Parallel regulation of thyroid hormone transporters OATP1c1 and MCT8 during and after endotoxemia at the blood-brain barrier of male rodents

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Abstract

There is increasing evidence that local thyroid hormone (TH) availability changes profoundly in inflammatory conditions due to altered expression of deiodinases that metabolize TH. It is largely unknown, however, how inflammation affects TH availability via the expression of TH transporters. In this study we examined the effect of bacterial lipopolysaccharide (LPS) administration on two TH transporters that are critically important for brain TH homeostasis, organic anion-transporting polypeptide 1c1 (OATP1c1) and monocarboxylate transporter 8 (MCT8). Messenger RNA levels were studied by in situ hybridization and quantitative PCR, and protein levels by immunofluorescence in both the rat and mouse forebrain. The mRNA of both transporters decreased robustly in the first 9h after LPS injection, selectively in brain blood vessels; OATP1c1 mRNA in astrocytes and MCT8 mRNA in neurons remained unchanged. At 24 and/or 48h after LPS administration, OATP1c1 and MCT8 mRNAs increased markedly above control levels in brain vessels. OATP1c1 protein decreased markedly in vessels by 24h, whereas MCT8 protein levels did not decrease significantly. These changes were highly similar in mice and rats. The data demonstrate that OATP1c1 and MCT8 expression are regulated in a parallel manner during inflammation at the blood-brain barrier of rodents. Given the indispensable role of both transporters in allowing TH access to the brain, the results suggest reduced brain TH uptake during systemic inflammation.
Introduction

It is well known that thyroid hormone (TH) homeostasis is altered in association with acute and chronic illnesses, a disorder commonly referred to as the nonthyroidal illness syndrome and manifested by reduced serum TH levels and an inappropriately low TSH (1-3). However, there is limited understanding about how TH availability changes at the level of specific target cells and tissues under these conditions. Local TH availability is largely determined by the presence of TH transporters that enable the passage of TH across cellular membranes, and deiodinases that metabolize TH (4, 5). Deiodinases have been studied extensively in animal models of inflammatory conditions (1), particularly type 2 deiodinase (D2) that activates T4 to the biologically more potent T3 (4). In these models, D2 expression increases in several organs (6-15), including novel expression in cells and tissues where it is not normally present (6, 8, 15), suggesting a local anti-inflammatory role for TH (6, 8, 16-18). In contrast, it is largely unknown how inflammation affects TH availability via TH transporters. Quantitative PCR studies reported decreased monocarboxylate transporter 8 (MCT8, coded by Slc16a2) expression in several organs of septic pigs (19) and in the subcutaneous fat of septic patients (20), while organic anion-transporting polypeptide 1c1 (OATP1c1, coded by Slco1c1) and monocarboxylate transporter 10 (MCT10) mRNAs increased in the hypothalamus of rabbits in prolonged critical illness (21). Details of these responses, such as cell-type specificity, temporal characteristics, and whether mRNA changes translate into protein levels, have not yet been studied. Shedding light on these aspects of TH handling would be important to better understand TH availability during disease at the cellular level.

Previously, we described how inflammation affects D2 expression in the brain of different rodent species, and demonstrated inflammation-induced D2 expression in the leptomeninges, choroid plexus and a subset of brain blood vessels (15). The present study was conducted to obtain a detailed, cell-type-specific insight into the inflammatory regulation of TH transporters. Of the several proteins capable of TH transport and present in the rodent brain (22, 23), we focused on OATP1c1 and MCT8 because they are critically important for brain TH homeostasis (24). Both transporters facilitate TH traffic in multiple cell
types (25-30), and are indispensable for TH entry into the brain via the blood-brain and/or blood-cerebrospinal fluid barrier (24, 31-34). In fact, almost the entire T3 uptake into the brain is facilitated by MCT8 (32), while both MCT8 and OATP1c1 contribute to T4 uptake (24). Importantly, a recent study by Mayerl et al. (24) demonstrated that the lack of both transporters results in a severely hypothyroid brain, with T3 and T4 contents being of only 10% of wild type levels.

In this study, we examined the effect of bacterial lipopolysaccharide (LPS) administration, an acute systemic inflammatory challenge, on OATP1c1 and MCT8 mRNA and protein expression in the brain using in situ hybridization, quantitative PCR, and immunofluorescence. Since we previously found major differences in the LPS-induced D2 expression between rats and mice (15), we performed the experiments in both species, with special attention to the rat meninges.

Materials and Methods

Animals

The experiments were carried out on adult, male, Sprague-Dawley (Taconic Farms) and Wistar rats (TOXI-COOP KKT, Hungary) weighing 220-260g, and adult, male C57Bl/6 mice (Taconic and Charles River), weighing 19-21 g. Animals were housed under standard conditions (lights on between 0600 and 1800 h, temperature 22 ± 1 C, rodent chow and water ad libitum). All experimental protocols were reviewed and approved by the Institutional Animal Care and Use Committee at Tufts Medical Center and the Animal Welfare Committee at the Institute of Experimental Medicine of the Hungarian Academy of Sciences.

