

Effects of vitamin D₃ in clinically isolated syndrome and healthy control participants: A double-blind randomised controlled trial

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Abstract

Background: Lowserum vitamin D levels are associated with susceptibility to, and severity of, multiple sclerosis. High dose vitamin D has been proposed as a potential immunomodulator in multiple sclerosis.

Objectives: We performed a single centre, investigator-led, exploratory, double-blind, randomised, placebo controlled, trial of vitamin D₃ in clinically isolated syndrome and healthy control participants to assess its immunological effects. Secondary end-points included clinical and magnetic resonance imaging outcomes and safety.

Methods: Clinically isolated syndrome patients and healthy control participants were randomised to: placebo, 5000 IU or 10,000 IU vitamin D₃/day (Vigantol oil). Study duration was 24 weeks.

Results: The trial did not meet its primary end point, with no difference in the frequency of pro-inflammatory CD4⁺ T cells (interleukin (IL)-17⁺/interferon (IFN)-γ⁺) seen. A higher level of disease freedom (67% versus 50%) was seen in those with serum 1,25 (OH) vitamin D levels >100 nmol/l but this did not reach significance. High dose vitamin D₃ was well tolerated with no safety signal.

Conclusions: High dose vitamin D₃ over 24 weeks was well tolerated but without immunological, magnetic resonance imaging or clinical evidence of benefit. The hypothesised therapeutic effects in clinically isolated syndrome or multiple sclerosis patients may require longer periods of administration or may only be seen in patients treated with vitamin D₃ as an adjunct to established disease modifying therapies.

Keywords: Multiple sclerosis, clinically isolated syndrome, vitamin D, randomised controlled trial, immunology, cholecalciferol

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Introduction

Multiple sclerosis (MS) is a relapsing inflammatory disorder affecting the central nervous system causing demyelination, axonal and neuronal injury resulting in cerebral and spinal atrophy. Its pathogenesis is not fully understood but involves a complex interplay between genetic and environmental factors.¹ There is a well recognised geographical distribution of MS with increased incidence and prevalence associated with latitude from the equator.² The association of both MS susceptibility and disease activity with low serum vitamin D levels has emerged from a number of studies showing

increased prevalence with lower levels of ultraviolet (UV) radiation^{3,4}, a higher risk of MS in offspring of women who are deficient in vitamin D^{5,6} which is further supported by month of birth studies;⁷ also, higher serum levels of vitamin D are associated with lower risk of relapse^{8,9} and the number of new lesions on magnetic resonance imaging (MRI) scanning.¹⁰

Vitamin D has been shown to modulate both the innate and adaptive immune systems.¹¹ The active metabolite, 1,25 (OH) vitamin D (25(OH)D), mediates its effects through the intracellular vitamin D receptor (VDR),¹²

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which is present in monocytes, dendritic cells, B-cells and CD4⁺ T cells.^{11,13} Activation of VDR alters transcription, proliferation and differentiation of immune cells¹² and both indirectly modulates immune response by reducing the activation of pro-inflammatory T cells by antigen presenting cells¹⁴ and directly inhibiting T cell and B cell proliferation.^{11,15} This results in a Th2 cell driven anti-inflammatory state.^{13,15,16} The diverse effects seen within the immune system in response to vitamin D supplementation may explain the conflicting results from the small number of human studies carried out to date.^{17–23}

Clinically isolated syndrome (CIS) refers to patients presenting with a first-ever symptom consistent with demyelination and without evidence of dissemination in time.²⁴ CIS patients have a well-established natural history, with an increased risk of MS associated with the number of asymptomatic MRI T2 lesions at presentation.²⁵ In many countries, including Ireland, a conservative approach of 'watchful waiting', rather than the immediate introduction of disease-modifying therapy, is taken. Vitamin D deficiency is common in Ireland,²⁶ as in many northern European countries and may contribute to both disease susceptibility and disease severity. We hypothesised that treatment of CIS patients with vitamin D₃ supplementation might reduce the risk of progression to either clinically- or MRI-confirmed MS. Therefore we carried out a randomised, double-blind, placebo-controlled trial of vitamin D₃ supplementation in CIS patients to assess the immunological and therapeutic effects, if any, of vitamin D₃ supplementation in these patients. Healthy control participants were also prospectively recruited to investigate if the immunological responses to vitamin D₃ supplementation differed between control participants and those with CIS. We also wanted to determine whether high dose vitamin D₃ is a safe and effective therapy for patients with CIS.

Participants and methods

Study participants

Inclusion criteria

1. CIS patients aged between 18–55 years with symptom onset within three months of screening, having two or more asymptomatic T2 lesions on brain MRI consistent with demyelination, not treated with steroids within 30 days of screening and not on any other disease-modifying therapy.

2. Healthy control participants (HCPs) were aged between 20–40 years with a female: male ratio of 2:1, broadly in line with the gender ratio of CIS.

Exclusion criteria

1. Exclusion criteria in both CIS patients and HCPs were: no history or evidence of hypercalcaemia, renal impairment, vitamin D intolerance, parathyroid dysfunction, sarcoidosis, pregnancy or refusal to use contraception, prior or current treatment with thiazide diuretics or vitamin D supplementation of greater than 1000 IU/day.
2. Exclusion criteria in patients with CIS were: (a) patients whose symptoms might be explained by a diagnosis other than MS; (b) patients with occurrence of an exacerbation less than six weeks prior to entry to the study; (c) previous treatment with any immunomodulating therapy in the last three months, steroids in the last four weeks or any previous treatment with mitoxantrone or other immunosuppressant.

