (red; 400 x magnification). White bars represent wild type controls (WT) and black bars GRK2^{+/-} chimeras (GRK2^{+/-}). ** $p < 0.01$.

Macrophage GRK2 Deficiency does not affect Phagocytosis, Apoptosis, or Foam Cell Formation

The major differences in plaque composition, regarding necrotic core and intimal macrophage content, prompted us to study the effect of GRK2 deficiency on macrophage function. Reduced GRK2 expression did not affect phagocytosis of apoptotic Jurkat T cells by BMDMs (Fig. 2A), rendering it unlikely that the reduced necrotic core size is attributable to increased phagocytosis.

In line with the unchanged TUNEL⁺ cell density observed in plaques from WT and GRK2^{-/-} chimeras, *in vitro* we observed no effects of GRK2

heterozygosity ($n=9$) on the susceptibility for apoptosis of BMDMs at baseline or in the presence of CPT (Fig. 2B), nor did we observe differences in necrosis between BMDMs at baseline and after camptothecin (CPT) treatment (Fig. 2C). Notably, oxLDL-induced necrosis was markedly attenuated in *GRK2*^{-/-} compared with WT BMDMs (30.1 ± 7.2 vs. $57.2 \pm 7.6\%$, $P=0.019$; Fig. 2C). Thus, partial GRK2 deficiency was seen to reduce oxLDL-induced necrosis, which could have contributed to the phenotypic differences in necrotic core size *in vivo*.

Figure 3 - *GRK2*^{+/-} chimeras have perturbed leukocyte patterns in circulation.

(A) Absolute number of white blood cells (WBC) were measured using a Sysmex differential cell counter. The relative fractions of leukocyte subsets were measured using flow cytometry analysis on a FACS Canto II (BD). (B) Granulocytes, (C) monocytes, (D) monocyte Ly6C subsets, (E) B-cells, and (F) T-cells were measured in the circulation of WT control (white bars) and *GRK2*^{+/-} chimeras (black bars). (G) Myeloid subsets were measured in bone marrow of donor mice. (H) Representative histograms of Ly6C^{high} and Ly6C^{low} distribution in blood of WT and *GRK2*^{+/-} chimeras. * $p < 0.05$ and ** $p < 0.01$.

Because macrophage and foam cell macrophage content was significantly increased in the $GRK2^{+/-}$ lesions compared with the WT lesions, we investigated *in vitro* whether GRK2 heterozygosity by itself influences foam cell formation. However, we were unable to detect any effects on the number of macrophage-derived foam cells nor on the level of lipid accumulation per cell after VLDL incubation (Fig. 2D), indicating that the observed plaque foam cell effects are not caused by GRK2 deficiency-associated changes in macrophage lipid handling.

Circulating Leukocytes are affected by GRK2 Deficiency

Peripheral blood analysis revealed increased circulating leukocyte numbers in $GRK2^{-/-}$ chimeras ($5.9 \pm 0.6 \times 10^6$ vs. $4.1 \pm 0.4 \times 10^6$ cells/mL, $P=0.03$; Fig. 3A). In-depth analysis of peripheral blood samples from WT and $GRK2^{-/-}$ chimeras fed a WTD (0.25% cholesterol) by flow cytometry indicated that there was no relative difference in granulocyte ($CD11b^+/Ly6G^+$) abundance (Fig. 3B). Circulating monocyte ($CD11b^+/Ly6G^-$) numbers were significantly lowered in $GRK2^{-/-}$ compared with those in WT chimeras (12.09 ± 0.61 vs. $16.09 \pm 1.58\%$ monocytes, $P=0.039$; Fig. 3C) and displayed a decreased $Ly6C^{high}/Ly6C^{low}$ ratio (Fig. 3D). No genotype-related differences were observed in B cells (B220+; Fig. 3E) and T cells ($CD3\epsilon^+$; Fig. 3F) or in T-cell subsets ($CD4^+/CD8^+$; data not shown). Spleen resident dendritic cell content ($CD11c^+$) and resident dendritic cell composition ($CD4^+/CD8^+$ /double-negative; data not shown) did not reveal any differences between WT and $GRK2^{-/-}$ chimeras. In spleen, we did not observe any significant differences other than a relative decrease in the number of granulocytes in the $GRK2^{-/-}$ chimeras compared with that in WT controls (3.30 ± 0.32 vs. $2.15 \pm 0.21\%$ granulocytes; Supplemental Fig. S3A). B cells and T cells did not show any differences in spleen (Supplemental Fig. S3B, C), while splenic monocytes, and in particular the $Ly6C^{high}$ subset, showed a tendency toward reduced numbers, as observed in peripheral blood (Supplemental Fig. S3D, E). In bone marrow, on the other hand, we observed no differences in monocytes or monocyte subsets (Fig. 3G). This result also indicates that impaired egress might be underlying the observed monocytopenia. Granulocytes are accumulating in the bone marrow of GRK2-deficient mice (Fig. 3G), but this does not translate into any effects on the atherosclerotic lesion. In short, these findings indicate that the effects of GRK2 deficiency on plaque formation and on circulating leukocytes mainly affect the myeloid lineage, monocytes and granulocytes, in blood, spleen, and bone marrow.

Figure 4 – Chemokine receptor expression on leukocytes from WT and GRK2^{+/-} chimeras.

Chemokine receptor (CCR5 and CCR7) expression was measured on circulating leukocytes from WT control (white bars) and GRK2^{+/-} chimeras (black bars). **(A)** CCR5 expression on T-cells, B-cells, NK-cells, granulocytes, and **(B)** monocytes. **(C)** Ly6C monocyte subsets showed differential CCR5 expression. **(D)** CCR7 expression on T-cells, B-cells, NK-cells, and granulocytes.

The reported role of GRK2 in chemokine receptor desensitization led us to investigate effects of GRK2^{-/-} deficiency on leukocyte expression of relevant chemokine receptors such as CCR5 and CCR7 by flow cytometry. No statistically significant increase in CCR5 expression could be observed in any of the leukocyte subsets at baseline (Fig. 4A). However, CCL5-stimulated monocytes and, in particular, the proinflammatory Ly6C^{high} subset, showed borderline significance toward increased CCR5 expression in GRK2^{-/-} chimeras compared with WT controls ($P=0.051$; Fig. 4B, C). For CCR7 we could not detect any significant differences in expression for any leukocyte subset tested (Fig. 4D).

Baseline Proliferation and CCL5-mediated Migration are increased in GRK2^{-/-} Animals

Proliferation was measured by [³H]thymidine incorporation in total splenocytes under baseline conditions (PBS) or after stimulation for 16 h with CCL5 (RANTES) or with the general mitogen ConA. It was observed that baseline proliferation (control) was increased in GRK2^{-/-} splenocytes compared with that in WT controls ($P<0.05$; Fig. 5A). However, the proliferation index in CCL5 or ConA stimulated cells, corrected for baseline proliferation, was unaffected in GRK2^{-/-} splenocytes (Fig. 5B).

Figure 5 – Baseline proliferation and migration upon CCL5 stimulation is increased in GRK2^{+/-} chimeras

(A) Baseline proliferation (unstimulated) was determined in WT and GRK2^{+/-} splenocytes. (B) Proliferation index (corrected proliferation for control condition) was determined for CCL5 (100 ng/mL; RANTES) or Concanavalin A (100 ng/mL; ConA) stimulation. (C) Total cell migration and (D) monocyte migration in WT control and GRK2^{+/-} chimeras upon i.p. CCL5 (200 ng; RANTES) challenge. * $p < 0.05$ and # $p < 0.05$ compared to GRK2^{+/-} PBS-treated.

Peritonitis is considered an established model to study leukocyte chemotaxis and activation and expansion outside of the atherosclerotic lesion. We opted to use CCL5-dependent peritonitis, because CCL5 is one of the chemokines desensitized by GRK2, which at the same time is considered to be relevant in atherosclerosis. *In vivo* migration was measured after intraperitoneal injection of PBS (control) or CCL5 (RANTES; 200 ng/mouse). CCL5 challenge led to increased peritoneal leukocyte presence compared with that in PBS treated animals for GRK2^{+/-} chimeras ($P < 0.05$) but not for WT animals (Fig. 5C). In-depth analysis of monocyte migration into the peritoneal cavity of GRK2^{+/-} chimeras showed a significant increased migration in CCL5-challenged animals compared with that in PBS-treated animals (Fig. 5D).

GRK2 Deficiency decreases Myeloid Hematopoiesis

The observed effects of decreased GRK2 levels on the myeloid lineage and in particular on granulocytes and macrophages could reflect altered myelopoiesis. CFU assays with bone marrow cells from WT ($n=6$) and GRK2^{+/-} mice ($n=5$) showed that bone marrow from GRK2^{+/-} mice contained fewer CFUs (43.0 ± 4.2 vs. 57.2 ± 2.8 CFUs, $P=0.017$; Fig. 6A). This result was solely caused by a decreased number of G-CFUs in GRK2^{+/-} chimeras (21.4 ± 1.0 vs. 33.2 ± 2.1 G-CFUs, $P < 0.001$; Fig. 6B); GM- and M-CFUs were unaffected. Thus, GRK2^{+/-} bone marrow cells show an attenuated G-colony stimulating factor (CSF) response, explaining the decreased myeloid (monocytes/granulocytes) over lymphoid (B cells/T cells/NK cells) cell ratio in the circulation (Fig. 6C).

Figure 6 - Myeloid progenitor cells are decreased in GRK2^{+/-} bone marrow; which is associated with reduced blood monocyte and granulocyte numbers.

Colony forming unit (CFU) assays were performed on bone marrow from wild type and GRK2^{+/-} donors. **(A)** The total number of colonies in WT control compared to heterozygous GRK2 chimeras. Based on morphological analysis Granulocyte/Macrophage-CFU (GM-CFU), Macrophage-CFU (M-CFU), and Granulocyte-CFU (G-CFU) were differentiated **(C)** Myeloid (granulocytes + monocytes) over lymphoid (B-cells + T-cells) ratio in the circulation of GRK2^{+/-} deficient mice compared to controls. **(D)** Representative pictures showing Macrophage-CFU (M-CFU), Granulocyte-CFU (G-CFU), and Granulocyte/Macrophage-CFU (GM-CFU) colonies (100x magnification). Wild type controls (white bars) and GRK2^{+/-} chimeras (black bars). * $p < 0.05$; *** $p < 0.001$

Granulocyte and Macrophage GRK2 Deficiency does not explain Phenotype

Because our results pointed to GRK2-deficient myeloid cells as the underlying cause for the phenotypic changes observed in the atherosclerotic lesion in GRK2^{-/-} chimeras, we studied the effect of granulocyte-and macrophage-specific GRK2 deficiency in a *LysM-Cre GRK2^{flox/flox}* bone marrow transplant model in *LDLr^{-/-}* mice ($n=8/9$). To our surprise, no difference was observed in total plaque area between *LysM-Cre GRK2^{flox/flox}* and WT chimeras (Fig. 7A). In addition, necrotic core size was unchanged in these myeloid cell-specific GRK2-deficient chimeras compared with that in controls (Fig. 7B), as were plaque collagen levels (Fig. 7C). Because the *LysM-Cre GRK2^{flox/flox}* model results in lowered macrophage GRK2 levels, we quantified intimal macrophage content in the aortic root lesions, revealing no differences between groups (Fig. 7D).

Figure 7 - Atherosclerotic plaque development and intimal macrophage accumulation are unaffected in LysM-Cre GRK2^{flox/flox} chimeras.

(A) Plaque area and (B) necrotic core size were determined based on haematoxylin and eosin (H&E) staining. (C) Collagen content was measured by analysis of Sirius Red staining under polarized light. (D) Macrophage content was measured in sections stained for the monocyte/macrophage marker MAC-3. (E) Representative pictures of H&E, Movat's Pentachrome, and MAC-3 staining are shown for wild type and LysM Cre GRK2^{flox/flox} chimeras. The top panels represent WT and the bottom panels LysM Cre GRK2^{flox/flox} chimeras. Wild type controls (WT) are shown in white and LysM Cre GRK2^{flox/flox} chimeras are shown in grey; scale bars are 100 μm.

In analogy with the *GRK2*^{-/-} chimera study, flow cytometry of peripheral blood showed marked changes in granulocytes and Ly6C monocyte subsets in the granulocyte- and macrophage-specific GRK2-deficient chimeras (Table 1). These differences did not translate to plaque phenotypic effects. It appears that granulocyte- and macrophage-specific GRK2 deficiency does not affect atherogenesis, despite perturbed leukocyte patterns in circulation. The involvement of GRK2-deficient leukocytes, other than monocytes, cannot be excluded. However, total monocyte counts were not affected in *LysM-Cre GRK2*^{flox/flox} chimeras but were decreased in *GRK2*^{-/-} chimeras, indicating that monocytes, rather than granulocytes or macrophages, were probably the cell type responsible for the atheroprotective phenotype in hematopoietic GRK2 deficiency.

Table 1 - Blood leukocyte composition is affected in macrophage/granulocyte specific GRK2 deficiency.

Leukocyte populations were determined in the blood from wild type (WT) and LysM-Cre GRK2^{flox/flox} chimeras. Numbers shown are the relative percentages of living cells, except for CD4 and CD8 (% of CD3e⁺ cells) and Ly6C high/low monocytes (% of monocytes). * p<0.05 compared to WT.

	WT	LysM-Cre GRK2f/f
% T-cells	20.2 ± 1.3	18.6 ± 0.9
% CD4+ T-cells	67.9 ± 0.5	69.4 ± 0.6
% CD8+ T-cells	25.7 ± 0.6	24.7 ± 0.2
% B-cells	22.1 ± 2	31 ± 1.3 *
% Granulocytes	29.4 ± 2	24.5 ± 0.8 *
% Monocytes	16.1 ± 1.6	15.8 ± 0.5
% Ly6C high monocytes	56.5 ± 1.3	46.2 ± 1.2 *
% Ly6C low monocytes	33.8 ± 1.9	44.2 ± 0.9 *

Discussion

A role of GRK2 in atherosclerosis has not yet been documented but could be inferred from its key regulatory role in CCL2 (monocyte chemotactic protein-1) and CCL5 (RANTES) function, which are instrumental in leukocyte chemotaxis and transmigration during atherogenesis^{20,31,38}. Moreover, GRK2 levels were shown to be decreased in rheumatoid arthritis, an inflammatory disease sharing many features with atherosclerosis. Further, GRK2 expression was shown to be responsive to atherogenic cytokines such as IFN- γ and IL-6^{ref 25,39}. Our data identify a role for hematopoietic GRK2 in atherogenesis, excluding GRK2 deficiency in granulocytes and macrophages being responsible for the observed effects.

First, we mapped the effect of heterozygous hematopoietic GRK2 deficiency on atherosclerosis in *LDL*^{-/-} mice. The donor mice were on a C57BL/6 background but were *LDLr*^{-/-}. *LDLr* expression in hematopoietic cells did not affect either repopulation of the hematopoietic lineage or atherosclerosis development and is widely accepted as the donor genotype in atherosclerosis research⁴⁰. The *GRK2*^{-/-} genotype results in an approximately 50% reduction in GRK2 protein levels as shown before⁴¹. Homozygous GRK2-deficient mice could not be used in these experiments because GRK2 deficiency is embryonically lethal, and embryos do not survive beyond gestational day 15.5. Most likely the *in utero* mortality is caused by cardiac dysfunction³³. Partial GRK2 deficiency resulted in reduced plaque burden and delayed plaque progression. In fact, *GRK2*^{-/-} plaques did not progress beyond a fatty streak stage and contained more macrophages and intimal smooth muscle cells but

showed profoundly diminished necrotic core formation. The latter pointed to a direct effect of GRK2 deficiency on macrophage phagocytosis or apoptosis. Indeed, GRK2 was previously shown to influence cytoskeleton function^{42,43}, thereby potentially modifying macrophage phagocytic capacity. As shown previously, disrupted apoptotic cell clearance can accelerate atherosclerosis in mice⁴⁴. However, analysis did not support the notion that the *GRK2*^{-/-}-associated reduction in necrotic core size is due to increased phagocytosis, because apoptotic cell handling by macrophages was not affected.

Impaired necrotic core expansion is not caused by altered apoptosis of plaque macrophages. However, *GRK2*^{-/-} macrophage necrosis *in vitro* was sharply attenuated after oxLDL exposure. The actual effect of macrophage death on atherosclerosis is still a matter of debate. Several studies have shown that in early and intermediate plaques, enhanced macrophage apoptosis is beneficial^{45,46}, albeit others suggested otherwise⁴⁷. In advanced atherosclerosis, the consensus is that macrophage death promotes necrotic core expansion and plaque destabilization^{48,49}.

Enhanced plaque macrophage content and reduced lesion burden in *GRK2*^{-/-} chimeras could reflect changes in peripheral inflammation. *GRK2*^{-/-} chimeras showed decreased monocyte counts. Total leukocyte numbers were increased, an effect that was mainly attributable to augmented granulocyte counts in circulation. Myelopoiesis, assessed by CFU assay, was attenuated in *GRK2*^{-/-} bone marrow, and G-CFUs, in particular, were decreased. Although it remains to be established how GRK2 exactly interferes with G-CSF response at a molecular level, this process may be linked to granulocyte release or life span because granulocyte numbers also increased in the bone marrow. At a functional level, mobilization of macrophages to inflammatory sites seemed to be increased with partial GRK2 deficiency, which is compatible with the desensitizing effect of GRK2 on chemokine receptor activity⁵⁰. Indeed, we could show that CCR5 chemokine receptor expression was increased on peripheral monocytes from *GRK2*^{-/-} chimeras. At a functional level, this increase was accompanied by increased monocyte migration in response to a CCL5 (RANTES) challenge, as shown in a model of chemokine-induced peritonitis. However, the roles of other cell types such as T lymphocytes, dendritic cells, and macrophages cannot be excluded, and they might contribute to the increased monocyte migration indirectly, because relevant,

potentially GRK2-responsive chemokine receptors (CCR1, CCR2, and CCR7) are also expressed by lymphocyte subsets^{51,52}.

The involvement of the chemokine receptor and ligand axis in regulating stromal release of Ly6C^{high} monocytes from the bone marrow has already been shown for the CCL2/CCR2 axis⁵³. However, the decreased Ly6C^{high} monocyte levels in circulation and spleen are not likely to be due to GRK2-dependent tuning of CCR2 egress because in that case increased monocyte levels would be expected. We did, however, observe that mobilization of monocytes is affected at some level because there are no differences observed in the bone marrow. On the other hand, mice deficient for CCL2 or CX3CR1 had decreased atherogenesis. Inhibition of the CCL5/CCR5 axis by treatment with a CCR5 antagonist (Met-CCL5) resulted in an even more dramatic reduction of atherogenesis⁵⁴. A similar reduction in atherosclerosis development was observed in this study, indicating that GRK2-dependent processes other than receptor desensitization may be responsible for the phenotype observed.