Injections and tissue preparation

LPS-injections: Rats and mice were injected ip with LPS from Escherichia coli (serotype O55:B5; Sigma-Aldrich Co), dissolved in saline, at a dose of 2.5 mg/kg body weight to all animals according to the Guide of the American Thyroid Association (35). Control animals received the same volume of saline.
Experiment for quantitative PCR analysis: Groups of Wistar rats (n=6 per group) and C57Bl/6 mice (n=8 per group) were injected with LPS or saline, and were decapitated 9 or 48h later. The brains were dissected and samples from the cerebral cortex collected and stored at −80°C until subjected to real-time PCR. In another experiment, leptomeninges of the basal forebrain were collected from 6 control and 8 LPS-treated Sprague-Dawley rats 9h after the injections.

Time course experiment for in situ hybridization and immunofluorescence studies: Groups of Sprague-Dawley rats and C57Bl/6 mice (n=4 or 5 in each group) were injected with LPS and 2, 4, 9, 24 or 48h later were anesthetized with ketamine-xylazine, then decapitated. Control mice (n=5) were euthanized at 9h, and control rats were euthanized at 9h (n=3) or 24h (n=2), as in situ hybridization and immunofluorescent signals for OATP1c1 and MCT8 were of the same intensity in 9h and 24h control rat brains. The brains were removed, snap-frozen on powdered dry ice and 16 μm thick coronal sections were cut using a Leica CM3050 S cryostat (Leica Microsystems). Sections were thaw-mounted on Superfrost Plus slides (Fisher Scientific Co), air-dried and stored at -80°C until processed for in situ hybridization.

RNA isolation and Real time PCR analyses
RNA was isolated using the RNeasy Lipid Tissue Mini Kit (Qiagen) from cortex samples and with RNeasy Tissue Mini Kit (Qiagen) from leptomeninges according to the manufacturer’s instructions. The purity and concentration of the RNA were analyzed using Spectrophotometer (Bio-Rad, SmartSpec Plus). Reverse transcription was performed with 1 μg of RNA to convert the total RNA to cDNA using the High Capacity cDNA Reverse Transcriptase Kit (Applied Biosystems by Life Technologies). Concentration of the generated cDNA was determined using the Qubit 2.0 Fluorometer with the Qubit ssDNA Assay Kit (Life Technologies). Expression of OATP1c1, MCT8 and D2 was measured by real-time quantitative TaqMan RT PCR reaction with a ViiA 7 Real-Time PCR System (Life Technologies), using commercially available TaqMan probes (code numbers are summarized in Supplementary Table 1.) on 10
ng cDNA template in duplicates. Glyceraldehyde-3-phosphate dehydrogenase (Gapdh) was used as housekeeping gene. Expression of the housekeeping gene did not vary between the experimental groups.

*Generation of hybridization probes*

Template cDNA fragments were generated with RT-PCR using standard procedures. Amplification was performed on cDNA synthesized from rat brain for OATP1c1 and mouse or rat liver for MCT8. Fragments were cloned into pGemT vector (Promega) and confirmed by sequencing. Probe sequences were as follows: rat OATP1c1 probe corresponds to nt 375-2525 of NM_053441.1 and has 93% homology with the corresponding mouse sequence, nt 368-2522 of NM_021471.2; mouse MCT8 probe nt 2569-3428 of NM_009197.2; rat MCT8 probe nt 41-648 of NM_147216.1, and has 98% homology with the mouse sequence NM_009197.2 between nt 227-834; rat D2 probe was reported previously (15). Antisense riboprobes were synthesized using SP6 or T7 RNA polymerase (Promega) in the presence of [35S]-uridine 5′-(alpha-thio) triphosphate (PerkinElmer), and purified with Mini Quick Spin RNA columns (Roche Applied Sciences). For fluorescent *in situ* hybridization, the OATP1c1 probe was labeled with digoxigenin-11-UTP (Roche).

*Isotopic in situ hybridization*

Isotopic *in situ* hybridization was performed as previously described (15, 36) using 50,000 cpm/μl radiolabeled probe concentrations. The same probe was used to detect OATP1c1 mRNA in both species; the rat MCT8 probe was used for rat sections, while both the rat and mouse MCT8 probes that are non-overlapping were used on mouse sections to increase hybridization sensitivity. Following stringency washes, sections were dehydrated in ascending ethanol series, air-dried, and placed on Amersham Hyperfilm autoradiography film (GE Healthcare Biosciences) for 6 days (rat OATP1c1) or 8 days (mouse OATP1c1 and MCT8). Slides were then dipped in Kodak NTB autoradiography emulsion (Carestream Health Inc), and stored at 4°C until developed. Exposure times were as follows: 9 days for mouse
OATP1c1; 24 days for mouse MCT8; two exposure times, 5 days and 9 days, were used for two different hybridizations for rat OATP1c1; 35 days for rat MCT8; and 14 days for rat D2. Autoradiograms were developed with Kodak D19 developer (Eastman Kodak Co). Sections were immersed in 0.0005% cresyl violet acetate (Sigma-Aldrich) for 2 min to obtain fluorescent labeling of cell nuclei, dehydrated in ascending ethanol series and xylenes, and coverslipped with DPX (Sigma-Aldrich). Darkfield images of the emulsion autoradiographs and fluorescent images of the cresyl violet signal were captured using a Zeiss Axioplan 2 microscope equipped with a SPOT Slider digital camera (Diagnostic Instruments).