Trial design

This was a single-centre, investigator-led, exploratory, double-blind, dose-ranging, randomised, placebo-controlled trial. A full detailed trial protocol has been previously published.²⁷ Detailed procedures carried out at each study visit are available in the Supplementary Material, Methods section. Participants were enrolled from November 2012 and last study visit was June 2015. Both HCPs and CIS patients were recruited over the same time period and studied in parallel. Both the HCPs and CIS patients were randomised to one of three interventions: placebo oil, 5000 IU and 10,000 IU vitamin D₃ per day. The vitamin D₃ was in the form of Vigantol Oil, supplied by Merck KgaA (Darmstadt, Germany). Both placebo and the active product were identical in appearance.

Endpoints

The primary endpoint was the effect of oral vitamin D₃ at 5000 IU and 10,000 IU per day compared to placebo on the frequency of CD4⁺ T cell subsets (interleukin (IL)-17⁺CD4⁺ T cells and interferon (IFN)- γ ⁺CD4⁺ T cells) and cytokine (IL-17, IL-10 and IFN- γ) responses in peripheral blood mononuclear cells (PBMCs) in patients with CIS and HCPs at 16 and 24 weeks compared to baseline.

Secondary endpoints included determining if, in CIS patients, there was a difference in the response to

5000 IU and 10,000 IU vitamin D₃, compared to placebo in:

1. Radiological activity as measured by the number of new or enlarging T2 lesions and gadolinium-enhancing lesions on brain MRI at 24 weeks compared to baseline.
2. Clinical activity as measured by the annualised relapse rate (ARR), percentage of patients relapse-free and time to first relapse.
3. Disease freedom as measured by the percentage of patients with no evidence of either radiological or clinical activity.

We also wished to determine the dose response and safety and tolerability of high dose vitamin D₃ supplementation in both CIS patients and HCPs.

Study procedures

Study visits were performed at baseline and then at four-weekly intervals over a 24-week period. A follow-up visit was arranged at week 28–30. At each visit, changes in medical and neurological history or medications were recorded. All patients had a full physical examination and CIS participants had their Expanded Disability Status Scale (EDSS) score recorded and were assessed for any evidence of a relapse. Treatment adherence, as measured by the volume of study drug returned at each visit, and any adverse events were also noted.

Vitamin D and parathyroid hormone

Plasma parathyroid hormone (PTH) levels were measured at baseline, week 4, 8, 16, 24 and 28–30 using the Roche intact PTH assay on the Cobas E411 Electrochemiluminescent Immunoassay Analyser (Roche Diagnostics GmbH, Mannheim, Germany). Serum 25(OH)D levels were measured by liquid chromatography mass spectrometry (LC-MS/MS) using a Waters Acquity ultra performance liquid chromatography (UPLC) coupled to a triple quadrupole mass (TQD) (Waters Corp., Milford, USA) with a semi-automated solid phase extraction (SPE) preparation of patient serum samples (Tecan Freedom Evo 100, Switzerland). A more detailed description is available in the Supplementary Material, Methods section.

Biochemistry, including urea and electrolytes, serum creatinine, liver function and serum calcium levels adjusted for albumin level, was measured at each study visit.

Immunological outcome measures

PBMCs were isolated and cryopreserved, and immunological analyses were performed upon completion of the study. PBMCs from all time-points were

thawed in batches, labelled with cell tracker dye CellTrace Violet (CTV) (Invivogen), and cultured at 2×10^6 cells/ml in a 48-well plate (0.5×10^6 cells in total) in complete Roswell Park memorial institute (RPMI) medium and were left unstimulated or stimulated with the following: 1 µg/ml anti-CD3 (eBioscience), 0.5×10^6 irradiated allogeneic PBMCs, 1 µg/ml each of purified protein derivative (PPD; Serum Staten Institute) and tetanus toxoid (TT) as recall antigens, and 0.625 µg/ml/0.375 nM each of myelin oligodendrocyte protein (MOG) and myelin basic protein (MBP) PepTivator overlapping peptides (Miltenyi Biotech). In parallel with each batch of samples the same standardised PBMC sample was included in order to monitor reproducibility between batches. After three days for anti-CD3, or seven days for antigen-specific stimuli, cell culture supernatants were removed and the concentrations of IFN-γ, IL-17 and IL-10 were analysed by enzyme-linked immunosorbent assay (ELISA) (eBioscience). In addition, the cells were restimulated with 50 ng/ml phorbol myristate acetate (PMA), 500 ng/ml ionomycin and 5 µg/ml of Brefeldin A for five hours (all Sigma), followed by staining for cell-surface CD3 and CD8 and intracellular IL-17 and IFN-γ (eBioscience). Cells were analysed by flow cytometry using a BD LSRFortessa Cell Analyser (BD Biosciences) and FlowJo software. T cell proliferation was analysed by CTV dilution.

MRI acquisition

Brain MRI was acquired at baseline and 24 weeks with contiguous axial 3 mm thick slices. Sequences included: axial and coronal T1-weighted spin echo (pre and post gadolinium), axial and sagittal PD/T2-weighted and axial fast fluid-attenuated inversion recovery (FLAIR). Baseline and 24-week scans were reviewed by two radiologists who were blinded to treatment group. MRI assessment included: (a) number of gadolinium-enhancing lesions; (b) number of new and enlarging T2 lesions; (c) total number of new lesions = number of gadolinium-enhancing lesions plus number of new and enlarging T2 lesions.

