The Development of Cladribine Tablets for the Treatment of Multiple Sclerosis: A Comprehensive Review

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Abstract
Cladribine is a purine nucleoside analog initially developed in the 1970s as a treatment for various blood cancers. Due to the molecule’s ability to preferentially reduce T and B lymphocytes, it has been developed into an oral formulation for the treatment of multiple sclerosis (MS). The unique proposed mechanism of action of cladribine allows for the therapy to be delivered orally over two treatment-week cycles per year, one cycle at the beginning of the first month and one cycle at the beginning of the second month of years 1 and 2, with the potential for no further cladribine treatment required in years 3 and 4. This review summarizes the clinical development program for cladribine tablets in patients with MS, including the efficacy endpoints and results from the 2-year phase III CLARITY study in patients with relapsing–remitting MS (RRMS), the 2-year CLARITY EXTENSION study, and the phase III ORACLE-MS study in patients with a first clinical demyelinating event at risk for developing MS. Efficacy results from the phase II ONWARD study, in which cladribine tablets were administered as an add-on to interferon-β therapy in patients with RRMS, are also summarized. A review of all safety data, including lymphopenia, infections, and malignancies, is provided based on data from all trials in patients with MS, including the initial parenteral formulation studies. Based on these data, cladribine tablets administered at 3.5 mg/kg over 2 years have been approved across the globe for various forms of relapsing MS.

1 Introduction
Multiple sclerosis (MS) is a chronic, inflammatory, immune-mediated demyelinating and neurodegenerative disease of the central nervous system (CNS). It is one of the most common causes of serious neurological disability in young adults, and one of the most prevalent neurological disorders in the world [1]. In 2016, the estimated global prevalence of MS was 2.2 million, an increase of 10.4% from the age-standardized MS prevalence in 1990 [2]. Among the more than 2 million people with MS, more than 900,000 are thought to reside in the United States (US) [3]. About 85% of patients with MS present with relapsing–remitting MS (RRMS), which is characterized by periodic acute exacerbations of disease activity (relapses) punctuated by periods of clinical stability. Relapses may be associated with partial or complete recovery [4].

MS is associated with poor health-related quality of life and can have a profound effect on social functioning, employment status, and healthcare costs [5, 6]. The mean age at onset of symptoms is 30 years, and approximately 75% of patients are female [7]. Therefore, many patients are women of childbearing age, and MS can impact family planning due to risks associated with treatments that may affect pregnancy [8]. While there are no adequately controlled studies of disease-modifying drugs (DMDs) and pregnancy, there is a significant body of evidence on the safety of DMDs during pregnancy, generated primarily...
from clinical experience and pregnancy registries. Large interferon (IFN)-β registry studies have found no evidence for differences in infant size [9], congenital anomalies, or miscarriages [10], and recently updated IFN-β Summaries of Product Characteristics recommend to continue treatment during pregnancy and while breastfeeding if clinically needed [11, 12]. Glatiramer acetate is also generally considered to be safe for use during pregnancy [13, 14]. However, the labels for the majority of DMDs for patients with MS either state to use ‘only if clearly needed’ [15], to avoid these therapies unless the benefits outweigh the risks [16–19], or not to use at all, and a washout period is recommended prior to pregnancy [20–26]. Longitudinal studies indicate that life expectancy is shorter for patients with MS compared with the general population (approximately 7 years shorter in a Norwegian population studied over a 60-year period) [27], but rising prevalence without parallel changes in incidence, and a shift in peak age towards older age groups, suggest that survival is improving (based on Canadian data collected over 2 decades) [28]. An aging MS population may be associated with significant disability; in a self-reporting postal survey conducted in a Canadian cohort of MS patients aged > 55 years who had been living with MS for a mean of approximately 33 years, 28% either required a wheelchair or were bedridden [29].

Lymphocytes play a central role in the pathogenesis of MS, with the actions of both B and T cells in the periphery and in the CNS implicated from an early stage in the disease process. Immune cells activated in the periphery enter the CNS, facilitated by chemokines [30, 31]. Autoreactive T cells are present in acute CNS lesions in early MS [30, 32]; their reaction to myelin protein-derived antigens [32] contributes to direct cytotoxic effects and stimulation of macrophages [31, 33]. B cells have a role in the proliferation and reactivation of T cells, acting as antigen-presenting cells, and fully differentiated B cells further contribute to demyelination via the production of antibodies to myelin by plasma cells [30, 31, 33, 34]. Thus, the combined actions of T cells, B cells, and macrophages contribute to demyelination and axonal damage, triggering neurodegenerative processes from disease onset [30, 31]. Later in the course of the disease, chronic CNS inflammation drives ongoing neurodegeneration, possibly via neurotoxic inflammatory mediators produced by activated microglia and astrocytes [32].

Treatment options for relapsing forms of MS (RMS) comprise a number of immunosuppressive and immunomodulatory agents. The first DMD, IFN-β-1b, was approved by the US Food and Drug Administration (FDA) for RMS in 1993 [19] and by the European Medicines Agency (EMA) in 1995 [35]. Alternative forms of IFN-β (subcutaneous [SC] and intramuscular [IM] IFN-β-1a) and glatiramer acetate entered the market between 1996 and 2002, becoming standard treatment over the ensuing years [36]. The introduction of natalizumab, a recombinant monoclonal antibody against cell adhesion molecule α4-integrin [23] (initially approved in 2004, withdrawn from the market, then reintroduced in 2006 [37]), marked a shift in the treatment paradigm for MS [38]. While this molecule was perceived to be more effective than the IFNs and glatiramer acetate [39], it is associated with greater risks, including progressive multifocal leukoencephalopathy (PML), which is rare but carries significant morbidity and a mortality rate of approximately 23% [40]. After a span of nearly 20 years of injectable therapies, oral DMDs became available. While patients have generally preferred oral DMDs over injections when given the choice [41], most approved oral DMDs, including fingolimod, teriflunomide, dimethyl fumarate and siponimod, require regular adherence via cytolysis and complement-mediated lysis, their reaction to myelin protein-derived antigens [32] contributes to direct cytotoxic effects and stimulation of macrophages [31, 33]. B cells have a role in the proliferation and reactivation of T cells, acting as antigen-presenting cells, and fully differentiated B cells further contribute to demyelination via the production of antibodies to myelin by plasma cells [30, 31, 33, 34]. Thus, the combined actions of T cells, B cells, and macrophages contribute to demyelination and axonal damage, triggering neurodegenerative processes from disease onset [30, 31]. Later in the course of the disease, chronic CNS inflammation drives ongoing neurodegeneration, possibly via neurotoxic inflammatory mediators produced by activated microglia and astrocytes [32].

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target antigens on B cells, T cells, and other immune cells [24, 25, 44–46].

Cladribine tablets are the first oral therapy with an infrequent dosing schedule, administered in two yearly treatment courses, each divided into two treatment cycles comprising 4–5 days of treatment [12, 26]. Cladribine tablets are thought to exert their clinical effects via a transient reduction of selective lymphocyte subtypes, followed by a recovery period during which cell numbers return to the normal range [47, 48]. While immune function is restored as cell numbers recover, cell subtype ratios are thought to be altered, resulting in a reduction in autoreactive lymphocytes [47–49]. Thus, a short dosing period is thought to achieve sustained effects on lymphocytes that persist for an extended period after the treatment period has ended.

Cladribine tablets have been approved worldwide by many different regulatory authorities, including the EMA in 2017 [50] and the FDA in 2019 [51]. In this review, we provide an overview of the development of cladribine tablets as a therapy for patients with MS, from initial discovery and development of the molecule to its putative MOA in MS. We also review the pertinent efficacy data from clinical studies, and an integrated safety analysis of > 3700 patient-years’ clinical trial experience with cladribine tablets monotherapy at the recommended dosage.

1.1 Discovery and Development

Cladribine (2-chlorodeoxyadenosine [2-CdA]) was first synthesized nearly 50 years ago [52] (Fig. 1). It is a synthetic purine nucleoside analog of deoxyadenosine that was developed to selectively target lymphocytes in lymphoproliferative diseases (e.g. hairy cell leukemia) and autoimmune disorders [53–55]. The design of cladribine was inspired by the consequences of adenosine deaminase (ADA) deficiency in children [55, 56]. The selective vulnerability of lymphocytes in this autosomal recessive genetic disorder was described by Carson and colleagues as resulting from the preferential accumulation of cytotoxic deoxyadenosine nucleotides in lymphocytes, causing lymphocytopenia [54, 57]. Based on this understanding, several nucleoside analogs were synthesized, including cladribine in the early 1970s [54, 58, 59]. Following the efficacy observed in lymphoid leukemias and the ability of cladribine to selectively target lymphocyte populations, the drug was considered for potential use in MS [53].

1.2 Mechanism of Action

Cladribine (2-CdA) is a small molecule (molecular mass 285.7 g/mol [60]) prodrug that is taken up by cells via nucleoside transporter proteins and becomes active in certain cells upon phosphorylation to 2-chlorodeoxyadenosine triphosphate (2-Cd-ATP). 2-CdA undergoes sequential intracellular phosphorylation, first to 2-chlorodeoxyadenosine monophosphate (2-Cd-AMP) mediated by deoxycytidine kinase (DCK), and subsequently to 2-chlorodeoxyadenosine diphosphate (2-Cd-ADP) and 2-Cd-ATP by other kinases. De-phosphorylation by 5′-nucleotidases (5′-NTases) prevents accumulation of 2-Cd-ATP in most cells [31]. Phosphorylation preferentially occurs in B and T lymphocytes due to their unique constitutively high DCK and relatively low 5′-NTase levels, compared with other cell types. DCK is a rate-limiting enzyme in the nucleoside salvage pathway that provides deoxyribonucleosides (dNTPs). The high DCK level is thought to be important for lymphocyte clonal expansion during development and immune reactions [61, 62]. Lymphocytes are therefore susceptible to accumulation of 2-Cd-ATP (Fig. 2). In cells where 2-Cd-ATP accumulates, it incorporates into deoxyribonucleic acid (DNA) strands, disrupting DNA synthesis and cell cycle progression, and inhibits enzymes involved in DNA synthesis, leading to cell death in both proliferating and quiescent lymphocytes [31, 59, 63].

While the precise mechanisms by which cladribine exerts its therapeutic effects in patients with MS is not known, its effects on B and T lymphocytes are thought to play a central role. Cladribine preferentially reduces cells of the adaptive immune system, while leaving the innate immune system relatively spared. Absolute lymphocyte counts (ALC) and B- and T-cell subset counts rapidly reach nadir following administration of cladribine tablets, and gradual lymphocyte count recovery begins soon after treatment and continues for months afterwards [64]. In the phase III CLARITY and CLARITY Extension studies [65, 66], median ALC reached nadir, at the lower limit of normal (LLN) at week 13, and was followed by recovery of cell counts back to the normal range (Fig. 3). In year 2, median ALC fell below the LLN but recovered into the normal range by week 84, or approximately 30 weeks postdose after completion of the second treatment cycle in year 2. Reduction of CD19+ B cells in year 1 also reached nadir at about week 13 (below LLN and approximately 80% change from baseline), followed by recovery towards baseline values (Fig. 4). CD19+ B-cell counts also fell below LLN in year 2, and recovered to LLN by week 84. CD4+ T-cell median counts showed a lesser decline from baseline to nadir (approximately 50% change from baseline, also reached at week 13) in year 1, followed by a more gradual recovery towards baseline (Fig. 5). While CD4+ T cells did not reach the LLN in year 1, they fell below the LLN in year 2 and reached threshold values for recovery by week 96, approximately 43 weeks post last therapy dose. Median CD8+ T cells never dropped below LLN (Fig. 6) [64]. Analysis of CD4+ T-cell subtypes in ORACLE-MS showed that subpopulations displayed different repopulation dynamics, leading to changes in the
relative proportions of the CD4+ T-cell subpopulations [67]. Median monocyte and neutrophil counts remained within the normal range throughout the CLARITY and CLARITY Extension studies (Figs. 7, 8), while the effects on natural killer (NK) cells were moderate and transient, with a 30–44% median decrease in NK cells followed by a recovery towards pretreatment levels by week 24 (Fig. 9). These findings support the view that the impact of cladribine on the innate immune system is relatively minor [67]. The preferential reduction of lymphocyte subpopulations, followed by the pattern of lymphocyte count recovery (termed immune reconstitution), may ‘reset’ the immune system to a less autoreactive state [64], a putative mechanism with considerable potential in the long-term treatment of MS [48].

1.3 Approvals

Cladribine tablets were developed as a potential treatment for MS in response to the efficacy seen in early studies of parenteral cladribine and the need for additional treatment options [53, 68, 69]. The pivotal CLARITY trial was completed in 2009 and published in 2010; however, some regulatory authorities indicated in 2011 that improved understanding of safety risks and the overall benefit–risk profile was required. Phase II and III studies that were ongoing at the time were completed, and a long-term safety registry was continued. The additional data served to support the thorough characterization of the safety profile of cladribine tablets in MS. Evaluation of these additional data and analyses of the compound’s longer-term benefit–risk profile supported new submissions to regulatory authorities. Cladribine tablets received EMA approval in August 2017 [50] and FDA approval in March 2019 [26]. As of July 2020, cladribine tablets have gained marketing authorization in more than 75 countries for the treatment of patients with various forms of RMS.

1.4 Dosing Schedule

Two dosages were investigated during the clinical development of cladribine tablets: 3.5 mg/kg and 5.25 mg/kg (cumulative doses over 2 years). In addition, in certain study arms, patients received retreatment in years 3 and 4 (cumulative doses over 4 years of 7.0 mg/kg and 8.75 mg/kg). The 3.5 mg/kg and 5.25 mg/kg doses appeared to be equally efficacious, but the 5.25 mg/kg dose was associated with an increased rate of higher-grade lymphopenia [65]. The 3.5 mg/kg dose was considered to have the most favorable benefit–risk profile.

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**Clinical Development of Cladribine Tablets**

**CLINICAL MILESTONES**

- 1972: IV formulation approved by FDA as an orphan drug for the treatment of hairy cell leukemia.
- 2010: Approved in Russia and Australia for RRMS but later withdrawn.
- 2011: CRL issued by the FDA for US approval.
- 2014: Marketing authorization application resubmitted to EMA with new clinical data from CLARITY-EXT, ORACLE-MS, and PREMIERE.
- 2016: NDA resubmitted to FDA with additional safety clinical data and analyses (no new efficacy data was submitted).
- 2017: EMA approval.
- 2018: FDA approval.
- 2019: Approved in more than 75 countries as of July 2020.

