

Development of new radio-labeled tracers for imaging and quantification of brown adipose tissue

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**Development of new radio-labeled tracers for imaging and
quantification of brown adipose tissue**

Andreas Paulus

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Development of new radio-labeled tracers for imaging and quantification of brown adipose tissue

DISSERTATION

to obtain the degree of Doctor at the Maastricht University,

on the authority of the Rector Magnificus,

Prof. Dr. Rianne Letschert

in accordance with the decision of the Board of Deans,

to be defended in public on Wednesday 24.04.2019, at 12:00 o'clock

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Table of Content

	Abbreviations	6
Chapter 1	Introduction	9
Chapter 2	Brown adipose tissue and lipid metabolism imaging	27
Chapter 3	Characterization of BAT activity in rats using invasive and non-invasive techniques	55
Chapter 4	Synthesis, radiosynthesis and in vitro evaluation of [¹⁸ F]Bodipy-C16/triglyceride as a dual modal imaging agent for brown adipose tissue	87
Chapter 5	[¹⁸ F]BODIPY-triglyceride-containing chylomicron-like particles as an imaging agent for brown adipose tissue in vivo	113
Chapter 6	Triglyceride-rich lipoprotein derived fatty acid uptake by brown adipose tissue in control and diabetic mice at room temperature, acute cold exposure and cold acclimation	139
Chapter 7	Discussion	165
	Valorization	177
	Acknowledgement	183
	Curriculum Vitae	187
	List of publications	189

Abbreviations

Chronological appearance in the thesis:

BAT	Brown Adipose Tissue
WAT	White Adipose Tissue
UCP1	Uncoupling Protein 1
TG	Triglyceride
FA	Fatty Acid
CD36	Cluster of Differentiation 36
FATP	Fatty Acid Transport Proteins
PET	Positron Emission Tomography
[¹⁸ F]FDG	[¹⁸ F]Fluoro-2-deoxy-2-D-glucose
LCFA	Long Chain Fatty Acid
TRL	Triglyceride Rich Lipoprotein
LPL	Lipoprotein Lipase
SPECT	Single Photon Emission Computed Tomography
MRI	Magnetic Resonance Imaging
CT	Computed Tomography
ACSL	Fatty acyl-CoA Synthetase
CoA	Coenzyme A
PTHrP	Parathyroid Hormone-related Protein
[¹⁸ F]FTHA	14(R,S)-[¹⁸ F]fluoro-6-thia-heptadecanoic acid
[¹¹ C]palmitate	[¹¹ C]hexadecanoic acid
[¹²³ I]BMIPP	15-(4-[¹²³ I]iodophenyl)-3-methyl-pentadecanoic acid
BODIPY- C ₁₆	4,4-Difluoro-5,7-Dimethyl-4-Bora-3a,4a-Diaza-s-Indacene-3-Hexadecanoic acid
[¹²³ I]IHXA	[¹²³ I]iodohexadecanoic acid

[¹²³ I]IHDA	[¹²³ I]iodoheptadecanoic acid
[¹²³ I]IPPA	15-(p-[¹²³ I]iodophenyl)-pentadecanoic acid
BODIPY	Boron-dipyrrromethene
QD	Quantum Dots
iBAT	interscapular Brown Adipose Tissue
ADIFAB	Acrylodan labelled rat intestinal fatty acid binding protein
[^{99m} Tc]TcMIBI	[^{99m} Tc]Tc-2-methoxy-isobutyl-isonitrile
[¹²³ I]MIBG	meta-[¹²³ I]iodobenzylguanidine
[¹²³ I]IPA	[¹²³ I]iodophenylalanine
MRS	Magnetic Resonance Spectroscopy
TR-1	Temperature Regime 1
MIBI	Sestamibi
VOI	Volume of Interest
SUV	Standardized Uptake Value
FLASH	Fast Low Angle SHot
iAT	interscapular Adipose Tissue
cBAT	cervical Brown Adipose Tissue
ipWAT	intraperitoneal White Adipose Tissue
BDP-FA	Bodipy-C ₁₆
BDP-TG	Bodipy-triglyceride
ESI-Q	Electrospray Ionization Quadrupole
Ppm	parts per million
RCY	Radiochemical Yield
NE	Norepinephrine
SO	Sulfosuccinimidyl-oleat
r.t.	room temperature
p	probability leve

PVAT

Perivascular Adipose Tissue

Chapter 1: Introduction

Brown Adipose Tissue (BAT) in mammals

BAT is uniquely found in mammals and enables them to produce heat to maintain their body temperature below their thermoneutral zone ¹. In contrast to typical white adipose tissue (WAT), BAT consists of multiple cellular lipid droplets for storage purposes which facilitates the opportunity of fast lipolysis due to its enhanced lipid surface ². BAT is also distinguished by its high number of mitochondria where its brown color originates from. Its high degree of vascularisation provides heat transport throughout the whole body in times of thermogenesis ³.

BATs thermogenic potential results from the BAT specific uncoupling protein 1 (UCP1) ⁴⁻⁷. During activation high amounts of internal triglycerides (TGs) are lipolysed and transported towards the mitochondria where usually β -oxidation leads to ATP synthesis. UCP1 uncouples this process by permeabilizing the inner mitochondrial membrane for protons whereby heat is produced ^{1,8}.

The ability to “burn” fatty acids (FAs) instead of storing energy makes BAT an interesting target in understanding the development of obesity and potentially fighting its related metabolic syndromes.

FA metabolism

Obesity is considered as a worldwide epidemic which will become even more severe in the near future. Already in 2016 1.9 billion adults were overweight from which 650 million were obese. Obesity results from a positive energy balance and is correlated to other diseases such as diabetes type-II ^{9,10}, atherosclerosis ¹¹ and cancer ¹², together referred to as metabolic syndrome.

FAs are the major source of energy and either derived from food intake or lipogenesis that includes glycolysis and the citric acid cycle ¹³. In animals predominantly FAs with even numbers can be found and FAs are grouped by their chain length, short chain (2–6 carbon atoms), medium chain (8–12 carbon atoms), long chain (14–18 carbon atoms) and very long chain (20–26 carbon atoms). The main type of circulating FAs are long-chain and very long-

chain with varying degrees of saturation. FAs are stored as TGs in adipocytes and set free in times of starvation or fasting ¹⁴.

Most types of tissue can switch between glucose and FA metabolism. A major tissue for energy expenditure is muscle because of its relative large size ^{15,16}. Other tissues such as heart also show this ability but because of its relative small size the heart can only increase its contribution to whole body energy expenditure to 5-10% ¹⁷. Significant amounts of glucose are cleared by the liver after a meal but also FAs are taken up, oxidised and incorporated as TGs into lipoproteins to transport them into other tissues for storage purposes ^{18,19}.

Uptake of FAs into cells was long thought to be the result of simple diffusion. Recently it was found that FA uptake is dependent on specific transport proteins such as cluster of differentiation 36 (CD36) or fatty acid transport proteins (FATP) ²⁰. Once in the cell FAs can be stored in lipid droplets or used in the mitochondria. The main pathway for FA degradation is mitochondrial β -oxidation ²¹ by which ATP is produced. It is important to mention that by a complete oxidation of one FA molecule e.g. palmitate more molecules ATP are produced than by an oxidation of one glucose molecule (105 vs. 31) ²².

Even though a high number of guidelines by national and international authorities has been published on healthy diets and physical activity, to fight against obesity and related metabolic syndromes, studies could show that patients only adhered short times and a recidivism to old behaviour patterns was the common case ²³⁻²⁶

Therefore BAT with its ability to uncouple its ATP production is an interesting target to increase overall body energy expenditure and to reduce weight of obese subjects.

Brown adipose tissue (BAT)

BAT research evolved within the last 20-40 years, revived in 2009 and is now an important topic in the field of endocrine research. Active BAT in adult humans was found using retrospective positron emission tomography (PET) studies with [¹⁸F]Fluoro-2-deoxy-2-D-glucose ([¹⁸F]FDG) ²⁷⁻²⁹. As already mentioned BAT has the unique ability to uncouple its

ATP production in its mitochondria by the BAT specific protein UCP1 and to produce heat during this process ⁴⁻⁷.

In the absence of heat production protons which are transported against the concentration gradient cause a voltage difference across the inner mitochondrial membrane. Protons which reenter the matrix by ATPase release their energy, which is necessary for ATP synthesis. During BAT activation protons enter the matrix by UCP1 ³⁰ and energy is released as heat ^{1,8}. This process is responsible for nonshivering thermogenesis ^{1,31}.

It was shown that long chain fatty acids (LCFA) (> 12 carbon atoms ³²) are of immense importance for the uncoupling process as they work as a carrier for protons through UCP1 ³³. Purine nucleotides like ATP inhibit UCP1 from the cytosolic side ^{8,34-36}. LCFA can overcome the blocking of UCP1 and support the activation of the uncoupling process ^{1,33,37}.

BAT is activated by cold exposure of the subject ³⁸⁻⁴⁰. Thermoreceptors in the skin are stimulated by cold, those stimuli are transmitted by the somatic nervous system and neurons are activated in the ventromedial nucleus of the hypothalamus resulting in release of norepinephrine by the postsynaptic nerve endings, triggering BAT activation ^{1,41}. Norepinephrine binding to β 3-adrenoceptors on BAT activates cyclic adenosine monophosphate, which in the next step stimulates protein kinase. Hormone sensitive lipase is activated resulting in TG lipolysis and FAs are released which are necessary for UCP1 activation ^{1,42,43}. By norepinephrine activation TGs stored in intracellular lipid droplets are lipolyzed to single FAs and metabolized in the mitochondria. Lipid droplets can then be potentially replenished in three different manners: glucose uptake with following *de novo* lipogenesis, free FA uptake facilitated by albumin binding, and uptake of triglyceride-rich lipoprotein (TRL)-derived FAs from the plasma with subsequent esterification to TGs ^{1,44-46}. TRL-derived FAs were identified as the main supply of TGs in BAT ⁴⁴ and FA internalization was shown to be dependent on the presence of lipoprotein lipase (LPL) ^{47,48}, cluster of CD36 ⁴⁹ and FATP ⁵⁰ which need to be located at the outer cellular membrane.

Increased [¹⁸F]FDG uptake as a measure for glucose consumption in BAT was recently observed in obese cold acclimated humans concluding that significant amounts of BAT can

be recruited during repeated cold exposure⁵¹. Glucose is mainly taken up by protein transporters of the GLUT family, mainly the fat muscle specific isoform GLUT4^{52,53}. Glucose is processed in different pathways e.g. in the citric acid cycle⁵⁴ or it is converted to FAs^{55,56}. The main metabolized substance class in BAT is FAs. Therefore BAT activity can be largely underestimated by [¹⁸F]FDG scans, which is till now the most often used technique to quantify BAT activity and volume. The aim of this work was to develop a lipid based tracer to gain more information and to quantify FA metabolism in BAT because till now it is not known to what extent activated BAT contributes to energy consumption in obese.

PET and single photon emission computed tomography (SPECT)

The era of tracer based non-invasive imaging was a huge progress for the society of nuclear medicine. Two prominent examples, which are frequently used to visualize processes *in vivo*, are PET and SPECT. Both rely on the same idea of radioactive material which is injected into a living organism and its distribution in the body can be followed up at later time stages. In PET β^+ -emitting radionuclides are used where the following pair of annihilation radiation can be measured in coincidence by a detector ring. By this technique a high amount of unspecific signals can be filtered out which makes it possible to precisely localise the point of the annihilation and has a positive effect on resolution and sensitivity. In SPECT γ -emitting radionuclides are used. Here a collimator is necessary to gain information about the direction of the emitted γ -ray. Because of this filter only a minor amount of the emission reaches the camera which makes SPECT less efficient in comparison to PET. A more recent approach is the combination of PET or SPECT with other imaging modalities such as computer tomography or magnetic resonance imaging. Next to the tracer distribution also information about morphological structures are obtained which help to better localize the site of the tracer accumulation.

Tracers and Isotopes

As described in the previous section PET and SPECT imaging offer the opportunity to gain information about the biodistribution of a radioactive drug by non-invasive imaging. This can help to identify for example disease patterns, cancer, injuries or metabolic activity. The design of the radioactive drug or also called tracer is of immense importance and varies for different tasks. In all cases a carrier molecule, which should target the region of interest is modified by a radionuclide. Sometimes the addition of the radionuclide is not trivial and can only be achieved by precursor molecules or chelators. Usually the radiolabeled drug is applied in amounts which have no pharmacological effect. This is the reason why they are also called tracer or radio-tracer.

PET radionuclides and tracer

Various PET suitable radionuclides exist and most of them have a half life comparable to the biological half life of the radiotracer. In Tab. 1 the most common radionuclides are listed. The primary mode of decay should be positron emission. Although many different radionuclides could be used in theory, carbon-11 and fluorine-18 are the most prominent ones. All organic compounds in nature consist of carbon atoms. Therefore an exchange of stable carbon-12 by carbon-11 has no effect on the pharmacological properties of a compound. Fluorine usually cannot be found in biomolecules but, its size is similar to the size of the hydroxyl group which implies only a minor change in steric properties. Fluorine is the atom with the highest electronegativity. Therefore introduction of fluorine can cause a change in electronic properties which might have an advantageous effect on the binding affinity⁵⁷. Its relative long half life compared to e.g. carbon-11 and the lowest decay energy of all PET radionuclides makes fluorine-18 the radionuclide with the best nuclear properties.

Table 1: PET radionuclides

Radionuclide	Half Life	Decay Mode	Max. Decay Energie [MeV]
¹¹ C	20.3 min	β ⁺	0.97
¹³ N	10 min	β ⁺	1.20
¹⁵ O	2 min	β ⁺	1.74
¹⁸ F	110 min	β ⁺	0.64
⁶⁴ Cu	12.7 h	β ⁺ /electron capture	0.66
⁶⁸ Ga	67.6 min	β ⁺ /electron capture	1.90
⁷⁶ Br	16.2 h	β ⁺ /electron capture	4.00
¹²⁴ I	4.18 d	β ⁺ /electron capture	2.14

For diagnostics, the most often used tracer is [¹⁸F]FDG. This glucose analog was developed in the 1970's ⁵⁸ and its synthesis was improved further a decade later ⁵⁹. Its field of application reaches from oncologic questions to neurology. Because of its broad application, [¹⁸F]FDG can have some limitations such as high uptake in brain, bladder and in inflamed tissue. Therefore more dedicated tracers have been developed e.g. [⁶⁸Ga]GaPSMA ⁶⁰, [¹⁸F]FLT ⁶¹ or [¹¹C]PK11195 ⁶². This selection of tracers is only a small list and should represent important fields of PET imaging. It does not reflect the enormous number of tracers which are available today.

SPECT radionuclides and tracer

In SPECT imaging radionuclides which directly emit γ-rays are used. In Tab. 2 the most important nuclides used in SPECT are listed. It is to mention that SPECT isotopes have in general a longer half life than PET isotopes. Therefore longer and excessive radiosynthesis can be conducted and longer processes in vivo can be visualized. The longer half life offers also the possibility to transport the radiopharmaceuticals to other institutions, adding an important commercial aspect to imaging with SPECT isotopes. The most important radionuclide is technetium-99m. Different to the other isotopes it does not need to be produced in a cyclotron because it is produced during the decay of its mother nuclide

molybdenum-99. This generator approach led to the label “workhouse of nuclear medicine” for technetium-99m.

Table 2: SPECT radionuclides

Radionuclide	Half Life	Decay Mode	Principal Photon emission
			energy [MeV]
¹²³ I	13.2 h	Electron capture	0.16
^{99m} Tc	6 h	Isomeric transition	0.14
¹¹¹ In	67.9 h	Electron capture	0.17/0.25
⁶⁷ Ga	78.3 h	Electron capture	0.09/0.19/0.30
²⁰¹ Tl	73.1 h	Electron capture	0.17

To label a biomarker with technetium-99m a chelating agent is necessary. Technetium-99m chemistry relies on complex chemistry to design a precursor molecule which has the ability to allow labelling with technetium-99m but also retain its binding affinity *in vivo*. The use of bulky chelating agents limits the application of technetium-99m-labeled compounds e.g. in neuroscience because the blood brain barrier could preclude target binding.

Another important SPECT radionuclide is iodine-123. Similar to fluorine-18, iodine forms covalent bonds with carbon atoms, making it easy to incorporate it into small molecules. A disadvantage compared to incorporation of fluorine-18 into a molecule is that iodine has a larger van der Waal’s radius and therefore higher steric alteration can be expected, especially for smaller molecules. The fact that iodine has several isotopes that can be used in nuclear medicine (β and γ emitters) offers the advantage to use iodine labeled compounds for imaging and therapy with the same molecule. An important agent for this purpose is MIBG where [¹²³I]MIBG is used to visualize neuroendocrine tumors and patients with MIBG positive areas obtain a therapy with [¹³¹I]MIBG ⁶³. Other important SPECT tracers are [¹²³I]FP-CIT ⁶⁴ a tracer to target the dopamine transporter in the brain or [^{99m}Tc]TcMAG3 which is used for renography ⁶⁵.

Tracers for BAT imaging

PET suitable tracer

As already mentioned [¹⁸F]FDG is the most frequently used tracer so far which might be a result of a high number of retrospective studies as well as its availability. It was found in dedicated experiments that age, obesity and various pathological conditions have a negative effect on detectable BAT volume^{38,66-69}. Uptake of [¹⁸F]FDG is also influenced by outdoor temperatures, fasting, and different medications e.g. β -blockers^{28,43,70}. Apart from quantification of BAT [¹⁸F]FDG is also used to explore BATs role in glucose metabolism. Lower amounts of active BAT were found in hyperglycaemic and diabetic patients^{71,72}.

Different fatty acid based tracers such as [¹⁸F]FTHA⁷³, [¹¹C]palmitate⁷⁴ and others⁷⁵⁻⁷⁷ have been developed. Most of them have been used for myocardial imaging, before their suitability to visualize BAT was considered to be useful. A more detailed discussion about lipid based tracers can be found in chapter 2.

[¹¹C]acetate is able to quantify FA synthase activity by visualizing BATs oxidative metabolism⁷³. In studies with healthy test subjects exposed to cold a significant [¹¹C]acetate increase in uptake was observed⁷⁸. In addition oxygen-15 tracers are also used to determine BATs oxidative capacity.

Studies with [¹⁵O]O₂ showed an increased metabolic BAT activity in cold exposed subjects⁷⁹. [¹⁵O]H₂O as a perfusion tracer, was able to show an increase in perfusion due to cold⁸⁰. Obesity attenuates this effect⁸¹.

[¹¹C]MRB as a highly selective ligand for the norepinephrine transporter was found to accumulate in BAT. This elevated uptake values could be already reached at room temperature while cold exposure only slightly increased [¹¹C]MRB uptake⁸².

In a retrospective study with patients scanned for pheochromocytoma by [¹⁸F]F-DA, a dopamine analog, it was found that BAT could be visualized in 18% of the patients⁶⁹. A similar occurrence was achieved by [¹⁸F]FDG scans. However it is to mention that those

patients might have a larger BAT tissue mass or activation because of increased levels of catecholamines excreted from their tumor.

Apart from the here presented tracer with a clinical application, a high number of promising pre-clinical tracers exist. Prominent examples are radiolabeled triphenylphosphonium based tracers targeting the voltage difference of the inner mitochondrial membrane^{83,84}.

SPECT suitable tracer

[¹²³I]MIBG is an FDA approved tracer for pheochromocytoma, neuroblastoma and cardiac imaging. Its biodistribution is known to be dependent on catecholamine production⁸⁵. In retrospective studies a prevalence of finding fat depots which are linked to BAT was 18%⁶⁹ and in preclinical studies BAT uptake was increased by 3 fold due to cold exposure⁸⁶. In dedicated clinical studies in healthy adults [¹⁸F]FDG uptake correlated with [¹²³I]MIBG in BAT⁸⁷.

[^{99m}Tc]TcMIBI is taken up in tissue with an increased blood flow and high mitochondrial density due to its lipophilic and cationic properties. An increased [^{99m}Tc]TcMIBI uptake in BAT vs. WAT was reported in a preclinical study where uptake in BAT was increased by adrenergic stimulation. In retrospective human studies BAT was detectable^{88,89} and BATs presence in [^{99m}Tc]TcMIBI positive samples was confirmed by immunohistochemistry⁸⁸.

[¹²³I]BMIPP^{90,91} is another FA based tracer which was initially developed for myocardial imaging. Similar to [¹⁸F]FTHA it is now used to study BAT uptake of FAs.

Another lipophilic tracer which originally was developed for cardiac imaging is [^{99m}Tc]TcTetrofosmin. Due to its lipophilic properties it is enriched in tissue with high amounts of mitochondria. In a retrospective study it was found that its prevalence to detect BAT depots in humans was 17%⁹² making it a suitable tracer to explore BATs mitochondrial function.

Other BAT detection techniques

Apart from the presented nuclear tracer there are other techniques suitable to visualize BAT and its metabolism. The possibilities to image BAT are here presented very briefly. More detailed information are given in chapter 2.

A high number of preclinical work is performed with fluorescent *in vitro* uptake experiments⁹³⁻⁹⁷. The chance of analyzing BAT uptake on a sub-cellular scale, working without patients or animals and without exposure to radioactivity makes this method suitable for a high throughput of experiments.

Next to imaging with radioactive tracers or fluorescent imaging there is also the possibility to image BAT with labelled particles. This particle visualization can consist of fluorescent labeled particles⁴⁵, radioactive labeled particles^{46,98} or MR functional particles⁴⁵. Incorporation of the tracer in particles might have beneficial effects on the biodistribution of the tracer itself. E.g. a bare FA based tracer relies on several uptake mechanism before it can be transported via the bloodstream and might therefore not reflect the real processes *in vivo*. For more detailed information see chapter 2.

Other methods to determine BATs volume are Magnetic Resonance Imaging (MRI) and Computed Tomography (CT). MRI is able to determine the water/lipid content in BAT by detecting a radio-frequency of excited protons in a magnetic field. This frequency is dependent on the chemical surrounding of the excited protons and is therefore tissue specific. By comparing it to previous records it is possible to gain information about BATs activation state and lipid consumption^{99,100}. In comparison to PET, MRI has the advantage that it can detect following properties of BAT: temporal and quantitative measurements of tissue fat content, changes in blood flow, volume, and oxygenation of active BAT¹⁰⁰. Additionally MR Spectroscopy is able to detect the relative abundance of chemicals within a single localized region and therefore helps to quantify changes in BATs lipid composition due to stimulation. CT on its own is almost not able to detect BAT. As BAT and its surrounding tissue do not differ too much in its Hounsfield units a standalone CT cannot distinguish between e.g. BAT and WAT next to each other. Also the interpretation of Hounsfield units is

biased ¹⁰¹⁻¹⁰⁴. Anyway, CT and MRI are useful additions to the common used ways to visualize BAT by adding morphological information to the image.

Aim of this dissertation

Multiple imaging and quantification approaches for BAT are available. The most often used technique is PET/CT with [¹⁸F]FDG. Its broad field of application as well as its use in clinical routine makes this tracer very popular, also in the field of BAT imaging. Nevertheless [¹⁸F]FDG has a huge disadvantage: it is a glucose based tracer. Although [¹⁸F]FDG was able to visualize BAT and to show increased metabolic activity in cold exposed animals and humans, lipids in the form of FAs are the main metabolized substance class. Therefore it is impossible to calculate BAT's total metabolic activity.

At first FA based tracers, such as [¹⁸F]FTHA or [¹²³I]BMIPP, were used to quantify BATs its demand of FAs. However, this method does not completely reflect the situation *in vivo* where FAs are transported as TGs incorporated in TRLs towards BAT.

Therefore the aim of this dissertation was to synthesize a radiolabeled TG and to incorporate it into a TRL which than can be applied *in vivo*. To our understanding this would come closest to the *in vivo* situation and will probably help to understand BATs role in whole body energy expenditure. Additionally such a tracer would also be useful for other applications (e.g. fatty liver disease, obesity and fatty acid metabolism in general).

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Chapter 2: Brown adipose tissue and lipid metabolism imaging

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Abstract

Purpose: Brown adipose tissue (BAT) research has evolved from an underestimated to a fast developing field. Its assumed curing properties for the world wide epidemic obesity, and its related diseases, makes this tissue an interesting target for a broad amount of non-invasive molecular BAT tracers. Apart from [¹⁸F]FDG PET/CT there are several methods to detect BAT and measure its metabolism in a more appropriate way. Especially interesting is the measure of lipid turnover, because fatty acids comprise the main fuel for active BAT. This review outlines different imaging modalities suitable for BAT imaging with the overall goal to explain the yet not completely understood mechanism in BAT and its quantitative contribution to whole body lipid and energy metabolism.

Methods: Publications with focus on brown adipose tissue and lipid metabolism imaging are analyzed, different imaging approaches are introduced and promising BAT tracers are presented.

Results: Radiolabelled and fluorescent fatty acids, labelled particles, [³H]Triolein and ADIFAB staining can give information about the inflow and therefore about the utilization of fatty acids which represents the activation state *in vivo/in vitro*. Non-invasive scanning with CT or MRI is a useful addition to those techniques.

Conclusion: Lipid metabolism imaging offers the opportunity to visualize and quantify yet undiscovered aspects of BAT metabolic activities and is key to completely clarify its role in whole body lipid and energy metabolism.

Introduction

Background:

The function and presence of BAT in adults was neglected until two decades ago. Now the investigation of BAT using molecular imaging has matured to one of the most interesting and fast developing research topics in endocrine research. Because obesity - and its related metabolic syndrome - is reaching epidemic proportions in the western world ¹ and may even become a more severe problem in the near future for the worldwide population, more attention is drawn to adipose tissue metabolism. A turning point was the discovery that WAT, apart from storing energy (fat), is able to secrete leptin, an important hormone controlling the energy balance ². Other substances ³ (autocrine and endocrine) released by WAT have been found and by that the consideration of the impact of adipose tissue on whole body metabolism rose continuously. In retrospective PET studies with [¹⁸F]FDG, it could be shown that besides WAT, another form of adipose tissue exists in adult humans ⁴⁻⁶. In these studies, symmetrical accumulations of [¹⁸F]FDG appeared in the supraclavicular region, which were originally thought to be attributed to uptake in cervical muscles. Later, scans with PET/CT indicated that these “artifacts” correlate with Hounsfield units of fat ⁶. By these [¹⁸F]FDG studies it could be proven that BAT is functionally present in adults and is metabolically active. BAT, named after its darker color resulting from higher mitochondria expression within the cell and increased blood circulation ⁷, was previously thought to be absent in adult humans and only be present in newborns to maintain their body temperature ⁸. The results of the PET scans indicate a chance of observing BAT in 5 - 8% of standard clinical routine PET scans ^{4,9} and a total contribution to body mass of 0,05 – 0,01% ¹⁰. These findings could be confirmed later by dedicated cold exposure studies where a direct correlation between cold exposure and BAT metabolic activity, measured through [¹⁸F]FDG uptake, was reported ¹¹⁻¹³. Assuming a fixed relative contribution of glucose and fatty acids and that mainly fatty acids and glucose contribute to energy expenditure ^{12,14}, an increase in metabolic activity of BAT would result in an increase of total body energy expenditure of 2-28% ¹⁵. Therefore activation

of BAT with unchanged food uptake, may lead to significant weight loss, offering an additional treatment option to obese patients. Another field of application would be in patients with pheochromocytoma. It was found that catecholamine secreting tumors activate BAT and lead to an increase in metabolic activity characterized by FA and glucose uptake¹⁶⁻¹⁸. In these studies BAT activity was inversely correlated to body mass index and in general patients with cancer cachexia are often suffering from body weight loss and depletion of muscular and adipose tissue¹⁹.

However, in order to actually calculate the metabolic activity of BAT in humans, in addition to [¹⁸F]FDG PET/CT, other tracers and techniques are needed. Through the upcoming interest in this tissue, several new activation, targeting and imaging strategies have been developed to visualize BAT's functions. [¹⁸F]FDG has set the stage; several attributions of BAT have been defined. Non-invasive visualization of lipid metabolism could give more insight since lipid turnover is one of the major features of BAT. This review summarizes early as well as the newest inventions of lipid metabolism imaging linked to BAT and describes how visualization of lipid fate and adipose tissue activity has evolved.

Activation of brown adipose tissue and fatty acid metabolism:

First observations of BAT in humans were performed with [¹⁸F]FDG⁴⁻⁶ which visualizes metabolic active tissue in terms of glucose uptake. It was observed that through cold exposure the chance of visualizing BAT increased in animals and lean subjects¹¹⁻¹³. Besides, it was shown that BAT is activated through binding to β 3-adrenoceptor^{20,21}. Recently it was observed for the first time that BAT takes up an increased amount of glucose in obese humans after cold acclimation. This led to the conclusion that significant amounts of BAT can be recruited during repeated cold exposure²². Glucose is mainly taken up by protein transporters of the GLUT family, mainly the fat muscle specific isoform GLUT4^{23,24}. Glucose is processed in different pathways e.g. in citric acid cycle²⁵ or it is converted to FA^{26,27}. Nevertheless LCFA are the main "fuel" for BAT. Tracers based on fatty acids, quantify the metabolic activity of BAT in a better way than glucose does.

Through the norepinephrine activation process FA stored as TG in lipid droplets are consumed in the mitochondria to produce heat and new FA are taken up^{28,29}. This identifies FA as the main metabolized substance in BAT and makes them and other compounds targeting lipid metabolism a powerful tool to visualize BAT and its functions within the body.

Usually mitochondria oxidize fatty acids and ATP is produced to store the nascent energy but BAT mitochondria contain the BAT specific UCP1 which gives them the ability to uncouple the oxidation process and to produce heat instead of ATP³⁰⁻³³. This process is responsible for nonshivering thermogenesis^{15,34}. Fedorenko et al. showed that LCFA (> 12 carbon atoms³⁵) are essential for the uncoupling process as they work as a carrier for H⁺ through the UCP1 and that LCFA are also produced within the inner mitochondria membrane by PLA2³⁶. In BAT cells UCP1 is inhibited by purine nucleotides, mainly ATP. The UCP1 channel is blocked by the nucleotides from the cytosolic side³⁷⁻⁴⁰. It was shown that LCFA can overcome the blocking of UCP1 and support the activation of the uncoupling process^{15,36,41}. Activation through LCFA also confirms the observation that lipolysis activates thermogenesis in the same order of magnitude than norepinephrine does^{42,43}.

As quoted before, stored TGs in BAT can be utilized for combustion or FA can be taken up after lipolysis by BAT. Uptake of FA from external sources by a brown adipocyte is realized by protein mediated transport⁴⁴, nevertheless the exact mechanism of FA uptake is unclear. FA are transported in TRLs (Triglyceride-rich lipoproteins) by the blood. BAT uptake of FA via internalization of these particles, via direct FA uptake after lipolysis related breakup of TRLs or a combination of both is possible. Khedoe et al. equipped TRLs of different size with [³H]oleate and [¹⁴C]cholesteryl oleate⁴⁵. At room temperature a lipolysis supported uptake could be observed as cholesteryl oleate stays within the particle core due to its hydrophobicity and most taken up activity in BAT resulted from [³H]oleate. Left over remnant particles, containing a higher ratio of [¹⁴C]cholesteryl oleate are cleared by the liver which explains why a much higher carbon-14-activity compared to tritium-activity can be observed in the liver. Also cold exposure or thermoneutrality did not promote internalization of TRLs in

BAT cells in an important amount but of course changed the amount of FA taken up by BAT⁴⁵.

In contrast to these experiments is the finding of Bartelt et al. They labelled TRLs with [³H]-oleate and hydrophobic [⁵⁹Fe]SPIO nanocrystals⁴⁶. As a result of lipolysis [³H]oleate can enter the cell whereas [⁵⁹Fe]SPIO nanocrystals stay inside of the TRL's. In cold exposed mice a significant faster clearance of [³H]oleate and [⁵⁹Fe]SPIO and a tenfold higher uptake of both imaging agents in BAT is reported. Through intravital microscopy a rapid attachment to the endothelium, followed by an internalization of the TRLs could be observed⁴⁶. Bartelt et al. explain their observations that through lipolysis products the endothelial barrier function is decreased and TRLs can enter the cell⁴⁷.

The discrepancy between both experiments may be explained by different imaging agents used and therefore different uptake mechanisms are needed to explain the results. Nevertheless, the dominant uptake mechanism (lipolysis assisted uptake of FA or internalization of TRLs) is not determined at the moment and needs to be established by other imaging approaches and experiments.

In terms of lipolysis, LPL is the first of multiple proteins involved in the process of FA uptake into BAT^{29,48,49}. Other important proteins related to FA uptake of BAT are fatty acid translocase (FAT/CD36)⁵⁰, fatty acid transport proteins (FATPs)⁵¹ and fatty acyl-CoA synthetases (ACSL)⁵². The common way of FA uptake is described as in figure 1, according to⁴⁸.

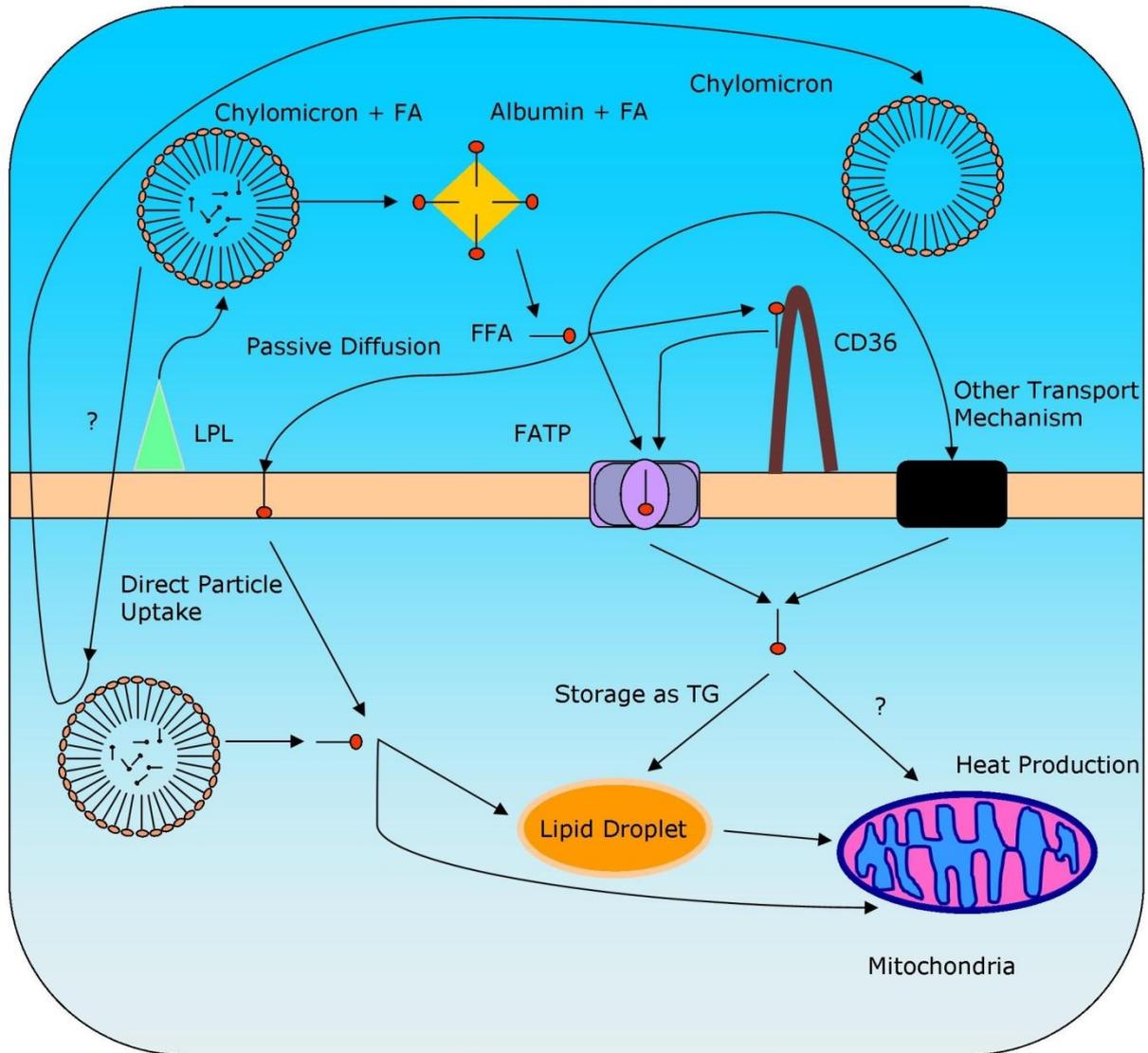


Figure 1: TRLs are hydrolyzed by LPL and FA can bind to albumin. After dissociation from albumin, FA bind to membrane proteins such as CD36 or directly to FATP. A direct diffusion through the membrane is possible but dependent on free FA concentrations and FA already taken up by the cells⁵³. FA from CD36 are handed to FATP and are transported through the membrane where they are either processed as FA or activated and coupled to coenzyme A (CoA) by ACSL. Free FA can then be stored in lipid droplets as TG after esterification or directly processed in the mitochondria.