Statistical methods

Power calculations were based on a pilot study²⁸ using α of 5% and β of 80%. A detailed description is available in the Supplementary Material, Methods section. All serum 25(OH)D levels were adjusted for season²⁹ and these values were used in all further analysis. All data was inspected for normality using histograms, skewness and kurtosis and normality tests (Kolmogorov-Smirnov and Shapiro-Wilk). Values displayed include mean and standard deviation (SD) or median and interquartile range where

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appropriate. Within group differences between baseline and follow-up were assessed using the Wilcoxon Rank test as outlined in the protocol. Log₁₀ ratio of week 16/baseline and week 24/baseline was calculated and adjusted for baseline measurements (including vitamin D level) and between group differences were compared using the *t*-test or Mann Whitney U test where indicated. All analysis was carried out in SPSS version 20.

Ethical approval

Ethical approval was granted by St Vincent’s University Hospital Ethics and Medical Research Committee and the study was approved by the Health Products Regulatory Authority (HPRA). The trial was registered with ClinicalTrials.gov (NCT01728922) and the European Union Clinical Trials Register (EudraCT: 2012-000635-68).

Results

Study participants

Recruitment to the study is outlined in Figure 1. Baseline characteristics of subjects in each treatment

arm are outlined in Table 1. All participants were Caucasian. No significant differences were seen between each of the three treatment arms in either CIS patients or HCPs. Mean baseline EDSS was 0.9 (SD: 1), 0.9 (SD: 1.2) and 0.4 (SD: 0.5) in each of the 10,000 IU, 5000 IU and placebo groups respectively. No patients were on disease modifying treatment as per protocol.

Serum 25(OH)D, PTH and calcium levels

There was no difference in mean baseline serum 25(OH)D levels between CIS patients and HCP (53 vs 52 nmol/l, *p* = 0.8). For the CIS group, in the placebo, 5000 IU and 10,000 IU treatment arms, mean changes in serum 25(OH)D levels from baseline to week 16 were 7 nmol/l (SD: 26), 81 nmol/l (SD: 63) and 121 nmol/l (SD: 35); from baseline to week 24, mean changes were 18 nmol/l (SD: 34), 76 nmol/l (SD: 57) and 115 nmol/l (SD: 51) respectively. For the HCP group, in the placebo, 5000 IU and 10,000 IU treatment arms, mean changes in serum 25(OH)D levels from baseline to week 16 were 4 nmol/l (SD: 12), 83 nmol/l (SD: 27) and 152 nmol/l (SD: 71) and from

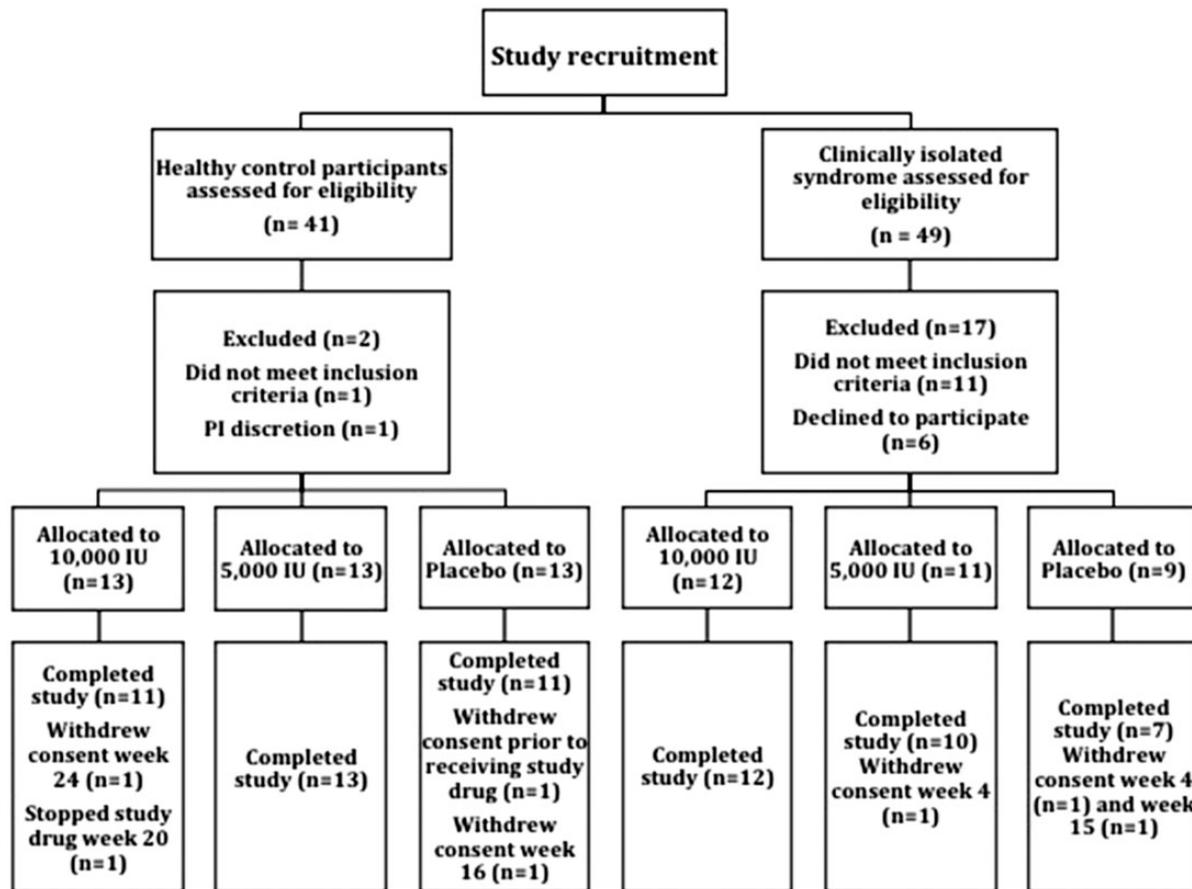
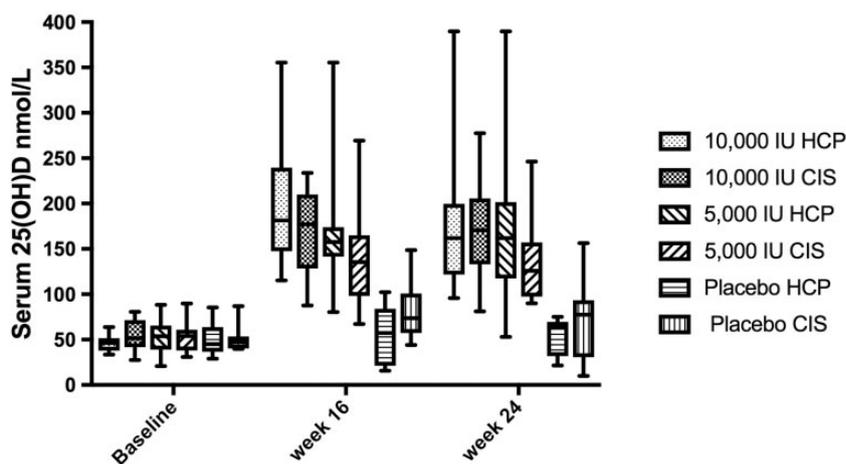


