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Inflammatory cytokine oncostatin M induces endothelial activation in macro- and microvascular endothelial cells and in APOE*3Leiden.CETP mice

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Abstract

Aims

Endothelial activation is involved in many chronic inflammatory diseases, such as atherosclerosis, and is often initiated by cytokines. Oncostatin M (OSM) is a relatively unknown cytokine that has been suggested to play a role in both endothelial activation and atherosclerosis. We comprehensively investigated the effect of OSM on endothelial cell activation from different vascular beds and in APOE*3Leiden.CETP mice.

Methods and results

Human umbilical vein endothelial cells, human aortic endothelial cells and human microvascular endothelial cells cultured in the presence of OSM express elevated MCP-1, IL-6 and ICAM-1 mRNA levels. Human umbilical vein endothelial cells and human aortic endothelial cells additionally expressed increased VCAM-1 and E-selectin mRNA levels. Moreover, ICAM-1 membrane expression is increased as well as MCP-1, IL-6 and E-selectin protein release. A marked increase was observed in STAT1 and STAT3 phosphorylation indicating that the JAK/STAT pathway is involved in OSM signaling. OSM signals through the LIF receptor alfa (LIFR) and the OSM receptor (OSMR). siRNA knockdown of the LIFR and the OSMR revealed that simultaneous knockdown is necessary to significantly reduce MCP-1 and IL-6 secretion, VCAM-1 and E-selectin shedding and STAT1 and STAT3 phosphorylation after OSM stimulation. Moreover, OSM administration to APOE*3Leiden.CETP mice enhances plasma E-selectin levels and increases ICAM-1 expression and monocyte adhesion in the aortic root area. Furthermore, IL-6 mRNA expression was elevated in the aorta of OSM treated mice.
Conclusion

OSM induces endothelial activation in vitro in endothelial cells from different vascular beds through activation of the JAK/STAT cascade and in vivo in APOE*3Leiden.CETP mice. Since endothelial activation is an initial step in atherosclerosis development, OSM may play a role in the initiation of atherosclerotic lesion formation.

1. Introduction

The endothelium is involved in many processes including maintenance of the endothelial barrier function, prevention of spontaneous blood clot formation, inflammatory cell recruitment upon injury and regulation of the vascular tone[1–3]. Impairment of one or more of these functions is often referred to as endothelial dysfunction, and may lead to the development of atherosclerosis, angiogenesis in cancer, vascular leakage, infectious diseases or stroke[4].

Although endothelial dysfunction is often described as the inability to dilate vessels, endothelial dysfunction is also characterized by endothelial activation, which is marked by increased cytokine release, adhesion molecule expression and endothelial permeability. The released cytokines attract leukocytes to the site of the activated endothelium, where the leukocytes bind to the endothelial barrier, which is enabled by enhanced adhesion molecule expression. Firmly adhered leukocytes then migrate through the endothelial barrier into the underlying tissue[5].

The process of endothelial activation can occur both, locally on well-known predilection sites and systemically, and is often triggered by traditional cardiovascular risk factors such as hypercholesterolemia, hypertension, smoking or diabetes and is initiated by inflammatory cytokines. One such a cytokine, which was first discovered in the cancer field, is oncostatin M (OSM). This relatively unexplored cytokine is an interleukin-6 family member that can signal through the LIFR and the OSMR, which are both dependent on heterodimerization with the gp130 receptor to form a functional receptor complex[6]. OSM is upregulated in multiple chronic inflammatory diseases including periodontitis, rheumatoid arthritis and inflammatory bowel diseases and is known to induce angiogenesis and smooth muscle cell proliferation and migration, both processes that are involved in atherosclerosis development[7–16]. Other pro-inflammatory cytokines that promote angiogenesis, smooth muscle cell proliferation and endothelial activation, such as TNFα and IL-18, have already been proven to accelerate atherosclerosis[17–24]. Furthermore, OSM is found in human carotid atherosclerotic plaques and in the intima and media of atherosclerotic mice[16].