Major effects of GRK2 deficiency on myeloid differentiation and mobilization have prompted us to address the contribution of granulocyte and macrophage GRK2 deficiency to the atheroprotection. The *LysM-Cre GRK2^{flox/flox}* transgenic mouse was previously shown to confer effective deletion efficiency in macrophages and granulocytes^{55,56}. Contrary to our expectations, the *LysM-Cre GRK2^{flox/flox}* bone marrow chimeras did not mirror the previously observed *GRK2^{-/-}* phenotype. In fact, *LysM-Cre GRK2^{flox/flox}* chimeras completely lacked a plaque phenotype, disqualifying granulocyte and macrophage GRK2 deficiency as the major culprit, although the possibility that the phenotype of homozygous GRK2 deficiency differs from that of *GRK2^{-/-}* cannot be excluded. On the other hand, in sepsis, it was recently shown that macrophage- and granulocyte-specific deficiency resulted in no clear phenotype attributable to these cells⁵⁷. Flow cytometry of the *LysM-Cre GRK2^{flox/flox}* chimeras and controls revealed a reduction in relative granulocyte abundance in blood of the *GRK2^{flox/flox}* animals and in keeping the granulocyte- and macrophage-specific deletion, monocyte levels in circulation were not affected in the *LysM-Cre GRK2^{flox/flox}* model. It has previously been shown that the LysM expression in circulating monocytes is not sufficient to get the CRE levels necessary for effective deletion of the floxed gene, in this case GRK2^{ref 58}. In contrast to this ineffective deletion in F4/80-negative immature macrophages or monocytes, the construct is highly efficient in the deletion of the floxed genes in F4/80-

positive macrophages and in granulocytes^{59,60}. Taken together, these data point to monocytes or potentially even T cells (albeit we did not observe overt effects of GRK2 deficiency on the number of circulating or plaque T cells) as responsible factors in the profound *GRK2*^{-/-} plaque phenotype.

Overall, our data indicate an equally complex and critical role of GRK2 in monocyte homeostasis and mobilization in the context of atherosclerosis. Partial hematopoietic GRK2 deficiency appeared to have a profound effect on atherogenesis. Whereas all data seem to pinpoint the myeloid lineage as the major effectors, reduced GRK2 levels in granulocytes and macrophages cannot be held directly accountable for this effect. Cumulatively, these data lead us to propose a major role for monocyte GRK2 in the observed phenotypic changes. It should be noted, however, that although we did not observe any major effects of *GRK2*^{-/-} on other leukocyte subsets known to affect atherogenesis, such as T lymphocytes and dendritic cells, the possibility that these subsets may nevertheless have modified the atherogenic response in an indirect manner cannot be excluded. Because even partial inhibition of GRK2 function already suffices to halt plaque progression at a fatty streak stage, our studies warrant further investigation into the perspective of GRK2 antagonists in atherosclerosis treatment and in determining the cell type responsible for effects on atherosclerosis development.

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Supplemental Figures

Supplemental figure 1 – Body weight development and cholesterol levels are unaffected in GRK2^{+/-} chimeras.

(A) Body weight was monitored during the course of the experiment for WT (○) and GRK2^{+/-} (●) chimeras. (B) Cholesterol levels were determined at baseline (week 0), week 9, and prior to sacrifice (week 18), for WT (white bars) and GRK2^{+/-} (black bars) chimeras.

Supplemental figure 2 – Morphological scoring of atherosclerotic lesions.

Atherosclerotic lesions from WT and GRK2^{+/-} chimeras were scored for (A) necrosis, (B) foam cell content, (C) neutrophil content, (D) cell adhesion, and (E) collagen by an independent operator based on Movat's Pentachrome staining. (F) Representative pictures for Movat's Pentachrome staining. WT controls (white bars) and GRK2^{+/-} chimeras (black bars); scale bars are 100 μm. * $p < 0.05$, ** $p < 0.01$

Supplemental figure 3 – Splenic leukocyte subsets.

Splenic (A) granulocytes, (B) B-cells, (C) T-cells, (D) monocytes, (E) Ly6C^{high} monocytes, and (F) Ly6C^{low} monocytes were determined in spleens from GRK2^{+/-} chimeras and WT controls. WT controls (white bars) and GRK2^{+/-} chimeras (black bars). * $p < 0.05$

Supplemental figure 4 – Gating strategies for flow cytometry analyses.
(A) Monocytes/granulocyte gating in blood and spleen. (B) T-cell and B-cell gating in blood and spleen. (C) Resident dendritic cells (rDC) gating in spleen. (D) Chemokine receptor gating.

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Chapter 04

Hypercholesterolemia exposed circulating monocytes ingest lipids and induce pro-inflammatory pathways

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Abstract

Increased cholesterol levels are a known risk factors for cardiovascular disease. Plaque macrophages accumulate oxidized cholesterol present in the atherosclerotic lesion. Despite the extensively described role of cholesterol in atherogenesis, little is known about the interaction between plasma cholesterol and circulating monocytes. Nowhere we analyzed not only the accumulation of lipoproteins and lipids in peripheral monocytes, but we also studied the effects of intracellular cholesterol accumulation on monocyte function to unravel processes in circulating monocytes contributing to atherogenesis. *LDLr^{-/-}* mice (n = 8) were fed a Western type diet (WTD) for up to three weeks to induce hypercholesterolemia and study the effects on early atherogenesis. After 0, 0.5, 1, and 3 weeks of WTD diet animals were sacrificed and cholesterol accumulation was determined. Flow cytometry data showed increased granularity, which was caused by intracellular cholesterol/lipid accumulation as determined by electron microscopy and Oil-red-O staining. Plasma cytokine levels (IL-10, IL12p70, TNF α , mKC, IL-6, and IL-1 β) were increased, with the most pronounced increase occurring in the first week of WTD. Increased granularity, and thus cholesterol accumulation, resulted in enhanced ROS production and increased monocyte rolling along and adherence to activated endothelium in mice on WTD. Taken together, we show that hypercholesterolemia induces lipid accumulation in circulating monocytes, thereby increasing both ROS production in these monocytes as well as stimulating monocyte-endothelium interactions. Overall, our data indicates that hypercholesterolemia in early atherosclerosis affects circulating monocytes and might contribute to unraveling novel pathways for intimal cholesterol accumulation through circulating monocytes.

Introduction

Hypercholesterolemia is a well-known risk factor for the development of cardiovascular disease (CVD) and more in particular atherosclerosis, the main cause of CVD ^{1,2}. Indeed, increased levels of low density lipoprotein (LDL) cholesterol have been associated not only with increased CVD risk, but also plaque burden ³. Classically, LDL is thought to permeate into the atherosclerotic plaque and after modification into oxidized LDL (oxLDL). Cells from the myeloid lineage, in particular monocyte-derived macrophages, will scavenge the oxLDL present in the lesion, identifying infiltrating monocytes and plaque macrophages as a major player in atherosclerosis.

Monocytes can be divided into two distinct subsets based on the expression of specific surface markers. In humans inflammatory CD14^{high}/CD16⁻ and patrolling CD14^{low}/CD16⁺⁺ monocytes have been described ⁴. In mice monocytes are separated based on Ly6C expression into inflammatory Ly6C^{high} and patrolling Ly6C⁻ monocytes ^{5,6}. CC-motif chemokine receptor 2 (CCR2) is expressed at high levels by inflammatory monocytes ⁷, whereas CX₃C-motif chemokine receptor 1 (CX₃CR1), or fractalkine receptor, is more abundant on patrolling monocytes ^{8,9}. Inflammatory monocytes are generally thought to be the most important subtype in early atherogenesis, being efficiently recruited to the lesion in a CCR2 dependent manner ¹⁰. However experimental loss-of-function studies as well as expression studies were supportive of a role for patrolling monocytes as well, at later stages of disease progression ¹¹. Nevertheless, the exact role of both subsets in plaque macrophage accumulation and plaque progression has not yet been elucidated completely.

Accumulation of cholesterol in macrophages is mediated by scavenger receptors such as SR-A and CD36 ^{12,13}. In atherosclerosis, the efflux of oxLDL from macrophages via ATP binding cassette (ABC) transporters, ABCA1 and ABCG1, is hampered causing progressive intracellular buildup of free cholesterol and eventually cholesterol crystals ^{14,15}. Both intracellular cholesterol, as well as free cholesterol in the lesion, can trigger several pro-inflammatory pathways, contributing to the progression of early atherosclerotic lesions. Main downstream signaling pathways triggered upon (ox)LDL dependent macrophage activation are NF-κβ signaling via toll-like receptor (TLR) 4 and TLR 6 ¹⁶, ERK signaling via TLR2/TLR4, and ER stress ¹⁷⁻¹⁹. Besides the effects described for oxLDL, also cholesterol crystals can trigger inflammatory signaling via the inflammasome through lysosomal damage ^{20,21}.

LDL cholesterol can also affect arterial endothelial cell function, upregulating integrins like intercellular adhesion molecules (ICAMs)²² and vascular cell adhesion molecule (VCAM)¹²³ as well as inducing monocyte chemotactic molecule secretion (e.g. CXCL1 and CCL2)^{19,24,25}. Although ample evidence points towards increased monocyte adhesion and transmigration under hypercholesterolemic conditions^{26,27}, this effect was generally attributed to endothelial activation, not to direct impact of hyperlipidemia on monocytes.

Nevertheless, hypercholesterolemia, in combination with reduced cholesterol efflux, is known to induce monocytosis in animal models such as LDL receptor deficient (*LDLr*^{-/-}) and apolipoprotein E deficient (*ApoE*^{-/-}) mice, but also in patients with familial hypercholesterolemia (FH)²⁸, or non-genetic hypercholesterolemia²⁹. This phenomenon could be attributable to intracellular cholesterol accumulation induced lipid raft formation, stimulating proliferation of myeloid progenitors via IL-3 and granulocyte/macrophage-colony stimulating factor (GM-CSF) signaling^{30,31}. In the current study we investigated implications of acute hypercholesterolemia on differentiation, function, and dynamics of circulating monocytic cells. Pro-atherogenic (*LDLr*^{-/-}) mice were monitored after onset of Western type diet (WTD) feeding and monocyte subsets were analyzed. Our study shows that circulating monocytes rapidly accumulate cholesterol in situ, resulting in diet-induced functional aberrations.

Materials & Methods

Animals

Male low density *LDLr*^{-/-} mice were obtained from the animal breeding facility at Maastricht University (Maastricht, The Netherlands). Mice were fed a regular chow diet (RM3; Special Diet Services [SDS], Essex, UK) or a Western-type diet (WTD; 0.25% cholesterol, 15% cocoa butter, 1% corn oil, 40.5% sucrose, 10% corn starch, and 5.95% cellulose; SDS). Drinking water and food were provided *ad libitum*. Experiments were performed at the Maastricht University animal facility. All experimental protocols were approved by the Maastricht University ethics committee for animal experiments.

***In vivo* monocyte-endothelial cell interaction measurement**

Monocyte-endothelial cell interactions were measured in male *LDLr^{-/-}* mice either on normal chow (n = 3) or WTD (n = 3). Alexa568-coupled anti-CD115 antibody (5 µg/mouse, eBioscience) and FITC-conjugated anti-CD31 antibody (35 µg/mouse, BD Bioscience) were administered intravenously. Immediately hereafter, mice were anesthetized by a dorsal subcutaneous injection with ketamine (100 mg/kg body weight, Nematek) and xylazine (10 mg/kg body weight, Sedamun). Subsequently, mice were prepared for multiphoton laser scanning microscopy (MPLSM) by exposing the sternohyoid muscle by dissection. MPLSM imaging was performed on a Leica SP5 imaging platform (Leica Microsystems, Germany) that integrates multiphoton microscopy with fast resonant scanning, and uses a Compact Ultrafast Ti:Sapphire Laser (Chameleon, Coherent, USA). An excitation wavelength of 820 nm was used in all experiments. Tissue was observed using a 20x NA 1.0 water immersion objective (HCX PL APO L, Leica Microsystems) with a numerical aperture of 1.00 and an integrated optical zoom mechanism allowing magnification up to 60x. Photo-multiplier tubes (PMT) were used to detect three spectral regions: PMT1: 500 - 550; PMT2: 565 - 605. *In vivo* MPLSM imaging was performed at a frame size of 400 x 400 pixels (pixel size: 1.0882 x 1.0882 µm), scanning at 8000 Hz for fast recording (20 frames/sec) and using a line average of 2 (10 frames/sec) for more detailed scans. Per time series a total of 500 frames was recorded. After *in vivo* imaging, the animal was sacrificed and additional *in situ* imaging was performed to examine the vasculature at higher resolution and an increased signal-to-noise ratio without motion artifacts. *In situ* imaging was performed at a frame size of 512 x 512 pixels (pixel size: 0.847 x 0.847 µm), scanning at 200 Hz, and an interplanar distance of 1.00 µm.

Flow cytometry

Blood, spleen, and bone marrow were collected upon sacrifice. Absolute leukocyte counts in blood were obtained using BD Trucount tubes, according to the manufacturers protocol (BD Biosciences). In short, Fc-receptor blocking antibody was added to the Trucount tubes. Subsequently, 50 µL anti-coagulated whole blood was added and the tube was gently vortexed. After incubation for ten minutes, the antibody cocktail (supplemental table 1) was added and the suspension was incubated at room temperature in the dark for

an additional 20 minutes. Finally, hypotonic lysis buffer was added and after 15 minutes samples were analyzed.

Myeloid progenitor densities were measured in spleen and bone marrow. Single cell suspensions were made from spleen by crushing the tissue over a 70 μm cell strainer (BD Biosciences, NJ, USA). One femur and one tibia per mouse were flushed with ice-cold PBS and single cell suspensions were prepared using a 70 μm cell strainer (BD). subsequently, spleen and bone marrow cells were stained with biotin-conjugated antibodies against lineage (Lin) markers (CD5, CD45RA [B220], CD11b, Gr-1 [Ly6C/G], 7-4, and Ter-119; Miltenyi Biotec) and Lin⁺ positive cells were depleted using streptavidin conjugated magnetic beads (Miltenyi Biotec) and LS columns (Miltenyi Biotec). Lin⁻ cell pools were stained with antibodies for Sca-1, c-Kit (CD117), CD34, CD16/32, CD115, and lineage markers (CD5, CD45RA [B220], CD11b, Ly6C, Ly6G, Ter-119).

Mature leukocytes in spleen and bone marrow were analyzed after initial removal of erythrocytes by incubation with hypotonic lysis buffer (8.4 g NH_4Cl and 0.84 g NaHCO_3 per liter distilled water). Non-specific Fc-receptor binding was blocked by the addition of anti-CD16/CD32 antibody (eBioscience, San Diego, USA). Subsequently, single cell pools were stained with antibody mix (supplemental table 1).

Intracellular interleukin-1 production was measured in blood leukocyte subsets. After lysis of erythrocytes, leukocyte pools were incubated for 4 hours in RPMI-1640 medium (supplemented with 10% fetal calf serum and 0.1% GolgiPlug [BD]) to induce intracellular cytokine accumulation. Cells were resuspended in FACS buffer and after FC-receptor block, surface markers were labeled with antibodies against CD3 ϵ , CD45RA (B220), NK1.1, CD11b, Ly6G, and Ly6C. Subsequently, cells were fixed using Fix/Perm solution (eBioscience). Next, cells were permeabilized and stained with interleukin (IL) 1 α , IL-1 β , and pro-IL-1 antibodies.

Intracellular reactive oxygen species accumulation in blood and splenic leukocyte subsets were determined by staining of cells with 100 μL 20 μM DCFDA in PBS and incubated for 20 minutes at 37°C. Cells were washed once in PBS and resuspended in FACS buffer. After Fc-receptor block, surface markers were labeled with antibodies against CD3 ϵ , CD45RA (B220), NK1.1, CD11b, Ly6G, and Ly6C and analyzed.

All samples and buffers were kept on ice throughout the experiment unless indicated otherwise. All measurements were performed on a FACS Canto II (BD Biosciences) and analysis of acquired data was performed using FACS Diva software (BD Biosciences).

FACS Sorting

Upon sacrifice blood and spleen were collected and single cell suspensions were prepared. Fc-receptor blocking antibody was added to the cell suspension. Subsequently, antibodies against CD3 ϵ , CD45RA (B220), NK1.1, CD11b, Ly6G, and Ly6C were added and incubated at 4°C in the dark for 30 minutes. Finally, monocyte Ly6C subsets were sorted in RPMI-1640 medium supplemented with 10% FCS on a FACS ARIA flow cytometer (BD Biosciences).

Cytokine multiplex

Plasma samples were frozen and stored at -80°C until further use. Samples were thawed and selected cytokines mKC (CXCL1), tumor necrosis factor (TNF) α , IL-1 β , IL-6, IL-10, IL-12p70, and interferon (IFN) γ were measured using Multi-array electrochemiluminescence platform (K15012A-5, detection range 2.4 pg/ml to 10,000 pg/ml; MesoScaleDiscovery, Gaithersburg, MD, USA). Analysis was performed as described by Van Bussel *et al*³². Each sample was analyzed in duplicate on the same array plate.

Electron microscopy

Monocytes were FACS sorted and fixed in 3% glutaraldehyde (GTA; Ted Pella, Redding, CA, USA) in 0.1M phosphate buffer (pH = 7.6) for 15 minutes. After several washes, cells were post-fixed in 1% osmium tetroxide solution and routinely dehydrated through 100% ethanol, cleared with propylene oxide, and embedded in epoxy resin. Ultra-thin sections (70 - 90 nm) were cut on an ultra-microtome, mounted on Formvar (1595 E, Merck) coated 75 mesh copper grids, and counterstained with uranyl acetate and lead citrate before analysis on a Philips CM100 transmission electron microscope.

Oil-red-O staining

FACS sorted monocytes were spotted on glass slides using Cyto-Tek centrifuge (Sakura Finetek, Torrance, CA, USA). Cells were fixed in 4% formalin for 10 minutes and subsequently washed twice in PBS. Next, slides were incubated for 15 minutes in 60% isopropanol, followed by 60% Oil-red-O solution for 20

minutes. Cells were washed gently with 60% isopropanol and once with water. Finally, the cells were counterstained with haematoxylin and slides were mounted with aquamount. Pictures were made at 40x magnification (Leica).

Statistical Analysis

Student's *t* test was used; nonparametric data were analyzed using a Mann-Whitney *U* test. All analyses were performed using GraphPad Prism 5 software (GraphPad Software Inc., LA Jolla, CA, USA), and values of $P < 0.05$ were considered statistically significant.

Results

Monocyte intra-cellular complexity is increased upon WTD

Flow cytometry analysis of whole blood from mice on WTD showed that the monocyte fraction displayed markedly increased side scatter (SSC) compared to chow fed control animals as early as half a week of WTD feeding (figure 1A). As expected, plasma cholesterol levels progressively increased upon onset of WTD feeding to plateau at 3 weeks (figure 1B). Of note, the average monocyte SSC correlated to the plasma cholesterol levels ($R^2 = 0.726$; $p < 0.0001$; data not shown).

Figure 1 - Monocyte granularity correlates to plasma cholesterol levels

(A) Representative dot plots indicating increased side scatter profile of monocytes (orange and blue) from $LDLr^{-/-}$ mice on normal chow or 3 weeks Western type diet (WTD). (B) Plasma cholesterol levels measured at baseline (normal chow), and after 0.5, 1, and 3 weeks of WTD feeding. ** $p < 0.01$ and *** $p < 0.001$

Next, we investigated whether WTD induced SSC changes manifested in both inflammatory Ly6C^{high} and patrolling Ly6C^{low} monocytes. Interestingly, the enhanced scattering was predominantly observed in Ly6C^{low} monocytes (figure 2A+B). Inflammatory Ly6C^{high} monocytes displayed significantly increased side scattering (granularity), albeit much less pronounced, than that of Ly6C^{low} monocytes (figure 2A+B).