Figure 1. Schematic diagram outlining recruitment to the study, randomization and final analysis.

Table 1. Baseline characteristics of both clinically isolated syndrome (CIS) patients and healthy control participants (HCPs).

	Clinically isolated syndrome (<i>n</i> = 29)			Healthy control participants (<i>n</i> = 38)		
	10,000 IU (<i>n</i> = 12)	5000 IU (<i>n</i> = 10)	Placebo (<i>n</i> = 7)	10,000 IU (<i>n</i> = 13)	5000 IU (<i>n</i> = 13)	Placebo (<i>n</i> = 12)
Female <i>n</i> (%)	8 (67)	5 (50)	6 (86)	6 (46)	10 (77)	10 (83)
Age (years) mean (SD)	37.2 (8.7)	32.7 (4.6)	34.3 (10.6)	30.5 (5.1)	30.3 (3.7)	29.1 (4.7)
BMI (kg/m ²) mean (SD)	30.6 (5.6)	26.3 (3.4)	26.7 (4.2)	25.4 (3.2)	23.5 (3.3)	27.6 (4.3)
Ever smoked <i>n</i> (%)	7 (58)	5 (50)	3 (43)	2 (15)	3 (23)	4 (33)

BMI: body mass index; SD: standard deviation.

**Figure 2.** Graph showing median (boxes: 25–75% interquartile range and error bars: minimum–maximum) seasonally adjusted serum 1,25 (OH) vitamin D (25(OH)D) levels (nmol/l) achieved in each of the treatment arms at baseline, week 16 and week 24. CIS: clinically isolated syndrome; HCP: healthy control participant.

baseline to week 24 were 2 nmol/l (SD: 14), 92 nmol/l (SD: 35) and 136 nmol/l (SD: 71) respectively (Figure 2).

These changes were significant in both the 10,000 IU and 5000 IU treatment arms when compared to placebo in both CIS patients and HCPs ($p < 0.01$). Adjusting for baseline serum 25(OH)D levels, no significant differences in serum 25(OH)D levels achieved were seen between CIS patients and HCPs in any treatment arm. No cases of hypercalcaemia (reference range: 2.2–2.6 mmol/l) were recorded in any participant. Mean plasma PTH levels were lower in both treatment arms compared to placebo at all time-points in both CIS patients and HCPs.

Immunological responses to supplementation

The primary outcome of the study was the change in T cell subsets (IFN- γ ⁺CD4⁺/IL-17⁺CD4⁺ T cells) in response to supplementation. No significant reduction in either IFN- γ ⁺CD4⁺/IL-17⁺CD4⁺ T cells was seen in any treatment arm at any time-point regardless of method of stimulation (polyclonal or antigen specific with MOG/MBP peptides, recall antigens (PPD/TT) or irradiated allogeneic PBMCs). Results of polyclonal stimulation in both CIS patients and HCPs are presented in Table 2 and antigen-specific stimulation in CIS patients in Table 3. Similarly no significant differences were observed in concentrations as measured by ELISA of IL-10, IL-17 and IFN- γ produced following stimulation of PBMCs in any treatment arm (Supplementary

Table 2. Paired comparison between baseline and week 16 and baseline and week 24 of percentage interferon (IFN)- γ^+ CD4 $^+$ proliferating T cells and interleukin (IL)-17 $^+$ CD4 $^+$ proliferating T cells amongst each treatment arm in both clinically isolated syndrome (CIS) and healthy control participants (HCPs). Values are displayed as median and interquartile range.