**REGULATORY MILESTONES**

- Cladribine tablets were approved by the EMA in 2017 and the FDA in 2019.

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Fig. 1 Cladribine tablets development milestones [51, 52, 65, 66, 94, 100, 102, 122–131]. a Naive and treatment-experienced patients with RRMS. b Treatment-naive patients at high risk for developing MS. c Patients with active RRMS in combination with IFN-β. d Including patients from CLARITY, CLARITY–EXT, ORACLE-MS, and PREMIERE. ADA adenine deaminase, 2-CdA 2-chlorodeoxyadenosine deaminase, CRL complete response letter, EMA European Medicines Agency, FDA Food and Drug Administration, IFN interferon, IV intravenous, MS multiple sclerosis, NDA new drug application, RRMS relapsing–remitting MS.
In countries where approved, the recommended cumulative dose of cladribine tablets is 3.5 mg/kg given over 2 years (one treatment course is 1.75 mg/kg/year) [12, 26]. Each course consists of two treatment weeks or cycles—one cycle at the beginning of the first month and one cycle at the beginning of the second month. Each treatment cycle lasts 4 or 5 consecutive days, depending on the patient’s weight, and patients receive one or two 10 mg tablets per day. Lymphocytes must be within normal limits prior to initiating the first course and at least 800 cells/mm³ prior to initiating the second course. Modeling simulations suggest that 92% of patients would not require a delay in receiving the second treatment course and < 1% would be ineligible for treatment in year 2 due to a delay in recovery of more than 6 months [70]. No further treatment with cladribine tablets may be required in years 3 and 4; a patient cohort

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**Fig. 2** Ratios of DCK to 5′-NTase mRNA expression in T cells and various non-hematologic cells [58]. 5′-NTase 5′-nucleotidases, DCK deoxycytidine kinase, mRNA messenger RNA. *Calculated using data from the BioGPS website (available at https://biogps.org/#goto=welcome). Adapted with permission from Giovannoni [58].
that received additional treatments in years 3 and 4 had comparable clinical efficacy compared with a study cohort that received placebo in years 3 and 4 [66]. The safety and efficacy of additional treatment courses after year 4 have not been studied. There are no current definitions of treatment failure with cladribine tablets. Any definition would need to be on a case-by-case basis, most likely taking into account disease activity prior to commencing cladribine tablets [71].

1.5 Clinical Pharmacology

Following oral administration, cladribine is rapidly absorbed, reaching maximal plasma concentrations after approximately 0.5 h (in the fasted state). Food delays absorption but does not affect overall exposure [72]. Oral bioavailability is approximately 40% compared with parenteral administration; reduced bioavailability is likely due to gastrointestinal efflux mediated by the breast cancer resistance
protein (BCRP), a transporter with affinity for cladribine [72]. Cladribine is rapidly taken up by lymphocytes, where it or its phosphorylated products accumulate, and reach intracellular concentrations that are approximately 30- to 40-fold greater than in plasma within 1 h of administration [72]. No measurable accumulation of cladribine in plasma has been observed following repeated once-daily oral dosing [72]. The estimated terminal half-life is approximately 1 day [26]. About 50% of cladribine is cleared renally. Non-renal clearance occurs largely in lymphocytes; accumulated 2-Cd-AMP and 2-Cd-ATP are cleared via lymphocyte elimination pathways and their intracellular half-life is approximately 10–15 h [72].

Due to the risk of additive effects on the immune system, concomitant immunosuppressive or myelosuppressive therapy (e.g. cyclosporine, methotrexate) with cladribine is contraindicated. Previous immunomodulatory or immunosuppressive therapy use, including washout periods, should be considered when initiating cladribine tablets [12, 26]. Cladribine pharmacokinetics was not altered when used concomitantly with the proton pump inhibitor pantopra- zole or with IFN-β. However, the risk of lymphopenia may
increase with concomitant use of IFN-β. Cladribine may also result in hematological adverse events (AEs) when administered with hematotoxic drugs (e.g. carbamazepine). As a substrate of BCRP, equilibrative nucleoside transporter 1 (ENT1) and concentrative nucleoside transporter 3 (CNT3), coadministration with cladribine may interfere with cladribine exposure. A decrease in cladribine exposure is possible when coadministered with potent inducers of BCRP (e.g. corticosteroids) or P-glycoprotein (e.g. rifampicin); however, acute short-term corticosteroid therapy can be concomitantly administered. Antivirals/antiretrovirals that require intracellular phosphorylation to become active (e.g. lamivudine, ribavirin) could potentially compete with cladribine over phosphorylation, affecting both cladribine and competing compound activity. The hydroxypropyl betadex component in cladribine may interact with active ingredients of other drugs to increase bioavailability, therefore a gap of at least 3 h between administrations is recommended. Cladribine is
Development of Cladribine Tablets

not a substrate of the cytochrome P450 (CYP) pathway and has no known inductive effect on CYP1A2, CYP2B6, and CYP3A4 enzymes [12, 26].

It is not known whether cladribine can reduce the effectiveness of hormonal contraceptives; however, a clinical trial to examine this is ongoing (ClinicalTrials.gov identifier: NCT03745144). A barrier method is currently recommended during treatment with cladribine tablets and for at least 4 weeks after the last dose in each treatment course [26]. In humans, cladribine has a half-life of < 24 h and is rapidly eliminated [73]. However, in supratherapeutic doses, teratogenicity has been observed in mice and rabbits, and short-term effects have been observed in male mice germ cells [74]. Cladribine tablets should therefore not be administered to pregnant women; pregnancy should be prevented using effective contraception during treatment and 6 months after the last dose in each treatment course in women and men of reproductive potential [26].

As a small molecule, cladribine has been demonstrated to cross the blood–brain barrier (BBB) in animal [75] and human studies [76, 77]. In a study of parenteral cladribine in children with acute myeloid leukemia, cerebrospinal fluid (CSF) concentrations reached approximately 25% of those in plasma [76]. CNS penetration of cladribine may allow for a treatment effect on lymphocytes that have migrated into the CNS, possibly also affecting lymphoid follicles in the meninges of patients and reducing intrathecal immunoglobulin G (IgG) synthesis. In a study of 29 patients with RRMS who had received parenteral cladribine, 55% of patients tested negative for oligoclonal bands (OCBs) in CSF post-treatment, despite all patients testing positive for OCB and demonstrating raised Ig concentrations in CSF at baseline [78]. Furthermore, in a 10-year follow-up, Expanded Disability Status Scale (EDSS) progression was significantly delayed in the OCB-negative patients despite both cohorts demonstrating equal characteristics at baseline.

2 Clinical Studies in Relapsing Forms of MS (RMS): Efficacy

2.1 CLARITY (CLAdRibe Tablets Treating MS orally): NCT00213135

CLARITY was a 96-week, phase III, double-blind, randomized, placebo-controlled, parallel-group, multicenter study that evaluated the safety and efficacy of cladribine tablets 3.5 mg/kg and 5.25 mg/kg (cumulative dose) in patients with RRMS (Table 1) [65]. Patients with RRMS (according to the McDonald 2001 criteria [79]) at the 2005 trial initiation date were eligible if they had lesions on magnetic resonance imaging (MRI) consistent with MS (according to the criteria of Fazekas et al. [80]), had at least one relapse within 12 months before study entry, and had an EDSS score of no more than 5.5 (on a 0–10 scale, with higher scores indicating a greater degree of disability). The primary endpoint of the study was annualized relapse rate (ARR) at 96 weeks. Key secondary endpoints included the proportion of patients who were relapse-free and time to 3-month confirmed disability progression (CDP). CDP was defined as the time to a sustained increase, confirmed at 3 months, of at least 1

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EDSS point, or an increase of ≥1.5 if the EDSS score at baseline was 0. Other assessments included time to first relapse, proportion of patients receiving rescue therapy with IFN-β, mean number of lesions at 96 weeks for gadolinium-enhancing (Gd+) T1-weighted lesions, active T2-weighted lesions, and combined unique active (CUA) lesions, defined as new Gd+ T1-weighted lesions or new non-enhancing or enlarging T2-weighted lesions [65].

Among the 1326 patients in the intent-to-treat (ITT) population, 1184 patients (89.3%) completed the 96-week study (91.9% in the cladribine tablets 3.5 mg/kg group and 87.0% in the placebo group), and 1165 completed treatment courses. Baseline patient demographics and disease characteristics were generally well-balanced across treatment groups, although patients receiving cladribine tablets 3.5 mg/kg had a shorter mean duration of disease (7.9, 9.3, and 8.9 mean years for cladribine tablets 3.5 mg/kg and 5.25 mg/kg, and placebo, respectively; \( p = 0.005 \) for the overall comparison among the three groups). Mean (range) age in the cladribine tablets 3.5 mg/kg and placebo groups was 37.9 years (18–65) and 38.7 years (18–64), respectively. The percentage of patients who had received a prior DMD was 26.1% (\( n = 113 \)) in the cladribine tablets 3.5 mg/kg group and 32.5% (\( n = 142 \)) in the placebo group. Mean (standard deviation [SD]) EDSS score at baseline was similar among groups (2.8 [1.2] for the cladribine tablets 3.5 mg/kg group and 2.9 [1.3] for the placebo group) [65].

Treatment with cladribine tablets 3.5 mg/kg significantly reduced disease activity versus placebo (Fig. 10; Table 2). ARR was 0.14 versus 0.33, respectively (\( p < 0.001 \) for a relative reduction of 57.6%) (Fig. 10). The proportion of patients who remained relapse-free at week 96 was significantly higher in the cladribine tablets 3.5 mg/kg group compared with the placebo group (80% vs. 61% \( p < 0.001 \)) with imputation of missing values for patients who withdrew early [65]; 81% vs. 63% \( p < 0.05 \) without imputation [26]). Additionally, time to first relapse was significantly longer in the cladribine tablets 3.5 mg/kg group versus placebo [hazard ratio (HR) 0.44, 95% confidence interval (CI) 0.34–0.58; \( p < 0.001 \)] [65]. A post-hoc analysis in patients stratified by baseline EDSS severity (with an EDSS of ≥3.5 or ≤3.0) at baseline revealed similar benefits in clinical outcomes compared with placebo across subgroups, with a 57.1% reduction in ARR relative to placebo in the EDSS ≥3.5 subgroup, and a 56.3% reduction in the EDSS ≤3.0 subgroup (\( p < 0.001 \) for both groups) [81]. An EDSS of >3.5 has been used as a proxy for secondary progressive MS (SPMS) [82–84], as there are no definitive accepted measures indicating when a patient switches from RRMS to SPMS [85].

Cladribine tablets 3.5 mg/kg significantly reduced the risk of 3-month CDP at 2 years, with a 33% reduction versus placebo (HR 0.67, 95% CI 0.48–0.93; \( p = 0.02 \)) in the primary analysis, which imputed patients who used rescue therapy with SC IFN-β-1a (11 [2.5%] and 27 [6.2%] patients in the 3.5 mg/kg and placebo groups, respectively) from the time that rescue therapy was started [65]. A post-hoc analysis including all patients (regardless of rescue therapy use) confirmed a significant reduction in the risk of 3-month CDP with cladribine tablets 3.5 mg/kg versus placebo (41% risk reduction; HR 0.59, 95% CI 0.43–0.82; \( p = 0.0013 \)) [86]. In this analysis, the risk of 6-month CDP was reduced by 47% with cladribine tablets 3.5 mg/kg compared with placebo (HR 0.53, 95% CI 0.36–0.79; \( p = 0.0016 \)) [86]. Among patients treated with cladribine tablets 3.5 mg/kg, 86% were free from 3-month CDP at 2 years, versus 79% with placebo (odds ratio [OR] 1.55, 95% CI 1.09–2.22, \( p = 0.02 \), in the primary analysis with imputation for patients with incomplete follow-up [65]; 87% vs. 81% without imputation [26]). A post-hoc analysis that removed patients with incomplete follow-up also found 86% of patients treated with cladribine tablets 3.5 mg/kg versus 79% with placebo were free from 3-month CDP at 2 years (OR 1.62, 95% CI 1.12–2.35; \( p = 0.0105 \)), and 91% of patients treated with cladribine tablets 3.5 mg/kg versus 85% with placebo were free from 6-month CDP at 2 years (OR 1.87, 95% CI 1.19–2.94; \( p = 0.0064 \)) [87]. A post-hoc analysis in a subgroup of 261 patients with high disease activity (HDA), or rapidly evolving severe RRMS, defined as two or more relapses in the year prior to study entry, found that this group had greater reduction in both 3-month CDP (72% risk reduction; HR 0.28, 95% CI 0.15–0.54; \( p = 0.0061 \)) and 6-month CDP (82% risk reduction; HR 0.18, 95% CI 0.08–0.44; \( p = 0.0036 \)) than the overall study population [88].