*Isotopic OATP1c1 in situ hybridization combined with GFAP immunofluorescence*

Sections from the control and 9h LPS groups from both species were hybridized for OATP1c1 as above, then processed further for immunofluorescence. The sections were treated with the mixture of 0.5% Triton X-100 and 0.5% H2O2 for 15 min, rinsed in PBS (3×10 min), immersed in maleate buffer (pH 7.5) for 10 min, and then in 1% blocking reagent for nucleic acid hybridization (Roche). The sections were incubated overnight in a mouse monoclonal antibody against the astrocyte marker, glial fibrillary acidic protein (GFAP) (Cat# MAB360, Millipore, diluted 1:1,000), and subsequently in Alexa Fluor 488-conjugated donkey anti-mouse IgG (1:400; Life Technologies) for 2 hours. Sections were dehydrated and dipped in Kodak NTB autoradiography emulsion. The autoradiograms were developed after 9 days. The fluorescent signal of Alexa Fluor 488 was pseudo-colored to red for better visibility of dual-labeled cells.

*Fluorescent OATP1c1 in situ hybridization combined with GFAP immunofluorescence*

Hybridizations were performed on both fresh-frozen control rat sections, and sections from four normal paraformaldehyde-perfused rats. Paraformaldehyde-perfusion was performed as previously described (36). Briefly, rats were anesthetized with an overdose of pentobarbital (50 mg/kg) and perfused transcardially with PBS followed by 4% paraformaldehyde. The brains were postfixed by immersion in 4% paraformaldehyde, cryoprotected in 20% sucrose in PBS overnight, then snap-frozen on dry ice. Serial 20 μm coronal sections were cut on a cryostat, collected in cryoprotectant solution and stored at -
20°C until used. Fluorescent in situ hybridization for fresh-frozen and paraformaldehyde-perfused sections was performed as previously described (15, 36). The digoxigenin-labeled probe was detected with peroxidase-conjugated Fab fragments of sheep anti-digoxigenin antibody (1:100, Roche). The hybridization signal was amplified with biotinylated tyramide for 30 min using the TSA amplification kit (Perkin Elmer), and visualized by Alexa 488-conjugated Streptavidin (1:500; Life Technologies). Sections were reacted with murine GFAP antibody as above, and detected with Cy3-conjugated donkey anti-mouse IgG (Jackson ImmunoResearch). Sections were then coverslipped with Vectashield containing 4′,6-diamidino-2-phenylindole (DAPI) (Vector Laboratories).

**Immunofluorescence for OATP1c1 and MCT8**

For MCT8 immunofluorescence, mounted fresh-frozen sections were fixed with 4% paraformaldehyde in 0.1M phosphate buffer (pH 7.4) for 15 min. For OATP1c1 immunofluorescence, sections were fixed with methanol at −20°C for 5 min. Sections were permeabilized with 0.25% Triton-X-100 for 30 min, blocked with 2% normal horse serum in PBS, and incubated overnight in a rabbit antiserum against human MCT8 (raised against amino acids 527-539; #1306, gift from Dr. Theo J. Visser) or a rabbit antiserum against human OATP1c1 (raised against amino acids 697-712; #3516, gift from Dr. Theo J. Visser), both diluted at 1:1,000. The primary antisera were detected with Alexa Fluor 488-conjugated donkey anti-rabbit IgG diluted at 1:400 (Jackson ImmunoResearch). Immunostaining with both antisera resulted in the same patterns reported earlier (26, 37).

**Image analysis**

In situ hybridization signals were compared between sections from a single hybridization experiment where all conditions and exposure times were identical. Darkfield images of OATP1c1 in situ hybridization were analyzed with ImageJ software (public domain at [http://rsb.info.nih.gov/ij](http://rsb.info.nih.gov/ij)). To maximize the analysis of OATP1c1 signal in blood vessels, hybridizations were analyzed in the hypothalamus where no significant signal was detected in astrocytes after 9 days exposure in mice or 5
days exposure in rats. Images were taken with a 5× objective at the level of the paraventricular nucleus (mice) or the dorsomedial nucleus (rats), and integrated density values were calculated. From the two rat OATP1c1 hybridizations, only the 5 day exposure experiment was used for quantitative image analysis. Images from both the 9 and 5 days exposure experiments were used to illustrate the magnitude of changes in OATP1c1 hybridization signal. To quantify vascular MCT8 hybridization in mice, blood vessel segments with clear hybridization signal were counted in the thalamus and hypothalamus in 3 sections per mouse at the rostro-caudal level of the median eminence. In these regions, the low neuronal hybridization signal allowed easier identification of labeled blood vessels than in cortical regions. OATP1c1 immunofluorescence in blood vessels was quantified in rats using ImageJ. Images were taken with a 20× objective of an area with high density of blood vessels in the midline thalamus. The area covered by immunofluorescently labeled vessels (pixel number) was measured by subtracting non-labeled areas (i.e., dark pixels) from the image using the threshold tool and the same threshold value for all images. Hematoxylin and eosin staining was used to facilitate the identification of arachnoid veins in rats.