To pinpoint the cause of this increased monocyte granularity, Ly6C^{low} and Ly6C^{high} monocytes were sorted and electron microscopic (EM) images were made. EM analysis indicated the overt presence of lipid droplets and even cholesterol crystals in circulating monocytes from WTD fed animals, but not chow fed animals (figure 2C). Additional, Oil-red-O staining confirmed that monocytes, both Ly6C^{low} and Ly6C^{high}, contained lipid droplets (figure 2C).

Hypercholesterolemia induces leukocytosis

Total leukocyte levels were up by almost 45% in WTD compared to chow fed mice (figure 3A). This increment could be ascribed to elevated granulocyte (figure 3B) and monocyte levels (figure 3C), which in fact more than doubled in WTD fed animals, confirmative of previous data on hypercholesterolemia induced monocytosis^{30,31}. Furthermore, the Ly6C^{high} to Ly6C^{low} ratio was doubled after 3 weeks of WTD (figure 3D+E). Marked leukocytosis and increased monocyte granularity were also observed in spleen (supplemental figure 1A-E) and bone marrow (supplemental figure 2A-D) of *LDLr*^{-/-} animals after 3 weeks of WTD. Despite a small reduction in LK-cells, no significant effects could be observed on erythrocyte/megakaryocyte progenitors (EMP), common myeloid progenitors (CMP), or granulocyte/macrophage progenitors (GMP) after 3 weeks WTD feeding compared to normal chow (supplemental figure 2E+F).

Figure 2 - Lipid accumulation is responsible for increased monocyte granularity
(A) Monocyte granularity was determined for Ly6C⁻ and Ly6C^{high} monocytes at baseline (normal chow, and after 0.5, 1, and 3 weeks WTD feeding). **(B)** Representative dot plots indicative of Ly6C⁻ and Ly6C^{high} monocyte granularity. Circles depict outline of populations under chow conditions. **(C)** Representative electron microscopy (EM) images for Ly6C⁻ (left column) and Ly6C^{high} (right column) monocytes on normal chow (top row) or WTD (middle row). Bottom panels show representative images for Oil-Red-O staining.

Figure 3 - Circulating leukocyte levels are increased upon cholesterolemia

Absolute leukocyte levels were measured in blood upon sacrifice using BD TruCount tubes. (A) Total CD45⁺ leukocytes, (B) CD11b⁺/Ly6G⁺ granulocytes, (C) CD11b⁺/Ly6G⁻ monocytes, and (D) Ly6C monocyte subsets were measured. (E) Representative histograms indicating Ly6C⁻ and Ly6C^{high} monocytes in chow fed and WTD fed animals. * $p < 0.05$ and *** $p < 0.001$

Cytokine profile indicates systemic diet-induced inflammatory profile

Plasma cytokine levels increased rapidly after the onset of WTD. After three to seven days, cytokine levels of IL-10, IL-12p70, IFN- γ , IL-1 β , mKC, and IL-6 on average doubled compared to baseline values (normal chow) (figure 4A-F). TNF plasma levels were either not detectable or just above the detection limit. While elevated IL-10, IL12p70, and IFN- γ levels, and to a lesser extent IL-6, persisted for 3 weeks (figure 4A-D). Plasma levels of the pro-inflammatory cytokines IL-1 β and mKC almost reverted to baseline after 3 weeks of WTD feeding (figure 4E+F).

Figure 4 - Cytokine levels increase in response to hypercholesterolemia

Plasma cytokine levels were measured for (A) IL-10, (B) IL-12p70, (C) IFN- γ , (D) IL-6 (E) IL-1 β , and (F) mKC at baseline (normal chow; open circles) and after 0.5 week (pale blue), 1 week (light blue), and 3 weeks (dark blue) of WTD. * $p < 0.05$ ** $p < 0.01$ and *** $p < 0.001$

Intra-cellular lipoproteins trigger ROS production

In macrophages intra-cellular lipid accumulation was previously reported to trigger the production of reactive oxygen species in an ER stress and ERK-signaling dependent manner¹⁹. Since, we observed similar lipid accumulation in circulating monocytes from WTD fed *LDLr*^{-/-} mice, we analyzed ROS production in these monocytes. Indeed, both high granular Ly6C⁻ and Ly6C^{high} monocytes had significantly increased levels of ROS compared to their low granular counterparts (figure 5A+B). The amount of intra-cellular lipids did not appear to affect the ROS production as the levels in Ly6C⁻ and Ly6C^{high} high granular monocytes was comparable. As Ly6C^{high} monocytes showed less accumulation of intra-cellular lipids this subset conceivably is more susceptible to lipid associated ROS formation.

Figure 5 - Reactive oxygen species are augmented in all high granular monocytes, while IL-1 β levels are only affected in high granular Ly6C⁻ monocytes

Reactive oxygen species (ROS) were measured in (A) Ly6C⁻ and (B) Ly6C^{high} monocytes for both the low granular and high granular fractions, based on normal chow gating control. In the dot plots the gating for the low and high granular monocyte subsets are indicated. IL-1 β levels were measured in (C) Ly6C⁻ and (D) Ly6C^{high} monocytes for both the low granular and high granular fractions. * $p < 0.05$ ** $p < 0.01$ and *** $p < 0.001$

As shown by Latz *et al*, macrophage cholesterol accumulation induces inflammasome activation. We therefore assessed intracellular production of the inflammasome cytokine IL-1 β in monocytes from mice on WTD for 3 weeks. High granular Ly6C⁺ monocytes had significantly increased IL-1 β levels compared to low granular counterparts (figure 5C). In contrast, Ly6C^{high} monocytes did not show any difference in intra-cellular IL-1 β levels between low and high granular fractions (figure 5D). Unfortunately, we were unable to reliably determine intra-cellular pro-IL-1 levels. Overall, increased granularity, and as such intracellular lipids, did only mildly affect IL-1 β levels in Ly6C⁺ monocytes, suggesting that monocyte lipid ingestion is not or only marginally associated with inflammasome activation.

Hypercholesterolemia increases monocyte-endothelial cell interactions

Increased plasma cholesterol levels were shown to induce inflammatory cytokine levels already after 1 week of WTD feeding. Further, ROS have been implicated to increase monocyte-endothelial cell adhesion *in vitro*^{33,34}. To address the combined effect of lipid accumulation, pro-inflammatory milieu, and ROS production on monocyte function we performed *in vivo* monocyte-endothelium adhesion experiments. At baseline, no CD115⁺ cells adhered to the endothelium of the vessels in the sternohyoid muscle in either mice on normal chow or 1 week WTD (data not shown).

After activation of the local vasculature with 50 μ L LPS (100 ng/mL) a clear increase in rolling and adhering CD115⁺ monocytes could be observed compared to baseline condition, where only an occasional rolling or adhering cells could be observed. CD115⁺ cells were considered rolling if their speed was reduced by at least two-fold and showed direct contact to the endothelial layer, adhering cells did not change position for at least 280 frames (30 seconds). Due to the small number of animals ($n = 3$ per group) examined no statistical analyses were performed. The increase in monocyte-endothelial cell interaction was most pronounced in animals on WTD. Animals on WTD showed increased adhering (figure 6A) and rolling CD115⁺ cells (figure 6B). Compared to baseline both animals on normal chow and WTD displayed more adhering and rolling cells, yet this increment was more pronounced in animals on WTD (figure 6C).

Figure 6 - *In vivo* monocyte-endothelial cell interaction is stimulated by intracellular cholesterol accumulation

Monocyte interaction with the endothelial layer for $LDLr^{-/-}$ animals was measured *in vivo* in the sternohyoid muscle after activating the endothelium with 100 ng/mL LPS. **(A)** Number of steadily adhering cells (> 30 seconds) was determined and corrected for the area of vascular endothelium examined. **(B)** Number of rolling cells along the vascular endothelium was determined and corrected for the total time of imaging. **(C)** Representative 3D reconstructions of the vasculature for animals on normal chow (top panel) and WTD (bottom panel) are shown. Endothelial cells are labeled with anti-CD31-FITC (green) and monocytes are labeled with anti-CD115-Alexa568 (red).

Discussion

The role of circulating monocytes in the development of atherosclerosis has been extensively studied over the last few years. Monocyte subsets have been reported to play differential roles in atherogenesis. Ly6C^{high} monocytes are widely considered to be the most important progenitor of plaque macrophages^{10,35}, while Ly6C^{low} monocytes are attributed mainly vascular patrolling functions and are thought to give rise to CD11c⁺ dendritic cells in the lesion³⁶⁻³⁸. Recently, Wu *et al* have demonstrated that cholesterol can induce CD11c expression in monocytes, and enhance their adhesion to the endothelium³⁹. This might implicate a more important role for Ly6C^{low} monocytes in atherogenesis. In this study we show that monocytes, both Ly6C^{high} and Ly6C^{low}, respond differentially to the hypercholesterolemia induction.

We demonstrate that circulating, next to splenic and bone marrow, monocytes rapidly accumulate lipids under hypercholesterolemic conditions, before entering the atherosclerotic lesion, an effect accompanied by generally transient elevation in plasma cytokine levels. Moreover we show that hypercholesterolemia not only induces ROS buildup, but also upregulates IL-1 β production by Ly6C^{high} and Ly6C^{low} monocytes, the latter likely without activating the inflammasome pathway. The progressive accumulation of cholesterol in these peripheral monocytes could also induce the pro-inflammatory milieu observed during atherogenesis, however additional studies are necessary to support this notion. We do observe increased interaction of circulating monocytes with (activated) endothelium under hypercholesterolemic conditions.

The initial observation of progressively increased side scattering of circulating monocyte subsets, and in particular the Ly6C^{low} subset, upon WTD feeding of *LDLr*^{-/-} mice, has led us to investigate the cause of increased granularity in more detail. Here, we present three lines of evidence that increased granularity reflects monocytic lipid accumulation: (I) plasma cholesterol correlated strongly to the increased granularity of monocytes; (II) electron microscopy showed the overt presence of intracellular vacuoles; and (III) Oil-red-O staining established the presence of intracellular lipids in monocytes from WTD-fed *LDLr*^{-/-} mice. Interestingly, this effect was more pronounced in Ly6C^{low} monocytes, however the responsible mechanism needs to be determined. As monocyte development is thought to develop from Ly6C^{high}

towards Ly6C⁺ monocytes⁴⁰, the turnover from lipid-laden Ly6C^{high} to Ly6C⁺ monocytes could add to the more distinct lipid accumulation in the downstream Ly6C⁺ monocyte pool. On the other hand, we cannot exclude that passive or active cholesterol/lipid accumulation, but also difference in efflux, between monocyte subsets contribute to the differences observed.

Lipid accumulation in circulating monocytes appears to have many similarities to scavenging of (ox)LDL by plaque macrophages. As lipid accumulation in macrophages activates the inflammasome, due to intracellular cholesterol crystals²⁰, the question arose whether lipid ingestion by circulating monocytes would in analogy to macrophages also here result in inflammasome activation. Despite the presence of cholesterol crystals in circulating monocytes and the transient increase of plasma IL-1 β levels upon diet change, involvement of the inflammasome in circulating monocytes could not be confirmed conclusively. Preliminary data supported that the IL-1 β effects were inflammasome independent, as pro-IL-1 levels seemed to increase, reflective of transcriptional induction rather than inflammasome activation. This was reinforced by unaltered caspase 1 activity in these monocyte subsets (data not shown). Nevertheless, inflammasome-independent IL-1 β has been implicated to be essential for sustaining extra-medullary monocytopoiesis⁴¹.

Reactive oxygen species did increase significantly in high granular monocytes carrying intracellular cholesterol deposits. While in macrophages cholesterol accumulation has already been linked to ROS production, only one single study has investigated cholesterol-dependent ROS production in monocytes⁴². Although the actual molecular pathway linking lipid to ROS accumulation in monocytes remains to be determined, as shown by Huang *et al* indicated that in humans ROS production might be mediated via C-reactive protein expression and the subsequent upregulation of NADPH oxidase (NOX) 2 and MCP-1 levels³³. As increased NOX2 and MCP-1 levels might affect chemotaxis of monocytes, hypercholesterolemia induced ROS production could well have effects on the migratory behavior towards activated endothelium.

Indeed our intravital microscopy studies showed that monocyte adhesion to and rolling along activated endothelium was substantially increased in WTD fed *LDLr*^{-/-} mice. Apart from a ROS associated phenomenon, the increased endothelial cell adhesion could also result from lipid accumulation induced upregulation of integrins on circulating monocytes

^{26,43,44}. However, as we did not determine a broad set of membrane markers, such as ICAM-1, very late antigen (VLA) 4, and CD62L, on these circulatory monocytes, the actual contribution of this pathway to the increased endothelial adhesion remains unclear. Nevertheless, our study clearly shows that hyperlipidemia impacts not only circulating monocyte constitution but also function, already within 1 week of WTD exposure.

To what extent this is relevant to for early atherogenesis is yet to be determined, given that our *in vivo* studies involving LPS activated small vessels rather than major arteries. However, since endothelial cells covering the (early) atherosclerotic lesion do express TLRs ^{45,46}, the LPS stimulation used in our experiments is representative for atherosclerotic endothelium. Especially since TLR4 polymorphisms as well as MyD88 deficiency have been shown to decrease atherogenesis ^{47,48}. Therefore, we expect cholesterol-loaden monocytes to adhere as efficiently to the endothelium covering the atherosclerotic plaques under hypercholesterolemic conditions. Atherosclerosis development might be further stimulated by ROS generation in circulating monocytes, as this contributes to the formation of oxLDL ⁴².

Indeed monocytes in compartments relevant for myeloipoiesis, such as spleen and bone marrow, also showed pronounced lipid accumulation, although at present we cannot exclude that these monocytes had compartmentalized upon lipid ingestion or whether they ingestion had occurred after monocyte grafting in these tissues ^{10,36,41}. Nagareddy *et al* recently showed that monocytosis might in part result from hyperglycemia, next to hypercholesterolemia ⁴⁹. However, this was mediated via increased proliferation and expansion of granulocyte/macrophage progenitors in the bone marrow, which we did not observe in our study.

Monocyte development is often considered to be a linear process, inflammatory (Ly6C^{high}) monocytes produced in the bone marrow migrate towards the spleen and at least a subset will mature towards patrolling (Ly6C⁻) monocytes ¹⁰. In atherosclerosis Ly6C^{high} monocytes appear to be most critical as precursor for plaque macrophages ³⁵, however the role of Ly6C⁻ monocytes is still debated. In human atherosclerosis, the role of inflammatory CD14^{high}CD16⁻ monocytes and patrolling CD14⁻CD16⁺ monocytes is even less defined, as no trafficking studies have been performed in human disease. Nevertheless, especially patrolling Ly6C⁻ monocytes show massive lipid accumulation, suggesting an important role in hypercholesterolemia and

possibly a role in atherosclerosis development, which is more important than currently thought.

Downregulation of the cholesterol transporters ABCA1 and Niemann-Pick disease type C protein 1 (NPC1) was recently identified as the central actor responsible for lipid accumulation in human monocytes in FH patients, with manifest hypercholesterolemia⁵⁰ and in familial coronary artery disease patients⁵¹. ABCA1/G1 deficiency results in hypercholesterolemia-associated monocytosis⁵², however ABCA1/G1 deficiency also results in lipid accumulation in macrophages⁵³, and potentially circulating monocytes. Not only accumulation of lipoprotein fractions in circulating monocytes have been observed before, but also accumulation of fatty acids⁵⁴, in our study we have not yet discriminated between these subsets. In contrast to Swirski *et al*³⁵, the current study has shown that Ly6C⁺ monocytosis might be an additional process contributing to atherosclerosis, at least in *LDLr*^{-/-} mice.

In conclusion, we show that hypercholesterolemia not only results in a rapid, but transient pro-inflammatory adaptive response in plasma. Furthermore, almost immediate intracellular changes in circulating as well as tissue resident monocyte subsets, including lipid accumulation and ROS production, which functionally translate in enhanced monocyte adhesion to activated endothelium. Altogether, our study reveal a novel pathway through which circulating monocytes contribute to atherosclerosis not only by fostering the subendothelial macrophage invasion but also potentially by conveying cholesterol into the atherosclerotic lesion.

Supplemental data

Supplemental table 1 - Overview of FACS antibodies used for these studies

Antibody	Conjugate	Company	Tissue
CD3ε	eFluor450	eBioscience	Blood
CD3ε	PerCP-Cy5.5	BD	Bone marrow
CD4	APC-H7	BD	Blood, bone marrow
CD8α	FITC	eBioscience	Blood
CD8α	eFluor450	eBioscience	Bone marrow
CD8α	APC-H7	BD	Spleen
CD11b	PE-Cy7	BD	Blood, spleen
CD11b	BV510	BD	Bone marrow
CD11b	eFluor450	eBioscience	Spleen
CD11c	PE-Cy7	BD	Bone marrow, spleen
CD45	PerCP	Biolegend	Blood
CD45RA (B220)	PerCP	BD	Bone marrow
CD45RA (B220)	V500	BD	Blood, spleen
CD115	PE	BD	Bone marrow
CD317 (PDCA-1)	PE	BD	Spleen
Ly6C	APC	Miltenyi	Blood, spleen, bone marrow
Ly6G	APC-Cy7	BD	Blood, spleen, bone marrow
MHC II	FITC	BD	Bone marrow, spleen
NK1.1	PE	BD	Blood, spleen
NK1.1	PerCP-Cy5.5	BD	Bone marrow

Supplemental figure 1 - Splenic leukocyte subsets increase upon WTD comparable to circulating leukocytes

Absolute leukocyte levels were measured in spleen upon sacrifice using BD TruCount tubes. (A) Total CD45+ leukocytes, (B) CD11c+ cDC, (C) pDCA1+ pDC, (D) CD11b+/Ly6G+ granulocytes, (E) CD11b+/Ly6G- monocytes, and (F) Ly6C monocyte subsets, and (G) monocyte granularity per subset were measured. * $p < 0.05$, ** $p < 0.01$, and *** $p < 0.001$

Supplemental figure 2 - Bone marrow leukocyte subsets increase upon WTD comparable to circulating leukocytes

Relative leukocyte levels were measured in bone marrow after 3 weeks of normal chow or WTD. (A) CD11b+/Ly6G+ granulocytes, (B) CD11b+/Ly6G- monocytes, (C) monocyte subsets, (D) granularity monocyte subsets, (E) progenitor cell population, and (F) LK subsets were measured. * $p < 0.05$, ** $p < 0.01$, and *** $p < 0.001$

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Chapter 05

Phage display identifies novel circulating and plaque monocyte markers for advanced atherosclerosis

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Abstract

Unstable angina pectoris is generally diagnosed based on clinical parameters, in default of accessible, sensitive, and selective biomarkers. In this study, we sought to identify novel and unstable angina pectoris-specific biomarkers on circulating monocytes, by phage display V_{HH} antibody repertoire screening. Hereto, a dedicated antibody library was generated from plasma cells isolated from llamas, immunized with unstable angina pectoris patient monocytes. Affinity selections on human LPS-activated THP-1 monocytes, afforded several clones with increased affinity to LPS-activated THP-1 monocytes. Subsequent validation of selected clones on CD14 positive monocytes derived from unstable angina pectoris patients versus monocytes derived from healthy volunteers, identified clones which preferentially recognized unstable angina pectoris patient monocytes. Finally, selected clones specifically bound circulating human monocytes (ELISA) as well as monocytic cells in the vulnerable atherosclerotic lesion (immunohistochemistry). In conclusion, we have identified novel monocyte binding phage clones and provide proof of concept for their potential as biomarker in unstable angina pectoris diagnosis and in molecular imaging approaches for vulnerable plaque detection.