Based on these findings and on the knowledge that endothelial cells are very high expressers of OSM receptors[25], we hypothesized that OSM may be involved in atherosclerosis development partially by inducing endothelial activation as a first step in the development of atherosclerosis. In this study, we incubated human endothelial cells with OSM to investigate if OSM induces systemic or local endothelial activation. As the cell heterogeneity among endothelial cells is huge[26,27] and endothelial cells from different vascular beds show different responses/behavior different to physiological stimuli[28,29], we tested the effect of OSM in endothelial cells derived from multiple vascular beds, human umbilical vein endothelial cells (HUVECs), human aortic endothelial cells (HAECs) and human microvascular endothelial cells (HMEC-1). Of which HAECs are the most suitable endothelial cell type to study atherosclerosis development as atherosclerosis mainly affects the medium and large-sized arteries[30]. To validate our findings in cultured endothelial cells in vivo, we administered OSM to APOE*3Leiden.CETP mice, a translational mouse model for hyperlipidemia and atherosclerosis[31,32].
mildly pro-inflammatory state that is present in this animal model of hyperlipidemia makes it a suitable model to investigate the role of OSM in atherosclerosis prone conditions. We found that OSM induces endothelial activation in all different investigated human endothelial cell types and in mice after chronic administration and identified the JAK/STAT pathway as a key player in this process.

2. Materials and methods

2.1 Cell culture

2 different batches of pooled primary human umbilical vein endothelial cells (HUVECs, Lonza, the Netherlands), a single batch of primary human aortic endothelial cells from one single donor (HAECs, ATCC, Manassas, VA, USA) and a human dermal microvascular endothelial cell line (HMEC-1, ATCC, Manassas, VA, USA) were cultured in EBM-2 medium (Lonza, Walkersville, MD) supplemented with EGM-2 SingleQuots (Lonza, Walkersville, MD) under normoxic conditions (21% O2). Throughout the study, passage 6 was used for HUVECs and HAECs, while passage 27 was used for the HMEC-1 cell line. All experiments were performed in 70% subconfluent HUVECs, HAECs, or HMEC-1 cells. After each experiment, cells and conditioned medium were collected for subsequent RNA or protein analysis. Repetitive experiments were only started if the previous experiment had been finished.

2.2 In vitro RNA expression

Human OSM (R&D systems, Minneapolis, MN) was added to HUVECs, HAECs and HMEC-1 cells in a concentration range from 0–20 ng/mL. After 3 or 6 hours, RNA was isolated with the NucleoSpin RNA kit (Macherey-Nagel, Düren, Germany) according to the manufacturer’s protocol. Isolated RNA (500 ng) was reverse transcribed into cDNA with the qSCript cDNA Synthesis Kit (Quanta Biosciences, Beverly, MA) and analyzed by real-time fluorescence assessment of SYBR Green signal (iQ SYBR Green Supermix, Bio-Rad, Hercules, CA) in the CFX96 Real-Time Detection System (Bio-Rad, Hercules, CA). Each sample was measured in duplicates. Primers were designed for the human genes of interest, sequences are listed in Table 1. MRNA levels were analyzed and corrected for the housekeeping gene ACTB. Experiments were repeated 4–7 times.

2.3 In vitro cytokine release

To determine the effect of OSM on endothelial activation, HUVECs, HAECs or HMEC-1 cells were incubated with 5 ng/mL OSM. 3h and 6h after OSM treatment, conditioned medium was collected. To investigate the effect of OSM on endothelial activation after siRNA knockdown of the LIFR and OSMR, siRNA transfected HUVECs were treated with 5 ng/mL OSM 48h post transfection. 6h after OSM treatment conditioned medium was collected. Conditioned medium was analyzed with the ProcartaPlex Mix&Match Human 6-plex (Thermo Fisher, Waltham, MA) according to the manufacturer’s protocol and measured on the Bio-plex 200 system (Bio-Rad, Hercules, CA) to determine the release of MCP-1, IL-6, soluble E-selectin, soluble P-selectin and soluble VCAM-1. Experiments were repeated 3–7 times.

2.4 Flow cytometry

5 ng OSM was added to HUVECs, HAECs, or HMEC-1 cells for 18h. Cells were washed with PBS and detached with accutase. Subsequently, cells were fixed with 1% PFA and incubated with 2.5 μL antibodies/1,000,000 cells against VCAM-1, ICAM-1, P-selectin and, E-selectin all obtained from Thermo Fisher (S1 Table). The experiment was repeated 3 times.
2.5 siRNA transfection

Knockdown of LIFR and OSMR was achieved by transfection with a mix of 4 specific siRNA sequences directed against the human mRNA sequence (SMARTpool siGENOME, GE Dharmacon, Lafayette, CO) in 70% subconfluent HUVEC cultures. Cells were incubated for 1 hour in a small volume of EGM-2 medium supplemented with DharmaFECT 1 (GE Dharmacon, Lafayette, CO) according to manufacturer’s instructions. After 2 hours cells were supplemented with extra EGM-2 medium to complement medium volumes. As controls, HUVECs were transfected with a mix of 4 scrambled, non-targeting siRNAs (siSham Smartpool; GE Dharmacon, Lafayette, CO). siRNA transfected HUVECs were treated with OSM 48h after siRNA transfection.

