Is there a link between inflammation and fatigue in multiple sclerosis?

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Purpose: Among autoimmune diseases of the central nervous system stands multiple sclerosis (MS), which is characterized by demyelination, synaptopathy, and neurodegeneration. MS fatigue can affect up to 90% of patients and be very disabling, with a drastic impact on their quality of life. To date, the evaluation of MS fatigue has relied mainly on subjective scales, and actual therapeutic interventions are challenged by modest efficacy and numerous undesirable effects. Therefore, finding biomarkers of MS fatigue might help in optimizing evaluation and treatment strategies. The main objective here was to assess the relationship between MS fatigue and inflammatory or other immunomediated markers.

Methods: Research was conducted according to PRISMA guidelines. Computerized databases (ie, PubMed/Medline and Scopus) were consulted till February 2018 aiming to identify articles that addressed inflammation and MS fatigue. Studies in English and French published at any time were considered.

Results: A total of 27 studies matched the research criteria. Inconsistency existed regarding the relationship between fatigue and the orexin A system, hypothalamus–pituitary–adrenal axis, and cerebrospinal fluid inflammatory markers. As for peripheral markers, although there was scarcity in the available data, serum proinflammatory cytokines (ie, IL6, TNFα, and IFNγ) seem to be associated with MS fatigue. Finally, no link was found between MS fatigue and T-cell populations (ie, CD3+CD4+ T lymphocytes, regulatory T cells) or other peripheral markers of inflammation (ie, CRP, erythrocyte-sedimentation rate, soluble ICAM1).

Conclusion: Future large-scale studies would benefit from comparing the relationship between fatigue and immune measures in patients with different disease phenotypes with and without disease-modifying drugs. With the subjective nature of fatigue scales, finding objective biomarkers for fatigue would be of great help.

Keywords: pathophysiology, cytokines, interleukins, cerebrospinal fluids, inflammatory markers

Introduction

Among autoimmune diseases of the central nervous system (CNS) stands multiple sclerosis (MS), which is the second-most common cause of physical handicap in young individuals.1–3 Its pathophysiological hallmarks are demyelination, synaptopathy, and neurodegeneration.4,5 Throughout the disease course, MS patients can experience periods of acute symptom emergence separated by symptom-free intervals. This characterizes the relapsing–remitting (RR) MS phenotype, which usually converts to a secondary progressive phase where patients can experience steady clinical deterioration.6 Primary progressive MS represents a third disease phenotype, where patients...
witness an evolutionary pattern of their disease from onset. Demyelination appears to be the fingerprint of the first type (ie, RRMS), whereas neurodegeneration/axonal loss seems to be the backbone of progressive types. While an immunomediated attack by blood-borne autoreactive T lymphocytes would dictate the occurrence of demyelination, immunomediated processes involving immune cells and soluble cytokines could also lead to excitotoxic changes and neurodegeneration, based on experiments involving the animal model of MS.5

Regardless of the disease phenotype, patients may experience a panel of symptoms involving the sensory, motor, cerebellar, emotional, cognitive, and behavioral domains. Among the frequently encountered symptoms, MS fatigue can affect up to 90% of patients and be very disabling, with a drastic impact on their quality of life.6 It is a challenging symptom that is described by patients as “malaise”, “excessive tiredness”, or “weakness” that seems to worsen throughout the day, as well as with hot and humid environments.4 From a scientific perspective, MS fatigue is considered a multidimensional symptom with physical, cognitive, and psychosocial components. Among the available definitions, some authors consider fatigue a lack of physical and/or mental energy.4,6 For another group of authors, fatigue designates a failure to initiate and/or maintain physical or mental activities requiring self-motivation in the absence of or not related to physical or cognitive dysfunction.4,6

Ever since the original work of Freal et al in 1984,7 there has been growing interest in understanding the pathophysiology of MS fatigue, especially the fact that this symptom remains difficult to be reported by patients and managed by physicians. To date, the evaluation of MS fatigue has relied mainly on subjective scales, such as the Modified Fatigue Impact Scale (MFIS), Fatigue Severity Scale (FSS), and Fatigue Scale for Motor and Cognitive Functions (FSMC), among others, and actual therapeutic interventions are challenged by their modest efficacy in face of their numerous undesirable effects.8 From this perspective, understanding the underlying mechanisms of this symptom might be of help in easing its evaluation and optimizing patient care. In a previous work, we addressed the cerebral anatomical correlates of MS fatigue.4 Based on neuroimaging studies, pathological findings were observed in a corticostriatothalamocortical loop that was linked to MS fatigue. These findings included regional gray- and white-matter pathologies, as well as abnormal patterns of brain activation. The inflammatory and immune medium might be implicated as well in the context of MS fatigue. Therefore, the main aim of the current work was to assess the role of MS-related central and peripheral inflammation and immunomediated endocrine dysregulation in the development of this symptom.

