Abstract  Treatment of multiple sclerosis (MS) is challenging: disease-modifying treatments (DMTs) must both limit unwanted immune responses associated with disease initiation and propagation (as T and B lymphocytes are critical cellular mediators in the pathophysiology of relapsing MS), and also have minimal adverse impact on normal protective immune responses. In this review, we summarize key preclinical and clinical data relating to the proposed mechanism of action of the recently approved DMT teriflunomide in MS. Teriflunomide selectively and reversibly inhibits dihydro-orotate dehydrogenase, a key mitochondrial enzyme in the de novo pyrimidine synthesis pathway, leading to a reduction in proliferation of activated T and B lymphocytes without causing cell death. Results from animal experiments modelling the immune activation implicated in MS demonstrate reductions in disease symptoms with teriflunomide treatment, accompanied by reduced central nervous system lymphocyte infiltration, reduced axonal loss, and preserved neurological functioning. In agreement with the results obtained in these model systems, phase 3 clinical trials of teriflunomide in patients with MS have consistently shown that teriflunomide provides a therapeutic benefit, and importantly, does not cause clinical immune suppression. Taken together, these data demonstrate how teriflunomide acts as a selective immune therapy for patients with MS.

1 Introduction

Multiple sclerosis (MS) is a chronic, progressive demyelinating disease of the central nervous system (CNS). MS typically emerges in young adulthood, and its incidence is highest in Northern Europe and North America, where it occurs in up to 1 in 1,000 individuals [1, 2]. In approximately 85% of cases, MS initially manifests as a relapsing–remitting form (RRMS), which is characterized by episodes of neurological worsening followed by at least partial recovery [2].

Disease-modifying treatments (DMTs) that reduce damage to the CNS are being investigated for the treatment of patients with MS. CNS damage is assumed to result from disturbances in immune tolerance networks [3]. Multiple perivascular inflammatory foci are seen in the CNS of patients with MS, and these become sites of demyelination and axonal injury [4]. The lesions are associated with infiltrating T cells and monocytes, and occasional B cells and plasma cells: T cells may target neurons directly [5, 6]. Additionally, there is evidence for subpial cortical injury, which appears not to be in perivascular distribution [7, 8], implicating further as yet unidentified mechanism(s).
Given the dominant role of abnormal immune activation in MS pathogenesis, DMTs must limit MS-associated immune responses to be effective. On the other hand, in order not to compromise responses to pathogens, DMTs should have minimal effects on normal protective immune responses.

Teriflunomide, a once-daily oral immunomodulatory DMT, is approved in several regions, including the United States and the European Union, for the treatment of RRMS; further regulatory reviews are ongoing in several other territories. Approvals were supported by placebo-controlled phase 3 clinical trials of teriflunomide that have demonstrated a favourable benefit/risk profile in this disease [9, 10]. In this review, we describe the hypothesized primary mechanism of action (MoA) of teriflunomide together with the preclinical evidence supporting this hypothesis in the context of MS pathogenesis. We also discuss the clinical evidence supporting preservation of protective immunity during teriflunomide treatment. Finally, we consider the place of teriflunomide in an expanding armamentarium of MS therapies.

2 MS Immunopathogenesis

MS is hypothesized to be primarily a T-helper cell-mediated autoimmune disease. Mounting evidence also supports the involvement of various other cells of the immune system. Immune cells cause demyelination and axonal/neuronal injury, and MS disease progression is considered to result partly from degenerative mechanisms as well as from ongoing (CNS-compartmentalized) inflammatory activity likely involving both T and B cells [6, 11–13].

2.1 T Cells

Various subsets of T cells exist, which are classified based on the expression of cell surface markers CD4 or CD8, and by their cytokine-secreting capacities. During a normal immune response, CD8+ T cells kill infected cells, while in MS they can mediate damage to CNS cells. CD4+ T cells, sometimes called “helper T” or “Th” cells, secrete cytokines that influence the activities of other immune cells, such as antibody-producing B cells. The roles of different immune cells in MS are illustrated in Fig. 1. According to the autoimmune hypothesis of MS pathogenesis, T cells reactive to CNS proteins (such as myelin basic protein [MBP] and myelin oligodendrocyte glycoprotein [MOG], among others [14]) must first be activated in the periphery before they gain the potential to cause damage within the CNS. Activated T cells up-regulate adhesion molecules and chemokine receptors that allow them to more efficiently cross the blood–brain barrier (BBB) [15]. Animal studies suggest that the potential to cross the BBB may depend on the specific antigen for each T cell being presented by cells of the vascular endothelium—this interaction initiates local BBB compromise and allows the T cell to enter into the CNS [16, 17]. The interleukin (IL)-17-secreting CD4+ T-cell subset (Th17 cells) has been implicated in animal models of MS and Th17 cells up-regulate the CCR6 receptor upon activation to gain entry into the CNS [18]. Recent data also indicate that neuronal activity can regulate entry of T cells into the spinal cord [19].