Introduction

Acute coronary syndromes continue to be a major cause of morbidity and mortality in the Western society ^{1,2}. The clinical manifestations generally are caused by rupture of an unstable atherosclerotic lesion ³. Nowadays, there is increasing interest in the characteristics of such unstable, rupture-prone lesions, not only from a mechanistic, but also from a diagnostic point of view, as unstable plaque markers are deemed very useful for molecular imaging purposes ^{4,5}. However, many of these proposed markers are non-selective or difficult to monitor using non- or minimally invasive methods ⁶. Growing evidence suggests that information on localized events in the vasculature or even focal processes in the vulnerable atherosclerotic lesion can probably be deduced from information accessible in the periphery ^{5,7,8}. The current study explores the possibilities of employing circulating monocytes as a read-out for diagnostic and potentially prognostic markers for unstable angina pectoris and acute, but also future, cardiovascular events.

Monocytes may be a particular relevant source for novel biomarkers, as monocyte-derived macrophages are instrumental in plaque inflammation, a major risk factor in plaque rupture ⁹. Furthermore, membrane protein expression on circulating leukocytes, and especially monocytes, are highly responsive to external influences, such as pathogens ¹⁰, trauma released factors, and inflammatory disease ^{11,12}. In response to these pathological processes foreign material can be presented on specialized cells. Cell types most famous for presenting antigens on MHC class I or II molecules are dendritic cells ¹³ and macrophages ¹⁴. Moreover, many of the differentially expressed membrane proteins, such as MHC class II, TLR4, and CD14 are associated with functional alterations or adaptations of the monocytes as they trigger or dampen the activity of downstream effectors. Overall, the modifications of surface marker expression on circulating monocytes due to local vascular events are of interest as these may harbor biomarker potential.

Leukocyte membrane markers, or the expression level of these markers, have repeatedly been shown to be affected by localized vascular events. Acute cardiovascular events was seen to result in altered receptor expression profiles on monocytes ^{15,16}. Conceivably, the CD14⁺CD16⁺ subset of monocytes which have a patrolling function, would be among the first to react to vascular injury or trauma. The information on local pathophysiological processes may therefore be directly analyzed from peripherally sampled blood

specimen, or possibly even using (non-invasive) imaging techniques. The latter could be further optimized using novel, highly specific markers for vulnerable atherosclerotic lesions. At the moment knowledge about the relation between monocyte expression profiles and cardiovascular disease are largely lacking.

We therefore opted for an unbiased approach to screen for antibodies that target membrane markers which are differentially expressed between monocytes from healthy controls and patients with unstable angina pectoris. The advantages of phage display derived V_{HH} antibodies are the direct application of the encoded antibody for targeted monocyte-based biomarker assays or imaging approaches. Further, the ability of these antibodies to bind targets non-accessible with conventional antibodies, such as pockets, could reveal targets unknown so far. Hereto, phage display was used as a tool to determine differences between monocyte populations, without any a priori knowledge. Normally, phage display is used to obtain (V_{HH}) antibodies against known targets, preferentially purified or recombinant proteins¹⁷, as well as cellular or plasma derived (purified) proteins^{18,19}. In this study the protocol was adapted to be able to determine novel targets specific for unstable atherosclerotic disease on circulating mononuclear cells²⁰. Screening was performed using an immune library, produced by immunization of a llama with UAP patient-derived circulating monocytes. We now present data on phage/antibody selections on UAP CD14⁺ monocytes specific phage clones, and show that selected clones preferentially bind monocytic cells in the advanced, vulnerable lesion. The high affinity for UAP monocytes indicates prospective potential for use in the clinics as diagnostic marker in blood or for detection of vulnerable plaques by imaging.

Materials & Methods

Subjects

Monocytes (CD14⁺ fraction) isolated from seven patients admitted to the University Medical Center Maastricht with unstable angina pectoris, were included for the analysis performed. One patient was later excluded as clinical parameters indicated no aberrations the ECG, all other patients were confirmed to have unstable angina pectoris by the clinician. For an optimal signal-to-noise ratio compared to non-diseased monocytes we included CD14⁺ monocytes from four healthy volunteers, without any known cardiovascular disease, as reference population. All participants provided written informed consent prior to participation. This study was approved by the local ethics committee.

Monocyte Isolation

Whole blood samples isolated from patients and healthy controls were centrifuged at 156 x g for 15 min without break. Next, plasma was removed and stored for future analysis at -80°C. A leucosep tube (Greiner) was filled with 15 mL Ficoll (Sigma) at room temperature. Tubes were centrifugated for 30 seconds at 1,000 x g and 25 to 30 mL of PBS diluted blood was added on top of the filter and centrifuged for 20 min at 1,000 x g without breaks. Liquid above the peripheral blood mononuclear cells (PBMC) enriched interphase was removed and the PBMC fraction was transferred to a fresh 50 mL tube and washed twice with cell selection buffer (PBS; 0.5% BSA, 2 mM EDTA, 0.09% sodium azide). Cells were resuspended in 1 mL cell selection buffer and a 20 µL aliquot was saved as reference. After centrifugation the pellet was resuspended in 75 µL anti-human CD14 antibody conjugated magnetic beads (BD) and incubated for 60 min on ice. Subsequently, 1 mL cell selection buffer was added and cell suspension was placed on the BD Imagnet (BD) for 10 min. With the tube on the magnet the CD14 negative fraction was aspirated and saved for other analyses. The CD14 positive fraction was removed from the magnet resuspended in cell selection buffer and incubated for an additional 4 min on the magnet. The positive cell fraction was resuspended in freezing medium and stored at -80°C until further use.

Cell lines

Unless stated otherwise, THP-1 monocytes, either lipopolysaccharide (LPS) stimulated or non-stimulated, were used for phage selection assays and affinity screenings. Cells were cultured in RPMI-1640 medium supplemented with glutamine and HEPES (Life Technologies) at 37°C and 5% CO₂. Cells were incubated for 1 hours with LPS (100 ng/mL; activated THP-1) or PBS (non-activated THP-1), prior to the experiment, to induce cell activation and emulate the inflammatory conditions as observed during unstable angina pectoris.

Phage library

The immune library was produced by injecting a llama (*Lama glama*) with whole cell CD14⁺ monocytes from unstable angina pectoris (UAP) patients. Two llamas were immunized at weekly intervals for a period of 6 weeks with CD14⁺ monocyte pools from 4 UAP patients. Peripheral blood lymphocytes (PBL) were isolated from immunized animals to create an immune V_{HH} library, as previously described ²¹. In short, PBL were isolated by Ficoll gradient and subsequently total RNA was isolated and transcribed into cDNA. Next, the repertoire of Ig heavy chain-encoding gene segments were amplified with the use of a framework 1 (FR1) specific primer and an oligo-dT primer. Finally, selected and purified cDNA fragments were ligated in a phagemid vector for display on filamentous bacteriophage ²² and electrotransformed to *Escherichia coli* TG1 (K12, *_(lac-pro)*, *supE*, *thi*, *hsdD5/F'traD36*, *proA+B+*, *lacIq*, *lacZ_M15*). This resulted in a immune phage library with a diversity of approximately 8 x 10⁷. The animal experiments were reviewed by the ethical committee and approved by the board of Utrecht University.

Whole Cell V_{HH} Selections

The phage library was rescued by inoculation in 2xTY medium (supplemented with ampicillin) and grown for 75 min at 37° C and 270 rpm. Subsequently, M13-helperphage (with kanamycin selection vector) was added in 10:1 ratio and incubated another 30 min. Cells were pelleted and grown overnight in 2xTY medium supplemented with ampicillin and kanamycin at 30° C and 270 rpm. Next day, phages were PEG-precipitated and blocked using 2% non-skimmed milk powder (Marvel) in PBS to prevent non-specific binding or sticking of phages to cells. Cell suspensions were also blocked for 30 min in

2%Marvel-PBS prior to selection. Blocked phages were added to control and CFSE-labeled, LPS-activated THP-1 cells (10:1 ratio), or CD14⁺ monocytes from healthy controls and CFSE-labeled CD14⁺ monocytes from UAP patients (10:1 ratio) and incubated for 30 min rotating at RT. Activated THP-1 cells or CD14⁺ monocytes from UAP patients were isolated by FACS sorting using a BD FACS Aria (BD) cell sorter based on the CFSE expression. Sorted cells were resuspended in 200 mM triethylamine (TEA; pH=12), incubated for approximately 10 min and subsequently neutralized with an equal volume 1M Tris-HCl (pH=6.8). Isolated phages were used to infect exponential TG1 (OD=0.5) and masterplates and titrations of input and output titer were determined, to determine enrichment for each selection round. Masterplates were used to prepare new glycerol solution, which was used as starting library for subsequent selection round (figure 1).

Whole Cell V_{HH} Screenings

Screenings were performed on either libraries from complete selection rounds or on individually selected clones. Marvel blocked control and LPS activated cells were incubated with 120 μ L phage suspension in blocked 96-wells V-shaped ELISA plates (Greiner) for 30 minutes at RT. Subsequently, 100 μ L of either control cells or activated cells were added to 100 μ L phage suspension and incubated for 60 minutes at 37°C. Plates were washed with Marvel-PBS three times and PBS two times.

Phages bound were detected using an HRP-conjugated anti-M13 antibody which was incubated with the cells for 30 minutes at 37°C. After three more wash steps bound phages were detected using TMB (KPL Inc., MD, USA) and analyzed in a plate reader (BioRad) at 450 nm (figure 1).

Figure 1 - Selection and screening using whole cell phage display

Phage display selections were performed according to the steps 1 – 5. In short, 1. B-cells were isolated from immunized llamas; 2. Heavy chain (V_{HH}) coding RNA was isolated and amplified; 3. V_{HH} cDNA was cloned/transfected into phagemid vector; 4. Selection on whole cells; 5. Recovery of selected phages. Subsequent affinity screening (step 6) was performed according to the steps 7 – 10 of the procedure shown. In short, 7. production of complete library of individual clone followed by binding to whole cells; 8. Washing to remove unbound phages; 9. Labeling of bound phages using anti-M13-HRP; 10. Detection of bound phages. Adapted in part from Hoogenboom (2005)¹⁷.

Flow Cytometry

Purity of isolated CD14 positive monocytes was analyzed using anti-CD14 PC7 and anti-CD16 PC5 (both BD). Measurements were performed on a BD Canto II flow cytometer (BD). In short, cells were pelleted and incubated with 1:25 diluted anti-CD14 PC7 and 1:25 diluted anti-CD16 PC5 antibodies for 30 min on ice in the dark. Subsequently cells were washed twice, dissolved in 200 μ L FACS buffer, and measured.

CD14 positive monocytes from unstable angina pectoris patients were labeled for 15 min with 1 μ M carboxyfluorescein diacetate succinimidyl ester (CFSE; Invitrogen) in serum-free medium. Subsequently, cells were washed twice in medium supplemented with 10% fetal calf serum. Phages were added to control monocytes and CFSE-labeled UAP monocytes. After incubation for 60 minutes, monocytes from UAP patients were recovered by FACS sorting based on CFSE signal. In short, cells were spun down after incubation with the phages and washed twice in PBS-2%Marvel and twice in PBS. Next cells were resuspended in PBS and sorted on a BD FACS Aria Cell sorter (BD). Both CFSE positive and CFSE negative fractions were isolated to determine efficiency and enrichment of the selection rounds.

Immunohistochemistry

Formalin fixed paraffin embedded sections from human carotid endarterectomy samples were deparaffinized and rehydrated. Sections were blocked for 30 minutes using 2% Marvel-PBS. Pre-blocked phages were added at a concentration of approximately 10^7 phages/mL in a total volume of 150 μ L per section and incubated for 60 minutes at RT. Sections were washed in subsequently 2%Marvel-PBS and PBS. Pre-blocked mouse anti-M13-HRP antibody (1:5000) was added to the sections and incubated for 30 minutes at 37°C. Next, slides were washed in PBS and Envision anti-mouse-HRP was added for 30 minutes to amplify the signal. After an additional wash step staining was developed using DAB and counterstained with haematoxylin. Finally, slides were dehydrated and coverslips were placed using Entellan.

Statistical analysis

Data are expressed as mean $\pm$ SEM. To compare individual groups, 2-tailed Students t-test was used; non-parametric data were analyzed using Mann-Whitney U test. All analyses were performed using GraphPad Prism 5 software (GraphPad Software Inc, La Jolla, CA, USA) and p-values below 0.05 were considered statistically significant.

Results

Figure 2 - THP-1 selection rounds reveal subtle yet increasing binding affinity
(A) Enrichment of the complete library after each selection round (R), indicating increased binding affinity towards activated THP-1 cells over control THP-1 cells. (B) Enrichment after selection round 4 showed heterogeneous affinity to activated cells for 19 different clones selected from this selection round. Results are based on a minimum of three screenings per clone. Results are shown as mean $\pm$ SEM, solid line indicates differential OD = 1 and dashed line indicates differential OD = 2. (C) Sequencing analysis of 20 clones showed variability in CDR1 (red), CDR2 (green), and mainly CDR3 (blue) region of V_{HH} antibodies.

THP-1 selection rounds revealed small number of high affinity clones

Initial selection rounds were performed on the THP-1 human monocytic cells, which were LPS-activated to simulate the pro-inflammatory milieu in CVD patients. After 4 selection rounds we analyzed the complete phage libraries of each selection round (figure 2A) as well as a number of individual clones from selection round 4 (figure 2B) to assess the general enrichment in monocyte

binding phages in the library as well as to pinpoint individual clones with highest affinity for LPS-activated monocytes. We observed a small increase in the affinity of the whole library. However, when analyzing the individual clones a large variety in affinity was observed for all selection rounds.

To determine whether the variety of clones present after the different selection rounds was affected between the different selection rounds we performed sequence analysis of the variable region for randomly selected clones from multiple selection rounds. Overall, enrichment resulted in diminished sequence variety. Still, at least some variety could be observed indicating the absence of a, possibly non-specific, dominant clone (figure 2C).

Figure 3 - Monocyte isolation is highly efficient and does not result in monocyte activation

(A) Monocytes were isolated from PBMC fractions. Quantification indicated that the hardly any monocytes were left in the CD14 negative (-) fraction, while the CD14 positive (+) fraction was very pure. This is also shown in the flow cytometry dot plots for the total PBMC and CD14+ fraction. (B) Histograms showing fluorescence intensity for CD11b of isolated monocytes at baseline (left panel) and after activation with 100 μM LPS (right panel).

CD14 isolation was effective and reproducible

Isolation of monocytes resulted in highly pure CD14⁺ monocyte pools (96.3% ± 2.1%) at a recovery of more than 80% for all samples. Recovery was determined by calculating the residual CD14⁺ population in the CD14 negative

fraction. Of note, there was hardly any spillover of CD14 negative cells in the isolated CD14⁺ monocyte pools (figure 3A).

The efficiency and purity observed for the isolation of fresh PBMCs did not differ from that obtained from frozen PBMCs. Freezing and thawing, either before or after CD14 isolation, did not show any effects on the CD14⁺ monocyte surface expression of CD11b, CD62L, and CCR2 (data not shown), illustrating that membrane markers are well preserved during the experimental procedure. Activation potential of isolated monocytes was neither influenced by the isolation procedure nor by the freezing/thawing step, as all cells still could be efficiently activated and showed no baseline increase of CD11b expression (figure 3B).

To verify whether the clones selected on activated THP-1 monocytes also had a higher affinity for monocytes derived from UAP patients, whole cell ELISA was performed comparing phage binding to control and UAP monocytes, isolated by CD14⁺ bead selection. Indeed, one clone (C10) showed significantly increased affinity for UAP-derived monocytes compared to monocytes from healthy individuals (figure 4A).

Figure 4 – Screenings on THP-1 cells, monocytes, and atherosclerotic lesions reveals that markers selected have biomarker potential

(A) Whole cell ELISA on human monocytes isolated from healthy controls and UAP patients showed increased affinity for UAP monocytes for some clones, compared to a control (G8) clone (B) Optical density (OD) of selected phages for binding on LPS-activated cells compared to control cells. Three clones were selected: G8C (Control; black bar), A12 (high affinity; blue bar), and C10 (intermediate affinity; orange bar). (C) Immunohistochemistry pictures for control (G8C), A12, and C10 clones on advanced atherosclerotic lesions. Inserts show positive cells for A12 and C10 staining.

Despite, the enrichment observed for the total phage libraries per selection round (figure 2A), as well as the enrichment for the individual clones was rather limited (figure 2B), we investigated the capacity of selected clones to bind monocytic cells in atherosclerotic lesions. We selected two clones for further testing; clone A12 which showed high affinity for activated THP-1 cells and clone C10 displaying moderate affinity for activated cells but increased affinity for UAP monocytes, as well as clone G8 (control) which had no increased affinity for binding to activated THP-1 cells or UAP monocytes (figure 4A+B). Early lesions (intimal thickening), advanced lesions (thick fibrous cap atheroma), or advanced, vulnerable lesions (intraplaque hemorrhage/plaque rupture) were stained with selected clones, after which binding was detected using a HRP-conjugated anti-M13 antibody. Early lesions did not stain positive for any of the tested clones (data not shown). However, clone A12 as well as C10, but not G8 (control), gave detectable staining signal for advanced atherosclerotic lesions (figure 4C).

Moreover, clones A12 and C10 were found to specifically stain cells in highly vascularized regions of the atherosclerotic plaque. Cells that showed positive staining were morphologically assigned to be monocytic cells, based on their size and nuclear shape. Interestingly, clones A12 and C10 showed a different staining pattern, with intra-cellular localized staining for A12, and clear plasma membrane staining for clone C10. Nevertheless, for both clones staining was very specific with regard to binding to monocytic cells in highly vascularized regions of advanced, vulnerable lesions.

Discussion

Diagnostic and prognostic markers used are generally based on levels of soluble proteins, lipids, carbohydrates, miRNAs, or levels of specific cell types in bodily fluids^{7,23}. One of the most frequently used markers in the field of cardiovascular medicine is troponin²⁴. This marker has very high specificity and also sensitivity for the diagnosis of an acute myocardial infarction (AMI)^{24,25}. For atherosclerosis and more in particular unstable angina pectoris no (protein) markers have been defined until now and therefore diagnosis is merely based on clinical parameters and exclusion of AMI⁵. In this study we show that circulating monocytes carry information on local vascular disease status which could be applicable to diagnose or even predict (future) acute cardiovascular syndromes.

Phage display is a powerful technique that yields high efficiency for the unbiased discovery of novel antibodies against known as well as unknown epitopes¹⁷. The technique has already been successfully applied in the cardiovascular field, to identify plaque proteins targeting autoantibodies²⁶ and to design binders of known candidates for molecular imaging and intervention such as P-selectin²⁷ and SR-AI²⁸, or of whole cells^{29,30}. In this study we applied phage display to identify antibodies able to differentiate between activated and non-activated on monocytes, for use in molecular diagnosis of unstable angina pectoris. Standard selection and screening protocols²⁰ were adapted to be able to accommodate whole cells as template. One of the challenges to be met, especially in the validation phase, was that, contrary to most phage display efforts thus far, antibody and antigen both were unknown at the start of the study. We opted for activated THP-1 monocyte cells as selection platform, to circumvent interpatient variability; taken advantage of the recent findings that UAP and AMI patient monocytes show altered TLR4 response and/or basal activation status³¹⁻³⁴, patient disease status was emulated by pre-activating monocytes with the TLR4 agonist lipopolysaccharide (LPS).