**Study selection**

Research was conducted according to the PRISMA guidelines.9 First, computerized databases that index peer-reviewed journals (PubMed/Medline and Scopus) were consulted till the end of February 2018. The research aimed to identify articles that addressed the relationship between MS fatigue and inflammatory, immune, and endocrine factors. Studies that were published at any time in English and French were considered. The following research terms were combined: (“fatigue” OR “fatigue severity scale” OR “FSS” OR “Modified Fatigue Impact Scale” OR “MFIS” or “Fatigue Scale for Motor and Cognitive Functions” or “FSMC”) AND (“multiple sclerosis” OR “MS”) AND (“inflammation” OR “inflammatory” OR “immune” OR “cytokine” OR “interleukin” OR “cerebrospinal fluid” OR “CSF” OR “lymphocytes” OR “blood cells” OR “endocrine”). In addition, both authors independently checked the references of the articles obtained, aiming to obtain additional sources. The initial search identified 503 articles in PubMed/Medline and 258 articles in Scopus. After removal of duplicates and excluding reviews, opinions, editorials, commentaries, viewpoints, and research articles involving healthy volunteers or patients with autoimmune diseases other than MS, 25 articles were retained. An additional two references were retrieved from the articles’ reference lists, yielding a total of 27 articles that were considered in the qualitative synthesis. These comprised information on MS fatigue and inflammatory or neuroendocrine markers and addressed the relationship between fatigue and hypothalamic function (two about the orexin A system, eight about the hypothalamic-pituitary–adrenal [HPA] axis), cerebrospinal fluid (CSF) markers (one about humoral and cellular CSF markers, one about CSF cytokines), serum-cytokine or blood-cell expression (15), or other peripheral inflammatory markers (three). In addition, six studies assessed changes in fatigue and cytokine profiles following exercise (four) or pharmacological (two) interventions. For the sake of this work, data of the selected studies are classified as central inflammation and neuroendocrine dysregulation and peripheral inflammation.

**Central inflammation, neuroendocrine dysregulation, and MS fatigue**

The exploration of inflammatory patterns within the CNS is possible by means of CSF analysis. However, the procedure...
consists of performing lumbar puncture, a procedure that is not only difficult to perform but also traumatizing for patients. This explains the scarcity of existing literature in this field. Available works on CSF analysis and MS fatigue focused on studying inflammation-related neuroendocrine dysregulation, humoral and cellular components, and cytokine levels. The remaining literature employed serum and salivary hormonal tests to assess specific hypothalamic functions and their relationship with fatigue.

To start, the hypothalamus plays a role in controlling several homeostatic functions. Some researchers were interested in assessing the relationship between MS fatigue and CSF levels of orexin A (also known as hypocretin 1), a hypothalamic peptide involved in arousal, motivation, energy, and circadian rhythm. In fact, consolidating night sleep and keeping adequate daytime activity seem to be respectively promoted by low and high orexin A levels. Therefore, the rationale behind these works lay in the fact that neuroinflammation, such as that seen in MS, may impact the orexin A system. As such, one can speculate that downregulation of the latter system might happen in the course of MS and result in sleep disorders and/or fatigue. The first insight on orexin A-system status in MS derived from case reports on patients suffering from hypersomnia displaying low CSF levels of orexin A. Afterward, Papuć et al studied orexin A levels in MS patients and healthy controls. In the absence of group difference (MS vs healthy controls) with regard to this peptide, significant positive correlation was found between fatigue severity and orexin A levels in the whole patient group. Although this positive relationship was unexpected, the authors of this work hypothesized that this might have occurred due to the activation of endogenous compensatory mechanisms. One year later, Constantinescu et al were not able to replicate this correlation. Here, the authors found neither a group difference in orexin A levels between MS patients and other patients with inflammatory and noninflammatory neurological disease nor a correlation between orexin A levels and fatigue scores. Given the impact of daytime and season on orexin A levels, there is a good chance that these two studies were performed in different seasons and/or at different times of the day, a finding that could provide an explanation for the difference in the reported results.