Early evidence supporting CD4+ T-cell involvement in MS came from experimental autoimmune encephalomyelitis (EAE) rodent models. EAE is induced by immunization with CNS-derived proteins or peptides, such as MBP [20, 21]. Transfer of MBP-specific CD4+ T cells from EAE animals can be sufficient to induce EAE in recipient animals, so long as the transferred cells are activated [22]. Transfer of T cells reactive to a mycobacterium derivative, which is used as an adjuvant, did not induce EAE in this experiment [22]. Further support for the involvement of CD4+ T cells in MS comes from the reported increased risk of MS associated with expression of specific major histocompatibility complex (MHC) class II alleles involved in antigen presentation to T lymphocytes [23].

Th1-type CD4+ T cells, characterized by production of interferon (IFN)γ, IL-2, and tumor necrosis factor alpha (TNFα), were initially believed to be the main drivers of MS [24, 25]. More recently, the role of Th17 cells has been recognized [26]. Interruption of Th17 differentiation using IL-23-blocking antibody in EAE mice restricts development of EAE [27], and IL-17 levels are elevated in blood and brain tissue of patients with MS [28]. Abrogation of disease activity in patients with aggressive MS following ablative chemotherapy and hematopoietic stem cell transplantation was associated with a diminished Th17 response, despite the reemergence of myelin-reactive Th1 and Th2 cells [29].

Various lines of evidence also indicate a role for CD8+ T cells in MS pathogenesis [13, 16, 30]. CD8+ T cells are able to cause EAE [31] and are present in MS lesions in patients [32], and some patients with MS have increased numbers of circulating MBP-reactive CD8+ T cells [33, 34]. Dysfunction or impaired maturation of both natural and inducible regulatory T cells (Treg), a type of CD4+ T cell, have been described in MS. Because Treg are critical for the maintenance of immune tolerance, this dysfunction or impaired maturation could possibly contribute to the disrupted tolerance found in patients with MS [35, 36].

2.2 B Cells

B-cell numbers are increased within the CNS of patients with MS [37], and B cells are considered to play an
important role in MS pathogenesis through both antibody-dependent and antibody-independent functions. Antibody abnormalities within the CNS of patients include increased levels of cerebrospinal fluid (CSF) immunoglobulin (Ig) and Ig synthesis rates, as well as the presence of CSF-restricted oligoclonal bands. The target(s) of such antibodies have not been fully characterized, although several antigens have been suggested [38, 39].

B cells described within the MS brain appear to have undergone maturation within the CNS, possibly within specialized tertiary lymphoid tissue described in the pia/meninges of patients [40, 41]. Lineage analysis has shown that B cells from patients with MS can mature on either side of the BBB and traffic across it [42].

Independently of their antibody-producing capability, B cells may also affect T-cell activity in MS through cytokine production. Aberrant profiles of both pro-inflammatory (lymphotoxin, TNFα, IL-6) and anti-inflammatory (IL-10) cytokine responses have been observed in B cells from patients with MS [43]. B-cell activation in patients with MS can thus drive a pro-inflammatory T cell response, potentially resulting in new disease activity [44]. This hypothesis is supported by the observation that following B-cell depletion with the anti-CD20 monoclonal antibody rituximab, pro-inflammatory T-cell responses in patients with MS are diminished [45].

Regulatory B cells (Breg) have been shown to inhibit the progression of EAE in mice [46]. Evidence from patients also supports a role for Breg in MS, or an effect of MS on Breg. Increased numbers of Breg in MS have been reported in patients with MS [47, 48], as have deficiencies in Breg functions, such as in IL-10 secretion [43, 44, 49].

2.3 Other Cells of the Immune System

Myeloid cells such as dendritic cells (DCs), either through their cytokine-secreting capacity or through their antigen-presenting or cell contact-mediated properties, are also likely involved in MS. In patients with MS, myeloid cells (both resident microglia and infiltrating macrophage/DCs) are abundant in CNS lesions and exhibit alterations in phenotype and function (expressing more activation markers and producing greater amounts of pro-inflammatory cytokines), supportive of a role in MS pathogenesis [50].