	CIS					
	Week 0	Week 16	<i>p</i> -Value	Week 0	Week 24	<i>p</i> -Value
IFN- γ^+ CD4 $^+$ proliferating T cells						
10,000 IU	7.3 (1.2–10.3)	7 (2.3–8.6)	0.646	7.3 (1.2–10.3)	5.1 (2.5–7.6)	0.169
5,000 IU	5.8 (3.3–6.8)	4.9 (3.6–10.2)	0.594	5.8 (3.3–6.8)	6.4 (4.7–8.6)	0.674
Placebo	4.7 (4.3–5.6)	7 (4.3–9)	0.043	4.7 (4.3–5.6)	6 (1.1–7.3)	0.6
IL-17 $^+$ CD4 $^+$ proliferating T cells						
10,000 IU	0.8 (0.1–2)	1.3 (0.6–3.2)	0.015 ^a	0.8 (0.1–2)	0.7 (0.3–2.9)	0.484
5000 IU	1.1 (0.6–2.9)	1.4 (0.8–2.9)	0.374	1.1 (0.6–2.9)	1.7 (0.5–2.4)	0.767
Placebo	0.3 (0.2–0.9)	0.6 (0.2–1.9)	0.046 ^a	0.3 (0.2–0.9)	0.8 (0–13)	0.173
	HCPs					
	Week 0	Week 16	<i>p</i> -Value	Week 0	Week 24	<i>p</i> -Value
IFN- γ^+ CD4 $^+$ proliferating T cells						
10,000 IU	9.3 (7–12.6)	11.3 (3.3–13.1)	0.94	9.3 (7–12.6)	8.9 (4.5–12.3)	0.29
5000 IU	8.5 (5–10.9)	6.5 (3.1–11.5)	0.59	8.5 (5–10.9)	8.2 (4.8–12.3)	0.93
Placebo	4.9 (2.5–7.5)	5.9 (1.4–12.6)	0.9	4.9 (2.5–7.5)	6.2 (3.8–12.6)	0.04 ^b
IL-17 $^+$ CD4 $^+$ proliferating T cells						
10,000 IU	0.6 (0.4–0.9)	0.5 (0.3–1.2)	0.37	0.6 (0.4–0.9)	0.7 (0.5–1.6)	0.46
5000 IU	1 (0.5–1.6)	0.6 (0.3–0.9)	0.18	1 (0.5–1.6)	0.5 (0.3–1.7)	0.86
Placebo	0.8 (0.3–1.5)	0.8 (0.3–1.9)	0.75	0.8 (0.3–1.5)	1.3 (0.4–2.8)	0.64

^aA significant increase in IL-17 $^+$ CD4 $^+$ T cells seen at these time-points in the 10,000 IU and placebo groups; ^ba significant increase in IFN- γ^+ CD4 $^+$ T-cells was seen at this time-points in the placebo group.

Material, Table 1). Stratifying the groups based on the increase in serum 25(OH)D level (<50 nmol/L/ \geq 50 nmol/L) seen at each time-point showed a trend towards lower IFN- γ^+ CD4 $^+$ proliferating T cells in both CIS patients and HCPs with significance seen at week 16 in CIS ($p=0.01$) and week 16 ($p=0.03$) and week 24 ($p=0.02$) in HCP (Figure 3). All values were adjusted for baseline value but not for multiple comparisons.

MRI results

Baseline and 24-week MRI scans were available in 26 CIS patients. An incorrect or incomplete protocol was carried out in three patients. Baseline T2 lesions and changes on follow-up MRI are outlined in Table 4. No significant differences were seen between the number of new lesions, or the proportion of participants with new activity on follow-up MRI, between groups, controlling for both baseline T2 and gadolinium-enhancing lesions.

Clinical outcomes

One patient had a relapse during the course of the study. Her symptoms began one day prior to her week 24 visit. She had been allocated to the 5000 IU vitamin D₃ treatment group equating to mean ARR of 0.2 in this group. No patient fulfilled standard EDSS criteria for evidence of disability progression. No significant difference in EDSS between groups was seen at any time-point.

Combined clinical and radiological outcomes

Disease freedom at the end of the study as measured by percentage of participants with no evidence of MRI or clinical activity was 50% (5/10), 33% (3/9) and 57% (4/7) in the 10,000 IU, 5000 IU vitamin D₃ and placebo arms respectively. Adjusting for baseline serum 25(OH)D level and, comparing CIS patients with a serum 25(OH)D level of <100 nmol/l at week 24 to those with

Table 3. Paired comparison between baseline and week 16 and baseline and week 24 of percentage interferon (IFN)- γ^+ CD4 $^+$ proliferating T cells and interleukin (IL)-17 $^+$ CD4 $^+$ proliferating T cells from antigen-specific stimulated peripheral blood mononuclear cells (PBMCs) from clinically isolated syndrome (CIS) patients. Values are displayed as median and interquartile range.