Besides regulating the orexin A system, the hypothalamus is involved in many axes, of which the most studied is the HPA axis. Facing physiological and stressful situations, the hypothalamus secretes the corticotropin-releasing hormone (CRH) to stimulate the activity of the anterior pituitary gland. The latter responds by producing the adrenocorticotropic hormone (ACTH) which in turn activates the adrenal glands (ie, zona fasciculata) yielding cortisol production. Proinflammatory cytokines can influence the activity of the HPA axis. This might provide an explanation for the HPA-axis dysregulation that appears to occur in patients with MS. In fact, the majority of studies on this topic have shown a hyperactive HPA axis, with fewer reports suggesting a hypoactivity pattern. Few studies have assessed the relationship between MS fatigue and HPA-axis activity.

Heesen et al employed combined dexamethasone–CRH challenge and low-dose dexamethasone-suppression tests. Both of these are used widely to measure activity of the HPA axis. The dexamethasone-suppression test consists of orally administering dexamethasone, a synthetic glucocorticoid, the night before blood sampling, in order to check the suppression of cortisol production (which is the normal physiological reaction). The combined dexamethasone–CRH-challenge test resembles the first, but CRH is also given intravenously the day of blood sampling and blood withdrawn at regular intervals to determine plasma levels of ACTH and cortisol levels at different times. In their four studies, the authors did not detect any significant association between MS fatigue and HPA-axis activity. In line with these results, Akcali et al employed a more comprehensive neuroendocrine evaluation that included plasma levels of ACTH, cortisol, and other pituitary products, namely corticotropin-like intermediate-lobe peptide (CLIP), which is an ACTH variant, and melanocyte-stimulating hormone (α-MSH, β-MSH, γ-MSH), produced in the anterior pituitary gland and previously found to be implicated in chronic fatigue syndrome (ie, α-MSH). Although abnormal HPA measures were observed in MS patients compared to healthy controls (ie, higher ACTH, cortisol and α-MSH and lower CLIP levels among patients), these measures did not differ between fatigued and unfatigued MS patients, suggesting the absence of any relationship between HPA-axis activity and MS fatigue. Conversely, a third study by Gottschalk et al employed combined dexamethasone–CRH-suppression tests and found significantly higher ACTH plasma levels among fatigued compared to unfatigued counterparts. The discrepancy in the results of the aforementioned works might have resulted from differences in clinical characteristics and treatments of the MS cohorts studied. Contrarily to Heesen et al and Akcali et al, mostly enrolled patients receiving disease-modifying drugs, Gottschalk et al recruited drug-naïve patients. Here, it is worth noting that immunotherapy may impact cytokine-expression levels and thus might influence HPA-axis activity.
In addition to the previously mentioned studies, Powell et al and Gold et al focused on the assessment of the cortisol awakening response (CAR) in MS patients, using a salivary test.²⁷,³³ CAR is a spike in serum cortisol around 30–45 minutes after awakening, and is crucial for sustaining normal circadian rhythm and wakefulness.³⁴ In the former trial, baseline fatigue scores, but not those obtained at the same day of CAR testing, were correlated with CAR.³³ In the latter, CAR did not predict MS fatigue, as per regression-analysis results.²⁷

Among the other adrenal products stands dehydroepiandrosterone (DHEA) and its sulfated ester (DHEAS). Low DHEA and DHEAS levels have been linked to fatigue in some autoimmune diseases, such as systemic lupus erythematosus and rheumatoid arthritis.³⁵,³⁶ In the only available study addressing serum levels of DHEA and DHEAS in MS patients, lower levels of both components were detected in fatigued compared to unfatigued patients.³⁷ Interestingly, these results support those of an earlier pharmacological study, in which fatigue improvement was obtained following DHEA hormone replacement.³⁸ However, the results of the latter work should be interpreted with caution, mainly because of its nonrandomized design. These preliminary findings warrant further research on this matter.

In addition to studies on HPA axis and MS fatigue, some researchers were interested in evaluating humoral, cellular, and other immune CSF markers. For instance, Biberacher et al included an exploratory and a validation phase that contained several evaluations. Of interest, they assessed the relationship between fatigue and several cellular and humoral CSF markers.³⁹ No correlation was found between fatigue scores and any of the CSF markers. More interestingly, fatigue scores tended to correlate negatively with CSF CD4:CD8 ratio in the discovery group and correlate positively with the former ratio in the validation group. However, the multivariate model failed to detect associations between fatigue and CSF parameters in either group (exploratory vs validation). A recent work aimed to understand the relationship between MS-fatigue and CSF-interleukin levels, particularly IL6 and IL8. While IL6 took part in innate and adaptive immune responses, including differentiation of T helper 17 cells, IL8 was mainly implicated in innate immunoresponses and had cytokine- and chemokine-like functions. In this work, Brenner et al documented a significant correlation between fatigue scores and IL6 levels.⁴⁰ This relationship was only seen among patients not receiving MS treatment. This might explain the absence of association in Biberacher et al, where >75% of patients were under MS therapies. Another difference between the studies lies in their methodological approach which consisted of cellular and humoral markers in the first versus interleukins in the second. The different clinical and demographic characteristics between the cohorts might present a third plausible explanation for the inconsistency in their results. Table 1 provides a summary of these studies.