Roles in MS for other regulatory immune cells, including subsets of natural killer (NK) cells and resident CNS cells, such as astrocytes, have been proposed because their numbers or functions are altered in patients with MS or in response to immunomodulatory therapy. For example, NK cell cytolytic activity is reduced in patients with MS, and the DMT fingolimod is believed to influence the balance of NK cell subsets [51]. Astrocytes perform multiple roles in the evolution of changes encountered in MS depending upon the stage of development of their associated lesion [52], while microglia, also found in MS lesions, support the activity of CNS-reactive T cells [53].

3 Teriflunomide Proposed MoA: Overview of Current Hypothesis

The primary MoA of teriflunomide in MS is hypothesized to relate to its effects on the proliferation of stimulated lymphocytes. Upon activation, lymphocytes undergo
division via the cell cycle, that comprises an S (‘synthesis’) phase (during which deoxyribonucleic acid [DNA] is copied), and M (‘mitosis’) phase (during which the dividing cell splits into daughter cells), separated by G (‘gap’) phases [54]. DNA synthesis in S phase requires the provision of new building blocks, including pyrimidine and purine bases. Teriflunomide blocks de novo pyrimidine synthesis by specific non-competitive, reversible inhibition of the mitochondrial enzyme dihydro-orotate dehydrogenase (DHODH) [55–57], an enzyme expressed at high levels in proliferating lymphocytes [57, 58]. Teriflunomide-mediated blockade of pyrimidine synthesis interrupts the cell cycle in S phase and exerts a cytostatic effect on proliferating T and B cells, limiting their involvement in the inflammatory processes involved in the MS pathogenesis (Fig. 2) [59].

Cells that do not proliferate in response to activation, such as resting lymphocytes, can self-renew through homeostatic proliferation [60], in which pyrimidine requirements are met through the salvage pathway [61]. Thus, teriflunomide inhibition of DHODH only affects activated, rapidly proliferating lymphocytes (Fig. 2). Rapid, antigen-induced proliferative expansion is unique to lymphocytes [62]. Other proliferating cells, such as gastrointestinal mucosal cells, express lower levels of DHODH [58] and are not specifically stimulated for rapid expansion; therefore, teriflunomide is far less able to inhibit their proliferation.

Of note, teriflunomide is not a nucleotide analogue, and the Ames test shows that it is not DNA intercalating [63]. Furthermore, teriflunomide has no effect on cell viability [64], in contrast to mitoxantrone (Table 1) and the range of immunosuppressive drugs (Table 2) used in MS treatments that interact with DNA (azathioprine, cyclophosphamide) and/or induce apoptosis (azathioprine, cyclophosphamide, methotrexate, mycophenolate mofetil).

4 Supportive In Vitro Evidence

4.1 Lymphocyte Proliferation

A variety of in vitro studies have demonstrated that teriflunomide causes cell cycle arrest in early S phase without apoptosis; these include studies of a human B-cell line stimulated with CpG oligonucleotides and IL-2 [65] and of mitogen-stimulated rat lymphocytes [66]. Similarly, using cultured primary human peripheral blood mononuclear cells (PBMCs) from normal donors, Li et al. [64] demonstrated that teriflunomide inhibits proliferation of CpG-stimulated B cells and anti-CD3-stimulated T cells (both CD4+ and CD8+ T cells) in a dose-dependent manner without causing cell death (Fig. 3a–c). In each case, the inhibitory effects of teriflunomide on DHODH and de novo pyrimidine synthesis were completely abrogated by the addition of exogenous uridine [64].

In B cells, decreased proliferation in the presence of teriflunomide was associated with a decrease in levels of cyclin-dependent kinase 2 (cdk2), which is required for S phase [67, 68]. These observations are consistent with reports that pyrimidine availability controls progression from early to intermediate S phase [69].

Intriguingly, the anti-proliferative effects of teriflunomide appear to be most pronounced in T cells bearing T-cell receptors (TCRs) with a high affinity for a presented peptide antigen [70]. Individual T cells each express a unique TCR, assembled by recombination at multiple TCR-encoding loci. Each TCR has a different affinity for a peptide antigen, and the mature T cell repertoire is selected on the basis of T-cell avidity for a peptide antigen [71]. Thus, this repertoire includes T cells with a range of avidities for a peptide antigen. In an in vitro setting, using CD4+ T cells specific for an ovalbumin peptide as a model,
Table 1  Reported MoA of MS therapies