Statistical analysis

Data are presented as mean ± SEM. Quantitative PCR data were expressed as RQ (mean +/- SEM). Quantitative PCR data from cortex and image analysis data were compared between groups by one-way ANOVA and Newman-Keuls multiple comparison post-hoc test. Quantitative PCR data from the leptomeninges were compared by Student’s t test.

Results

Effect of LPS on OATP1c1 expression in the brain

OATP1c1 mRNA. LPS administration similarly affected OATP1c1 expression in mice and rats. OATP1c1 mRNA levels decreased profoundly within 9h after LPS injection, as demonstrated by a striking reduction in hybridization signal across the forebrain (Fig. 1A, 1B). Emulsion autoradiography revealed that this reduction was due to a loss of hybridization signal specifically in blood vessels, but not
astrocytes. Vascular OATP1c1 expression decreased uniformly in the forebrain, but the course of this response could be followed most clearly in the hypothalamus (Fig. 2A) where hybridization signal in astrocytes was the lowest among forebrain regions. Hybridization signal in vessels decreased significantly as soon as 2h after LPS, and markedly by 4h (Fig. 2A). Vascular hybridization signal was virtually absent in mice 9h after LPS injection, except light labeling of the highly vascularized hypothalamic paraventricular nucleus (Fig. 2A). A similar reduction was observed in rats, where only scattered blood vessels with light hybridization signal were detected at 9h after LPS (in the hybridization experiment with the longer, 9 day autoradiography exposure). Moderate labeling remained only in some vessels of the paraventricular and supraoptic nuclei (Fig. 3A; also in Fig 1A); in fact, vessels in these two nuclei were more intensely labeled than elsewhere, even in control rats. The loss of OATP1c1 hybridization signal in blood vessels is also demonstrated in images from the cerebral cortex in Figs. 2B and 3A.

Hybridization signal in astrocytes did not change noticeably in either species, and remained comparable to control levels 9h after LPS, as demonstrated in images from the mouse and rat cortex (Fig. 2B, 3A). Notably, OATP1c1 mRNA in astrocytes varied greatly between brain regions, confirming recent studies (23, 27). In mice, OATP1c1 mRNA was easily detected in hippocampal and cortical astrocytes, while it was not detected in hypothalamic astrocytes with the exposure time used in this experiment. In rats, differences in astrocytic OATP1c1 expression were confirmed by both fluorescent (Fig. 3B) and radioactive in situ hybridization. Hybridization signal was moderate to intense in astrocytes of the cortex, hippocampus, striatum, and ventrolateral thalamus, whereas it was much lighter in astrocytes of the hypothalamus, such as in the ventromedial nucleus (Fig. 3B).

In both species, vascular OATP1c1 hybridization signal increased above control levels during the recovery phase from endotoxemia. In mice, OATP1c1 mRNA was still below control levels at 24h, but increased markedly at 48h after LPS (Fig. 2A). In rats, upregulation of OATP1c1 mRNA in blood vessels could be observed at both 24h and 48h after LPS (Fig. 3A). Conspicuously increased signal in vessels was observed in one out of four rats in the 24h LPS group, and three out of four rats in the 48h
LPS group. Markedly increased OATP1c1 mRNA at 24h after LPS was confirmed in another experiment (2 out of 2 rats; data not shown). This indicates that the timing of post-endotoxemic increase in vascular OATP1c1 mRNA can vary among individual rats. Image analysis results from mice and rats representing vascular OATP1c1 mRNA expression at each time point following LPS injection are presented in Fig. 4A. Quantitative RT-PCR data from cortical samples confirmed the in situ hybridization results, as OATP1c1 mRNA was significantly decreased 9h after LPS injection in both species (Fig. 4B). At 48h after LPS, OATP1c1 mRNA levels were not different from control levels by quantitative PCR (Fig. 4B).

In the choroid plexus, OATP1c1 hybridization signal was consistently reduced 9h after LPS in mice (Fig. 1B), but not rats. In tanycytes, OATP1c1 mRNA expression was extremely intense in rats (Supplemental Fig. 1), but conspicuously absent in mice (Supplemental Fig. 2). LPS did not affect OATP1c1 expression in rat tanycytes.