Assessment of active brain lesions on MRI showed that patients treated with cladribine tablets 3.5 mg/kg had a reduction of 85.7% in the mean number of T1 Gd+ lesions versus placebo (0.12 vs. 0.91; \( p < 0.001 \)) at 2 years (Table 2), with consistent reductions seen in the first and second treatment years [89]. At 2 years, 87.2% of patients treated with cladribine tablets 3.5 mg/kg were free of new T1 Gd+ lesions, compared with 47.4% of patients receiving placebo (OR 8.14, 95% CI 5.73–11.57; \( p < 0.0001 \)) [87]. Patients treated with cladribine tablets 3.5 mg/kg also showed a 73.4% reduction in the mean number of active T2-weighted lesions versus placebo at 2 years (0.38 vs. 1.43; \( p < 0.001 \)) [89], and 61.8% of patients treated with cladribine tablets 3.5 mg/kg were free of active T2 lesions versus 27.6% treated with placebo at 2 years (OR 5.34, 95% CI 3.60–7.91; \( p < 0.0001 \)) [87]. Cladribine tablets 3.5 mg/kg also reduced

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1 The 47% reduction is based on a Cox proportional hazards model with EDSS at baseline as the fixed effect. Other analyses utilize a Cox regression that adjusts for treatment group and geographic region, and result in a 46% reduction in risk of 6-month CDP.
| Study        | Design                                      | Objective                                                                 | Patient population                                                                 | Primary endpoint                                                                 |
|--------------|---------------------------------------------|---------------------------------------------------------------------------|------------------------------------------------------------------------------------|----------------------------------------------------------------------------------|
| CLARITY      | Phase III<br>96-week, randomized, double-blind, placebo-controlled, multicenter | Investigate the efficacy and safety of cladribine tablets                  | RRMS (McDonald criteria 2001), had lesions consistent with MS, had ≥ 1 relapse within 12 months before study entry, and a Kurtzke EDSS score ≤ 5.5 [N = 1303] | Annualized relapse rate at 96 weeks                                               |
| CLARITY Extension | Phase IIIb<br>96-week, randomized, double-blind, placebo-controlled, multicenter extension of CLARITY | To assess the safety and tolerability of cladribine tablets for an additional 96 weeks in patients who completed the CLARITY study | Definite RRMS (McDonald criteria 2001) [N = 806]                                     | Safety evaluations<br>AEs<br>Clinical laboratory test results<br>ECGs               |
| ORACLE-MS    | Phase III<br>Randomized, double-blind, placebo-controlled study with three groups and three treatment periods: 96-week initial treatment, followed by 96 weeks of either IFN-β maintenance or long-term follow-up (cladribine tablets or no therapy) | To evaluate the efficacy of two doses of cladribine tablets vs. placebo in delaying conversion to CDMS (Poser criteria) and to McDonald MS | Treatment-naïve patients with an FCDE at high risk of converting to MS, EDSS score of 0–5.0 [N = 617 (616 administered)] | Time to conversion to CDMS (Poser criteria) from randomization                   |
| ONWARD       | Phase IIb<br>96-week, randomized, double-blind, placebo-controlled, multicenter study with two parallel groups | To assess the safety and tolerability of cladribine tablets when added to IFN-β therapy; ≥ 1 relapse within 48 weeks prior to screening, EDSS score of 1.0–5.5 [N = 214] | Patients with active RMS despite ongoing IFN-β therapy, ≥ 1 relapse within 48 weeks prior to screening, EDSS score of 1.0–5.5 [N = 214] | Safety and tolerability extension study safety evaluations:<br>AEs<br>Clinical laboratory tests<br>ECGs |

*AEs* adverse events, *CDMS* clinically definite MS, *ECGs* electrocardiograms, *EDSS* Expanded Disability Status Scale, *FCDE* first clinical demyelinating event, *IFN* interferon, *MS* multiple sclerosis, *RMS* relapsing multiple sclerosis, *RRMS* relapsing–remitting MS
were 67%, 54%, and 44%, respectively, in the cladribine tablets 3.5 mg/kg group, compared with 39%, 24%, and 16%, respectively, in the placebo group (p < 0.0001 for cladribine tablets 3.5 mg/kg vs. placebo at each time point). Results were similar when analyzed using 6-month CDP for the disability component of NEDA (47% vs. 17%, cladribine tablets 3.5 mg/kg vs. placebo, at 2 years; OR 4.25, 95% CI 3.03–5.96; p < 0.0001) [87].

NEDA was also evaluated in subgroups of patients with HDA at baseline. Subgroups of interest included patients with high relapse activity (two or more relapses in the year prior to study entry), and an expanded group including those with high relapse activity (regardless of treatment status) plus patients with persistent disease activity despite treatment with a DMD (one or more relapses in the year prior to study entry while taking a DMD, and one or more T1 Gd+ lesion(s) or nine or more T2 lesions). In all subgroups evaluated (both HDA subgroups and their counterparts with lower disease activity), the proportions of patients achieving NEDA were significantly higher with cladribine tablets 3.5 mg/kg compared with placebo, but ORs were greater for HDA than non-HDA subgroups (ORs 7.8–8.0 for HDA subgroups, compared with 3.5–3.6 for the respective non-HDA subgroups), indicating that the greatest treatment benefit, relative to placebo, was attained by patients with HDA [88].

NEDA was further evaluated in subgroups of patients with a baseline EDSS severity of ≥ 3.5 or ≤ 3.0. The percentage of patients achieving NEDA using 3-month CDP at 2 years was 48.6% versus 17.3% in the EDSS ≥ 3.5 subgroup, and 41.8% versus 14.8% in the EDSS ≤ 3.0 subgroup (cladribine tablets 3.5 mg/kg vs. placebo, respectively). ORs (95% CI) were similar across subgroups and favored cladribine tablets 3.5 mg/kg over placebo, at 2 years; OR 4.25, 95% CI 3.03–5.96; p < 0.0001 for both groups) [87]. Patients had active disease as a criterion for enrollment into the trials.

2.2 CLARITY Extension: NCT00641537

CLARITY Extension was a 96-week, phase IIIb, double-blind, randomized, parallel-group, multicenter, extension study investigating long-term safety, tolerability, and efficacy of cladribine tablets for an additional 2 years beyond the 2-year CLARITY study (Table 1) [66]. Patients were eligible for the extension study if they completed the 2-year study period in CLARITY and had normal lymphocyte counts and other normal hematologic results within 28 days of the first planned additional dose [66]. Eligible patients who received placebo in CLARITY were assigned to cladribine tablets 3.5 mg/kg, and patients treated with cladribine tablets in CLARITY were re-randomized (2:1) to an additional 2-year course of cladribine tablets 3.5 mg/kg or placebo (Fig. 11). Clinical endpoints were considered exploratory, and included ARR, proportion of patients free from relapses, time to first relapse, and time to CDp [66]. MRI

![Fig. 10 CLARITY: annualized qualified relapse rate [65]. EDSS Expanded Disability Status Scale.](image-url)
Table 2 CLARITY efficacy: clinical and imaging endpoints and relapses during the 96-week study (ITT population) [65]

| Endpoint | Placebo [n = 437] | Cladribine tablets 3.5 mg/kg [n = 433] | Cladribine tablets 5.25 mg/kg [n = 433] |
|----------|------------------|--------------------------------------|--------------------------------------|
| Relapse rate (primary endpoint) | | | |
| ARR (95% CI) | 0.33 (0.29–0.38) | 0.14 (0.12–0.17) | 0.15 (0.12–0.17) |
| Relative reduction in ARR for cladribine vs. placebo, %<sup>a</sup> | | 57.6 | 54.5 |
| p value<sup>b</sup> | <0.001 | <0.001 | |
| Relapse-free rate | | | |
| Patients without relapse [n (%)] | 266 (60.9) | 345 (79.7) | 360 (78.9) |
| OR for cladribine vs. placebo (95% CI)<sup>c</sup> | 2.53 (1.87–3.42) | 2.43 (1.81–3.27) | |
| p value<sup>d</sup> | <0.001 | <0.001 | |
| Relapse at 96 weeks | | | |
| Number of relapses [n (%)] | | | |
| 0 | 266 (60.9) | 345 (79.7) | 360 (78.9) |
| 1 | 109 (24.9) | 69 (15.9) | 77 (16.9) |
| 2 | 44 (10.1) | 13 (3.0) | 13 (2.9) |
| 3 | 15 (3.4) | 5 (1.2) | 5 (1.1) |
| ≥4 | 3 (0.7) | 1 (0.2) | 1 (0.2) |
| p value<sup>e</sup> | <0.001 | <0.001 | |
| Need for rescue therapy | | | |
| Patients receiving rescue therapy [n (%)] | 27 (6.2) | 11 (2.5) | 9 (2.0) |
| OR for cladribine vs. placebo (95% CI)<sup>c</sup> | 0.40 (0.19–0.81) | 0.31 (0.14–0.66) | |
| p value<sup>d</sup> | 0.01 | 0.003 | |
| Time to first relapse | | | |
| 15th percentile of time to event, months<sup>f</sup> | 4.6 | 13.4 | 13.3 |
| HR for cladribine vs. placebo (95% CI)<sup>g</sup> | 0.44 (0.34–0.58) | 0.46 (0.36–0.60) | |
| p value<sup>g</sup> | <0.001 | <0.001 | |
| Time to 3-month CDP | | | |
| 10th percentile of time to event, months<sup>f</sup> | 10.8 | 13.6 | 13.6 |
| HR for cladribine vs. placebo (95% CI)<sup>g</sup> | 0.67 (0.48–0.93) | 0.69 (0.49–0.96) | |
| p value<sup>g</sup> | 0.02 | 0.03 | |
| Patients without a 3-month CDP | | | |
| Patients with no change [n (%)] | 347 (79.4) | 371 (85.7) | 387 (84.9) |
| OR for cladribine vs. placebo (95% CI)<sup>c</sup> | 1.55 (1.09–2.22) | 1.46 (1.03–2.07) | |
| p value<sup>d</sup> | 0.02 | 0.03 | |
| Lesion activity on brain MRI | | | |
| Gd+ T1-weighted lesions | | | |
| Mean number | 0.91 | 0.12 | 0.11 |
| Relative reduction, % | 85.7 | 87.9 | |
| Active T2-weighted lesions | | | |
| Mean number | 1.43 | 0.38 | 0.33 |
| Relative reduction, % | 73.4 | 76.9 | |
| Combined unique lesions | | | |
| Mean number | 1.72 | 0.43 | 0.38 |
| Relative reduction, % | 74.4 | 77.9 | |
| p value<sup>h</sup> | <0.001 | <0.001 | |

ARR annualized relapse rate, CDP confirmed disability progression, CI confidence interval, Gd+ gadolinium-enhancing, HR hazard ratio, ITT intent-to-treat, MRI magnetic resonance imaging, OR odds ratio

<sup>a</sup>The relative reduction in the ARR was calculated as the ratio of the difference in the ARR between the placebo group and the cladribine group to the ARR rate in the placebo group.

<sup>b</sup>The p value was based on a Wald Chi-square test from an analysis of the number of relapses with the use of a Poisson regression model with
endpoints included number of T1 Gd+ lesions, number of active T2 lesions, total T2 lesion volume, and proportions of patients with no T1 Gd+ lesions or no active T2 lesions [91]. Patients who received cladribine tablets in CLARITY and CLARITY EXT did so as a result of randomization and not as a result of evidence of disease activity following the CLARITY study, which differs from the clinical design of other phase III extension studies [66, 92]. It could be of further interest to study such patients.

Of the 1184 patients who completed CLARITY, 867 (73.2%) were enrolled in CLARITY Extension. As the first patients completed the CLARITY study, the CLARITY Extension study was not yet ready to start. Consequently, there was some variability in the interval between completing CLARITY and entering the Extension (median 40.3 weeks, range 0.1–118.0 weeks) [66]. There were five treatment groups: patients who received cladribine tablets 3.5 mg/kg in CLARITY followed by placebo in CLARITY Extension (CP3.5), n = 98; patients who received cladribine tablets 5.25 mg/kg in CLARITY followed by placebo in CLARITY Extension (CP5.25), n = 92; patients who received cladribine tablets 3.5 mg/kg in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension (CC7.0), n = 186; patients who received cladribine tablets 5.25 mg/kg in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension (CC8.75), n = 186; and patients who received placebo in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension (PC3.5), n = 244 [66]. In CLARITY Extension, 806 patients were treated, of whom 738 (91.6%) completed the 2-year study. Demographic and disease characteristics at the start of CLARITY Extension were generally similar across groups, except the PC3.5 group (patients who received placebo in CLARITY), which showed evidence of greater disease activity than those who received cladribine tablets in CLARITY [66]. In particular, patients in the PC3.5 group had greater numbers and volume of T1 Gd+ lesions versus those who received cladribine tablets 3.5 mg/kg in CLARITY [66].

Treatment with cladribine tablets 3.5 mg/kg for 2 years followed by 2 years of treatment with placebo (CP3.5) demonstrated durable clinical benefits at year 4; these efficacy results were similar to those of patients who had 4 years of cladribine tablets 3.5 mg/kg treatment (CC7.0), with no significant differences observed between CP3.5 and CC7.0 across clinical efficacy endpoints (Table 3) [66]. MRI endpoints at the end of year 4 consistently showed high percentages of patients with no new T1 Gd+ lesions in both the CP3.5 and CC7.0 groups, although significantly more patients who received additional cladribine (4 years’ active treatment) remained free from T1 Gd+ lesions (73.0% vs. 88.9%, CP3.5 vs. CC7.0, respectively; p = 0.001) [91].

Exploratory post-hoc analysis suggested that NEDA status was durable in patients treated with cladribine tablets 3.5 mg/kg: overall, 42% and 48% of patients in the CP3.5 and CC7.0 groups (p = 0.31), respectively, had NEDA during the first year of the CLARITY Extension study [93]. The proportion of patients reaching NEDA status over the long-term with oral cladribine merits further research.

### 2.3 ORACLE-MS (ORAL CLadribine for Early MS): NCT00725985

ORACLE-MS was a 96-week, phase III, double-blind, randomized, placebo-controlled, multicenter study investigating the effect of cladribine tablets on conversion to clinically definite MS (CDMS) in patients with early signs of disease as evidenced by a first clinical demyelinating event (Table 1) [94]. Patients with a first clinical demyelinating event ≤75 days before screening were randomized to cladribine tablets 3.5 or 5.25 mg/kg (cumulative dose over 2 years) or placebo. The planned number of subjects was 600 (200 per treatment group) and the primary endpoint was time to conversion to CDMS [94].