2.6 Western blot

HUVECs were lysed with cOmplete™ Lysis-M, EDTA-free reagent (Sigma Aldrich, Saint Louis, MO) for 15 minutes on ice. Next, protein concentration was determined with the Pierce™ BCA protein Assay Kit (Thermo Scientific, Waltham, MA). The protein sample was treated with NuPAGE™ Sample Reducing Agent (Thermo Scientific, Waltham, MA) and NuPAGE™ LDS Sample Buffer (Thermo Scientific, Waltham, MA). Subsequently, the solution was boiled at 70°C for 10 minutes. Samples were loaded on a Bolt™ 4–12% Bis-Tris Plus gel (Thermo Scientific, Waltham, MA), run for 50 minutes at 160V and transferred to an iBlot®2 PVDF Stack (Thermo Scientific, Waltham, MA) with the iBlot®2 Gel Transfer Device (Thermo Scientific, Waltham, MA). Blots were incubated with the primary antibody overnight at 4°C (S1 Table). Subsequently, blots were incubated with the appropriate secondary antibody conjugated with horseradish peroxidase (HRP) for 1h at RT (S1 Table). Peroxidase labeled antibodies were detected with Chemiluminescent Peroxidase Substrate (Sigma, Saint Louis, MO).

### Table 1. Primer sets for qPCR analysis.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Species</th>
<th>Direction</th>
<th>Primer sequence (5’-3’)</th>
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<tr>
<td>MCP-1</td>
<td>Human</td>
<td>Forward</td>
<td>TGGGATCTTGAACCCCAGCTCT</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Reverse</td>
<td>CAGCCAGATGCAAATCAAGGCC</td>
</tr>
<tr>
<td>IL-6</td>
<td>Human</td>
<td>Forward</td>
<td>AGTGAGGAAACAAGCCAGAGG</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Reverse</td>
<td>GTCAGGGGTTATGATGATGAT</td>
</tr>
<tr>
<td>ICAM-1</td>
<td>Human</td>
<td>Forward</td>
<td>TTGAAACCCACAGTCACCTAT</td>
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<tr>
<td></td>
<td></td>
<td>Reverse</td>
<td>CCTCTGACGGTCGATACCA</td>
</tr>
<tr>
<td>VCAM-1</td>
<td>Human</td>
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<td></td>
<td>Reverse</td>
<td>GTCTCCAACTGAGCCAGCAA</td>
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<tr>
<td>E-SELECTIN</td>
<td>Human</td>
<td>Forward</td>
<td>AAGCCTCTGACTGACGGAAG</td>
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<tr>
<td></td>
<td></td>
<td>Reverse</td>
<td>TCCCTCTAGTTCCACAGAGT</td>
</tr>
<tr>
<td>ACTB</td>
<td>Human</td>
<td>Forward</td>
<td>GATCGGCGGCTCCATCTTG</td>
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<td>Reverse</td>
<td>GACTGAGTACACTCTCTGCC</td>
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<tr>
<td>Msp-1</td>
<td>Murine</td>
<td>Forward</td>
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<td>Icam-1</td>
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<td></td>
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<td>Reverse</td>
<td>CAGCCAAAGCTGCTGAAACA</td>
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2.7 Animals and treatments