**Table 1** Studies assessing the relationship between multiple sclerosis fatigue and central inflammatory or neuroendocrine markers

<table>
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<tr>
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<td><strong>Studies assessing the HPA axis</strong></td>
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<tr>
<td>Heesen et al²⁴</td>
<td>40 patients (8 RR, 19 SP, 13 PP, sex details NA, treated, but details NA); 11 HCs (sex-matched, details NA)</td>
<td>MFIS, FSS</td>
<td>Dexamethasone-suppression test</td>
<td>HPA-axis hyperactivity among progressive patients (but not RR patients) No correlation between fatigue and HPA-axis activity</td>
</tr>
<tr>
<td>Gottschalk et al²⁹</td>
<td>15 fatigued MS patients (all RR, 12F/3M); 16 unfatigued MS patients (all RR, 10F/6M); 19.4% of whole cohort treated</td>
<td>FSS, MFIS, VAS</td>
<td>Dexamethasone-suppression test</td>
<td>Higher adrenocorticotropin (but not cortisol) levels in the Dexam-CRH test among fatigued patients Correlation analysis NA No influence of sex on fatigue</td>
</tr>
<tr>
<td>Heesen et al²⁵</td>
<td>15 fatigued MS patients (6 RR, 8 SP, 1 PP, 9/6M, 66.7% treated); 15 unfatigued MS patients (11 RR, 2 SP, 2 PP, sex-matched, 9/6M, 60% treated)</td>
<td>MFIS, FSS</td>
<td>Dexamethasone-suppression test</td>
<td>No group difference in Dexam-CRH test Correlation between fatigue and HPA-axis activity</td>
</tr>
<tr>
<td>Heesen et al²⁶</td>
<td>50 MS patients (27 RR, 23 SP, 29F/21M, 48% treated)</td>
<td>MFIS, FSS</td>
<td>Dexamethasone-suppression test</td>
<td>Group comparison NA No correlation between fatigue and HPA-axis activity</td>
</tr>
<tr>
<td>Gold et al²⁷</td>
<td>44 MS patients (all RR, all female, 59% treated)</td>
<td>MFIS, FSS</td>
<td>Dexamethasone-suppression test and salivary CAR test</td>
<td>Group comparison NA Dexamethasone-suppression test and CAR data did not predict fatigue</td>
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Peripheral inflammation and MS fatigue

Immunodysregulation constitutes the core of the disease process in MS. The role of peripheral inflammation in the development of MS fatigue has been considered in few immunological studies that assessed serum levels of cytokines, cytokine-producing cells, or other inflammatory markers. The first evidence on this matter dates back to 1990. In a series of eight fatigued MS patients, the authors assessed serum levels of IL2 and its soluble receptor, an interleukin that was suggested to intervene in CNS demyelination in MS and was previously found to be associated with the disease state in an animal MS model. The authors reported that in all patients, the variables studied were below the level of sensitivity of the test used (enzyme-linked immunosorbent assay [ELISA]), and thus denied the association between MS fatigue and
IL2 levels. Afterward, Flachenecker and colleagues applied real-time polymerase chain reaction (RT-PCR) to compare the expression (mRNA) of circulating proinflammatory (ie, TNFα, IFNG) and anti-inflammatory cytokines (IL10) in fatigued and unfatigued MS patients.\textsuperscript{45} TNFα expression was heightened among fatigued patients, with no group differences regarding IFNG and IL10 expression. The role of TNFα has also been suggested in some studies that documented a decrease in TNFα levels following exercise therapy, a finding that was paralleled by an improvement in MS fatigue.\textsuperscript{46–48}

In a similar way to these studies, the same cytokines were assessed in two trials by Heesen et al.\textsuperscript{25,49} ELISA was employed in both works. In the first, only fatigued MS patients were recruited and compared to healthy controls at baseline following a cognitive task that assessed psychological stress.\textsuperscript{49} No group difference was observed at baseline with regard to cytokine levels but following the cognitive task the MS group had relatively diminished IFNG response compared to the healthy group. No significant correlations were observed between fatigue scores and cytokine levels. In the second study, the authors recruited two groups of MS patients with and without fatigue and obtained positive findings.\textsuperscript{25} That is to say that, compared to unfatigued patients, the fatigued ones had higher proinflammatory cytokines (TNFα and IFNG), with no group differences observed with regard to the anti-inflammatory cytokine (IL10). A correlation was also found between fatigue scores (MFIS and FSS) and TNFα and IFNG levels. The difference between the studies might lie in differences in the study populations, where the first considered only fatigued patients, whereas the second also considered an unfatigued patient group.