Of the 617 patients enrolled and randomized, 206 were assigned to cladribine tablets 3.5 mg/kg, 205 were assigned to cladribine tablets 5.25 mg/kg, and 206 were assigned to placebo (ITT = 616; one patient in the 5.25 mg/kg group discontinued after randomization but before receiving study drug). Baseline patient demographics and disease characteristics were well-balanced across treatment groups; mean age across groups was between 31.7 and 32.2 years, and each group was 63–67% female. At the end of year 2, 363 patients (59%) had completed all treatments, and 149 patients (24%) had permanently discontinued treatment. Cladribine tablets
Development of Cladribine Tablets

**Fig. 11 CLARITY Extension: study schema** [66]. CP 3.5 mg/kg cladribine tablets 3.5 mg/kg in CLARITY followed by placebo in CLARITY Extension, CP 5.25 mg/kg cladribine tablets 5.25 mg/kg in CLARITY followed by placebo in CLARITY Extension, CC 7 mg/kg cladribine tablets 3.5 mg/kg in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension, CC 8.75 mg/kg cladribine tablets 5.25 mg/kg in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension. DMD disease-modifying drug, MRI magnetic resonance imaging, PC 3.5 mg/kg placebo in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension, RRMS relapsing–remitting multiple sclerosis. Each short course of treatment comprised one or two 10-mg cladribine tablets taken once daily for 4 or 5 consecutive days, or an equivalent number of matching placebo tablets. As the first patients completed the CLARITY study, the CLARITY EXTENSION study was not yet ready to start. Consequently, after completing CLARITY, there was a variable gap period (bridging interval) before patients entered the Extension (the median gap duration for the overall population was 40.3 weeks). Patients who had received interferon-β or glatiramer acetate during the gap period had to discontinue their DMD therapy at least 3 months before the first study day of the Extension [66]. Adapted with permission from Giovannoni et al. [66]

**Table 3 CLARITY Extension efficacy** [66]

|                  | PC 3.5 mg/kg [n = 244] | Cladribine tablets 3.5 mg/kg | Cladribine tablets 5.25 mg/kg |
|------------------|------------------------|-------------------------------|-------------------------------|
| CP 3.5 mg/kg [n = 98] | ARR (97.5% CI)         | 0.10 (0.07–0.13)              | 0.15 (0.09–0.21)              | 0.10 (0.06–0.13) |
| CC 7.0 mg/kg [n = 186] | Proportion of patients qualifying relapse-free [n (%)] | 180 (79.6)                    | 68 (75.6)                     | 134 (81.2)         |
| CP 5.25 mg/kg [n = 92] | Proportion of patients who remained free from 3-month CDP [n (%)] | 185 (75.8)                    | 71 (72.4)                     | 144 (77.4)         |
| CC 8.75 mg/kg [n = 186] |                         | 132 (76.7)                    | 72 (78.3)                     | 142 (76.3)         |

Qualified relapses were defined as a ≥ 2-point increase in at least one functional system or a ≥ 1-point increase in at least two functional systems of the EDSS, excluding bladder/bowel or cognition changes, lasting ≥ 24 h, in the absence of fever, and preceded by ≥ 30 days of clinical stability or improvement. ARR adjusted for time on study only. Data are mean (SD), unless otherwise stated. The CLARITY Extension data in this table covers the 96-week double-blind period and the 24-week supplemental follow-up (including the bridge between periods). ARR annualized relapse rate. CC 7 mg/kg cladribine tablets 3.5 mg/kg in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension, CC 8.75 mg/kg cladribine tablets 5.25 mg/kg in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension, CP 3.5 mg/kg cladribine tablets 3.5 mg/kg in CLARITY followed by placebo in CLARITY Extension, CP 5.25 mg/kg cladribine tablets 5.25 mg/kg in CLARITY followed by placebo in CLARITY Extension, CDP confirmed disability progression, CI confidence interval, EDSS Expanded Disability Status Scale, PC 3.5 mg/kg placebo in CLARITY followed by cladribine tablets 3.5 mg/kg in CLARITY Extension, SD standard deviation.
3.5 mg/kg were associated with a significant risk reduction versus placebo (67%) for conversion to CDMS (HR 0.33, 95% CI 0.21–0.51; \( p < 0.0001 \)) (Table 4) [94]. Compared with placebo, patients treated with cladribine tablets 3.5 mg/kg had significantly lower median numbers of new or persisting T1 Gd+ lesions, new or enlarging T2 lesions, and CUA lesions (\( p < 0.0001 \)) (Table 4) [94]. In total, 25 cladribine tablets 3.5 mg/kg and 60 placebo patients who progressed to CDMS chose to receive subsequent SC IFN-β1a therapy, and in these patients, 4% of those previously in the cladribine tablets 3.5 mg/kg treatment group and 3.3% of those previously in the placebo group experienced lymphopenia, suggesting that there was no additive effect of treatment switching on developing lymphopenia [94].

Diagnostic criteria for MS have been revised since enrollment of the ORACLE-MS study population [95]. More than one-third of patients in ORACLE-MS, originally considered to have clinically isolated syndrome (CIS) under the 2005 McDonald criteria [96], would receive a diagnosis of MS under the revised 2010 McDonald criteria [95, 97]. A post-hoc analysis conducted in patients with CIS per the 2010 McDonald criteria (\( n = 393 \)) was consistent with the original analysis in the ITT population: treatment with cladribine tablets 3.5 mg/kg (\( n = 138 \)) significantly reduced the risk of next clinical event or 3-month CDP by 63% versus placebo (\( n = 134; p = 0.0003 \)) [97]. Furthermore, in patients with MS defined per the McDonald 2010 criteria, cladribine tablets 3.5 mg/kg (\( n = 68 \)) significantly reduced the risk of another event or 3-month CDP by 74% versus placebo (\( n = 72; \text{HR} 0.26, 95\% \text{CI} 0.12–0.58; p = 0.0009 \)) [97]. Application of the most recent 2017 McDonald criteria [98] to the ORACLE-MS population would likely identify an even higher proportion of patients with RRMS [99]. In the US label, CIS is not a recommended indication for cladribine tablets [26].

### 2.4 Onward (Oral cladribine added on to IFN-β in patients with Active Relapsing Disease): NCT00436826

ONWARD was a 2-year, randomized, double-blind, phase IIb study assessing cladribine tablets as an add-on to IFN-β in patients with active RMS (one or more relapses during 48 weeks of treatment with IFN-β) (Table 1) [100]. Under the original study protocol, two cumulative doses of cladribine tablets (3.5 mg/kg and 5.25 mg/kg) added to existing IFN-β therapy (any form of IFN-β) were to be assessed.
Following a signal related to lymphopenia with cladribine tablets 5.25 mg/kg plus IFN-β in this study (15/17 [88.2%] patients had grade 3/4 lymphopenia [<500 cells/mm³]), this dose was discontinued and patients who had already been randomized to this dose continued in the trial but received IFN-β only and were followed up for safety. The increased risk of lymphopenia when cladribine tablets are used in combination with IFN-β is reflected in product labels, which state that concomitant treatment is not recommended [12, 26]. Under an amended protocol, patients in the ONWARD study were randomized 2:1 to cladribine tablets 3.5 mg/kg plus IFN-β, or placebo plus IFN-β [100].

Eligible patients were aged 18–65 years, had RRMS or SPMS with relapses (2005 McDonald criteria [96]), received treatment with IFN-β for ≥ 48 consecutive weeks before screening with one or more MS relapses during that period, had clinical stability (other than relapses) during the 28 days before screening, had an EDSS score of 1.0–5.5, and had normal hematologic parameters within 28 days of baseline (day 1 of randomization) [100]. The primary outcome was safety and tolerability of cladribine tablets added to IFN-β. Secondary objectives included the following efficacy evaluations: the number of qualifying relapses over 96 weeks; time to first qualifying relapse; time to 3-month CDP; mean numbers of new Gd+ T1, active T2, and CUA lesions (new Gd+ T1, new or enlarging T2 lesions, or both, without double counting) on MRI scans at 96 weeks. As ONWARD was designed as a phase II safety study, efficacy evaluations were exploratory and were not sufficiently powered to detect intergroup differences [100].

Under the amended protocol, 172 patients were randomized and analyzed for safety and efficacy; 42 patients were randomized under the original protocol and analyzed separately for safety [100]. Here, efficacy results for the 172 patients recruited under the amended protocol are discussed. IFN-βs administered in combination with cladribine tablets 3.5 mg/kg were SC IFN-β-1a 44 μg three times/week (Rebif®; n = 82), IM IFN-β-1a 30 μg once/week (Avonex®; n = 30), and SC IFN-β-1b 250 μg every other day (Betaseron®/Betaferon®; n = 60) [100]. Baseline demographics were similar in the cladribine tablets 3.5 mg/kg and IFN-β group and the placebo and IFN-β group; mean patient age was 38.5 and 40.1 years, and gender breakdown was 67.7% and 75.0% female, respectively [100].

Cladribine tablets at a cumulative dose of 3.5 mg/kg over 2 years plus IFN-β demonstrated a significant reduction in relapses and new T1 Gd+, active T2, and CUA lesions compared with IFN-β alone (Table 5) [100]. Patients treated with cladribine tablets 3.5 mg/kg were 63% less likely to have a qualifying relapse than patients receiving placebo and IFN-β (relative risk 0.37, 95% CI 0.22–0.63; p = 0.001) [100]. During the double-blind period, there was a significant reduction in the number of new T1 Gd+ lesions in the cladribine tablets 3.5 mg/kg and IFN-β group (mean 0.25 [SD 1.46]) compared with the placebo and IFN-β group (1.27 [SD 3.39]) [100]. Patients in the cladribine tablets 3.5 mg/kg and IFN-β group were 90% less likely to have a new T1 Gd+ lesion and 59% less likely to have a CUA lesion compared with the placebo and IFN-β group [100]. Fewer patients in the cladribine tablets 3.5 mg/kg and IFN-β group experienced a first qualifying relapse (23/124; 18.5%) versus the placebo and IFN-β group (16/48; 33.3%); however, the survival curves crossed, indicating that the assumptions on which the statistical analysis is based are not valid, therefore the result should be interpreted with caution [100]. The proportion of patients in the cladribine 3.5 mg/kg and IFN-β group with confirmed EDSS progression over 2 years (19/124; 15.3%) was similar to that in the placebo and IFN-β group (6/48; 12.5%) [100].

3 Clinical Studies in RMS: Safety

An integrated analysis of safety data from the phase III clinical trials (CLARITY, CLARITY Extension, and ORACLE-MS) and the PREMIERE registry (an observational subject registry designed to allow for further characterization of the long-term safety profile [101]) was performed to comprehensively assess the safety profile of cladribine tablets [102, 103]. The integrated analysis was performed using aggregated data with a cut-off date of 20 February 2015 [102], and subsequently updated using data gathered to May 2017 [103]. Data for patients treated with cladribine tablets 3.5 mg/kg monotherapy (n = 923; 3754 patient-years to May 2017) or placebo (n = 641; 2275 patient-years) in the CLARITY, CLARITY Extension, and ORACLE-MS phase III studies and the follow-up PREMIERE registry were aggregated to comprise the Monotherapy Oral Cohort. Studies that involved the use of parenteral cladribine in patients with MS, or cladribine tablets administered in combination with another DMD (the ONWARD study), were included in the All Exposed Cohort (cladribine n = 1976 and placebo n = 802) [102].

In the initial integrated analysis (cut-off February 2015), the incidence rate of treatment-emergent AEs (TEAEs) in patients receiving cladribine tablets 3.5 mg/kg as monotherapy compared with placebo-treated patients was 103.29 versus 94.26 adjusted AE incidence rates per 100 patient-years [102]. The most common TEAE was headache, with a similar incidence in patients exposed to cladribine tablets or placebo. Hematologic TEAEs such as lymphopenia and neutropenia were more common with cladribine tablets 3.5 mg/kg than placebo (Table 6) [102]. The updated safety analysis (cut-off date May 2017) demonstrated that AE rates remained stable, with no new major safety findings [103].
Table 5  ONWARD efficacy [100]

|                                | Placebo/IFN β [n = 48] | Cladribine tablets 3.5 mg/kg and IFN-β [n = 124] | Relative risk (95% CI)a | p value |
|--------------------------------|------------------------|--------------------------------------------------|------------------------|---------|
| Annualized qualifying relapse rate | 0.32                   | 0.12                                             | 0.37 (0.22–0.63)       | 0.001   |
| Number of new T1 Gd+ lesions [mean (SD)] | 1.27 (3.39)           | 0.25 (1.46)                                      | 0.10 (0.03–0.37)       | <0.001b |
| Total number of new T2 lesions [mean (SD)] | 3.65 (6.83)           | 2.05 (4.91)                                      | 0.54 (0.27–1.10)       | 0.090c  |
| Total number of combined unique lesions [mean (SD)] | 3.96 (7.38)           | 2.12 (4.95)                                      | 0.41 (0.21–0.81)       | 0.011d  |

Annualized qualifying relapse rate = (total number of qualifying relapses/total time on study during the double-blind period) × 365.25. Qualified relapses were defined as a ≥ 2-point increase in at least one functional system or a ≥ 1-point increase in at least two functional systems of the EDSS, excluding bladder/bowel or cognition changes, lasting ≥ 24 h, in the absence of fever, and preceded by ≥ 30 days of clinical stability or improvement. p value based on the Wald Chi-square test from the number of qualifying relapses using a Poisson regression model with fixed effects for the treatment group and IFN-β treatment and with log of time on study during the double-blind period as an offset. All efficacy analyses were exploratory, as the study was not powered to detect between-group differences for any endpoint. All p values < 0.05 are considered nominally significant

CI confidence interval, EDSS Expanded Disability Status Scale, Gd+ gadolinium-enhancing, IFN interferon, SD standard deviation

aRelative risk and associated 95% CI were estimated using a negative binomial model with fixed effects for the treatment group and IFN-β treatment, baseline T1 Gd+ lesions as a covariate, and log of number of scans as an offset variable

b p value based on the Wald Chi-square test from analysis of the total number of new T1 Gd+ lesions using a negative binomial model with fixed effects for the treatment group and IFN-β treatment, baseline T1 Gd+ lesions as a covariate, and log of number of scans as an offset variable

c p value based on the Wald Chi-square test from analysis of the total number of active T2 lesions using a negative binomial model with fixed effects for the treatment group and IFN-β treatment, with the log of number of scans as an offset variable

d p value based on the Wald Chi-square test from analysis of the total number of combined unique lesions using a negative binomial model with fixed effects for the treatment group and IFN-β treatment, baseline T1 Gd+ lesions as a covariate, and log of number of scans as an offset variable

Lymphopenia, infections, and malignancies were three AEs analyzed within the clinical program. Lymphopenia was commonly reported among patients receiving cladribine tablets. In the CLARITY and CLARITY Extension studies, grade 3 lymphopenia (ALC 0.2 to < 0.5 × 10⁹ cells/L) was experienced on at least one occasion by 25% of patients in the cladribine tablets 3.5 mg/kg treatment group; < 1% experienced grade 4 lymphopenia (ALC < 0.2 × 10⁹ cells/L) at any point during the studies [102]. Analysis of patients from CLARITY and CLARITY Extension, including those also enrolled in PREMIERE, showed that lymphocyte counts steadily increased in the weeks following the nadir (observed at about 2 months after treatment initiation in each year) and had reached normal range (ALC > 1.0 × 10⁹ cells/L) by study weeks 48 and 84 in treatment years 1 and 2, respectively [64].