For the verification stage we used patient-derived CD14+ monocytes. Importantly, the activation status of CD14+ monocytes was not impacted by cell isolation and CD14+ still retained full responsiveness to stimuli such as lipopolysaccharide (LPS). Thus, differential binding of phages to UAP monocytes versus monocytes from healthy volunteers was very likely reflecting disease state, although we cannot completely exclude that differences between patient groups in other parameters such as medication or diet could have had an effect on proteins expressed by circulating monocytes. Further validation in larger cohorts of risk factor matched patient groups will be required to address this point.

As TLR4 activation was applied for the initial biomarker discovery phase, the specificity of the binders found does not have to be restricted to monocytes in relation to advanced or unstable atherosclerosis. TLR4 has been implicated in rheumatoid arthritis³⁵ and systemic lupus erythromatosus, however TLR4 appears to be non-involved or downregulated in these autoimmune diseases^{36,37}. Still, TLR4 activation is implicated to be involved in multiple sclerosis³⁸ and possibly asthma^{39,40}. Therefore, TLR4 activation dependent biomarkers could contribute at least to some of the

aforementioned diseases. Nevertheless, TLR4 signaling is most often linked to CVD, underpinning the relevance of our approach in a cardiovascular setting.

Monocytes have a pivotal role in all phases of atherosclerosis development⁴¹ and interact with the vessel wall⁴². These cells are the first cells to enter the initial lesion upon injury⁴³, and the main source of macrophages which were seen to exert important functions in plaque progression⁴⁴ and eventual plaque destabilization, resulting in clinical manifestations such as myocardial infarction and stroke⁴⁵. This key role in disease ontogenesis, combined with their innate patrolling function renders them highly sensitive sentinels of local vascular disease. Despite the cognition of macrophages as biomarker source for CVD status⁴⁶, circulating monocytes have hardly been considered as carrier of information on plaque or vessel status, apart from their rolling and adhesion behavior. However, interaction with the diseased endothelium, as well as with blood containing cells and cytokines, can impact monocyte phenotype as we show in the current study. This notion concurs well with previous studies, which investigated disease-associated changes in monocyte expression and their potential as biomarker in cardiovascular diseases, as reviewed by Gratchev *et al*⁴⁷. Different types of monocyte markers are described including monocyte (subset) numbers, protein markers, and functional markers. Both total monocyte counts⁴⁸ as well as subpopulations^{49,50} (e.g. CD14⁺⁺/CD16⁻ monocytes) can function as biomarker for CVD. Further, activation markers as CD18, CD11b, and CXCR1 give information on morbidity in heart failure⁵¹. Finally, increased monocyte-endothelial cell interactions⁵² and increased activation potential upon IL-4 stimulus⁴⁷ can also give information on local atherogenic disease. The mere fact that our phage marker is able to distinguish between vulnerable and stable plaque monocytes is highly supportive of cross-interaction between lesion and monocyte.

To conclude we offer proof of concept of phage display aided identification of monocyte markers with increased affinity for circulating and plaque monocytes in unstable or advanced atherosclerosis. While holding promise, the actual target of the phage exposed antibody remains to be determined and its specificity assessed, before this observation can be translated into a clinically viable diagnostic test. Defining the exact epitope will also give a better insight in the disease associated downstream signaling events and its consequences for monocyte phenotype in vascular pathologies

such as investigated in this study, but potentially also in other inflammatory diseases, especially since TLR4 activation is present in acute inflammatory conditions. Taken together, we present novel markers for circulating and plaque monocytes in patients with unstable angina which could contribute to improvement of diagnosis of disease.

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The background of the entire page is a white surface covered with numerous red splatters and droplets of varying sizes, resembling blood. Some splatters are large and circular, while others are thin, elongated streaks or fine mist-like sprays. The distribution is dense, with a higher concentration of larger droplets in the lower half and more fine sprays in the upper half.

Chapter 06

Circulating unstable angina pectoris patient monocytes harbor no basal transcriptomic differences

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Abstract

Angina pectoris is considered to be one of the hallmarks of advanced cardiovascular disease and it often precedes ischemic events such as myocardial infarction and stroke. The current study investigates the differences between stable and unstable angina pectoris on a transcriptomics/genomic level to determine processes involved and to discover biomarkers to distinguish between stable and unstable angina pectoris. Circulating monocytes from patients included in the CTMM Circulating Cells cohort were isolated upon admission to the hospital. Monocytes were analyzed on a cellular level by flow cytometry and on a genomic level using micro-array analysis. Both the analysis on a proteomic and genomic level indicated the absence of distinct differences between the stable and unstable angina pectoris patients, however both analyses indicated the involvement of migratory pathways, as chemokine and cytokine signaling were limitedly affected. Despite major clinical differences between patients suffering from stable and unstable angina, we were unable to segregate these patient populations on a cellular or genetic level, indicative of only minor alterations of circulating monocytes in response to local atherosclerotic vessel disease.

Introduction

Cardiovascular diseases are one of the leading causes of death in Western societies and prevalence is increasing in developing countries ¹. Despite the growing public health efforts in preventive medicine, most patients will be referred to the hospital when already suffering clinical symptoms such as chest pain. These symptoms, also referred to as angina pectoris, are caused by reduced availability of oxygen (and nutrients) to the myocardium. Almost ubiquitously the underlying cause is represented by stenosis or impaired dynamics of the coronaries due to atherosclerotic disease ². Distinction between stable angina pectoris (SAP) and unstable angina pectoris (UAP) is made based on clinical history and semeiotics. Since this is a subjective process, and plaque stability in general is an important variable when it comes to the prognosis of a patient with angina, there is increasing need for alternative markers which can be used to classify angina pectoris into stable or unstable.

Preferably, such markers should also be able to quantitatively estimate the risk of developing future cardiovascular events, with high reproducibility, or at least reflect key processes in disease development. Additionally, it would be desirable to identify circulating biomarkers, easily measurable with an easy and stress-free blood sampling procedure, to minimize the physical burden for patients and to seamlessly combine testing within the already established clinical procedures (e.g. troponin measurement ³). Further, markers for SAP and UAP should have prognostic features to ensure optimal assessment of the long term risk of cardiovascular events and develop personalized medicine strategies.

In recent years, biomarker research has broadened its focus to also include cellular markers besides more classical (soluble) circulating proteins. Established protein biomarkers include C-reactive protein (CRP) for acute coronary syndrome ^{4,5}, but also cytokines like growth differentiation factor 15 (GDF15) for CVD in general ⁶. Recently, increasing numbers of various UAP specific biomarkers have been proposed, including serum levels of placental growth factor ⁷, deoxyribonuclease I ⁸, matrix metalloproteinase (MMP) ^{8 ref 9}. Cytokines, like soluble CD40 ^{ref 10}, pro-brain type natriuretic peptide (BNP) ¹¹, and myeloperoxidase (MPO) ¹², have also been suggested as biomarkers for CVD. Furthermore, monocyte subset (CD14⁺/CD16⁺/CX₃CR1⁺) counts ¹³, as well as levels of specific membrane markers as the fractalkine receptor (CX₃CR1) ¹⁴

have been linked to plaque stability in UAP patients. Nevertheless, most of these markers are either restricted to specific patients populations¹⁵, or influenced by drug use¹³, limiting the clinical application as biomarker.

Novel cellular markers for CVD are circulating endothelial cells, which could be promising prognostic biomarkers¹⁶. Also markers on circulating leukocytes have been shown to have potential as diagnostic or even prognostic CVD biomarker. In fact, CD14 expression on monocytes is commonly studied to determine monocyte activation and disease progression¹⁷⁻¹⁹, however the results are still contradictory. It has been shown that inflammatory membrane markers such as toll-like receptors^{20,21} and CD11b/CD18^{ref 22,23} are also associated with progression of CVD in general. However, many biomarker studies have a lack of discriminative power, are not specific to cardiovascular disease, or do not show diagnostic potential and risk prediction for individual patients.

These data nevertheless demand a more elaborate screening of proteome and transcriptome on relevant leukocyte species. As cells from the monocyte/macrophage lineage are accepted as one of the key players in atherosclerosis development, we focused on the potential of circulating monocytes as biomarker for angina pectoris. Furthermore, a transcriptome approach is very suitable for identifying novel biomarkers²⁴, and can also give insight in underlying pathway regulation^{25,26}. Here, we present data on transcriptomics analysis on UAP patient monocytes compared to SAP monocytes to identify novel targets for diagnosis and prognosis.

Materials & Methods

Patients

This study included 25 patients with stable angina pectoris and 25 patients with unstable angina pectoris from the Center for Translational Molecular Medicine (CTMM) – Circulating Cells Cohort that were presented to the Maastricht University Medical Center, The Netherlands between July 2009 and June 2011. Patient groups were matched for age, sex, smoking habit, dyslipidemia, and hypertension (table 1). As negative control, 11 healthy individuals without any known cardiovascular disease were included. These controls presented to the hospital with chest pain of suspected cardiovascular origin, but did not show any overt stenosis and scored negative on ST elevation and/or troponin T (TnT). All participants provided written informed consent

prior to participation. This study was approved by the local ethics committee. Exclusion criteria were active inflammatory conditions, autoimmune disease, malignancies, use of immunosuppressive drugs, and known hematological disorders. Patients with ST-elevation myocardial infarction (STEMI) were also excluded. Blood of patients with suspected unstable angina or non-STEMI was studied but retrospectively excluded from this analysis.

Cell Isolation

Upon inclusion, blood samples were collected in ethylenediaminetetraacetic acid (EDTA) anti-coagulated vacuum tubes and processed according to standardized procedures to analyze leukocyte subsets. In short, blood was transferred to a 50 mL tube (Greiner) and centrifuged for 15 minutes at 156x g without brake. Subsequently, plasma was removed and 30 mL PBS-diluted blood was added on top of 15 mL Ficoll-Paque plus (Sigma) and spun for 20 minutes at 1000x g without brake. The leukocyte-enriched interphase was collected and washed twice with PBS before CD14⁺ monocyte isolation according to the manufacturers protocol (BD Bioscience). Isolated CD14⁺ monocytes were gently frozen and stored at -80°C until further use.

Flow Cytometry

To assess expression of surface markers 50 µl of heparin anti-coagulated whole blood was incubated with fluorescent antibodies (supplemental table 1) for 30 minutes. After washing and whole blood erythrocyte lysis, the samples were analyzed by flow cytometry (Beckman Coulter FC 500). Granulocytes were gated based on their scatter properties, and confirmed based on CD16 expression. Monocytes were identified based on their scatter properties and monocyte subsets were quantified (percentage positive cells and mean fluorescence intensity [MFI]) based on surface expression of CD14, CD16, CCR2, and CX₃CR1.

RNA Isolation and Micro-Array Analysis

Monocyte samples were shipped to AROS (Aarhus, Denmark) for RNA isolation and subsequent micro-array analysis. RNA was isolated using Illumina TotalPrep RNA Amplification Kit (Illumina, San Diego, CA, USA) and cDNA was produced. The RNA quality and integrity was very good, as the lowest RNA integrity number (RIN) was 8.3 besides on outlier with a RIN of 2.4. The

average RIN was 9.22 ± 0.14 , indicative of very good quality of the RNA samples.

Next, labeled cRNA was prepared and used on the array for hybridization according to manufacturer's instructions. HumanHT-12 v4 Bead chips were scanned by Illumina BeadStation (Illumina Inc., San Diego, CA, USA). Raw image analysis and signal extraction was performed with Illumina Beadstudio Gene Expression software with default settings (no background subtraction) and no normalization. Data were exported as text files.

Computational methods

Raw data were imported in R ^{ref 27} and analyzed within the Bioconductor suite ²⁸. Quality control, background correction, and robust spline normalization have been performed by lumi ²⁹. Differential expression analysis was performed by fitting a linear model to the paired set of expression values, and by applying a Bayesian empirical method for shrinkage of the standard errors towards a common value, in limma ³⁰. P-values have been adjusted for False Detection Rate ³¹.

To evaluate the reproducibility of our differential expression analysis, the package GeneSelector ³² was used. First the differential expression test was repeated using a set of diverse statistical methods (Wilcoxon statistics empirical Bayes ³³, Wilcoxon statistics, Bayesian t-statistics ³⁴, and SAM ³⁵). Finally we repeated the analysis for altered (jackknife, label exchange, bootstrapping, and noise addition) datasets, to estimate the bias and variance of the data. A principal component analysis was ran to explore the relationship among the samples. Arbitrarily selecting the top 500 genes from the differential expression analysis results, pathway enrichment analysis was performed using signaling pathway impact analysis (SPIA) ³⁶ which leverages on a system biology approach to integrate information concerning gene ID, differential expression and pathway topology to calculate the pathway significance and its activation or inhibition.

Statistical Analysis

All data on patient characteristics and flow cytometry are presented as mean $\pm$ SEM. To compare individual groups, 2-tailed Students t-test was used; non-parametric data were analyzed using Mann-Whitney U test; discrete data was analyzed using Chi square test. All analyses on patient characteristics and flow cytometry were performed using GraphPad Prism 5 software (GraphPad Software Inc, La Jolla, CA, USA) or IBM SPSS statistics 20 (IBM, Amsterdam, The Netherlands). *p*-values < 0.05 were considered statistically significant.

Results

Patient characteristics

In total 25 stable angina pectoris (SAP) patients, 25 unstable angina pectoris (UAP) patients, and 11 controls were included in this study. However one UAP patient was reclassified as SAP by the clinicians after initial selection procedure was performed. A second UAP patient was excluded as no adequate monocyte sample was available. As a result the study comprised 23 UAP patients, 26 risk factor-matched SAP patients, and 11 controls for final analysis.

Table 1 – Patient characteristics

Baseline characteristics for healthy controls (controls), stable angina pectoris patients (SAP), and unstable angina pectoris patients (UAP). Indicated is: number (percentage of group), except for Age (mean); BMI = Body Mass Index, MI = Myocardial Infarction

	Controls	SAP	UAP	<i>p</i> -value
	N = 11	N = 26	N = 23	
Age	59.55	63.23	63.7	0.50
Sex	7 (63.6)	14 (53.8)	13 (56.5)	0.86
BMI	28.34	26.92	26.97	0.70
Smoker	3 (27.3)	9 (34.6)	11 (47.8)	0.45
Hypertension	4 (36.4)	15 (57.7)	16 (69.6)	0.18
Diabetes Mellitus	2 (18.2)	3 (11.5)	7 (30.4)	0.25
Dyslipidemia	5 (45.5)	21 (80.8)	16 (69.6)	0.10
Peripheral artery disease	2 (18.2)	3 (11.5)	3 (13.0)	0.86
Previous MI	0 (0.0)	7 (26.9)	3 (13.0)	0.11
Cerebrovascular attack	0 (0.0)	2 (7.7)	5 (21.7)	0.13

No statistically significant differences in patient characteristics were detected for age, sex, and body mass index (BMI) between healthy controls, SAP and UAP patients (table 1). In the healthy control group 7 patients were male (63.6%) and the mean age was 59.6 years (59.6 ± 3.2 years). It should be noted that accepted risk factors as smoking, hypertension, diabetes mellitus, and dyslipidemia, additional matching criteria for patient group buildup, did not differ between groups (table 1). Despite the absence of previous myocardial infarction and cerebrovascular attacks in the control group, no statistically significant difference could be observed between healthy controls, SAP, and UAP patients (table 1).

Leukocyte counts are unaltered between SAP and UAP patients

No statistically significant differences were observed in total leukocyte counts between controls, SAP, and UAP patients (table 2). Also lymphocyte and granulocyte counts were unchanged between controls and patient groups (table 2). Both lymphocytes and granulocytes tended to increase with severity of disease (*controls* $\rightarrow$ *SAP* $\rightarrow$ *UAP*), albeit these results did not reach significance. Of note, monocyte counts were significantly reduced in SAP patients compared to both healthy controls and UAP patients, but did not differ between the latter two groups (table 2).

Table 2 – Leukocyte profile of patients included

Absolute cell counts from healthy controls (controls), stable angina pectoris patients (SAP), and unstable angina pectoris patients (UAP) upon inclusion. Indicated is mean $\pm$ SEM. * $p < 0.05$ vs UAP

	Controls	SAP	UAP	<i>p-value</i>
	N = 11	N = 26	N = 23	
Leukocytes	7.00 (± 0.42)	6.90 (± 0.66)	7.47 (± 0.80)	0.82
Lymphocytes	1.66 (± 0.17)	1.83 (± 0.19)	2.27 (± 0.20)	0.15
Monocytes	0.55 (± 0.04)	0.39 (± 0.03)*	0.51 (± 0.04)	< 0.01
Neutrophils	4.68 (± 0.65)	5.35 (± 0.35)	5.99 (± 0.42)	0.23

Flow cytometry excludes polarization bias between patient groups

Flow cytometry analysis confirmed the above findings, in that there were no differences in granulocytes between stable and unstable angina (figure 1A). Relative total monocyte counts measured by flow cytometry (figure 1B), as well as CD14^{high}CD16⁻ and CD14⁺CD16⁺ monocyte subsets were unaltered in UAP versus SAP patients (figure 1C+D). The CCR2 and CX₃CR1 chemokine receptor density on CD14^{high}CD16⁻ and CD14⁺CD16⁺ was not affected as mean fluorescence intensity (MFI) for these chemokine receptors on both monocyte subsets was unaltered in SAP and UAP patients (table 3).

Figure 1 - Flow cytometry indicates no differences in chemokine receptor expression.

Flow cytometry data showing (A) granulocytes, (B) total monocytes, (C) CD14^{high}CD16⁻ monocytes, and (D) CD14⁺CD16⁺ monocytes for SAP and UAP patients. White bars are stable angina pectoris patients (SAP) and black bars are unstable angina pectoris (UAP) patients. * $p < 0.05$ and ** $p < 0.01$

Table 3 – Corrected mean fluorescence intensity for chemokine receptor expression on monocyte subsets

Corrected MFI		Controls	SAP	UAP	<i>p-value</i>
		N = 11	N = 26	N = 23	
CD14 ^{high} CD16 ⁻	CCR2	0.283 (± 0.011)	0.288 (± 0.007)	0.284 (± 0.009)	0.93
	CX ₃ CR1	0.290 (± 0.003)	0.293 (± 0.004)	0.289 (± 0.003)	0.52
CD14 ⁺ CD16 ⁺	CCR2	0.309 (± 0.022)	0.297 (± 0.013)	0.304 (± 0.010)	0.84
	CX ₃ CR1	0.339 (± 0.005)	0.344 (± 0.004)	0.344 (± 0.003)	0.56

Principal component analysis does not reveal any segregation between patient groups

Flow cytometry based quality control for the monocyte isolation confirmed that CD14 positive monocytes could be isolated at high purity (> 99.5%) and moderate to high recovery (> 80%; figure 2A). Bead-purified monocytes were still viable and still able to react to subsequent lipopolysaccharide activation, indicating that the selection method did not (completely) activate the monocytes (figure 2B).

Density histogram of raw log-intensities from patient samples before (upper panel) and after (lower panel) preprocessing revealed that the micro array analyses were of good quality. This was confirmed by the overlap of samples after normalization, which indicates that batch effects have been eradicated and that all the signals share the same scale (figure 3A).