The same group of authors performed a third study that highlighted the role of IFNG-producing CD8+ T cells as the only significant predictor of fatigue scores.\textsuperscript{27} The contribution of IFNG to the pathophysiology of MS fatigue was also assessed by Pokryszko-Dragan et al.\textsuperscript{50} Using flow cytometry, they studied the production of this cytokine by CD3+CD4+ T lymphocytes. Heightened IFNG production was observed among fatigued patients compared to unfatigued patients and healthy controls, a finding that also tended to correlate with fatigue scores (FSS and MFIS). However, when multiple regression analysis was run, fatigue scores were not linked to IFNG. The role of interferon signaling was also highlighted in a pilot study by Mulero et al.\textsuperscript{51} Here, compared to healthy controls, fatigued MS patients had significant activation in genes participating in the systemic interferon response.

Another important factor in the context of MS is IL17 which appears to be implicated in glutamate-mediated excitotoxicity and thus may explain the potential link between inflammation and neurodegeneration in MS.\textsuperscript{52} The relationship between fatigue and IL17 levels has been addressed by three works that yielded inconsistent outcomes, with two studies confirming such a correlation\textsuperscript{47,53} and one denying it.\textsuperscript{56}

Other circulating cytokines were the subject of a study by Malekzadeh et al, who compared serum levels of a panel of proinflammatory (IL1β, IL2, IL6, IL8, IL12p70, IL17, TNFα, and IFNG) and anti-inflammatory cytokines (IL4, IL5, IL10, and IL13) in fatigued and unfatigued MS patients using an electrochemiluminescence-based multiplex immunoassay.\textsuperscript{53} In the absence of group differences with regard to these variables, a significant correlation was found between IL6 levels and fatigue scores. Of interest, the IL6 levels were found to diminish in another work following the administration of antifatigue pharmacological therapies such as amantadine and pemoline, a finding that was paralleled by fatigue improvement.\textsuperscript{54}

An additional study by Akcali et al evaluated serum levels of TNFα, IL10, and other interleukins (IL1β, IL2, and IL35) in MS patients and healthy controls.\textsuperscript{28} Compared to healthy controls, the only group difference was observed with regard to IL35 and IL2, which were higher in the patient group. However, neither was there a difference between fatigued and unfatigued patients nor was there any correlation between fatigue scores and the markers considered.

Again, inconsistency in results across studies might have been related to cohort characteristics and sample-size difference, but also to other plausible factors. For instance, levels of pro/anti-inflammatory markers can fluctuate during the disease course and disease-modifying therapies can impact cytokine expression.\textsuperscript{28,55} To overcome this limitation, recent studies by Alvarenga-Filho et al enrolled drug-naive MS patients.\textsuperscript{47,56} Here, higher IL6 and TNFα levels were observed among fatigued patients,\textsuperscript{47} and fatigue scores were correlated with IL6 and TNFα levels\textsuperscript{47,56} and tended to correlate with IFNG levels.\textsuperscript{57} Another factor to consider is differences in methods adopted in measuring cytokine levels. This is obviously illustrated with regard to IFNG. In reality, IFNG was found to be unrelated to fatigue when using RT-PCR\textsuperscript{45} and a multiplex kit\textsuperscript{53} and associated or tended to associate with fatigue when using ELISA\textsuperscript{25,56} and flow cytometry.\textsuperscript{50} As such, these tests seem to have different sensitivity/specificity profiles, and this would hamper the possibility of drawing formal conclusions from the existing literature. Moreover, even when using ELISA, results might vary between in vivo and in vitro approaches. The best documentation of this variability can be found in the
studies on IL17 (ie, serum-cytokine levels in vivo and versus stimulated cytokines production in vitro). In fact, one of these works simultaneously adopted both approaches, but only documented a significant correlation between fatigue and in vitro IL17 production. In addition, differences in statistical approaches might explain differences in results reported. While positive studies IFN-γ employed group-comparison and correlation analysis, studies that failed to demonstrate this relationship adopted group comparison without correlation analysis or multiple regression analysis.