Thirty-two female APOE*3Leiden.CETP transgenic mice (15–22 weeks of age) were used. The number of animals per group was calculated with Java Applets for Power and Sample Size [Computer software], from http://homepage.stat.uio.wa.edu/~rlenth/Power/index.html using a one-way ANOVA with a probability of 0.05 and a Dunnett’s correction, a SD of 20%, a power of 80% and a minimal expected difference of 35%. Mice were housed under standard conditions with a 12h light-dark cycle and had free access to food and water. Body weight, food intake and clinical signs of behavior were monitored regularly during the study. Mice received a Western type diet (WTD) (a semi-synthetic diet containing 15 w/w% cacao butter and 0.15% dietary cholesterol, Altromin, Tiel, the Netherlands). At T = 0 weeks, after a run-in period of 3 weeks, mice were matched based on plasma total cholesterol levels, plasma triglyceride levels, body weight, and age in 4 groups of 8 mice. Two mice died during the diet intervention period, 1 in the 1μg/kg/day OSM group and 1 in the 10μg/kg/day OSM group. At T = 7 weeks, an ALZET® Osmotic Pump Type 1004 (4-week release duration, Durect, Cupertino, CA) containing either 1, 3 or 10μg/kg/day murine OSM (R&D systems, Minneapolis, MN) or PBS was placed subcutaneously in the flank. Doses were based on previous studies, which gave a single or double injection of 5–50 μg/kg OSM resulting in local increased permeability, edema, swelling, infiltration of immune cells, increased serum VEGF levels and increased angiopoetin 2 expression[33–36]. All solutions, also PBS of control group, contained 1% mouse serum to prevent OSM from sticking to plastics. Prior to surgery, mice received the analgesic Carprofen (5 mg/kg s.c.) and were anesthetized with isoflurane (induction 4%, maintenance 2%). At T = 10 weeks, mice were euthanized by gradual CO2 inhalation (6 L/min in a 20 Liter container). CO2 flow was maintained for a minimum of 1 minute after respiration ceased (as observed by lack of respiration and faded eye color). Death was confirmed by exsanguination (via heart puncture). Hearts were isolated for immunohistochemistry in the aortic root and aortas were isolated for RNA expression analysis. EDTA blood samples were drawn after a 4 hour fast at T = 0 and T = 10 weeks. All animal experiments were performed conform the guidelines from Directive 2010/63/EU of the European Parliament on the protection of animals used for scientific purposes or the NIH guidelines. The care and use of all mice in this study was carried out at the animal facility of The Netherlands Organization for Applied Research (TNO) in accordance with the ethical review committee “TNO-DEC” under the registration number 3683. Animal experiments were approved by the Institutional Animal Care and Use Committee of TNO under registration number TNO-202.

2.8 Plasma parameters

Plasma cholesterol and triglycerides were measured spectrophotometrically with enzymatic assays (Roche diagnostics). The inflammatory markers, E-selectin and MCP-1 were measured with ELISA kits from R&D. Plasma ALT and AST were determined using a spectrophotometric assay (Boehringer Reflotron system) in group wise-pooled samples from sacrifice plasma. All assays were performed according to the manufacturer’s instruction.

2.9 Histological assessment of vascular inflammation

Vascular inflammation was assessed in the aortic root area as reported previously by Landlinger et al[37] in control mice and mice receiving 10 μg/kg/day OSM. Briefly, the aortic root was identified by the appearance of aortic valve leaflets and serial cross-sections of the entire aortic root area (5 μm thick with intervals of 50 μm) were mounted on 3-aminopropyl triethoxysilane-coated slides and stained with hematoxylin-phloxine-saffron (HPS). Each section consisted of 3 segments (separated by the valves) and in 4 sections ICAM-1 expression and the
number of monocytes adhering to the activated endothelium was counted after immunostaining with mouse monoclonal ICAM-1 antibody (Santa Cruz) and AIA 31240 antibody (Accurate Chemical and Scientific) respectively (S1 Table). One mouse from the control group was excluded from analysis due to a technical error, resulting in 7 and 8 mice per group.

### 2.10 RNA isolation murine tissue

To isolate RNA from aortic tissue, RA1 lysis buffer (Macherey-Nagel, Düren, Germany) containing 1% DTT was added to the tissue, which was cut in tiny pieces and subsequently minced. RNA was isolated with the RNeasy® Plus Micro Kit (Qiagen, Hilden, Germany) according to the RNeasy Fibrous Tissue Mini Kit protocol (Qiagen, Hilden, Germany). Isolated RNA (500 ng) was reverse transcribed into cDNA with the qScript™ cDNA Synthesis Kit (Quanta Biosciences, Beverly, MA) and analyzed by real-time fluorescence assessment of SYBR Green signal (iQ™ SYBR® Green Supermix, Bio-Rad, Hercules, CA) in the CFX96™ Real-Time Detection System (Bio-Rad, Hercules, CA). Each sample was measured in duplicates. Primers were designed for the murine genes of interest, sequences are listed in Table 1. mRNA levels were analyzed and corrected for the housekeeping gene Hprt. RNA isolation was unsuccessful in one mouse from the 3μg/kg/day OSM group resulting in 6, 7 and 8 mice per group.