Figure 2 - Monocyte are isolated efficiently and without massive activation.

(A) Representative dot plots for monocyte population based on CD14 and CD16 expression before (left panel) and after (right panel) CD14⁺ bead isolation. (B) Histograms showing CD11b expression on isolated monocytes before (upper panel) and after (lower panel) stimulation with 100 ng/mL lipopolysaccharide (LPS).

Initial principal component analysis (PCA) was performed on the raw data to identify whether the attributes of the data were able to show clustering of the datasets. PCA indicated that there was no segregation of the data based on the patients disease state (control, stable angina pectoris, unstable angina pectoris) before normalization and background correction (figure 3B). Normalization of the data and subsequent background correction, did not reveal additional information, as there was no clustering of patient groups (or controls) visible (figure 3C).

Monocyte show subtle differences on transcriptomics level

Differential expression analysis, after false discovery rate (FDR) adjustment (to correct for multiple testing), was used to determine genes of interest for further analysis. Analysis revealed the presence of both up- and down-regulated gene sets in UAP versus SAP patients after correction for the controls (figure 4A). However, after stringent multiple testing correction we could no longer observe any statistically significant gene sets (data not shown) for the used conditions ($p < 0.01$ and/or foldchange $> 30\%$).

To verify whether the results were influenced by the statistical test used for these analyses, multiple tests were compared, including Wilcoxon statistics empirical Bayes (WilcEbam), Wilcoxon statistics (Wilcoxon), Bayesian t-statistics (BaldiLongT), and statistical analysis of micro arrays (SAM). All tests, except for WilcEbam, resulted in similar outcomes regarding macroscopic clustering of the results as our default procedure linear models for micro-array data (Limma), suggesting that our analysis was not biased by the procedure used for analysis (figure 4B).

Figure 3 - Micro-array analysis does not result in clustering of patient groups. (A) Micro-array analysis quality was assessed using density histograms of log2 intensity. Density histograms before (upper panel) and after (lower panel) processing are shown. Overlay of the lines after processing is indicative of good quality. (B + C) Principal component analysis (PCA) showed the absence of clustering of patient populations before (B) and after (C) background correction and normalization of the data.

Figure 4 - Differential expression analysis indicate minute, yet significant differences. (A) Volcano plot indicating differentially expressed genes (blue dots) with fold change over 30% and/or p-value below 0.01. Red dots are genes not differentially expressed between SAP and UAP patients. (B) Macroscopical ordering of the results based on different statistical tests does not reveal test dependent bias of our results.

Monocyte chemokine pathways are minimally affected in UAP patients

Pathway analysis, arbitrarily using the top 500 genes differentially expressed between SAP and UAP patient monocytes, indicated seven pathways which were significantly inhibited in UAP patients compared to SAP patients after correction for the controls. Two additional pathways were activated in UAP versus SAP patients (figure 5A). Four pathways remained significant after Bonferroni correction, and all nine were significant after FDR correction. The top four pathways were all immune-related: chemokine signaling pathway (inhibited; $p < 0.001$), cytokine-cytokine receptor interaction (inhibited; $p < 0.01$), vibrio cholera infection (inhibited; $p < 0.05$), and epithelial cell signaling in *H. pylori* infection (inhibited; $p < 0.05$; figure 5B). The other five pathways, Shigellosis (inhibited), cell cycle (inhibited), salivary secretion (inhibited), pathogenic *E. coli* infection (activated), and Alzheimer's disease (activated), did not pass Bonferroni correction (figure 5A+B). Besides cell division control protein 42 homolog (Cdc42) and adenylyl cyclase (AC), chemokines and chemokine receptor genes were most predominantly affected in the top dysregulated pathway (chemokine signaling pathway; figure 5C).

Figure 5 - Pathway analysis reveals subtle differences between SAP and UAP patients.
(A) SPIA pathway analysis reveals differentially regulated pathways between SAP and UAP patients based on Bonferroni correction (red dots) and FDR (blue dots). The red line and blue line indicate $p < 0.05$ for Bonferroni and FDR correction, respectively. (B) Table indicating differentially regulated pathways including KEGG-ID, adjusted p-value, and status (activated/inhibited). (C) Chemokine pathway including the differentially expressed genes (red genes) within this pathway.

Discussion

Distinguishing between stable angina pectoris (SAP) and unstable angina pectoris (UAP) patients at an early time point is essential for treatment of cardiovascular disease and risk prediction for future cardiovascular events. Furthermore, it is also essential to exclude patients with non-cardiovascular chest pain early to avoid additional, time-consuming, and expensive examinations and procedures. Various studies have investigated circulating or even cellular biomarkers to determine the disease state of the patient³⁷⁻³⁹. So far, no biomarker has been defined that is superior to commonly used clinical parameters, in discriminating SAP from UAP patients^{40,41}. Cellular markers can not only give insight in the disease state of a patient, but can also contribute to understanding of underlying mechanisms.

The current study has explored the possibilities of using blood derived cellular markers for determination of disease state. The underlying notion is that circulating cells, and more in particular monocytes, can on the one hand sense the local and systemic milieu by either reacting to released cytokines⁴² and chemokines⁴³, and on the other hand directly interact with the vessel wall^{5,43-45}, activating (patrolling) monocytes due to increased integrins in response to advanced atherosclerotic vessel disease^{44,46}. Conceivably, the transient accumulation of activated platelets at lesioned foci in coronary arteries⁴⁷, one of the processes underlying angina, will be sensed at an early stage by circulating monocytes. Moreover, dysregulated plasma cytokine and chemokine levels constitute a markedly altered microenvironment to which circulating monocytes are exposed. Overall, monocytes have the capacity to detect multiple targets associated with the pathogenesis of angina pectoris, and to respond to such stimuli on a cellular level.

The possibilities of using circulating leukocytes as target for biomarker discovery is not completely novel, as a limited number of studies has investigated peripheral blood mononuclear cells (PBMC) as messenger of atherosclerotic vessel disease using a transcriptomics approach⁴⁸. Moore *et al* found a panel of differentially expressed genes, including genes involved in monocyte activation (CD14, TLR2) and scavenger receptors (CD163, CD36), which could identify stroke patients⁴⁹. Furthermore, Patino *et al* identified Finkel-Biskis-Jenkins osteosarcoma (FOS) gene to be upregulated in circulating CD14⁺ monocytes from patients undergoing carotid endarterectomy compared

to healthy controls⁵⁰. However, in contrast to our study, these studies focused on endpoints (surgery and stroke) rather than diagnosis of patients at risk.

In this study, the SAP and UAP patients selected were matched for risk factors related to cardiovascular disease. The healthy controls did not suffer from cardiovascular disease before or upon inclusion. Leukocyte analysis indicated that total leukocytes, as well as lymphocytes and granulocytes were not different between healthy controls, SAP and UAP patients. Unexpectedly, circulating monocytes were slightly decreased in SAP patients, but not UAP patients, compared to healthy controls. Overall, patient demographics are not expected to have any influence on our analysis as parameters were very comparable between controls, SAP and UAP patients.

Flow cytometry did not show any differences in monocyte numbers, both total monocytes and subsets, as well as in chemokine receptor expression. With regard to monocytes, inflammatory $CD14^{high}CD16^{-}$ and patrolling $CD14^{+}CD16^{+}$ monocyte subsets, we could not detect any differences in relative abundance. Recently $CD14^{+}CD16^{+}$ monocyte enrichment was proposed as a measure of disease severity⁵¹, however this study did not differentiate between stable and unstable angina. On the protein level, we did not observe any changes in expression of two prototypic chemokine receptors instrumental in $CD14^{high}CD16^{-}$ and $CD14^{+}CD16^{+}$ trafficking, CCR2 and CX3CR1 as examined by flow cytometry. These data indicate that our transcriptomics analysis is not confounded by basal differences in SAP or UAP monocyte skewing. This supports, at least in part, the subtle differences in monocyte phenotype between SAP and UAP patients.

More detailed analysis of our transcriptomics data by principal component analysis failed to reveal clustering of patient groups. Previously, individual genes were suggested to be useful as a biomarker for coronary artery disease^{52,53}, or as prognostic marker for disease progression⁵⁴. However, as Venet *et al* showed individual genes tested might be as predictive for a specific disease as randomly chosen genes⁵⁵. To prevent detection of non-specific genes, we conducted in-depth genome wide analysis to specifically detect gene clusters predictive for disease state. Differential expression analysis unveiled, in contrast to our flow cytometry data, subsets of genes to be up- or downregulated in unstable angina pectoris patients compared to stable angina pectoris patients after correction for the healthy controls, demonstrating that transcriptomic differences, though subtle, do

exist between the patient populations. Pathway analysis indicated chemokine and cytokine pathways, as well as pathways related to infection and inflammation to be inhibited in UAP patients. This confirmed the notion that local ischemia and/or unstable atherosclerosis, and their resulting signals can impact circulating monocyte functions such as chemotaxis. The initially unexpected inhibition of chemokine – chemokine receptor interaction in UAP patients may also explain the elevated monocyte levels in this group compared to SAP patients, as the most response monocytes in UAP patients might be recruited to the site of injury, in this case the atherosclerotic lesion.

In conclusion, we were able to show that circulating cells do harbor hallmarks of local atherosclerotic disease, as UAP monocytes showed subtle transcriptomics differences on e.g. chemokine and cytokine receptor regulation compared to SAP monocytes. However, the detected effects are too small to justify circulating monocyte transcriptomics for biomarker derivation. Additional research is necessary to investigate the best approach to amplify or detect the subtle genomic or even downstream proteomic differences between SAP and UAP in a bed-side setting.

Supplemental data

Supplemental table 1 - Antibodies for flow cytometry

Antibody	Company	Dilution
CD14 PC7	BD Biosciences	1:10
CD16 PC5	BD Biosciences	1:10
CCR2 PE	BD Biosciences	1:5
CX₃CR1 FITC	BD Biosciences	1:5
CD3 ECD	BD Biosciences	1:10
CD4 PE-Cy7	BD Biosciences	1:10
CD8 FITC	BD Biosciences	1:5

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Chapter 07

General Discussion & Future Perspectives

Main findings in perspective

The role of the monocyte/macrophage lineage has already been considered detrimental in atherosclerosis development for many years. Especially plaque macrophages have been studied extensively in relation to pathogenesis ¹. Thereby, the focus was not only on unraveling processes involved, but also on the determination of biomarkers for diagnosis and prognosis of cardiovascular disease (CVD) in general ^{2,3}. Over the last years, research has shifted its focus from plaque macrophages to circulating monocytes in relation to atherogenesis. In fact, it is acknowledged nowadays that these circulating monocytes might be as important for disease as the macrophages present in the atherosclerotic plaques ⁴⁻⁷. Nevertheless, many functional aspects of circulating monocytes are not yet elucidated completely, e.g. the functional effects of hypercholesterolemia on circulating monocytes. Furthermore, these cells might yield so far unknown potential as biomarker, due to their monitoring capabilities of the vasculature.

This thesis focused on the role of circulating monocytes in atherosclerosis, and the question whether these cells exert effects locally in the lesion or systemically in the periphery. We set out to assess the effects of (I) depleting monocytes/macrophages from the circulation and lesion, as well as (II) the role of migration of circulating myeloid cells in relation to atherogenesis. Further, we studied (III) the functional effects of hypercholesterolemia on monocytes, and (IV) set out to gain more insight in the potential of circulating monocytes as biomarker for cardiovascular disease. The main findings of this thesis are:

1. Depletion of both CD115⁺ circulating monocytes and intimal macrophages attenuates plaque progression (**Chapter 2**)
2. Increased monocyte chemotaxis towards the lesion, and thus augmented intimal macrophage numbers, reduces atherogenesis (**Chapter 3**)
3. Hypercholesterolemia induces excessive cholesterol accumulation and functional changes in circulating monocytes (**Chapter 4**)
4. Activated circulating monocytes harbor potential as (diagnostic and imaging) biomarkers in human atherosclerotic vessel disease (**Chapter 5 and 6**)

The effects of monocyte influx on plaque progression

In **Chapter 2**, we investigated the effects of depleting both plaque-resident and systemic myeloid cells on atherosclerosis development. Here, a clear role for both plaque macrophages and circulating monocytes could be deduced from our results. In the first study we assessed the impact of whole body CD115⁺ leukocyte ablation, as CD115 (or M-CSF receptor) is essential to monocyte/macrophage development. In contrast to our expectations, no effects on plaque development could be observed, which was only in part comparable to other studies on monocyte/macrophage ablation as reviewed by Croons *et al*⁸. These studies used genetic approaches in which human diphtheria toxin receptor (hDTR) expression was controlled by the myeloid specific promotor CD11b^{9,10}, pharmacological ablation using clodronate liposomes^{11,12}, and several other approaches. Even though the approaches differ substantially, the level of depletion of myeloid cells achieved in our study was at least comparable to these studies.

The difference between earlier studies and the study we performed was the combination of local apoptosis of plaque macrophages combined with systemic apoptosis of monocytes and granulocytes, but also osteoclasts. Calin *et al* already reported that depletion of monocytes and macrophages can act as a two-edged sword¹². On the one hand, depletion of the monocyte/macrophage reduces the proteolytic and pro-inflammatory signals in the lesion. On the other hand, secondary necrosis accompanied by the release of cellular cholesterol and lipid deposits into the lesion contributes to plaque progression. Indeed, Döring and Soehnlein *et al* reported that interferon regulatory factor (IRF) 8 deficiency affects monocyte/macrophage function, resulting in defective intracellular lipid accumulation and disturbed efferocytosis. In combination with increased lesional neutrophil influx this led to accelerated atherosclerosis¹³.

Unexpectedly, CD115⁺ leukocyte ablation affected hematopoiesis in which osteoclasts and bone marrow macrophages were depleted. This ablation resulted in uncontrolled release of immature leukocytes and progenitor cells from the bone marrow, as macrophages and osteoclasts are important regulators of hematopoiesis¹⁴⁻¹⁶. This contributed to the extramedullary myelopoiesis and monocytosis, in particular in spleen. Colony forming unit assays confirmed a shift in hematopoiesis from bone marrow to spleen.

Overall, not only plaque monocytes/macrophages, but also peripheral ablation of CD115⁺ cells affected atherogenesis.

Furthermore, local induction of apoptosis in the atherosclerotic lesion resulted in plaque progression rather than attenuation of plaque development. Also here an important role for circulating monocytes could explain for the effects observed. Massive intimal macrophage apoptosis will lead to the build-up of inefficiently cleared apoptotic cellular debris¹⁷. Ineffective efferocytosis indeed was shown to result in secondary necrosis and increased necrotic core formation, which has major implications for atherosclerotic plaque progression¹⁷⁻¹⁹. Consequently, a more pro-inflammatory, rather than anti-inflammatory milieu will develop resulting in the attraction of monocytes from the blood and enhanced plaque development²⁰. In conclusion, our study identified an important contribution of CD115⁺ myeloid cells outside the atherosclerotic plaque affecting lesion progression. Nevertheless, localized modulation of monocytes/macrophages in the lesion could potentially be a valid target for therapeutic intervention as circulating monocytes do affect atherogenesis.

Next to ablation of monocytes/macrophages, the role of chemokine receptor desensitization was investigated in **Chapter 3**. In this study, we addressed the effects of hematopoietic GRK2 deficiency in atherosclerosis. GRK2 is a receptor kinase mediating desensitization and receptor recycling of G-protein coupled receptors for monocyte function and chemotaxis, such as CCR2 and CCR5^{21,22}. Inhibited desensitization of chemokine receptors in monocytes resulted in expanded plaque macrophage content, but decreased plaque size. Chemokine receptor signaling is a central process in atherogenesis²³, but also in hematopoiesis, in particular for regulating stromal release of mature monocytes²⁴. Indeed, inhibition of the CCL2, CX₃CR1, or the CCL5/CCR5 axis results in dramatically reduced atherogenesis, indicative of the importance of both the chemokine receptor and ligands²⁵. Besides involvement of GRK2 in chemokine receptor desensitization, the GRK2 interactome has been extended to endocytic proteins, but also non-receptor targets, e.g. growth arrest specific (Gas) 8^{ref 26}. GRK2 dysfunction is involved in several diseases, including cardiovascular pathologies (e.g. heart failure)²⁷ and chronic inflammatory disease (e.g. rheumatoid arthritis)²⁸. From the target receptors in combination with diseases with comparable pathogenesis, we deduced that GRK2 may also play an important role in atherosclerosis.

Hematopoietic GRK2 deficiency led to enhanced monocyte migration, but attenuated plaque development. The role of circulating monocytes was inferred by excluding granulocytes and macrophages being responsible for the plaque phenotype. GRK2 deficiency resulted in increased proliferation as well as monocyte migration. Overall, increased monocyte chemotaxis resulted in plaque attenuation, most likely due to more efficient efferocytosis of apoptotic material, explaining the dramatic reduction in necrotic core size. These data show that stimulating monocyte chemotaxis towards the atherosclerotic lesion might be a promising target for intervention, especially in early atherosclerosis.

Hypercholesterolemia modifies circulating monocytes and their function

Next to inflammation-induced recruitment of monocytes towards the atherosclerotic lesion, hypercholesterolemia is the second hallmark of, and risk factor for, atherosclerosis²⁹. Despite, the well-known association between hypercholesterolemia, especially increased LDL cholesterol levels, and the risk for cardiovascular disease, still little is known about the impact of hypercholesterolemia on circulating monocytes in the context of early atherosclerosis development. Most research focuses on the effects of long term (> 10 weeks) hypercholesterolemia on monocytes^{30,31}. In our study we analyzed the short term (< 3 weeks) effects on monocyte function upon Western type diet feeding and subsequent hypercholesterolemia.

In **Chapter 4** we analyzed the early effects of diet-induced hypercholesterolemia on circulating monocyte subsets. Interestingly, and in contrast to the current paradigm, cholesterol does not only accumulate in plaque macrophages, but is also engulfed by circulating and splenic monocytes. Despite earlier observations of this phenomenon in murine and human monocytes by Mosig *et al*^{32,33} and Tolani *et al*³⁴ the functional implications of this intracellular lipoprotein accumulation and its consequences for atherosclerosis development are not recognized completely.

Traditionally, LDL cholesterol is thought to preferentially accumulate in the arterial intima at places with reduced shear stress³⁵. This process is followed by the influx of monocytes/macrophages, which scavenge the subendothelial oxLDL. We could now show that next to the cholesterol accumulation, monocyte-endothelial cell interaction is also enhanced in animals with increased cholesterol levels. As this increment was only present

after activation of the endothelium by LPS stimulation, it is believed to reflect an intrinsically increased adhesive capacity of lipid-laden monocytes and this increased recruitment of monocytes to inflammatory vascular loci (e.g. atherosclerotic lesions). Nevertheless, hypercholesterolemia leads to endothelial cell dysfunction³⁶, potentially contributing to increased monocyte rolling and adhesion. Our findings thus point out that cholesterol might also enter the atherosclerotic lesion via migration of lipid-laden circulating monocytes into the atherosclerotic lesion.