Different studies have identified LPL and CD36 as indispensable factors for FA uptake and a loss of one of these proteins results in an impaired adipose metabolism leading to hypertriglyceridemia, increased glucose uptake and hypothermia during fasting and cold exposure^{54,55}.

LCFA can also be produced through lipogenesis within the cell by fatty acid synthase up to a carbon chain length of 16 carbon atoms ²⁹. Elongation to very long chain fatty acids is done by very long chain fatty acids enzymes (ELOVL) where ELOVL3 was found to be overexpressed in cold exposed BAT and mice lacking of ELOVL3 were only able to survive cold by shivering thermogenesis ⁵⁶. These findings indicate lipogenesis as an essential feature to ensure the TG amount necessary during cold exposure.

Its high amount of mitochondria and multilocular lipid droplets make BAT appear more like muscle than adipose tissue ¹⁵.

White adipose tissue, fat storage and endocrine function

The main function of WAT is fat storage and fatty acid secretion. When energy intake is higher than consumption, fatty acids are stored in WAT. In times of fasting or cold exposure, WAT is activated to release its fuel (fatty acids) by stimulation of lipolysis through the sympathetic nervous system ⁵⁷. Through the 90's of the last century the interest in WAT changed from a tissue only responsible for energy storage to a tissue that also takes part in body metabolism. Its physiological role has changed through the discovery of the first protein secreted from WAT: leptin ². More and more proteins have been discovered that are responsible for different functions in the body ^{3,58,59}. Leptin e.g. is the critical hormone in energy balance and is produced principally by white fat, giving the tissue an endocrine function. Other proteins are: angiotensinogen, adiponin, acylation-stimulating protein, adiponectin, retinol-binding protein, tumour necrosis factor α , interleukin 6, plasminogen activator inhibitor-1 and tissue factor. These proteins can work as inflammatory cytokines, some influence the lipid metabolism and others are involved in vascular haemostasis ⁶⁰.

Beige/brite adipose tissue:

A special type of adipose tissue is the so called beige (or brite) adipose tissue. Characteristics of the white adipocyte are a large lipid droplet and a small amount of mitochondria. BAT has multiple small lipid droplets, its transcriptional profile is similar to

skeletal muscle cells ⁶¹ and BAT has a large amount of mitochondria which overexpress UCP1. Beige adipocytes can differentiate *in vivo* from WAT or WAT precursor cells through activation processes (e.g. β 3-adrenergic receptor agonists) and then contain multiple lipid droplets and express UCP1 within their mitochondria, which potentially makes them an important player in increasing energy expenditure and fighting against obesity ⁶². This clusters of beige cells appear e.g. through prolonged cold exposure in WAT depots close to noradrenergic nerve fibers which trigger the beige adipocyte recruitment ⁶³. Typically norepinephrine is released from the sympathetic nerves and activates adipocyte (brown and beige) thermogenesis. This process normally induces UCP1 expression through phosphorylation of PPAR γ coactivator 1 α and other activators ^{64,65}. Sidossis *et al.* investigated the hypermetabolism in patients with burn injuries and found increased circulating levels of epinephrine and norepinephrine as well as elevated expression of UCP1 in the according WAT-areas as a result of adrenergic stress ⁶⁶. This proves that epinephrine and norepinephrine are responsible for browning of WAT which makes this tissue able to participate in the process of non-shivering thermogenesis. Alternatively, M2 macrophages are recruited to subcutaneous WAT during cold exposure and secret catecholamines to activate WAT browning ^{67,68}. Irisin, an exercise-induced myokine, can also induce browning of WAT ^{69,70}. This paradox finding could be shown *in vitro* and *in vivo* ⁶⁹ and it has been hypothesized that this mechanism evolved from shivering muscle contraction to support non-shivering thermogenesis by an increase in BAT volume and has thereby a therapeutic potential to treat obesity ^{69,70}. Another important player in WAT browning is cancer cachexia. Through tumor-derived parathyroid hormone-related protein (PTHrP) WAT browning is promoted and more energy is needed in a Lewis lung cancer model ⁷¹. Deactivation of PTHrP inhibits WAT browning and prohibits energy wastage ⁷¹. Besides of the listed factors for browning there are also environmental factors, endocrine hormones and transcriptional mediators which can cause browning of WAT ⁶³. Beige adipose tissue cells are only temporarily available for thermogenesis as they develop back to WAT-cells after activation.

By other studies it could be confirmed that beige adipocytes almost reached UCP1 levels of BAT and that beige fat mitochondria are thermogenically active ⁶¹.

Form of Fatty acids (free fatty acid/ Triglyceride/ LDL-particle)

The main suppliers of FAs taken up by BAT are TRLs (i.e. chylomicrons or very-low-density-lipoprotein-particles) circulating in the blood ⁷². After incorporation into adipocytes, FAs are stored as TGs in lipid droplets. Lately LPL-activity was found to be important for whole body TRL clearance through BAT and WAT ^{46,73}. Savonen et al. recently found a correlation between LPL distribution and type of tissue where the FA are taken up and identified LPL as a mandatory binding side for LDL particles to be processed (lipolysis with exposure of FA) ⁷⁴.

Fatty acid imaging

As already mentioned, fatty acids are the main metabolized substance in BAT. For this reason they are used as tracers to visualize adipose tissue and its metabolism.

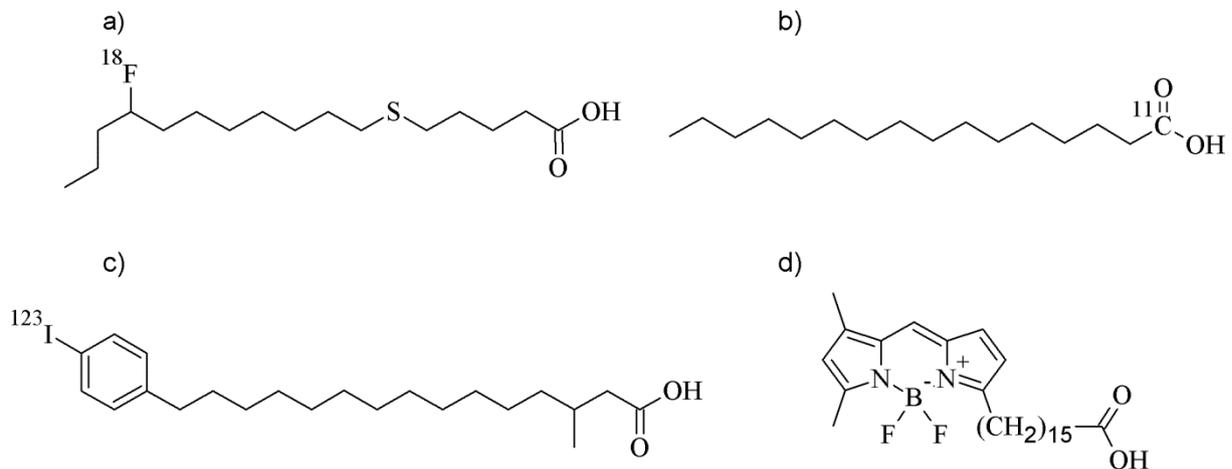


Figure 2: common used fatty acids for lipid metabolism imaging. a) 14(R,S)-[¹⁸F]fluoro-6-thia-heptadecanoic acid ([¹⁸F]FTHA), b) [¹¹C]hexadecanoic acid ([¹¹C]palmitate), c) 15-(4-[¹²³I]iodophenyl)-3-methyl-pentadecanoic acid ([¹²³I]BMIPP), d) 4,4-Difluoro-5,7-Dimethyl-4-Bora-3a,4a-Diaza-s-Indacene-3-Hexadecanoic Acid (BODIPY- C₁₆)

[¹⁸F]FTHA

FTHA (Fig 2 a)) was developed as a myocardial imaging agent because a high amount of LCFA is processed in the tricarboxylic acid cycle. It can be labelled by fluoride-18 through isotope exchange which makes this compound an easily available PET imaging agent. It is β-oxidized in the mitochondria and is bound irreversibly to mitochondria proteins once the sulphur group is free⁷⁵. Maximal uptake of [¹⁸F]FTHA is similar to [¹¹C]palmitate showing that neither the sulphur atom at the sixth position nor the introduction of fluorine-18 has influence on the uptake kinetics^{76,77}. 13F3THA analog showed significant lower uptake and faster washout, beta-oxidation is completely blocked by the sulfur atom at the odd position of 13F3THA^{78,79}. Radioactivity taken up by BAT was specific to only the even-substituted thia-[¹⁸F]FTHA and transport and metabolism steps are hindered for 13F3THA because the carboxyl group and the sulphur are located very close to each other. The fatty acid is internalized through assistance of several binding proteins in the endothelial, interstitial, and

intracellular spaces ⁷⁶. A similar mechanism for BAT as for WAT can be assumed which makes this compound suitable for BAT imaging. In first attempts of imaging BAT with [¹⁸F]FTHA Ouellet ⁸⁰ could show increased FA uptake in combination with an elevated [¹¹C]acetate uptake, as an oxidative tissue marker and therefore metabolism indicator, in cold exposed humans. Additionally a higher [¹⁸F]FDG uptake was reported indicating a higher energy turnover as a result of a higher metabolic activity during cold exposure ⁸⁰. CT gave information about the lipid density, which indicates the amount of intracellular TGs, and a utilization of these stored TG identifies them as the main fuel during cold exposure. These observations are in agreement with findings in other animal models ^{80,81}.

[¹¹C]palmitate

Another important PET tracer is [¹¹C]palmitate (Fig 2 b)). Its short half life of 20 min. makes an in-house cyclotron mandatory which restricts its use significantly. Apart from that restriction palmitate would be the optimal choice with respect to binding affinity as no big bulky chelator molecule needs to be coupled for imaging purposes. It behaves like any other natural non-synthesized fatty acid, justifying why it is always compared to modified molecules as a gold standard ⁸². [¹¹C]palmitate has been used as a myocardial imaging agent in clinical examinations ⁸³⁻⁸⁵ but as adipose tissue imaging becomes more and more important it is also used as a fat imaging agent in clinical experiments ⁸⁶. In comparison to [¹²³I]BMIPP (see below) and FTHA this FA is fully metabolized and metabolism products including [¹¹C]CO₂ are secreted by adipocytes which is decreasing the image quality.

[¹²³I]BMIPP

Like [¹⁸F]FTHA, [¹²³I]BMIPP (Fig 2 c)) started as a myocardial imaging agent ⁸⁷. It is labelled with iodine-123 through an isotopic exchange reaction to produce a SPECT- imaging agent. Commercial availability of BMIPP (up to 2012) and iodine-123 as well as the easy to perform exchange reaction ⁸⁸ made this compound favourable for adipose tissue imaging. A dependence on the isomeric confirmation was shown by Lin et al. ⁸⁷ and the [¹²³I]-3(R)-

BMIPP version showed uptake suitable for clinical trials. This implies that internalization is not hindered by the bulky benzol ring at the end of the FA. In comparison with palmitic acid under fasting conditions, [¹²³I]BMIPP had lower uptake into hepatocytes but similar characteristic changes were observed over time⁸². So far [¹²³I]BMIPP has only been used by one group to image BAT and its protein contribution in FA uptake^{54,89}.

Other radiolabelled fatty acids

As already mentioned, radiolabelled fatty acids have been often used as myocardial imaging agents in the past. The already described fatty acids have been shown good characteristics in terms of stability, binding affinity, pharmacokinetics etc. but there have been others developed which will be mentioned in the following paragraph.

[¹²³I]iodohexadecanoic acid ([¹²³I]IHX) and [¹²³I]iodoheptadecanoic acid ([¹²³I]IHDA) have been developed as myocardial perfusion agents^{90,91}. Clearance kinetics for both agents were compared with [¹¹C]palmitate and a similar clearance was found for the fast clearance components. The slow clearance components differed in that way that the half time for IHXA and IHDA was only approximately one fourth of that of [¹¹C]palmitate^{90,92}. These two components represent β -oxidation and fatty acid storage in lipid pools. This fast elimination half time makes this compound unsuitable for SPECT as decreased image quality can be expected. In addition a deiodination process could be observed which requires special corrections to differentiate between myocardial and blood pool activity⁹³.

15-(p-[¹²³I]iodophenyl)-pentadecanoic acid ([¹²³I]IPPA) showed a faster myocardial clearance compared to [¹²³I]BMIPP making it unfavorable for SPECT as this results in image distortions and loss of quantitative image accuracy⁹³. This problem is caused mainly because straight chain fatty acids are faster metabolized and cleared via β -oxidation. The methy-group at the β -position of BMIPP slows down the oxidation process and improves clearance and image accuracy in that way⁹⁴.

Boron-dipyrromethene (Bodipy)

A different imaging approach is fluorescent imaging with Bodipy. Because of its hydrophobic properties, Bodipy dyes can be used for staining of lipids, membranes and other lipophilic substance classes. High fluorescence quantum yields can be achieved due to its wide spread π -electron system. Fluorescence of this molecule is achieved through excitation at 503 nm and emission at 512 nm. The extension of the crude dye determines its field of usage for imaging purposes. There are many approaches from tumor imaging to adipose tissue detection with a fatty acid modified Bodipy. It has been proven that Bodipy C-16 (Fig 2 d)) binds specifically to fatty acid binding proteins⁹⁵ with a similar affinity as oleic acid, is incorporated into lipid droplets in hepatocytes⁹⁶, can be quantified by fluorescent cell scanners or plate readers⁹⁷ and can be followed downstream in metabolic processes after uptake⁹⁸. In a first try we were able to visualize WAT and BAT with Bodipy-C16 specifically *in vitro* with a 2 μ M Bodipy C-16 solution (unpublished results), like already reported for hepatocytes⁹⁶. Uptake could be identified as specific uptake as the lipid droplets were filled by Bodipy C-16 and no uptake in the cell nucleus can be observed. Incorporation of the fatty acid works as already described through protein mediation and the Bodipy dye does not hinder the uptake qualitatively. As penetration depth of excitation and emitted light cannot overcome the tissue barrier for BAT imaging, fluorescence imaging is restricted to *in vitro* applications in this approach. With its high spatial resolution it is a powerful tool to image on a cellular level^{95,97}.

Luciferin

By imaging adipose tissue with bioluminescence, problems of PET and SPECT imaging, like short half life and un-physiological uptake can be overcome⁹⁹. To a non-esterified fatty acid probe a luciferine derivative is coupled via a disulfide linker. This disulfide bridge is stable outside of the cells¹⁰⁰ and through reduction by glutathione within the cell a thiol is produced which undergoes cyclization to release free luciferine¹⁰¹. Free luciferine is then converted to oxyluciferine and a photon of light by luciferase¹⁰². Henkin et al. investigated the imaging

abilities of this compound and reported specific uptake as it could be blocked by oleate *in vitro*. Additionally luciferine uptake from micelles containing FA-luciferine could be observed and BAT specific uptake *in vivo* could be proven as uptake could be increased through BAT stimulation¹⁰¹.

Particle Imaging

To image lipid metabolism there are also other pathways available apart from imaging FA. Some interesting approaches with indirect measures for FA uptake have been reported and will be described in this section

Fluorescence labelled nanocrystals

FA are mostly transported in TRLs through the body. At the moment there are two assumptions about the delivery of FA through TRLs. FA can be taken up by adipocytes after TRLs are lipolyzed outside of the cell⁴⁵ or TRLs are internalized before they are lipolyzed⁴⁶. To investigate this process Bartelt et al. labelled TRLs with hydrophobic fluorescent nanocrystals (QD-TRL)⁴⁶. Quantum dots (QD) have a broad excitation and a narrow emission range and are therefore best suited for fluorescence imaging¹⁰³. With QD-TRLs an internalization of lipoproteins due to an increase in permeability of the endothelial barrier function could be observed⁴⁶. Other applications of this imaging agent can be thought of for quantitative and qualitative lipoprotein incorporation.

Functional MRI

Similar to QD-TRLs, Bartelt et al. labelled lipoproteins with hydrophobic superparamagnetic [⁵⁹Fe]oxide. The intention is the same as for QD-TGLs. The hydrophobic iron nanocrystals stay inside of the lipoprotein and indicate if the particle is incorporated or not. In addition the particle is filled with [³H]oleate to control whether FA are taken up. Bartelt et al. observed a faster clearance for the FA as well as for the nanocrystals after cold exposure indicating a contribution of BAT. Total oleate and Fe-oxide concentrations in cold activated mice of BAT and liver were comparable and omitting cold exposure lead to a decrease in BAT uptake for both. These results and the fact that [⁵⁹Fe]oxide concentrations were still increased one week after injection indicate an internalization of the whole lipoprotein particle⁴⁶.

In contrast to these observations, the study of Khedoe et al. is indicating an extracellular lipolysis followed by protein mediated uptake of FA ⁴⁵.

Other imaging techniques

MRI

With the possibility to image with a very high resolution and no radiation dose for the patient, MRI is an important player in adipose tissue imaging. By measuring the lipid content in interscapular BAT (iBAT), Grimpo et al. investigated the response to cold exposure and noradrenaline stimulation for wild type and UCP1 knock-out mice. Lipid content was the same for both groups under all temperature conditions with less FA stored in lipid droplets after cold exposure. A faster noradrenaline response could be observed in wild type mice ¹⁰⁴. Additionally in the study of Holstila et al. MRI was identified as a more adequate method to identify and quantify BAT, independently of its activation state, compared to PET/CT with [¹⁸F]FDG. BAT masses calculated with PET/CT were significant smaller than the masses obtained through dissection. MRI provided more accurate information even if a small overestimation has been observed. Advantageous is the independence of the activation state as this method is measuring the water/lipid content and compares it with previous records. For the same reason no cold exposure is necessary to activate BAT. A successful human pilot study was performed too by this group ¹⁰⁵. Additionally first studies have been done in adult humans with [¹⁸F]FDG-PET/CT as a reference standard by van Rooijen et al. ¹⁰⁶. In comparison to the mentioned studies in animal models ¹⁰⁷⁻¹¹⁰, they could not discern active BAT on the basis of tissue fat content. Dynamic BAT activation could be identified by T2*-weighted MRI during cold stimulation. Eventhough PET provides images of active BAT with a higher contrast, MRI is capable in gaining additional information such as: temporal and quantitative measurements of tissue fat content, changes in blood flow, volume, and oxygenation of active BAT ¹⁰⁶.

CT

A standalone computed tomography to identify BAT seems in terms of a lower radiation dose compared to PET/CT a good choice but without PET it is almost impossible to identify BAT

by the difference in Hounsfield units and gets even more complicated when WAT is located next to it. So most of the recent studies are PET/CT related and some indicate PET/CT with [¹⁸F]FDG not as the optimal choice ¹¹¹. Another problem is the difference in interpreting the Hounsfield units as adipocytes. Gortel et al. used a threshold of -200 to -30, Maurovic-Horvat et al. used a threshold from -149 to -30 and Mahabadi et al a threshold of -195 to -45 ¹¹²⁻¹¹⁵. By having these differences an under or overestimation is easily possible and does not really describe the amount of BAT. Because of the listed reasons a CT on its own is not capable in identifying BAT and also a PET/CT with [¹⁸F]FDG is not the best choice anymore. There are several better compounds which target BAT without activation and also MRI seems to have more advantages and a more accurate way to identify adipocytes.

Ex vivo methods

[³H]Triolein

Tritium does not have the optimal properties for non-invasive imaging. Its really soft radiation makes it impossible to detect it from outside of the body because the emitted electrons are stopped after a few micrometers in tissue. Its field of application is supported by its favourable biodistribution and plasma extraction. By that method the uptake at certain time points can be tested by the fraction of Triolein still present in the blood ^{46,77}. An impact of cold exposure or BAT activation through noradrenaline can be observed but not a clear quantification of adipocyte uptake can be achieved as lipids can be taken up by other tissues or cleared from the blood.

[³H]triolein is also used in autoradiography studies. Stein et al. investigated metabolism steps of starved rat fat cells and found that after exposure with [³H]triolein containing blood, 9-17% of the esterified fatty acids were in the form of diglycerides, less than 1% were monoglycerides and the rest were TG. They assumed therefore the esterification of diglycerides to TG as the rate limiting step in the TG synthesis in adipose tissue. Four fifth of the signals observed were found within the lipid droplets of the cell identifying them as the primary destination for newly taken up FA ¹¹⁶.

ADIFAB staining

ADIFAB is a stain for unbound fatty acids and is only applicable in *in vitro* work. Acrylodan labelled rat intestinal fatty acid binding protein (ADIFAB) is fluorescent if it is bound to a fatty acid. Kampf et al. ¹¹⁷ investigated the change in cytosolic unbound FA in response to a change in extracellular unbound FA. They discovered an influx concentration dependence of membrane carrier proteins which could be saturated and a strong influence of translocation for the fatty acid uptake ratio ¹¹⁷. Other experiments have been done to examine the fatty acid uptake mechanism *in vitro* with ADIFAB and identified flip-flop as the rate limiting step for oleate transport through the lipid bilayer membrane ¹¹⁸.

Conclusion

Lipid metabolism imaging has a broad field of application. It is generally accepted in terms of myocardial imaging, hepatic imaging etc. Our focus was to outline its broad function in fields within the whole body and specifically for the use to visualize adipose tissue and its metabolism. Its fast developing field within the niche of adipose tissue imaging draws attention to new imaging approaches and agents and helps to understand more and more this, for a long term, underestimated tissue. We expect radiolabelled fatty acids to be the most promising agents in clinical research to unravel adipose tissue processes in BAT for the next years but also fluorescent imaging, with its ability to image on subcellular level, and MRI should not be underestimated and will contribute to the exploration of adipose tissue and lipid metabolism.

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Chapter 3: Characterization of BAT activity in rats using invasive and non-invasive techniques

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Abstract

Brown adipose tissue (BAT) is widely considered as a potential target for combating obesity in humans where active BAT metabolizes glucose and fatty acids as fuel resulting in heat production. Several prospective studies in humans have been set up to further study the presence and metabolic activity of BAT mostly using PET imaging in cold-stimulated conditions with the radiolabeled glucose derivative [^{18}F]FDG. However, radiotracers beyond [^{18}F]FDG have been proposed to investigate BAT activity, targeting various aspects of BAT metabolism. It remains questionable which tracer is best suited to detect metabolic BAT activity and to what extent the generated results correlate with *ex vivo* metabolic BAT activity.

Methods: Dynamic PET and SPECT imaging, targeting different aspects of BAT activation such as glucose metabolism, fatty acid metabolism, noradrenergic stimulation, blood perfusion and amino acid transport system, was performed immediately after injection of the tracer in rats under different temperatures: room temperature, acute cold (4 °C for 4 h) or acclimated to cold (4 °C for 6 h per day during 28 days). Furthermore, MRS-derived BAT temperature was measured in control and cold-acclimated rats.

Results: In rats housed at room temperature, only [^{18}F]FDG visualized BAT. Glucose metabolism, fatty acid metabolism, noradrenergic stimulation and blood perfusion showed a clear tracer-dependent twofold increase in BAT uptake upon cold exposure. Only the tracer for the amino acid transport system did not show BAT specific uptake under any of the experimental conditions. MRS demonstrated that cold-acclimated animals had BAT with a stronger heat-production compared to control animals.

Conclusion: BAT activity following cold exposure in rats was visualized by several tracers, while only [^{18}F]FDG was also able to show BAT activity under non-stimulated conditions (room temperature). The variances in uptake of the different tracers should be taken into account when developing future clinical applications in humans.

Introduction

Brown adipose tissue (BAT) has gained considerable attention over the last decade, as its appearance and function in humans is becoming more elucidated. The main function of BAT is to dissipate energy in the form of heat, a process driven by the mitochondrial uncoupling protein 1 (UCP1), in response to cold exposure^{1,2}. BAT is highly vascularized, densely innervated by the sympathetic nervous system and has thermogenic capacity that can significantly influence homeostasis³⁻⁵.

The presence and/or activity of BAT can be non-invasively visualized in humans using magnetic resonance imaging (MRI), computed tomography (CT), thermography and molecular imaging⁶⁻⁹. Thermography is a relatively cheap technique, able to quantify elevations in skin temperature resulting from increased BAT activity^{9,10}. Since skin temperature is only indirectly correlated to BAT abundance ($R^2 < 0.3$)¹⁰, this technique is not optimally suited for directly monitoring BAT activity. MRI and CT are capable of visualizing soft tissue with a high spatial resolution but struggle to distinguish between white adipose tissue (WAT) and BAT. These techniques quantify the decrease in fat content during cold exposure as a reflection of BAT activation. Molecular imaging, using either positron emission tomography (PET) or single photon emission computed tomography (SPECT) is performed with different radiopharmaceuticals. In this study we employed 2-^[18F]fluorodeoxyglucose (^[18F]FDG), 14-^[18F]fluoro-6-thia-heptadecanoic acid (^[18F]FTHA), ^[99mTc]Tc-2-methoxy-isobutyl-isonitrile (^[99mTc]TcMIBI) and ^[123I]-metaiodobenzylguanidine (^[123I]MIBG). These radiopharmaceuticals are able to visualize and quantify BAT activity in humans¹¹⁻¹⁶. ^[18F]FDG mainly visualizes glucose transport and is the most frequently used tracer to image BAT due to its high availability. However fatty acids are the main metabolized substance class in BAT^{17,18} and therefore ^[18F]FDG might largely underestimate BAT activity. In addition ^[18F]FDG BAT uptake is verifiably reduced in diabetic patients¹⁹ due to their insulin resistance. ^[18F]FTHA visualizes uptake of free fatty acids (probably mainly via CD36²⁰). Fatty acids are the main metabolized substance class in BAT and might therefore be a good

measure for BAT activation state. [¹²³I]MIBG visualizes the density of sympathetic nerve endings. Norepinephrine, which is released by the sympathetic nervous system is a known activator for BAT ^{21,22} and therefore imaging with [¹²³I]MIBG is a good indication of the susceptibility of BAT to be activated. [^{99m}Tc]TcMIBI visualizes perfusion of tissue by binding to mitochondria rich cells. BAT cells have a high number of mitochondria and in previous studies it was shown that [^{99m}Tc]TcMIBI is able to visualize BAT under basal conditions ^{11,12}. In acute cold conditions uptake was only slightly pronounced and it was speculated that [^{99m}Tc]TcMIBI only shows increased uptake after cold acclimation. Additionally [¹²³I]-Phenylalanine ([¹²³I]IPA) (targeting LAT1-4 amino acid transport system density) was used. [¹²³I]IPA has never been reported to visualize BAT clinically or preclinically, but as it correlates to a key amino acid transport system we feel it is important to include it in our study.

PET and SPECT images indicating presumed regions of BAT can also be selectively identified on CT and MRI images, and mRNA and protein analysis from tissue samples from these regions confirmed BAT-characteristics ^{23,24}. It has also been shown that [¹⁸F]FDG uptake on PET images correlates positively to cold outside temperatures and negatively to a subjects BMI, in correspondence with predictions from in vitro work ²⁵. However, ex vivo and in vitro data can vary greatly within a specific tissue or type of cell culture ²⁶, and it remains an open question just to what extent intra-tissue measurements from each of these techniques truly represents its metabolic activity.

The most straightforward method to activate BAT *in vivo* is exposure to acute cold but also cold acclimation further stimulates [¹⁸F]FDG PET uptake in humans ^{6,27,28}. However no other radiopharmaceuticals have been used in a clinical setting to evaluate the effect of cold acclimation, although Baba *et al.* compared a number of tracers ([²⁰¹Tl]thalliumchloride, [¹²³I]MIBG, ([^{99m}Tc]TcMIBI, [¹⁸F]- or [³H]FDG, [³H]-I-methionine, and [³H]thymidine) to assess BAT uptake in rodents after acute cold exposure ¹¹.

Therefore 3 questions remain: 1) which aspects of BAT metabolism can best be used to investigate BAT activity upon acute cold exposure and cold acclimation? 2) which

radiopharmaceutical is most suitable to evaluate this aspects? 3) to what extent does data acquired using radiopharmaceuticals correlate with ex vivo findings and data acquired by other modalities such as MR Spectroscopy (MRS)?

In our study, we primarily investigated to what extent different radiopharmaceuticals are suitable to quantify response of BAT to acute cold and cold acclimation in rats. For this reason, we used a rat model which was acutely exposed to cold or acclimated to cold. We compared tracers that were previously described to visualize BAT in clinical setting, namely [¹⁸F]FDG, [¹⁸F]FTHA, [¹²³I]MIBG, [^{99m}Tc]TcMIBI and ([¹²³I]IPA). Furthermore, using MRS, we visualized and quantified BAT in rat following acute cold or cold acclimation by means of gradient-echo sequence (FLASH – Fast Low Angle SHot).

Materials & Methods

Animal model

All animal studies were approved by the local animal ethical committee of the Maastricht University; with the internal permit number: DEC 2012-001. Male 12-week old Wistar rats were acquired from Harlan and housed under controlled temperatures of $22\text{ }^{\circ}\text{C} \pm 1\text{ }^{\circ}\text{C}$ and 55-75% air humidity, in a 12 h light–dark cycle with water and food chow ad libitum. 36 animals were, after an initial week of normal housing, placed at $4\text{ }^{\circ}\text{C}$ for 6 h per day (9 am-3 pm) (food/drink ad libitum) during 28 days, see also figure 1. At day -2 animals were scanned with one of the listed radiotracer to determine the control (room temperature) condition. To measure the acute cold conditions animals were scanned at day 0 after 4 h cold exposure and cold acclimated conditions were obtained after 28 days of cold exposure for 6 h/day. At day 30 cold exposed animals were injected in the cold with the PET/SPECT tracer and were further exposed to cold for 30 min before they were scanned to see if cold exposure during injection has an effect. 8 animals served as controls and were maintained at room temperature throughout the experiment. The physiological impact of the experiment was assessed by comparing food uptake, weight gain and discomfort in cold-acclimated and room temperature groups of animals. In addition, supraclavicular BAT was dissected and weighed in sacrificed animals.

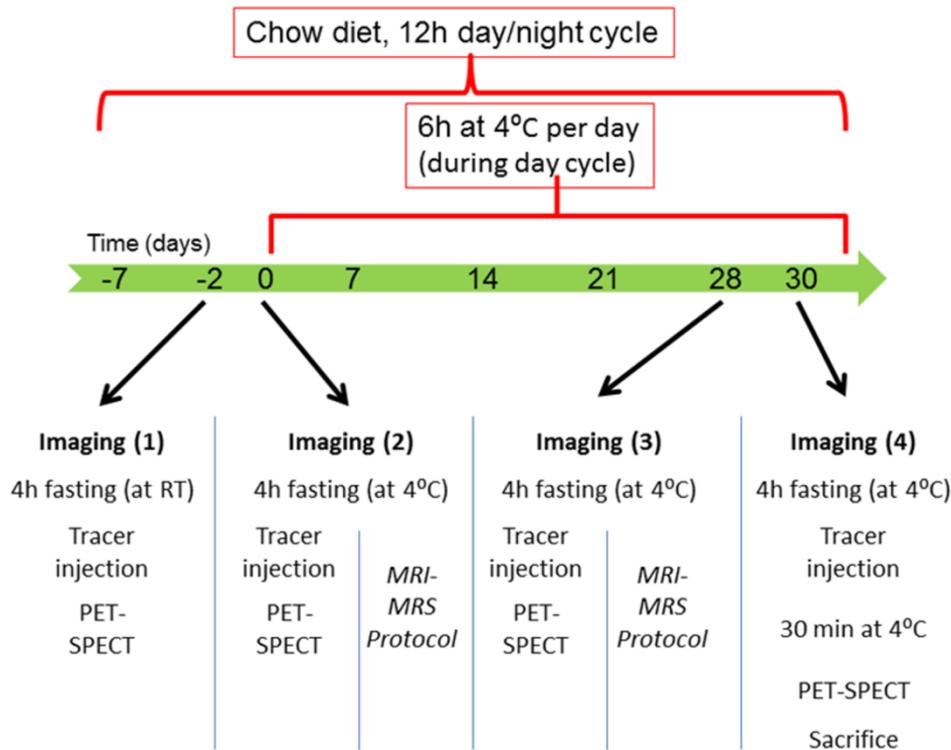


Fig 1: Study design. Imaging with each tracer was performed sequentially in baseline conditions (day -2, temperature regime 1 (TR-1)), after acute cold exposure (day 0, TR-2), after cold acclimation (day 28, TR-3) and after tracer distribution in the cold (day 30, TR-4).

Tracers: radiosynthesis and formulation

[¹⁸F]FDG was purchased from GE Healthcare. Sestamibi (MIBI) was labeled with technetium-99m according to manufacturer guidelines (Mallinckrodt Medical). Non-carrier added [¹²³I]MIBG was prepared as suggested by the manufacturer of the precursor²⁹. In short, synthesis was achieved by reacting iodine-123 with a polystyrene resin of dibutylstannyl benzylguanidine (Molecular Insight) in an oxidizing solution, transferring the [¹²³I]MIBG over a cation exchange filter, rinsing with saline solution and finally collecting the non-carrier-added tracer by a phosphoric acid/ascorbic acid solution. After neutralization with NaOH a final volume of about 2ml is reached and the solution is ready for injection. The synthesis of ([¹²³I]IPA) was adapted from earlier publications^{30,31}. Non-carrier added [¹⁸F]FTHA was

prepared according to an adapted procedure from ³², starting from benzyl-14-(R,S)-tosyloxy-6-thiaheptadecanoate (ABX) as a precursor. The final purified compound was formulated using a 0.1% rat serum albumin solution to ensure solubility. All tracers showed a radiochemical purity of at least 98%.

Imaging – general and anesthesia

PET, SPECT and MRI was performed according to the timeline represented in figure 1. The animals were divided into subgroups, with group A, B, C, D and E (n=4 or more per group) being scanned with respectively [¹⁸F]FDG, [¹⁸F]FTHA, [¹²³I]MIBG, [^{99m}Tc]TcMIBI and [¹²³I]IPA and group F was used for MRI (n=8). Imaging started at 1 pm. Prior to any scan or animal sacrifice, animals were fasted for 4 hours at either room temperature (baseline control conditions) or at 4 °C (exposure to cold). PET/SPECT imaging was performed under pentobarbital anesthesia (0.1 ml of a 60 mg/ml solution per 100 g body weight, i.p.), as this sedative was reported to show the least side effects on BAT activity ³³. MRI/MRS imaging, due to its study duration (2.5-3 h per animal) and difficulty to access the animal, did not allow pentobarbital, so isoflurane was used (O₂ as carrier, 3% for initial sedation, 1.8% for maintenance).

Imaging (PET, SPECT) and biodistribution

Animals were sedated and subsequently injected in a tail vein under the camera (μ PET Focus 120, Siemens, with a 1.4 mm spatial resolution or U-SPECT, MiLabs, with a 0.6 mm resolution) with 20-50 MBq [¹⁸F]FDG (group A), 20-50 MBq [¹⁸F]FTHA (group B), 100MBq [¹²³I]MIBG (group C), 100 MBq [^{99m}Tc]TcMIBI (group D) or 100 MBq [¹²³I]IPA (group E). The imaging room was conditioned to room temperature (21 °C).

On day -2, 0 and 28 dynamic imaging of the rat upper torso was performed immediately after injection for 25 minutes (8x15 sec, 6x30 sec, 5x 60 sec, 3x300 sec for PET, and 15x 3 min for SPECT), while on day 30 a static image was acquired at 30-55 minutes after injection. On day -2, the rats were placed on a heating pad, while this pad was omitted on day 0, 28 and

30. Body core temperature was monitored using a rectal probe during imaging. At day 30, immediately after imaging, animals were sacrificed and relevant organs/tissues dissected, weighed and counted (automated NaI(Tl) gamma counter (Wallac Wizard)).

Image analysis (PET/SPECT)

After smoothing to 3x3x3 mm voxel size, image analysis was performed by drawing a volume of interest (VOI) around the interscapular BAT (iBAT), part of the myocardium and part of the liver (PMOD 3.0). The VOI around iBAT was drawn around the visible iBAT (on PET or SPECT image), combined with knowledge of anatomical location. Each VOI was then limited by applying a cut-off value of 30% of the (maximum– minimum) value in the VOI, thereby maintaining only tissue with true uptake. A cut-off value of 30% was found to be optimal in previous studies (unpublished), and lead to the final metabolically active volume (metabolic volume). For each VOI, the Standardized Uptake Values (SUV_{mean} , Bq/cc in the region of interest, divided by the injected dose per animal weight), as well as the metabolic volume and the total metabolic activity (SUV_{mean} multiplied by metabolic volume) were calculated. For PET, this SUV_{mean} value could be calculated directly using the output parameters from the μ PET. For SPECT, the output parameter (“counts/cc”) was converted to Bq/cc using a previously determined phantom-based conversion factor of 635 (Bq per count) for both technetium-99m and iodine-123.