| Drug                        | MoA (included in label)                                                                 | MoA (evidence for additional mechanisms)                                                                 |
|-----------------------------|----------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------|
| **Approved therapies**     |                                                                                        |                                                                                                          |
| Alemtuzumab                 | Binds CD52, expressed at high levels on the surface of T and B cells, to cause their depletion from circulation | Repopulation of lymphocytes rebalances the immune system to reduce MS disease activity                    |
| Dimethyl fumarate (BG-12)   | Protects against oxidative stress-induced cellular injury and loss in neurons and astrocytes via up-regulation of an Nrf2-dependent antioxidant response | Neuroprotective effects are immune mediated                                                                |
| Fingolimod                 | Binds the S1P receptor, causing lymphoid cell retention in secondary lymphoid tissue   | Promotes a Th1–Th2 shift in cytokine response                                                                 |
| Glatiramer acetate (GA)     | Modifies many immune processes                                                         | Promotes a Th1–Th2 shift in T-cell responses through effects on DCs                                       |
| Interferon beta (IFNβ)      | Decreases T-cell activation through binding to IFN receptors                            | Promotes migration of Th2 cells into the CNS                                                               |
| Mitoxantrone                | Intercalates with DNA and causes single- and double-strand breaks, and inhibits DNA repair through inhibition of DNA topoisomerase II | May exert direct neurotrophic effects and promote remyelination through induction of BDNF                  |
| Natalizumab                | Targets integrin α4β1 (VLA-4), restricts T-cell and NK-cell extravasation and migration into the CNS | May down-modulate VLA-4 ligand VCAM-1                                                                      |
| Teriflunomide               | Inhibits proliferation of activated T and B lymphocytes through inhibition of DHODH     | Mild pro-inflammatory stimulatory effect on CD4+ helper T cells, leading to increases in IL-2, IFN-γ, and IL-17 expression |
| **Therapies in phase 3 development** |                                                                                         |                                                                                                          |
| Daclizumab                  | Promotes differentiation and expansion of CD56 (bright) immunoregulatory NK cells      | Promotes differentiation and expansion of autoreactive T cells through binding to CD25                      |

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| Table 2 | MoA of immunosuppressive therapies occasionally or historically used off-label in patients with MS | Safety |
| --- | --- | --- |
| **Drug** | **MoA (included in label)
MoA (evidence for additional mechanisms)** | **Safety** |
| Azathioprine [169] | A purine analogue that is incorporated into DNA and blocks the de novo purine synthesis pathway [170] | Induces apoptosis in stimulated T cells [107] | Over 10% of patients experience leukopenia, cancer risk increases with treatment duration and cumulative dose [171] |
| Cyclophosphamide [172] | Active metabolites of cyclophosphamide alkylate DNA during cell division, impairing DNA synthesis and leading to apoptosis in actively proliferating cells [173] | Inhibits the suppressive capability of Treg (at low doses) [174] | Side effects include risk of malignancy [175] |
| Methotrexate [176] | Competitive inhibitor of DHFR, leading to a deficiency of nucleic acid precursors and impaired DNA synthesis [177] | Impairs lymphocyte activation and adhesion [178] | Rare cases of severe pancytopenia [180] |
| Mycophenolate mofetil [181] | Reversible inhibitor of IMD type II, an enzyme mainly found in lymphocytes and responsible for de novo synthesis of the purine nucleotide guanine [182] | Impairs DNA synthesis and increases apoptosis in activated T cells and impairs proliferation in B and T cells [183] | Inhibits antigen presentation by DCs [183] | Risk of transitory leukopenia, digestive disorders, and benign infections [184] |

**MoA mechanism of action, MS multiple sclerosis, Th T helper, DC dendritic cell, CNS central nervous system, NK natural killer, MHC major histocompatibility complex, MBP myelin basic peptide, BDNF brain-derived neurotrophic factor, IFN interferon, APC antigen-presenting cell, DNA deoxyribonucleic acid, VLA very late activation, VCAM vascular cell adhesion molecule, IL interleukin, DHODH dihydro-orotate dehydrogenase, MCP-1 monocyte-derived chemotactic protein 1, Treg T regulatory cell, TGF transforming growth factor beta, RRMS relapsing-remitting multiple sclerosis**

* The MoA of these therapies is not completely understood and the listed mechanisms are hypotheses
* Included in labels where drug is approved
* Trials in patients with RRMS as of October 2013

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**Table 1 continued**

| Drug | MoA (included in label)
MoA (evidence for additional mechanisms) |
| --- | --- |
| Laquinimod | Decreases Treg numbers in the CNS [162] |
| | Decreases lymphocyte migration [162, 163] |
| | Decreases IL-17 production [163] |
| | Increases B-cell expression of markers of regulatory activity, such as CD25, IL-10, and CD86 [164] |
| | Increases IL-10 and TGFβ production by T cells (B-cell mediated) [164] |
| | Augments BDNF expression [162] |
| | Induces type II monocytes [165] |
| | Causes a Th1–Th2 shift in CD4+ T cells [166] |
| Ocrelizumab | Binds CD20 to cause complement-dependent lysis of and/or antibody-dependent cytotoxicity towards B cells [167, 168] |