### 2.11 Statistical analysis

qPCR data was analyzed according to the ΔΔCt method, statistical tests were performed on ΔCt values. Two-way-anova was used to analyze in vitro data to take into account day-to-day variation of the experiments. Not normally (Gaussian) distributed parameters were transformed with the natural logarithm or in case of undetectable values analyzed with the appropriate non-parametric test. Dose-dependency was determined by a Pearson correlation. All statistical analyses were performed in SPSS statistics version 21.0. A two-tailed p-value of 0.05 was regarded statistically significant in all analyses. Graphs were made in GraphPad Prism version 7.02 for Windows, GraphPad Software, La Jolla California USA, www.graphpad.com

### 3. Results

#### 3.1 OSM induces endothelial activation in human endothelial cells

To investigate whether OSM induces endothelial activation, we first examined cytokine mRNA expression in HUVECs, HAECs and HMEC-1 cells treated with 5 ng/mL OSM for 3 or 6 hours. OSM treatment was found to increase mRNA expression of the cytokines MCP-1 (p<0.01) and IL-6 (p<0.001) in HUVECs, HAECs (p<0.001) and HMEC-1 cells (p<0.001) at both 3h and 6h time points (Fig 1A–1F). Since these cytokines are released by activated endothelial cells, we next measured MCP-1 and IL-6 protein concentrations in conditioned medium of OSM treated HUVECs, HAECs and HMEC-1 cells. Both MCP-1 (p<0.05) and IL-6 (<0.001) release were increased in OSM treated HUVECs, HAECs (p<0.05 and p<0.01 respectively) and HMEC-1 cells (p<0.001) at both time points (Fig 1G–1L). Subsequently, we measured adhesion molecule expression, which is another feature of endothelial activation. ICAM-1 mRNA expression was increased by OSM treatment in HUVECs (p<0.001) and HAECs (p<0.01) again at both 3h and 6h time points and in HMEC-1 cells 3h after addition of OSM (p<0.01)(Fig 2A–2C). VCAM-1 mRNA expression was upregulated in HUVECs at 3h (p = 0.008) and in HAECs at both 3h and 6h (p<0.001)(Fig 2D and 2E). Moreover, we observed an upregulation in E-selectin mRNA expression in both HUVECs and HAECs at both 3h and 6h (p<0.001 and p<0.05)(Fig 2F and 2G), while VCAM-1 and E-selectin mRNA
Inflammatory cytokine oncostatin M induces endothelial activation

**A**

HUVECs

Relative MCP-1 mRNA expression

Time (h)

**B**

HAECs

Relative MCP-1 mRNA expression

Time (h)

**C**

HMEC-1

Relative MCP-1 mRNA expression

Time (h)

**D**

HUVECs

Relative IL-8 mRNA expression

Time (h)

**E**

HAECs

Relative IL-8 mRNA expression

Time (h)

**F**

HMEC-1

Relative IL-8 mRNA expression

Time (h)

**G**

HUVECs

MCP-1 (pg/mL)

Control OSM Control OSM

3h 6h

**H**

HAECs

MCP-1 (pg/mL)

Control OSM Control OSM

3h 6h

**I**

HMEC-1

MCP-1 (pg/mL)

Control OSM Control OSM

3h 6h

**J**

HUVECs

IL-8 (pg/mL)

Control OSM Control OSM

3h 6h

**K**

HAECs

IL-8 (pg/mL)

Control OSM Control OSM

3h 6h

**L**

HMEC-1

IL-8 (pg/mL)

Control OSM Control OSM

3h 6h
levels were too low expressed in HMEC-1 cells. In addition, ICAM-1 membrane expression was increased in HUVECs (p < 0.05), HAECs (p < 0.05) and HMEC-1 cells (p < 0.05) (Fig 2H–2J), but not membrane expression of VCAM-1, P-selectin or E-selectin (S1 Fig). Since these adhesion molecules can also be shed upon endothelial activation[38], we measured P-selectin, E-selectin, soluble VCAM-1 and soluble ICAM-1 levels in conditioned medium. Soluble VCAM-1 was upregulated in conditioned medium of HUVECs 6h after OSM addition (p < 0.05) and in HAECs at both 3h and 6h post OSM addition (p < 0.01) (Fig 2K and 2L). Soluble VCAM-1 was not detectable in conditioned medium of HMEC-1 cells. Additionally, E-selectin levels were upregulated at both time points in conditioned medium of OSM treated HUVECs (p < 0.05) and HAECs (p < 0.01) and 6h post OSM addition in HMEC-1 cells (p < 0.05) (Fig 2M–2O). P-selectin levels were not detectable. Overall, these results indicate that OSM consistently induces endothelial activation in vitro in the different human endothelial cell types. Therefore, subsequent mechanistic studies were conducted in HUVECs.

