MerTK-mediated regulation of myelin phagocytosis by macrophages generated from patients with MS

**ABSTRACT**

**Objective:** To document functional differences between monocyte-derived macrophages (MDMs) of patients with MS and the ability of age/sex-matched healthy donor cells to phagocytose human myelin and to investigate the molecular mechanisms that underlie this.

**Methods:** MDMs were derived from peripheral blood monocytes of 25 untreated patients with relapsing-remitting MS and secondary progressive MS and age/sex-matched healthy controls (HCs). Phagocytosis was assessed by flow cytometry using fluorescently labeled human myelin. Quantification of messenger RNA and protein expression of Tyro3, Axl, and MerTK family molecules was determined by quantitative PCR, Western blotting, and flow cytometry.

**Results:** Cells of patients with MS display a reduced ability to phagocytose human myelin but not red blood cells as compared to matched HCs. These cells express significantly lower levels of the phagocytic tyrosine kinase receptor, MerTK, and its natural ligand, growth arrest-specific 6, independently of the activation state of the cells. Increased expression of interleukin 10 following myelin uptake by healthy donor cells is lost in MDMs of patients with MS; this effect is mediated through the MerTK pathway. Treatment of MS cells with transforming growth factor β (TGFβ) restored both phagocytosis and expression deficits.

**Conclusions:** We describe a molecular mechanism that underlies a defect in myelin phagocytosis by macrophages generated from patients with MS. This abnormality involves decreased expression of MerTK and its ligands and can be rescued by treatment with TGFβ. Neurol Neuroimmunol Neuroinflamm 2017;4:e402; doi: 10.1212/NXI.0000000000000402

**GLOSSARY**

FBS = fetal bovine serum; Gas6 = growth arrest-specific 6; HC = healthy control; IL-10 = interleukin 10; MDM = monocyte-derived macrophage; mRNA = messenger RNA; PBS = phosphate-buffered saline; PPAR-γ = peroxisome proliferator-activated receptor gamma; ProS = protein S; PS = phosphatidylserine; RBC = red blood cell; RRMS = relapsing-remitting MS; RXR = retinoid X receptor; SPMS = secondary progressive MS; TAM = Tyro3, Axl, and MerTK; TGFβ = transforming growth factor β.

Clearance of myelin debris following CNS tissue injury in MS is required for subsequent tissue repair.1 Myeloid cells within MS lesions capable of phagocytosis include both microglia and macrophages that are derived from circulating monocytes.2–4 We previously observed that the tyrosine kinase receptor, MerTK, a member of the TAM family (Tyro3, Axl, and MerTK) family of tyrosine kinase receptors, is important in mediating myelin phagocytosis by human myeloid cells.5 MerTK expression correlated with increased phagocytosis and was dependent on the myeloid cell activation state, with transforming growth factor β (TGFβ) enhancing expression of the TAM family receptors and pro-inflammatory cytokines decreasing their expression.5

Myeloid cell–mediated myelin phagocytosis was inhibited in vitro by selective MerTK antagonism.5

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TAM receptors mediate their action via their 2 natural ligands, growth arrest-specific 6 (Gas6) and protein S (ProS). Receptor-bound Gas6 and ProS concurrently bind to phosphatidylserine (PS), a phospholipid, exposed on the outer membrane of apoptotic cells.6,7 PS is also found in lipid-rich myelin.8 Both receptors and ligands are needed for efficient phagocytosis.9 TAM receptor signaling is implicated in regulating inflammation because deficiencies in signaling have been associated with sustained immune activation and chronic inflammation.10 Recently identified and characterized polymorphisms in the MerTK gene have established MerTK as a novel risk gene for MS susceptibility.11,12

A recent study demonstrated that peripheral blood monocyte-derived macrophages (MDMs) of patients with MS display impaired myelin phagocytosis and an enhanced pro-inflammatory phenotype.13 In the current study, we confirm that MDMs of patients with MS are functionally abnormal in their ability to phagocytose and that this deficit is relatively selective to myelin. This functional deficit is linked to altered MS myeloid cell expression of MerTK and its ligands, where both expression and functional deficits can be subsequently rescued by exposure to the TGFβ. Our results indicate that the TAM receptor-signaling axis may be amenable to the therapeutic intervention.