**OATP1c1 protein.** Immunofluorescence for OATP1c1 gave clear labeling of blood vessels in the rat brain with low background, whereas in mice the labeling was less sensitive and less clear due to higher background levels. The antiserum did not label astrocytes. Immunostaining of blood vessels did not differ between control, 2h and 4h LPS groups, but visibly decreased at 9h and 24h (5A, 5B). At 24h, when the decrease was significant by image analysis (Fig. 4A), only scattered major vessel segments were labeled in both species (Fig. 5A, 5B). OATP1c1 immunostaining in vessels at 48h after LPS was similar to controls (Fig. 5A, 5B). OATP1c1 immunostaining was intense in rat tanycytes (Supplemental Fig. 1), whereas in mice it was observed selectively in lateral beta (beta-1) tanycytes in the floor of the third ventricle (Supplemental Fig. 2). The presence of OATP1c1 immunoreactivity in mouse tanycytes is in agreement with a previous report by Roberts et al. (26) that used highly specific antibodies.

**Effect of LPS on MCT8 expression in the brain**

**MCT8 mRNA.** LPS similarly affected MCT8 expression in mice and rats. The hybridization patterns in control brains were identical to what was previously described (25). Several neuronal populations were labeled with variable, light to intense signal, and scattered larger blood vessels were labeled with
moderate or light signal (Fig. 1C). Rat tanycytes were intensely labeled, and extremely intense signal labeled mouse tanycytes and the choroid plexus in both species (Fig. 1C). Neuronal MCT8 expression did not noticeably change at any time after LPS injection in both species (Fig. 1C). In contrast, MCT8 mRNA expression in blood vessels changed markedly in response to LPS. In mice, the intensity of vascular labeling decreased gradually from 2h until 9h, when hybridization signal was no longer detected in vessels (Fig. 6A). A similar trend was observed in rats, with noticeably less numerous and less intensely labeled blood vessels at 4h, and only occasional, lightly labeled vessels visible at 9h (Fig. 6B). In mice, both the number of detected vessel segments and their signal intensity increased at 24h after LPS compared to control mice, and even more strikingly at 48h (Fig. 1C, 6A). The quantification of labeled vessels in mice is presented in Fig. 4A. In rats, labeling of blood vessels was comparable to control levels at 24h, but increased markedly by 48h after LPS (Fig. 6B). Quantitative PCR from the mouse cortex did not detect significant changes in MCT8 mRNA after LPS (Fig. 4B). In the rat cortex, MCT8 mRNA tended to decrease at 9h albeit not significantly, but significantly increased 48h after LPS (Fig. 4B).

MCT8 protein. MCT8 immunolabeling in blood vessels was not different from control levels at 2, 4, 9 or 48h after LPS in both species. Only a slight decrease in labeling was observed at 24h in mice (Fig. 5C). Intense MCT8 immunofluorescence was observed in axons in both species, in accordance with our previous description of specific axonal MCT8 labeling (37). MCT8-containing axons were present in most brain regions, but unaffected by LPS.

Effect of LPS on OATP1c1, MCT8 and D2 mRNAs in the rat leptomeninges

Since we previously observed that LPS administration induces robust D2 expression in the leptomeninges of rats (15), we also studied the expression of TH transporters in this tissue. OATP1c1 mRNA-expressing cells were observed in arachnoid veins, but not arteries or in the arachnoid tissue itself (Fig. 7A). Following LPS, OATP1c1 hybridization signal could no longer be detected in the wall of the veins at 2, 4 or 9h later (Fig. 7A), but returned to control levels by 24h, and remained normal at 48h. Conversely, D2 mRNA was induced in the wall of the arachnoid veins but not arteries following LPS (Fig
which we did not previously recognize (15). Moderate to intense MCT8 hybridization signal was observed in the leptomeningeal layers between the hippocampus and thalamus (Fig. 7A), and the dorsal third ventricle adjacent to the choroid plexus. This signal did not change significantly following LPS administration. MCT8 hybridization signal in the arachnoid covering the outer brain surface was uncertain, as control hybridizations with the sense transcript also yielded labeling in this location, although less intensely than the antisense probe. By quantitative PCR from leptomeninges removed from the basal forebrain, OATP1c1 mRNA was reduced to approximately 10% of control level 9h after LPS injection, MCT8 mRNA did not change, while D2 mRNA increased 86-fold on average (Fig. 7B).

Discussion

While data have been accumulating on cell-type specific regulation of TH action in the brain, regulation of TH transport is poorly understood. In the present study, regulation of TH transporters was studied in the forebrain using a model of non-thyroidal illness syndrome, a condition known to evoke marked changes in tissue TH availability (1, 2). In particular, we report changes in OATP1c1 and MCT8 expression in the rodent brain in response to LPS administration, with the following main findings: 1) OATP1c and MCT8 mRNAs decrease rapidly and robustly in blood vessels; 2) this effect is cell-type specific as OATP1c1 and MCT8 mRNA levels remain unaltered in astrocytes and neurons; 3) a robust decrease in OATP1c1 but not MCT8 protein levels in vessels follows hours later; 4) during recovery from endotoxemia, OATP1c1 and MCT8 mRNA levels increase markedly in blood vessels above control levels; 5) these changes occur in both mice and rats. In addition, we describe OATP1c1 expression in arachnoid veins but not arteries that respond to LPS in a similar way as parenchymal blood vessels, and MCT8 mRNA in part of the rat leptomeninges.