Incidence rates for overall infection, severe infections, infections leading to discontinuation, or opportunistic infections were comparable between the cladribine tablets 3.5 mg/kg treatment group and the placebo group. Herpes zoster was the most common herpetic infection reported in patients in the cladribine tablets 3.5 mg/kg treatment group (Table 7). Herpes zoster infections occurred most frequently during periods of grade 3/4 lymphopenia (adjusted incidence rate 4.50 per 100 patient-years, compared with 0.73 per 100 patient-years outside periods of grade 3/4 lymphopenia) [102]. For patients who are antibody-negative for varicella zoster virus (VZV), vaccination against VZV is recommended at least 4–6 weeks prior to initiation of cladribine tablets therapy. Due to the risk of active vaccine infections, cladribine therapy must not be initiated within 4–6 weeks after vaccination with live or live-attenuated vaccines. Vaccinations with live or live-attenuated vaccines during and after a treatment course (when white blood cells counts are not within normal limits) should also be avoided [26]. During the clinical development program of cladribine tablets in MS, a total of three cases of tuberculosis (TB) were reported [26]. All three cases occurred in regions where TB is endemic and in patients who were enrolled in the trials prior to implementation of a protocol amendment requiring screening for TB [26]. TB is included as an adverse reaction in the product labeling of cladribine tablets [12, 26].

The overall incidence of malignancies in the integrated safety analysis was 0.27 per 100 patient-years (10 events in 3754 patient-years) in patients receiving cladribine tablets 3.5 mg/kg monotherapy, and 0.13 malignancies per 100 patient-years (3 events in 2275 patient-years) in patients receiving placebo, as of May 2017 [26]. Malignancies in the cladribine tablets 3.5 mg/kg oral monotherapy group included melanoma (n = 2), basal cell carcinoma, squamous cell carcinoma, pancreatic, breast, ovarian, rectal, and papillary thyroid cancers, and bile duct adenocarcinoma (n = 1 each) (data on file); in the placebo group, there were two cases of cervical carcinoma and one basal cell carcinoma [26]. This indicates no clustering of malignancies of any particular type and no increase in malignancies commonly
Table 6 Integrated safety data: summary of AEs for the monotherapy oral cohort (cut-off February 2015) [102]

|                                      | Placebo [N = 641] | Cladribine tablets 3.5 mg/kg [N = 923] |
|--------------------------------------|-------------------|----------------------------------------|
|                                      | n                | T     | Adj-AE per 100 PY | n    | T     | Adj-AE per 100 PY |
| Number of patients with ≥ 1 TEAE     | 515              | 546.3 | 94.26             | 773  | 748.4 | 103.29          |
| Number of patients with ≥ 1 TEAE related to study drug | 291              | 1162.8 | 25.03             | 542  | 1605.5 | 33.76          |
| Number of patients with ≥ 1 serious TEAEa | 67              | 1876.3 | 3.57              | 124  | 3096.8 | 4.00          |
| Number of patients with TEAE leading to treatment discontinuation | 21              | 1993.7 | 1.05              | 67   | 3229.0 | 2.07          |
| Number of patients with ≥ 1 severe TEAE reported in ≥ 2 patients | 57              | 1912.5 | 2.98              | 115  | 3111.2 | 3.70          |
| Number of patients with TEAE leading to death | 5              | 2024.7 | 0.25              | 9    | 3431.0 | 0.26          |

Most commonly reported TEAEs (Adj-AE per 100 PY of ≥ 1.0 in either group)²

| Nervous system disorders            | 226              | 1429.5 | 15.81             | 327  | 2346.7 | 13.93          |
|-------------------------------------|-------------------|--------|--------------------|-------|--------|-----------------|
| Headache                            | 144              | 1631.9 | 8.82              | 230  | 2641.9 | 8.71            |
| Dizziness                           | 36               | 1944.4 | 1.85              | 47   | 3268.0 | 1.44            |
| MS relapse                          | 11               | 2014.0 | 0.55              | 21   | 3404.9 | 0.62            |
| Gastrointestinal disorders          | 197              | 1454.9 | 13.54             | 278  | 2482.0 | 11.20           |
| Nausea                              | 62               | 1845.6 | 3.36              | 86   | 3134.4 | 2.74            |
| Diarrhea                            | 44               | 1915.9 | 2.30              | 68   | 3202.0 | 2.12            |
| Abdominal pain, upper               | 22               | 1969.2 | 1.12              | 42   | 3283.1 | 1.28            |
| Toothache                           | 22               | 1957.5 | 1.12              | 35   | 3335.8 | 1.05            |
| Abdominal pain                      | 23               | 1966.2 | 1.17              | 30   | 3346.2 | 0.90            |
| Constipation                        | 22               | 1968.7 | 1.12              | 24   | 3368.3 | 0.71            |
| Vomiting                            | 24               | 1982.6 | 1.21              | 21   | 3637.3 | 0.62            |
| Blood and lymphatic system disorders| 47               | 1901.6 | 2.47              | 276  | 2543.4 | 10.85           |
| Lymphopenia                         | 21               | 1985.0 | 1.06              | 217  | 2731.8 | 7.94            |
| Leukopenia                          | 8                | 2008.4 | 0.40              | 43   | 3276.9 | 1.31            |
| Neutropenia                         | 4                | 2015.0 | 0.20              | 27   | 3362.8 | 0.80            |
| Musculoskeletal and connective tissue disorders | 153           | 1608.0 | 9.51              | 245  | 2654.5 | 9.23           |
| Back pain                           | 46               | 1890.0 | 2.43              | 102  | 3115.8 | 3.27            |
| Arthralgia                          | 38               | 1938.8 | 1.96              | 63   | 3236.0 | 1.95            |
| Pain in extremity                   | 33               | 1965.6 | 1.68              | 50   | 3283.6 | 1.52            |
| Myalgia                             | 21               | 1966.3 | 1.07              | 28   | 3356.9 | 0.83            |
| Musculoskeletal pain                | 16               | 1989.1 | 0.80              | 19   | 3375.9 | 0.56            |
| General disorders and administration site conditions | 168           | 1558.5 | 10.78             | 213  | 2717.1 | 7.84           |
| Influenza-like illness              | 61               | 1857.4 | 3.28              | 75   | 3167.6 | 2.37            |
| Fatigue                             | 47               | 1897.2 | 2.48              | 54   | 3252.1 | 1.66            |
| Pyrexia                             | 20               | 1980.8 | 1.01              | 36   | 3341.5 | 1.08            |
| Asthenia                            | 25               | 1941.8 | 1.29              | 33   | 3311.5 | 1.00            |
| Investigations                      | 83               | 1802.4 | 4.61              | 145  | 2939.6 | 4.93            |
| Lymphocyte count decreased          | 2                | 2023.1 | 0.10              | 26   | 3337.4 | 0.78            |
| Psychiatric disorders               | 85               | 1787.8 | 4.75              | 121  | 3009.3 | 4.02            |
| Insomnia                            | 32               | 1938.7 | 1.65              | 46   | 3285.5 | 1.40            |
| Depression                          | 23               | 1960.6 | 1.17              | 46   | 3298.1 | 1.39            |
| Anxiety                             | 12               | 2002.3 | 0.60              | 37   | 3312.1 | 1.12            |
| Respiratory, thoracic, and mediastinal disorders | 95           | 1770.8 | 5.36              | 118  | 3065.2 | 3.85           |
| Oropharyngeal pain                  | 35               | 1929.2 | 1.81              | 44   | 3278.8 | 1.34            |
| Cough                               | 27               | 1968.3 | 1.37              | 34   | 3336.6 | 1.02            |
| Vascular disorders                  | 45               | 1909.6 | 2.36              | 72   | 3231.2 | 2.23            |
| Hypertension                        | 25               | 1965.3 | 1.27              | 35   | 3338.5 | 1.05            |
associated with immunosuppression (hematologic, virally induced, or non-melanoma skin cancers) in patients treated with cladribine tablets 3.5 mg/kg [102]. Meta-analysis using CLARITY data showed that the zero incidence of malignancies in the placebo group was significantly lower than the expected rate seen in placebo groups in other trials in similar RMS populations, while the malignancy rate with cladribine tablets (n = 3 in the cladribine tablets 3.5 mg/kg group [65]) was not significantly different to rates in the active treatment groups in trials of other DMDs [104]. Additionally, results from epidemiologic analyses showed no increase in the incidence of malignancies in the clinical trial program compared with matched reference populations (standardized incidence ratio [SIR] 0.91, 95% CI 0.41–1.75, for the 3.5 mg/kg oral monotherapy cohort vs. GLOBOCAN matched reference population; SIR 0.44, 95% CI 0.13–1.41, for placebo vs. reference population) [105]. Analysis of malignancy rates over time showed that the incidence of malignancies remained constant in follow-up for years 1–4 (0.29 per 100 patient-years in the oral monotherapy cohort, as of the February 2015 cut-off) and years 5–8 + (0.28 per 100 patient-years). In contrast, the malignancy rates in the placebo group were higher in years 5–8 + (0.60 per 100 patient-years) than years 1–4 (0.06 per 100 patient-years), although the overall number of events was small [102]. This suggests that the low level of malignancies observed in the placebo group in the short-term was a chance observation.

In addition to continued monitoring of the safety population described above, a long-term, multi-country, prospective, observational post-authorization safety study is underway to further characterize the long-term safety profile of cladribine tablets [102]. The CLARION study is designed to follow 8000 patients who are starting treatment with cladribine tablets or fingolimod (an oral comparator that depletes lymphocytes) for a period of 10 years to evaluate AEs [102, 106].

4 Discussion

Despite the recent approval of several new DMDs for RMS, there still remains a need for therapies that provide improved efficacy, address issues with treatment adherence and compliance, reduce the need for invasive procedures and intensive monitoring, and avoid the potential for treatment-related fatigue observed with daily long-term administration. Due to the observed efficacy of parenteral cladribine in earlier pilot studies of patients with MS, the cladribine tablets formulation was designed and developed to address some of these needs.
### Table 7  Integrated safety data: summary of infections and infestations for the monotherapy oral cohort (cut-off February 2015) [102]

|                                      | Placebo [N = 641] | Cladribine tablets 3.5 mg/kg [N = 923] |
|--------------------------------------|-------------------|----------------------------------------|
|                                      | n   | Total PY | Adj-AE per 100 PY | n   | Total PY | Adj-AE per 100 PY |
| Most commonly reported TEAEs in SOC infections and infestations \(^a\) (Adj-AE per 100 PY of ≥ 1.0 in either group) | 314  | 1160.8 | 27.05  | 478  | 1917.5 | 24.93  |
| Nasopharyngitis                      | 97   | 1764.5  | 5.50    | 158  | 2951.0  | 5.35    |
| Upper respiratory tract infection    | 61   | 1869.1  | 3.26    | 109  | 3112.0  | 3.50    |
| Influenza                            | 51   | 1898.1  | 2.69    | 87   | 3169.5  | 2.74    |
| Bronchitis                           | 22   | 1964.4  | 1.12    | 55   | 3234.1  | 1.70    |
| Urinary tract infection              | 46   | 1916.8  | 2.40    | 55   | 3249.7  | 1.69    |
| Herpes zoster                        | 4    | 2019.0  | 0.20    | 28   | 3360.2  | 0.83    |
| Pharyngitis                          | 31   | 1961.4  | 1.58    | 27   | 3348.0  | 0.81    |
| Rhinitis                             | 22   | 1962.3  | 1.12    | 24   | 3354.0  | 0.72    |
| **Serious TEAEs in SOC infections and infestations** \(^b\) (Adj-AE of ≥ 0.05 in either group) | 10   | 2003.4  | 0.50    | 23   | 3357.6  | 0.69    |
| Appendicitis                         | 2    | 2023.5  | 0.10    | 1    | 3426.8  | 0.03    |
| Chronic hepatitis C                  | 1    | 2019.4  | 0.05    | 0    | 3432.7  | 0       |
| Chronic sinusitis                    | 1    | 2024.1  | 0.05    | 0    | 3432.7  | 0       |
| Erysipelas                           | 1    | 2022.1  | 0.05    | 0    | 3432.7  | 0       |
| Herpes zoster                        | 0    | 2026.0  | 0       | 2    | 3425.6  | 0.06    |
| Myocarditis bacterial                | 1    | 2024.2  | 0.05    | 0    | 3432.7  | 0       |
| Pneumonia                            | 3    | 2019.7  | 0.15    | 6    | 3403.4  | 0.18    |
| Pyelonephritis                       | 0    | 2026.0  | 0       | 2    | 3423.1  | 0.06    |
| Urethral abscess                     | 1    | 2025.9  | 0.05    | 1    | 3430.3  | 0.03    |
| Urinary tract infection              | 1    | 2024.4  | 0.05    | 4    | 3419.4  | 0.12    |
| **Infections and infestations SOC leading to treatment discontinuation** (Adj-AE of ≥ 0.01 in either group) | 3    | 2020.8  | 0.15    | 4    | 3415.5  | 0.12    |
| Hepatitis B                          | 0    | 2026.0  | 0       | 1    | 3427.9  | 0.03    |
| Herpes zoster                        | 0    | 2026.0  | 0       | 1    | 3424.8  | 0.03    |
| Pneumonia bacterial                  | 0    | 2026.0  | 0       | 1    | 3428.2  | 0.03    |
| Urinary tract infection              | 0    | 2026.0  | 0       | 1    | 3432.6  | 0.03    |
| Appendicitis                         | 1    | 2025.8  | 0.05    | 0    | 3432.7  | 0       |
| *Gardnerella* infection              | 1    | 2025.8  | 0.05    | 0    | 3432.7  | 0       |
| Varicella                            | 1    | 2021.0  | 0.05    | 0    | 3432.7  | 0       |
| **Severe infection** \(^c\) (Adj-AE of ≥ 0.05 in any group) | 17   | 1983.4  | 0.86    | 28   | 3336.2  | 0.84    |
| Pneumonia                            | 3    | 2109.7  | 0.15    | 6    | 3403.4  | 0.18    |
| Urinary tract infection              | 2    | 2023.4  | 0.10    | 4    | 3419.4  | 0.12    |
| Herpes zoster                        | 1    | 2024.4  | 0.05    | 3    | 3424.1  | 0.09    |
| Gastroenteritis                      | 0    | 2026.0  | 0       | 2    | 3423.8  | 0.06    |
| Pyelonephritis                       | 0    | 2026.0  | 0       | 2    | 3423.1  | 0.06    |
| **Opportunistic infections** \(^d\) | 23   | 1965.5  | 1.17    | 36   | 3321.9  | 1.08    |
| Axillary candidiasis                 | 1    | 2018.3  | 0.05    | 0    | 3432.7  | 0       |
| Fungal infection                     | 2    | 2023.4  | 0.10    | 8    | 3398.3  | 0.24    |
| Fungal skin infection                | 2    | 2018.4  | 0.10    | 3    | 3430.8  | 0.09    |
| Mastitis fungal                      | 1    | 2025.6  | 0.05    | 0    | 3432.7  | 0       |
| Onychomycosis                        | 3    | 2023.1  | 0.15    | 4    | 3419.0  | 0.12    |
| Oral candidiasis                     | 1    | 2021.6  | 0.05    | 1    | 3431.0  | 0.03    |
| Oral fungal infection                | 2    | 2013.9  | 0.10    | 1    | 3431.5  | 0.03    |
| Pulmonary tuberculosis              | 0    | 2026.0  | 0       | 1    | 3429.6  | 0.03    |
Cladribine tablets are generally approved for the treatment of patients with RMS; however, indication wordings differ in several countries based on the individual assessment of the same clinical data by regulatory agencies. In some regions, including the EU and Switzerland, cladribine tablets are approved for use in adult patients with highly active disease [12, 107], whereas in other countries, including Canada and Australia, cladribine tablets are approved for the treatment of patients with RRMS [108, 109]. In the US, cladribine tablets are approved in adults with RMS, including RRMS and active SPMS, and are generally recommended for patients who have had an inadequate response to, or are unable to tolerate, an alternate drug indicated for MS [26].