	MOG/MBP stimulation					
	Week 0	Week 16	<i>p</i> -Value	Week 0	Week 24	<i>p</i> -Value
IFN- γ^+ CD4 $^+$ proliferating T cells						
10,000 IU	0.7 (0.1–1.1)	0.55 (0.2–1.9)	<i>p</i> = 0.45	0.7 (0.1–1.1)	1.1 (0.5–1.9)	<i>p</i> = 0.04 ^a
5000 IU	0.95 (0.3–4.4)	1.13 (0.7–1.9)	<i>p</i> = 0.39	0.95 (0.3–4.4)	1.3 (0.7–3.0)	<i>p</i> = 0.29
Placebo	0.54 (0.2–1.4)	0.6 (0.2–1.1)	<i>p</i> = 0.23	0.54 (0.2–1.4)	0.38 (0.2–0.9)	<i>p</i> = 0.17
IL-17 $^+$ CD4 $^+$ proliferating T cells						
10,000 IU	0.42 (0.1–1.0)	0.21 (0.1–0.7)	<i>p</i> = 0.24	0.42 (0.1–1.0)	0.36 (0.1–0.9)	<i>p</i> = 0.37
5000 IU	0.06 (0.03–0.2)	0.07 (0.03–0.3)	<i>p</i> = 0.4	0.06 (0.03–0.2)	0.29 (0.05–0.4)	<i>p</i> = 0.5
Placebo	0.1 (0.07–0.3)	0.37 (0.05–0.4)	<i>p</i> = 0.23	0.1 (0.07–0.3)	0.14 (0.1–0.7)	<i>p</i> = 0.9
	PPD stimulation					
	Week 0	Week 16	<i>p</i> -Value	Week 0	Week 24	<i>p</i> -Value
IFN- γ^+ CD4 $^+$ proliferating T cells						
10,000 IU	0.45 (0.2–2.1)	0.87 (0.2–2.6)	<i>p</i> = 0.44	0.45 (0.2–2.1)	1.2 (0.3–3.3)	<i>p</i> = 0.04 ^a
5000 IU	0.61 (0.4–1.7)	1.21 (0.6–2.1)	<i>p</i> = 0.29	0.61 (0.4–1.7)	1.3 (0.6–2.6)	<i>p</i> = 0.61
Placebo	0.69 (0.05–1.6)	1.19 (0.4–1.9)	<i>p</i> = 0.04 ^a	0.69 (0.05–1.6)	0.37 (0.1–2.9)	<i>p</i> = 0.46
IL-17 $^+$ CD4 $^+$ proliferating T cells						
10,000 IU	0.27 (0.7–3.0)	0.27 (0.1–1.6)	<i>p</i> = 0.7	0.27 (0.7–3.0)	0.38 (0.1–2.0)	<i>p</i> = 0.7
5000 IU	0.13 (0.01–0.4)	0.12 (0.04–0.3)	<i>p</i> = 0.88	0.13 (0.01–0.4)	0.22 (0.04–0.4)	<i>p</i> = 0.8
Placebo	0.49 (0.2–1.4)	0.56 (0.3–0.9)	<i>p</i> = 0.47	0.49 (0.2–1.4)	0.29 (0.1–1.1)	<i>p</i> = 0.23
	Allogeneic stimulation					
	Week 0	Week 16	<i>p</i> -Value	Week 0	Week 24	<i>p</i> -Value
IFN- γ^+ CD4 $^+$ proliferating T cells						
10,000 IU	1.24 (0.7–2.7)	1.1 (0.9–3.1)	<i>p</i> = 0.89	1.24 (0.7–2.7)	1.74 (0.6–2.9)	<i>p</i> = 0.95
5000 IU	4.5 (1.3 (1.3–7.7))	1.5 (0.9–2.7)	<i>p</i> = 0.02 ^b	4.5 (1.3–7.7)	1.75 (0.8–2.8)	<i>p</i> = 0.008 ^b
Placebo	0.89 (0.3–1.3)	1.1 (0.6–1.7)	<i>p</i> = 0.69	0.89 (0.3–1.3)	1.6 (0.1–1.7)	<i>p</i> = 0.35
IL-17 $^+$ CD4 $^+$ proliferating T cells						
10,000 IU	1.15 (0.3–1.8)	0.73 (0.2–1.6)	<i>p</i> = 0.31	1.15 (0.3–1.8)	0.14 (0.1–1.4)	<i>p</i> = 0.07
5000 IU	0.52 (0.2–1.2)	0.33 (0.2–0.5)	<i>p</i> = 0.14	0.52 (0.2–1.2)	0.43 (0.1–1.8)	<i>p</i> = 0.77
Placebo	0.2 (0.1–0.6)	0.51 (0.3–0.7)	<i>p</i> = 0.27	0.2 (0.1–0.6)	0.56 (0.3–0.8)	<i>p</i> = 0.35

^aA significant increase in IFN- γ^+ CD4 $^+$ proliferating T cells at that time point; ^ba significant reduction in IFN- γ^+ CD4 $^+$ proliferating T cells at that time point.

a level ≥ 100 nmol/l showed a disease freedom rate of 50% (7/14) compared to 67% (8/12) (*p* = 0.4).

Safety and adverse events

There was one serious adverse event recorded during the study; a patient required hospitalisation for a lumbar discectomy. This event was reported

immediately to the study sponsor and the patient withdrew from the study. However, the event was felt to be unrelated to the study drug. At the end of the study the patient was identified as being in the placebo group. A full adverse event listing is included as Supplementary Material, Tables 2 and 3.