Imaging (MRI/MRS)

During the MR exam the rats were placed in a cradle, which was positioned in the center of a quadrature volume coil (\varnothing 72 mm, transmit-receive) in a 7 Tesla MR System (Bruker Biospin, Ettlingen, Germany). Rectal temperature was measured using a fiber probe. Whole-body cooling was achieved by placing of the rats on a waterbed which was connected to a heat exchanger. The water was circulated by a pump and cooled from to 45 °C to ~25 °C, allowing rats to be cooled by three degrees Celsius from their initial rectal temperature. After

this temperature loss, animals were warmed again using the waterbed until rectal temperature reached at least 36 °C.

In this study we use a gradient-echo sequence with a fat suppression pulse to visualize the position of BAT. The FLASH (Fast Low Angle SHot) sequence makes use of a moderate flip angle of 40 degrees, short TE of 3.9 msec and long TR of 2050 msec. When also applying a fat suppression pulse, the tissues containing relatively more fat will appear darker, allowing to precisely locate iBAT. Temperature-dependent chemical shift between H₂O and the CH₂ peak of fat was measured using point resolved spectroscopy (PRESS, TE=14 ms, TR=4500 ms, NSA=64). Spectra were acquired from a 3x2x3 mm voxel positioned in BAT, every 4 minutes for at least 2.5 hours. Spectra were fitted using an in-house program. The chemical shift of the resonance of H₂O and of the CH₂ resonance of fat were determined and temperature change was quantified by assuming a temperature-dependent frequency shift of H₂O of 0.01 ppm/°C.

Western blot

All animals (control animals and cold-acclimated animals) were sacrificed at day 30 of the experiment. iBAT and visceral WAT were dissected and snap-frozen in liquid nitrogen. Western blot has been performed as stated previously³⁴. In short, adipose tissue samples were incubated with RIPA buffer, frozen and after thawing lysates were passed through a 25-G needle. Lysates were separated using SDS-PAGE prior to electrophoretic transfer onto nitrocellulose membranes. The UCP1 antibody was from Abcam (Cambridge, UK). CD36 and GLUT4 antibodies were purchased from Santa Cruz (Dallas, TX).

Quantitative real-time PCR

All animals (control animals and cold-acclimated animals) were sacrificed at day 30 of the experiment. iBAT and visceral WAT were dissected stored in RNA-later prior to further mRNA expression analysis. Rat mRNA primer sets were developed and optimized for PPAR γ , C/EBP α , Sirtuin1, UCP1, UCP2, ADRB3, DIO2, GLUT4, ATGL, LPL and PRDM16

transcripts. All gene expression data were normalized to beta-actin (see supplemental materials for detailed information on mRNA analysis).

Statistics

Imaging data from animal groups were compared using ANOVA (with Bonferroni correction) for inter-group comparison, and a paired student's t-test for intra-group comparison. mRNA data and protein data were compared using an unpaired t-test. $P < 0.05$ was considered to be statistically significant. All statistical tests were performed using GraphPad Prism (GraphPad Software).

Results

Physiological impact of the study

Animals exposed to cold showed no signs of severe discomfort. Only during the first 2-3 days of the study the animals showed mild discomfort (shivering when exposed to cold). Animals acclimated to cold tended to gain less weight compared to animals in control conditions (1.10 ± 0.15 g/day vs 1.43 ± 0.12 g/day, $p=0.09$), however the cold acclimated animals did eat more (29.4 ± 1.1 g/day vs 26.3 ± 0.9 , $p=0.05$). In addition, upon dissection, total interscapular adipose tissue (iAT) showed a tendency to be larger in cold-acclimated animals (1.66 ± 0.13 g vs 1.35 ± 0.08 g, $p=0.13$). iAT consisted of brownish adipose tissue, covered by a layer of white-pale adipose tissue. iWAT reduced in weight by cold acclimation from 0.65 ± 0.08 to 0.50 ± 0.06 ($p=0.14$), while iBAT increased in weight from 0.70 ± 0.07 g to 1.16 ± 0.10 g ($p=0.01$) (Fig 2).

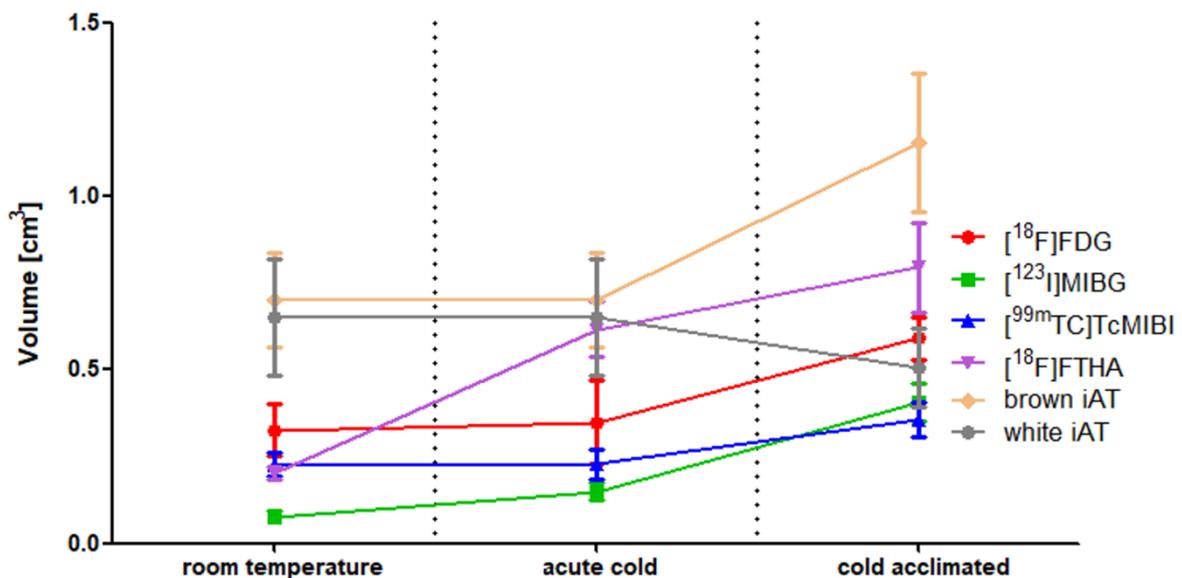


Fig 2: iBAT metabolic volumes of [¹⁸F]FDG, [¹⁸F]FTHA, [¹²³I]MIBG and [^{99m}Tc]TcMIBI, as well as iBAT and iWAT volume determined by dissection, in different temperature conditions.

PET and SPECT imaging

In baseline conditions, [^{18}F]FDG images showed pronounced uptake in heart and brain, and to some extent in iBAT (Fig 3). Upon acute cold exposure, uptake was still increased in the heart and the brain (although less pronounced), but now iBAT and cervical brown adipose tissue (cBAT) were also clearly visible (image not shown). After acclimation, this pattern was very well-defined (Fig 4). For [^{18}F]FTHA, baseline condition images showed a high uptake in the liver and the heart, with only minor uptake in the iBAT. Upon cold exposure, uptake was clear again in the heart and the liver, however, now iBAT was also clearly visible. After acclimation, this pattern was even more pronounced, also showing cBAT (Fig 3 and 4). For [^{123}I]MIBG, a pattern similar as compared to [^{18}F]FTHA was observed, although uptake in BAT was less prominent (Fig 3 and 4). For [$^{99\text{m}}\text{Tc}$]TcMIBI, baseline condition images showed again a high uptake in the liver and the heart, with only minor uptake in iBAT. Upon cold exposure or cold acclimation, uptake in iBAT increased mildly (Fig 3 and 4). Finally, [^{123}I]IPA showed no specific uptake anywhere in the upper torso in any condition (Fig 3 and 4).

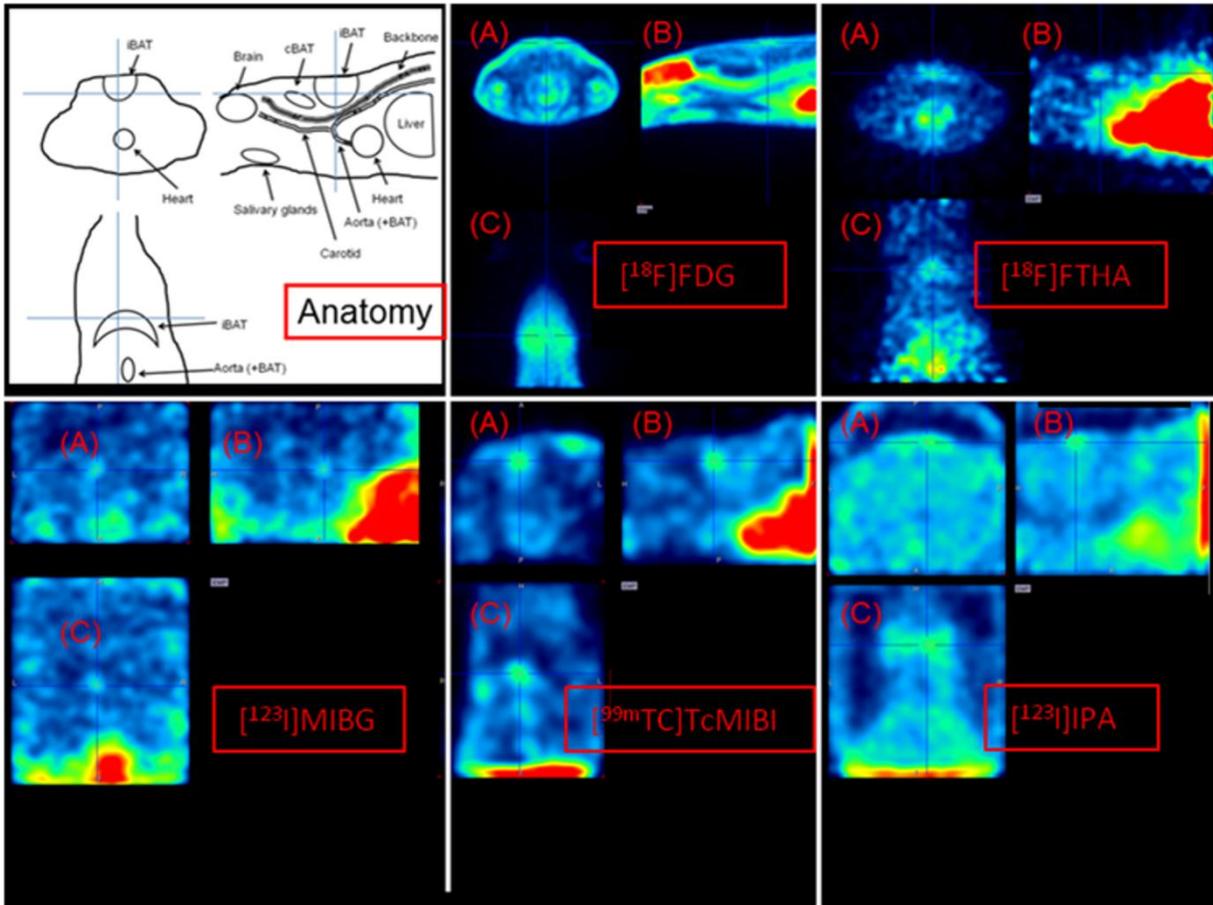


Fig 3: Transverse (A), sagittal (B) and coronal (C) slices of images from a room-temperature housed rat, centered on iBAT, depicting the anatomy, $[^{18}\text{F}]$ -FDG distribution, $[^{18}\text{F}]$ -FTHA distribution, $[^{123}\text{I}]$ -MIBG distribution, $[^{99\text{m}}\text{Tc}]$ -MIBI and $[^{123}\text{I}]$ -IPA distribution. Uptake in BAT is low for all tracers.

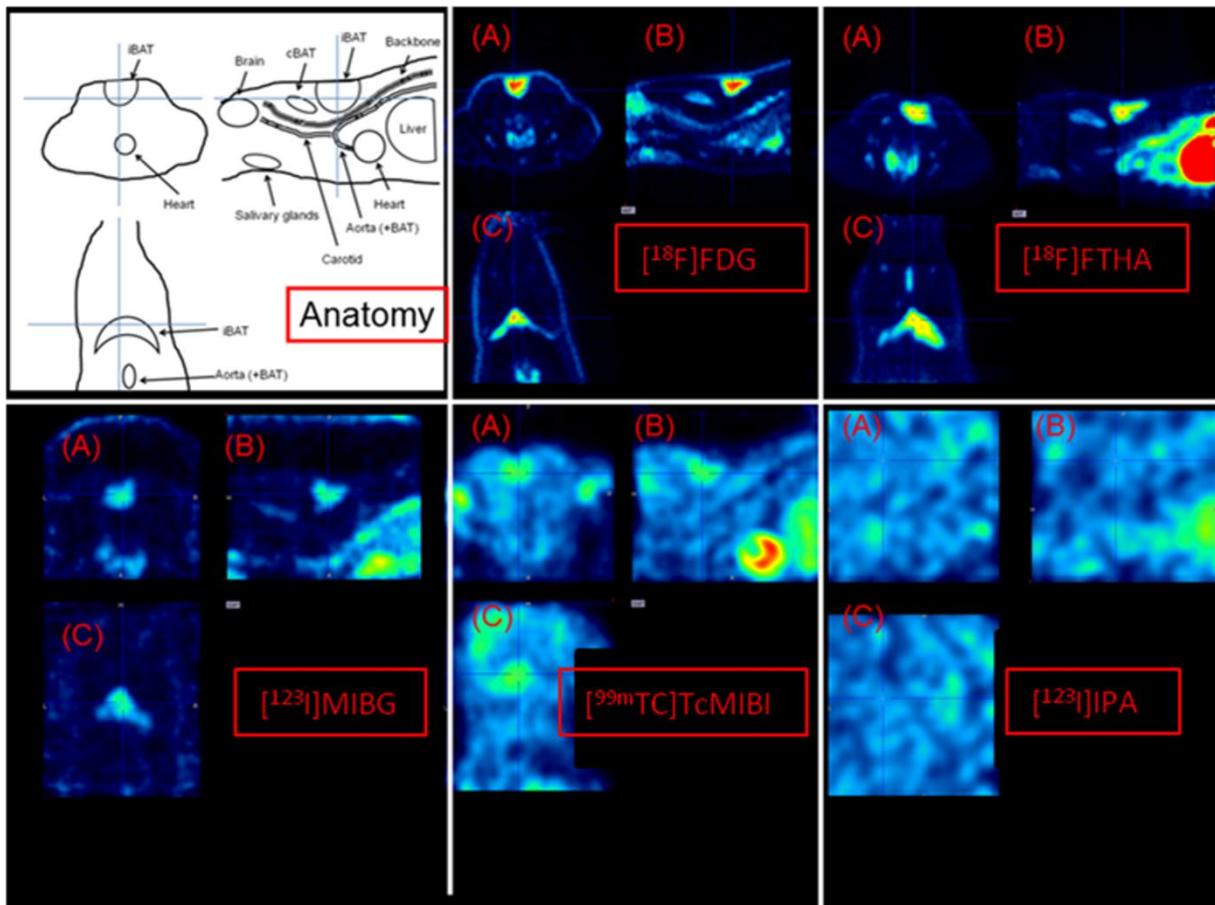


Fig 4: Transverse (A), sagittal (B) and coronal (C) slices of images from a cold-acclimated rat, centered on iBAT, depicting the anatomy, $[^{18}\text{F}]$ -FDG distribution, $[^{18}\text{F}]$ -FTHA distribution, $[^{123}\text{I}]$ -MIBG distribution, $[^{99\text{m}}\text{Tc}]$ -MIBI and $[^{123}\text{I}]$ -IPA distribution. iBAT, and to some extent cBAT show a high uptake for all tracers except $[^{123}\text{I}]$ -IPA.

Figure 5 shows the SUV_{mean} values for each tracer in iBAT in different conditions. It can be seen that acute cold exposure results in an increased SUV_{mean} value for $[^{18}\text{F}]$ FDG, $[^{18}\text{F}]$ FTHA, $[^{123}\text{I}]$ MIBG and $[^{99\text{m}}\text{Tc}]$ TcMIBI in iBAT when compared to baseline conditions (p -values are respectively 0.11; 0.03; 0.001 and 0.02) (also see Table 1). Cold acclimation results in similar increases when compared to room temperature values. As shown in figure 2, the metabolically active tissue volume in iBAT is increased after cold acclimation when investigated with $[^{18}\text{F}]$ FDG, $[^{18}\text{F}]$ FTHA, $[^{123}\text{I}]$ MIBG and $[^{99\text{m}}\text{Tc}]$ TcMIBI. Surprisingly, this increase in labeled tissue is already present after single acute cold exposure for $[^{18}\text{F}]$ FTHA and to some extent also $[^{123}\text{I}]$ MIBG. $[^{123}\text{I}]$ IPA shows poor uptake in iBAT, preventing accurate VOI drawing. As a result, it is only possible to estimate SUV_{mean} $[^{123}\text{I}]$ IPA values in iBAT, but

not the metabolic volume or total metabolic activity. The SUV_{mean} values show no change over the course over the experiment.

Table 1: SUV, metabolic volume and total metabolic activity of [^{18}F]FDG, [^{18}F]FTHA, [^{123}I]MIBG, [^{99m}Tc]TcMIBI and [^{123}I]IPA in iBAT in baseline conditions, after acute cold exposure, after cold acclimation and after uptake in the cold. \uparrow indicates a p-value < 0,10 when compared to baseline conditions, while \dagger indicates a p-value < 0,10 when compared to the value after acute cold exposure. NQ indicates “not quantifiable”, while NA means these data are not available.

iBAT	Baseline	Acute cold exposure	Acclimated to cold	Uptake in cold
Tracer	SUV (g/cm ³)			
[^{18}F]FDG	1.54±0.26	3.24±0.88	3.10±1.39	3.23±1.51
[^{18}F]FTHA	0.92±0.07	2.14±0.20 \uparrow	1.78±0.22 \uparrow	2.36±0.24 \uparrow
[^{123}I]MIBG	0.86±0.24	3.15±0.20 \uparrow	2.06±0.42 \uparrow	1.82±0.12 \uparrow
[^{99m}Tc]TcMIBI	1.99±0.31	2.69±0.34 \uparrow	2.20±0.40	2.83±0.56
[^{123}I]IPA	1.27±0.04	1.21±0.09	1.22±0.01	NA
	Metabolic volume (cm ³)			
[^{18}F]FDG	0.32±0.07	0.35±0.12	0.59±0.06 \uparrow	0.54±0.03 \uparrow
[^{18}F]FTHA	0.20±0.01	0.62±0.04 \uparrow	0.80±0.06 \uparrow, \dagger	0.64±0.12 \uparrow
[^{123}I]MIBG	0.08±0.02	0.15±0.02 \uparrow	0.41±0.05 \uparrow, \dagger	0.30±0.03 \uparrow, \dagger
[^{99m}Tc]TcMIBI	0.23±0.03	0.23±0.05	0.36±0.05 \uparrow, \dagger	0.42±0.05 \uparrow, \dagger
[^{123}I]IPA	NQ	NQ	NQ	NA
	Total metabolic activity (SUV x Volume) (cm ³)			
[^{18}F]FDG	0.48±0.09	0.89±0.11 \uparrow	1.71±0.52 \uparrow	1.78±0.66 \uparrow
[^{18}F]FTHA	0.18±0.01	1.41±0.16 \uparrow	1.41±0.17 \uparrow	1.54±0.38 \uparrow
[^{123}I]MIBG	0.07±0.04	0.46±0.06 \uparrow	0.79±0.12 \uparrow, \dagger	0.54±0.02 \uparrow, \dagger
[^{99m}Tc]TcMIBI	0.43±0.06	0.59±0.08	0.74±0.06 \uparrow	1.25±0.34 \uparrow, \dagger
[^{123}I]IPA	NQ	NQ	NQ	NA

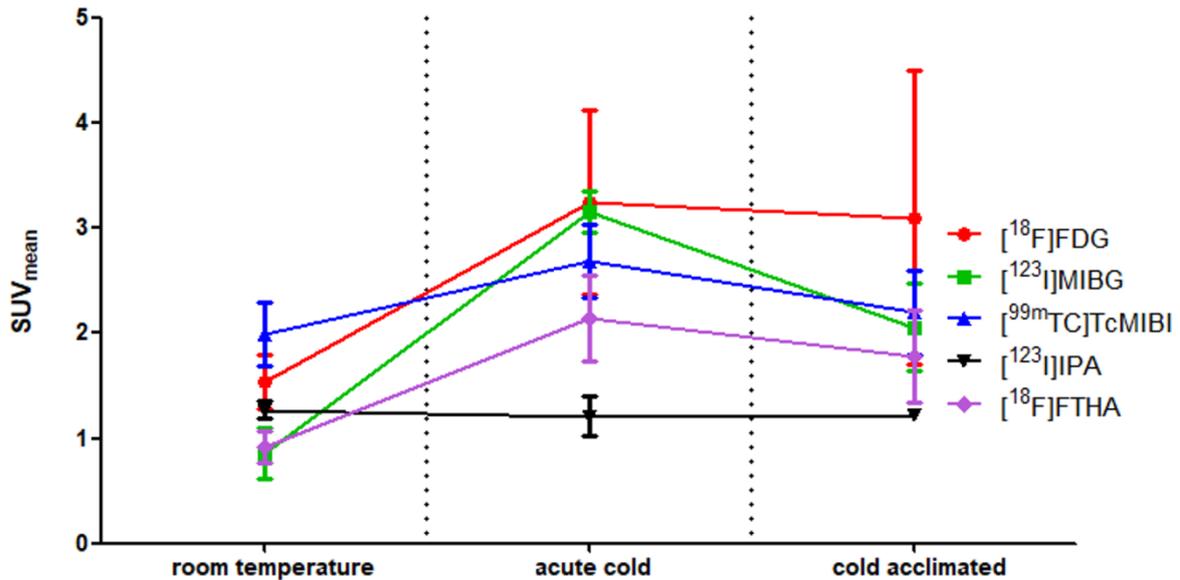


Fig 5: iBAT SUV_{mean} values of [¹⁸F]-FDG, [¹⁸F]-FTHA, [¹²³I]-MIBG, [^{99m}Tc]-MIBI and [¹²³I]-IPA in different temperature conditions.

In comparison, the myocardium and the liver showed no significant changes in SUV_{mean} values throughout the study for any of the tracers, nor did their (metabolic) volume change.

In addition, there was no difference in data obtained from images that were acquired by injecting the tracer in cold-acclimated animals under the camera, in a room where the animal was no longer exposed to cold during tracer distribution, and images that were acquired after the tracer had been given the chance to distribute in the cold-acclimated animal for 30 minutes in a cold environment (also see Table 1).

Analysis of biodistribution data obtained from dissected tissue (Table 2) confirms the data obtained from images. Dissection data did allow quantification of a number of aspects that were impossible to derive from imaging data, such as the discrimination between iWAT and iBAT, thereby clearly showing that only iBAT has tracer uptake. Perivascular BAT, often visible but never quantifiable on images due to its small size and structure, showed a similar uptake to that in iBAT when corrected for weight.

Table 2: Biodistribution (determined by dissection) of [^{18}F]FDG, [^{18}F]FTHA, [^{123}I]MIBG, [$^{99\text{m}}\text{Tc}$]TcMIBI and [^{123}I]IPA in selected organs 1 h after tracer uptake in the cold.

%ID/g	[^{18}F]FDG	[^{18}F]FTHA	[^{123}I]MIBG	[$^{99\text{m}}\text{Tc}$]TcMIBI	[^{123}I]IPA
iBAT	2.89±0.66	1.36±0.39	0.93±0.23	0.90±0.10	0.37±0.04
iWAT	0.41±0.12	0.16±0.06	0.18±0.03	0.16±0.02	0.26±0.03
Perivascular BAT	3.65±1.55	1.08±0.30	2.02±0.54	1.49±0.37	0.52±0.05
Intestinal WAT	0.19±0.01	0.14±0.05	0.12±0.03	0.05±0.01	0.22±0.10
Subcutaneous WAT	0.18±0.02	0.08±0.01	0.17±0.05	0.10±0.03	0.04±0.07
Liver	0.37±0.04	3.16±0.38	1.64±0.14	1.25±0.26	0.68±0.04
Lung	0.36±0.05	0.53±0.11	3.45±0.39	0.55±0.02	0.57±0.02
Heart	2.34±0.25	1.01±0.08	4.35±0.34	3.00±0.23	0.58±0.01
Muscle	0.10±0.01	0.07±0.01	0.11±0.01	0.12±0.02	0.40±0.04
Salivary glands	0.51±0.05	0.43±0.06	1.87±0.28	1.93±0.30	0.34±0.23
Blood	0.34±0.06	0.12±0.01	0.16±0.01	0.02±0.01	0.93±0.02

MR Spectroscopy

The use of the gradient-echo sequence with a fat suppression pulse made it possible to visualize the position of BAT in rat (Fig 6). The FLASH sequence combined with a fat suppression pulse, makes the WAT darker in comparison to the BAT. Temperature was determined in BAT, and as a reference, also in muscle tissue by MRS and furthermore, rectal temperature was determined. The one-hour cooling protocol resulted in a decreased temperature of 1.9 ± 0.9 °C in non-acclimated animals and a decrease of 1.8 ± 0.9 °C in cold-acclimated animals (also see Table 3). During this period of cooling, MRS-derived temperature in BAT decreased by 1.2 ± 0.9 °C in non-acclimated animals but increased by 0.3 ± 1.5 °C in cold-acclimated animals ($p=0.22$). MRS-derived temperature in muscular tissue (including a minor amount of subcutaneous adipose tissue) decreased by 2.4 ± 1.5 °C in non-acclimated animals and by 1.4 ± 1.8 °C in cold-acclimated animals ($p=0.18$). In cold-acclimated animals the decrease in BAT temperature is less pronounced than that in

subcutaneous muscular tissue ($p = 0.08$) or in the rectum ($p = 0.004$), while temperature in non-acclimated animals showed a similar time course in BAT vs. muscle and rectal temperature ($p > 0.15$ and > 0.17 respectively).

Table 3: MRS-derived temperature differences in BAT and Muscle and rectal temperature difference in control and cold-acclimated animals. There is no statistically significant difference between cold-acclimated and control animals.

Temperature change upon 1 hour of cooling	Control	Cold-acclimated
Rectal	$-1.9 \pm 0.9 \text{ } ^\circ\text{C}$	$-1.8 \pm 0.9 \text{ } ^\circ\text{C}$
MRS-derived muscle	$-2.4 \pm 1.5 \text{ } ^\circ\text{C}$	$-1.4 \pm 1.8 \text{ } ^\circ\text{C}$
MRS-derived BAT	$-1.2 \pm 0.9 \text{ } ^\circ\text{C}$	$+0.3 \pm 1.5 \text{ } ^\circ\text{C}$

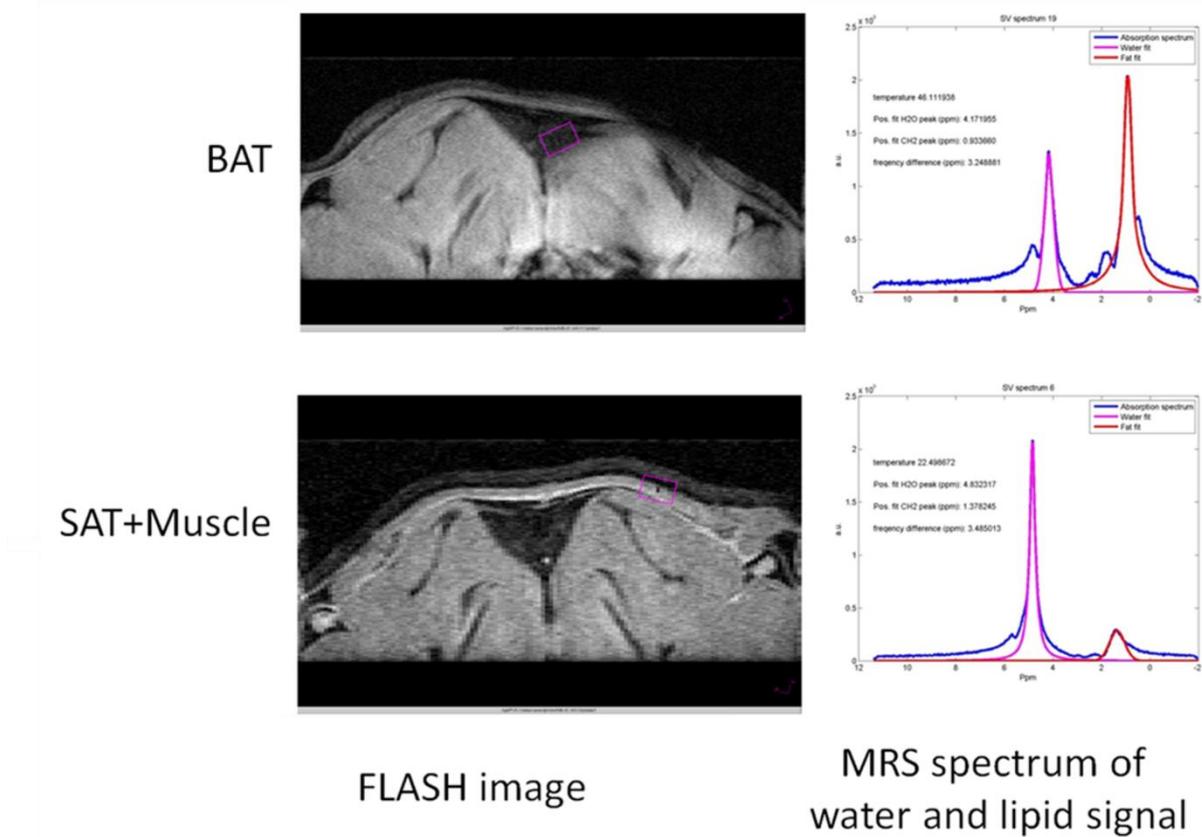


Fig 6: Images and their corresponding spectra from different positions in the body. BAT = Brown Adipose tissue, SAT = Subcutaneous Adipose Tissue.

mRNA expression and protein expression

mRNA expression of genes related to BAT metabolism (PPAR γ 2, C/EBP α , UCP1, ADRB3, DIO2, ATGL, PRDM16) were higher in iBAT compared to iWAT. Similar results were observed for GLUT4 mRNA expression when comparing iBAT to iWAT although the results did not reach statistical significance (data not shown). When examining protein expression, control animals showed higher levels of UCP1 and GLUT4 protein expression in iBAT compared to WAT, however CD36 protein expression was similar between iBAT and WAT ($p=0.43$).

Cold acclimation did not alter mRNA expression of the selected genes in BAT. PRDM16 mRNA expression decreased following cold acclimation in BAT ($p=0.013$). No changes in protein levels of UCP1, GLUT4 or CD36 in BAT were detected following cold acclimation using western blot (data not shown).

Discussion

In this study, we used a rat model to assess BAT molecular imaging with [¹⁸F]FDG, [¹⁸F]FTHA, [¹²³I]MIBG, [^{99m}Tc]TcMIBI and [¹²³I]IPA in baseline and different cold-stimulated conditions. Additionally we investigated temperature of BAT and muscle as a negative control in cold stimulated conditions by MRS and analyzed mRNA and protein expression of a number of relevant genes.

In our model, cold acclimation caused a clear growth of iBAT volume, while white interscapular adipose tissue decreased in volume. mRNA and protein levels easily allowed to distinguish brown from white adipose tissue in our study, but did not show any significant differences between control and cold-acclimated animals. Unlike reported in literature ^{26,35-37} we cannot report an up regulation of BAT specific genes (e.g. UCP1, PPAR γ , LPL). This can be caused by several reasons e.g. the biological half-life of certain mRNAs is in the order of hours and therefore dependent on diurnal variations. Timing of sampling might therefore be of high importance ^{38,39}. Furthermore, the choice of housekeeping genes is a topic of discussion. In our experiment, a broadly used housekeeping gene, GAPDH was variable throughout the experiment and therefore not suitable as reference. As we expose the animals to cold over a long period, cell type alterations in the whole fat depot are possible which can be caused e.g. by cell differentiation. In such experiments, the choice of housekeeping gene is especially difficult, as these alterations might affect the transcriptional apparatus and by that the absolute expression of the housekeeping gene might be changed ²⁶.

Protein expression levels of relevant genes did allow, just like mRNA expression levels, to distinguish BAT from WAT. However, just like mRNA expression levels, cold-acclimation did not lead to a significant change in protein expression levels.

Although we were not able to report any differences in mRNA/protein levels of BAT in cold acclimated vs. control animals as reported in literature (e.g. UCP1 ²⁶, PPAR γ ³⁵, CD36 ^{36,37})

we were able to distinguish between BAT and WAT by these techniques, which was important for *ex vivo* analysis of tissue samples.

MRS showed that the temperature drop in BAT was less pronounced after cold acclimation, which can be interpreted as an increase in BAT activity. In cold acclimated animals BAT temperature differed significantly in comparison to rectal and muscle temperature. A comparison between BAT temperatures of non cold acclimated vs. cold acclimated animals showed a clear trend towards higher temperatures in cold acclimated animals but did not reach significant differences. This is also in accordance with the results obtained from the biodistribution where cold acclimation significantly increased iBAT volume. A larger BAT depot, which results from cold acclimation, would be better suited to maintain or even increase its temperature during cold due to more active BAT.

[¹⁸F]FDG allowed iBAT visualization in warm conditions as well as in animals exposed to acute cold and cold-acclimated animals. If animals were exposed to cold, cBAT was visible as well. [¹⁸F]FTHA, [¹²³I]MIBG and [^{99m}Tc]TcMIBI poorly visualized iBAT in warm conditions, but clearly showed iBAT and to some extent also cBAT in animals exposed to acute cold and when animals were acclimated to cold. [¹²³I]IPA did not visualize iBAT in any setting and is therefore not suited as a BAT imaging tracer.

The analysis of tracer SUV_{mean} values in iBAT showed that, when compared to warm conditions, the focal uptake of [¹⁸F]FDG, [¹⁸F]FTHA and [¹²³I]MIBG in animals exposed to acute cold was about double, while for [^{99m}Tc]TcMIBI the increase was by a factor of 1.3. Considering the time window of 4 hours cold exposure, the factor of 1.3 is probably due to an increase in perfusion / blood flow and not to an increase in mitochondrial density. This confirms earlier findings of increased blood flow due to cold exposure with other tracers such as [²⁰¹Tl]thallium chloride ¹¹.

Cold-acclimation did not further increase focal uptake of any of the tracers, indicating that one single 4 h cold exposure is sufficient to reach a maximum tracer uptake in (the metabolically active part of) iBAT in our rodent model. This is in contrast to what has been reported in humans, where the SUV_{mean} of active BAT continued to increase upon chronic

cold acclimation^{27,40}. We hypothesized that this effect is induced by the fact that room temperature (22 °C) is already below the thermoneutral zone of rats (~30 °C)⁴¹ so that a slight cold acclimation already takes place under control conditions.

The volume of tissue with a high tracer uptake shows a different pattern, as the volume after cold exposure is similar to that in warm conditions and is only increased upon cold-acclimation for [¹⁸F]FDG and [^{99m}Tc]MIBI. For [¹⁸F]FTHA and [¹²³I]MIBG the increase in volume already starts after acute cold exposure, and then increases further upon cold-acclimation. These findings are to some extent not congruent with those found in literature from clinical studies, in which [¹⁸F]FTHA showed a lack of uptake under acute cold exposure and less uptake compared to FDG upon cold stimulation¹⁴. This difference in [¹⁸F]FTHA may be due to a metabolic difference in animal species (rat vs. human), but may also be due to the higher difficulty of delineating brown adipose tissue in humans compared to rats. The volume increase of active BAT is in accordance with the results obtained from MRS and the biodistribution where increased BAT temperature or volume could only be found after cold acclimation.

The total BAT activity is increased firstly by an acute cold exposure which induced an increase in metabolic activity in a small volume of iBAT (i.e. activation of existing brown adipocytes), and secondly by a cold-acclimation induced increased in metabolically active volume (i.e. increased number of activated brown adipocytes). This is different from humans, where cold acclimation leads to both an increased volume and increased SUV_{mean} value²⁷. It is possible that, compared to the necessary presence of frequently activated BAT in rodents, human BAT may be more facultative and therefore can be stimulated to a greater extent when compared to baseline room temperature conditions.

All tracer except for [¹²³I]IPA were able to visualize BAT under acute and acclimated cold conditions. With the support of the different tracer we were able to investigate different aspects of BAT metabolism and features during cold exposure and compare these results to other measurement techniques. Invasive measurements of BAT volume were coherent with results obtained by MRS and dissection of iBAT.