Given the role of T cells in the pathophysiology of MS, few works have addressed the relationship between circulating T-cell populations and MS fatigue and failed to document any association. Fatigued and unfatigued MS patients did not differ with regards to the amount of IFN-γ-producing CD3+CD4+ T lymphocytes in one study or the number of leukocyte and lymphocyte subsets including regulatory T cells and its suppressive function in another study.

Finally, three studies included markers of peripheral inflammation in the assessment of MS fatigue. In the first, Giovannoni et al failed to demonstrate any relationship between fatigue and serum (ie, CRP, soluble ICAM1) or urinary (daily urinary neopterin excretion measured over 2 weeks) markers. Similarly, in a second study by Flachenecker et al, the erythrocyte-sedimentation rate, a marker of systemic inflammation, did not differ between fatigued and unfatigued MS patients. Finally, in a third study by Adamczyk-Sowa et al, no correlation was found between MS fatigue and plasma lipid hydroperoxides or homocysteine concentration, which are markers of oxidation. Table 2 summarizes these studies.

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<th>Study</th>
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<th>Fatigue assessment</th>
<th>Method</th>
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<td>Rudick and Barma</td>
<td>Eight fatigued MS patients (disease details NA, 6F/2M), 50 HCs</td>
<td>Serum levels of IL2 and its receptor (using ELISA)</td>
<td>No group difference with regard to IL2 or its receptor level Correlation NA</td>
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<td>Flachenecker et al</td>
<td>26 fatigued MS patients; 11 unfatigued MS patients; cohort characteristics, 29 RR, 8 SP, 27F/10M, 54% treated</td>
<td>Serum mRNA expression of IFNγ, TNFα, and IL10 (using RT-PCR)</td>
<td>Higher TNFα (but not IFNγ or IL10) mRNA expression among fatigued patients Correlation analysis between fatigue and cytokine mRNA expression NA</td>
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<tr>
<td>Heesen et al</td>
<td>23 fatigued MS patients (19 RR, 3 SP, 1 PP, 18F/5M, 60.9% treated); 25 HCs (sex-matched, 20F/5M)</td>
<td>Whole-blood stimulatory capacity for TNFα, IFNγ, and IL10 (using ELISA); cognitive task to examine the immunoresponse (cytokines) to psychological stress</td>
<td>No significant group difference in baseline cytokines Blunted response of IFNγ among MS patients following psychological stress (no group difference in TNFα or IL10 responses) Correlation between fatigue and cytokine levels</td>
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<td>Heesen et al</td>
<td>15 fatigued MS patients (6 RR, 8 SP, 1 PP, 10 F/6 M, 66.7% treated); 15 fatigued MS patients (11 RR, 2 SP, 2 PP, sex-matched, 9F/6M, 60% treated)</td>
<td>Whole-blood stimulatory capacity for TNFα, IFNγ, and IL10 (using ELISA)</td>
<td>Higher levels of TNFα and IFNγ (but not IL10 levels) in fatigued MS patients Correlation between fatigue scores of TNFα and IFNγ</td>
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<tr>
<td>Gold et al</td>
<td>44 MS patients (all RR, all female, 59% treated)</td>
<td>Serum intracellular levels of cytokines IFNγ and TNFα (using flow cytometry)</td>
<td>Frequency of IFNγ-producing CD8+ T cells predicted of fatigue scores (regression analysis)</td>
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<td>Pokryszko-Dragan et al</td>
<td>20 fatigued MS patients; 20 unfatigued MS patients; cohort characteristics, 30 RR, 10 SP, 30F/10M, untreated; 25 HCs (sex NA)</td>
<td>Levels of IFNγ (using flow cytometry)</td>
<td>Higher IFNγ production among fatigued MS A trend toward correlation between fatigue and IFNγ</td>
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<tr>
<td>Malekzadeh et al</td>
<td>21 fatigued MS patients (15 RR, 5 PP/SP, 1 missing, 10F/7M, 47.6% treated); 14 unfatigued MS patients (11 RR, 3 PP/SP, sex-matched, 10F/4M, 50% treated)</td>
<td>Self-reported checklist: individual strength, fatigue subscale</td>
<td>No group differences with regard to variables measured Association between fatigue and IL6 levels (regression analysis)</td>
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Table 2 (Continued)