METHODS Standard protocol approvals, registrations, and patient consents. All institutional ethics approval was obtained according to Canadian Institutes of Health Research guidelines. All patients provided written informed consent in accordance with the Declaration of Helsinki, and approval was obtained from the Institutional Review Board of McGill University.

Patient selection. A total of 25 patients with MS (18 relapsing-remitting MS [RRMS] and 7 secondary progressive MS [SPMS]) were recruited at the Montreal Neurological Hospital MS Clinic (figure 1A). Of the 18 patients with RRMS in this study, treatment history was as follows: treatment naive (7), Copaxone (5), Gilenya (2), Avonex (1), Rebif (2), and Tecfidera (1). All treatment history was as follows: treatment naive (7), Copaxone (3), Gilenya (2), Avonex (1), Rebif (2), and Tecfidera (1). All patients had a minimum 3-month washout period prior to contributing to this study, and we did not observe any difference between treatment-naïve patients and those currently off drug, Healthy age- (± 5 years) and sex-matched controls were used for comparison.

Cell culture (human MDMs). Peripheral blood mononuclear cells were isolated from the venous blood using Ficoll-Paque density gradient centrifugation (GE Healthcare). Monocytes were positively selected using CD14+ immunomagnetic beads (according to manufacturer’s instructions; Miltenyi Biotec, Bergisch Gladbach, Germany) to routinely achieve 95%–99% purity. Monocytes were cultured (5 × 10⁵ cells/mL) in Roswell Park Memorial Institute 1640 media supplemented with 10% fetal bovine serum (FBS), 0.1% penicillin/streptomycin, and 0.1% glutamine. MDMs were differentiated in vitro with macrophage-colony stimulating factor (25 ng/mL) for 6 days. For select experiments, TGFβ (20 ng/mL) was added on days 1 and 5.

Phagocytosis assay. Endotoxin-free human myelin was prepared as previously described.14 To evaluate myelin phagocytosis, MDMs were plated in 12-well plates. Myelin was conjugated to a pH-sensitive dye (pH-Rodamine; Invitrogen, Carlsbad, CA) for 1 hour in phosphate-buffered saline (PBS) (pH 8) and added to the culture wells (20 μg/mL) for 1 hour; cells were washed, detached, pelleted, and passed through a flow cytometer (FACS Fortessa; BD Biosciences, San Jose, CA) to quantify the percentage of myeloid cells containing internalized fluorescent myelin. A similar protocol was used to study phagocytosis of red blood cells (RBCs) acquired from Ficoll-Paque density gradient centrifugation and diluted in PBS at a ratio of 1:2,500. RBCs were then incubated with opsonizing antibody (20 μg/mL: Cederlane) and pH-rodamine for 30 minutes at room temperature. MDMs were exposed to opsonized RBCs for 1 hour at a 4:1 (RBC:MDM) ratio, and internalized RBCs were quantified as an increase in mean fluorescent intensity in the PE channel.

Quantification of TAM receptor/ligand and cytokine mRNA expression. Cells were lysed in TRizol reagent (Invitrogen) and stored at −80°C. Total RNA extraction and reverse transcription were performed according to manufacturer’s instructions (Qiagen, Hilden, Germany). Individual gene expression was analyzed by PCR using specific TaqMan probes to assess expression relative to Gapdh (housekeeping gene). Fold change calculations were done using the −ΔΔ threshold cycle method.

Quantification of protein expression. For quantitation of MerTK protein expression, MDMs were detached using EDTA/PBS (2 mM). Cells were blocked in 10% human serum and normal mouse immunoglobulin G (3 ng/mL) in serum and normal mouse immunoglobulin G (3 ng/mL) in fetal bovine serum (FBS) (pH –80°C. Total RNA extraction and reverse transcription were performed according to manufacturer’s instructions (Qiagen, Hilden, Germany). Individual gene expression was analyzed by PCR using specific TaqMan probes to assess expression relative to Gapdh (housekeeping gene). Fold change calculations were done using the −ΔΔ threshold cycle method.

Western blotting Polyvinylidene difluoride membranes were probed with anti-TGFβ1 antibody (polyclonal rabbit anti-human antibody 1:5,000; Abcam, Cambridge, United Kingdom) and detected using horseradish peroxidase-conjugated secondary antibodies (1:1,000: Califiochem, San Diego, CA). Membranes were stripped with ReBlot (Millipore, Billerica, MA) and re-probed for loading control (anti-β-actin; 1:1,000; Invitrogen). Protein band intensities were quantified using ImageJ software (NIH), and data were expressed as relative fold change to untreated controls.