The current imaging procedures in humans, with different cooling protocols and different tracers in different centers, is known to confer significantly different results in terms of total BAT activity, especially when comparing different tracers ^{1,42-46}. Our data may offer an explanation for this phenomenon, as tracer uptake can be dependent on the amount of cold exposure, and the nature of this dependence seems different for each tracer.

Conclusion

$[^{18}\text{F}]\text{FDG}$, $[^{18}\text{F}]\text{FTHA}$, $[^{123}\text{I}]\text{MIBG}$ and $[^{99\text{m}}\text{Tc}]\text{TcMIBI}$, but not $[^{123}\text{I}]\text{IPA}$, are suitable for imaging aspects of BAT activity using non-invasive molecular imaging, while MRS is able to quantify heat production in BAT, even in an animal model where classical ex vivo techniques fail to show significant trends. The uptake in BAT of each tracer responds differently to acute cold exposure, and cold-acclimation does not increase BAT tracer uptake as strongly in our animal model as it does in humans. Those differences in tracer uptake characteristics should be considered during study planning as well as it should be translated to clinical applications.

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Supplementary Information

mRNA quantification

Selected tissues were harvested from 16-week old Wistar rats that were housed continuously at room temperature (n=3) and frozen in RNA-later (Qiagen) at -80 °C for later RNA isolation and gene expression analyses. The selected tissues were brown interscapular adipose tissue (iBAT), white interscapular adipose tissue (iWAT) and intraperitoneal WAT (ipWAT). Rat mRNA primer sets were PPAR γ , C/EBP α , Sirtuin1, UCP1, UCP2, ADRB3, DIO2, GLUT4, ATGL, LPL and PRDM16.

Tissues were lysed, disrupted and homogenized using the Mini-BeadBeater by violently agitating 2 ml screw-cap microvials containing small glass beads and RLT buffer (Qiagen) supplemented with 1% 2-mercaptoethanol (Sigma). Total cellular RNA was isolated from lysed cells using RNeasy Mini kit (Qiagen) according to manufacturer's instructions. Genomic DNA was removed by DNase I (Invitrogen) treatment followed by reverse transcription using random hexamer primers (Invitrogen) and Superscript III Reverse Transcriptase (Invitrogen) according to standard procedures. Primers for quantitative PCR analysis were generated using primer-BLAST software (NCBI). All primers were selected to span exon-exon boundaries with a maximal amplification length of 300 bp. Primer specificity was confirmed by melting curve analyses and product size confirmation by gel-based PCR analyses. Real-time PCR was performed with Sybr green detection (SensiMix SYBR & Fluorescein Kit, Bioline) using an iCycler iQ (Biorad) and primers (10pmol) specific for the different genes. The PCR program consisted of 10 min. initial heating at 95°C (*hot start* polymerase), followed by 35 cycles amplification (30 sec. at 95°C, 20 sec. at the optimized annealing temperature and 20 sec. at 72°C) and a final heating up to 90°C (increasing 0.5°C/ 30 sec.) for the generation of a melting curve. Quantification of a gene of interest was done by generating a standard curve using serial dilutions of a reference sample (mixed cDNA of BAT of 3 different rats). Relative expression levels were obtained by normalizing to the expression of β -actin in the corresponding sample. Each data point of the graphs was generated by

determining expression of the gene of interest on the different rats in duplicate and normalizing these data to the corresponding expression of β -actin used as housekeeping gene in our assay (β -actin was chosen as it was the most stable housekeeping gene in our experiment).

S1 Table: Relative mRNA expression levels (shown as a percentage of beta-actin expression) of different genes in iBAT, iWAT and ipWAT. Cold-acclimation = animals exposed to cold for 6h per day for 4 weeks. – indicates a value below 1%. (*) indicates a statistically significant difference between animals housed at RT or in cold-acclimated conditions, while (†) indicates a statistically significant difference between brown adipose tissue and white adipose tissue.

mRNA gene	Room temperature housing			Cold acclimation		
	iBAT	iWAT	ipWAT	iBAT	iWAT	ipWAT
PPAR γ 2 (†)	139 \pm 25	35 \pm 12	50 \pm 7	111 \pm 11	32 \pm 12	39 \pm 6
C/EBP α (†)	237 \pm 43	77 \pm 24	93 \pm 20	110 \pm 28 (*)	55 \pm 19	80 \pm 22
FOXo1	139 \pm 38	123 \pm 7	71 \pm 19	144 \pm 20	84 \pm 21	85 \pm 10
Sirtuin1	198 \pm 31	145 \pm 15	125 \pm 24	139 \pm 27	96 \pm 2	84 \pm 8
UCP1 (†)	238 \pm 43	77 \pm 24	93 \pm 20	110 \pm 28 (*)	55 \pm 19	80 \pm 22
UCP2	4 \pm 3	13 \pm 8	10 \pm 2	6 \pm 1	14 \pm 4	20 \pm 9
ADRB3 (†)	517 \pm 175	124 \pm 77	157 \pm 63	193 \pm 64(*)	50 \pm 30	25 \pm 18(*)
DIO2 (†)	5 \pm 3	-	-	6 \pm 1	-	-
GLUT4 (†)	29 \pm 5	3 \pm 1	2 \pm 1	24 \pm 5	4 \pm 1	6 \pm 1
ATGL (†)	295 \pm 78	66 \pm 55	51 \pm 21	183 \pm 48	20 \pm 11	17 \pm 5
LPL	120 \pm 46	83 \pm 38	136 \pm 20	129 \pm 18	86 \pm 41	130 \pm 30
PRDM16 (†)	125 \pm 15	11 \pm 4	7 \pm 1	55 \pm 3(*)	4 \pm 1	8 \pm 5

S2 Table: Primer sequences for RT-PCR. For each gene, a minimum of 20 bases was chosen for the primer length, which was increased where necessary to ensure specificity.

mRNA gene	Forward	Reverse
PPAR γ 2	GCCTGCGGAAGCCCTTTGGT	CAGCAAGCCTGGGCGGTCTC
C/EBP α	GGTACGGCGGGAACGCAACA	GAAGATGCCCCGCAGCGTGT
FOXo1	GGACAGCCGCGCAAGACCAG	TTGAATTCTTCCAGCCCGCCGA
Sirtuin1	GCTCGCCTTGCGGTGGACTT	GACGGCTGGAAGTGTCCGGG
UCP1	ACCTTCCCCTGGACTGAC	GCCAGGGTGGTATGGTCCC
UCP2	GCTGGGACAGCTGCCTGCAT	CGGTGCGCACTAGCCCTTGA
ADRB3	ACGAGATGGCTCCGTGGCCT	CAGCAGTTGCCTCCCACCG
DIO2	GGGCTGCGCTGTGTCTGGAA	CGGCCCATCAGCGGTCTTC
GLUT4	ACCCACCGGCAGCCTCTGAT	TAGGCTGGCTGTCCCACCCC
ATGL	TGGACGCCTGGGCATCTCCC	TGTGCTGCAGACATTGGCCTGG
LPL	AGTCTGGCTGACACTGGACAAACA	CACCCTGGGTTAGCCACCGTTTA
PRDM16	CCAATCAGGCGGGGTCTGGC	GCACCAACAGTTCCTCTCCAGGC
Beta-actin	CGACAACGGCTCCGGCATGT	TAGGGCGGCCACGATGGAG

Chapter 4:
**Synthesis, radiosynthesis and *in vitro* evaluation of
[¹⁸F]Bodipy-C₁₆/triglyceride as a dual modal imaging agent
for brown adipose tissue**

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Abstract

Background: Brown adipose tissue research is in the focus in the field of endocrinology. We designed a dual-modal fluorescent/PET fatty acid based tracer on commercially available Bodipy-C₁₆, which can be synthesized to its corresponding triglyceride and which combines the benefits of fluorescent and PET imaging.

Methods: Bodipy-C₁₆ was coupled to 1,3-diolein resulting in Bodipy-triglyceride. Bodipy-C₁₆ and Bodipy-triglyceride compounds were radiolabeled with fluorine-18 using an ¹⁸F/¹⁹F exchange reaction to yield a dual-modal imaging molecule. Uptake of radiolabeled and non-labeled Bodipy-C₁₆ and Bodipy-triglyceride was analyzed by fluorescence imaging and radioactive uptake in cultured adipocytes derived from human brown adipose tissue and white adipose tissue.

Results: Bodipy-C₁₆ and Bodipy-triglyceride were successfully radiolabeled and Bodipy-C₁₆ showed high shelf life and blood plasma stability (99% from 0-4 h). The uptake of Bodipy-C₁₆ increased over time in cultured adipocytes, which was further enhanced after beta-adrenergic stimulation with norepinephrine. The uptake of Bodipy-C₁₆ was inhibited by oleic acid and CD36 inhibitor sulfosuccinimidyl-oleate. The poor solubility of Bodipy-triglyceride did not allow stability or in vitro experiments.

Conclusion: The new developed dual modal fatty acid based tracers Bodipy-C₁₆ and Bodipy-triglyceride showed promising results to stimulate further *in vivo* evaluation and will help to understand brown adipose tissues role in whole body energy expenditure.

Introduction

Positron emission tomography (PET) allows non-invasive whole body imaging for different purposes by detecting pairs of annihilation rays. The positron emitter fluorine-18 is frequently used due to its short half-life (109 min), which makes it suitable for imaging purposes, and its broad domain in chemical reactions. Nevertheless PET is limited, already by physical laws (pathway of the positron), in spatial resolution and therefore cannot detect microscopic or subcellular structures. On the other hand optical fluorescence imaging has a high spatial resolution, making it an interesting topic for intraoperative imaging as well as *in vitro* evaluation of tracers^{1,2}. However, fluorescence imaging is lacking of high penetration depths. By combining PET and fluorescence imaging it is possible to overcome the disadvantages of both techniques and to create a new powerful tool to image from the whole-body down to sub-cellular level with the same imaging agent. The increased complexity and the effect of the fluorescent dye on the biodistribution are the major challenges when it comes to the development of a dual-modal imaging agent.

Brown adipose tissue (BAT) research has evolved vastly within endocrine research. For a long time it was thought that BAT was only present in infants but retrospective PET/CT studies with [¹⁸F]Fluoro-2-deoxy-2-D-glucose ([¹⁸F]FDG) identified active BAT in adult humans³⁻⁵. These findings could be confirmed later by dedicated cold exposure studies where a direct correlation between cold exposure and BAT metabolic activity, measured through [¹⁸F]FDG uptake, was reported⁶⁻⁸. The potential of BAT to combat obesity and obesity-associated diseases makes BAT an interesting target⁹.

The variety of quantification approaches of BAT volume and metabolic activity reaches from *in vitro* experiments¹⁰ over invasive imaging with fluorescence probes¹¹ or tritiated compounds¹² to non-invasive experiments with PET^{3-5,13,14}, SPECT^{15,16} and MRI¹⁷⁻¹⁹. Even though fatty acids (FAs) are the main fuel source for adipocytes, [¹⁸F]FDG is mostly used in studies to quantify BAT activity^{4,20,21}. FA uptake is more difficult to quantify because there is a large variety of different FAs and triglycerides (TGs) present in the human body which

makes the uptake dependent on the affinity of the single FA and not on the substance class itself. Nevertheless FAs are the major metabolized substances in BAT and therefore it is possible that BAT activity and lipid uptake is largely underestimated by FDG scans (which only show glucose-related uptake) ¹³. Therefore the need exists to use a FA-based BAT tracer to quantify BAT activity and FA uptake, to study uptake dynamics and to exclude the chance of underestimating BATs metabolic activity with FDG scans.

Radiolabeled FAs in general have been developed in several variations for imaging purposes (e.g. FTHA (14(R,S)-[¹⁸F]fluoro-6-thia-heptadecanoic acid), BMIPP (beta-methyl-p-[¹²³I]iodophenylpentadecanoic acid), [¹¹C]palmitate ²²⁻²⁶). We here report the development of a FA-based tracer which is suitable for both, PET and fluorescence imaging from the fluorescent FA Bodipy-C₁₆ (BDP-FA) with which it is possible to image from whole body to sub-cellular level. Bodipy dyes have been already used to image brown adipose tissue ¹¹ and it has been proven that fatty acid transport proteins (FATP) have a preference for Bodipy-FL coupled to a long carbon chain (C ≥ 8) ²⁷. Furthermore, downstream metabolic reactions in white adipocytes were already visualized ²⁸. Since we did not want to decrease the good binding properties of BDP-FA by introduction of another chelator molecule, we got interested in ¹⁸F/¹⁹F exchange reactions used to transform fluorescent dyes into dual-modality PET/fluorescent imaging dyes ²⁹⁻³³. Because the FA is only modified at the end of the carbon chain, neither an increased steric demand, nor lowering of the targeting efficiency is expected. In comparison to previous reports, we want to go a step further and synthesize also the, *in vivo* predominant, triglyceride form of the [¹⁸F]BDP-FA. Here we describe a synthetic approach resulting in a dual-modal molecule to visualize BAT *in vivo* and *in vitro* with the same tracer which should help to understand this not completely evaluated tissue, its functions and metabolism.

Materials and Methods

Commercially available compounds were used without further purification unless otherwise stated. BODIPY-FL-C₁₆ was purchased from Thermo Fischer Scientific (99%) (Netherlands). 1,3-diolein was purchased from Sigma Aldrich ($\geq 99\%$). DMEM/F-12 was purchased from ThermoFischer (Waltham, MA).

All HPLC purifications (1.0 mL/min, solvent A; 0.1% TFA in water, solvent B; CH₃CN, 50°C) were performed on a Shimadzu UFLC HPLC system equipped with a DGU-20A₅ degasser, a SPD-M20A UV detector, a LC-20AT pump system, a CBM-20A communication BUS module, a CTO-20AC column oven, and a Scan-RAM radio-TLC/HPLC-detector from LabLogic using an Aeris™Widepore column (XB-C18, 3.6 μ m, 4.6 mm \times 250 mm) for the BDP-FA or an Aeris™Widepore column (C4, 3.6 μ m, 4.6 mm \times 250 mm) for the Bodipy-triglyceride (BDP-TG). ESI-MS was performed on a Applied Biosystems SCIEX API 150 EX electrospray ionization quadrupole (ESI-Q) mass spectrometer with the method of McAnoy et al.³⁴. Briefly, 0.1M aqueous ammonium acetate solution was added to the probe to observe the ammonium salt in the MS.

¹H-NMR spectra were carried out on a Bruker Ultrashield *TH 400 plus* at 400 MHz. Tol-d₈ was used as solvent with TMS as internal standard. Chemical shifts are reported in parts per million (ppm) relative to the internal standard.

Synthesis of BDP-TG 2

BDP-FA **1** (300 μ g, 0.6 μ mol) in acetonitrile was evaporated to complete dryness before the reactant was reconstituted in toluene (100 μ L). To the resulting solution SOCl₂ in toluene (100 μ L, 4 vol.-%) was added, incubated for 5 min at 70°C in a closed vial and evaporated. The product was reconstituted in toluene (50 μ L) containing 1,3-diolein (2 μ L, 2.8 μ mol) and heated to 100°C for 30 min. After the reaction time, purification by HPLC (1 mL/min, 30% to 15% A in 5 min, 15% to 0% A from 5 to 6 min, 0% A to 20 min) yielded **2** (225 μ g, 75%) as a red solid; t_R = 12.3 min. ESI-MS(+): m/z (%) = 1058 (100) [M - F]⁺, 1095 (82) [M + NH₄]⁺. ¹H

NMR (400 MHz, Tol- d_8); δ (ppm) = 5.46 (m, 4H), 4.26 (m, 2H), 4.06 (m, 2H), 3.13 (m, 1H), 1.75 (s, 3H).

Additional experiments were performed using BMIPP (beta-methyl-p-iodophenylpentadecanoic acid) as starting FA. Different ways of synthesis were evaluated, where reaction time, temperature and chlorinating agent were changed (see results section).

Radiolabeling of Bodipy-C₁₆ 3

Aqueous fluorine-18 solution was loaded on a QMA-cartridge which was preconditioned with 15 mL K_2CO_3 in water and 20 mL water. Fluorine-18 (42 MBq) was eluted with a mixture of 600 μ L acetonitrile, 300 μ L H_2O and 100 μ L K_2CO_3 solution (5 mg/mL). Fluorine-18 solution was transferred into a drying vessel containing tetra-n-butylammonium bromide (80 μ L) as a phase transfer agent. Acetonitrile (3 \times 1.0 mL) was added and the solution of fluorine-18 was dried by heating to 100 °C with a continuous flow of argon. After reconstitution of fluorine-18 in anhydrous acetonitrile (100 μ L), a solution of **1** (50 μ g, 0.1 μ mol) and $SnCl_4$ (0.2 M in acetonitrile, 100 μ L) was added to the activity solution and the reaction was stirred at r.t. for 30 min. After addition of water (200 μ L) and filtration (Millex, hydrophile PVDF 0.22 μ m) a quality control was performed by HPLC (1 mL/min, 30% to 15% A in 5 min, 15% to 0% A from 5 to 6 min, 0% A to 20 min) and afforded **3** (decay corrected radiochemical yield (RCY): 76%, 22 MBq) with a decay corrected specific activity of 220 MBq/ μ mol and a radiochemical purity of \geq 99%; t_R = 13.3 min. In addition a TLC with Toluene, $CHCl_3$ and MeOH (80.9%, 14.3%, 4.8%) was performed.

Radiolabeling of Bodipy-TG 4

Drying process was performed as mentioned in previous section. After reconstitution of fluorine-18 (83 MBq) in anhydrous acetonitrile (100 μ L), a solution of **2** in toluene (107 μ g, 0.1 μ mol in 50 μ L) and $SnCl_4$ (0.2 M in acetonitrile, 100 μ L) was added to the solution with the activity and the reaction solution was stirred at room temperature (r.t.) for 30 min. After addition of water (200 μ L), filtration and washing with water (2 \times 200 μ L) a quality control was

performed by HPLC (1 mL/min, 30% to 15% A in 5 min, 15% to 0% A from 5 to 6 min, 0% A to 20 min) and afforded **4** (decay corrected RCY: 44%, 25 MBq) with a decay corrected specific activity of 250 MBq/μmol and a radiochemical purity of > 95%; t_R = 12.5 min. In addition TLC with toluene, CHCl₃ and MeOH (80.9%, 14.3%, 4.8%) was performed.

Human primary adipocyte cultures derived from BAT and white adipocytes (WAT)

The isolation and differentiation of human adipocytes has been described before³⁵. The study was reviewed and approved by the ethics committee of Maastricht University Medical Center (METC 10-3-012, NL31367.068.10). Informed consent was obtained prior to surgery. The precursor cells were obtained from a stromal vascular fraction of BAT and WAT tissue. The sample was taken from a 34 year old male with a benign formation in the left thyroid gland. The cells have been metabolically characterized³⁵. In short, the stromal vascular fraction was obtained from BAT and subcutaneous WAT biopsies from the same individual undergoing deep neck surgery. Collected cells were grown to confluence prior to start of differentiation in DMEM/Ham's F-12 (Gibco) supplemented with 10% Fetal Bovine Serum (Bodinco BV). Differentiation medium composed of biotin (33 μM), D-pantothenate (17 μM), h-insulin (100 nM), dexamethasone (100 nM), IBMX (250 μM), rosiglitazone (5 μM), T3 (2 nM), transferrin (10 μg/ml). After 7 days of differentiation, the medium was exchanged for maintenance medium composed of biotin (33 μM), pantothenate (17 μM), insulin (100 nM), dexamethasone (10 nM), T3 (2 nM), and transferrin (10 μg/ml).

In vitro uptake of **3 and **4** in cultured adipocytes derived from human BAT and WAT**

Adipocytes derived from human BAT and WAT were incubated with **3** and **4** (1 nM, 100 μL) for 1 h. **3** was added in 0.5% fatty acid free bovine albumin in DMEM/F-12. **4** was dissolved in 50% PEG in DMEM/F-12. Adipocytes were exposed to the tracer or preincubated with sulfosuccinimidyl oleate (500 μM, 100 μL, 30 min) or norepinephrine (1 μM, 100 μL, 30 min). Radioactivity was quantified by measuring washed, detached cells with a WIZARD² automatic γ-counter from Perkin Elmer. Number of cells was determined by three extra wells

which underwent a standard cell count protocol using a LUNA II (Logosbio) automated cell counter.

In vitro imaging of cultured adipocytes derived from human BAT and WAT

Adipocytes derived from human BAT and WAT were incubated with **1** and **2** (0-4 μM , 500 μL) for different time points (1-24 h) in experimental medium. **1** was added in 0,5% fatty acid free bovine albumin in DMEM/F-12. **2** was dissolved in 50% PEG in DMEM/F-12. Adipocytes were exposed to the tracer or coincubated with oleic acid (400 μM , 500 μL) or preincubated with sulfosuccinimidyl-oleate (500 μM , 500 μL , 30 min). After incubation adipocytes were fixed in 3.7% formaldehyde (4°C, 30 min) and stained with DAPI (4',6-diamidino-2-phenylindole) (RT, 5 min). The adipocytes were imaged with a Sony Eclipse e800 fluorescence camera. The signal was measured in the FITC channel (460-490 nm excitation, 510-550 nm emission) and in the DAPI channel (385-415 nm excitation, 450-470 nm emission). Quantification of signal was performed with Fiji ³⁶ as well as correction for exposure time, background and cell auto-fluorescence. For every data point at least three cells were analyzed. Graphs and statistical analysis was performed using GraphPad Prism 6.

Fluorescence measurements (microplate reader)

Adipocytes derived from human BAT cultured on 96 well chamber slides were incubated with **1** and **2** (2 μM , 100 μL) for different time points (1-4 h). **1** was added in 0,5% fatty acid free bovine albumin in DMEM/F-12. **2** was dissolved in 50% PEG in DMEM/F-12. Adipocytes were exposed to the tracer or preincubated with norepinephrine (1 μM , 500 μL , 30 min). After incubation adipocytes were washed 3 times (30 seconds each) with cold PBS and suspended in PBS. The signal was measured using a SpectraMax M2 plate reader (molecular devices) (excitation 485 nm, emission 520 nm). Graphs and statistical analysis was performed using GraphPad Prism 6.

Ex vivo stability

Shelf life of **3** was investigated under following conditions: 24 MBq (60 μ L) were added to H₂O (1 mL) and integrity was examined by HPLC (1-4 h). Areas under the peak of free fluorine-18 and metabolites were compared to the area under the peak of **3**.

Plasma stability was examined by adding 28 MBq (60 μ L) of **3** to 1 mL of human plasma (37°C). After certain time points (20, 40, 60 min and 2-4 h) integrity of the compound was analyzed by HPLC.

Determining the aqueous or plasma stability of the radiolabeled TG **4** failed due to the insolubility of the compound in aqueous medium.

Results

Different conditions for the esterification of the BDP-C₁₆ to BDP-TG were tested (Table 1). The best yield, analyzed by HPLC injection of the crude reaction mixture, was achieved with BDP-C₁₆ in Toluene (50 μ L) and the strong chlorinating agent thionylchloride at 70°C in 30 min. More hydrophilic solvents, larger reaction volumes, lower temperatures or milder chlorination conditions resulted in lower yields and/or byproducts. In optimal conditions, after reaction and purification **2** was obtained in a yield of 75%. After purification of **2** further analysis of quenching effects was performed by HPLC. Both, FA and TG, showed similar absorption when the same molar amount was injected. NMR of **2** confirmed the identity of the TG (Fig 1a and S1 Fig 4). The disappearance of the alcohol function (3.75 ppm) with the simultaneous formation of a third ester bond (3.15 ppm) plus the additional BDP peaks (1.75 ppm and 2.47 ppm) proved the successful synthesis of **2**. Additionally ESI-MS of **2** supports these findings as peaks found in the spectrum correspond with the calculated molecular masses (Fig 1b).

Table 1: Different esterification conditions for BMIPP or BDP-C16. Yields tested by injection of the crude reaction mixture into HPLC and comparison of the esterified vs. the non-esterified FA signal plus byproducts

Fatty acid	Coupling agent	Solvent	Rec. volume [μ L]	Activator	Temperature [°C]	Reaction time [min]	Yield [%]	n
BMIPP	1,3-Diolein	Toluene	50	SOCl ₂	70	30	72	4
BMIPP	1,3-Diolein	Acetonitrile	50	SOCl ₂	70	30	0.6	1
BMIPP	1,3-Diolein	Diethylether	50	SOCl ₂	70	30	67	1
BMIPP	1,3-Diolein	Tetrachlormethane	50	SOCl ₂	70	30	8	1
BMIPP	1,3-Diolein	Benzene	50	SOCl ₂	70	30	38	1
BMIPP	1,3-Diolein	Toluene	50	Oxalyl Chloride	70	30	43	1
BMIPP	1,3-Diolein	Toluene	50	Oxalyl Chloride	70	60	44	1
BMIPP	1,3-Diolein	Toluene	50	SOCl ₂	70	90	54	2
BMIPP	1,3-Diolein	Toluene	50	SOCl ₂	70	120	67	2
BMIPP	1,3-Diolein	Toluene	50	SOCl ₂	70	150	86	2
BMIPP	EtOH	MeCN	100	SOCl ₂	0	30	1	1
BMIPP	EtOH	MeCN	100	SOCl ₂	22	30	64	1
BMIPP	EtOH	MeCN	100	SOCl ₂	70	30	88	1
Bodipy C ₁₆	1,3-Diolein	Toluene	200	SOCl ₂	70	30	59	1
Bodipy C ₁₆	1,3-Diolein	Toluene	50	SOCl ₂	70	30	95	8

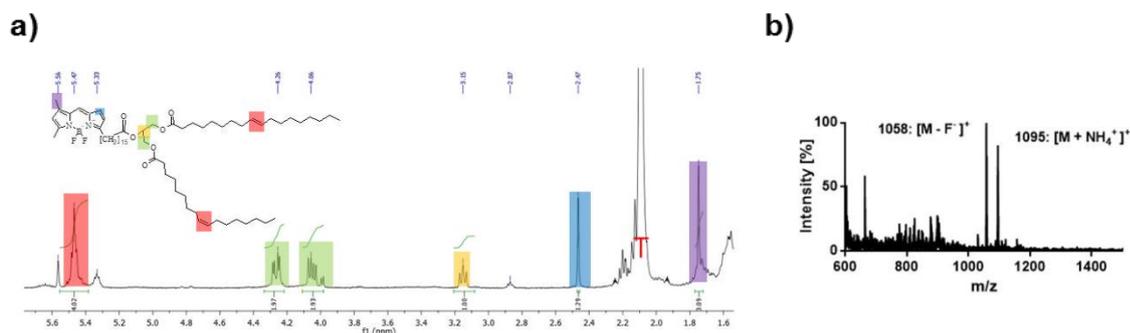


Fig 1: BDP-TG analysis. a) $^1\text{H-NMR}$ of BD-TG in toluene- d_5 , red T corresponds to the toluene peak b) ESI-MS of BDP-TG with NH_4OAc in Chloroform/MeOH 50:50

The radiolabeled dual-modality imaging agent [^{18}F]BDP-TG was synthesized in a two-step procedure. First the Bodipy-FA **1** was esterified to BDP-TG **2** and then radiolabeled with fluorine-18 using the strong Lewis acid SnCl_4 in dry acetonitrile/toluene. After 30 min of incubation, [^{18}F]BODIPY-TG **4** was isolated by washing 3 times with water allowing 44% radiochemical yield, a specific activity of 250 MBq/ μmol and a radiochemical purity of > 95% (Fig 2b). First approaches to radiolabel the BDP-FA and perform the esterification afterwards resulted in higher labeling yields (76%) (Fig 2a) in the first step but the following esterification afforded only small amounts of radiolabeled BDP-TG. The radiolabeling was performed in acetonitrile but in previous cold esterification attempts it was shown that only small yields can be reached in this solvent (Table 1). A complete exchange of acetonitrile to toluene is time consuming as well as the problem of [^{18}F]BDP-FA getting attached to the glass vial arises. Different reaction conditions for the radiolabeling can be found in Table 2.

Table 2: radiolabeling of BDP-FA **3** and BDP-TG **4**. Different conditions tested for both compounds.

Radiochemical yields calculated after purification

Agent	V (SnCl ₄) [μL]	Solvent	Volume [μL]	Reaction time [min]	RCY [%]	n
Bodipy C ₁₆	100 (0.1M)	MeCN + K ₂₂₂ + K ₂ CO ₃	250	30	7.3	4
Bodipy C ₁₆	30 (0.1M)	MeCN + K ₂₂₂ + K ₂ CO ₃	180	30	13.8	2
Bodipy C ₁₆	100 (0.2M)	MeCN + K ₂₂₂ + K ₂ CO ₃	250	30	28	4
Bodipy C ₁₆	100 (0.3M)	MeCN + K ₂₂₂ + K ₂ CO ₃	250	30	0	1
Bodipy C ₁₆	100 (0.2M)	MeCN + K ₂ CO ₃	250	30	76	9
Bodipy-Tg	100 (0.2M)	MeCN + K ₂ CO ₃ /DMA	250	30	0	1
Bodipy-Tg	100 (0.2M)	MeCN + K ₂ CO ₃ /Toluene	250	30	44	7

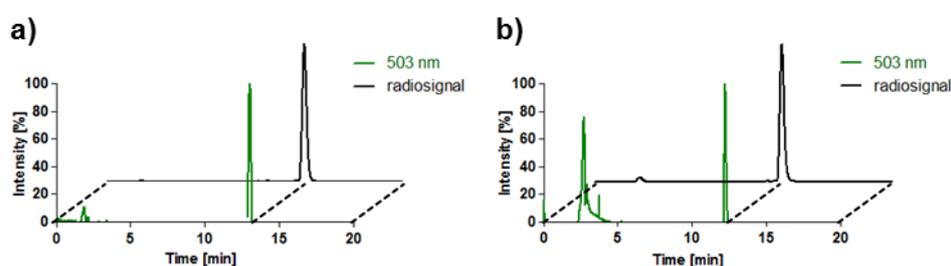


Fig 2: chromatograms for BDP-FA/TG a) HPLC chromatograms (radio trace and absorbance at 503 nm) of ¹⁸F-BDP-FA **3** on a C18 column. b) HPLC chromatograms (radio trace and absorbance at 503 nm) of [¹⁸F]BDP-TG **4** (peak at 3 min corresponds to toluene which was used to solubilize the probe) on a C4 column.

Shelf life and plasma stability showed 99% intact radiolabeled compound **3** after 4 h (S1 Fig 5). Fluorescent uptake of **1** was evaluated *in vitro* with cultured adipocytes derived from human BAT. Adipocytes were incubated with BDP-FA (2 μM) for 1 to 4 h and fluorescence was measured with a microplate reader (SpectraMax M2) (Fig 3a). An increase over time was observed where BAT took up 274%, 362% and 697% more after 2 h, 3 h and 4 h compared to the 1 h time point. Activation with norepinephrine (1 μM, 30 min before incubation with BDP-FA) increased BAT uptake of **1** by 134%, 199% and 143% after 2, 3 and 4 h compared to their basal uptake values at these time points. Only after 1 h no significant increase was observed. Single cell uptake was analyzed and quantified with a fluorescence microscope (Sony Eclipse e800) under different conditions (S1 Fig 1). Concentration dependent uptake was examined within a 2 h time period (Fig 3b). A clear Michaelis -

Menten like kinetic was observed in uptake-positive BAT cells (with $k_m = 1.15 \mu\text{M}$, $R^2 = 0.93$). Coincubation with $400 \mu\text{M}$ oleic acid reduced uptake of BDP-FA by 46%.

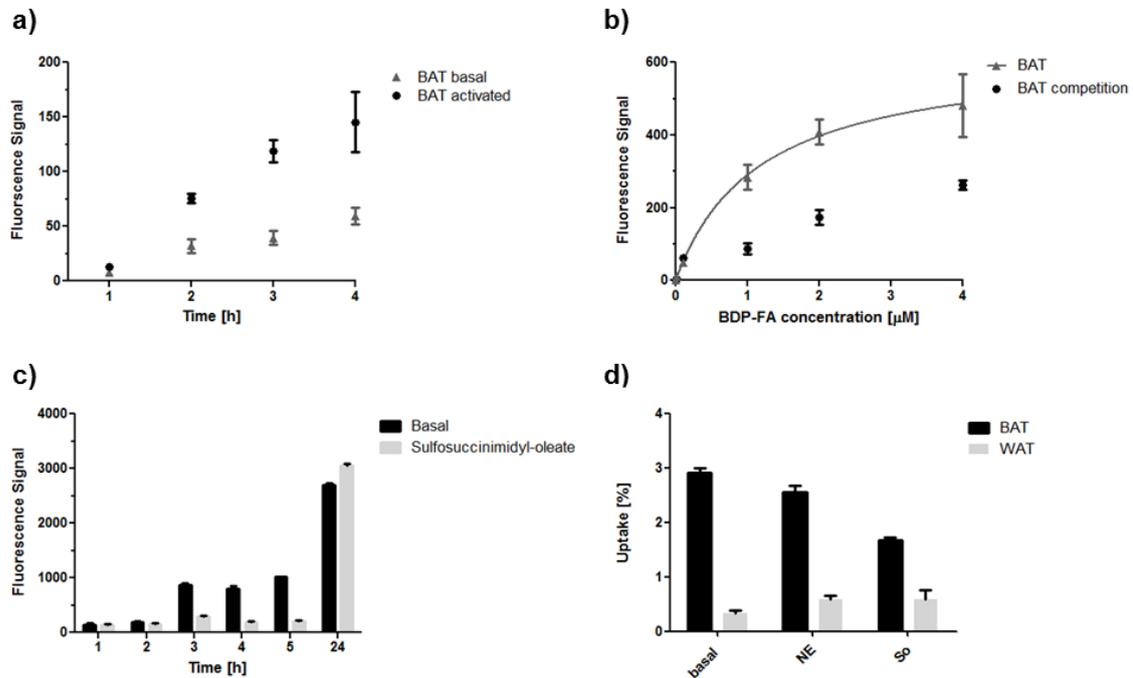


Fig 3: *in vitro cell* uptake a) uptake time dependence measured with $2 \mu\text{M}$ 1 in BAT with and without activation with norepinephrine (microplate) b) uptake concentration dependence measured in BAT over 2 h with and without coincubation with oleic acid (fluorescence microscope) c) BAT blocking study over time with sulfosuccinimidyl-oleat (fluorescence microscope) d) uptake of 3 in BAT and WAT cells under basal conditions, activation with Norepinephrine (NE) and blocking with sulfosuccinimidyl-oleat (So) after 1 h incubation time

Time dependence with simultaneous inhibition of the cluster of differentiation 36 (CD36), which is a scavenger protein for long chain fatty acids³⁷ and was shown to be essential for thermogenesis in animals during fasting conditions¹⁶, was additionally quantified on a cellular level (Fig 3c) (S1 Fig 1). By preincubation with sulfosuccinimidyl-oleate a blocking of $73\% \pm 6\%$ was reached after 3 – 5 h. After 24 h the uptake recovered to basal conditions. No uptake was observed by incubation with 2 ($2 \mu\text{M}$, 24 h) (S1 Fig 3).

Cell uptake experiments were also performed in WAT cells to further analyze the suitability of the BDP-FA as a tracer for adipocytes and to compare BAT and WAT cells. The WAT cells were obtained from the same human sample as the BAT cells.

In first microplate activation experiments it was shown that, similar to BAT, the uptake of **1** increased over time. After 2 h, 3 h and 4 h the uptake increased by 221%, 294% and 357%. Activation of the cells by norepinephrine resulted in an increase in uptake between 10% and 48% and was significant lower compared to BAT cells.

In WAT cells incubated with **1** (0-4 μM) for different concentrations, a Michaelis - Menten like kinetic was observed in uptake-positive cells (with $k_m = 1.58 \mu\text{M}$ and $R^2 = 0.97$). The uptake was also reduced with an excess of oleic acid (400 μM) by 53%.

CD36 dependency on uptake kinetics was also investigated in WAT cells (S1 Fig 2). Sulfosuccinimidyl-oleat only had a significant negative effect on the uptake of the 2 h and 3 h group ($51\% \pm 16\%$) but in general the uptake kinetics of the preincubated cells with sulfosuccinimidyl-oleat followed more the kinetics of the cells under basal conditions.

BAT and WAT uptake of **3** and **4** was investigated to prove the functionality of the radiolabeled compounds and to quantify the total uptake of both tracers. Interestingly, in this experimental setup, BAT took up in all cases more FAs than WAT (Fig 3d). After 1 h WAT took up $0.3\% \pm 0.05\%$ and BAT $2.9\% \pm 0.09\%$. Activation of WAT with norepinephrine showed a small increase in uptake ($0.6\% \pm 0.07\%$ vs. $0.3\% \pm 0.05\%$) whereas the uptake in BAT was slightly decreased ($2.6\% \pm 0.11\%$ vs. $2.9\% \pm 0.09\%$). An effect could be observed for preincubation with sulfosuccinimidyl-oleat in BAT after 1 h ($1.7\% \pm 0.04\%$ vs. $2.9\% \pm 0.09\%$). WAT showed no significant decrease ($0.58\% \pm 0.18\%$ vs. $0.3\% \pm 0.05\%$).