**MoA mechanism of action, MS multiple sclerosis, Th T helper, DC dendritic cell, CNS central nervous system, NK natural killer, MHC major histocompatibility complex, MBP myelin basic peptide, BDNF brain-derived neurotrophic factor, IFN interferon, APC antigen-presenting cell, DNA deoxyribonucleic acid, VLA very late activation, VCAM vascular cell adhesion molecule, IL interleukin, DHODH dihydro-orotate dehydrogenase, MCP-1 monocyte-derived chemotactic protein 1, Treg T regulatory cell, TGF transforming growth factor beta, RRMS relapsing-remitting multiple sclerosis**

* The MoA of these therapies is not completely understood and the listed mechanisms are hypotheses
* Included in labels where drug is approved
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Proliferation of high-avidity T cells was strongly inhibited by teriflunomide, whereas proliferation of low-avidity T cells was less affected [70]. This is of particular interest because high-avidity T cells are thought to mediate many autoimmune diseases, including MS [72]. A selective effect of teriflunomide on autoreactive high-avidity T cells would leave lower-avidity T cells available for normal responses to pathogens.

**Fig. 3** In vitro, preclinical, and clinical evidence for the selective immunomodulatory MoA of teriflunomide. Teriflunomide inhibits the proliferation of a CD4+ and b CD8+ T-cell subsets, and c B cells, in vitro, in a DHODH-dependent (uridine-reversible) manner. Graphs show the degree of inhibition of proliferation in response to anti-CD3 antibody (a, b) or CpG oligonucleotide (c), measured by CFSE dye dilution and flow cytometry; data presented as mean ± standard error. d In the Dark Agouti EAE rat model of MS, teriflunomide treatment attenuates the number of T cells (identified by anti-CD3 staining and flow cytometry) in cervical spinal cord at all phases of disease; data presented as least square mean ± standard error of the mean. e White blood cell counts in patients treated with teriflunomide 14 mg remain within the normal range and stabilize after the first 3 months of treatment. f Responses to seasonal influenza vaccine, by influenza strain, in patients treated with teriflunomide 14 mg. Dotted line shows European criteria for vaccine efficacy (70 % of patients with post-vaccination titres ≥40), error bars show 90 % confidence interval. MoA mechanism of action, EAE experimental autoimmune encephalomyelitis, DHODH dihydro-orotate dehydrogenase, CFSE carboxyfluorescein succinimidyl ester, SEM standard error of the mean. Reproduced with permission from [64] (a–c), [81] (d), [94] (f).
4.2 Immune Cell Function

In vitro studies have suggested that teriflunomide treatment can impede formation of the immune synapse (necessary for activation of T cells) via impaired integrin activation through a DHODH-independent mechanism [73]. Decreased protein aggregation caused by teriflunomide may further weaken the interaction between T cells and antigen-presenting cells [74].

Teriflunomide treatment significantly decreased release of IL-6, IL-8, and monocyte chemotactic protein-1 (MCP-1) from lipopolysaccharide (LPS)-stimulated human PBMCs, attributed to the monocyte compartment. These effects were not reversible by uridine and therefore unlikely to be DHODH-dependent [64]. This decrease in pro-inflammatory cytokine release is consistent with earlier reports from rodent studies of impaired Th1 differentiation and enhanced Th2 differentiation in the presence of teriflunomide [75, 76]. Li et al. also showed that teriflunomide did not impair LPS-induced maturation of monocyte-derived DCs and had no impact on the ability of LPS-matured DCs to induce allogeneic T-cell responses, suggesting that teriflunomide does not broadly impair the capacity of DCs to regulate adaptive immunity [77].

Of note, many of the earlier studies reporting DHODH-independent effects of teriflunomide used relatively high in vitro concentrations of the drug (~100 μM) when the half maximal concentration for inhibition of DHODH was in the high nanomolar range (657 nM in human splenocytes) [55]. Thus, some of the phenomena observed in these studies, such as modulation of tyrosine kinase activity (reviewed in Claussen MC, Korn T [78]), may not be pertinent to teriflunomide’s in vivo activity in patients with MS.

5 In Vivo (Preclinical) Evidence to Support Current Hypothesis of Teriflunomide MoA

5.1 Rat EAE Model

The Dark Agouti (DA) rat EAE model mimics the inflammatory features of RRMS [79] and has been used in several studies to elucidate the actions of teriflunomide. DA rats develop EAE after immunization with a syngeneic spinal cord homogenate and Freund’s adjuvant. Immunization induces a range of neurological symptoms progressing from tail paralysis, disturbed gait, and paresis to paralysis. These defects are accompanied by demyelinating inflammatory lesions in the spinal cord with infiltrating immune cells.