Results MDMs of patients with MS exhibit abnormally reduced phagocytosis of human myelin (but not RBCs), which can be restored by TGFβ. A lower percent of MS (both RRMS and SPMS) patient–derived MDMs phagocytosed myelin compared with their matched healthy controls (HCs) (figure 1C). TGFβ treatment increased myelin phagocytosis in both HCs and RRMS MDMs to levels that were no longer
distinguishable (figure 1D). MDMs of patients with MS exhibited normal phagocytosis of opsonized human RBCs (figure 1E), which was not enhanced by TGFβ (figure 1F).

Reduced expression of MerTK and its ligand in MDMs of patients with MS can be restored by TGFβ. messenger RNA (mRNA) levels of MerTK were significantly downregulated in patients with RRMS as compared to HCs (figure 2A) and were restored following TGFβ exposure (figure 2B), findings that were confirmed when assessing MerTK cell-surface protein expression prior to (figure 2C) and following (figure 2D) TGFβ exposure. Expression of the TAM receptor ligands Gas6 (figure 2E) and ProS (figure 2G) was also examined at the mRNA level. MDMs from patients with RRMS display significantly decreased levels of Gas6 expression, with levels of ProS expression comparable between the 2 populations. Expression of Gas6 in MDMs of untreated patients with MS was restored to HC levels following TGFβ exposure with TGFβ having no effect on ProS (figure 2, F and H).
Increased expression of TGFβ receptors in MS patient-derived MDMs compared with HC cells. Expression of all 3 TGFβ receptors was significantly increased in monocytes of patients with MS as compared to HC monocytes (figure 3A). Expression of TGFβR1 protein was examined by Western blotting and mirrored the mRNA expression profile with higher expression of TGFβR1 in monocytes of patients with RRMS as compared to matched HCs. TGF-β1 and 2 expression was also increased in MDMs of patients with MS (data not shown). Monocytes of healthy controls and patients with MS did not differ in expression of CD80, HLA-DR, CD274, and CCR7 (figure 3B).

MDMs of patients with MS are unable to adopt an anti-inflammatory profile following myelin phagocytosis. To examine the consequence of reduced myelin uptake in these MDM populations, we measured interleukin 10 (IL-10) mRNA and protein expression following myelin uptake both in the presence and absence of TGFβ. TGFβ increased IL-10 mRNA expression in both cell populations with myelin ingestion causing a significant increase in expression in MDMs of HCs but not in MDMs of RRMS (figure 4, A and B). This increase was inhibited following pre-incubation with the MerTK-specific antagonist (UNC2025) (figure 4A). IL-10 protein was measured by ELISA; MDMs of patients with RRMS release significantly less IL-10 protein following phagocytosis as compared to matched healthy donor cells.

Discussion In line with a recent study, we report a selective deficit in MS patient-derived MDMs to phagocytose myelin debris.13 We propose that this deficit is linked to an abnormal reduction in expression of the TAM receptor MerTK and its natural ligand as observed at both mRNA and protein levels. The MS MDM phagocytosis deficit could be rescued with exposure to TGFβ, restoring both expression of the molecular machinery (MerTK and Gas6) necessary for the TAM receptor-mediated clearance of myelin, and, at the functional level, restoring the phagocytic ability of MDMs.
We also observed that MDMs of HCs but not MDMs of patients with MS were able to upregulate expression of the anti-inflammatory cytokine, IL-10, following myelin ingestion. This reduced expression of IL-10 in cells of patients with MS may be due to their aforementioned reduced ability to phagocytose, which