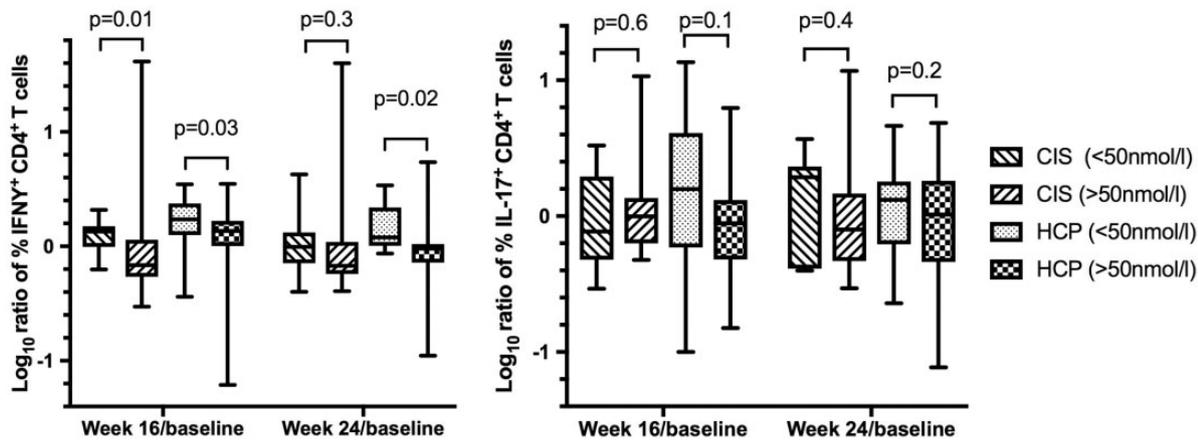


Figure 3. Comparison between change in interferon (IFN)- γ +CD4+ proliferating T cells, and interleukin (IL)-17+CD4+ proliferating T cells between baseline and week 16 and baseline and week 24 between CIS and HCP. Values are expressed as a log₁₀ ratio of week 16/baseline and week 24/baseline. The clinically isolated syndrome (CIS) and healthy control participant (HCP) groups have been stratified into two groups based on a change in serum 1,25 (OH) vitamin D (25(OH)D) of <50 nmol/l and \geq 50 nmol/l seen at each time-point.

Table 4. Baseline T2 lesions on magnetic resonance imaging (MRI) and change in follow-up scan at 24 weeks per randomised group.

	10,000 IU VD (n = 10)	5000 IU VD (n = 9)	Placebo (n = 7)
Baseline T2 lesions Mean (SD)	18 (17)	23 (11)	13 (10)
Baseline GD-enhancing lesions Mean (SD)	0.3 (0.7)	0.3 (0.5)	0.1 (0.4)
New T2 lesions Mean (SD)	0.4 (0.5)	1.4 (1.7)	0.6 (0.8)
New GD-enhancing lesions Mean (SD)	0	0.1 (0.3)	0.1 (0.4)
Number of patients with new disease activity by MRI (%)	5 (50%)	5 (56%)	3 (43%)

GD: gadolinium; SD: standard deviation; VD: vitamin D.

Discussion

This trial showed that vitamin D₃ supplementation at doses of 5000 IU and 10,000 IU per day is both safe and tolerable and results in significant increases in serum 25(OH)D levels in both healthy controls and CIS patients. It seems probable that serum 25(OH)D levels of greater than 100 nmol/l are required to produce the immunological effects of vitamin D, although no consensus level exists.²⁹ A diminished serologic response to vitamin D₃ supplementation has been described in MS patients compared to controls raising the possibility of different vitamin D pharmacokinetics in people with MS.³⁰ In our study, a higher mean change in serum 25(OH)D levels was seen in the control group but this was not statistically significant. Given the relatively small sample size it

was not possible to adjust for a number of confounders including age, sex and body mass index (s) across each of the treatment arms.

Analysis of the primary outcome showed no difference in the frequency of pro-inflammatory CD4⁺ T cells (IL-17⁺CD4⁺/IFN- γ ⁺CD4⁺) between baseline and either time-point in any of the treatment arms. No difference was seen between randomisation groups or in response to varying methods of PBMC stimulation (polyclonal or antigen-specific). A significant reduction in proliferating IFN- γ ⁺CD4⁺ T-cells was seen in both control participants (week 16 and 24) and CIS patients (week 16) when the groups were stratified according to change in serum 25(OH)D level but these results were not

adjusted for multiple comparisons and need to be interpreted with caution. Measurement of anti-inflammatory (IL-10) and pro-inflammatory (IL-17/IFN- γ) cytokines from stimulated PBMCs by ELISA also failed to show any significant changes in response to supplementation or between randomisation groups.

The role of IL-17⁺CD4⁺ T cells in MS pathogenesis has been shown in the experimental autoimmune encephalitis (EAE) model, and is also supported by high levels of IL-17 in MS plaques and cerebrospinal fluid.³¹ *In vitro* studies have suggested that vitamin D significantly inhibits IL-17 production and increases IL-10 in relapsing remitting MS (RRMS) patients and controls.¹¹ However, the results of a small number of published clinical trials exploring various immunological outcomes in response to vitamin D supplementation in MS have been conflicting and somewhat disappointing in identifying the underlying molecular immunological mechanism of vitamin D supplementation.^{17–23} A recent study by Sotirchos *et al.* showed a reduction in IL-17⁺CD4⁺ T cells at week 24 in RRMS patients receiving high dose (10,400 IU/day) vitamin D₃.¹⁷ These effects were not observed in the low dose group (800 IU/day) group and no significant change was seen in IFN- γ ⁺CD4⁺ T cells in either group. A Dutch sub-study of a larger clinical trial exploring high dose (up to 14,000 IU/day) vitamin D₃ supplementation, as an add-on therapy to interferon β -1a in patients with RRMS, showed no difference in either IL-17⁺CD4⁺ or IFN- γ ⁺CD4⁺ T cells at 48 weeks follow-up.²² The investigators found a reduction in the proportion of anti-inflammatory IL-4⁺CD4⁺ T cells in the placebo group and suggested that supplementation may contribute to immune homeostasis early in the disease course.