3.2 JAK/STAT signaling is involved in OSM induced endothelial activation

IL-6 family members signal through the Janus kinase/signal transducers and activators of transcription (JAK/STAT) pathway, a pathway that is often involved in cytokine and growth factor signaling[39–41]. Therefore, we investigated whether this pathway is also involved in OSM induced endothelial activation. STAT1 and STAT3 phosphorylation were markedly increased (p < 0.05) (Fig 3) upon addition of OSM indicating that the JAK/STAT pathway is involved in OSM induced endothelial activation as well.

3.3 OSM induces endothelial activation by simultaneous signaling through the LIFR and OSMR

As OSM can signal through both the OSMR and the LIFR, a siRNA knockdown was performed to investigate which of these receptors is involved in OSM induced endothelial activation. LIFR mRNA expression was decreased to 25 ± 6% (mean ± SD), and OSMR mRNA expression to 52 ± 15%. Simultaneous knockdown resulted in a decrease of the LIFR to 31 ± 8% and of the OSMR to 45 ± 11% (Fig 4A and 4B). Single knockdown of LIFR did significantly decrease MCP-1 (p = 0.019) and IL-6 secretion (p = 0.005), but not VCAM-1 or E-selectin shedding. Single knockdown of OSMR did only decrease IL-6 secretion (p < 0.001), while MCP-1 secretion was significantly increased (p = 0.007). VCAM-1 and E-selectin shedding were both not significantly changed. Double knockdown did not only decrease IL-6 (p < 0.001) and MCP-1 (p < 0.001) secretion, but also VCAM-1 (p = 0.009) and E-selectin (p < 0.001) shedding compared to non-targeting siRNA treated cells (Fig 4C and 4D). A similar effect was observed for STAT1 and STAT3 phosphorylation, which was only reduced by double knockdown (p < 0.05) compared to control (Fig 4E and 4F). Altogether, these data indicate that OSM signals through LIFR and OSMR simultaneously in human endothelial cells.
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A | B | C

HUVeCs | HAECs | HMEC-1

Relative ICAM-1 mRNA expression

Time (h)

D | E | F

HUVeCs | HAECs | HMEC-1

Relative VCAM-1 mRNA expression

Time (h)

G | H | I

HAECs | HMEC-1

Relative E-selectin mRNA expression

Time (h)

J | K | L

ICAM-1

Relative cell number

M | N | O

HUVeCs | HAECs | HMEC-1

E-selectin (ng/mL)

Control OSM
0 3h 6h

Control OSM
0 3h 6h

Control OSM
0 3h 6h
3.4 OSM induces an inflammatory response in APOE<sup>3</sup>Leiden.CETP mice

To investigate whether OSM activates the endothelium in vivo as well, hyperlipidemic APOE<sup>3</sup>Leiden.CETP mice were administered OSM for 3 weeks. No clinical signs of deviant behavior and no significant effects on food intake were noted in any treatment group as compared to control. Plasma ALT and AST, measured at end-point as safety markers, showed no aberrant results (S2 Table). Also, no significant difference in body weight, triglyceride, or cholesterol levels were observed compared to control (Fig 5A–5C). As endothelial activation goes hand in hand with a pro-inflammatory response, plasma levels of inflammatory markers MCP-1 and E-selectin were measured. Plasma MCP-1 tended to be increased (p = 0.107) and plasma E-selectin was increased (p < 0.001) in mice treated with 10 μg/kg/day OSM compared to the control group (Fig 5D and 5E).

3.5 OSM induces endothelial activation in the vasculature of APOE<sup>3</sup>Leiden.CETP mice

To further investigate if OSM is able to induce endothelial activation, the aortic root area was examined for relevant markers. ICAM-1 protein expression tended to be elevated from 39 ± 15% (mean ± SD) to 59 ± 22% (p = 0.067) and an increase in monocyte adhesion to the activated endothelium was observed from 5.7 ± 3.0 to 10.3 ± 4.7 monocytes (mean ± SD, p < 0.05) in mice treated with 10μg/kg/day OSM (Fig 6). Furthermore, aortic mRNA expression analysis revealed a dose-dependent increase in Il-6 expression (p < 0.001) and Icam-1 expression tended to be increased in the 1μg/kg/day and 10μg/kg/day OSM treated groups (p = 0.101 and p = 0.133 respectively) compared to control. Mcp-1 mRNA expression was not enhanced (Fig 7). These results show that OSM does not only induce endothelial activation in vitro, but also in vivo in a hyperlipidemic mouse model.

4. Discussion

The present study demonstrates that OSM induces endothelial activation in cultured human endothelial cells as well as in vivo in APOE<sup>3</sup>Leiden.CETP mice. The data show increased release of inflammatory markers and adhesion molecule expression, both features of endothelial activation. Furthermore, OSM increased monocyte adhesion in the aortic root area, as functional marker of endothelial activation.