MCT8 and OATP1c1 have complementary functions in the blood-brain barrier, and work in tandem to allow TH access to the brain (24). The parallel changes in OATP1c1 and MCT8 mRNAs in brain vessels during endotoxemia further corroborates the functional link between these transporters, and suggest a common mechanism regulating the expression of these genes during inflammation. The mechanisms by
which rapid downregulation of both transporter mRNAs occurs is unknown, but could be a direct effect of cytokines or other associated inflammatory signals that are associated with endotoxin administration. However, this response was specific to cells comprising the brain vasculature, such as endothelial cells known to express these transporters (26, 34, 38) and possibly also pericytes, while OATP1c1 and MCT8 mRNA levels remained unchanged in astrocytes and neurons. One possibility to explain the latter phenomenon is that the extracellular signal that down-regulates MCT8 and OATP1c1 expression in vessels has limited access to astrocytes and neurons inside the blood-brain barrier. Alternatively, the signal may be present in the brain parenchyma, but due to intrinsic properties of astrocytes and neurons, has no effect on OATP1c1 or MCT8 mRNA. The latter might be explained by the lack of receptors for the signal molecule, or that the signaling pathway activated is not coupled to the mechanism that changes the transcription of the transporter genes.

The marked increase in OATP1c1 and MCT8 mRNAs during the recovery period from endotoxemia may be secondary to low intracellular TH levels in microvascular cells, supported by data that OATP1c1 in rat brain microvessels is upregulated by hypothyroidism (28). However, regulation of OATP1c1 and MCT8 mRNAs by TH was not observed in congenitally hypothyroid mice (25), raising the possibility that increased OATP1c1 and MCT8 mRNA expression may be a rebound effect independent of TH concentration.

At the protein level, OATP1c1 decreased in brain vessels in a similarly robust manner as OATP1c1 mRNA, but only several hours later. This delay likely reflects the turnover rate of OATP1c1 protein in cells of the blood-brain barrier. In contrast, only a modest decrease in vascular MCT8 immunostaining was observed in mice. A possible explanation is that the half-life of MCT8 may be substantially longer than that of OATP1c1, and in our model, the inflammatory state may have been too transitory to see a significant decrease in the MCT8 protein. Thus, an inflammation model of longer duration may be necessary to more definitively address whether MCT8 decreases similar to OATP1c1. Since our analysis did not provide information on the subcellular location of the transporters, it is also conceivable that

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internalization of OATP1c1 and/or MCT8 from the cell membrane could result in a more rapid decrease in TH transport before a decrease in protein levels is realized.

Access of TH to most of the brain parenchyma primarily occurs via the blood-brain barrier, and much less via the blood-cerebrospinal fluid barrier (39). Since deficiency in either OATP1c1 or MCT8 is sufficient to reduce brain TH uptake (24, 32), downregulation of these transporters in brain vessels during inflammation suggests diminished TH uptake into the brain parenchyma and may contribute to common symptoms associated with illness such as fatigue, depressed mood, and impaired neurocognitive function.

In support, a recent clinical study reported that certain OATP1c1 polymorphisms are associated with fatigue and depression in a population of patients with adequately treated hypothyroidism (40).

Inflammatory regulation of OATP1c1 and MCT8 was strikingly different from that observed for D2 that is highly inducible by the inflammatory transcription factor, nuclear factor-kappa B (NF-κB) (11, 41). LPS induces D2 via NF-κB in tanycytes (7, 15), and probably in meningeal fibroblasts (15). In contrast, OATP1c1 and MCT8 expression did not increase following LPS in any cell type in the brain, and markedly decreased in cells of brain blood vessels. The opposite regulation of D2 and OATP1c1/MCT8 by inflammation raises the possibility that during inflammation, TH availability may not be uniform throughout the brain. The downregulation of OATP1c1 and MCT8 at the blood-brain barrier suggests decreased TH availability within the brain parenchyma, whereas induction of D2 suggests increased TH availability in the leptomeninges and choroid plexus (15). Therefore, inflammation may result in both “hypothyroid” and “hyperthyroid” compartments in the rat brain. As we previously hypothesized (15), increased TH levels in the leptomeninges and choroid plexus, where the proinflammatory reaction is the most intense following LPS administration (42-47), may serve to control inflammatory processes and improve macrophage function (6, 8, 16-18). It can be also speculated, however, that at the same time decreased TH levels might be adaptive for cells in the brain parenchyma. While further research will be necessary to understand the pathophysiological role of these changes, the present findings underscore that TH availability may differ significantly for different cell types of a single organ during the non-thyroidal illness syndrome.
In conclusion, OATP1c1 and MCT8 expression is downregulated in the blood-brain barrier during inflammation, suggesting decreased TH uptake into the rodent brain. TH uptake studies in inflammation models will be essential in the future to test the importance of TH transporter downregulation on TH delivery to the brain.