Expression of TGFβ receptors 1–3 was examined by quantitative PCR in freshly isolated monocytes of HCs (circle) and patients with MS (square). (A) Levels of all 3 receptors were significantly increased in MDMs of patients with MS as compared to matched controls. (B) Analysis of TGFβR1 protein expression was performed by Western blotting, showing a significant reduction in expression in monocytes of patients with MS as compared to matched controls (n = 4). (C) Expression of known monocyte activation markers (CD80, HLA-DR, CD274, and CCR7) did not differ between monocytes of HCs and patients with MS. mRNA fold change was calculated using the 2-ΔΔ threshold cycle method, relative to Gapdh (housekeeping gene). (A and B) Student’s paired t-test. HC = healthy control; MDM = monocyte-derived macrophage; mRNA = messenger RNA; RRMS = relapsing-remitting MS; TGFβ = transforming growth factor β; TGFβR1 = transforming growth factor β receptor 1.
has been shown to have an anti-inflammatory effect on macrophages and microglia.\textsuperscript{15,16} We previously showed that the MerTK-specific antagonist UNC2025 significantly inhibited IL-10 induction in MDMs of HCs.\textsuperscript{5} Of note, the defective MS MDM phagocytosis did not appear to simply reflect an “activated” myeloid phenotype because their expression of classic activation markers was not increased, albeit a nonsignificant increase in the homing receptor CCR7 may reflect an enhanced ability of these cells to traffic to the CNS.\textsuperscript{17} Of interest, we observed increased expression of all 3 TGF\(\beta\) receptors in MS patient–derived monocytes. TGF\(\beta\)R1 in particular was seen to be upregulated at both mRNA and protein levels in monocytes of patients with RRMS. This contrasts with a decrease in expression of these receptors following exposure to classic pro-inflammatory activating stimuli such as lipopolysaccharide and interferon gamma (data not shown), again suggesting that these cells are not classically activated, “M1-skewed” cells. It is difficult to comment on whether our myeloid populations contain CNS-experienced monocytes that have returned to the periphery; this increased expression of TGF\(\beta\) receptors may be in response to either interaction with activated lymphoid cells in draining lymph nodes or possibly in response to blood-borne inflammatory markers reflecting the ongoing CNS pathology.

We previously reported that TGF\(\beta\) can cause increased expression of MerTK and its ligands in both human MDMs and microglia.\textsuperscript{5} In the current study, activation of MDMs of patients with MS with TGF\(\beta\) was able to overcome the functional abnormality in myelin phagocytosis, suggesting that this deficit could be targeted therapeutically. It is also reported that the decreased ability of MDMs of patients with MS to phagocytose myelin can be ameliorated by treatment with pioglitazone, an agonist for the nuclear receptor, peroxisome proliferator–activated receptor gamma (PPAR-\(\gamma\)) and its binding partner, the retinoid X receptor (RXR).\textsuperscript{13} Of interest, signaling via PPAR-\(\gamma\)-RXR and other nuclear hormone receptors has been shown to induce MerTK expression that promoted the clearance of apoptotic bodies by dendritic cells.\textsuperscript{18–20} MerTK has been identified as a gene associated with increased MS susceptibility in 2 large, independent discovery and replication cohorts.\textsuperscript{11,12,21} Susceptibility single nucleotide polymorphisms in the MerTK gene were significantly associated with altered MerTK gene expression in purified monocytes but not in lymphocytes.\textsuperscript{12} Our study is not powered to resolve whether differences in myelin phagocytic activity reflect MerTK gene polymorphisms.

In the periphery, reduced myeloid cell expression of MerTK has been shown to directly affect the clearance of apoptotic lymphocytes in lymphoid organs.\textsuperscript{22} In TAM receptor knockout mice, this leads to a buildup of apoptotic cells in these structures causing swelling and leading to a broad spectrum of autoimmune disease, characterized by lymphocyte infiltration in most major organs and high titers of circulating auto-antibodies.\textsuperscript{23} This has a direct relevance to MS where defective T-cell tolerance can lead to expansion of self-reactive T cells.\textsuperscript{24} We described a molecular mechanism that underlies a selective, phagocytic deficit in macrophages.
generated from peripheral monocytes of untreated patients with MS. This reduced ability to phagocytose myelin involves decreased expression of MerTK and its ligand, Gas6, previously implicated in myelin phagocytosis by myeloid cells, and may be amenable to the therapeutic intervention.

**AUTHOR CONTRIBUTIONS**
L.M.H., J.P.A., and A.B.-O. conceived the project. L.M.H. designed the studies, performed the experiments, analysis, and data interpretation, and wrote the manuscript. J.H.J. and S.-Y.W. performed experiments and analysis of data. Y.H.L. performed experiments and proofread the manuscript. S.A. analyzed data. H.T. recruited patients. All authors critically revised the manuscript.

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