Intriguingly, our data indicate that hypercholesterolemia leads to a rapid monocytoxis response, as well as an increase in plasma levels of pro-inflammatory cytokines within days after introduction of WTD. However, after three weeks of WTD plasma levels for the majority of cytokines appeared to normalize at least in part, suggesting that sustained hypercholesterolemia does not lead to prolonged hyper-inflammatory status. This indicates that hyperinflammation represents an early adaptive response to new metabolic conditions. In human disease, postprandial cholesterol and lipid levels are associated with cardiovascular events^{37,38}. As we did not measure time points beyond three weeks or included reversal of diet to normal chow until now, it is still unclear to what extent cellular and systemic inflammatory and metabolic pathways affect systemic, or possibly local, inflammatory processes involved in early atherogenesis. Furthermore, the effects of these interventions on the regulation of cytokine levels in persistent hypercholesterolemia are to be determined. Recent evidence suggests that even mild reduction in cholesterol levels of hypercholesterolemic subjects can reduce plasma cytokine levels back to normal³⁹. Nevertheless, more research on the interaction between cholesterol and cytokine levels is necessary to unravel the exact pathways and cell types responsible.

Circulating monocytes as biomarker for cardiovascular disease

Monocytes and monocyte-derived proteins are still underrepresented amongst biomarkers for disease and this holds in particular for CVD, despite the critical roles exerted by this subset in its pathology, as we have shown in our murine atherosclerosis studies. As indicated in the introduction only a limited number of monocyte-derived biomarkers have been suggested thus far, such as CD163 and IL-6 secreted by peripheral blood mononuclear cells ^{40,41}. Besides these secreted proteins, non-secreted membrane markers also have been proposed to have diagnostic value, e.g. monocyte HLA-DR expression in acute liver failure ⁴² and sialic acid-binding Ig-like lectin 1 (Siglec-1) and Fcγ receptor I (CD64) expression in systemic lupus erythematosus ^{43,44}. However, for cardiovascular disease, hardly any monocyte biomarkers have been described, apart from plakoglobin ⁴⁵. As monocytes play a critical role in the development of atherosclerosis and also have a patrolling function of the vasculature, they conceivably represent a rich and important source of diagnostic markers of disease status and prospect which legitimates more in depth searches for monocyte-derived biomarkers. In this thesis, we show that both on protein level (Chapter 5) and RNA level (Chapter 6) activated monocytes are encouraging sources of novel biomarkers for cardiovascular disease.

In **Chapter 5** a whole cell phage display approach was applied for the discovery of novel membrane bound biomarkers on circulating monocytes from unstable angina pectoris patients. Although whole cell phage display is not a standardized technique, it has already been successfully applied for biomarker discovery ⁴⁶, especially in cancer research, where small tumor specimens were used as target ⁴⁷⁻⁵⁰. However, also cell lines ⁵¹ and isolated platelets ⁵² have been considered as scaffold for phage display selections. Here, we applied whole cell phage display selection on LPS-activated THP-1 monocytes and validated the markers found on isolated CD14⁺ monocytes from unstable angina pectoris patients. To favor selection of relevant clones, we incorporated a depletion step using monocytes from healthy volunteers. Despite limited enrichment for clones binding UAP monocytes, monocyte-based selections were proved to be efficient as we identified two clones with increased affinity for UAP monocytes. In addition, these clones showed their value for imaging, since they specifically bound monocytes in advanced, vulnerable atherosclerotic lesions, but not early lesions. Overall, from our study, monocytes emerge as a promising target for antibody discovery, in spite

the large variety in surface proteins expressed on these cells. Our study also implies that circulating monocytes indeed can reflect local vascular processes such as atherosclerosis.

Inspired by these promising findings, we zoomed in on differences in RNA expression profiles between SAP, UAP, and healthy control CD14⁺ monocytes in **Chapter 6**. Surprisingly, we could not detect any major differences between the study populations based on micro-array analysis. This is surprising as several groups have previously identified UAP-specific markers⁵³⁻⁵⁵, albeit that none of these markers were cell-bound markers. These data suggest that secretome, rather than transcriptome, difference are present between patient populations. Still, Patel *et al* revealed that monocyte-platelet complexes did also not differ between SAP and UAP patients⁵⁶ and differential expression of membrane receptors was also shown for UAP^{57,58}. Even though our results did not show major differences in the transcriptome of UAP versus SAP monocytes, we are the first to investigate mRNA expression profiles of monocytes from SAP and UAP patients in a large patient cohort. In contrast to previous publications, we have studied differences in circulating CD14⁺ monocytes in a patient cohort without any known cardiovascular disease, whereas others have studied total peripheral blood mononuclear cells⁵⁹, specific patient populations (e.g. systemic lupus erythromatous patients⁵⁷), or plaque macrophages⁶⁰. Taken together, our study primarily focused on the discovery of diagnostic markers for discrimination between SAP and UAP patients, based on peripheral blood monocytes.

Interestingly, our studies on the proteome (chapter 5) and transcriptome (chapter 6) do not accord completely. On the protein level we were able to identify altered surface expression on UAP monocytes compared to control monocytes using phage display. However, our phage display approach used LPS-activated THP-1 monocytes for target discovery and UAP monocytes for validation. This approach might have boosted the discrimination power of our phage display analysis. Nevertheless, LPS is a strong stimulus for TLR-signaling, which is not specific to UAP, but also is linked to various infectious and non-infectious diseases⁶¹. Preliminary data on micro-array analysis from LPS-activated SAP monocytes with and without a CVD event during follow-up indicates that differences are present between the two study populations. However, the observation of differences within the

population of SAP patients might also point to heterogeneity of SAP and UAP patients, contributing to the lack of separation between SAP and UAP patient monocyte transcriptomes. Still, the above suggests that activation potential, rather than basal expression levels, of UAP monocytes might be different from SAP monocytes. Indeed, first analysis of LPS activated monocytes from patients with event during follow-up compared to event-free controls, revealed an extensive list of differentially expressed genes. More in depth research on SAP and UAP patient monocytes using a broad 'omics' approach should reveal novel UAP specific monocytic biomarkers useful for imaging, diagnosis, and potentially even risk prediction.

Figure 1 - Overview of the role of circulating monocytes in atherosclerosis

Circulating monocytes contribute to atherosclerosis development at different levels. In **Chapter 2** we showed the role of circulating monocytes in combination with plaque macrophages in regulating plaque progression, by induction of CD115⁺ cell apoptosis. Regulation of circulating monocyte chemotaxis via GRK2 towards the atherosclerotic lesion was discussed in **Chapter 3**. Interaction of circulating monocytes with cholesterol and the atherosclerotic lesion as well as the effects thereof on proteome and transcriptome biomarkers were discussed in **Chapter 4** and **Chapters 5 and 6**, respectively.

Concluding Remarks and Future Perspectives

The role of circulating monocytes in atherosclerosis was investigated in this thesis. We were able to show that monocyte migration as well as monocyte/macrophage apoptosis are indeed essential processes in atherogenesis. Next, hypercholesterolemia was proven to influence circulating monocyte function rapidly after onset hypercholesterolemia. Finally, we have explored circulating monocytes as a novel target for current, but also future biomarker research. Here, we identified activated monocytes as a potential marker for diagnosis and imaging of patients at risk for CVD, both on protein and RNA level. Taken together, monocytes not only appear to play a central role in regulating plaque progression, but are also useful biomarkers for diagnosis, imaging, and possibly even prognosis of cardiovascular disease.

Circulating monocytes are thus central actors in atherogenesis. We were able to identify these cells not only being essential for plaque initiation and progression, but also pinpoint novel opportunities for CVD biomarkers. Nevertheless, additional research on activation potential of monocytes will contribute to both unraveling mechanism involved in monocyte function and additional information on current and future disease and risk for cardiovascular events, such as myocardial infarction and stroke. Future research will need to focus on more in depth knowledge regarding the translation from local information in the atherosclerotic lesion to the circulating monocytes, to determine additional markers for diagnosis, risk prediction, and imaging. Further, screening methods which can reliably and successfully identify these subtle yet important differences need to be optimized facilitating minimally invasive and rapid measurements of these circulatory biomarkers.

Circulating monocytes in atherosclerosis: Local or systemic actors?

The central question of this thesis was whether circulating monocytes only impact atherosclerosis at a local level or also systemically. Our data indicate that indeed both peripheral and local monocyte functions are relevant for atherosclerosis development. Locally, monocyte chemotaxis and monocyte/macrophage ablation influence plaque development as we have shown. Lipid-loading of circulating monocytes not only could contribute to cholesterol accumulation in the lesions, but also alter monocyte function systemically. Nevertheless, influencing monocytes systemically will not only affect plaque development, but also monocyte functionality (e.g. patrolling behavior). In the end circulating monocytes have the potential to be used as novel biomarker for CVD, as we have shown in this thesis, indicating the link between local vascular pathology and systemic effects on monocytes.

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Summary

Cardiovascular diseases are still one of the leading causes of death in the Western societies and account for over 17 million deaths world-wide annually. Atherosclerosis is the predominant underlying pathology responsible for various cardiovascular diseases.

Atherosclerosis is characterized by the accumulation of low density lipoprotein (LDL) cholesterol in the intima of arteries. Oxidation of intimal cholesterol will result in activation of the vessel wall and will trigger a pro-inflammatory cascade. Due to the release of chemokines and upregulation of integrins on the endothelial cells covering the early lesion leukocytes will be attracted in an effort to resolve the immune response and remove the cholesterol. In the first stage, predominantly monocytes will migrate into the early atherosclerotic lesion, and give rise to plaque macrophages which will sequester the local oxidized cholesterol. Excessive intra-cellular cholesterol accumulation combined with ineffective excretion results cholesterol filled foamy macrophages also known as foam cells. Apoptosis of foam cells combined with ineffective clearance of apoptotic cells, will result in buildup of cell debris and (oxidized) cholesterol in the lesion, forming a lipid-rich, necrotic core, covered by a fibrous cap of smooth muscle cells and collagen. Destabilization of the fibrous cap by proteases secreted by macrophages and other inflammatory cells in the lesion, can eventually result in plaque rupture, exposing the pro-thrombotic content of the lesion to the circulation, potentially resulting in clinical complications, such as myocardial infarction and stroke.

Monocytes and plaque macrophages are instrumental in the development of atherosclerosis, from initial early until advanced vulnerable lesion. The role of circulating monocytes in relation to atherosclerosis development is not completely elucidated, despite the importance of these subsets in the development of the disease. Therefore, this thesis investigated the role of circulating monocytes in atherosclerosis, to determine (I) whether apart from plaque residing monocytic cell pools, the peripheral monocytes are able to impact disease onset and progression, and (II) whether these cells have potential as biomarker for cardiovascular disease in humans.

The function of circulating monocytes as well as plaque macrophages can be studied by specific depletion of these cell types. In **Chapter 2**, we studied the impact of systemic versus plaque-targeted ablation of myeloid cells on atherosclerosis in ApoE^{-/-} mice, bearing a Macrophage FAS Induced

Apoptosis (MaFIA) suicide gene. Local ablation of CD115 positive plaque macrophages aggravated atherosclerosis, as both necrotic core and plaque size were expanded significantly by promoting either clonal expansion of plaque macrophages or monocyte influx. In contrast, systemic ablation of CD115 positive myeloid cells did not result in changes in plaque burden but promoted a more vulnerable plaque phenotype. Interestingly, it was associated with enhanced extramedullary hematopoiesis and mobilization of immature myeloid cells into the circulation. Our data suggest that the augmented extra-medullary myelopoiesis and the increased mobilization of immature myeloid cells into the circulation, upon systemic induction of monocyte/macrophage apoptosis may have underlain the observed increase in plaque vulnerability.

As migration of leukocytes towards the lesion is a key process in atherogenesis, we describe, in **Chapter 3**, the role of chemotaxis regulation on atherosclerosis development. Chemokine receptors are G protein-coupled receptors (GPCR) and are responsible for the directional migration of cells along a chemokine gradient. The activity of GPCRs is controlled by receptor kinases, such as GRK2. We here studied the effect of partial hematopoietic GRK2 deficiency on atherosclerosis development in LDL receptor knock-out mice. Hematopoietic GRK2^{+/-} deficiency resulted in marked attenuation of plaque development. To exclude that macrophage and granulocyte GRK2 was responsible for the effects observed, we used a LysM-cre GRK2^{flox/flox} conditional knock-out model which results in macrophage and granulocyte specific GRK2 deficiency. As atherosclerosis development was not affected in this model, we concluded that monocytes were most likely responsible for the phenotype observed in the GRK2^{+/-} study and that even partial GRK2 deficiency already was able to prevent atherosclerotic lesion development beyond the fatty streak stage.

Next to inflammation, hypercholesterolemia is one of the key processes in atherosclerosis development and a well-known risk factor for cardiovascular disease, and evidence is culminating that both actors are tightly associated. In **Chapter 4**, we analyzed the interaction between plasma cholesterol and circulating monocytes in LDL receptor deficient mice to unravel the effects of hypercholesterolemia on monocyte function and numbers in early atherosclerosis. We could not only show that under hyperlipidemic conditions, circulating monocytes carry lipoprotein depositions already before entering the lesion, and were able to link this to alterations in monocyte

function. In fact, cholesterol-laden monocytes exhibited increased reactive oxygen species production and rolling along and adhesion to activated endothelium. Combined with the inflammatory cytokine profile observed in these animals, hypercholesterolemia was shown to affect circulating monocytes already in very early phases of atherosclerosis development and might reveal novel pathways contributing to intimal cholesterol accumulation.

As circulating monocyte subsets have a patrolling function in the vasculature we studied the biomarker potential of human circulating monocytes. We analyzed the clinical perspective of using circulating monocytes for disease detection, as well as therapy, both on a proteome and transcriptome level.

In **Chapter 5**, we identified at least two markers which may yield potential as diagnostic biomarker in unstable angina pectoris, as well as in molecular imaging approaches. Phage display affinity selections were performed on activated THP-1 monocytes to identify clones of interest. Subsequently, these selected clones were validated on CD14 positive monocytes isolated from healthy controls and unstable angina pectoris patients, as well as on early stable and advanced vulnerable plaque tissue, identifying at least two clones with high specificity.

In **Chapter 6**, we analyzed the transcriptome of monocytes isolated from stable and unstable angina pectoris patients to identify biomarkers specific for unstable angina pectoris. The patient populations did not differ in terms of monocyte markers or chemokine receptor expression as determined by flow cytometry. Surprisingly, we could also not detect any major differences in the transcriptome of these monocytes between stable and unstable angina pectoris, despite major differences in clinical parameters, and as reported plasma cytokine patterns. These data indicate that at baseline mild acute cardiovascular syndromes such as unstable angina pectoris do not impact the transcriptome of circulating monocytes.

Finally, in **Chapter 7** the findings presented in this thesis are discussed and future perspectives are indicated. In conclusion, we were able to confirm that monocytes play an important role in atherosclerotic lesion development. Monocyte migration towards the lesion as well as numbers and function of plaque macrophages were shown to be critical in atherogenesis, but also the interaction between plasma cholesterol and circulating monocytes was identified to have importance in disease development. Furthermore,

circulating monocytes convey information from local sites, such as atherosclerotic plaques, to the periphery. Indeed, we were able to show that circulating monocytes harbor potential as diagnostic biomarker for at least some cardiovascular pathologies. Future research will need to focus on identification of specific targets as well as on investigation of the prospective of activation potential of circulating monocytes in relation to biomarker discovery.

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Vereenvoudigde Nederlandse Samenvatting

Hart- en vaatziekten zijn nog altijd een van de meest voorkomende doodsoorzaken in de Westerse landen en zijn verantwoordelijk voor meer dan 17 miljoen doden per jaar wereldwijd. Aderverkalking, ook wel atherosclerose genoemd, is één van de belangrijkste onderliggende oorzaken verantwoordelijk voor de gevolgen van hart- en vaatziekten, zoals hersen- en hartinfarct.

Atherosclerose is een ziekte van de middelgrote en grote slagaders, waarbij cholesterol zich opstapelt in de vaatwand door een ontregeling van de cellen die de vaatwand bedekken op plaatsen waar de bloedstroom verstoord is, zoals bochten en vertakkingen van de vaatboom. Door de ophoping van cholesterol ontstaat er een ontstekingsreactie in de vaatwand die ervoor zorgt dat witte bloedcellen worden aangetrokken. In eerste instantie zullen vooral monocyten, dit is een type witte bloedcel, naar de vaatwand migreren. In de vaatwand zullen deze cellen zich verder ontwikkelen en vormen ze zogenaamde macrofagen, die het cholesterol dat aanwezig is in de vaatwand zullen gaan opnemen. Door de grote hoeveelheid cholesterol die zich ophoopt in de cellen zullen deze een schuimachtig uiterlijk krijgen, waardoor ze schuimcel genoemd worden. Als deze schuimcellen te veel cholesterol hebben opgenomen, zullen ze dood gaan en zal het opgenomen cholesterol, maar ook de resten van de gestorven cellen achterblijven in de vaatwand. Er ontstaat hierdoor uiteindelijk een cholesterol-rijke kern die middels een kapsel bestaande uit collageen en gladde spiercellen wordt afgescheiden van het bloed: de atherosclerotische plaque. Afbraak van het kapsel dat over de plaque ligt, kan resulteren in het scheuren van het kapsel, waardoor het prothrombotische materiaal in de plaque wordt blootgesteld aan het bloed en er een bloedstolsel zal ontstaan. Het is dit bloedstolsel dat door afsluiting van het bloedvat kan leiden tot zuurstofgebrek in het achterliggende weefsel, waardoor een hart- of herseninfarct ontstaat.

Monocyten in het bloed en macrofagen in de plaque zijn belangrijke cellen in de ontwikkeling van atherosclerose, zowel in beginnende plaques als ook in meer gevorderde stadia van het ziekteproces. De rol van monocyten in het bloed in relatie tot de ontwikkeling van atherosclerose is nog niet volledig opgehelderd, ondanks dat deze cellen worden gezien als een zeer belangrijk celtype tijdens de ontwikkeling van deze ziekte. Mijn onderzoek richtte zich op de rol van deze in het bloed circulerende monocyten in de ontwikkeling van atherosclerose, waarbij ik getracht heb om te achterhalen hoe deze cellen de

onderliggende processen beïnvloeden die ten grondslag liggen aan atherosclerose. Het uiteindelijke doel van dit onderzoek was te bepalen of deze cellen gebruikt kunnen worden als biomarker voor hart- en vaatziekten.

De functie van circulerende monocytten, maar ook macrofagen in de plaque, kan effectief worden bestudeerd door deze cellen te elimineren. In **Hoofdstuk 2** hebben we aangetoond dat circulerende monocytten plaque ontwikkeling kunnen afremmen in een muismodel waarin bepaalde witte bloedcellen, zoals monocytten en macrofagen, na toediening van een chemische stof, verwijderd worden. Eliminatie van zowel plaque macrofagen alsook circulerende monocytten resulteerde niet in veranderingen in plaque ontwikkeling vergeleken met de controle. Wel werd er een verhoogde productie van witte bloedcellen (hematopoëse) waargenomen, waarbij er mogelijk sprake was van een toegenomen migratie van monocytten naar de plaque. Als alleen de plaque macrofagen werden verwijderd, resulteerde dit in een toename van de plaque ontwikkeling. Samenvattend hebben we kunnen aantonen dat plaque ontwikkeling beïnvloed kan worden door de beschikbaarheid van circulerende monocytten alsook de aanwezigheid van macrofagen in de plaque, waarbij de verstoorde productie van deze cellen een belangrijke rol speelt.