Incubation experiments with the radiolabeled TG **2** were conducted additionally but showed no consistent and reproducible results (data not shown).

Discussion

Imaging of BAT has evolved to an interesting and fast developing topic in endocrine research. Several imaging approaches have been used in the past to visualize and quantify BAT and its metabolic activity reaching from *in vitro* experiments¹⁰ over invasive imaging with fluorescence probes¹¹ or tritiated compounds¹² to non-invasive experiments with PET^{3-5,13,14}, SPECT^{15,16} and MRI¹⁷⁻¹⁹. Most often [¹⁸F]FDG scans are used for BAT imaging. This has the big disadvantage that it only shows glucose-related uptake and has the chance to underestimate BAT activity because the major metabolized substance class in BAT is FAs. There is therefore a need for a FA BAT tracer to quantify metabolic activity in a more precise way. In studies with [¹⁸F]FTHA it was observed that radiolabeled FAs showed an increased uptake in BAT under cold stimulation in humans¹³. With our developed tracer it is possible to examine uptake characteristics with both PET and fluorescence imaging. Therefore scans from subcellular level, to determine the localization of the probe within the cell, to whole body scans are possible.

In vivo FAs are transported as TGs in lipoproteins towards adipocytes^{38,39}. We were able to produce BDP-TG in a very reasonable yield (77%). Both, the FA and the TG analog, were successfully radiolabeled (RCY: 76% for [¹⁸F]BDP-FA and 44% for [¹⁸F]BDP-TG), comparable to previously achieved yields with other modified Bodipy-dyes by other groups^{31,40}. Activation of fluorine-18 by K₂CO₃ and K₂₂₂ (4,7,13,16,21,24-Hexaoxa-1,10-diazabicyclo[8.8.8]hexacosane) resulted in side products and decomposition of the Bodipy-dye. Reactions without addition of a base, such as K₂CO₃, gave lower labeling yields. Shelf life and plasma stability of [¹⁸F]BDP-FA indicated its suitability as an *in vivo* imaging agent, although these values could not be determined for [¹⁸F]BDP-TG due to its insolubility in aqueous medium. Nevertheless no large stability deviations are expected for [¹⁸F]BDP-FA and [¹⁸F]BDP-TG, as the lipid backbone should not affect the boron-fluoride bond in a large extend. Next to our synthesis work we put effort in first *in vitro* application of our cold tracer. Uptake experiments proved that the BDP-FA is taken up with the characteristics of a regular

FA ($k_m = 1.15 \mu\text{M}$ compared to $k_m = 0.2 \mu\text{M}$ of oleate⁴¹ and cis-parinaric acid: $k_m = 1.5 \mu\text{M}$ in 3T3 fibroblasts⁴²) and uptake could be decreased by coincubation with oleic acid in excess. BAT cells were sensitive to norepinephrine and uptake was increased by a preincubation with this hormone. Sulfosuccinimidyl-oleat as an antagonist blocked CD36 and overall uptake of BDP-FA was decreased to a minimum. This shows that CD36 is essential for the lipid uptake in BAT cells and is in accordance to already published *in vivo* results^{43,44}. Only after 24 h the cells were able to overcome this blocking procedure and showed comparable uptake to basal conditions. Stahl et al. found over 90% of CD36 in serum starved adipocytes on the cell membrane⁴². Therefore it can be speculated that the increase in uptake resulted in a displacement of sulfosuccinimidyl-oleat at this time point.

The uptake in WAT cells of BDP-FA was also analyzed. Comparable to BAT cells fluorescence signal increased over time and uptake was partially blocked by coincubation with an excess of oleic acid. Neither the positive uptake effect in cells preincubated with norepinephrine nor the negative uptake effect in cells preincubated with sulfosuccinimidyl-oleat was observed in such an extent in WAT compared to BAT. Therefore a smaller amount of β 3-adrenoreceptors on the cell surface⁴⁵ and a CD36 independent uptake mechanism⁴⁶ is proposed for WAT cells.

In former experiments (data not shown) it was observed that the FA and the TG signal are significantly quenched in an aqueous medium. This is in accordance with other published results for the BDP-FA⁴⁷. A self-quenching effect for different Bodipy-dyes depending on their solvent solubility has also been reported before⁴⁸. This makes fluorescent imaging with this tracer susceptible for misinterpretation because one can underestimate the total uptake. Nevertheless we could easily and without any dose characterize, but not quantify, *in vitro* uptake kinetics of this fluorescent FA.

Even though first indications on uptake mechanism in brown and white adipocytes were obtained by fluorescent experiments, radiolabeled compounds were necessary to quantify cell uptake and to exclude the chance of a quenched fluorescent signal. Cell uptake experiments with the radiolabeled FA showed that after correction for different cell numbers,

BAT took up significantly more [^{18}F]BDP-FA than WAT. Sulfosuccinimidyl-oleat decreased the uptake significantly after 1 h in the BAT group. By preincubation with norepinephrine the uptake in BAT was not affected (as seen in the fluorescent experiments, increased uptake could only be observed 2 h after stimulation). In WAT no significant effect was observed for the preincubation with sulfosuccinimidyl-oleat. Preincubation with norepinephrine resulted in the same uptake value as preincubation with sulfosuccinimidyl-oleat, which denotes to a norepinephrine insensitive and CD36 independent uptake mechanism. Overall, the results obtained with the radiolabeled BDP-FA are in accordance with the results obtained in the fluorescent experiments. By this method it was possible to quantify uptake values in percent uptake and a comparison between both cell types could be performed.

For the TG no uptake was observed with the presented methods. This might be due to the insolubility of TG in aqueous solutions and it is therefore not available for uptake by adipocytes. Alternatively, another reason could be that lipoprotein lipase, which is responsible for TG hydrolysis, is only activated by lipoproteins, which was not present in our *in vitro* experiments.

A possible solution for the inaccessibility of the BDP-TG tracer, non-radiolabeled and radiolabeled, for our *in vitro* cell model would be the incorporation of this tracer into lipoproteins. As TG are transported *in vivo* in these water soluble particles an incorporation of our tracer would 1) simulate the physiologic state how TG are transported and 2) overcome the limited solubility of the compound, which also limits the *in vivo* application. With other FA-tracers (mainly tritiated) this has already been achieved and *in vivo* experiments have been performed^{11,12}. In comparison to tritiated compounds, the here presented tracer has the advantage that it can be visualized and quantified by PET *in vivo* and its uptake mechanism and kinetics can be followed by fluorescence microscopy *in vitro*.

To conclude, we present the successful development of a multimodal FA/TG BAT tracer. We were able to conjugate BDP-C₁₆ to 1,3-diolein and radiolabel this triglyceride. We then investigated its uptake characteristics *in vitro* with fluorescence imaging in a human BAT and WAT sample, and observed CD36 mediated uptake which was sensitive to norepinephrine in

BAT cells. Additionally, we showed that BAT takes up significantly more FAs than WAT in our *in vitro* radioactive uptake experiments.

Future *in vitro/vivo* experiments are required with [¹⁸F]Bodipy-TG, incorporated into a chylomicron, where these first insights in brown adipose tissue metabolism will help to specify the role of this interesting tissue for whole body energy metabolism.

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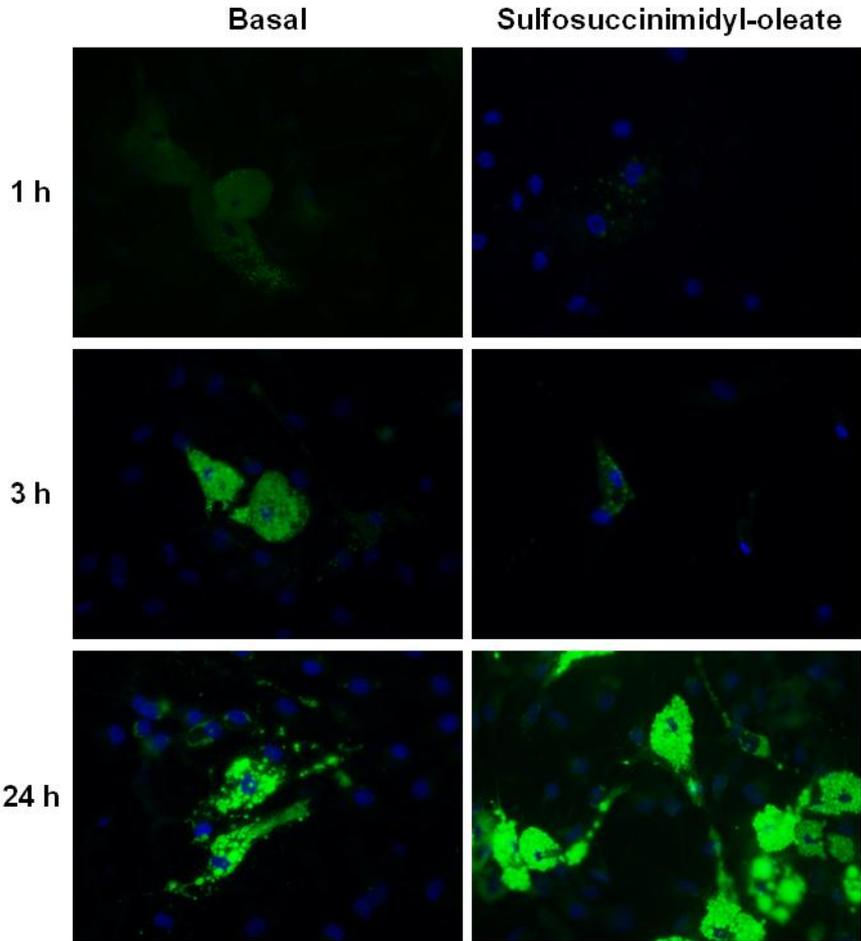
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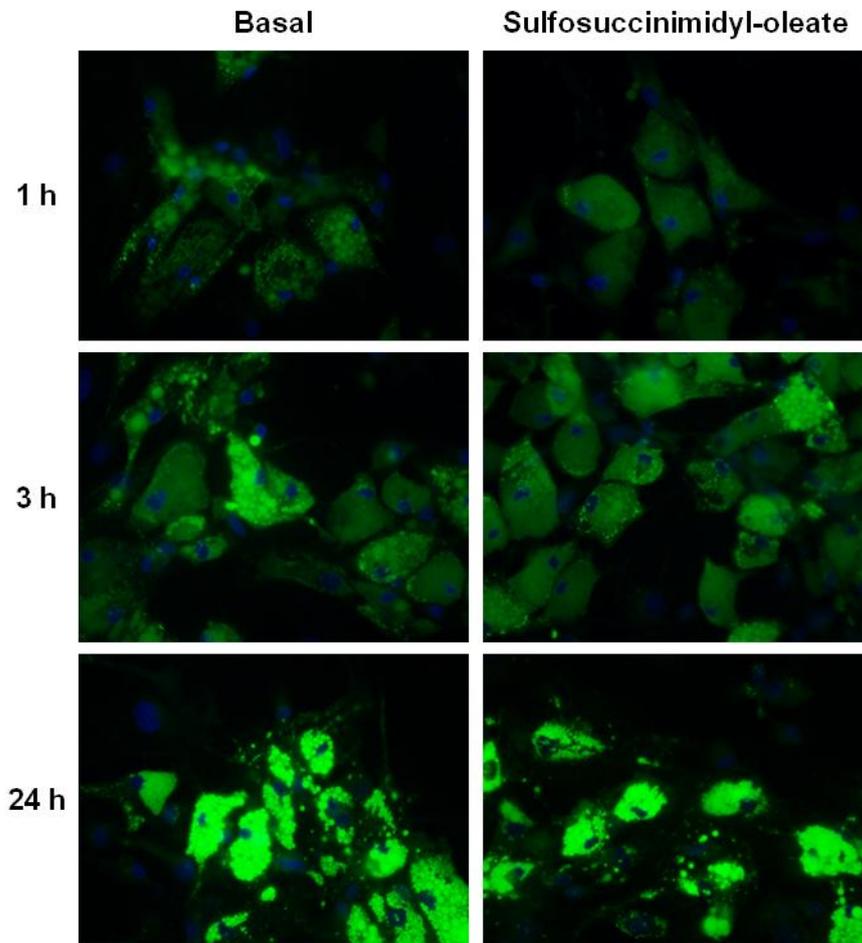
Supplementary Information

BAT

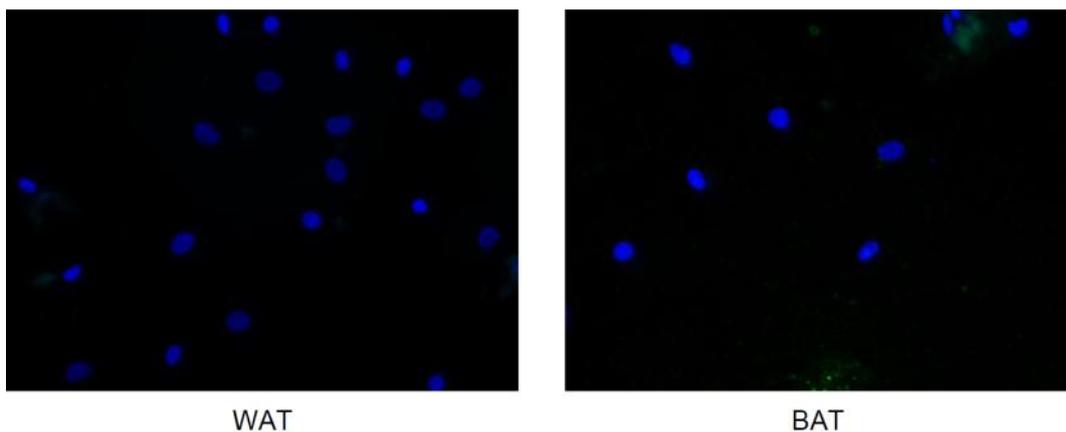


S1 Fig 1: uptake of 1 in BAT images obtained from fluorescence microscope by incubation with BDP-FA under basal conditions, and sulfosuccinimidyl – oleat incubation. Green (Bodipy-signal): 460-490 excitation, 510-550 emission, Blue (Dapi-signal): 385-415 excitation, 450-470 emmision.

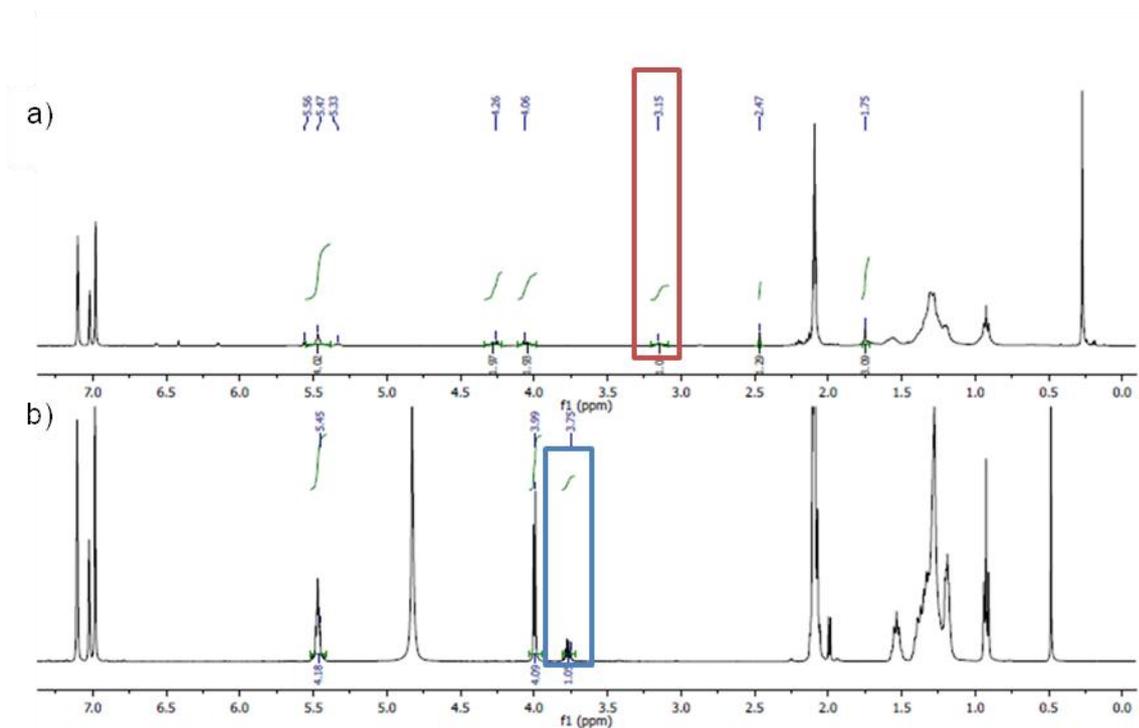
WAT



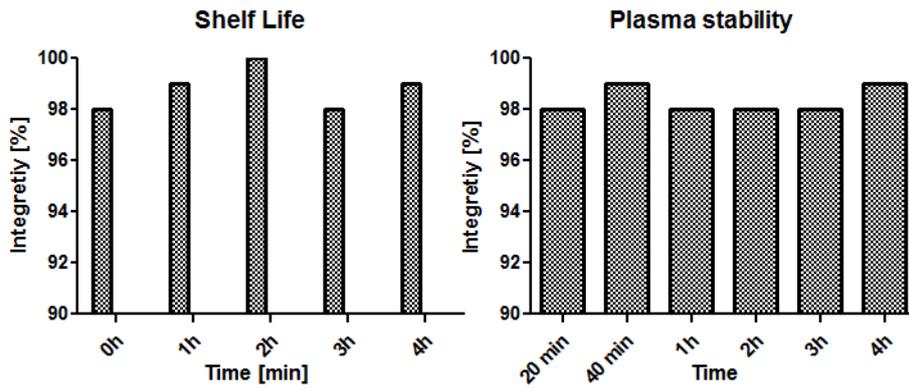
S1 Fig 2: uptake of 1 in WAT images obtained from fluorescence microscope by incubation with BDP-FA under basal conditions and sulfosuccinimidyl – oleat incubation. Green (Bodipy-signal): 460-490 excitation, 510-550 emission, Blue (Dapi-signal): 385-415 excitation, 450-470 emmision.



S1 Fig 3: uptake of 2 in WAT and BAT images obtained from fluorescence microscope by incubation with BDP-TG (2 μ M, 24 h) under basal conditions. Green (Bodipy-signal): 460-490 excitation, 510-550 emission, Blue (Dapi-signal): 385-415 excitation, 450-470 emission.



S1 Fig 4: NMR analysis NMR spectra of BDP-TG 1(a) and 1,3-Diolein (b), alcohol function of the Diolein is shown in red square, formed triple ester bond is shown in blue square. No significant impurities were noted.



S1 Fig 5: Shelf Life and Plasma Stability of 3

Chapter 5:
[¹⁸F]BODIPY-triglyceride-containing chylomicron-like particles as an imaging agent for brown adipose tissue *in vivo*

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Abstract

Background: Brown adipose tissue (BAT) is present in human adults and the current gold standard to visualize and quantify BAT is [^{18}F]FDG PET-CT. However, this method fails to detect BAT under insulin-resistant conditions associated with ageing and weight gain, such as type 2 diabetes. The aim of this study was to develop a novel triglyceride-based tracer for BAT. For this purpose we designed a dual-modal fluorescent/PET fatty acid tracer based on commercially available BODIPY-FL- C_{16} , which can be esterified to its correspondent triglyceride, radiolabeled and incorporated into pre-synthesized chylomicron-like particles.

Methods: BODIPY-FL- C_{16} was coupled to 1,2-diolein with a subsequent radiolabeling step resulting in [^{18}F]BODIPY- C_{16} -triglyceride that was incorporated into chylomicron-like particles. Various quality control steps using fluorescent and radioactive methods were conducted before BAT visualization was tested in mice.

Results: Triglyceride synthesis, radiolabeling and subsequent incorporation into chylomicron-like particles was carried out in decent yields. This radiotracer appeared able to visualize BAT *in vivo*, and the uptake of the radiotracer was stimulated by cold exposure.

Conclusion: The here reported method can be used to incorporate radiolabeled triglycerides into pre-synthesized chylomicron-like particles. Our approach is feasible to visualize and quantify the uptake of triglyceride-derived fatty acids by BAT.

Introduction

Brown adipose tissue (BAT) research has evolved from an underestimated to a fast developing field in endocrine research and non-invasive imaging is an important technique to visualize and quantify BATs metabolic activity. Brown adipocytes have the ability to combust energy to heat by nonshivering thermogenesis in their mitochondria by virtue of the presence of uncoupling protein 1 (UCP1) ¹. It has been shown that BAT activation can be triggered by cold exposure, which induces a release of noradrenalin from nerve endings ². This neurotransmitter binds to adrenergic receptors on the BAT membrane and promotes an induction of intracellular lipolysis which leads to a release of fatty acids (FAs) from triglyceride (TG)-filled lipid droplets ³ (Fig 1).

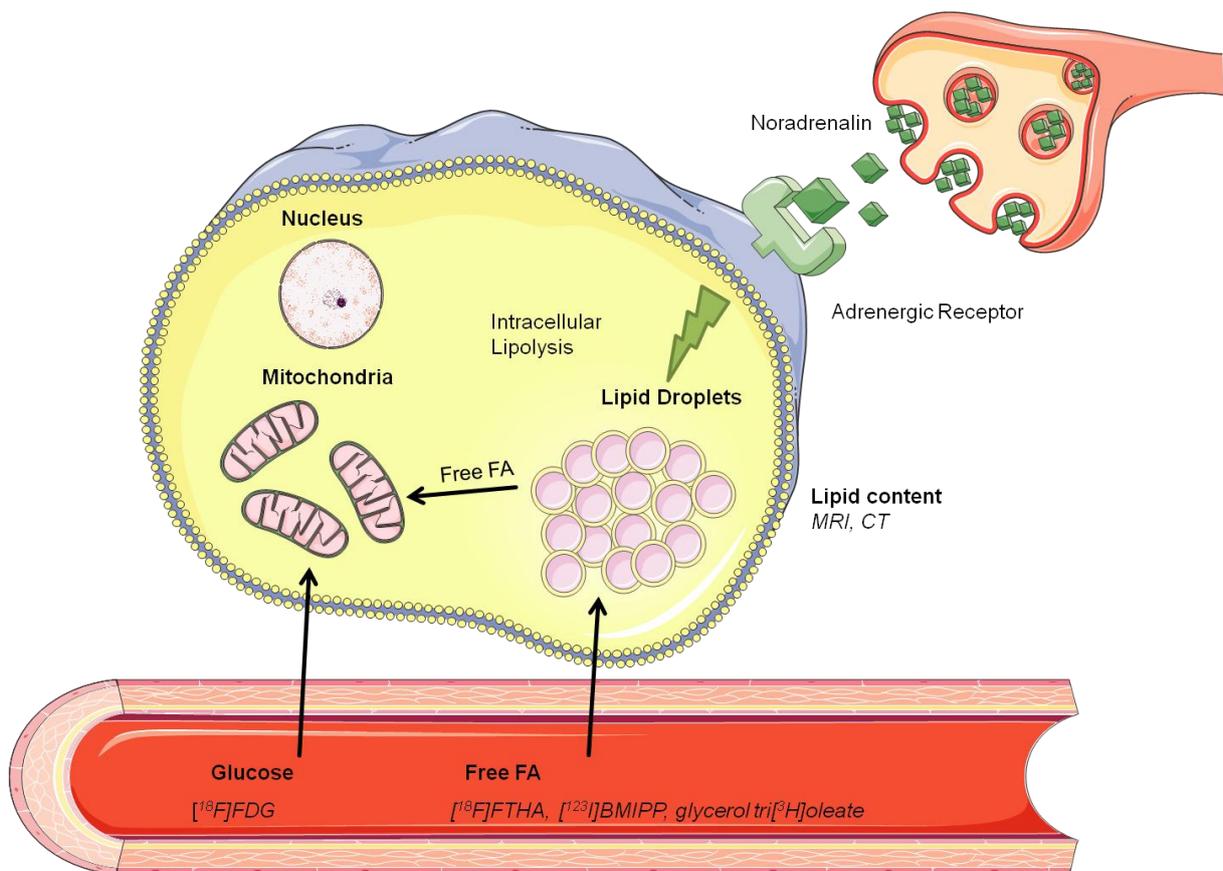
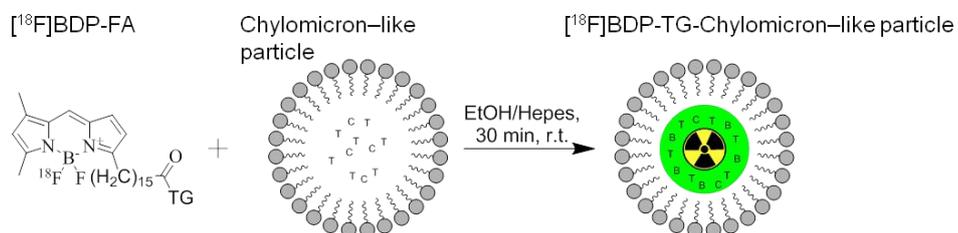


Fig 1: BAT activation by noradrenergic stimulation and possible quantification approaches by $[^{18}\text{F}]\text{FDG}$ (glucose consumption), $[^{18}\text{F}]\text{FTHA}$, $[^{123}\text{I}]\text{BMIPP}$, glycerol tri ^3H oleate-derived ^3H oleate (lipid uptake) and MRI, CT (lipid content).

Those FAs activate UCP1 within the inner mitochondrial membrane ² whereby the proton gradient across the membrane is dispersed and heat is produced.

In activated BAT, internal lipid droplets are replenished by nutrient uptake from plasma in three different ways: uptake of TG-rich lipoprotein (TRL)-derived FAs, glucose uptake followed by *de novo* lipogenesis, and uptake of circulating albumin-bound FAs ²⁻⁵. Although direct TRL particle uptake with adjacent FA release has been suggested ⁴, more recent findings using glycerol tri[³H]oleate and [¹⁴C]cholesteryl oleate double-labeled TRL-like particles showed an approx. 10-fold higher uptake of FAs compared to cholesteryl esters by BAT ^{5,6}, indicating that the majority of TG-derived FAs is internalized after liberation by lipoprotein lipase (LPL). In fact, TRL-derived FAs were identified as the main supply of TGs in BAT ^{3,7} and TG-derived FA internalization was shown to be dependent on the presence of lipoprotein lipase (LPL) ^{7,8}, cluster of differentiation 36 (CD36) ⁹ and fatty acid transport proteins (FATP) ¹⁰ (see also Fig 2 b).

a)



b)

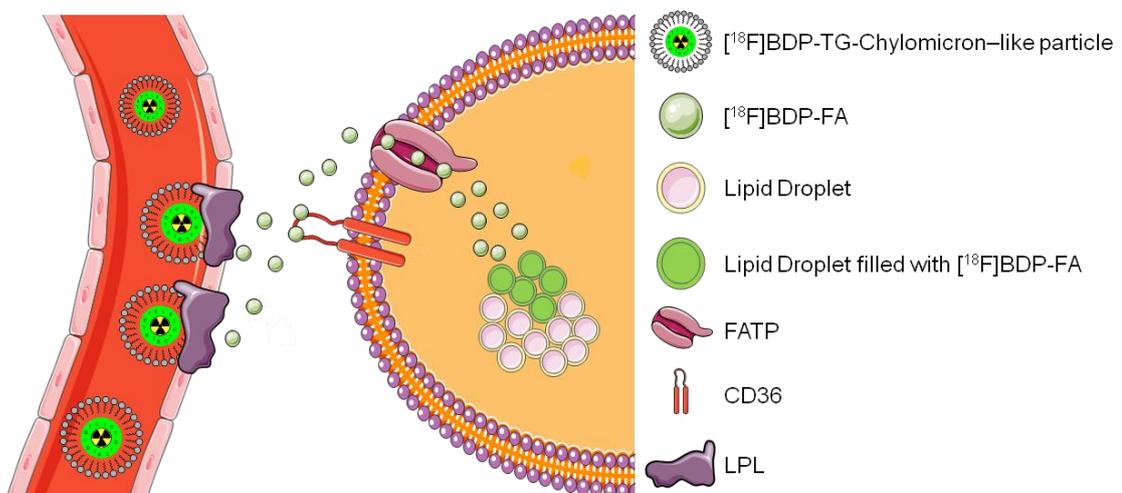


Fig 2: a) Incorporation of radiolabeled [¹⁸F]BDP-TG into pre-synthesized chylomicron-like particles. b) uptake of TG-rich lipoprotein (TRL)-derived FAs from the plasma facilitated by LPL, CD36 and FATP.

The variety of quantification approaches of BAT volume and metabolic activity reaches from invasive imaging with fluorescence probes⁴ or ³H-labeled TG⁵ to non-invasive experiments with PET¹¹⁻¹⁵, SPECT^{16,17} and MRI¹⁸⁻²⁰ and was reviewed recently by us²¹. [¹⁸F]FDG (as a measure for glucose uptake) is frequently used in studies to quantify BAT activity and is currently the gold standard^{12,22-24} (Fig 1). Nevertheless FAs are the main metabolized substance class in BAT, which is not simply mirrored by [¹⁸F]FDG scans¹⁴. In addition, insulin resistance such as type 2 diabetes of BAT occurring with ageing and weight gain will underestimate BAT activity as assessed with [¹⁸F]FDG scans, whereas FA and oxidative metabolism is not dependent on insulin sensitivity²⁵. It should be mentioned that LPL activity was decreased during insulin resistance in mice²⁶ and insulin was found to be essential for the lipolytic processing of TRLs by BAT²⁷. Therefore BAT visualization with TRLs could become difficult during diabetic conditions, but it might be the most precise way to gain information about BATs lipid turnover and metabolic activity. Taken together, we reasoned a TG-derived FA tracer is needed to visualize and quantify lipid uptake by BAT to better reflect the thermogenic potential of BAT compared to [¹⁸F]FDG.

Radiolabeled FAs in general have been developed in several variations for imaging purposes (e.g. [¹⁸F]FTHA and [¹²⁵I]BMIPP^{28,29}). We here report the development of a FA-tracer based on fluorescent FA BODIPY-FL-C₁₆ (BDP-FA), which is suitable for both PET and fluorescence imaging, and present a method to esterify the FA into a TG and subsequently incorporate it into a chylomicron-like particle. We hypothesized that by this approach the physiological situation is mimicked where TRL-derived TGs get lipolyzed locally before they are taken up by brown adipocytes. Additionally it will be possible to image from whole body to sub-cellular level and *in vitro* experiments can be carried out with the same molecule without any radiation dose. BODIPY dyes have been already used to image BAT⁴ and it is known that FA transport proteins (FATP) have a preference for BODIPY-FL coupled to a long carbon chain (C ≥ 8)³⁰. Furthermore, downstream metabolic reactions in white and brown adipocytes have already been visualized *in vitro*^{31,32}.

To avoid decreasing the FA characteristic properties of BDP-FA by introduction of another chelator molecule, we performed $^{18}\text{F}/^{19}\text{F}$ exchange reactions used to transform fluorescent dyes into dual-modality PET/fluorescent imaging dyes ^{4,33-36}. Neither an increased steric demand, nor lowering of the targeting efficiency of the FA towards proteins responsible for FA uptake is expected as BDP-FA is only modified at the end of the carbon chain and therefore keeps its FA characteristics. We recently described the synthesis and radiolabeling of [^{18}F]BDP-TG and conducted first *in vitro* experiments with primary human adipocytes where [^{18}F]BDP-FA uptake could be modified by different BAT activating and blocking agents ³². The aim of the presented study was to incorporate the [^{18}F]BDP-TG into chylomicron-like particles to obtain one of the first TG-based PET imaging agents and to test the new developed tracer in mice. Different to other used FA-based imaging tracers, the here presented TG will be first lipolyzed on-site before it can be internalized by BAT. This reflects the physiological situation in a better way than other FA-based tracers do and will help to quantify BATs consumption of lipids and its contribution to whole body energy expenditure.

Methods

Commercially available compounds were used without further purification unless otherwise stated. BDP-FA was purchased from Thermo Fischer Scientific (99%) (Netherlands). 1,3-diolein was purchased from Sigma Aldrich ($\geq 99\%$). 1,2-diolein was purchased from Cayman Chemicals (USA) ($\geq 95\%$). DMEM/F-12 was purchased from ThermoFischer (Waltham, MA). All HPLC purifications (1.0 mL/min, solvent A; 0.1% TFA in H₂O, solvent B; CH₃CN, 50 °C) were performed on a Shimadzu UFLC HPLC system equipped with a DGU-20A₅ degasser, a SPD-M20A UV detector, a LC-20AT pump system, a CBM-20A communication BUS module, a CTO-20AC column oven, and a Scan-RAM radio-TLC/HPLC-detector from LabLogic using an Aeris™Widepore column (C4, 3.6 μ m, 4.6 mm \times 250 mm) for the BDP-TG. ESI-MS was performed on an Applied Biosystems SCIEX API 150 EX electrospray ionization quadrupole (ESI-Q) mass spectrometer with the method of McAnoy et al.⁵³. Briefly, 0.1 M aqueous ammonium acetate solution was added to the sample to observe the ammonium salt of the synthesized TG in the MS.

¹H-NMR spectra were carried out on a Bruker Ultrashield *TH 400 plus* at 400 MHz. Tol-d₈ was used as solvent with TMS as internal standard. Chemical shifts are reported in parts per million (ppm) relative to the internal standard.

Gel electrophoresis was used to determine the amount of free fluorine-18 and [¹⁸F]BDP-TG in the solution containing [¹⁸F]BDP-TG incorporated in chylomicron-like particles. Gel electrophoresis was carried out under native running conditions where the sample was mixed (1:2) with native sample buffer and loaded into an any kD TGX gel (20 kBq per lane). For visualization phosphor screens were exposed for 10 h to the gel and analyzed by a Typhoon FLA 7000 phosphor imager (GE Healthcare).

Synthesis of chylomicron-like particles

Synthesis of chylomicron-like particles was performed as reported before^{37,54}. Briefly, emulsion particles were prepared from triolein (70 mg), egg yolk phosphatidylcholine (Lipoid)

(22.7 mg), lysophosphatidylcholine (2.3 mg), cholesteryl oleate (3.0 mg), and cholesterol (2.0 mg). Sonification was performed using a Soniprep 150 (MSE Scientific Instruments, UK) that was equipped with a water bath for temperature (54 °C) maintenance, at 10 µm output. The emulsion was fractionated by density gradient ultracentrifugation steps in a Beckman SW 40 Ti rotor. After centrifugation for 30 min at 17,850 rpm at 20 °C, an emulsion fraction containing chylomicron-like particles was removed from the top of the tube by aspiration. Characterization of chylomicron-like particles was done by DLS and transmission electron microscopy. Chylomicron-like particles were stored at 4 °C and were used within 5 days following preparation.

Dynamic Light Scattering

The particle sizes were measured by photon correlation spectroscopy performed at an angle of 90°; with a setup consisting of an ALV-SP8 goniometer, an ALV-SIPC photomultiplier, a multiple τ digital real-time ALV-7004 correlator, and a solid state laser (Koheras) with a red laser ($\lambda = 633$ nm) as light source. The time resolved signal of two Single Photon Counting Modules was cross-correlated. To prevent multiple scattering highly diluted chylomicron-like particle solutions of 0.1 mg/mL in bi-distilled and filtered H₂O (1.2 µm poly(tetrafluoroethylene) membrane filters) were prepared. Sample cuvettes were immersed in a toluene bath and tempered within an error of $\pm 0.1^\circ\text{C}$. Autocorrelation functions of intensity fluctuations $g_2(q, t)$ are converted by the Siegert relation and give the field autocorrelation function $f(q,t)$:

$$f(q, t) = \int_0^\infty G(D_0) \exp\{-D_0 q^2 \tau\} dD_0 \quad (1)$$

Where τ is the decay time, $G(D_0)$ is the distribution function of D_0 , the diffusion coefficient and q as the scattering vector defined as

$$q = \frac{4\pi n}{\lambda_0} \sin\left(\frac{\theta}{2}\right) \quad (2)$$

with θ being the scattering angle and λ_0 being the wavelength of the laser light in vacuum.

The intensity-weighted decay-time τ distributions obtained from the field autocorrelation function by cumulant analysis were analyzed in respect to multimodality. For each diffusive mode the decay rate $\Gamma = 1/\tau$ was plotted against the squared length of the scattering vector q^2 . The slope gave the Z-average translational diffusion coefficient D_0 and results in the hydrodynamic radius R_h after use of the Stokes Einstein equation:

$$D_0 = \frac{k_B T}{6\pi\eta R_h} \quad (3)$$

with q , k_B , T and η being the scattering vector, the Boltzmann constant, absolute temperature, and solvent viscosity, respectively. A hydrodynamic radius distribution was calculated from the regularized Laplace inversion of correlation functions with CONTIN algorithm.