In phase II and III clinical trials, cladribine tablets have demonstrated efficacy over placebo across the RMS spectrum, from treatment-naïve to treatment-experienced patients, and from patients with early MS to those with highly active disease. In the ORACLE-MS study, cladribine tablets reduced the risk of conversion from CIS to CDMS [97]. In patients with a confirmed diagnosis of RRMS in the CLARITY study, significant reductions in both clinical and MRI activity with cladribine tablets led to a significantly higher proportion of patients attaining NEDA compared with placebo patients [65]. This efficacy was maintained across all subgroups evaluated, with patients with HDA at baseline achieving the greatest improvements in NEDA rates compared with placebo [88]. Furthermore, patients with active, as evidenced by one or more relapses within 12 months of study entry, SPMS, as defined by EDSS ≥ 3.5, achieved similar efficacy results to the subgroup without SPMS [87].

Table 7 (continued)

| Condition                        | Placebo [N = 641] | Cladribine tablets 3.5 mg/kg [N=923] |
|----------------------------------|-------------------|-------------------------------------|
|                                  | n | Total PY | Adj-AE per 100 PY | n | Total PY | Adj-AE per 100 PY |
| Skin candida                     | 1 | 2025.4   | 0.05               | 0 | 3432.7   | 0                   |
| Tonsilitis fungal                | 1 | 2022.0   | 0.05               | 0 | 3432.7   | 0                   |
| Tuberculosis                     | 0 | 2026.0   | 0                  | 1 | 3432.7   | 0.03                |
| Upper respiratory fungal infection| 0 | 2026.0   | 0                  | 1 | 3431.7   | 0.03                |
| Urinary tract infection fungal   | 0 | 2026.0   | 0                  | 1 | 3424.1   | 0.03                |
| Vulvovaginal candidiasis         | 3 | 2013.3   | 0.15               | 7 | 3416.9   | 0.20                |
| Vulvovaginal mycotic infection   | 7 | 2016.4   | 0.35               | 9 | 3402.0   | 0.26                |
| **Herpetic infections**          |    |          |                    | 19| 1969.9   | **0.96**            |
| Herpes zoster                    | 4 | 2019.0   | 0.20               | 28| 3360.2   | 0.83                |
| Oral herpes                      | 11| 1996.9   | 0.55               | 20| 3381.3   | 0.59                |
| Herpes simplex                   | 1 | 2018.1   | 0.05               | 5 | 3416.6   | 0.15                |
| Herpes virus infection           | 1 | 2020.6   | 0.05               | 4 | 3416.9   | 0.12                |
| Varicella                        | 2 | 2020.6   | 0.10               | 3 | 3422.6   | 0.09                |
| Herpes zoster disseminated       | 1 | 2023.7   | 0.05               | 2 | 3427.5   | 0.06                |
| Genital herpes                   | 0 | 2026.0   | 0                  | 1 | 3429.0   | 0.03                |
| **Herpes zoster**                | 4 | **2017.7**| **0.20**           | 29| **3356.2**| **0.86**            |
| Herpes zoster                    | 4 | 2019.0   | 0.20               | 28| 3360.2   | 0.83                |
| Herpes zoster disseminated       | 1 | 2023.7   | 0.05               | 2 | 3427.5   | 0.06                |

If a patient has multiple events, the time to first event is considered. For a patient with no event the time is censored at the last follow-up time for that patient

Adj adjusted, AE adverse event, n number of patients with events, HLT higher level term, MedDRA Medical Dictionary for Regulatory Activities, PTs preferred terms, PY patient-years, SOC system organ class, TEAEs treatment-emergent adverse event, Total PY total patient time on study in years

*a System organ class and preferred terms are presented in descending order of Adj-AE per 100 PY rate in the cladribine 3.5 mg/kg group

*b ‘Serious’ was defined as resultant in death; life-threatening; required inpatient hospitalization; congenital anomaly or birth defect; or was otherwise considered as medically important

*c Severe infection is a custom grouping defined by any serious or severe event belonging to the MedDRA SOC ‘Infections and Infestations’

*d Medical Concept Opportunistic Infection is a custom query containing PTs belonging to the medical concept of opportunistic infections, excluding the PTs belonging to the HLT ‘Herpes viral infections’

*e Medical concept herpetic infection, which is a custom grouping defined by PTs with the term ‘herpes zoster’

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These findings are encouraging, as progressive forms of MS have historically been more difficult to manage and DMDs have often been found to be less efficacious in these cohorts [83, 110]. CLARITY Extension showed that the efficacy observed in CLARITY was durable—the magnitude of suppression of clinical and MRI activity was sustained, and the proportion of patients with NEDA remained high, even in patients who did not receive additional active treatment with cladribine tablets (those who were randomized to placebo in the extension study, after receiving cladribine tablets 3.5 mg/kg in CLARITY) [66, 91]. Therefore, further treatment in years 3 and 4 does not appear to be required [12, 26] and is not recommended based on current evidence [26]. The efficacy and safety of treatment re-initiation beyond year 4 has not been investigated.

The development program of cladribine tablets has resulted in a comprehensive characterization of its safety profile, which includes up to 10 years’ follow-up in some patients [111] and more than 12,000 patient-years of follow-up overall (for MS patients receiving oral or parenteral cladribine or placebo), including 9509 patient-years’ exposure to cladribine in any formulation at any dosage, and >3700 patients-years’ follow-up (as of May 2017) for patients receiving cladribine tablets 3.5 mg/kg monotherapy in CLARITY, CLARITY EXTENSION, ORACLE-MS, and the PREMIERE registry [103]. In these clinical studies, lymphopenia was the most common TEAE, occurring more frequently with cladribine tablets 3.5 mg/kg compared with placebo as expected due to the mechanism of action of cladribine tablets. The majority of patients experienced grade 1 or 2 lymphopenia; grade 3 lymphopenia was observed in approximately 25% of patients, and <1% of patients experienced grade 4 lymphopenia at any time over 2 years of treatment [102]. Lymphocyte recovery began within weeks after each treatment course in years 1 and 2, and stabilized on return to baseline [64]. There was no evidence of B-cell hyper-repopulation, which has been observed with alemtuzumab (which depletes lymphocytes and has a different proposed MOA, targeting the CD52 epitope) and has been speculated to mediate autoimmune adverse effects (recorded in >20% of patients treated with alemtuzumab, most commonly thyroid disease) [112, 113]. The gradual recovery of lymphocyte populations following cladribine tablets occurs without triggering secondary autoimmunity [102]. In patients receiving cladribine tablets, no increase in opportunistic infections was shown. Herpes zoster, oral herpes, and TB were identified as adverse reactions with cladribine tablets [102]. Given the observed increase in herpes zoster cases during periods of grade 3/4 lymphopenia, the product label carries a recommendation to administer antiviral prophylaxis if a patient’s lymphocyte count drops below 200 cells/µL (i.e. grade 4 lymphopenia, defined as ALC < 0.2 × 10⁹ cells/L) [12, 26]. Analysis of clinical trial data showed an imbalance in the number of malignancies between cladribine tablets and placebo; however, the incidence was not dose-related, did not increase over time, and there was no clustering of malignancies of any particular type. Malignancies are considered a potential risk with cladribine because of the limited number of observations possible in a phase III study. The USPI includes a boxed warning stating that cladribine tablets may increase the risk of malignancy [26], and this potential risk is being further examined in a large post-approval study.

Most DMDs can be classified as maintenance therapies, based on their dosing regimens [15–23, 114]. Maintenance therapies are defined as being administered continuously without an intended interruption in dosing, because clinical efficacy is tightly linked to the period of dosing. The effects of these drugs are closely related to their short half-lives and pharmacokinetic and pharmacodynamic profiles; therefore, while some need to be administered more frequently than others, all require regular dosing. Over the past 30 years, multiple maintenance DMDs have been approved for the treatment of MS; however, the continuous immunosuppression of maintenance therapies may interfere with immune surveillance [115].

Drugs with MOA-producing effects that persist long after the drug is eliminated from the body provide an alternative to ongoing maintenance therapy. Cladribine tablets are postulated to be an immune reconstitution therapy (IRT) [64, 116]. Characteristics of IRTs include transient reductions of B and T lymphocyte counts and/or select lymphocyte subtypes, followed by a recovery period during which cells gradually repopulate and immune function is restored. The reconstituted lymphocyte population is hypothesized to be less autoreactive towards the CNS component, resulting in prolonged efficacy, as demonstrated in the cladribine tablets clinical studies [47, 48, 66, 91]. In these studies, there was no rebound or return of clinical or MRI activity even after recovery of lymphocyte counts, and this is also consistent with a qualitative change in the adaptive immune response after treatment with cladribine tablets [64].

Cladribine tablets are the first oral MS therapy with a non-continuous dosing regimen, with extremely short treatment courses. This may be beneficial to patients, since the small total number of treatment days could potentially improve treatment adherence [117, 118]. In a survey designed to assess the state of MS care in the US and EU, 51% (total N = 982) of patients who self-identified as having MS reported skipping doses, and the most common reason given was ‘forgot’ (43%) [119]. Furthermore, patients receiving long-term therapy for MS often experience treatment fatigue, leading to a decline in adherence over time [120]. Studies have found that the serious consequences of non-adherence may include a greater risk of relapse or progression of disease compared with patients who adhere to...
treatment [121]. The dosing regimen of cladribine tablets limits the potential for non-adherence, which may reduce the risk of relapse and disease progression in real-world settings. In addition, the short and non-continuous treatment regimen may allow for the planning of life events, including pregnancy; cladribine tablets allow for pregnancy 6 months after the last dose of medication has been administered [12, 26].

5 Conclusion

Important therapeutic advances in the past several years have provided numerous therapeutic options for patients with MS, allowing selection depending on their particular disease characteristics, expressed desires, and lifestyle choices. The recent introduction of cladribine tablets provides a new high-efficacy therapeutic option for treating patients with MS, with a low treatment and monitoring burden. In clinical studies, cladribine tablets (3.5 mg/kg over 2 years) have demonstrated efficacy across the RMS spectrum: from early MS to patients with RMS, including those with HDA. Potentially due to the molecule’s unique proposed MOA, cladribine tablets provide durable efficacy, with a maximum of 20 total days of treatment over 2 years. Cladribine has an established safety and tolerability profile, with a clinical development program that included more than 12,000 patient-years’ overall clinical and observational trial experience (all formulations and dosages), including more than 3700 patient-years for patients receiving cladribine tablets at the approved dosage (3.5 mg/kg taken over 2 years). Furthermore, cladribine tablets’ weight-based and unique posology provides an alternative dosing schedule that allows for reduced treatment and management burden for patients and providers. This new addition to the existing arsenal of DMDs provides another treatment option to further tailor disease management in patients with MS.

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Declarations

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References

1. Browne P, Chandraratna D, Angood C, Tremlett H, Baker C, Taylor BV, et al. Atlas of multiple sclerosis 2013: a growing global problem with widespread inequity. Neurology. 2014;83(11):1022–4. https://doi.org/10.1212/WNL.000000000000768.

2. GBD 2016 Multiple Sclerosis Collaborators. Global, regional, and national burden of multiple sclerosis 1990–2016: a systematic analysis for the Global Burden of Disease Study 2016. Lancet Neurol. 2019;18(3):269–85. https://doi.org/10.1016/S1474-4422(18)30443-5.

3. Wallin MT, Culpepper WJ, Campbell J, Nelson LM, Langer-Gould A, Marrie RA et al. The prevalence of multiple sclerosis in the United States: a population-based healthcare database approach. Paris, France: European Committee for Treatment and Research in Multiple Sclerosis (ECTRIMS); 25–28 October 2017. p 344.
Development of Cladribine Tablets

4. National Multiple Sclerosis Society. Relapsing-remitting MS. https://www.nationalmssociety.org/What-is-MS/Types-of-MS/Relapsing-remitting-MS. Accessed 24 Jul 2020.

5. Miller A, Dishon S. Health-related quality of life in multiple sclerosis: the impact of disability, gender and employment status. Qual Life Res. 2006;15(2):259–71. https://doi.org/10.1007/s11136-005-0891-6.

6. Owens GM. Economic burden of multiple sclerosis and the role of managed care organizations in multiple sclerosis management. Am J Manag Care. 2016;22(6 Suppl):s151–s15818.

7. Tilling K, Lawton M, Robertson N, Tremlett H, Zhu F, Harding K, et al. Modelling disease progression in relapsing-remitting onset multiple sclerosis using multilevel models applied to longitudinal data from two natural history cohorts and one treated cohort. Health Technol Assess. 2016;20(81):1–48. https://doi.org/10.3310/hta20810.

8. Dobson R, Dassan P, Roberts M, Giovannoni G, Nelson-Piercy C, Brex PA. UK consensus on pregnancy in multiple sclerosis: ‘Association of British Neurologists’ guidelines. Pract Neurol. 2019;19(2):106–14. https://doi.org/10.1136/practneurol-2018-002060.

9. Burkill S, Vattulainen P, Geissbuehler Y, Sabido Espin M, Popescu C, Suzart-Woischnik K, et al. The association between exposure to interferon-beta during pregnancy and birth measurements in offspring of women with multiple sclerosis. PLoS One. 2019;14(12):e0227120. https://doi.org/10.1371/journal.pone.0227120.