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**Figure legends**

**Figure 1.** X-ray film autoradiograms of isotopic *in situ* hybridizations for rat and mouse OATP1c1, and mouse MCT8. (A) OATP1c1 mRNA expression in the rat forebrain in the function of time following LPS injection. Hybridization signal decreases after LPS injection; at 4 and 9h, the remaining visible signal in the hypothalamic paraventricular and supraoptic nuclei is vascular, while in other regions, such as the hippocampus and cortex, is primarily astrocytic. (B) LPS has the same effect on OATP1c1 mRNA in mice as in rats. In mice, OATP1c1 mRNA decreases even in the choroid plexus at 9h. (C) MCT8 mRNA in the mouse brain following LPS administration. Neuronal labeling remains unchanged. The bottom panel shows magnified views of the diencephalic area, where several labeled blood vessels (arrowheads) are visible at 24h and 48h, but none at 9h after LPS. cp, choroid plexus; Hip, Hippocampus; PVH, hypothalamic paraventricular nucleus; tan, tanycytes; SO, supraoptic nucleus; VMH, hypothalamic ventromedial nucleus. Scale bars = 2mm.

**Figure 2.** (A) Darkfield emulsion autoradiography images demonstrate the time course of OATP1c1 mRNA expression (silver grain accumulation, white) in the mouse hypothalamus after LPS injection. Hybridization signal labeled only blood vessels but not astrocytes in the hypothalamus. The signal virtually disappears by 9h, when only light signal remains visible in the paraventricular nucleus (PVH). At 48h, OATP1c1 mRNA increases significantly. Scale bar = 200µm. (B) Combination of *in situ* hybridization for OATP1c1 (silver grains) and GFAP immunofluorescence (red) demonstrates that LPS did not affect OATP1c1 mRNA expression in astrocytes. Images were taken from the piriform cortex (control) and adjacent cortical amygdala (LPS). White arrows in the overlay images indicate astrocytes.
with OATP1c1 hybridization signal, open arrows point to blood vessel segments. Note that hybridization signal labels vessel segments in the control, but not in LPS-injected brain. Scale bar = 50µm.

**Figure 3.** (A) Darkfield emulsion autoradiography images illustrate the effect of LPS on OATP1c1 mRNA in the rat brain. The decrease in OATP1c1 mRNA levels at 9h is demonstrated in hybridizations with longer exposure time, while shorter exposure was necessary to visualize the increase at 48h. The top panel shows the dramatic decrease of OATP1c1 mRNA in vessels of the hypothalamic paraventricular nucleus (PVH). Middle panel demonstrates the effect in the cortex, where hybridization in vessels is almost completely absent 9h after LPS, and the remaining signal labels predominantly astrocytes. Arrows indicate longer vessel segments in the control cortex. The bottom panel demonstrates the robust increase in OATP1c1 mRNA levels in blood vessels at 48h after LPS; images were taken from the hypothalamic dorsomedial nucleus. PVH, hypothalamic paraventricular nucleus. Scale bar = 100 µm. (B) Fluorescent *in situ* hybridization demonstrates OATP1c1 mRNA in different astrocyte populations in the rat. Hybridization on paraformaldehyde-perfused sections shows high levels of OATP1c1 mRNA (green) in astrocytes of the hippocampus, cortex, and striatum. Note the numerous intense green puncta denoting OATP1c1 mRNA in the cytoplasm and proximal processes of astrocytes, which were identified by immunofluorescence for GFAP (red). In contrast, more sensitive hybridization using fresh-frozen sections was necessary to detect OATP1c1 mRNA in hypothalamic astrocytes. In these cells, hybridization signal concentrated in only a few, 5-8, smaller sized puncta. Cell nuclei are labeled by the blue fluorescence of DAPI. Scale bar = 20 µm.

**Figure 4.** (A) Results of semiquantitative image analysis of OATP1c1 and MCT8 *in situ* hybridization, and OATP1c1 immunofluorescence. OATP1c1 hybridization signal was quantified in the hypothalamus and represents mRNA expression specifically in blood vessels, as astrocytes were not labeled with the used exposure times. In the mouse MCT8 *in situ* hybridization experiment, the number of labeled blood vessels were counted in the diencephalic area in 3 sections per mouse, and the average number of vessels
per section was calculated and presented in the graph. OATP1c1 immunofluorescence in rats was quantified in images taken from the thalamus by measuring the area (pixel number) covered by immunolabeled vessels. Sample sizes: 4 or 5 rats or mice/group. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$ vs control. (B) Quantitative RT-PCR results from the mouse and rat cortex 9 and 48h after LPS injection. OATP1c1 mRNA is significantly reduced 9h after LPS injection in both species: In rats, MCT8 mRNA tended to decrease at 9h but did not reach significance ($p = 0.07$). Sample sizes: n=6 rats/group and 8 mice/group. * $P < 0.05$, *** $P < 0.001$ vs the corresponding controls.