Transmission electron microscopy

Samples were allowed to adsorb on glow discharged formvar-carbon-coated nickel grids (Maxtaform, 200 mesh, Plano, Wetzlar, Germany) for 3 min. Adhesive drops were removed by filter paper. Negative staining was performed with uranyl acetate (0.5% in H_2O , Science Services GmbH, Munich, Germany) for 1-3 seconds. Excess liquid was removed, samples were air dried and examined using a LEO 906 E transmission electron microscope (Zeiss, Oberkochen, Germany), operated at an acceleration voltage of 60 kV.

Synthesis of BDP-TG

Synthesis was performed as reported before³². Briefly, BDP-FL-C₁₆ (300 μ g, 0.6 μ mol) in acetonitrile was evaporated to complete dryness before the reactant was reconstituted in toluene (100 μ L). To the resulting solution $SOCl_2$ in toluene (100 μ L, 4 vol.-%) was added, incubated for 5 min at 70 °C in a closed vial and evaporated. The product was reconstituted in toluene (50 μ L) containing 1,2-diolein (2 μ L, 2.8 μ mol) and heated to 100 °C for 30 min. After the reaction time, purification by HPLC (1 mL/min, 30% to 15% A in 5 min, 15% to 0% A from 5 to 6 min, 0% A to 20 min) yielded **2** (225 μ g, 75%) as a red solid; $t_R = 12.3$ min. ESI-

MS (+) m/z (%) = 1058 (100) [M - F]⁺, 1095 (82) [M + NH₄]⁺. ¹H NMR (400 MHz, Tol-d₈);
δ (ppm) = 5.46 (m, 4H), 4.26 (m, 2H), 4.06 (m, 2H), 3.13 (m, 1H), 1.75 (s, 3H).

Radiolabeling of BDP-TG

Radiolabeling was performed as reported before³². Briefly, aqueous fluorine-18 solution was loaded on a QMA-cartridge which was preconditioned with 15 mL K₂CO₃ in H₂O and 20 mL H₂O. Fluoride (42 MBq) was eluted with a mixture of 600 μL acetonitrile, 400 μL H₂O and 6 mg Sodium p-toluenesulfonate (Sigma-Aldrich). Fluorine-18 solution was transferred into a drying vessel containing tetra-n-butylammonium bromide (80 μL) as a phase transfer agent. Acetonitrile (3 × 1.0 mL) was added and the solution of fluorine-18 was dried by heating to 100°C with a continuous flow of argon. After reconstitution of Fluorine-18 in anhydrous acetonitrile (100 μL), a solution of BDP-TG in toluene (107 μg, 0.1 μmol in 50 μL) and SnCl₄ (0.2 M in acetonitrile, 100 μL) was added to the solution with the activity and the reaction solution was stirred at room temperature (r.t.) for 30 min. [¹⁸F]BDP-TG was obtained (decay corrected RCY: 44%, 25 MBq) with a decay corrected specific activity of 250 MBq/μmol and a radiochemical purity of 45% determined by a radio-TLC with toluene, CHCl₃ and MeOH (80.9%, 14.3%, 4.8%) of the reaction solution.

Ex vivo incorporation of [¹⁸F]BDP-TG into chylomicron-like particles

To [¹⁸F]BDP-TG (233 MBq) in the reaction solution 500 μL H₂O were added and centrifuged for 5 min. The aqueous phase was aspirated and another 500 μL H₂O were added to precipitate the remaining SnCl₄. The mixture was heated to 100 °C, the organic phase was evaporated and the aqueous phase was taken off. [¹⁸F]BDP-TG was reconditioned in 20 μL EtOH and another radio – TLC was performed. [¹⁸F]BDP-TG could be obtained with a radiochemical purity of > 96% and an overall decay corrected radiochemical yield of 21%. 400 μL chylomicron-like particles in HEPES were added (1.5 mg TG content) and incubated for 1 h at r.t.. [¹⁸F]BDP-TG-chylomicron-like particles were obtained (overall decay corrected RCY: 18%, 19 MBq) with a radiochemical purity of > 99% analyzed by gel electrophoresis

and radio-TLC (Fig 3 e and f). Shorter incubation time points and different temperatures were tested by radio-TLC (Fig 3 b and c)

Fluorescence measurements of BDP-TG-chylomicron-like particles

BDP-TG-chylomicron-like particles (80 μ L) (0.1 μ mol BDP-TG dissolved in 20 μ L EtOH + chylomicron-like particles (750 μ g TG content in 200 μ L HEPES), BDP-TG (80 μ L) (0.1 μ mol dissolved in 20 μ L EtOH + 200 μ L HEPES) and chylomicron-like particles (750 μ g TG content in 200 μ L HEPES) are measured using a SpectraMax M2 plate reader (molecular devices) (excitation 485 nm, emission 520 nm).

Animal experiments

Experimental protocols were approved by the “Centrale Commissie Dierproeven” and all animal experiments and procedures were performed in accordance with the guidelines set of this institution. From 13:00 p.m. on, female C57Bl/6 mice (fasted for 4 h at r.t. or fasted and cold exposed at 4 °C for 4 h) were anesthetized (Isoflurane, 1.5-2.0% at 2 mL/min in oxygen), mice kept at r.t. were put on a heating pad and both groups were injected with [¹⁸F]BDP-TG-chylomicron-like particles (1-5 MBq) in HEPES (100 μ L) via the tail vein. Mice were scanned dynamically for 1 h on a microPET (Focus 120, Siemens). Images were analyzed using Pmod V3.707. After the scanning time animals were killed and organs harvested, weighed wet and counted using a WIZARD² automatic γ -counter from Perkin Elmer.

Statistical analyses

Data are presented as mean \pm SD, unless indicated otherwise. Differences at a probability level (p) of 0.05 were considered statistically significant. GraphPad Prism 5.01 (La Jolla, CA, USA) for Windows was used for statistical analyses.

Availability of materials and data

All data generated or analysed during this study are included in this published article (and its Supplementary Information files).

Results

Synthesis of BDP-TG

Synthesis of BDP-TG was conducted as described before³². The resulting TG was obtained with a yield of $45 \pm 8\%$ SD after HPLC purification. BDP-TG has a t_r of 12.3 min and NMR and ESI-MS confirmed the identity of BDP-TG.

Radiolabeling of BDP-TG

The radiolabeled dual-modality imaging agent [¹⁸F]BDP-TG was synthesized in a two-step procedure and was obtained with a decay corrected specific activity of 250 MBq/ μ mol and a decay corrected radiochemical yield of 44%. After washing with H₂O [¹⁸F]BDP-TG was obtained with an overall decay corrected radiochemical yield of 21% and a radiochemical purity >96%. Shelf life and plasma stability showed 99% intact [¹⁸F]BDP-FA after 4 h³². Because of the insolubility of [¹⁸F]BDP-TG in aqueous medium shelf life and plasma stability experiments needed to be performed with [¹⁸F]BDP-FA, however no difference in stability is expected.

Synthesis of chylomicron-like particles and characterization

Chylomicron-like particles were synthesized essentially as reported before^{5,6,37}. After synthesis size of the particles was determined by DLS, showing a mean diameter of 164 ± 20 nm and a polydispersity index of 0.181 (n=4). Additionally, particles were analyzed by transmission electron microscopy. Polydisperse particles could be visualized, all of which showed an encapsulated lipid core and a distinct surface shell with a mean particle diameter of 156 ± 55 nm (n=25) (Fig S1).

***Ex vivo* incorporation of [¹⁸F]BDP-TG into chylomicron-like particles and characterization**

After synthesis the chylomicron-like particles were loaded with BDP-TG or [¹⁸F]BDP-TG. Interestingly, once the BDP-TG is encapsulated, particles show a dark band within their lipid core (compare Fig 3 a with Fig S1). Incorporation of [¹⁸F]BDP-TG into chylomicron-like particles in time was analyzed by TLC. After 25 min more than 99% of the TG was incorporated (Fig 3 b). Different temperatures did not affect the incorporation speed, where r.t. showed the best result after 60 min (99.5%) compared to 0 °C (98.9%) and 38 °C (99.1%) (Fig 3 c). To test whether the [¹⁸F]BDP-TG was truly incorporated, chylomicron-like particles loaded with BDP-TG were measured on a fluorescence microplate reader. Excitation was compared to chylomicron-like particles or BDP-TG alone. A significant increase in intensity (>1,400 fold) was observed for the chylomicron-like particles loaded with BDP-TG in comparison to the particles or the BDP-TG alone (Fig 3 d). Additionally, chylomicron-like particles incubated with [¹⁸F]BDP-TG did not show any impurities of free fluorine-18 or free [¹⁸F]BDP-TG after 60 min of incubation, analyzed by gel electrophoresis and radio-TLC (Fig 3 e and f).

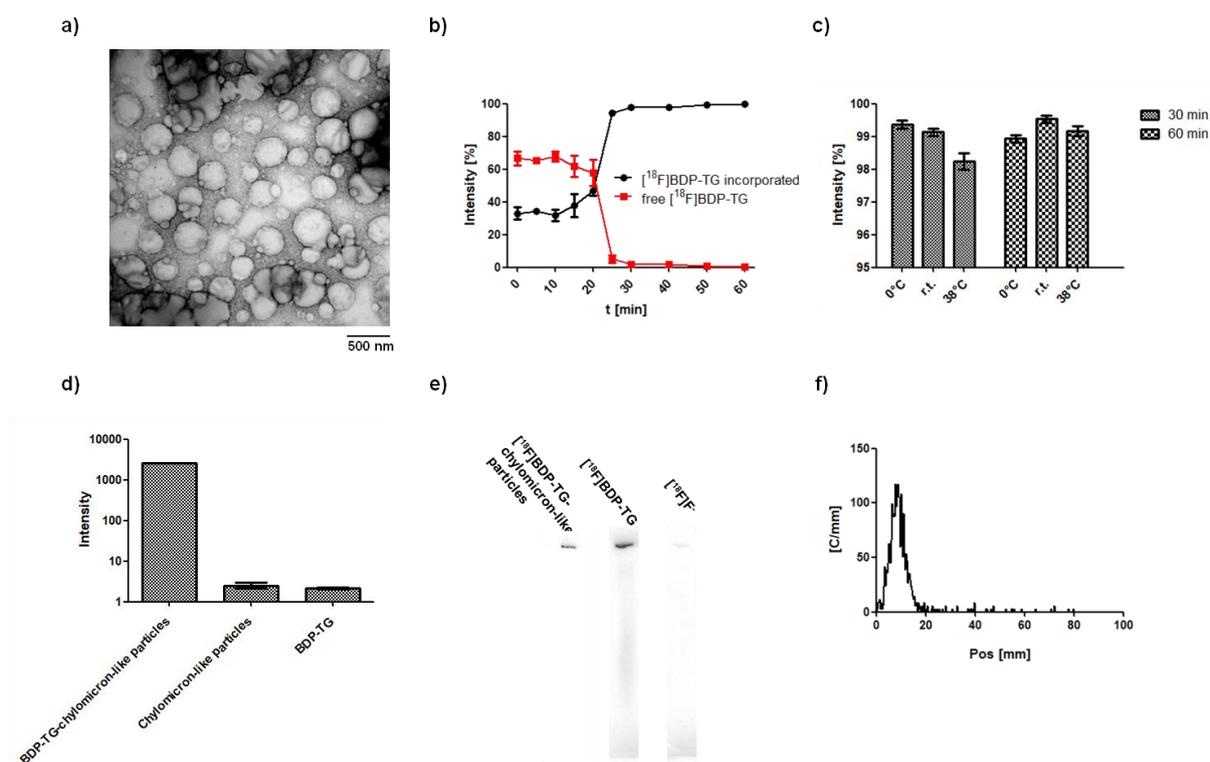


Fig 3: a) Transmission electron microscopy of chylomicron-like particles incubated with $[^{18}\text{F}]$ BDP-TG. b) Incorporation of $[^{18}\text{F}]$ BDP-TG into chylomicron-like particles. c) Temperature dependence of the incorporation of $[^{18}\text{F}]$ BDP-TG into chylomicron-like particles. d) Fluorescence measurement by a microplate reader of BDP-TG-chylomicron-like particles, chylomicron-like particles and BDP-TG. e) Phosphorimaging after gel electrophoresis of chylomicron-like particles labeled with $[^{18}\text{F}]$ BDP-TG (starting position), $[^{18}\text{F}]$ BDP-TG (mainly at starting position) and free fluorine-18 (end position). f) TLC of $[^{18}\text{F}]$ BDP-TG-chylomicron-like particles (Pos=10 mm); possible impurities: free fluorine-18 (Pos=10 mm), free $[^{18}\text{F}]$ BDP-TG (Pos=70 mm).

Animal experiments

$[^{18}\text{F}]$ BDP-TG-chylomicron-like particles (1-5 MBq) were injected into female C57Bl/6 mice, which were fasted for 4 h either at r.t. or at 4 °C. After scanning for 1 h the animals were euthanized and the organs were harvested. Analysis of the PET images showed highest uptake in liver and heart at r.t. and at 4 °C (Fig 4 a and b). A rapid increase with a slow washout in both organs could be visualized (Fig S2 a and b). In bone a constant increase in signal was observed (Fig S2 c and d), which probably indicates a defluorination process of the tracer *in vivo*, as reported in literature³⁶. Lung showed a fast increase with a fast washout and stayed constant at later time points under both temperature conditions. Brain as

a negative control showed negligible uptake. BAT uptake increased significantly when comparing cold exposed animals vs. control animals at 50-60 min p.i. (Mann-Whitney U, n=10, two tailed $p < 0.0001$) (Fig S2 c and d).

PET images are supported by the results of the biodistribution. Highest uptake values in animals kept at r.t. were found in heart ($17.9 \pm 3.3\%$ ID/g), liver ($16.8 \pm 2.6\%$ ID/g), bone ($9.0 \pm 3.0\%$ ID/g) and spleen ($8.5 \pm 2.2\%$ ID/g). After 1 h only $4.5 \pm 1.1\%$ ID/g were found in the blood, indicating a fast blood clearance during the scanning time. In total $6.9 \pm 1.3\%$ ID was found in the blood calculated by an assumed total blood volume of $5.85 \text{ mL}/100\text{g}$ ³⁸. Uptake by BAT ($3.5 \pm 0.7\%$ ID/g) was approximately 3-fold higher than uptake by WAT ($1.1 \pm 0.6\%$ ID/g; Mann-Whitney U, n=5, two tailed $p < 0.001$).

In fasted animals exposed to $4 \text{ }^\circ\text{C}$ highest uptake was reached in heart ($33.7 \pm 9.5\%$ ID/g), liver ($32.0 \pm 5.5\%$ ID/g) and spleen ($14.7 \pm 7.9\%$ ID/g). Uptake in BAT was found to be $6.9 \pm 1.1\%$ ID/g and was significantly higher (Mann-Whitney U, n=5, two tailed, $p < 0.001$) compared to WAT ($1.5 \pm 0.2\%$ ID/g). Total activity remaining in blood was calculated to be $4.9 \pm 0.8\%$ ID and was therefore significantly lower (Mann-Whitney U, n=10, two tailed, $p < 0.01$) compared to animals fasted at r.t. Organs which showed a significant difference between r.t. and cold exposed fasting were heart (Mann-Whitney U, n=10, two tailed, $p < 0.001$), liver (Mann-Whitney U, n=10, two tailed, $p < 0.001$), spleen (Mann-Whitney U, n=10, two tailed, $p < 0.01$), BAT (Mann-Whitney U, n=10, two tailed, $p < 0.001$) and perivascular adipose tissue (PVAT) (Mann-Whitney U, n=10, two tailed, $p < 0.001$). No difference could be found in WAT, bone, muscle and all other analysed organs (Fig 4 c and d).

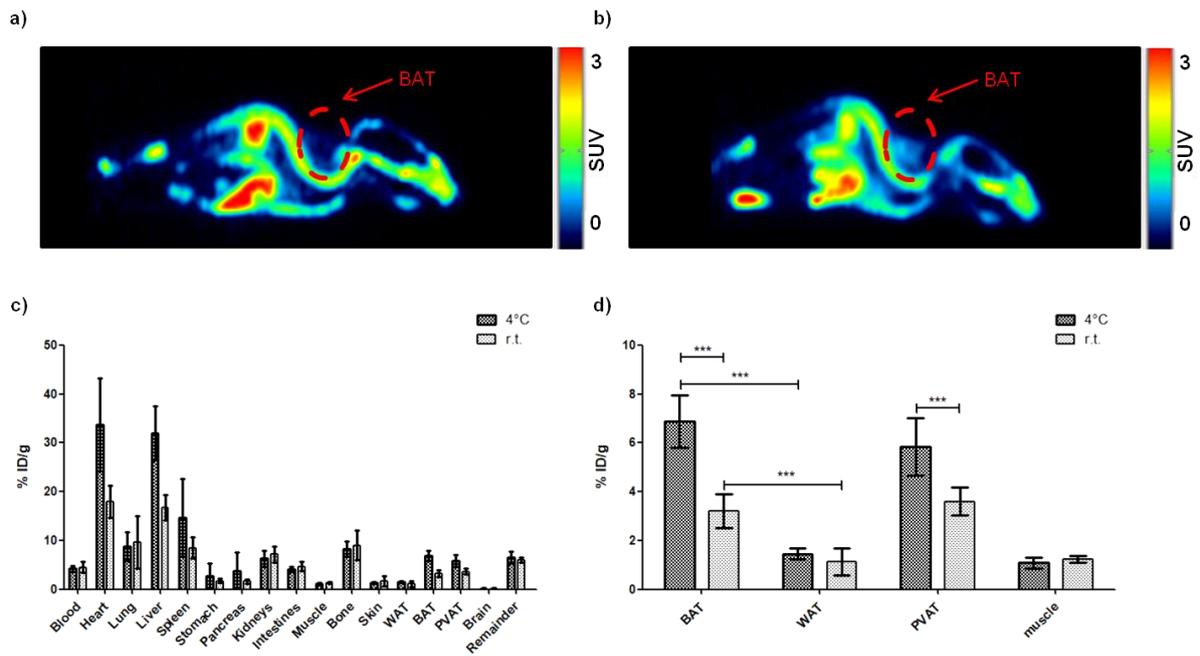


Fig 4: a) PET image (40-60 min) of $[^{18}\text{F}]$ BDP-TG-chylomicron-like particles in a r.t. fasted animal. b) PET image (40-60 min) of $[^{18}\text{F}]$ BDP-TG-chylomicron-like particles in a 4°C fasted animal. c) Biodistribution of $[^{18}\text{F}]$ BDP-TG-chylomicron-like particles 1 h after injection. d) BAT uptake in comparison to WAT, Aorta + PVAT and muscle.

Discussion

Exploring BAT and its metabolism has become an interesting and fast developing topic in endocrine research. A variety of different imaging approaches have been used in the past reaching from *in vitro* experiments³⁹ over invasive imaging with fluorescence probes⁴ or ³H-compounds⁵ to non-invasive experiments with PET¹¹⁻¹⁵, SPECT^{16,17} and MRI¹⁸⁻²⁰. In a clinical environment most often [¹⁸F]FDG scans are used for BAT imaging but it only shows glucose-related uptake and has therefore the chance to misinterpret BAT activity by underestimating lipid uptake and metabolism. Additionally, [¹⁸F]FDG uptake is dependent on insulin sensitivity and therefore might not reflect the real activation state of BAT. In studies with [¹⁸F]FTHA it was observed that radiolabeled FAs showed an increased uptake in BAT under cold stimulation in humans¹⁴. However, these results with a free FA-based tracer might be less relevant because the majority of FAs is TRL derived^{3,7} where FAs are transported as TGs. With our developed tracer [¹⁸F]BDP-TG which is incorporated into chylomicron-like particles we can overcome these limitations and gain new insights in BATs lipid metabolism.

BDP-TG was produced in a decent yield (45%) like previously published³² and radiolabeling was carried out with a decay corrected radiochemical yield of 44% which is in accordance to radiolabeling yields reported in literature^{35,40}.

Chylomicron-like particles have been synthesized with a mean diameter of 164 nm (DLS) and 156 nm (TEM). Those sizes are in accordance with previously described particles^{5,6,37}.

After synthesis of the particles, loading with either BDP-TG or [¹⁸F]BDP-TG was performed. A pre-purification of [¹⁸F]BDP-TG from free fluorine-18 and SnCl₄ before incubation with chylomicron-like particles is of immense importance. In a previous approach [¹⁸F]BDP-TG received a single wash with H₂O (500 µL) before it was incubated with the particles. After 60 min the sample was purified by centrifugal filtration with an Amicon Ultra centrifugal filter (10 KDa). This caused two problems: 1) remaining SnCl₄ caused a co-precipitation of the particles 2) filtration destroyed the particles yielding large lipid emulsions. This could be

overcome by an intensified washing procedure where all the SnCl_4 as well as free fluorine-18 was washed out before [^{18}F]BDP-TG was added to the chylomicron-like particles.

An incorporation of the TG during the formation of the chylomicron-like particles in the sonicator has been tested but resulted in a breakdown of the boron-fluoride bond. Incorporation speed of [^{18}F]BDP-TG into chylomicron-like particles was found to be fast (>99% after 60 min (Fig 3 b)) and temperature seemed to have no effect on the incorporation yield and speed (Fig 3 c). Purity of radiolabeled chylomicron-like particles was found to be >96% and therefore suitable for *in vivo* applications. (Fig 3 e and f).

Additionally, we evaluated the incorporation of the BDP-TG into chylomicron-like particles by fluorescence. We previously found that the fluorescence intensity is strongly related to the environment of the compound, meaning that only if the BDP-TG is dissolved it will give a fluorescence signal ³². BDP-TG incorporated into chylomicron-like particles in HEPES solution showed a huge increase in signal (1413 fold) compared to BDP-TG or chylomicron-like particles alone. This indicates that once the BDP-TG is incorporated into a chylomicron-like particle it regains its fluorescence because it is in a lipophilic environment.

In vivo preferential uptake of [^{18}F]BDP-TG in BAT compared to WAT was observed. Exposure to cold during fasting, thereby activating BAT ^{2,24}, pronounced the difference, indicating the ability of our tracer to visualize BATs metabolic activity and FA consumption. Although not visible on the microPET images, biodistribution data showed that PVAT followed a similar trend. Muscle tissue, which may become activated in the cold due to shivering ⁴¹ showed only low uptake of [^{18}F]BDP-TG and no increased uptake due to cold-exposure. This does not exclude a higher metabolic activity of muscle, as muscle may preferentially use glucose under conditions of cold exposure ⁴².

BAT was however not the only tissue with a high uptake of [^{18}F]BDP-TG. Uptake of the tracer was in fact highest in liver, heart, spleen and bone, and a significant increase by cold exposure is also demonstrated in liver, heart and spleen. Hepatocytes have a low LPL expression ⁶. High uptake values can be explained by increased uptake of remnants of the chylomicron-like particles, which still contain TGs ⁴³, or by spill over of FAs generated during

lipolysis⁴⁴. This might also explain the elevated uptake due to cold activation because more remnants and FA are produced in this situation. Since the heart has high LPL expression^{45,46} a high tracer accumulation was expected. Increased LPL activity due to cold stimulation was already reported in cardiomyocytes^{47,48} which explains the increased heart uptake. The spleen, as an organ of the mononuclear phagocytic system, contains high numbers of macrophages. Those macrophages are able to engulf large particles which might explain the marked uptake observed in the spleen⁴⁹. Lung uptake might be the result of a polydisperse particle distribution, as large particles are prone to get entrapped in lungs capillaries⁵⁰. Indeed, in previous experiments with filtrated particles, where particle coagulation was frequent and sizes >1000 nm diameter could easily be reached, we experienced very high uptake values of >200% ID/g in the lung.

We also noted a high uptake of radioactivity in the bone. Although *ex vivo* plasma stability tests with [¹⁸F]BDP-FA showed >99% intact compound after 4 h³², this may not be valid *in vivo*. We were unable to demonstrate free fluorine-18 *in vivo* in the plasma, due to the rapid blood clearance and the difficulty to perform analyses on radiochemical purity of ¹⁸F-labeled chylomicrons from within a blood sample. Still, in recent literature it was shown that *in vivo* the stability is not ensured for radiolabeled BODIPY-compounds, making it likely that our compound is defluorinated as well³⁶. In a former publication 1,3-diolein was coupled to BDP-FA³². We speculated that FAs on position 1 or 3 on the glycerol backbone might have a higher chance to get released during lipolysis. No significant differences in chemical and radiochemical yields during synthesis nor changes in the biodistribution can be reported for 1,3-diolein-BDP-FA vs. 1,2-diolein-BDP-FA (data not shown).

In comparison to our data, similar particles loaded with glycerol tri[³H]oleate previously showed higher uptake of [³H]oleate by BAT (approx. 5 fold⁶, 6 fold⁵ and 10 fold⁵¹) and lower [³H]oleate uptake by liver (approx. 0.6 fold⁶, 0.6 fold⁵ and 0.5 fold⁵¹) and heart (approx. 0.3 fold⁶, 0.3 fold⁵ and 0.5 fold⁵¹). Also no other organs showed an increased uptake when BAT was stimulated. Different experimental conditions (e.g. use of anesthetized vs. non-anesthetized mice) could be a reason for this difference in the biodistributions as gaseous

anesthetics such as isoflurane are known to suppress adrenergic signaling ⁵². In general these results may point to a reduced BAT LPL-activity due to anesthesia which results in increased uptake by liver and heart.

Conclusion

In the current manuscript, we presented a dual-modal fluorescent and PET active TG which was successfully incorporated into chylomicron-like particles. With different quality control methods we showed incorporation of the radiolabeled TG into chylomicron-like particles. *In vivo* animal studies showed that the resulting tracer was able to reach BAT but was also taken up by other tissues which employ LPL-mediated FA uptake. BAT uptake of the tracer was increased in cold exposed animals. The here presented technique is able to visualize TRL-derived FA BAT uptake after TG-lipolysis which is advantageous in comparison to conventional FA-based tracers which do not reflect the physiological situation and are mainly taken up by the liver. We anticipate that [¹⁸F]BDP-TG-chylomicron-like particles are a promising step forward to visualize and quantify BATs lipid metabolism and gain more information about BATs contribution to whole body energy expenditure in the future.

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Supplementary Information

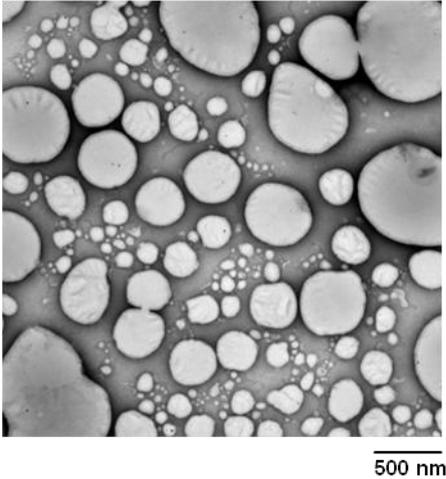


Fig S1: Representative picture of transmission electron microscopy of chylomicron-like particles without BDP-TG loading.

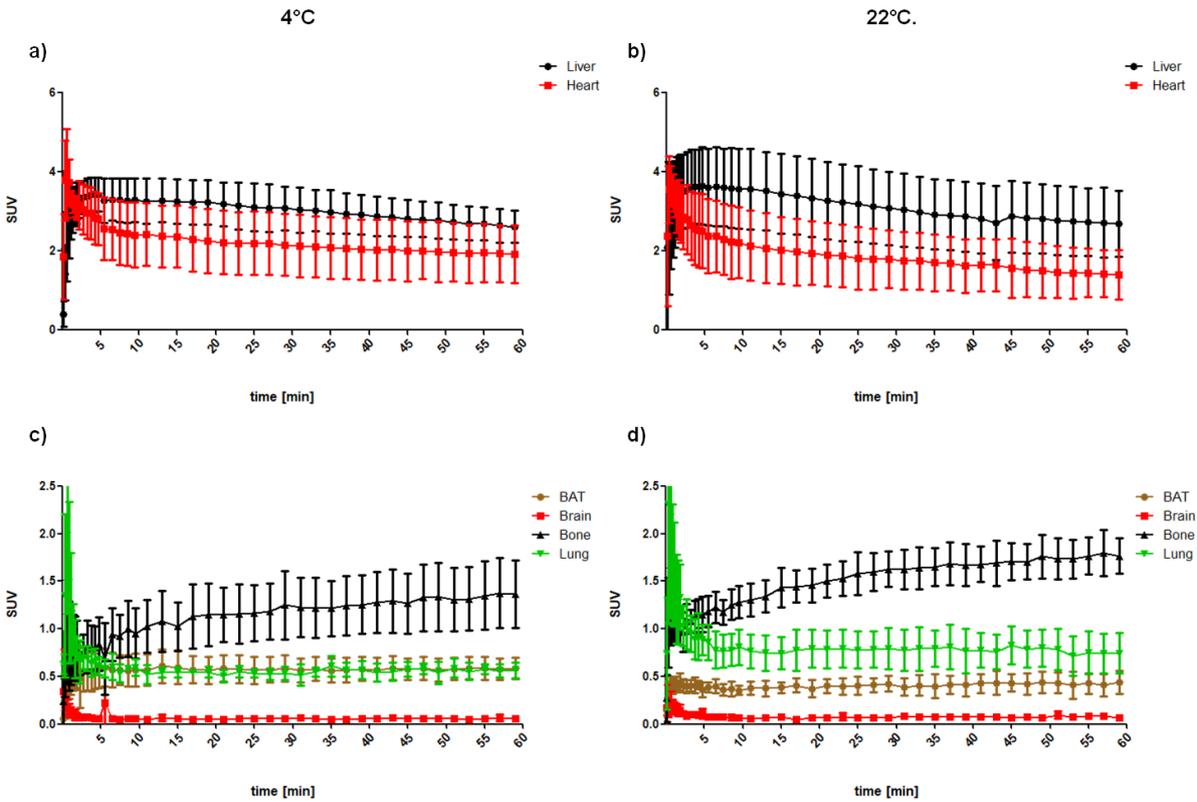


Fig S2: a) Time activity curves for specified organs in 4 °C fasted animals b) Time activity curves for specified organs in 22 °C fasted animals

**Chapter 6:
Triglyceride-rich lipoprotein derived fatty acid uptake by
brown adipose tissue in control and diabetic mice at room
temperature, acute cold exposure and cold acclimation**

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Brown adipose tissue uptake of triglyceride-rich lipoprotein derived fatty acids in diabetic or obese mice under different temperature conditions. *Molecular Imaging & Biology*, (2019)

Abstract

Background: Brown adipose tissue (BAT) is present in adult humans and contributes to their overall energy expenditure. The most often used tracer to visualize BAT and its metabolism is [¹⁸F]FDG which suffers from the problem that it is not able to visualize BATs complete metabolic activity. Under insulin resistant conditions associated with ageing and weight gain this method fails to detect BAT. We here used our novel developed triglyceride-based tracer to visualize BATs metabolic activity under different temperature conditions and in a mouse model with induced diabetes.

Methods: Control and streptozotocin-induced diabetic mice, housed at 21 °C, acutely exposed to cold or cold acclimated over 28 days, were injected with [¹⁸F]BDP-TG-chylomicron-like particles and scanned dynamically for 32.5 min.

Results: [¹⁸F]BDP-TG-chylomicron-like particles were able to visualize BAT under all conditions. Increased uptake was found in control mice acutely exposed to cold but not in other conditions. Diabetes reduced BAT uptake to levels of control mice housed at 21 °C.

Conclusion: BAT was visualized under conditions were commonly used techniques (e.g. [¹⁸F]FDG) fail. BAT activation by cold exposure under diabetic conditions could not be detected indicating an insulin dependent uptake mechanism.

Introduction

Brown adipose tissue (BAT) research has evolved rapidly within the last 2 decades to an important field in endocrine research. To visualize BAT and its metabolism, non-invasive imaging is a central technique. BAT has the ability to uncouple its ATP production and to produce heat instead of ATP. During this process protons enter the mitochondrial matrix by uncoupling protein 1 (UCP1)¹, a BAT specific protein, and release their energy as heat^{2,3}. This process is considered as nonshivering thermogenesis^{2,4}.

BAT activity is triggered by cold stimulation and the chance to visualize BAT in animals and lean humans raises⁵⁻⁷. During cold exposure, thermoreceptors in the skin are stimulated and activate neurons in the hypothalamus, resulting in a release of norepinephrine^{2,8}. Norepinephrine binding to β 3-adrenoceptors on BAT is activating a signalling cascade, which will lead to lipolysis of intracellular triglycerides (TG) and at the end result in UCP1 activation^{9,10}. Therefore BAT might be an interesting target in the fight against obesity as it “burns” lipids instead of storing them.

Most BAT scans were performed with [¹⁸F]FDG because of BATs history. For long, symmetrical accumulations in the supraclavicular area during [¹⁸F]FDG positron emission tomography (PET) (e.g. during cancer scans) were attributed to cervical muscle until later scans using PET/computed tomography (CT) could show that Hounsfield units are similar to them of adipose tissue¹¹⁻¹³. Therefore a high number of retrospective BAT studies have been published and until now [¹⁸F]FDG is used to visualize BAT. The prevalence to find active BAT depots during [¹⁸F]FDG/PET scans is strongly dependent on different variables such as age, BMI, or outdoor temperature^{14,15}.

BAT [¹⁸F]FDG/PET scans suffer from two severe problems: 1) Fatty acids (FAs) have been identified as the main metabolized substance class^{16,17} and therefore [¹⁸F]FDG/PET might largely underestimate BATs metabolic activity. 2) In clinical studies [¹⁸F]FDG BAT uptake was impaired in diabetic patients but uptake of [¹⁸F]FTHA (a radiolabeled FA) was not altered when compared to non-diabetic controls¹⁸. As a high number of obese patients suffer from

diabetes type II ¹⁹ their insulin resistance will additionally decrease [¹⁸F]FDG uptake in BAT and underestimate BATs metabolic activity.

Due to this facts BAT visualization with lipid tracers would be advantageous over [¹⁸F]FDG. FA based tracers, such as [¹⁸F]FTHA ²⁰ or [¹²⁵I]BMIPP ²¹ have been developed and can be found in clinical applications. But even FA tracer might not be the optimal choice.

During BAT activation internal lipid droplets are replenished by nutrient uptake from plasma in three different ways: uptake of FAs from triglyceride (TG)-rich lipoproteins (TRL) after external lipolysis, glucose uptake followed by *de novo* lipogenesis, and uptake of circulating albumin-bound FAs ^{2,22-24}. It was found that TRL-derived FAs are the main supply of TGs in BAT ^{24,25}. Due to this fact, free radiolabeled FAs applied *in vivo* rely on many different uncontrolled uptake and incorporation process before they are taken up by BAT, which we wanted to avoid. Therefore we have developed a radiolabeled TG ²⁶ which we were able to incorporate into a pre-synthesized chylomicron-like particle ²⁷. Additionally the tracer was tested *in vivo* and was able to visualize BAT at room temperature (21 °C) conditions as well as its uptake was increased by acute cold exposure.

In this manuscript we investigate the effect of induced diabetes as well as acute cold and cold-acclimation in a mouse model on BAT metabolism visualized by our recently developed lipid based tracer [¹⁸F]BODIPY ([¹⁸F]BDP)-TG-chylomicron-like particle.

Methods

Commercially available compounds were used without further purification unless otherwise stated. BDP-FA was purchased from Thermo Fischer Scientific (99%) (Netherlands). 1,2-diolein was purchased from Cayman Chemicals (USA) ($\geq 95\%$). All HPLC purifications (1.0 mL/min, solvent A; 0.1% TFA in H₂O, solvent B; CH₃CN, 50°C) were performed on a Shimadzu UFLC HPLC system equipped with a DGU-20A₅ degasser, a SPD-M20A UV detector, a LC-20AT pump system, a CBM-20A communication BUS module, a CTO-20AC column oven, and a Scan-RAM radio-TLC/HPLC-detector from LabLogic using an Aeris™Widepore column (C4, 3.6 μ m, 4.6 mm \times 250 mm) for the BODIPY-triglyceride (BDP-TG). ESI-MS was performed on an Applied Biosystems SCIEX API 150 EX electrospray ionization quadrupole (ESI-Q) mass spectrometer with the method of McAnoy et al.²⁸. Briefly, 0.1 M aqueous ammonium acetate solution was added to the sample to observe the ammonium salt of the synthesized TG in the MS.

¹H-NMR spectra were carried out on a Bruker Ultrashield *TH 400 plus* at 400 MHz. Tol-d₈ was used as solvent with TMS as internal standard. Chemical shifts are reported in parts per million (ppm) relative to the internal standard.