In **Hoofdstuk 3** beschrijven we de rol van gerichte migratie van cellen naar een stimulus, en de regulatie van dit proces in atherosclerose. Chemokine receptoren zijn verantwoordelijk voor deze migratie van cellen en ze behoren tot de familie van G eiwit gekoppelde receptoren. De activiteit van deze receptoren wordt gereguleerd door bepaalde enzymen, zoals GRK2. We hebben het effect van gedeeltelijke uitschakeling van GRK2 in witte bloedcellen onderzocht in pro-atherogene knock-out muizen. Gedeeltelijke uitschakeling van GRK2 resulteerde in zeer sterke remming van atherosclerose ontwikkeling. Uitschakeling van dit enzym in granulocyten en macrofagen, naast monocytten zijn dit andere witte bloedcellen die GRK2 aanmaken, liet geen effect zien op de ontwikkeling van atherosclerose, wat suggereert dat GRK2 in (circulerende) monocytten verantwoordelijk is voor de remming van atherosclerose.

Naast ontstekingsreacties is ook een verhoogd cholesterol niveau betrokken bij de ontwikkeling van atherosclerose en een risicofactor voor de ontwikkeling van hart- en vaatziekten. In **Hoofdstuk 4** hebben we de interactie tussen cholesterol in het bloed en circulerende monocytten onderzocht. We

konden aantonen dat naast het in de plaque opgehoopte cholesterol ook het in het bloed aanwezige cholesterol van belang is voor de ontwikkeling van atherosclerose, omdat dit wordt opgenomen door circulerende monocytten. Hierbij worden bepaalde functies van deze monocytten ook beïnvloed. Cholesterol-geladen monocytten produceren meer zuurstofradicalen en hechten nog beter dan normaal aan de ontstoken bloedvatwand. Samen met een verhoogde concentratie van bepaalde ontstekingsmediatoren in het bloed, die worden waargenomen na het volgen van een cholesterol/vetrijck dieet dragen deze veranderingen in de monocytten bij aan het ontstaan van atherosclerose. De toegenomen hechting van deze ontstekingscellen aan de vaatwand in combinatie met de ophoping van cholesterol in deze circulerende monocytten, zijn een nieuwe route waarlangs cholesterol vanuit het bloed in de vaatwand terecht kan komen, waardoor uiteindelijk een atherosclerotische plaque zal ontstaan.

Aangezien circulerende monocytten ook een functie hebben als verkenner van de bloedvaten, hebben we bestudeerd of deze cellen potentie hebben als marker voor hart- en vaatziekten in de mens. We hebben zowel het eiwit als het DNA niveau bestudeerd om deze nieuwe markers te vinden. In **Hoofdstuk 5** hebben we minstens twee markers kunnen identificeren die potentie hebben als diagnostische marker voor instabiele angina, oftewel pijn op de borst in rust, maar daarnaast ook mogelijkheden bieden om gebruikt te worden bij detectiemethodes die gebruik maken van beeldtechnieken. Bindende antilichamen werden geselecteerd met behulp van faag display selecties op geactiveerde humane monocytten (de THP-1 cellijn). Bij faag display worden antilichamen gekoppeld aan virusdeeltjes, waarmee de antilichamen die binden aan de monocytten geselecteerd kunnen worden door deze te laten reageren met de monocytten en niet-bindende antilichamen weg te wassen. Hierna werden de geselecteerde antilichamen getest op hun capaciteit om te binden aan monocytten geïsoleerd uit patiënten met instabiele angina, maar ook aan monocytten in vroege en meer gevorderde atherosclerotische plaques. Minstens twee antilichamen bleken met hoge voorkeur te binden aan deze cellen en zijn mogelijk geschikt voor de diagnose van hart- en vaatziekten en mogelijk zelfs voor het opsporen van gevorderde plaques.

In **Hoofdstuk 6** hebben we getracht om biomarkers te identificeren waarmee onderscheid gemaakt kan worden tussen stabiele angina (allen bij inspanning pijn-op-de-borst) en instabiele angina (ook tijdens rust pijn-op-de-borst) patiënten. Allereerst hebben we gekeken of er verschillen aanwezig waren tussen de hoeveelheid eiwitten die aanwezig zijn op de buitenkant van de monocytten. We konden aantonen dat de monocytten van deze twee patiëntgroepen hierin niet van elkaar verschilden. Vervolgens hebben we gekeken naar de mate waarin informatie van het DNA door de cel gebruikt wordt om eiwitten te produceren en of hierin verschillen aanwezig waren tussen de patiëntgroepen. Tegen onze verwachtingen in konden we ook voor deze eigenschappen geen verschillen ontdekken tussen monocytten van stabiele angina patiënten en instabiele angina patiënten. Onze studie toont daarmee aan dat er geen grote verschillen aanwezig zijn in circulerende monocytten van stabiele en instabiele angina patiënten.

Tot slot worden in **Hoofdstuk 7** de resultaten, gepresenteerd in dit proefschrift, bediscussieerd en worden toekomstige mogelijkheden voor onderzoek aangegeven. Samenvattend toont dit onderzoek aan dat circulerende monocytten essentieel zijn voor de ontwikkeling van atherosclerose. We hebben kunnen aantonen dat de migratie van monocytten naar de plaque, maar ook de hoeveelheid macrofagen in de plaque kritieke processen zijn in de ontwikkeling van atherosclerose. Daarnaast tonen onze studies aan dat cholesterol al in de bloedsomloop de functie van de circulerende monocytten ongunstig beïnvloedt, een tot nu onderbelicht mechanisme bij het ontstaan van atherosclerose. Circulerende monocytten weerspiegelen de lokale situatie zoals die in de plaque is. Deze informatie kan gebruikt worden, zoals we hebben laten zien in hoofdstuk 5, voor de diagnose van hart- en vaatziekten. Toekomstig onderzoek is noodzakelijk om niet alleen biomarkers te identificeren waaraan de door ons ontdekte antilichamen binden, maar ook om de perspectieven van het gebruik van andere eigenschappen van de monocytten zoals hun activatiepotentieel als marker voor hart- en vaatziekten in kaart te brengen.

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Curriculum Vitae

Curriculum Vitae

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Best Poster Presentation

3e Cardiovasculaire Conferentie
Nederlands Lipoproteïne Club Session
Noordwijkerhout, The Netherlands
14 - 15 March 2013

Oral Presentations**3e Cardiovasculaire Conferentie**

Nederlandse Lipoproteïne Club Session
Noordwijkerhout, The Netherlands
14 - 15 March 2013

Scandinavian Society for Atherosclerosis Research

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American Heart Association (AHA)

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Poster Presentations

European Society for Clinical Investigation

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17 - 20 April 2013

World Immune Regulation Meeting VI

Innate and Adaptive Immune Response and Role of
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Davos, Switzerland

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1st Cardiovasculaire Conferentie

Noordwijkerhout, The Netherlands

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ATVB Early Career Networking Reception

16 November 2010

Chicago, IL, USA

Publications

Otten JJ*, Medina I*, Bermudez B, Wolfs I, Rademakers T, Wijnands E, Schurgers LJ, Biessen EA. Local but not systemic CD115⁺ myelocyte ablation aggravates atherogenesis. *Submitted*.

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List of Abbreviations

ABC	ATP-binding cassette
ABCA	ATP-binding cassette transporter
AMI	Acute myocardial infarction
ANOVA	Analysis of variance
APC	Allophycocyanin
Apo	Apolipoprotein
BMDM	Bone marrow-derived macrophage
BMI	Body mass index
BNP	B-type natriuretic peptide
CCL	CC-motif chemokine ligand
CCR	CC-motif chemokine receptor
CD	Cluster of differentiation
CFSE	Carboxyfluorescein succinimidyl ester
CMP	Common myeloid progenitor
CPT	Camptothecin
CRP	C-reactive protein
CSF-1R	Colony stimulating factor 1 receptor
CTMM	Center for translational molecular medicine
CVD	Cardiovascular disease
CX3CR	CX3C-motif chemokine receptor
CXCL	CXC-motif chemokine ligand
CXCR	CXC-motif chemokine receptor
DAB	3,3-Diaminobenzidine
DCFDA	2',7' -dichlorofluorescein diacetate
DNA	Deoxyribonucleic acid
ECG	Electro cardiogram
EDTA	Ethylenediaminetetraacetic acid
Egr	Early growth response protein
ELISA	Enzyme-linked immunosorbent assay
EM	Electron microscopy
EMP	Erythrocyte/megakaryocyte progenitor
ER	Endoplasmatic Reticulum
FACS	Fluorescence-activated cell sorting
FCS	Fetal calf serum
FDR	False discovery rate
FITC	Fluorescein isothiocyanate
GAPDH	Glyceraldehyde 3-phosphate dehydrogenase

G-CFU	Granulocyte colony forming unit
G-CSF	Granulocyte colony stimulating factor
GDF	Growth differentiation factor
gDNA	Genomic deoxyribonucleic acid
GFP	Green fluorescent protein
GM-CFU	Granulocyte/macrophage colony forming unit
GM-CSF	Granulocyte/macrophage colony stimulating factor
GMP	Granulocyte/macrophage progenitor
GPCR	G protein-coupled receptor
GRK	G protein-coupled receptor kinase
H&E	Haematoxylin and eosin
HDL	High density lipoprotein
hDTR	Human diphtheria toxin receptor
HRP	Horse radish peroxidase
hsTNT	High sensitive troponin T
i.p.	Intraperitoneal
i.v.	Intravenous
ICAM	Intercellular cell adhesion molecule
IFN	Interferon
IL	Interleukin
IRF	Interferon regulating factor
Klf	Kruppel-like factor
LBP	LPS-binding protein
LCM	L929-conditioned medium
LDL	Low Density Lipoprotein
LDLr	Low density lipoprotein receptor
LPS	lipopolysaccharide
MaFIA	Macrophage FAS induced apoptosis
M-CFU	Macrophage colony forming unit
MCP1	Monocyte chemo-attractant protein 1
M-CSF	Macrophage colony stimulating factor
M-CSFr	Macrophage colony stimulating factor receptor
MHC	Major Histocompatibility Complex
MMP	Matrix metallo-protease
MPLSM	Multi-photon laser scanning microscopy
MPO	Myeloperoxidase
mRNA	Messenger RNA

NF-κB	Nuclear factor κB
NHF	Netherlands Heart Foundation
NOX	NADPH oxidase
NPC	Niemann-Pick disease type C protein
oxLDL	Oxidized low density lipoprotein
PBL	Peripheral blood lymphocytes
PBMC	Peripheral blood mononuclear cell
PBS	Phosphate buffered saline
PCA	Principal component analysis
PE	Phycoerythrin
PEG	Poly(ethylene) glycol
PerCP	Peridinin chlorophyll protein complex
PFA	Paraformaldehyde
PMT	Photo-multiplier tube
RIN	RNA integrity number
RNA	Ribonucleic acid
ROS	Reactive oxygen species
RT	Room temperature
SAP	Stable angina pectoris
sCD40L	Soluble CD40 ligand
SEM	Standard error of the mean
Siglec	Sialic acid-binding Ig-like lectin
SMA	Smooth muscle cell actin
SMC	Smooth muscle cell
sPLA ₂	Secreted phospholipases A2
SSC	Side scatter
STEMI	ST-elevated myocardial infarction
TEA	Triethylamine
TGF	Transforming growth factor
TLR	Toll-like receptor
TNF	Tumor necrosis factor
TnT	Troponin T
TRAP	Tartrate resistant acid phosphatase
TUNEL	Terminal deoxytransferase dUTP nick-end labeling
UAP	Unstable angina pectoris
VCAM	Vascular cell adhesion molecule
VLA	Very late antigen

VLDL	Very low density lipoprotein
VSMC	Vascular smooth muscle cell
WT	Wild type
WTD	Western type diet

The image features a white background heavily decorated with red ink splatters. These splatters vary in size, from tiny dots to large, elongated streaks and blotches. The distribution is dense, particularly in the lower half of the frame, where the red marks overlap and create a complex, textured appearance. The word "Dankwoord" is printed in a dark, bold, sans-serif font, positioned in the upper right quadrant of the image. The overall effect is one of dynamic, expressive movement, characteristic of a hand-drawn or spray-painted artwork.

Dankwoord

Na ruim 4 jaar werken en het schrijven van ongeveer 75.000 woorden in deze thesis, zijn dit voor veel van jullie waarschijnlijk de eerste woorden die je leest. Het is en blijft een apart idee dat iedereen het meest geïnteresseerd is in de persoonlijke verhalen aan het einde van een thesis, maar hier doe ik zelf eigenlijk ook altijd aan mee.

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Hoewel ik geen officiële co-promotor heb gehad tijdens mijn promotie, heb ik wel van één iemand in het bijzonder zeer veel steun en vriendschap mogen ontvangen, hè opper-paranimf! Beste Timo (of moet ik zeggen TimoPedia) jij bent een van de redenen geweest waardoor ik in deze helse wereld ben

beland die men ook wel “research” noemt. En ik moet zeggen, ik heb er geen moment spijt van gehad. Aangezien we het grootste gedeelte van de afgelopen 4 jaar op dezelfde kamer hebben doorgebracht, is onze vriendschap alleen maar hechter geworden, en de 7-jaar regel hebben we duidelijk gebroken. Zoals je ook al in een van je stellingen omschreef: “I never did give anybody hell. I just told the truth and they thought it was hell.” Ik weet zeker dat genoeg mensen die met ons samen een kamer hebben gedeeld, dit zeker beamen. Daarbij hebben we enkele keren samen een congres bezocht, waarbij Denemarken wel één van de mooiste congressen, lees: uitstapjes ;-) was. Helaas, zullen onze professionele werelden zich vanaf nu enigszins scheiden, maar ik weet zeker dat we in de toekomst nog vele dingen samen zullen beleven. Ik wens je veel succes met je weg naar Professor Timo, die je waarschijnlijk eerst naar het Hoge Noorden zal leiden, maar ik ben ervan overtuigd dat de toekomst je toch wel weer een keer naar het Zuiden terug zal brengen en anders weet ik je in mijn nieuwe functie ook wel weer te vinden!

Beste Stephan, we kennen elkaar al bijna 30 jaar en hoewel je niet uit het wetenschapswereldje komt, heb je het aangedurfd om mij als paranimf te ondersteunen, waar ik heel erg blij mee ben. We hebben in de lange tijd die we elkaar kennen zeer veel samen meegemaakt, in ieder geval veel te veel om zelfs maar in het kort hier neer te schrijven. Wel weet ik zeker dat ik op jou kan terugvallen, en dat is iets wat voor mij heel belangrijk is. Binnen de Ealèniers vormen we de laatste jaren ook al een mooi team binnen het bestuur, en drinken we allebei graag een pilsje. Je beland dankzij mij nu even in een wonderbaarlijke wereld, maar met ons carnavalsverleden zijn we natuurlijk heel wat gewend! Ik wens jou en Britta een hele mooie toekomst samen toe en kijk uit naar de volgende 30 jaar en alles wat die tijd ons zal brengen.

Binnen de EVP zijn er natuurlijk nog veel meer mensen die ik moet, maar vooral ook wil bedanken.

Ow FACS-koning...;-) Beste Erwin, jij was een van de personen binnen de afdeling die al wist wie ik was voordat ik ook maar was begonnen. Via alle kanalen die er zijn met het Orbis werd er veel informatie uitgewisseld, waardoor mijn komst al was aangekondigd. De afgelopen jaren hebben we extreem veel uren samen gebrainstormd over alles wat maar met FACS (of wat dan ook) te maken had. Hoewel ik denk dat ik je zeker ook wel één of twee

dingen heb kunnen leren, heb ik vooral ongelooflijk veel van jou geleerd. Uiteindelijk zijn we zelfs nog samen op vakantie geweest met Orbis naar Oostenrijk en voor het werk naar Zwitserland, waarbij we tussendoor zelfs nog een congres hebben bezocht. Ik weet zeker dat de monocyte-studie nog tot een mooie publicatie zal leiden. Voor de toekomst hoop ik dat we nog veel ski-tripjes kunnen maken, waarbij je wellicht in de toekomst iets minder lang op mij hoeft te wachten... Veel geluk gewenst voor de toekomst samen met Linda!

Beste Mat, hartelijk dank voor al je hulp met het ML-1 werk. Je was de afgelopen 4 jaar voor mij iemand op wie ik altijd kon rekenen als ik iets wilde weten over faag display, maar ook PCR en celkweek behoorden tot jouw kennisgebied. We zijn ook nog een paar keer naar Dortmund geweest om treintjes te bekijken, want dat is toch een mooie hobby die we delen. Ik hoop dat je binnenkort een kamer in huis mag gaan gebruiken om ook je eigen spoor weer te kunnen opbouwen. Ik reserveer in ieder geval voor komend jaar het dagje Duitsland weer in mijn agenda!

Beste Karen, we hebben ongeveer even lang op de afdeling rondgelopen en zijn nu ook bijna tegelijk klaar met onze thesis. Hoewel onze onderzoeken wat verder uit elkaar lagen, hebben we heel vaak samen bij de muizen of in het lab gewerkt. Ik hoop dat de toekomst jou vol geluk toelacht, maar ik weet zeker dat jij goed terecht komt, zowel privé als op het werk.

Lieve Lieve (sorry maar blijft leuk), ondanks dat je later dan een groot gedeelte van de andere mensen binnen de groep bent gekomen, denk ik dat je een van de personen bent die niet alleen voor top onderzoek zorgt, maar ook voor een hele hoop gezelligheid en plezier. Ik heb ook erg genoten van alle keren dat je samen met Lucas naar de borrels kwam, jong geleerd is oud gedaan, hè! Voor mij was verder het schminken in Aachen wel een hoogtepunt. Heel veel respect dat je me liet klieren! Veel geluk voor de toekomst samen met Pieter, Lucas en Marieke.

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im einer der top Labos der Welt arbeiten solltest. Ich hoffe das wir uns noch öfter sehen und vielleicht könnten wir nochmals Schifahren machen, oder am wenigsten ein Bier (und Jägermeister) trinken. Viel Glück und bis Bald!

Beste Ine, sinds je naar de patho gekomen bent, heb je een zware tijd gehad, geloof ik. Je moest immers een kamer delen met onder andere Timo en mij, en wij zijn nu eenmaal nooit rustig hè. Wel heb je met succes deelgenomen aan onze cursus, waarvoor nogmaals mijn complimenten. Ik wil je ook danken voor alle hulp die je hebt geboden, met het uitvoeren van allerlei experimenten. Zelfs al was je dag drie keer volgepland, dan nog vond je altijd ruimte om een ander en dus ook mij te helpen. Ik hoop dat je thesis snel volgt en geloof me maar als ik zeg dat dit allemaal goed gaat komen. Veel succes met je verdere carrière en ik kom graag nog eens BBQ-en.

Dear David, I still do not know which language I should choose when we are talking. However, most often we just stuck to English. We have spent a lot of time together in the lab but also in our room. And I have to say that it became quiet after you left, no more German swearing or loud music coming from your ears. Ich möchte Ihnen für die letzten Jahre bedanken und wünschen Ihnen viel Glück bei Ihrer nächsten Herausforderung in der Forschung. En voor een biertje ben ik altijd wel te bereiken!

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Onze kamer heeft de laatste jaren ook een behoorlijke buitenlandse invloed gehad. Naast de Belgische enclave, was er ook een duidelijk Duitse invloed.

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Jeroen

*Als je de richting van de wind niet kunt veranderen,
verander dan de stand van je zeilen*