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<th>Method</th>
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<td>Mulero et al51</td>
<td>Seven fatigued MS patients (all RR, 5F/2M, 85.7% treated); 7 HCs (details NA)</td>
<td>MFIS</td>
<td>Whole-blood gene expression (using microarrays and RT-PCR)</td>
<td>Activation of IFN-response genes among fatigued MS patients Correlation NA Higher IFNγ, IL6, TNFα, IL17, and IL22 among MS patients In vivo: correlation between fatigue and each of IL6 and TNFα and a trend toward a correlation with IFNγ In vitro: correlation between fatigue and of IL6, TNFα, IFNγ, and IL22 levels</td>
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<tr>
<td>Alverenga-Filho et al57</td>
<td>18 MS patients (all RR, 15F/3M, untreated); 10 HCs (age-matched 8F/2M)</td>
<td>FSS</td>
<td>In vivo and in vitro assessment of peripheral levels of IL6, IL10, IL21, IL22, IL17, TNFα, and IFNγ (using ELISA)</td>
<td>Higher IL6 and TNFα levels in fatigued MS patients In vivo: correlation between fatigue and IL6 and TNFα levels and a trend toward a correlation between fatigue and IFNγ In vitro: correlation between fatigue and IL1β, IL6, IL17, IL22, and IL23 levels</td>
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<td>Akcali et al58</td>
<td>15 fatigued MS patients (all RR, 11F/4M, untreated); 15 unfatigued MS patients (all RR, sex-matched, 10F/5M, untreated)</td>
<td>FSS, NFI-MS</td>
<td>Serum IL1β, TNFα, IL35, IL2, and IL10 (using ELISA)</td>
<td>No group differences between fatigued and unfatigued MS patients No group differences between fatigued and unfatigued patients for any measure No correlation between fatigue and any cytokines studied</td>
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<td>Yaldizli et al53</td>
<td>26 fatigued MS patients (14F/12M); 28 unfatigued MS patients (15F/13M); cohort characteristics, all RR, 87.1% treated; 26 HCs (13F/13M); sex-matched groups</td>
<td>MFIS</td>
<td>Lymphocyte subsets in peripheral blood mononuclear cell cultures (using flow cytometry); suppressive function of regulatory T cells (using antigen stimulation)</td>
<td>No difference in leukocyte and lymphocyte subsets, including regulatory T cells between fatigued and unfatigued MS patients The entire patient group tended to have lower suppressive regulatory T-cell activity compared to HCs, with no differences between fatigued and unfatigued patients Correlation NA</td>
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<tr>
<td>Pokryszko-Dragan et al59</td>
<td>20 fatigued MS patients; 20 unfatigued MS patients; cohort characteristics, 30 RR, 10 SP, 30F/10M, untreated; 25 HCs (sex NA)</td>
<td>MFIS, FSS</td>
<td>Percentage of IFNγ-positive CD3+CD4+ T lymphocytes (using flow cytometry)</td>
<td>No group difference with regard to percentage of IFNγ-positive CD3+CD4+ T lymphocytes No correlation between fatigue and percentage of IFNγ-positive CD3+CD4+ T lymphocytes</td>
</tr>
<tr>
<td>Giovannoni et al58</td>
<td>38 MS patients (16 RR, 9 SP, 13 PP, 17F/21M, all untreated)</td>
<td>FQS, FSS</td>
<td>Serum CRP and sICAM-1 levels; urinary neopterin excretion (measured daily for 2 weeks)</td>
<td>Patients with raised serum CRP had higher FSS (but not FQS) scores than patients with normal CRP levels No correlation between fatigue (FSS, FQS) and any variable measured Correlation analysis NA</td>
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<td>Flachenecker et al54</td>
<td>26 fatigued MS patients; 11 unfatigued MS patients; cohort characteristics, 29 RR, 8 SP, 27F/10M, 54% treated</td>
<td>FSS</td>
<td>Serum ESR</td>
<td>No group differences in ESR values Correlation analysis NA</td>
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<tr>
<td>Adamczyk-Sowa et al59</td>
<td>102 MS patients (85 RR, 17 PP/SP, 67F/35M, 79.4% treated); 20 HCs (sex-matched)</td>
<td>MFIS</td>
<td>Plasma lipid hydroxyperoxides and homocysteine concentrations</td>
<td>Higher lipid-hydroxyperoxide levels among MS patients compared to HCs No correlation between fatigue and biochemical measures</td>
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</table>

Abbreviations: ELISA, enzyme-linked immunosorbent assay; ESR, erythrocyte-sedimentation rate; F, female; FSS, Fatigue Severity Scale; FQS, Fatigue Questionnaire Scale; HCs, healthy controls; M, male; MFIS, Modified Fatigue Impact Scale; MS, multiple sclerosis; NA, not available; NFI, Neurological Fatigue Index; PP, primary progressive; RR, relapsing–remitting; RT-PCR, real-time polymerase chain reaction; SP, secondary progressive.
Conclusion