Synthesis of chylomicron-like particles

Synthesis of chylomicron-like particles was performed as reported before^{29,30}. Briefly, emulsion particles were prepared from triolein (70 mg), egg yolk phosphatidylcholine (Lipoid) (22.7 mg), lysophosphatidylcholine (2.3 mg), cholesteryl oleate (3.0 mg), and cholesterol (2.0 mg). Sonification was performed using a Soniprep 150 (MSE Scientific Instruments, UK) that was equipped with a water bath for temperature (54 °C) maintenance, at 10 μ m output. The emulsion was fractionated by density gradient ultracentrifugation steps in a Beckman SW 40 Ti rotor. After centrifugation for 30 min at 17,850 rpm at 20 °C, an emulsion fraction containing chylomicron-like particles was removed from the top of the tube by aspiration. Characterization of chylomicron-like particles was done by DLS and transmission electron

microscopy. Chylomicron-like particles were stored at 4 °C and were used within 5 days following preparation.

Synthesis of BDP-TG

Synthesis was performed as reported before ²⁶. Briefly, BDP-FL-C₁₆ (300 µg, 0.6 µmol) in acetonitrile was evaporated to complete dryness before the reactant was reconstituted in toluene (100 µL). To the resulting solution SOCl₂ in toluene (100 µL, 4 vol.-%) was added, incubated for 5 min at 70 °C in a closed vial and evaporated. The product was reconstituted in toluene (50 µL) containing 1,2-diolein (2 µL, 2.8 µmol) and heated to 100 °C for 30 min. After the reaction time, purification by HPLC (1 mL/min, 30% to 15% A in 5 min, 15% to 0% A from 5 to 6 min, 0% A to 20 min) yielded **2** (225 µg, 75%) as a red solid; *t_R* = 12.3 min. ESI-MS (+) *m/z* (%) = 1058 (100) [M - F]⁺, 1095 (82) [M + NH₄]⁺. ¹H NMR (400 MHz, Tol-d₈); δ (ppm) = 5.46 (m, 4H), 4.26 (m, 2H), 4.06 (m, 2H), 3.13 (m, 1H), 1.75 (s, 3H).

Radiolabeling of BDP-TG

Radiolabeling was performed as reported before ²⁶. Briefly, aqueous fluorine-18 solution was loaded on a QMA-cartridge which was preconditioned with 15 mL K₂CO₃ in H₂O and 20 mL H₂O. Fluoride (42 MBq) was eluted with a mixture of 600 µL acetonitrile, 400 µL H₂O and 6 mg Sodium p-toluenesulfonate (Sigma-Aldrich). Fluorine-18 solution was transferred into a drying vessel containing tetra-n-butylammonium bromide (80 µL) as a phase transfer agent. Acetonitrile (3 × 1.0 mL) was added and the solution of fluorine-18 was dried by heating to 100 °C with a continuous flow of argon. After reconstitution of Fluorine-18 in anhydrous acetonitrile (100 µL), a solution of BDP-TG in toluene (107 µg, 0.1 µmol in 50 µL) and SnCl₄ (0.2 M in acetonitrile, 100 µL) was added to the solution with the activity and the reaction solution was stirred at room temperature (r.t.) for 30 min. [¹⁸F]BDP-TG was obtained (decay corrected RCY: 44%, 25 MBq) with a decay corrected specific activity of 250 MBq/µmol and a radiochemical purity of 45% determined by a radio-TLC with toluene, CHCl₃ and MeOH (80.9%, 14.3%, 4.8%) of the reaction solution.

Ex vivo incorporation of [¹⁸F]BDP-TG into chylomicron-like particles

Incorporation of radiolabeled [¹⁸F]BDP-TG was performed as reported before ²⁷. Briefly, the [¹⁸F]BDP-TG solution (233 MBq) was quenched with 500 μL H₂O and centrifuged for 5 min. The organic phase was washed 3 times with 500 μL H₂O before [¹⁸F]BDP-TG was reconditioned in 20 μL EtOH. [¹⁸F]BDP-TG could be obtained with a radiochemical purity of > 96% and an overall decay corrected radiochemical yield of 21%. 400 μL chylomicron-like particles in HEPES were added (1.5 mg TG content) and incubated for 1 h at r.t.. [¹⁸F]BDP-TG-chylomicron-like particles were obtained (overall decay corrected RCY: 18%, 19 MBq) with a radiochemical purity of > 99% analyzed by gel electrophoresis and radio-TLC.

Animal experiments

Experimental protocols were approved by the “Centrale Commissie Dierproeven” and all animal experiments and procedures were performed in accordance with the guidelines set of this institution. 23 female C57Bl/6 mice were divided into 2 groups. Group 1 served as control group and group 2 was injected with Streptozocin 10 days before the experiment to destroy the β-cells in the pancreas ³¹. Mice from group 1 were fed with a normal diet (10% kcal fat content, Research Diets Inc.), while group 2 received a high fat diet ³² (45% kcal fat content, Research Diets Inc.) from the day of Streptozocin injection. An animal was considered as diabetic when the glucose level at the day of the experiment was > 10 mM.

Both groups were divided into 3 subgroups where subgroup 1 was fasted for 4 h at 21 °C before the imaging experiment. Subgroup 2 was housed at 21 °C but was fasted for 4 h during exposure to 6 °C prior to tracer injection. Subgroup 3 was exposed to cold for 6 h per day for 28 days and was fasted for 4 h before tracer injection (Fig 1).

Mice were anesthetized (Pentobarbital, 60 mg/kg i.p.), and injected with [¹⁸F]BDP-TG-chylomicron-like particles (1-10 MBq) in HEPES (100 μL) via the tail vein. Mice were scanned dynamically for 32.5 min on a microPET (Focus 120, Siemens). Images were analyzed using Pmod V3.707. After the scanning time animals were killed and organs

harvested, weighed wet and counted using a WIZARD² automatic γ -counter from Perkin Elmer.

Additionally mice (n=12) were divided into two groups. Group 1 was injected at day 0 either with [¹⁸F]FDG (n=4) or with [¹⁸F]FTHA (n=4) (10 ± 2 MBq) and scanned dynamically for 20 min. After 14 days of cold acclimation (4 °C for 6 h/day), group 2 was also injected with [¹⁸F]FDG (n=4) or with [¹⁸F]FTHA (n=4) (10 ± 2 MBq) and scanned dynamically for 20 min.

BAT uptake was analyzed by SUV values.

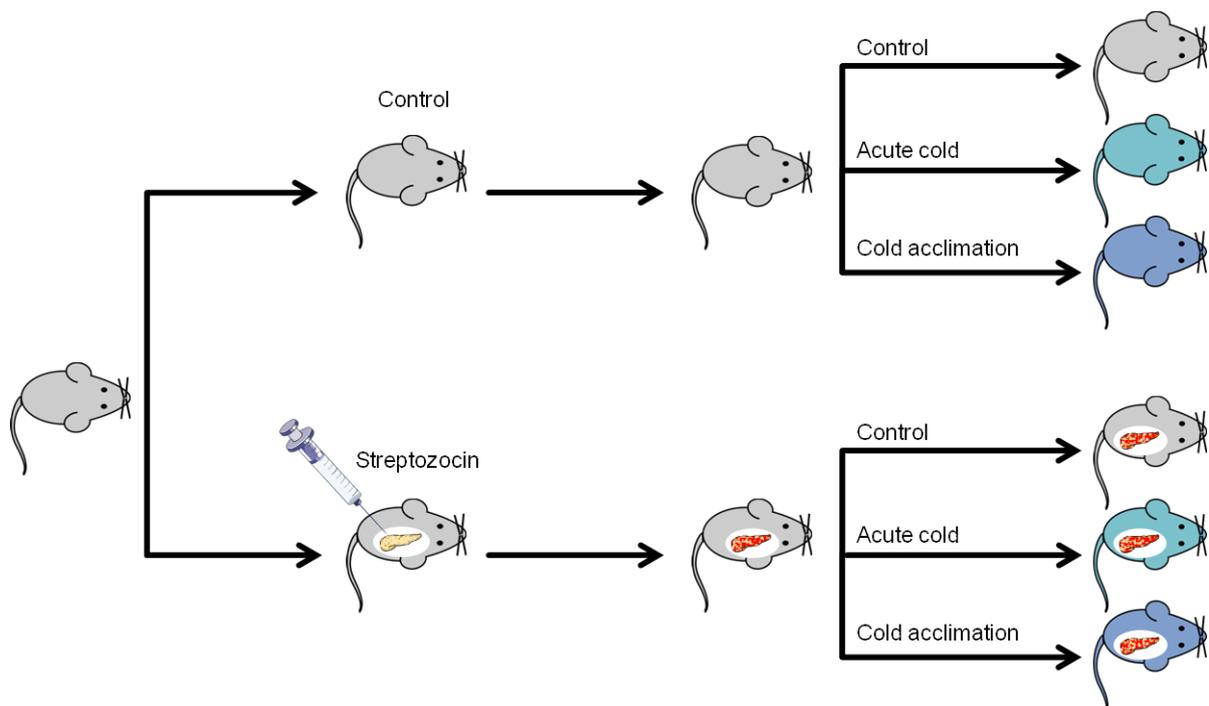


Fig 1: Mice were divided into 2 groups. Group 1 was used as a control group. Group 2 was injected with Streptozocin 10 days before the imaging experiment and afterwards fed with a high fat diet to induce diabetes. Both groups were divided into 3 subgroups where subgroup 1 was housed at 21 °C before the experiment. Subgroup 2 was housed at 21 °C but was exposed to 6 °C for 4 h prior to tracer injection. Subgroup 3 was exposed to cold for 6 h per day for 28 days before the imaging experiment.

Statistical analyses

Data are presented as mean \pm SD, unless indicated otherwise. Differences at a probability level (p) of 0.05 were considered statistically significant. GraphPad Prism 5.01 (La Jolla, CA, USA) for Windows was used for statistical analyses.

Availability of materials and data

All data generated or analysed during this study are included in this published article (and its Supplementary Information files).

Results

Synthesis of BDP-TG, radiolabeling and incorporation into chylomicron-like particles

Esterification of BDP-C₁₆ yielded BDP-TG (45 ± 8%) after HPLC purification (t_r 12.3 min) and its identity was confirmed by NMR and ESI-MS like reported before ²⁶. Radiolabeling of BDP-TG was conducted as described before (44% decay corrected rcy., 250 MBq/μmol corrected specific activity). After washing steps [¹⁸F]BDP-TG could be obtained with an overall rcy. of 21% (rcp. >96%).

Chylomicron-like particles were synthesized with a mean diameter of 164 ± 20 nm and a polydispersity index of 0.181 (n=4). Electron microscopy confirmed the identity of the particles.

[¹⁸F]BDP-TG was incorporated into chylomicron-like particles like reported before ²⁷. After 60 min at r.t. more than 99% of the [¹⁸F]BDP-TG was incorporated in the particle and no free fluorine-18 was found in solution.

Animal experiments

[¹⁸F]BDP-TG-chylomicron-like particles (1-10 MBq) were injected *i.v.* into female C57Bl/6 mice, which were fasted for 4 h either at 21 °C or at 4°C. After scanning for 32.5 min the animals were euthanized and the organs were harvested. Analysis of the PET images showed highest uptake in liver, and heart of mice exposed at 21 °C, acute cold and cold acclimation (Fig 4 and Fig S1-6 a) in both control and diabetic animals. A rapid increase with a slow washout in both organs could be visualized (Fig S1-6 a). In bone a constant increase in signal was observed (Fig S1-6 b), which probably indicates a defluorination process of the tracer *in vivo*, as reported in literature ³³. Lung showed a fast increase with a fast washout and stayed constant at later time points under all temperature conditions. Brain as a negative control showed negligible uptake (Fig S1-6 b).

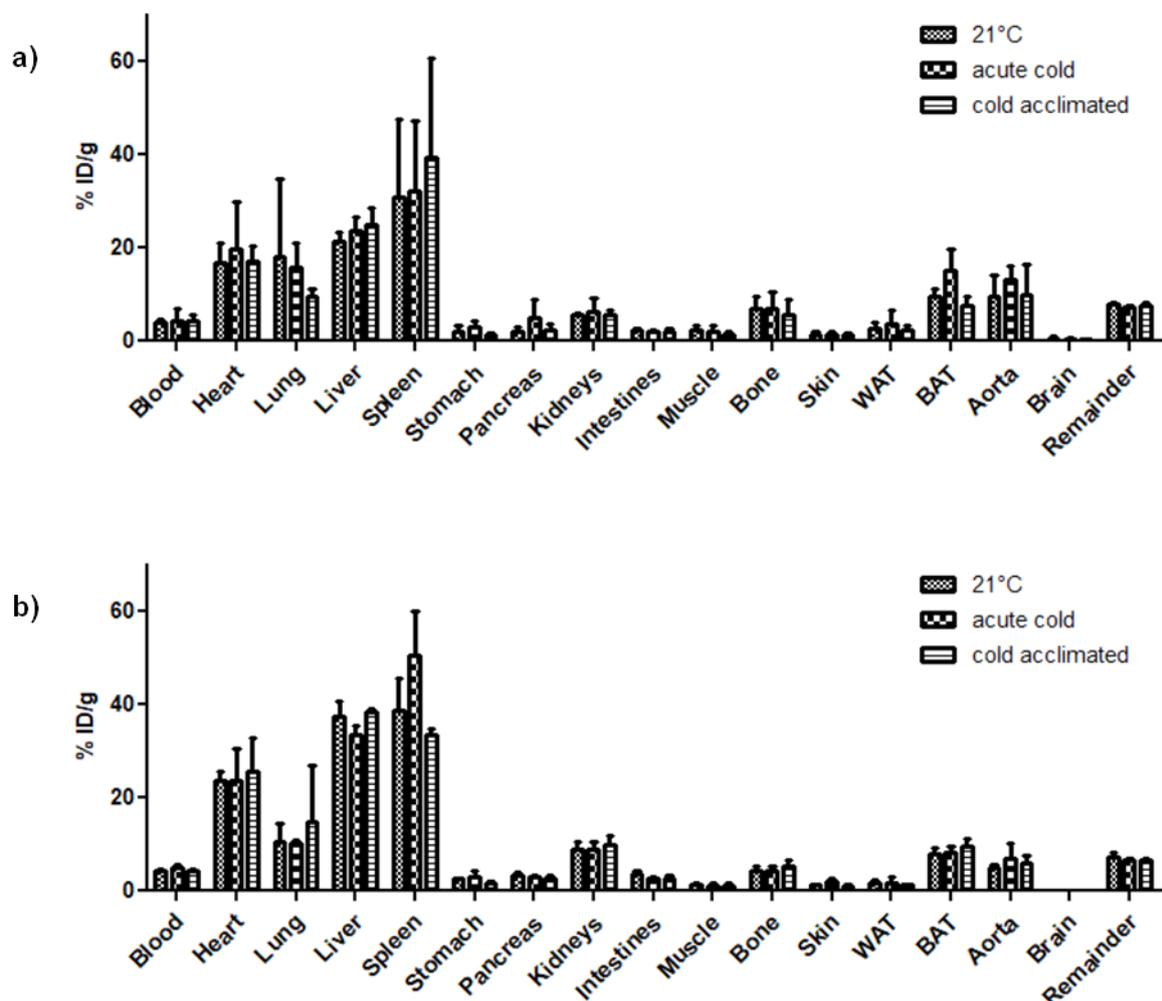


Fig 2: Biodistribution data of control (a) and diabetic (b) mice at 21 °C, under acute cold and cold acclimation 30 min after injection of [18F]BDP-TG-chylomicron-like particles.

BAT uptake was significantly increased under control conditions by acute cold vs. 21 °C (Mann-Whitney U, n=3, two tailed, $p < 0.05$) as well as by acute cold vs. cold acclimation (Mann-Whitney U, n=3, two tailed, $p < 0.05$) (Fig S1-3 b). No other effects due to cold exposure were observed in the control group and in the group with induced diabetes. Under acute cold exposure the control group showed significant increased BAT uptake vs. the diabetic group (Mann-Whitney U, n=6, two tailed, $p < 0.001$) (Fig S2 and 5 b). No other significant group differences can be reported.

PET images are supported by the results of the biodistribution. Highest uptake values in control animals housed at 21 °C were found in spleen ($30.1 \pm 16.9\%$ ID/g), liver ($21.5 \pm 2.0\%$ ID/g), lung ($18.0 \pm 2.0\%$ ID/g), and heart ($16.9 \pm 4.1\%$ ID/g) (Fig 3 a). After 30 min only $4.7 \pm 2.2\%$ ID/g were found in the blood, indicating a fast blood clearance during the scanning time. Uptake by BAT ($9.6 \pm 1.7\%$ ID/g) was approximately 3.5-fold higher than uptake by WAT ($2.8 \pm 1.4\%$ ID/g; Mann-Whitney U, n=4, two tailed $p < 0.001$) (Fig 2 a).

In animals exposed to acute cold highest uptake was reached in spleen ($32.2 \pm 14.8\%$ ID/g), liver ($23.8 \pm 3.9\%$ ID/g) and heart ($19.9 \pm 9.8\%$ ID/g). Uptake in BAT was found to be $15.1 \pm 4.7\%$ ID/g and was significantly higher (Mann-Whitney U, n=4, two tailed, $p < 0.05$) compared to WAT ($3.6 \pm 3.0\%$ ID/g) (Fig 2 a).

In cold acclimated animals highest uptake was reached in spleen ($39.3 \pm 21.3\%$ ID/g), liver ($24.9 \pm 3.6\%$ ID/g) and heart ($17.2 \pm 3.3\%$ ID/g). Uptake in BAT was found to be $7.6 \pm 1.9\%$ ID/g and was significantly higher (Mann-Whitney U, n=3, two tailed, $p < 0.05$) compared to WAT ($2.4 \pm 0.9\%$ ID/g) (Fig 2 a).

BAT uptake in animals exposed to acute cold showed a clear trend towards increased uptake compared to animals housed at 21 °C but could not reach significance (Mann-Whitney U, n=6, two tailed, $p = 0.07$). Acute cold vs. cold acclimation animals showed a significant increase in uptake in animals exposed to acute cold (Mann-Whitney U, n=6, two tailed, $p < 0.05$) where housing at 21 °C vs. cold acclimation did not show significant differences. Other organs did not show any response to acute cold or cold acclimation (2way Anova).

Highest uptake values in diabetic animals housed at 21°C were found in spleen ($38.7 \pm 6.9\%$ ID/g), liver ($37.5 \pm 3.3\%$ ID/g) and heart ($23.7 \pm 1.9\%$ ID/g). $4.4 \pm 0.3\%$ ID/g were found in the blood. Uptake by BAT ($8.0 \pm 1.3\%$ ID/g) was 4.2-fold higher than uptake by WAT ($1.9 \pm 0.5\%$ ID/g; Mann-Whitney U, n=4, two tailed $p < 0.001$) (Fig 2 b).

In diabetic cold exposed animals highest uptake was reached in spleen ($50.3 \pm 9.7\%$ ID/g), liver ($33.4 \pm 2.1\%$ ID/g) and heart ($23.8 \pm 6.6\%$ ID/g). Uptake in BAT was found to be $8.2 \pm$

1.4% ID/g and was significantly higher (Mann-Whitney U, n=4, two tailed, $p < 0.001$) compared to WAT ($1.8 \pm 1.2\%$ ID/g) (Fig 2 b).

In cold acclimated animals highest uptake was reached in liver ($38.4 \pm 0.6\%$ ID/g), spleen ($33.4 \pm 1.2\%$ ID/g) and heart ($25.5 \pm 7.2\%$ ID/g). Uptake in BAT was found to be $9.5 \pm 1.8\%$ ID/g and was significantly higher (Mann-Whitney U, n=3, two tailed, $p < 0.05$) compared to WAT ($1.4 \pm 0.1\%$ ID/g) (Fig 2 b).

All three conditions (22 °C, acute cold exposure and cold acclimation) in diabetic animals were compared to each other but only showed significant differences in spleen in 22 °C vs. acute cold ($p < 0.001$) and acute cold vs. cold acclimated ($p < 0.001$) (2way Anova).

When control and diabetic animals were compared at the same temperature significant difference was found at 22 °C in liver ($p < 0.001$), in acute cold animals in liver ($p < 0.05$), spleen ($p < 0.001$) and BAT ($p < 0.05$) and in cold acclimated animals in liver ($p < 0.01$).

BAT uptake of [^{18}F]BDP-TG was plotted against the glucose level and both groups differed significantly by their glucose levels (Mann-Whitney U, n=23, two tailed, $p < 0.05$) but not by their BAT uptake values (Fig 3).

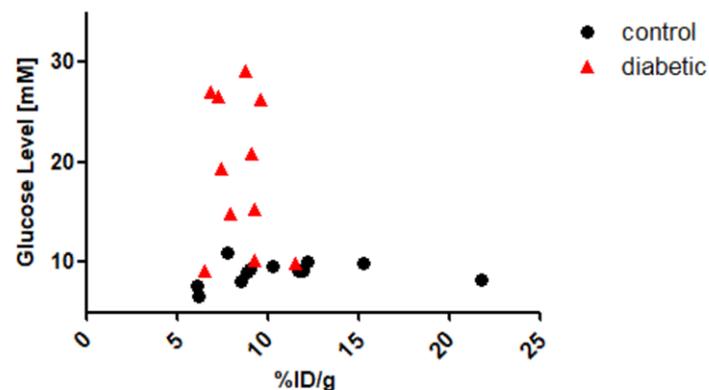


Fig 3: glucose level [mM] and %ID/g BAT uptake of [^{18}F]BDP-TG-chylomicron-like particles in control and diabetes induced mice.

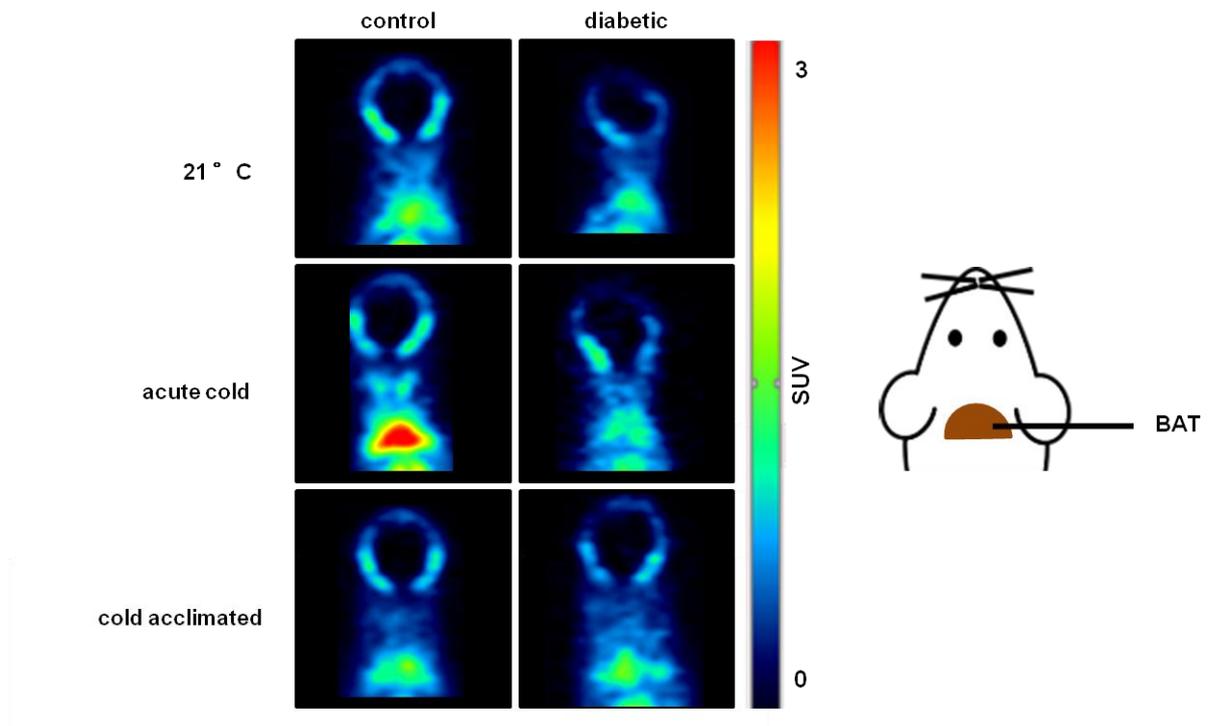


Fig 4: PET images (22-32.5 min) of [^{18}F]BDP-TG- chylomicron-like particles in control and diabetic mice under different temperature conditions.

BAT uptake of [^{18}F]FDG was significant higher compared to [^{18}F]FTHA in animals housed at 21 °C (2.5 ± 1.1 vs. 0.5 ± 0.1) (Mann-Whitney U, $n=8$, two tailed, $p<0.05$). The same difference in uptake can also be reported for animals acclimated to cold for 14 days (3.0 ± 1.0 vs. 0.4 ± 0.1) (Mann-Whitney U, $n=8$, two tailed, $p<0.05$). Animals scanned with the same radiotracer did not show a significant response to 14 days of cold exposure for [^{18}F]FDG (Mann-Whitney U, $n=8$, two tailed, $p>0.05$) and for [^{18}F]FTHA (Mann-Whitney U, $n=8$, two tailed, $p>0.05$).

Discussion

BAT and its metabolism has become an interesting topic in the field of endocrine research within the last two decades. A high amount of imaging and quantification approaches, reaching from *in vitro* experiments³⁴, over invasive²³ or fluorescent probes²² to non-invasive imaging techniques with PET^{11-13,35,36}, SPECT^{37,38} and MRI³⁹⁻⁴¹, has been adapted or especially developed to explore BAT. Till now [¹⁸F]FDG is the most often used technique for BAT imaging but it is suffering from the problem that it is only able to show glucose related uptake. Therefore [¹⁸F]FDG might largely underestimate BATs metabolic activity as FAs are the main metabolized substance class^{16,17}. Another disadvantage of BAT imaging with [¹⁸F]FDG is that it is strongly dependent on insulin sensitivity. In a clinical study it was shown that BATs glucose uptake was strongly impaired during insulin resistance where FA uptake was not affected under these conditions when compared to healthy controls¹⁸. These findings indicate that even during insulin resistance, BATs oxidative metabolism and FA uptake are not altered²⁵. TRL-derived TGs were identified as BATs main source of FAs^{24,25} and lipoprotein lipase (LPL) was shown to be crucial for processing TRLs *in vivo*^{25,42}. As LPL activity is decreased during insulin resistance⁴³, and insulin was found to be essential for the lipolysis of TRLs⁴⁴, uptake of TRL-derived FA is also altered under diabetic conditions which might complicate BAT detection. Nevertheless, to our opinion, radiolabeled TG encapsulated into TRLs might be the most accurate way to visualize BATs FA uptake and metabolism as it reflects the *in vivo* situation best.

In the presented study we put emphasis on different cold exposure protocols as well as on BAT lipid uptake during diabetes. The radiotracer was synthesized according to previous work^{26,27} but sedation of mice was performed with pentobarbital instead of isoflurane. Due to the different sedative used, tracer uptake in BAT was significantly increased at 21 °C but other characteristics previously discussed were still present²⁷. An increase in tracer uptake under control conditions due to acute cold exposure was observed but could not reach significance (Fig 2 a). Similar to experiments in humans, long term cold acclimation could not

increase tracer uptake in BAT compared to exposure to 21 °C⁴⁵. This effect was also observed in mice acclimated to cold and injected with [¹⁸F]FDG and [¹⁸F]FTHA.

In diabetic mice no difference between housing at 21 °C, acute cold exposure or cold acclimation could be found. Even the pronounced uptake after acute cold exposure vs. 21 °C (Fig 2 b) could not be observed anymore. In a recent publication Heine et al. described BATs dependence on insulin to maintain its lipolytic capacity to process TRLs⁴⁴. The here presented results are in accordance to this work. Nevertheless, BAT could be visualized under conditions where [¹⁸F]FDG would have failed⁴⁶. In general BAT uptake was comparable to uptake in control mice housed at 21 °C which indicates BATs metabolic activity under diabetic conditions.

When control and diabetic animals were compared at different temperature conditions increased uptake in liver, spleen and BAT was found which also points to an insulin dependent uptake mechanism.

As a high number of patients which suffer from obesity also suffer from diabetes, additional experiments with obese mice are planned to decrypt BATs metabolism under these conditions.

In general it is to note that a combination of the here presented tracer and MRI or CT would be an optimal choice to quantify BATs lipid consumption. With [¹⁸F]BDP-TG-chylomicron-like particles it is possible to measure lipid influx, where Magnet Resonance Imaging (MRI) and Computed Tomography (CT) are able to quantify intracellular lipid usage at the same time.

Possible drawbacks might be the influence of the diet on the pharmacokinetics of the tracer and the possibility to overestimate BATs activity because of blood pool activity. A high fat diet was necessary to induce diabetic conditions. Therefore higher amounts of TGs were expected in the blood which would reduce tracer uptake by competition. Blood perfusion in BAT was found to be increased due to cold by a factor of 2⁴⁷. Therefore we cannot rule out the possibility of measuring only blood pool activity. Nevertheless, the relative low amount of activity found in the blood and the clearance of the tracer point more towards an active tracer uptake.

Conclusion

We were able to visualize BAT uptake of TRL derived FAs under different metabolic and temperature conditions. Cold acclimation of control mice could not show increased FA uptake where acute cold exposure had an observable effect. Under diabetic conditions no difference between different temperatures was noted but uptake values were comparable to control mice housed at 21 °C. [¹⁸F]BDP-TG-chylomicron-like particles showed BAT uptake under diabetic conditions. A combination between the here presented tracer and MRI or CT would be advantageous to fully quantify BATs lipid consumption.

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Supplementary Information

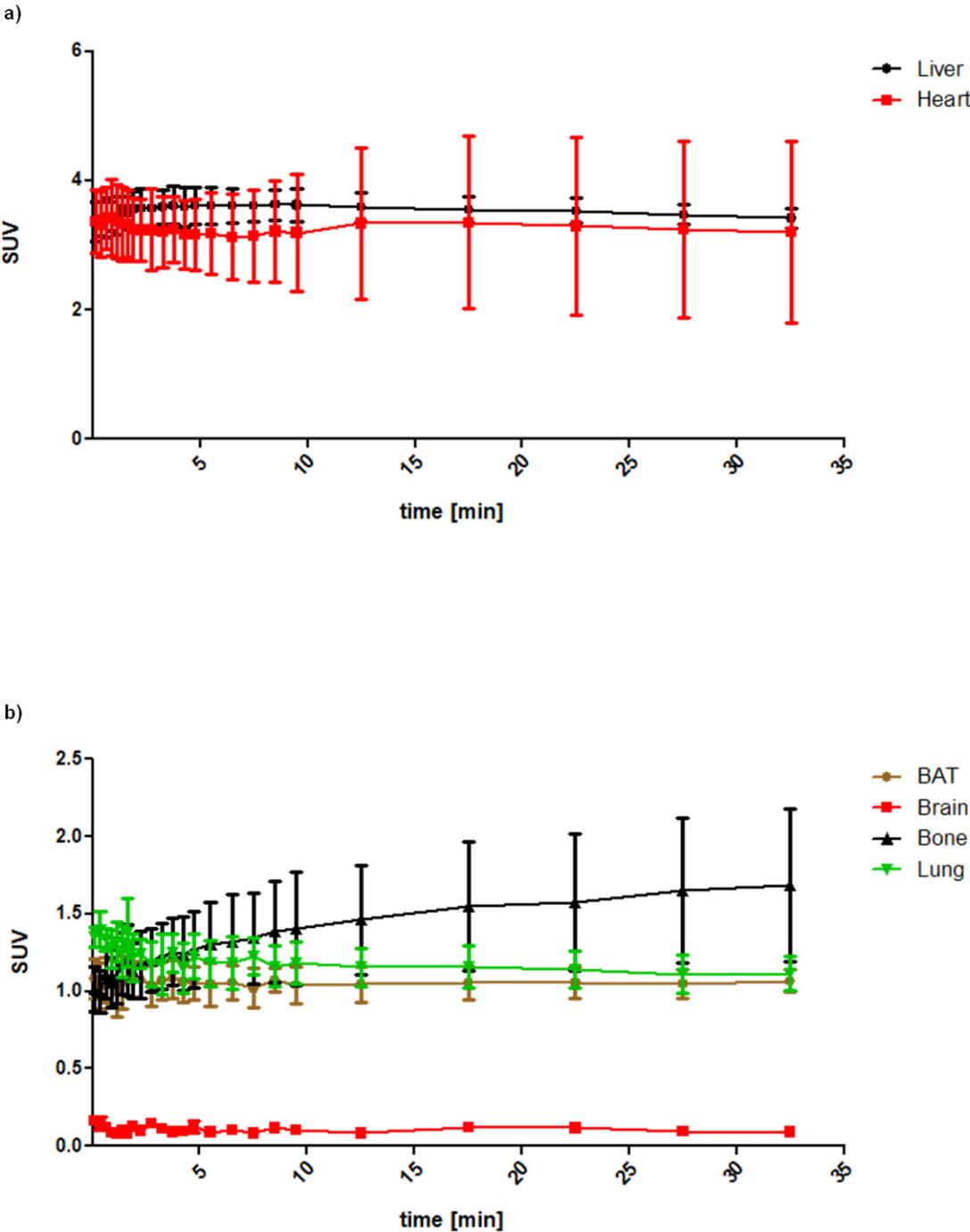


Fig S1: time activity curves in control mice housed 21 °C in a) liver and heart b) BAT, brain, bone, lung.

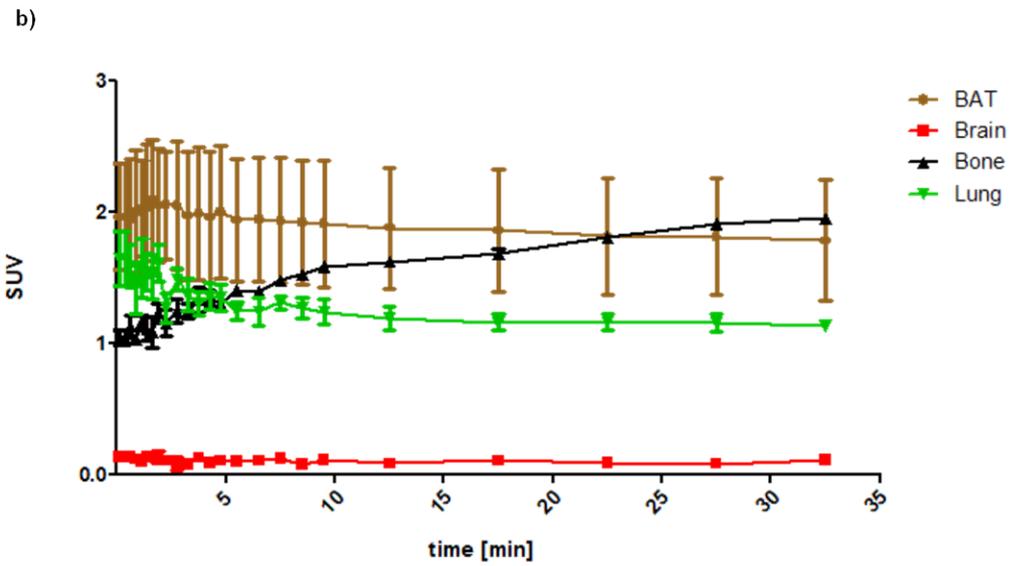
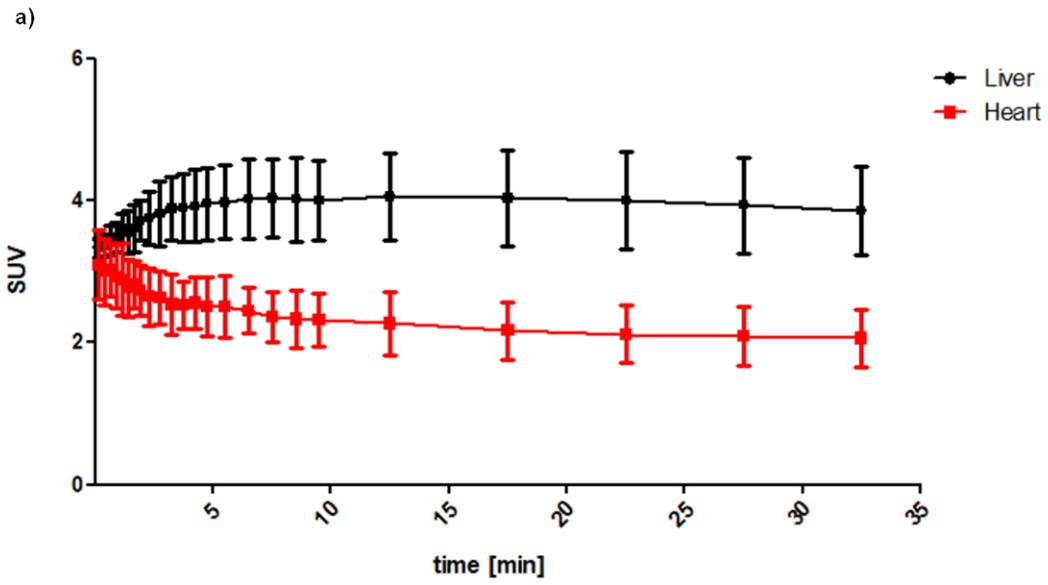


Fig S2: time activity curves in control acute cold exposed mice in a) liver and heart b) BAT, brain, bone, lung.