We studied OSM induced endothelial activation in vitro by investigating the effect of OSM in three different types of human endothelial cells. Our data add to and expand on previous data that showed that OSM increases IL-6, IL-8 and MCP-1 secretion, ICAM-1 and VCAM-1 membrane expression and PMN adhesion to endothelial cells in vitro [34,42,43]. Consistently, increased VCAM-1 and E-selectin shedding was observed in all three endothelial cell types. ICAM-1 is an important adhesion molecule in monocyte binding as ICAM-1<sup>−/−</sup> endothelial...
cells show a strong attenuation in monocyte binding compared to control endothelial cells [44]. Although we did not observe an increase in membrane E-selectin and VCAM-1 expression, OSM did increase soluble E-selectin and VCAM-1. Soluble VCAM-1 was previously shown to serve as a monocyte chemoattractant agent and soluble E-selectin enhances leukocyte migration and binding to endothelial cells[45,46]. Taken together, these observations show that OSM induces different biomarkers of endothelial activation in cultured endothelial cells.

Fig 3. JAK/STAT pathway is involved in OSM induced endothelial activation. HUVECs were incubated with 5 ng/mL OSM for 15 or 30 min. A representative picture shows STAT1 phosphorylation at Tyr701, STAT1, STAT3 phosphorylation at Tyr705, STAT3 and GAPDH (A). Bar graphs show relative STAT1 and STAT3 phosphorylation (B,C). A two-way ANOVA was performed to test for significance. Data represent mean ± SD (n = 5). *p<0.05 **p<0.01 ***p<0.001 compared to control.

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**A** LIFR mRNA

**B** OSMR mRNA

**C** MCP-1

**D** IL-6

**E** VCAM-1

**F** E-selectin

**G** Western blot analysis of pSTAT1, STAT1, and GAPDH

**H** Western blot analysis of pSTAT3, STAT3, and GAPDH

**I** PhosphoSTAT1/STAT1 ratio

**J** PhosphoSTAT3/STAT3 ratio
Previous short term in vivo studies in healthy wildtype mice with OSM administered for only 6 to 24 hours have shown signs of acute endothelial activation, such as increased angiotropin 2 expression in cardiac tissue, increased plasma VEGF levels and increased permeability and infiltration of inflammatory cells\[33–36\]. It is important to note that publicly available datasets show that Osmr and Lifr mRNA are expressed in aortic endothelial cells from mice as well (data accessible at NCBI GEO database [47], accession GSE114805 and [48], accession GSE115618).

The aim of the present study was to investigate the effect of chronic OSM exposure on endothelial activation in a hyperlipidemic mouse model, the APOE\(^+\)3Leiden.CETP mouse. This mouse model features elevated lipid levels, representing humans with hyperlipidemia and mild chronic inflammation who have an increased risk of developing atherosclerosis.
We found that OSM tended to increase plasma MCP-1 and significantly increased plasma E-selectin, both markers of activated or dysfunctional endothelium\[^{3,49}\] after 3 weeks of chronic OSM administration. Moreover, mRNA expression of \( \text{Il-6} \) was increased dose-dependently in aortic tissue of OSM treated mice. We also observed a trend towards increased ICAM-1 expression in the aortic root of OSM treated mice and a markedly enhanced monocyte binding as functional marker of activated endothelium, thus demonstrating augmented endothelial activation. ICAM-1 expression and adhesion of monocytes are strongly related, as previous studies show increased monocyte binding upon enhanced ICAM-1 expression and decreased monocyte binding upon reduced ICAM-1 expression\[^{44,50}\]. Collectively, these findings provide evidence that OSM does not only induce endothelial activation \textit{in vitro}, but

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**Fig 6.** OSM increases ICAM-1 expression and monocyte adherence in the aortic root area in OSM treated APOE\(^{\text{3Leiden.CETP}}\). Representative pictures showing the endothelial ICAM-1 expression (brown staining) in a control (A) and a 10 \( \mu \text{g/kg/day} \) OSM treated (B) mouse and monocyte adherence (arrows) in a control (C) and a 10 \( \mu \text{g/kg/day} \) OSM treated (D) mouse. Endothelial ICAM-1 expression was determined as percentage of the endothelial surface in the cross sections (E) and adhering monocytes were counted per cross-section after staining with AIA 31240 (F). Data represent mean \( \pm \) SD (n = 7–8). An independent t-test was used to test for significance. Data represent mean \( \pm \) SD. \(^*\) \( p<0.05 \).