This work has evaluated the contribution of central and peripheral inflammatory processes to MS fatigue. Among the selected studies dealing with central inflammatory and neuroendocrine processes, an inconsistency existed regarding the relationship between fatigue and orexin A system (present in one study, absent in one), the HPA axis (present in two studies, absent in five studies), and some CSF markers (present in one study assessing cytokines, absent in one study assessing cellular, humoral, and other CSF parameters). As for peripheral markers, although there was scarcity in the available data, serum proinflammatory cytokines (ie, IL-6, TNF-α, and IFN-γ) seemed to be associated with MS fatigue. However, given the existence of some conflicting data in this domain, such an association merits further investigation. Finally, concerning T-cell population (ie, CD3+CD4+ T lymphocytes or regulatory T cells) or peripheral markers of inflammation (ie, CRP, erythrocyte-sedimentation rate, and soluble ICAM1), few data were available, and these studies failed to find a link between MS fatigue and these measures. It is also worth noting that studies differed greatly in the clinical characteristics of their cohorts, especially concerning treatment profiles. The fact that MS treatment can modulate the inflammatory milieu would stand behind the differences observed in study outcomes, with studies including untreated patients yielding positive results on the relationship between fatigue and inflammation. Of note, several trials have documented differences in immune/inflammatory profiles between treated and naïve MS patients. That is to say, downregulation of proinflammatory cytokines was observed among MS patients treated with disease-modifying therapies such as IFNβ, glatiramer acetate, dimethyl fumarate, fingolimod, natalizumab, and teriflunomide.

Another issue to consider is the possible impact of MS treatments on fatigue per se. Few reports are available on this matter. In a cross-sectional study, higher fatigue rates were observed among MS patients treated with IFNβ or glatiramer acetate compared to age- and sex-matched patients receiving natalizumab. In other works, rituximab seemed to induce fatigue in MS patients, natalizumab appeared to improve fatigue, and fingolimod did not seem to modify symptom severity. Therefore, more research is needed to understand the potential effects of MS therapies on fatigue perception and cytokine profiles.

Another difference among studies concerned fatigue scales, which consisted of the FSS, MFIS, FSMC, visual analogue scale for fatigue, eleven-item Fatigue Scale, Neurological Fatigue Index – MS, self-reported checklist – individual strength (fatigue subscale), and Fatigue Questionnaire Scale. This adds more difficulty in comparing study outcomes. While some of these scales (eg, FSS) mainly address the physical component of fatigue, other scales (eg MFIS) reflect the physical, and, the cognitive and psychosocial dimensions of this symptom. This difference might not have had a large impact on group differences (fatigued versus unfatigued) but may have affected the correlation between fatigue severity and cytokine levels and could partly explain the discrepancies observed among studies.

Using different immunological techniques (ie, ELISA, RT-PCR, genetic analysis, flow cytometry, and electrochemiluminescence-based multiplex immunoassay) might have been behind interstudy differences, particularly those evaluating peripheral cytokines. Another point to consider is the relationship between sex, hormones, and immunodysregulation. Like many autoimmune diseases, MS is more prevalent in women than men, and hormones seem to exert an immunomodulatory effect and might influence damage repair in the CNS. Interestingly, sex dysmorphism was observed with regard to cytokine production in MS patients. In this context, it is of importance to note that although some studies controlled for sex effects by including sex-matched controls, only eight studies enrolled cohorts predominantly, or exclusively, composed of female patients or did not provide sufficient details on the matter. Therefore, future work could benefit from comparing fatigue and cytokine production between male and female patients.

Moreover, studying the impact of environmental, genetic, and epigenetic MS risk factors on MS fatigue would be of great interest. These factors include ultraviolet-radiation exposure, vitamin D intake, smoking, dietary, and exercise habits, and body-mass index. It is also of importance to control for some confounders that occur frequently in MS and can impact MS fatigue. These include physical disability, emotional symptoms, and sleep disorders. There is still a long way to go to define the utility and place of the aforementioned measures in clinical wards. Future large-scale studies are critically needed to conclude on this matter and would benefit from comparing the relationship between fatigue and inflammation in patients with different disease phenotypes (RR vs primary progressive vs secondary progressive) with and without disease-modifying drugs. From this perspective, applying different measures might help to decide on the optimal target to serve as an immunological surrogate of MS fatigue. Facing the subjective nature of fatigue scales, developing objective biological markers for fatigue, as those visited here, would be of great help.
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References