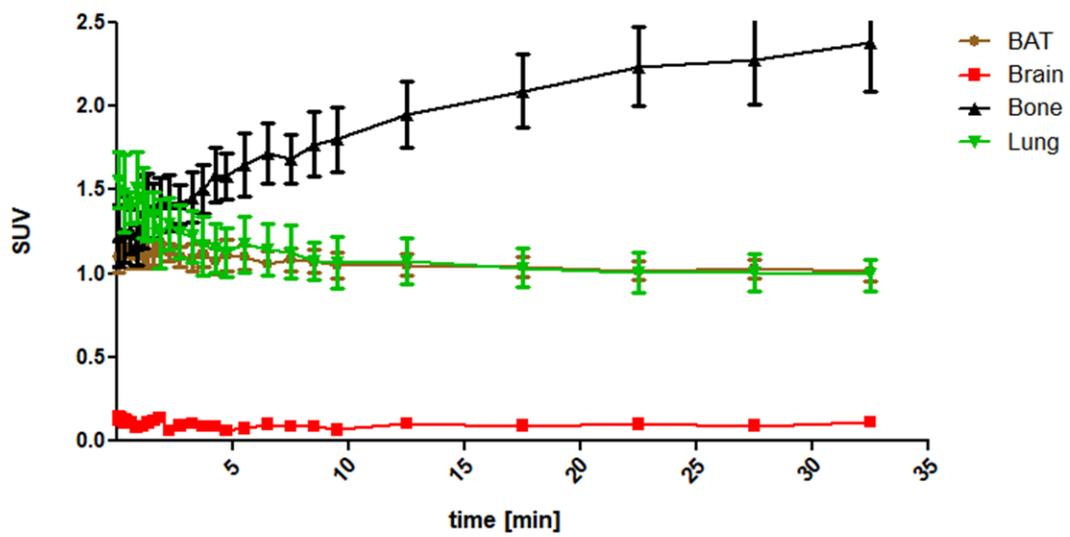
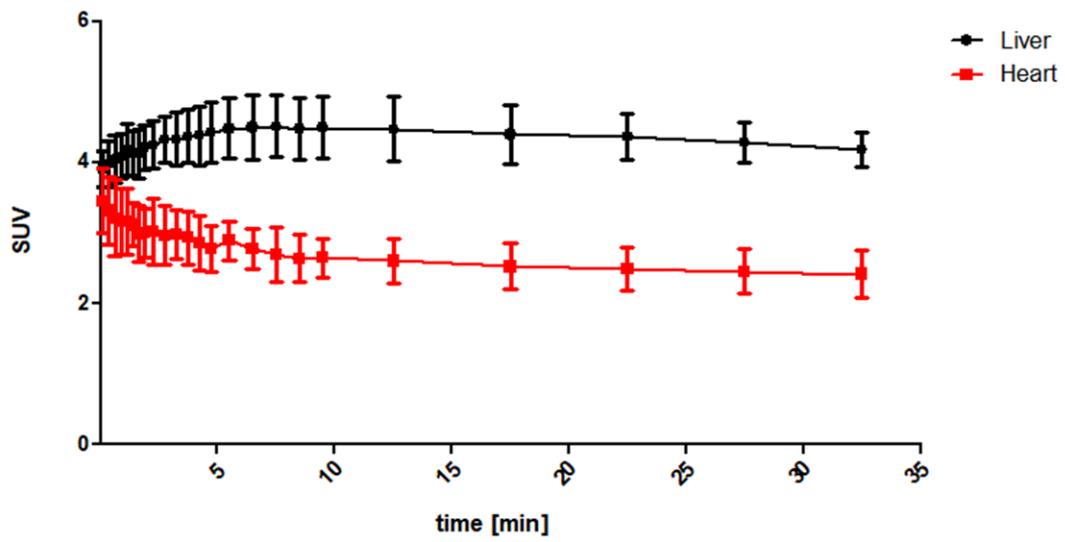


Fig S3: time activity curves in control cold acclimated mice in a) liver and heart b) BAT, brain, bone, lung.

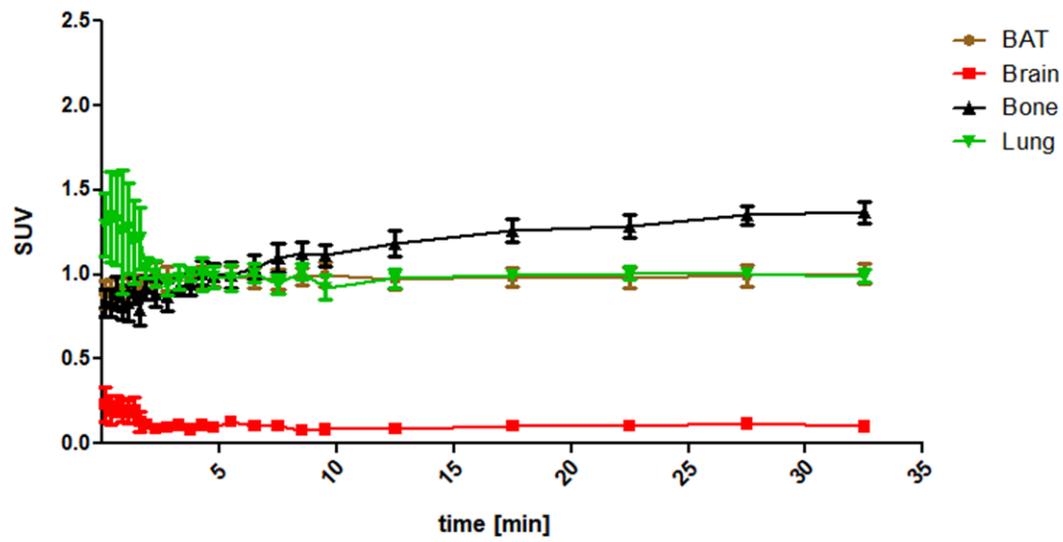
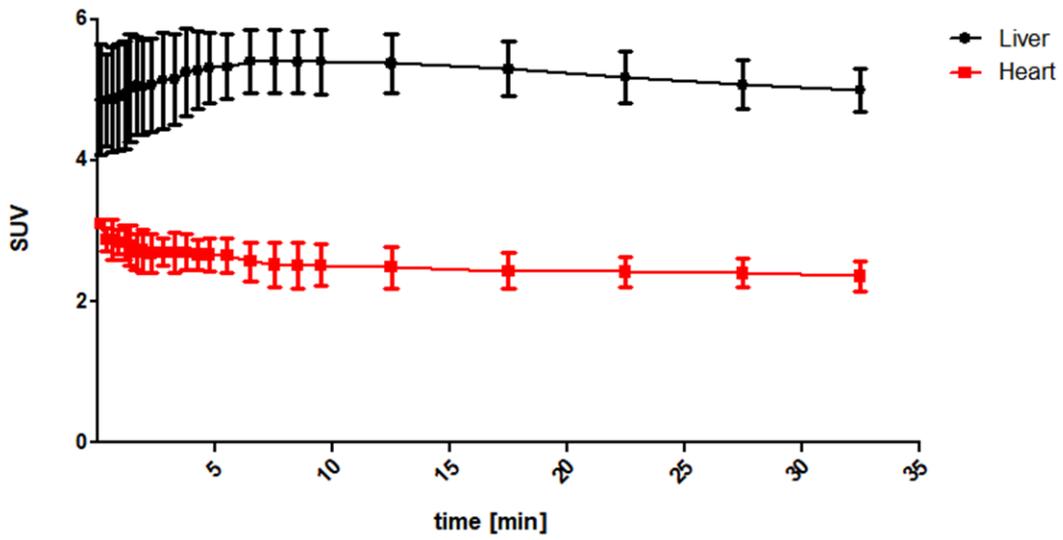


Fig S4: time activity curves in diabetic mice housed 21 °C in a) liver and heart b) BAT, brain, bone, lung.

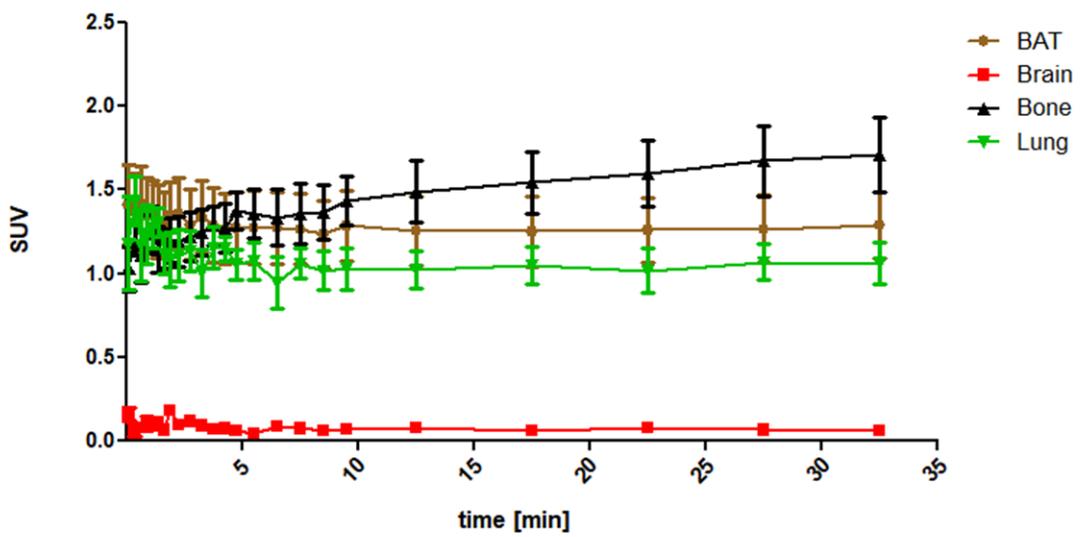
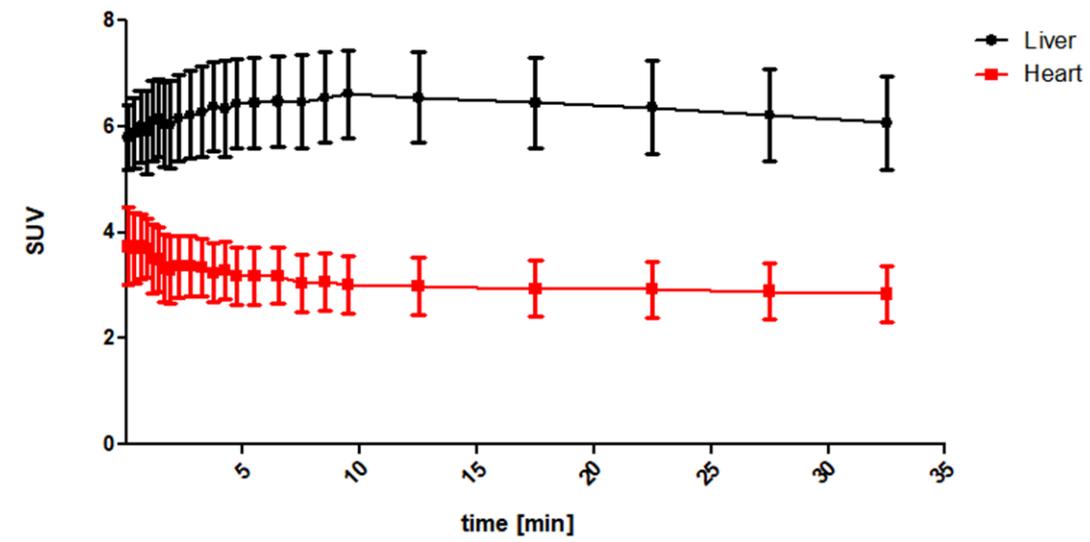


Fig S5: time activity curves in diabetic acute cold exposed mice in a) liver and heart b) BAT, brain, bone, lung.

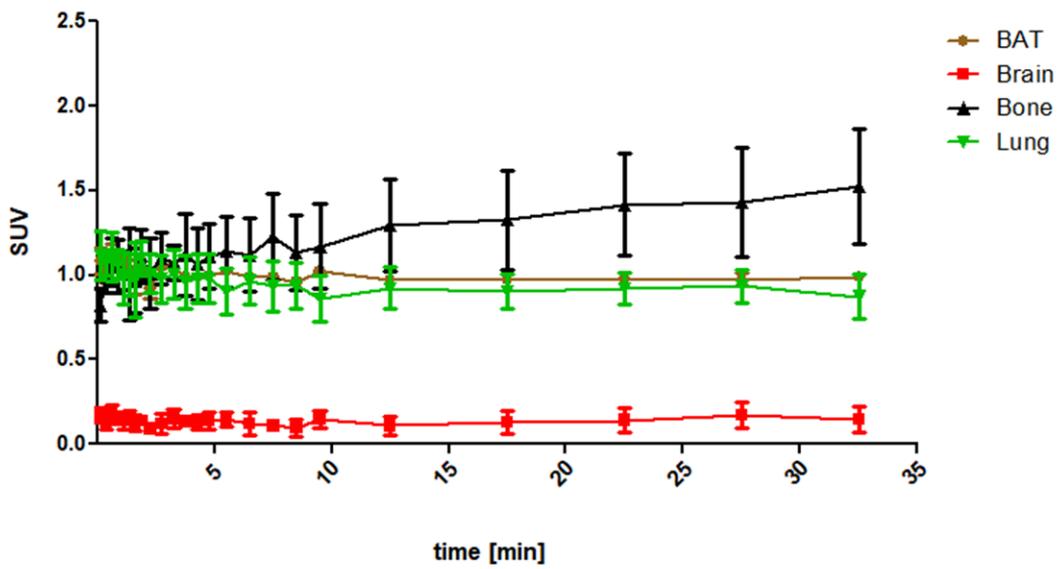
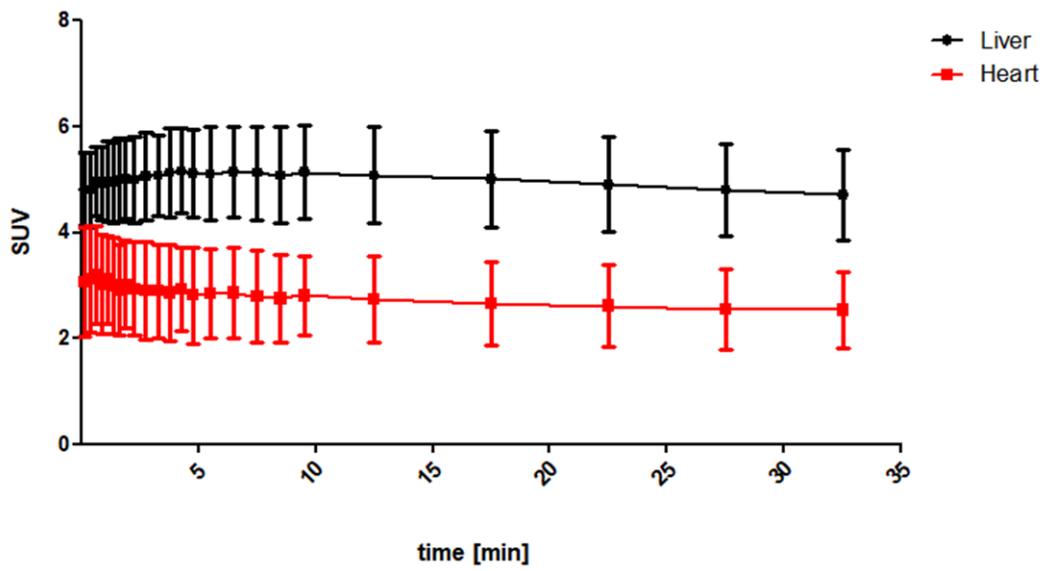


Fig S6: time activity curves in diabetic cold acclimated mice in a) liver and heart b) BAT, brain, bone, lung.

Chapter 7: Discussion

Brown adipose tissue (BAT) research started as an underestimated research field in the last century which was thought to have limited beneficial effect for adult humans. Already in the 20's ¹ brown adipocytes in humans were described and following studies ²⁻⁴ analyzed BATs function in newborns. Nevertheless, BATs size was found to decrease with aging and therefore it was thought to be irrelevant for adults ⁵⁻⁷. In the 80's of the last century significant amounts of BAT were found in outdoor workers ⁸ and since the 00's of this century active BAT was imaged and then verified in human adults based on [¹⁸F]Fluoro-2-deoxy-2-D-glucose ([¹⁸F]FDG) Positron Emission Tomography/Computed Tomography (PET/CT) ⁹⁻¹¹ and consecutive biopsies. Subsequent dedicated cold exposure experiments showed an increased [¹⁸F]FDG uptake ¹²⁻¹⁴ and short term cold exposure was able to recruit BAT even in obese subjects ¹⁵. BAT research gained more and more interest over time and new techniques and tracers to image BATs volume and metabolic activity have been developed. Nevertheless, the predominant PET tracer for BAT imaging is [¹⁸F]FDG. Its broad availability and relative long half-life offer the opportunity for non-in-house production and shipment to the place of usage. Anyways, the disadvantages of the use of a glucose tracer have been described earlier, i.e. this tracer offers only information about BAT volume and glucose consumption. As this reflects only a small portion of the energy metabolism, the overall metabolic BAT activity cannot be determined with [¹⁸F]FDG. Therefore there is need for new methods and techniques to unravel BATs contribution to whole body expenditure.

Reviewing the use of lipid metabolism imaging and its application in the field of BAT we concluded (chapter 2) that many different techniques are available to image BAT and its metabolism from which are some already translated to the clinic, some have the potential to be translated and some will have only preclinical application. MRI, SPECT and PET are all suitable to non-invasively image BAT but also the right tracer should be chosen to visualize different aspects of BAT. We expected PET imaging with radiolabeled FAs to be the most promising technique for BAT metabolism imaging because it reflects its main metabolized substance class ¹⁶ but also other imaging modalities such as fluorescence imaging and MRI

will have an impact on the exploration of BAT and its contribution to whole body energy expenditure.

To demonstrate the many different opportunities to visualize BAT we looked broader (chapter 3) and compared different imaging techniques, available in our institute, with regard to their ability to monitor BAT activity. We included MR Spectroscopy (MRS) but also PET and SPECT which was conducted with tracers which were already applied in the clinic but where no inter-tracer comparison was available. Additionally we investigated the effect of acute cold exposure and cold acclimation on the biodistribution of the different tracers as no data was available on this information. Here, we concluded that different modalities, like PET, SPECT and MRS were able to visualize BAT in rats and that BAT could also be identified by its protein and mRNA levels. With MRS we were able to quantify heat production of BAT. [^{18}F]FDG, [^{18}F]FTHA and [^{123}I]MIBG showed pronounced uptake under acute cold conditions, like reported in literature ¹⁷, but uptake was not further increased significantly by cold acclimation. In humans the situation is different. Here the SUV value increases up on chronic cold acclimation ^{18,19} but we hypothesized that BAT in rats housed at 22 °C is already acclimated to a small extend as this temperature is below their thermoneutral zone (~30 °C) ²⁰. The other tracers did not show significant increased uptake values up on cold stimulation or were not able to visualize BAT.

Based on the conclusions from chapter 2 and 3, we looked deeper into the use of radiolabelled FAs as a tool to assess BAT activity.

FA tracers like [^{18}F]FTHA and [^{123}I]BMIPP might already have advantages over [^{18}F]FDG but nevertheless FAs should not be applied as such because usually BAT takes up FAs after lipolysis of TGs resulting from triglyceride-rich lipoproteins (TRLs)²¹. Therefore a conversion of the discussed FAs to their corresponding TGs following and incorporation into TRLs would be necessary to mimic the *in vivo* situation best. We focused on this tasks and experienced several problems during the synthesis of such TGs (chapter 4) and carrier systems (chapter 5). Esterification of FAs to TG is a straight forward technique. However, most protocols focus on large amounts of precursor which is in most cases not necessary and too

expensive for the synthesis of radiotracers. Therefore, a downscaling of these reactions needed to be done. Also a radiolabeling of the TG might lead to several problems as most radiolabelings with halogens are performed in polar solvents where the TG-precursor molecule would not be soluble. An intermediate solvent system needed to be found as well as extensive purification steps to assure that, during the following incorporation of the TG into the lipoprotein-like particles, the particle itself is not altered. Quality controls had to be implemented before *in vivo/vitro* administration.

We developed a dual modal fluorescent/PET active TG (chapter 4) which was tested *in vitro* and compared to the crude FA. First we aimed to esterify FTHA but within this step all we observed was a major side-product. We hypothesised that due to the strong chlorinating agent thionyl chloride, which is used during the esterification, the thioether bond is not stable. Also by variations of the reaction conditions the product was not successfully synthesized. The next FA we choose was the commercially available BODIPY-C₁₆ (BDP-FA). This FA offers the opportunity to radiolabel it with fluorine-18 by a ¹⁹F/¹⁸F exchange reaction on its boron core but is also fluorescent which facilitates first *in vitro* testing. Esterification with 1,3-diolein was successfully performed and radiolabeling of the BDP-FA and the BDP-TG was carried out in decent yields which were in accordance to literature ^{22,23}. The following *in vitro* uptake experiments with primary human BAT and WAT cells revealed that BDP-FA is taken up with the characteristics of other FAs ($k_m = 1.15 \mu\text{M}$) ^{24,25} but not the BDP-TG. As BDP-TG is highly lipophilic it will not dissolve in the aqueous cell medium and will therefore not be available for any cells. We analyzed uptake characteristics by fluorescence and found a CD36 dependent uptake in BAT which could be stimulated by norepinephrine. WAT cells seemed to be resistant to the same interactions, which was also reported in literature ^{26,27} and therefore a different uptake mechanism was assumed. From literature ^{28,29} but also by our own experience we observed that BDP-dyes have a high potential for quenching effects as soon as they are not perfectly solubilised. Therefore, similar experiments with the [¹⁸F]BDP-FA were conducted and comparable results were found.

In chapter 4 a study describing the possibility of BDP-FA employed as a FA based tracer is presented. It is taken up with the characteristics of a FA and can be characterized by its fluorescence. It can be radiolabeled by a straight forward technique offering the opportunity for real, unquenched, uptake quantification. BDP-FA was esterified to its corresponding TG, which did not show any uptake in *in vitro* experiments due to its highly lipophilic character. Nevertheless BDP-TG could also be radiolabeled in a decent yield.

As already mentioned the main lipid source of BAT is lipolysed FAs from TRLs³⁰. Therefore, we focused (chapter 5) in a next step on the incorporation of the BDP-TG/[¹⁸F]BDP-TG into a chylomicron-like particle which we synthesized *ex vivo*. For this purpose we adapted a method from our collaborator Prof. Rensen who labelled particles with [³H]triolein³¹⁻³⁶. Different to his approach we intended to label the particles after formation, and reduce the total labelling time from 1 day to < 2 hours. A direct labeling during formation of the particles was tested but resulted only in free fluorine-18. We hypothesized that the high energies emitted from the sonicator, which is necessary for particle formation, is sufficient to break the boron-fluoride bond. Additionally long synthesis times and extensive workup would not be beneficial due to the relative short half-life of fluorine-18. We tested a variety of loading procedures (e.g. extruder, gentle sonication, heating) and incubation for 1 h at room temperature turned out to bring the highest incorporation yields. To our knowledge this is the first time chylomicron-like particles were loaded in such a way and this method might be beneficial for other attempts to label lipoproteins after their formation. Different quality control measures were taken to assure that the BDP-TG/[¹⁸F]BDP-TG is truly located in the lipid core of the particle. It was found that extensive washing of the [¹⁸F]BDP-TG is of immense importance as the SnCl₄ which is used during the radiolabeling step precipitates in aqueous medium and causes a co-precipitation of the chylomicron-like particles. Also a purification of [¹⁸F]BDP-TG from free fluorine-18 is ensured by multiple washing steps because a later purification of the [¹⁸F]BDP-TG-chylomicron-like particles by centrifugal filters caused an irreversible aggregation of the particles. *In vivo* data proved the uptake of [¹⁸F]BDP-TG-chylomicron-like particles by BAT but also high uptake values were observed in heart, liver

and spleen. Heart, as an organ with a known lipoprotein lipase expression, ^{37,38} is able to bind high numbers of lipoproteins and therefore this high uptake values are not remarkable. High uptake values for liver were explained by remnant particles which still contained [¹⁸F]BDP-TG ³⁹. The spleen contains high numbers of macrophages which are able to engulf large particles. This might explain the marked uptake observed in the spleen ⁴⁰. BAT uptake was in any case significantly increased in comparison to WAT and uptake in BAT could be significantly increased by cold exposure for 4 h before the experiment.

In chapter 5 we describe the synthesis from [¹⁸F]BDP-TG to the radiolabeled chylomicron-like particle. Rigorous washing steps are of importance and determine the quality of the final product. *In vivo* we were able to show [¹⁸F]BDP-TG-chylomicron-like particles suitability as a BAT imaging agent.

As we proved the suitability of [¹⁸F]BDP-TG-chylomicron-like particles to visualize BAT and its metabolism we went a step further (chapter 6) and investigated its behaviour under different cooling protocols and under diabetic conditions. Interestingly we could only observe an increase in [¹⁸F]BDP-TG uptake under acute cold stimulation in BAT where this effect was not visible after long term cold acclimation. Also no significant difference was detected in diabetic animals and both groups BAT uptake differed only under acute cold conditions.

As already discussed, most BAT scans in patients are performed with [¹⁸F]FDG. Here the same phenomenon was observed where [¹⁸F]FDG BAT uptake was impaired in diabetic patients but uptake of [¹⁸F]FTHA was not altered when compared to non-diabetic controls ⁴¹. This may lead to the assumption that glucose uptake is not an indicator for thermogenesis in BAT ⁴².

We compared the obtained results with data of [¹⁸F]FDG and [¹⁸F]FTHA under the same cooling protocol, but not in diabetic mice. Similar results were observed for [¹⁸F]FDG and [¹⁸F]FTHA where cold acclimation could not show an effect. These results are in accordance with the results we obtained in chapter 3 in a rat model where cold acclimation could not further increase any of the applied tracers, including [¹⁸F]FDG and [¹⁸F]FTHA in comparison to acute cold exposure, and where no elevated mRNA genes and protein levels related to

BAT metabolism could be found. Cold acclimation leads to browning⁴³ and therefore to a total increase of brown and beige BAT cells. We speculated that under long term cold exposure the single BAT depots might not be stimulated to such an extent anymore as under acute cold exposure because the total BAT mass has grown. An additional explanation would be that under acute cold the FA influx into BAT is up-regulated whereas by cold acclimation an equilibrium is reached where the influx is regulated to a constant level.

In the presented thesis we produced a radiolabeled TG which can be incorporated into lipoprotein-like particles for non-invasive *in vivo* imaging of lipid metabolism. This is to our knowledge the first time that PET active TGs were incorporated into chylomicron-like particles. Additionally those particles and also the BDP-FA can be used to quantify BAT uptake *in vitro* providing the opportunity for first testings without the use of radioactive material. We put emphasis on imaging the lipid uptake of BAT but this method can easily be adapted for other applications such as e.g. imaging of lipid uptake into the liver or heart. This methodology has the potential for *in vivo* imaging in humans but more work needs to be done, namely: 1) Exchange of the BDP-FA to a FA which can be used in humans. 2) Quality control of radiolabeled lipoproteins needs to be optimized.

Nevertheless, we have the opinion that imaging with radiolabeled TGs incorporated into lipoproteins mimics the *in vivo* situation, how lipids are transported in the body, better than other applied techniques so far and is therefore a huge step forward in the field of lipid metabolism imaging.

Conclusion

In our experiments we found that [¹⁸F]BDP-TG-chylomicron-like particles are able to reach BAT and its uptake is up-regulated by acute cold stimulation. We also found that BAT did not show high lipid influx by cold acclimation. This may implicate that the contribution of BAT may not be as high as in optimistic studies a decade ago ⁴⁴.

However, we have developed a new lipid based imaging agent for BAT, which will help to uncover BATs contribution to whole body energy expenditure but which also can easily be adapted for other applications. The developed dual-modal tracer combines several advantages and has therefore a broad field of applications ranging from first *in vitro* testings without any radiation dose to non-invasive PET imaging.

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Valorization

Social relevance

Obesity has long been a severe issue but newest reports from the WHO show alarming numbers on the development of obesity. In 2016 worldwide more than 1.9 billion adults were overweight from which 650 million were obese. Frighteningly also 41 million children under the age of 5 and over 340 million children between the age of 5 and 19 were overweight or obese. According to the Robert Koch Institute 66% men and 50% women were overweight in Germany in 2011. In total 25% were obese while this number has tripled between 1975 and 2014. Obesity is related to a number of other diseases such as diabetes type II, cardiovascular diseases and cancer. In 2015 worldwide deaths of 4 million people could be directly related to obesity but also people suffering from overweight have a decreased life expectancy. In the future the prevalence to develop obesity is predicted to rise to 33% (United States) within the next two decades. Therefore action needs to be taken to counteract this epidemic.

Obesity is the result of a positive energy balance. This can be overcome by a reduction of energy intake or an increase in energy expenditure. Reduction of energy intake is most often addressed by diets but in studies it could be shown that only a small number of patients could maintain their weight after a diet. The easiest way to increase energy expenditure might be physical activity. Similar like diets it was shown that people had problems following exercise plans on a regular level. Additionally, physical exercise might not be performed by extremely obese patients which attenuates its role on weight loss.

An alternative strategy to increase energy expenditure is the activation of brown adipose tissue (BAT). BATs contribution to overall energy expenditure is estimated to be 2 - 30% in humans but its role in lipid metabolism is not completely understood yet. Therefore this thesis is focused on BAT imaging to gain more information about its metabolism which should help to level out the impaired energy balance of obese people in the future.

Target group

As reported in the last section, obesity is a worldwide epidemic. Finding new strategies to prevent people from becoming obese and reducing body weight of already obese people will have a huge impact on our worldwide society. An enormous number of overweight and obese people (1.9 billion) will be affected where obese people will benefit most. Nevertheless, overweight people will also gain quality of life after weight reduction. BAT might be an interesting target to solve this problem. Even in the past, studies already showed that BAT can be activated by mild cold exposure which should have an impact on how we are tempering our houses and offices. Anyhow, more research has to be done to obtain coherent results of such interventions but on the other hand it is important to inform the general public to call attention to this subject. Even though this study is more related to basic research, it might help to understand BATs metabolism and will therefore have on a long term an effect on the general society.

Activities

The here presented work was carried out as a collaboration between the Department of Radiology and Nuclear Medicine, Department of Nutrition and Movement Sciences, Nuclear Medicine of Maastricht University Medical Center+ and Department of Nuclear Medicine of University Hospital RWTH Aachen. Additionally, the Institute for Experimental Molecular Imaging of the University Hospital RWTH Aachen and the Department of Medicine of Leiden University Medical Center were involved in parts of the project. Specialists in the field of chemistry, biology, radiochemistry, animal handling, positron emission tomography, electron microscopy and BAT metabolism were involved. This extensive collaboration between different institutions and different fields of science shows the amount of knowledge that is necessary to carry out such studies.

The work performed in this thesis was presented in original articles that were published in scientific journals with a background in radiochemistry, imaging and BAT metabolism. The articles are not only published in those journals but are also available online and some are

even open access. Therefore the obtained results are available to the broad scientific community. Results were also presented on national and international conferences. A new method to load lipoproteins with radiolabeled TGs was described and may improve BAT imaging in the near future.

Innovation

Active BAT in adult humans has first been described in the 00's of this century. A combination of positron emission tomography and computed tomography (PET/CT) helped to identify what has long thought to be artefacts as BAT depots. Till now most human scans were performed with 2-deoxy-2- ^{18}F fluoroglucose (^{18}F FDG) and a lot of retrospective studies with the same tracer were conducted. Even though ^{18}F FDG is able to visualize BAT and uptake was observed to be increased after cold exposure, it only reflects BATs glucose metabolism. In BAT mainly fatty acids (FA) are metabolised where *de novo* lipogenesis from glucose plays a minor role. Therefore ^{18}F FDG might largely underestimate BATs metabolic activity and a FA based tracer is needed to quantify BATs metabolic activity and potential. Some FA tracers are already in use for clinical test but even they do not completely reflect the situation *in vivo*. It was shown that BAT predominantly takes up lipids from triglyceride (TG)-rich lipoprotein (TRL)-derived FAs over circulating albumin-bound FA. Therefore application of radiolabeled free FA might not represent BATs metabolism completely.

The aim of this study was to develop a radiolabeled TG and to incorporate it into a lipoprotein particle which has been synthesized *ex vivo*. This method has the advantage that uptake of BAT does not rely on additional uptake mechanism of free FAs *in vivo* which would complicate data interpretation. To our knowledge *ex vivo* synthesized particles were only loaded with TGs radiolabeled with tritium which cannot be used for *in vivo* imaging. Therefore our approach is a step forward in the field of BAT visualization and quantification. In addition, the FA used in this work also offers the ability for fluorescence imaging. *In vitro* experiments could be carried out without any radiation and the behaviour could be characterized on a sub-cellular level.

Schedule & Implementation

In this thesis we developed a new lipid based radiotracer to image BAT and its metabolism. We could demonstrate that the radiotracer is taken up by BAT under control conditions and that the uptake could be increased significantly after cold exposure. To our knowledge this is the first time that radiolabeled TGs were incorporated into lipoprotein-like particles for *in vivo* imaging. We successfully gained information about BATs lipid metabolism under control and cold exposed conditions but also under diabetes, obesity and long term cold acclimation. The here presented methods can be applied to incorporate other radiolabeled FAs into lipoproteins and by that the current generation of FA tracer which are used in the clinic might be lifted up to another level because the *in vivo* situation is mimicked better than with only radiolabeled FA.

More animal experiments have to be performed in the future to gain more information about BAT and its metabolic activity. As large scale human experiments are restricted by ethical and radiation protection laws we need animals to understand BATs function and translate those results to humans.

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I cannot believe that these are my last words I am writing on my way to become a PhD. I am still thinking back at the time when I was in secondary school where I had large discussions with my parents and my grandmother about my school performance which was probably not the best ;). However, I always told them that the time I invested for learning was sufficient and that I do not need most of the subjects anyways. Probably nobody, including me, expected me to become a PhD but somehow I made it to this point. At the beginning, in 2015, I started here in Maastricht and thought that 4 years is a really long period but time was flying so quickly and I got to know so many great colleagues that I am thinking about the end of my PhD with a laughing but also with a crying eye. Thanks to all of you that I could learn so many techniques from you but also for working together as one big team. Without you this dissertation would not be possible.

Prof. Mottaghy thanks for offering me this super interesting PhD position in Maastricht and in Aachen. I still remember the day of my job interview in Aachen and you started with the sentence: "We do not have an open position here". I was confused but then you gave me the chance to work abroad in Maastricht. At first I was a bit sceptical but then it turned out to be a great decision as I could take advantage of both labs. Thank you for giving me the chance to visit that many interesting courses and conferences which also helped to develop myself as an independent researcher. In our meetings you always had good advices to bring the project a step further. Even though you are also working part time in Aachen and Maastricht I had the feeling that in case I experienced a problem I could always come to you to ask for help. I think that I had a great promotor in you.

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Curriculum Vitae

Andreas Paulus was born on the 27th of October 1988 in Düren, Germany. He finished his secondary school at Burgau Gymnasium in Düren in 2008. After receiving his Bachelor degree in Applied Chemistry in 2012 he continued with his Master in Nuclear Application at the Fachhochschule Aachen. During this time he spend 6 months as an undergraduate researcher at the Memorial Sloan Kettering Cancer Center in New York City working on the development of new clickable radiotracer for prostate cancer imaging. In 2015 he obtained his master degree and started as a PhD student under the supervision of Prof. Dr. F. M. Mottaghy and Prof. Dr. W.D van Marken Lichtenbelt at the Department of Radiology and Nuclear Medicine at Maastricht University. He focused on the development of new lipid based imaging agents for brown adipose tissue as demonstrated in this dissertation as well as in numerous publications in scientific journals.

List of publications

Published

- 1 Paulus, A., van Marken Lichtenbelt, W., Mottaghy, F. M. & Bauwens, M. Brown adipose tissue and lipid metabolism imaging. *Methods* **130**, 105-113, doi:10.1016/j.ymeth.2017.05.001 (2017).
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