[31,32,37].
also in vivo on top of the inflammatory state that is present in hyperlipidemic mice, resulting in increased monocyte recruitment and adherence.

Even though, endothelial cells are directly activated by OSM in vitro, it is important to note that the in vivo situation is much more complex and other cell types may have contributed to the observed effects as well. For instance, the increase in plasma MCP-1 and cardiac Il-6 expression can partly be caused by fibroblasts or smooth muscle cells, as these two cell types

Fig 7. OSM increases Il-6 mRNA expression in aortic tissue of APOE-3Leiden.CETP mice treated with OSM. After 3 weeks of OSM treatment, mRNA was isolated from the aorta and analyzed by qPCR. Il-6 (A), Mcp-1 (B) and Icam-1 (C) mRNA expression were quantified. All values are relative values compared to the control group, which was given an arbitrary value of 1. Values were normalized to HPRT and calculated with the ΔΔCt method. A one-way ANOVA with Dunnett’s test was performed on the ΔCt values to test for significance. Data represent geometric mean ± geometric SD (n = 6–8). All values were compared to control. *p<0.05 **p<0.001 compared to control, ns = not significant.

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also show increased IL-6 and MCP-1 expression upon OSM treatment \textit{in vitro}[51,52]. Furthermore, OSM can promote growth factor and cytokine release in cell types other than endothelial cells, these released growth factors and cytokines can in turn activate the endothelium, thereby inducing indirect endothelial activation[51–53]. An example of such a growth factor is vascular endothelial growth factor (VEGF), which can be upregulated by OSM in multiple cell types[35,54–56] and is known to induce endothelial activation by increasing adhesion molecule expression and leukocyte adhesion[57].

Although our \textit{in vivo} study was not aimed at and was too short to investigate whether chronic OSM exposure aggravates atherosclerosis, our results do give clues that OSM may be involved in the initiation of the atherosclerotic process. Some of the diverse hallmarks of endothelial activation that we observed, have previously been associated with atherosclerosis development in humans[49,58]. Further indications come from reports showing that OSM is present in both murine and human plaques[16], and higher mRNA expression levels of OSM in PBMCs derived from coronary artery disease patients compared to healthy individuals[59]. Moreover, a recent paper showed that prevention of OSM signaling, as opposed to stimulation of OSM signaling in our study, in OSMR-β/−ApoE−/− mice resulted in less and smaller atherosclerotic lesions and less macrophages compared to ApoE−/− mice[60].

Other studies have shown that partial inhibition of endothelial activation by knockdown of E-selectin, P-selectin, ICAM-1 or MCP-1 attenuates atherosclerosis development in mice [61,62]. Therefore, lowering of plasma OSM levels or intervention in OSM signaling might be worth investigating as a possible future approach in the treatment of atherosclerosis.

As it is currently unknown which of the OSM receptors is involved in OSM induced endothelial activation, we performed a siRNA knockdown of the LIFR and the OSMR. Single knockdown experiments showed that solely LIFR or OSMR downregulation is not sufficient to prevent OSM induced endothelial activation or JAK/STAT signaling. Only simultaneous knockdown of both receptors was able to dramatically decrease IL-6 and MCP-1 release, VCAM-1 and E-selectin shedding and STAT1 and STAT3 phosphorylation. Hence, it is essential to block both receptors simultaneously or to target OSM when considering intervening in OSM signaling as a possible future therapy. Targeting both receptors or OSM itself could be a relative safe approach since OSM−/− mice are viable and healthy[63].

Taken together, our comprehensive study provides new evidence that OSM induces activation of human endothelial cells from different vascular beds and in APOE 3Leiden.CETP mice chronically treated with OSM. Moreover, we provided data indicating both receptors for OSM as well as OSM itself as potential therapeutic targets in atherosclerosis and other chronic inflammatory diseases in which endothelial activation is involved such as rheumatoid arthritis, abnormal angiogenesis and thrombosis[64–67].

**Supporting information**

**S1 Fig. OSM does not affect membrane expression of VCAM-1, E-selectin and P-selectin in endothelial cells.** HUVECs, HAECs and HMEC-1 cells were incubated with 5 ng/mL OSM for 18h. A two-way ANOVA with Dunnett’s test was performed on the median to test for significance (n = 3). ns = not significant.

(TIF)

**S1 Table. List of antibodies used for flow cytometry, immunohistochemistry and western blot.**

(PDF)

**S2 Table. Average food intake and ALT and AST levels in mice.**

(PDF)
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References


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