10. Hellwig K, Geissbuehler Y, Sabido M, Popescu C, Adamo A, Klinger I, et al. Pregnancy outcomes in interferon-beta-exposed patients with multiple sclerosis: results from the European Interferon-beta Pregnancy Registry. J Neurol. 2020. https://doi.org/10.1007/s00415-020-09762-y.

11. European Medicines Agency. Rebif - Procedural steps taken and scientific information after the authorisation. Amsterdam, The Netherlands. 2020. https://www.eea.europa.eu/en/documents/procedural-steps-after-rebif-epar-procedural-steps-taken-scientific-information-after-approval_en.pdf. Accessed 11 Feb 2020.

12. Mavenclad [package insert]. Rockland, MA: EMD Serono, Inc.; 2019.

13. Lunde HMB, Assmus J, Myhr KM, Bo L, Grytten N. Survival and cause of death in multiple sclerosis: a 60-year longitudinal population study. J Neurol Neurosurg Psychiatry. 2017;88(8):621–5. https://doi.org/10.1136/jnnp-2016-315238.

14. Marrie RA, Yu N, Blanchard J, Leung S, Elliott L. The rising prevalence and changing age distribution of multiple sclerosis in Minnesota. Neurology. 2010;74(6):465–71. https://doi.org/10.1212/WNL.0b013e3181cf6e0c.

15. Ploughman M, Beaulieu S, Harris C, Hogan S, Manning OJ, Alderdice PW, et al. The Canadian survey of health, lifestyle and aging with multiple sclerosis: methodology and initial results. BMJ Open. 2014;4(4):e005718. https://doi.org/10.1136/bmjopen-2014-005718.

16. Baker D, Marta M, Pryce G, Giovannoni G, Schmierer K. Memory B cells are major targets for effective immunotherapy in relapsing multiple sclerosis. EBioMedicine. 2017;16:41–50. https://doi.org/10.1016/j.ebiom.2017.01.042.

17. Leist T, Weisssert R. Cladribine: mode of action and implications for treatment of multiple sclerosis. Clin Neuropharmacol. 2011;34(1):28–35. https://doi.org/10.1038/nni3871.

18. Neuhaus O, Archeullos JJ, Hartung HP. Immunomodulation in multiple sclerosis: from immunosuppression to neuroprotection. Trends Pharmacol Sci. 2003;24(3):131–8. https://doi.org/10.1016/S0165-6147(03)00028-2.

19. Jelcic I, Al Nimer F, Wang J, Lentsch V, Planas R, Jelcic I, et al. Memory B cells activate brain-homing, auto-antireactive CD4(+) T cells in multiple sclerosis. Cell. 2018;175(1):85–100. https://doi.org/10.1016/j.cell.2018.08.011.e23.

20. Betaferon [package insert]. Berlin, Germany: Bayer AG; 2018.

21. Goodin DS, Frohman EM, Garmany GP Jr, Halper J, Likosky WH, Lublin FD, et al. Disease modifying therapies in multiple sclerosis: report of the Therapeutics and Technology Assessment Subcommittee of the American Academy of Neurology and the MS Council for Clinical Practice Guidelines. Neurology. 2002;58(2):169–78.

22. CenterWatch. Tysabri (natalizumab). https://www.centerwatch.com/drug-information/fda-approved-drugs/drug/869/tysabri-natalizumab. Accessed 22 Jul 2019.

23. Fontoura P. Monoclonal antibody therapy in multiple sclerosis: paradigm shifts and emerging challenges. Mabs. 2010;2(6):810–6. https://doi.org/10.1017/S1352458515623366.

24. Copaxone [package insert]. North Wales, PA: Teva Pharmaceuticals USA, Inc.; 2016.

25. Ocrevus [summary of product characteristics]. Grenzach-Wyhlen: Roche Pharma AG; 2018.

26. Ocrevus [summary of product characteristics]. Haverhill: Genzyme Ltd.; 2018.

27. Aebischer P, Mironi A, Lehnert M, et al. Ocrelizumab depletes CD20(+) B cells in multiple sclerosis patients. Neurology. 2010;74(6):465–71. https://doi.org/10.1212/WNL.0b013e3181cf6e0c.

28. Eagle T, Stuart F, Chua AS, LaRussa A, Leclaire K, Cook SL, et al. Treatment satisfaction across injectable, infusion, and oral disease-modifying therapies for multiple sclerosis. Mult Scler Relat Disord. 2017;18:196–201. https://doi.org/10.1016/j.msard.2017.10.002.

29. Songar BA. The role of natalizumab in the treatment of multiple sclerosis: benefits and risks. Ther Adv Neurol Disord. 2017;10(9):327–36. https://doi.org/10.1177/1756285617716002.

30. Eagle T, Stuart F, Chua AS, LaRussa A, Leclaire K, Cook SL, et al. Treatment satisfaction across injectable, infusion, and oral disease-modifying therapies for multiple sclerosis. Mult Scler Relat Disord. 2017;18:196–201. https://doi.org/10.1016/j.msard.2017.10.002.
Wiendl H. Cladribine—an old newcomer for pulsed immune reconstitution therapy (IRT) in multiple sclerosis. J Inflamm Res. 2014;7:19–27. https://doi.org/10.2147/JIR.S38079.

Ceronie B, Jacobs BM, Baker D, Dubuisson N, Mao Z, Ammoscoto F, et al. Cladribine treatment of multiple sclerosis is associated with depletion of memory B cells. J Neurol. 2018;265(5):1199–209. https://doi.org/10.1007/s00415-018-8830-y.

Wiendl H. Cladribine—an old newcomer for pulsed immune reconstitution in MS. Nat Rev Neurol. 2017;13(10):573–4. https://doi.org/10.1038/nrneurol.2017.119.

Karussis D, Petrou P. Immune reconstitution therapy (IRT) in multiple sclerosis: the rationale. Immunol Res. 2018;66(6):642–8. https://doi.org/10.1007/s12026-018-9032-5.

Terranova N, Hicking C, Dangond F, Munafo A. Effects of post-transplant lymphoproliferative disease on survival and quality of life in patients with hematopoietic stem cell transplantation: a systematic review and meta-analysis. J Clin Transl Hematol. 2019;3(2):100016. https://doi.org/10.1007/s40262-019-00695-y.

Fernandez O. Alemtuzumab in the treatment of multiple sclerosis: current evidence and future prospects. Ther Adv Neurol Disord. 2016;9(1):44–52. https://doi.org/10.1177/1756285615601933.

Beutler E. Cladribine (2-chlorodeoxyadenosine). Lancet. 1994;344(8914):9–13. https://doi.org/10.1016/0140-6736(94)92826-2.

Carson DA, Wasson DB, Taeble R, Yu A. Specific toxicity of 2-chlorodeoxyadenosine toward resting and proliferating human lymphocytes. Blood. 1983;62(4):737–43.

Beutler E. Cladribine (2-chlorodeoxyadenosine). Lancet. 1992;340(8825):952–6. https://doi.org/10.1016/0140-6736(92)92826-2.

Carson DA, Kaye J, Seegmiller JE. Lymphospecific toxicity in adenosine deaminase deficiency and purine nucleoside phosphorylase deficiency: possible role of nucleoside kinase(s). Proc Natl Acad Sci USA. 1977;74(12):5677–81.

Giovannoni G. Cladribine to treat relapsing forms of multiple sclerosis. Neurotherapeutics. 2017;14(4):874–87. https://doi.org/10.1007/s13311-017-0573-4.

Carson DA, Wasson DB, Kaye J, Ullman B, Martin DW Jr, Robbins RK, et al. Deoxycoformycin-mediated toxicity of deoxycytidine analogs toward malignant human lymphoblasts in vitro and toward murine L1210 leukemia in vivo. Proc Natl Acad Sci USA. 1980;77(11):6865–9.

National Center for Biotechnology Information. PubChem Database. Cladribine, CID=20279. National Institutes of Health. 2019. https://pubchem.ncbi.nlm.nih.gov/compound/Cladribine. Accessed 19 Jun 2019.

Toy G, Austin WR, Liao HI, Cheng D, Singh A, Campbell DO, et al. Requirement for deoxycoformycin kinase in T and B lymphocyte development. Proc Natl Acad Sci USA. 2010;107(12):5551–6. https://doi.org/10.1073/pnas.0913900107.

Toy G, Austin WR, Liao HI, Cheng D, Singh A, Campbell DO, et al. Requirement for deoxycoformycin kinase in T and B lymphocyte development [supplement]. Proc Natl Acad Sci USA. 2010;107(12):5551–6. https://doi.org/10.1073/pnas.0913900107.

Sigal DS, Miller HJ, Schram ED, Saven A. Beyond hairy cell: the activity of cladribine in other hematologic malignancies. Blood. 2010;116(16):2884–966. https://doi.org/10.1182/blood-2010-02-246140.

Comi G, Cook S, Giovannoni G, Rieckmann P, Sorenson PS, Vermersch P, et al. Effect of cladribine tablets on lymphocyte reduction and repopulation dynamics in patients with relapsing multiple sclerosis. Mult Scler Relat Disord. 2019;29:168–74. https://doi.org/10.1016/j.msard.2019.01.038.

Giovannoni G, Comi G, Cook S, Rammohan K, Rieckmann P, Soelberg Sorenson P, et al. A placebo-controlled trial of oral cladribine for relapsing multiple sclerosis. N Engl J Med. 2010;362(5):416–26. https://doi.org/10.1056/NEJMoa0902553.

Giovannoni G, Soelberg Sorenson P, Cook S, Rammohan K, Rieckmann P, Comi G, et al. Safety and efficacy of cladribine tablets in patients with relapsing-remitting multiple sclerosis: an extended analysis of surface markers. Ther Adv Neurol Disord. 2019;12:17562849584986. https://doi.org/10.1080/1756285617727603.

Stuve O, Soelberg Sorenson P, Leist T, Giovannoni G, Hyvert Y, Damian D, et al. Effects of cladribine tablets on lymphocyte subsets in patients with multiple sclerosis: an extended analysis of surface markers. Ther Adv Neurol Disord. 2019;12:17562849584986. https://doi.org/10.1080/1756285617727603.

Beutler E, Sipe JC, Romine JS, Koziol JA, McMillan R, Zyroph J. The treatment of chronic progressive multiple sclerosis with cladribine. Proc Natl Acad Sci USA. 1996;93(4):1716–20. https://doi.org/10.1073/pnas.93.4.1716.

Romine JS, Sipe JC, Koziol JA, Zyroph J, Beutler E. A double-blind, placebo-controlled, randomized trial of cladribine in relapsing-remitting multiple sclerosis. Prog Assoc Am Physicians. 1999;111(1):35–44.

TERRANOVA N, Hicking C, Dangond F, MUNAFO A. Effects of postponing treatment in the second year of cladribine administration: Clinical trial simulation analysis of absolute lymphocyte counts and relapse rate in patients with relapsing-remitting multiple sclerosis. Clin Pharmacokinet. 2019;58(3):325–33. https://doi.org/10.1007/s40262-018-0693-x.

Sorenson PS, Centonze D, Giovannoni G, Montalban X, Selchen D, Vermersch P, et al. Expert opinion on the use of cladribine tablets in clinical practice. Ther Adv Neurol Disord. 2020;13:17562849584986. https://doi.org/10.1177/17562849584986.

Herrmann R, Karlsson MO, Novakovic AM, Terranova N, Fluck M, Munafò A. The clinical pharmacology of cladribine tablets for the treatment of multiple sclerosis. Clin Pharmacokinet. 2019;58(3):283–97. https://doi.org/10.1007/s40262-018-0695-9.

Savic RM, Novakovic AM, Ekbom M, Munafò A, Karlsson MO. Population pharmacokinetics of cladribine in patients with multiple sclerosis. Clin Pharmacokinet. 2017;56(10):1245–53. https://doi.org/10.1007/s40262-017-0516-6.

Giovannoni G, Galazka A, Schick R, Leist T, Comi G, Montalban X, et al. Pregnancy outcomes during the clinical development program of cladribine in multiple sclerosis: an integrated analysis of safety. Drug Saf. 2020;43(7):635–43. https://doi.org/10.1007/s40264-020-00948-x.

Singh V, Voss EV, Benardais K, Stangel M. Effects of 2-chlorodeoxyadenosine (Cladribine) on primary rat microglia. J Neuroimmun Pharmacol. 2012;7(4):939–50. https://doi.org/10.1007/s11481-012-9387-7.
Development of Cladribine Tablets

76. Kearsn CM, Blakney RL, Santana VM, Crom WR. Pharmacokinetics of cladribine (2-chlorodeoxyadenosine) in children with acute leukemia. Cancer Res. 1994;54(5):1235–9.

77. Lilemark J. The clinical pharmacokinetics of cladribine. Clin Pharmacokinet. 1997;32(2):120–31. https://doi.org/10.2165/00003088-199732020-00003.

78. Rejdak K, Stelmasiak Z, Grieb P. Cladribine induces long lasting oligoclonal bands disappearance in relapsing multiple sclerosis patients: 10-year observational study. Mult Scler Relat Disord. 2019;27:117–20. https://doi.org/10.1016/j.msard.2018.10.006.

79. McDonald WI, Compton A, Edan G, Goodkin D, Hartung HP, Lublin FD, et al. Recommended diagnostic criteria for multiple sclerosis: guidelines from the international panel on the diagnosis of multiple sclerosis. Ann Neurol. 2001;50(1):121–7.

80. Fazekas F, Barkhof F, Filippi M, Gromman RL, Li DK, McDonald WI, et al. The contribution of magnetic resonance imaging to the diagnosis of multiple sclerosis. Neurology. 1999;53(3):448–56.

81. Rammohan K, Giovannoni G, Comi G, Cook S, Rieckmann P, Soelberg Sorensen P, et al. Cladribine tablets for multiple sclerosis: efficacy across patient subgroups from the phase III CLARITY study. Mult Scler Relat Disord. 2012;1(1):49–54. https://doi.org/10.1016/j.msard.2011.08.006.

82. Lorscheider J, Buzzard K, Jukubaitis V, Spelman T, Havrdova E, Horakova D, et al. Defining secondary progressive multiple sclerosis. Brain. 2016;139(Pt 9):2395–405. https://doi.org/10.1093/brain/aww173.

83. Kapoor R, Ho PR, Campbell N, Chang I, Deykin A, Forrestal I, et al. Siponimod versus placebo in secondary progressive multiple sclerosis (EXPAND): a double-blind, randomised, phase 3 study. Lancet. 2018;391(10127):1263–7. https://doi.org/10.1016/S0140-6736(18)30475-6.