Administration of either prophylactic or therapeutic (post-disease onset) teriflunomide to DA EAE rats improved disease outcomes, delayed disease onset, and reduced maximal and cumulative disease scores [80]. Histopathology of teriflunomide-treated DA EAE rat spinal cord demonstrated a reduction in demyelination and axonal loss of up to 90%, with a decrease in inflammation of up to 70% [80]. Treatment with teriflunomide also attenuated levels of spinal cord-infiltrating T cells (Fig. 3d), NK cells, macrophages, and neutrophils [81]. It is worth noting that in patients with rheumatoid arthritis (RA) treated with leflunomide (the parent compound of teriflunomide), infiltration of macrophages and T cells into the synovial tissue was reduced, and expression patterns of lymphocyte adhesion molecules and matrix metalloproteinase were altered, suggesting an effect of teriflunomide on lymphocyte migration [82]. Teriflunomide treatment also appeared to protect against oligodendrocyte cell death in the CNS of DA EAE rats [83].

Teriflunomide treatment also improved sensory and motor functional outcomes in DA EAE rats. Studies of electrophysiological somatosensory-evoked potential showed that prophylactic teriflunomide treatment prevented both a decrease in waveform amplitude and an increase in the latency to waveform initiation in EAE animals. Therapeutic dosing had similar effects [80]. Measurement of transcranial magnetic motor-evoked potentials showed that teriflunomide prevented a delay in waveform latency and a decrease in waveform amplitude [84].

It is likely that observed reductions in maximal disease scores, decreased axonal loss, and decreased loss of nerve function in teriflunomide-treated DA EAE rats can be attributed to reduced inflammation in these animals, as a consequence of the direct effects of teriflunomide on immune cells. It remains to be determined whether local CNS effects may also contribute to the activity of teriflunomide.

Additional observations made in teriflunomide-treated DA EAE rats indicate that teriflunomide treatment at disease onset reversed immunopathological changes associated with the disease, including mitigation of reductions in spleen weight (spleen weights in teriflunomide-treated animals were only slightly lower than those observed in disease-free animals and significantly higher than those observed in untreated EAE animals), of increased levels of circulating monocytes at acute attack and relapse, and of increased ratios of CD4+/CD8+ T cells in the spleen at acute attack and remission [81].

5.2 Mouse Theiler’s Murine Encephalomyelitis Virus (TMEV) Model

Teriflunomide also showed efficacy in a mouse TMEV infection model used to simulate several pathophysiologic
features of MS. In susceptible mouse strains, TMEV induces biphasic disease. A week after infection, TMEV causes polioencephalomyelitis characterized by apoptosis of gray matter neurons. During the chronic phase, approximately 1 month after infection, the virus infects glial cells and macrophages and induces inflammatory demyelination with oligodendrocyte apoptosis and axonal degeneration in the white matter of the spinal cord [85]. Progression of neurological deficits in TMEV-infected mice is mitigated by therapeutic teriflunomide treatment, as shown by a higher neurological function index in teriflunomide-treated TMEV-induced demyelinating disease mice than in vehicle-treated mice [86].

Importantly, teriflunomide treatment in this viral model of MS only resulted in a mild delay in the production of antiviral antibodies and was associated with a transient, non-significant increase in viral load in the CNS 30 days post-infection. This finding supports the concept that protective immunity is maintained in the presence of teriflunomide, and also suggests that a significant increase in viral infections would not be expected in teriflunomide-treated patients with MS [87].

6 Clinical Evidence of Immune Competence and Protective Immunity in Patients Treated with Teriflunomide

Clinical data are consistent with evidence from preclinical in vivo studies suggesting that teriflunomide treatment does not significantly disrupt protective immunity, and its anti-inflammatory activity primarily affects the pathogenic immune processes associated with disease activity.