**Figure 5.** (A) OATP1c1 immunofluorescence shows the time course of OATP1c1 protein in blood vessels of rats following LPS injection. Images were taken from the thalamus; note the decrease in labeling at 9h, and that only scattered vessel segments are labeled at 24h. (B) The same phenomenon is shown in mice in images taken from the rostral perifornical area. (C) MCT8 immunofluorescence in blood vessels of the mouse thalamus; labeling intensity is only mildly reduced at 24h after LPS. Scale bar = 50 µm.

**Figure 6.** (A) Darkfield emulsion autoradiography images demonstrate the effect of LPS on MCT8 mRNA in brain blood vessels. Images were taken from the thalamus where labeled vessels (arrows) stand out due to virtually undetectable hybridization signal in neurons. The vascular signal vanishes completely by 9h after LPS. At 24h, and especially at 48h, more intense hybridization signal labels a higher number of vessels. Scale bar = 200 µm. (B) A similar phenomenon is illustrated in images taken from the rat hippocampus. At 9h after LPS only occasional vessels with faint hybridization signal were detected, while several vessels with intense signal were observed at 48h. Scale bar = 100 µm.

**Figure 7.** (A) *In situ* hybridization for OATP1c1, MCT8 and D2 in the rat leptomeninges. Red fluorescent cresyl violet counterstaining is overlaid on darkfield emulsion autoradiography images to help identify tissue locations (OATP1c1 and D2 images). Top: OATP1c1 mRNA is expressed specifically in
the wall of veins (v) that run in the arachnoid, but not in arteries (a), or the arachnoid tissue itself.

Labeling of veins is absent 9h after LPS injection. Bottom left: MCT8 mRNA is expressed in the
leptomeningeal layers between the hippocampus and thalamus (arrows). Bottom right: D2 mRNA is
expressed in both the arachnoid tissue and wall of arachnoid veins (v), but not arteries (a) 9h after LPS
injection. a, artery; opt, optic tract; SO, supraoptic nucleus; v, vein. Scale bar = 100µm in top, 200µm in
bottom panels. (B) Quantitative RT-PCR results from rat leptomeningeal samples obtained from 6 control
and 8 LPS-treated rats. OATP1c1 mRNA decreases to 10% of control value, while D2 increases ~86-fold
on average. * P < 0.05; *** P < 0.0001 vs control.
Fig 1

(A) OATP1c1 mRNA in rats

(B) OATP1c1 mRNA in mice

(C) MCT8 mRNA in mice
Fig 2

A. OATP1c1 mRNA in the mouse hypothalamus

Control | LPS, 2h | LPS, 4h
--- | --- | ---
PVH | PVH | PVH

Control | LPS, 9h | LPS, 24h | LPS, 48h
--- | --- | --- | ---
PVH | PVH | PVH | PVH

B. OATP1c1 mRNA in mouse cortical astrocytes

Control | LPS, 9h
--- | ---
GFAP | GFAP

GFAP/OATP1c1
---

GFAP labeled astrocytes (arrows) are increased after LPS treatment.
Fig 3

A  Rat OATP1c1 mRNA

9d exposure

PVH
Control  LPS, 9h

Cortex
Control  LPS, 9h

Hypothalamus
Control  LPS, 48h

5d exposure

B  OATP1c1 mRNA in rat astrocytes

Hippocampus  Striatum

Cortex  Hypothalamus

Bar scale: 0.1 mm
**A** OATP1c1 protein in rat brain blood vessels

<table>
<thead>
<tr>
<th>Control</th>
<th>LPS, 2h</th>
<th>LPS, 4h</th>
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<tr>
<td>LPS, 9h</td>
<td>LPS, 24h</td>
<td>LPS, 48h</td>
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**B** OATP1c1 protein in mouse brain blood vessels

| Control | LPS, 9h | LPS, 24h | LPS, 48h |

**C** MCT8 protein in mouse brain blood vessels

| Control | LPS, 9h | LPS, 24h |

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*Note: Figure 5 shows immunofluorescence images of OATP1c1 and MCT8 proteins in the blood vessels of rat and mouse brain under different LPS treatment conditions.*
Figure 6: MCT8 mRNA in blood vessels of mice and rats.
A

OATP1c1, MCT8 and D2 mRNAs in the rat leptomeninges

B

qPCR from rat leptomeninges

Relative mRNA levels

- **OATP1c1**
  - Control: 1.00 ± 0.05
  - LPS 9h: 0.25 ± 0.02

- **MCT8**
  - Control: 1.10 ± 0.03
  - LPS 9h: 1.25 ± 0.08

- **D2**
  - Control: 75 ± 5
  - LPS 9h: 125 ± 10

*P < 0.05, **P < 0.001