84. Lee DH, Peschke M, Utz KS, Linker RA. Diagnostic value of the McDonald criteria: results from the ORACLE-MS study. Mult Scler J Exp Transl Clin. 2017;3(4):2055217317732802. https://doi.org/10.1177/1756285617732802.

85. Thompson AJ, Banwell BL, Barkhof F, Carroll WM, Coetzee T, Comi G, et al. Diagnosis of multiple sclerosis: 2010 revisions to the McDonald Criteria. Ann Neurol. 2011;69(2):292–302. https://doi.org/10.1002/ana.22366.

86. Polman CH, Reingold SC, Banwell B, Clanet M, Cohen JA, Filippi M, et al. Diagnostic criteria for multiple sclerosis: 2010 revisions to the McDonald Criteria. Ann Neurol. 2011;69(2):292–302. https://doi.org/10.1002/ana.22366.

87. Lee DH, Peschke M, Utz KS, Linker RA. Diagnostic value of the "McDonald Criteria". Ann Neurol. 2005;58(6):840–6. https://doi.org/10.1002/ana.20703.

88. Miret M, Menezes H, et al, editor. Evaluation of the long-term safety of cladribine tablets in multiple sclerosis treated with cladribine tablets in the CLARITY study: a post-hoc subgroup analysis. American Academy of Neurology (AAN); 15–21 April 2016; Vancouver, BC: 1510.

89. Giovannoni G, Cook S, Rammohan K, Rieckmann P, Sorensen PS, Vermersch P, et al. Sustained disease-activity-free status in patients with relapsing-remitting multiple sclerosis treated with cladribine tablets in the CLARITY study: a post-hoc and subgroup analysis. Lancet Neurol. 2011;10(4):329–37. https://doi.org/10.1016/S1474-4422(11)70023-0.

90. Giovannoni G, Soelberg Sorensen P, Cook S, Rammohan KW, Rieckmann P, Comi G, et al. Efficacy of cladribine tablets in high disease activity subgroups of patients with relapsing multiple sclerosis: a post hoc analysis of the CLARITY study. Mult Scler. 2019;25(6):819–27. https://doi.org/10.1177/1352458518771875.

91. Comi G, Cook SD, Giovannoni G, Rammohan K, Rieckmann P, Sorensen PS, et al. MRI outcomes with cladribine tablets for multiple sclerosis in the CLARITY study. J Neurol. 2013;260(4):1136–46. https://doi.org/10.1007/s00415-012-7745-0.

92. De Stefano N, Girogio A, Battaglini M, De Leucio A, Hicking C, Dangond F, et al. Reduced brain atrophy rates are associated with lower risk of disability progression in patients with relapsing multiple sclerosis treated with cladribine tablets. Mult Scler. 2018;24(2):222–6. https://doi.org/10.1177/1352458517700269.

93. Comi G, Cook S, Rammohan K, Soelberg Sorensen P, Vermersch P, Adeniji AK, et al. Long-term effects of cladribine tablets on MRI activity outcomes in patients with relapsing-remitting multiple sclerosis: the CLARITY Extension study. Ther Adv Neurol Disord. 2018;11:1756285617753365. https://doi.org/10.1177/1756285617753365.

94. Leist TP, Comi G, Cree BA, Coyle PK, Freedman MS, Hartung HP, et al. Effect of oral cladribine on time to conversion to clinically definite multiple sclerosis in patients with a first demyelinating event (ORACLE MS): a phase 3 randomised trial. Lancet. 2014;13(3):257–67. https://doi.org/10.1016/S1474-4224(14)70005-5.

95. Polman CH, Reingold SC, Banwell B, Clanet M, Cohen JA, Filippi M, et al. Diagnostic criteria for multiple sclerosis: 2010 revisions to the McDonald Criteria. Ann Neurol. 2011;69(2):292–302. https://doi.org/10.1002/ana.22366.

96. Polman CH, Reingold SC, Edan G, Filippi M, Hartung HP, Kapoor R, et al. Diagnostic criteria for multiple sclerosis: 2005 revisions to the "McDonald Criteria". Ann Neurol. 2005;58(6):840–6. https://doi.org/10.1002/ana.20703.

97. Freedman MS, Leist TP, Comi G, Cree BA, Coyle PK, Hartung HP, et al. The efficacy of cladribine tablets in CIS patients retrospectively assigned the diagnosis of MS using modern criteria: results from the ORACLE-MS study. Mult Scler J Transp Clin. 2017;3(4):2055217317732802. https://doi.org/10.1177/1756285617732802.

98. Thompson AJ, Banwell BL, Barkhof F, Carroll WM, Coetzee T, Comi G, et al. Diagnosis of multiple sclerosis: 2017 revisions of the McDonald criteria. Lancet Neurol. 2018;17(2):162–73. https://doi.org/10.1016/S1474-4422(17)30470-2.

99. Thompson AJ, Banwell BL, Barkhof F, Carroll WM, Coetzee T, Comi G, et al. Diagnosis of multiple sclerosis: 2017 revisions of the McDonald criteria. Lancet Neurol. 2018;17(2):162–73. https://doi.org/10.1016/S1474-4422(17)30470-2.

100. Lee DH, Peschke M, Utz KS, Linker RA. Diagnostic value of the McDonald criteria. Lancet Neurol. 2018;17(2):162–73. https://doi.org/10.1016/S1474-4422(17)30470-2.

101. Lee DH, Peschke M, Utz KS, Linker RA. Diagnostic value of the 2017 McDonald criteria in patients with a first demyelinating event suggestive of relapsing-remitting multiple sclerosis. Eur J Neurol. 2019;26(3):540–5. https://doi.org/10.1111/ene.13853.

102. Montalban X, Leist TP, Cohen BA, Moses H, Campbell J, Hickson C, et al. Cladribine tablets added to IFN-beta in active relapsing MS: the ONWARD study. Neurol Immunmunol Neuroinflamm. 2018;5(5):e477. https://doi.org/10.1212/NXI.0000000000000477.

103. Miret M, WJ, Genney L, et al, editor. Evaluation of the long-term safety of cladribine tablets in multiple sclerosis design of PREMIERE, a prospective, observational, 8-year safety registry. Berlin, Germany: European Neurological Society (ENS); 19–23 June 2010.

104. Cook S, Leist T, Comi G, Montalban X, Giovannoni G, Nolting A, et al. Safety of cladribine tablets in the treatment of patients with multiple sclerosis: an integrated analysis. Mult Scler Relat Disord. 2019;29:157–67. https://doi.org/10.1016/j.msard.2018.11.021.

105. Cook S, Giovannoni G, Leist T, Syed S, Nolting A, Schick R. Updated safety analysis of cladribine tablets in the treatment of patients with multiple sclerosis. Berlin, Germany: European
Committee for Treatment and Research in Multiple Sclerosis (ECTRIMS); 10–12 October 2018. p 875.

104. Pakpour J, Disanto G, Altman DR, Pavith S, Turner BP, Marta M, et al. No evidence for higher risk of cancer in patients with multiple sclerosis taking cladribine. Neuroul Neuroimmunol Neuroinflamm. 2015;2(6):e158. https://doi.org/10.1210/NNI.2015-00000000000158.

105. Cook S, Giovanni G, Leist T, Syed S, Nolting A, Damian D et al. Updated safety analysis of cladribine tablets in the treatment of patients with multiple sclerosis. American Academy of Neurology (AAN); 4–10 May 2019, Philadelphia, PA: P2-048.

106. European Network of Centres for Pharmacoepidemiology and Pharmacovigilance. Long term, prospective, observational cohort study evaluating the safety profile in patients with highly active relapsing multiple sclerosis (RMS) newly started on oral cladribine (CLARION). Registration Number: EUPAS24484. European Medicines Agency. https://www.emea.europa.eu/en/press_release/rrdipmsp8otbkooaxq3yvijyq8bqrys/xwqzv11sxxqosqbb6th9/s379562573046?id=25122. Accessed 15 Jul 2019.

107. Mavenclad [product information]. Zug: Merck (Schweiz) AG; 2019.

108. Mavenclad [product monograph]. Mississauga, ON: EMD Serono, Inc.; 2017.

109. Mavenclad [product information]. Frenchs Forest, NSW: Merck Serono Australia; 2019.

110. Shirani A, Okuda DT, Stueve O. Therapeutic advances and future prospects in progressive forms of multiple sclerosis. Neurotherapeutics. 2016;13(1):58–69. https://doi.org/10.1007/s13311-015-0409-z.

111. PR Newswire. Up to 10-years of follow-up data reaffirm safety profile of investigational cladribine tablets. EMD Serono, Inc. 2018. https://www.prnewswire.com/news-releases/up-to-10-years-of-follow-up-data-reaffirm-safety-profile-of-investigational-cladribine-tablets-873347245.html. Accessed 15 Jul 2019.

112. Baker D, Herrod SS, Alvarez-Gonzalez C, Giovannoni G, Schmiere K. Interpreting lymphocyte reconstitution data from the pivotal Phase 3 trials of alemtuzumab. JAMA Neurol. 2017;74(8):961–9. https://doi.org/10.1001/jamaneurol.2017.0676.

113. Cosburn M, Pace AA, Jones J, Ali R, Ingrum G, Baker K et al. Autoimmune disease after alemtuzumab treatment for multiple sclerosis in a multicenter cohort. Neurology. 2011;77(6):573–9. https://doi.org/10.1212/WNL.0b013e318228bec5.

114. Glatopa [package insert]. Princeton, NJ: Sandoz Inc.; 2018.

115. Glatopa [package insert]. Princeton, NJ: Sandoz Inc.; 2018.

116. Peupot M, Rubin RR, Kruger DF, Travis LB. Correlates of insulin injection omission. Diabetes Care. 2010;33(2):240–5. https://doi.org/10.2337/dc09-1348.

117. Tintore M, Alexander M, Costello K, Duddy M, Jones DE, Law N, et al. The state of multiple sclerosis: current insight into the patient/health care provider relationship, treatment challenges, and satisfaction. Patient Prefer Adherence. 2017;11:33–45. https://doi.org/10.2147/PDD.115090.

118. Crawford A, Jewell S, Mara H, McCatty L, Pelfrey R. Managing treatment fatigue in patients with multiple sclerosis on long-term therapy: the role of multiple sclerosis nurses. Patient Prefer Adherence. 2014;8:1093–9. https://doi.org/10.2147/PDD.S67334.

119. Menzini J, Caon C, Nichols C, White LA, Friedman M, Pill MW. Narrative review of the literature on adherence to disease-modifying therapies among patients with multiple sclerosis. J Manag Care Pharm. 2013;19(1 Suppl A):S24–S40. https://doi.org/10.18553/jmcp.2013.19.s1.S24.

120. Christensen LF, Broom AD, Bloch A. Synthesis and biological activity of selected 2,6-disubstituted-(2-deoxy-α- and β-n-erythro-pentofuranosyl)purines. J Med Chem. 1972;15(7):735–9. https://doi.org/10.1021/jm00077a010.

121. Menzin J, Caon C, Nichols C, White LA, Friedman M, Pill MW. Narrative review of the literature on adherence to disease-modifying therapies among patients with multiple sclerosis. J Manag Care Pharm. 2013;19(1 Suppl A):S24–S40. https://doi.org/10.18553/jmcp.2013.19.s1.S24.

122. Christensen LF, Broom AD, Bloch A. Synthesis and biological activity of selected 2,6-disubstituted-(2-deoxy-α- and β-n-erythro-pentofuranosyl)purines. J Med Chem. 1972;15(7):735–9. https://doi.org/10.1021/jm00077a010.

123. EMD Serono, Inc. Prospective observational long-term safety registry of multiple sclerosis patients who have participated in cladribine clinical trials (PREMIERE). ClinicalTrials.gov. 2018. https://clinicaltrials.gov/ct2/show/NCT01013350. Accessed 15 Jul 2019.

124. US Food and Drug Administration. FDA Approved Drug Products—Leustatin. 1993. https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm?event=overview.process&ApplNo=020229. Accessed 15 Jul 2019.

125. Menzin J, Caon C, Nichols C, White LA, Friedman M, Pill MW. Narrative review of the literature on adherence to disease-modifying therapies among patients with multiple sclerosis. J Manag Care Pharm. 2013;19(1 Suppl A):S24–S40. https://doi.org/10.18553/jmcp.2013.19.s1.S24.

126. Merck Serono Europe Limited withdraws its marketing authorisation application for Movectro (cladribine) [press release]. European Medicines Agency; 11 February 2011. https://www.ema.europa.eu/en/documents/press-release/merck-serono-europe-limited-withdraws-its-marketing-authorisation-application-movectro-cladribine_en.pdf. Accessed 11 Feb 2020.

127. FDA grants priority review to cladribine tablets for the treatment of relapsing forms of MS [press release]. EMD Serono, Inc.; 28 July 2010. https://media.emdserono.com/2010-07-28-EMD SERONO INC FDA GRANTS PRIORITY REVIEW W-TO-CLADRIBINE-TABLETS-FOR-THE-TREATMENT-OF-RELAPSING-FORMS-OF-MS. Accessed 15 Jul 2019.

128. PR Newswire. EMD Serono receives complete response letter from FDA on cladribine tablets new drug application. 2011. https://www.prnewswire.com/news-releases/emd-serono-receives-complete-response-letter-from-fda-on-cladribine-tablets-new-drug-application-117220383.html. Accessed 15 Jul 2019.

129. The Pharma Letter. EMA accepts Merck’s MAA for cladribine tablets in multiple sclerosis. 2016. https://www.thepharmaletter.com/article/ema-accepts-merck-s-maa-for-cladribine-tablet-in-multiple-sclerosis. Accessed 15 Jul 2019.

130. European Commission grants approval for mavenclad (cladribine tablets) [press release]. Merck KGaA; 25 August 2017. https://www.mavgroup.com/en/news/mavenclad-25-08-2017.html. Accessed 15 Jul 2019.

131. FDA accepts file for cladribine tablets as potential treatment for relapsing forms of multiple sclerosis [press release]. EMD Serono, Inc.; 30 July 2018. https://media.emdserono.com/FDA-Accepts-File-For-Cladribine-Tablets. Accessed 11 Feb 2020.

132. Solaard-Borgersen P, Domant F, Hicking C, Giovannoni G. Long-term lymphocyte counts in patients with relapsing-remitting multiple sclerosis (RRMS) treated with cladribine tablets 3.5 mg/kg: total lymphocytes, b and t cell subsets. Paris, France: European Committee for Treatment and Research in Multiple Sclerosis (ECTRIMS); 25–28 October 2017.