Pooled data from placebo-controlled studies of teriflunomide, including over 2400 patients from two phase 3 studies (TEMSO [NCT00134563] [9] and TOWER [NCT00751881] [10]) and one phase 2 study (NCT01487096) [88], showed that teriflunomide treatment was associated with a mean decrease in leucocyte counts of approximately 15% from baseline (Fig. 3e) [89]. However, mean absolute counts remained within the normal range, decreases occurred within the first 3 months of treatment, and levels did not change further with continuing treatment. In the groups treated with teriflunomide from this analysis, a low incidence of serious infections that was not significantly different from placebo was observed, and the relative risk of any infection was not increased with teriflunomide treatment. There were four serious infections considered opportunistic (opportunistic infections were broadly defined in this analysis, and generally considered to be infections that are only pathogenic in immunocompromised individuals); two in patients receiving placebo (one patient developed herpes zoster infection that resolved following treatment discontinuation, and another patient developed hepatitis C and cytomegalovirus [CMV] infection leading to permanent treatment discontinuation), and two in patients treated with teriflunomide 14 mg (one patient with a case of gastrointestinal tuberculosis who recovered following study drug discontinuation and anti-tuberculosis therapy, and a second patient who developed hepatitis with CMV who recovered after treatment discontinuation [discontinuation was protocol mandated due to increased alanine aminotransferase]) [89]. In long-term extensions of the phase 2 and TEMSO studies (up to 9 years of teriflunomide treatment) and consistent with the double-blind phases of the studies, there was no signal for increased malignancies or serious opportunistic infections [90, 91]. Taken together, these findings provide evidence that teriflunomide does not have a negative impact on protective immunity over the observed treatment period.

To investigate whether the generation of recall antigen responses was preserved under teriflunomide treatment, immune responses to the 2011–2012 seasonal influenza vaccination (including strains H1N1, H3N2, and B) were measured in patients with MS treated with IFNβ and in patients exposed to teriflunomide for at least 6 months (TERIVA, NCT014033760) [92]. Most patients had antibodies against each strain detectable at baseline, which was attributed to the 2011–2012 vaccine containing the same strains as the 2010–2011 vaccine. More than 90% of patients achieved post-vaccination antibody titres ≥40 for H1N1 and B in all groups. For H3N2, titres ≥40 were achieved in ≥90% of patients in the teriflunomide 7-mg and IFNβ groups, and in 77% of the teriflunomide 14-mg group, respectively (Fig. 3f). Geometric mean titre ratios (post-/pre-vaccination) were ≥2.5 for all groups and strains, except for H1N1 in the 14-mg group (2.3), confirming vaccine efficacy as per European guidelines [92, 93]. Therefore, the mild quantitative and qualitative change in the immune response to vaccination induced with teriflunomide was not clinically significant, and the ability of patients to mount protective vaccine responses to recall antigens while receiving teriflunomide was considered to be preserved [92].

Rabies vaccination was used as a tool to assess the impact of teriflunomide on the immune responses to a neoantigen. Healthy volunteers who had never been vaccinated before with rabies vaccine received teriflunomide or placebo for 1 month prior to vaccination; a slightly lower mean antibody response in the teriflunomide group than in the placebo group was observed. However, adequate seroprotection (antibody titre ≥0.5 IU/mL) was achieved in all subjects [94].

In summary, data on the clinical consequences of treatment indicate that teriflunomide—while slightly

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reducing peripheral leucocyte counts does not compromise clinically relevant immune competence. In keeping with this, responses to both recall and neoantigen vaccinations, though mildly reduced, remained sufficient to afford seroprotection.

7 Safety Considerations with Teriflunomide

As discussed previously, the primary action of teriflunomide is proposed to be specific to activated lymphocytes as these cells express high levels of DHODH and undergo rapid proliferative expansion [58, 62]. However, all patients with MS treated with any DMT may experience a range of side effects that are not necessarily related to the proposed primary MoA of that therapy [95]. In this regard, clinical trials of teriflunomide have demonstrated its manageable safety profile (Table 3) [96, 97].

Elevations of serum aminotransferases in clinical trials and rare cases of liver injury in the post-marketing setting have been documented in patients treated with leflunomide [100] so as a precaution, patients receiving teriflunomide are subject to regular hepatic monitoring during the first 6 months of treatment [101, 102]. In some locations, less frequent monitoring continues throughout subsequent treatment [102].

Leflunomide is associated with embryo-lethality and teratogenicity in rats and rabbits, and therefore is contraindicated in women of childbearing potential not using reliable contraception [99]. However, two prospective studies conducted by the Organization of Teratology Information Specialists (OTIS), have found no increase in the rate of major structural defects in newborns of women with RA exposed to leflunomide [103, 104]. Additionally, in the period 2001–2013, there were more than 2.3 million patient years of leflunomide use without any signal for teratogenicity (Genzyme, data on file). Based on the occurrence of teratogenicity and embryo-lethality in the offspring of teriflunomide-treated rats and rabbits, teriflunomide is also contraindicated in pregnant women, or women of childbearing potential not using reliable contraception [101]. Preclinical teriflunomide studies showed no evidence for mutagenicity or clastogenicity [63] and consistent with the OTIS findings for leflunomide, all newborns born to mothers or fathers who received teriflunomide had no structural or functional abnormalities at birth [105].

In the event of pregnancy, patients receiving teriflunomide must undergo an accelerated elimination

Table 3 Overview of the safety profile of teriflunomide in three placebo-controlled studies [96]

|                              | Placebo | Teriflunomide (7 mg) | Teriflunomide (14 mg) |
|------------------------------|---------|----------------------|-----------------------|
| Number (%) of patients       | n = 806 | n = 838              | n = 786               |
| Any TEAE                     | 697 (86.5) | 734 (87.6)            | 702 (89.3)            |
| Any serious TEAE             | 101 (12.5) | 107 (12.8)              | 109 (13.9)          |
| Any TEAE leading to study discontinuation | 56 (6.9) | 92 (11.0)              | 107 (13.6)         |
| Maximum intensity of TEAEs   |         |                      |                      |
| Mild                         | 218 (27.0) | 225 (26.8)             | 201 (25.6)          |
| Moderate                     | 374 (46.4) | 400 (47.7)             | 385 (49.0)          |
| Severe                       | 105 (13.0) | 109 (13.0)             | 116 (14.8)          |
| Most common TEAEs            |         |                      |                      |
| Alanine aminotransferase increased | 62 (7.7)  | 100 (11.9)              | 110 (14.0)         |
| Diarrhea                     | 63 (7.8) | 109 (13.0)             | 113 (14.4)          |
| Nausea                       | 63 (7.8) | 74 (8.8)               | 97 (12.3)           |
| Headache                     | 125 (15.5) | 156 (18.6)             | 128 (16.3)          |
| Hair thinningb               | 35 (4.3) | 90 (10.7)              | 111 (14.1)         |

Pooled data from TEMSO, TOWER, and Phase 2 Study [96]

TEAE treatment-emergent adverse event

a Occurring in ≥ 10 % of patients in any treatment group and ≥ 2 % in either teriflunomide group compared with placebo

b Medical Dictionary for Regulatory Activities preferred term: alopecia

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△ Adis
procedure until teriflunomide plasma concentrations fall below 0.02 mg/L, a level predicted to have minimal risk to the fetus [101].

8 Teriflunomide as Part of the Growing MS Therapeutic Armamentarium

The MoAs of therapies in development or approved for use in RRMS are summarized in Table 1. Additional therapies are sometimes used off-label in patients with MS (Table 2), though it is important to note that use of these immunosuppressant medications may be associated with serious side effects such as leukopenia or increases in the risk of malignancy.

The immunomodulation exerted by teriflunomide is seemingly selective in a manner that is particularly useful in the treatment of RRMS, a disease mainly driven by activated T and B lymphocytes. Importantly, a specific cytostatic effect on proliferating lymphocytes as a primary MoA is unique to teriflunomide. None of the agents currently in clinical trials or being marketed for MS possess a similar specificity for activated lymphocytes inherent in their MoA. Other agents rely on mechanisms such as direct cytotoxicity [106], induction of apoptosis [107], or exclusion of lymphocytes from the CNS [108–110] to reduce the numbers of lymphocytes available to contribute to MS pathogenesis.

9 Summary and Conclusions

Teriflunomide is known to reversibly inhibit DHODH, a mitochondrial enzyme specifically required for de novo pyrimidine biosynthesis in activated lymphocytes, and as such has no direct effects on DNA. In keeping with this primary MoA, teriflunomide acts to impair the proliferation of activated T and B cells, and reduces their ability to participate in a potentially damaging immune attack on the CNS.

Emerging in vitro and in vivo experimental data increasingly support the view that teriflunomide exerts a selective effect on the immune system that strikes a favorable balance between efficacy and safety. This selectivity and balance is reinforced in observations from studies in patients with MS. Phase 3 studies showed that teriflunomide can restrict the type of limited immune activation relevant to MS disease activity; trial results showed significant reductions in disability progression, relapse rates, and magnetic resonance imaging measures of disease activity [9, 111]. At the same time, vaccination studies in patients with MS or healthy subjects have shown that teriflunomide treatment does not limit the immune activation required for maintaining immune competence: the capacity to mount functional vaccine responses (to both recall antigens and neoantigens) is preserved. In addition, the comparable incidence of infections and malignancies observed with teriflunomide and placebo also indicates maintenance of functional immune competence. Teriflunomide can be considered as an immune therapy targeting pathogenic inflammatory processes involved in the MS disease process without significant compromise of the patient’s protective immune functions. Given the selectivity of its effects on the immune system, teriflunomide represents a valuable addition to the list of approved therapies available to treat patients with MS.

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