There were methodological differences in the processing of PBMCs between these studies and our own which may have contributed towards the different percentages of IL-17⁺CD4⁺ and IFN- γ ⁺CD4⁺ T cells seen. The experimental design of our study was in part similar to that of Sotirchos *et al.* in that intracellular cytokines were measured by restimulation after an initial *in vitro* stimulation period using cryopreserved PBMC. However, we cultured cells for three rather than five days with anti-CD3, and included additional antigen-specific stimuli. The rationale for including antigen-specific stimuli was that vitamin D₃ is likely to exert at least some of its anti-inflammatory effects on T cells via antigen-presenting cells, and such effects may not be apparent after polyclonal stimulation. The allogeneic stimulation would be expected to stimulate naïve allo-specific

T cells and would therefore reveal possible effects of vitamin D₃ on T cell differentiation, whereas recall antigens PPD/TT or putative MS auto-antigens MOG/MBP would restimulate antigen-specific memory T cells. In contrast, Muris *et al.* measured intracellular cytokine production directly *ex vivo* without cryopreservation or an *in vitro* stimulation period.²² Although our study was designed to examine the effects of vitamin D₃ supplementation on T cell responses either directly or indirectly via the antigen presenting cell, it is possible that vitamin D₃ may also exert other immunologic effects that were not examined here. It has been suggested that vitamin D may have immunomodulatory effects on B cells³², regulatory T cells³³ and dendritic cells.³⁴

The discordance between published results to date may suggest alternative hypotheses as to the observed association between vitamin D deficiency and MS. It has been argued that low serum 25(OH)D levels seen early in the disease course are a result of the disease itself and supplementation is unlikely to affect disease activity.³⁵ Alternative environmental factors need to be considered. Ultraviolet exposure has been shown to have immunomodulating effects on skin lymphocytes³⁶ and thus, observed vitamin D deficiency may only be a marker of lack of sunlight exposure.

Exploration of secondary endpoints showed no difference between treatment arms and placebo with regard to either radiological or clinical activity. A higher level of disease freedom (67% versus 50%) was seen in patients with serum 25(OH)D levels >100 nmol/l but this was not significant. The trial was not powered to meet this endpoint and the trial duration was probably too short to detect a treatment effect of vitamin D₃ supplementation.

It is possible that high dose vitamin D₃ therapy on its own may not be sufficiently effective as an anti-inflammatory or immunomodulatory therapy in MS or CIS. There is convincing evidence, from a post-hoc analysis, that higher serum 25(OH)D levels in patients recruited into the BENEFIT and BEYOND studies had additional therapeutic effects on disease activity in patients on betaferon-1b.^{37,38} Recent results from the SOLAR study of high dose vitamin D₃ supplementation as an add-on therapy to interferon β -1a, showed a non-significant reduction in the annualised relapse rate of 30% in the vitamin D₃ group at 48 weeks.³⁹ A significant reduction in both unique new lesions and percentage change in T2 lesion volume on MRI was also seen suggesting a therapeutic benefit of supplementation. That study was hampered by difficulty with

recruitment leading to a reduced trial duration. Longer follow-up and larger numbers are needed to confirm these findings. A number of trials are ongoing to address this issue.^{40,41}

The strengths of this study are that it was a double-blind randomised controlled trial and the first clinical trial of vitamin D₃ therapy in treatment-naïve CIS patients. The HCP group allowed us to establish if immunological effects observed were unique to a CIS population. There were limitations to our study. Power calculations were based on a small pilot study²⁸ and may have resulted in the study being underpowered. Difficulty with recruitment in the CIS group also led to smaller numbers than originally anticipated. The CIS group had a higher mean age than controls, which is a potential confounder when comparing results of some of the secondary immunologic outcomes and vitamin D pharmacokinetics. Our results were adjusted for baseline values but not for multiple comparisons, which may limit their interpretability.

Conclusion

In conclusion, this study demonstrates that vitamin D₃ supplementation at doses up to 10,000 IU daily is safe and tolerable in both HPCs and CIS patients. High dose vitamin D₃ supplementation did not reduce pro-inflammatory IL-17⁺CD4⁺ T cells; there were possible minor effects on IFN- γ ⁺ CD4⁺ T cells, but no changes were observed in concentrations of IL-10, IL-17 or IFN- γ produced by stimulated PBMCs. The potential role of vitamin D₃ as an immunomodulator needs to be further clarified in larger randomised controlled trials.

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Conflicts of interest

The author(s) declared the following potential conflicts of interest with respect to the research, authorship, and/or publication of this article: Merck KGaA (Darmstadt, Germany) reviewed the manuscript for medical accuracy only before journal submission. The authors are fully responsible for the content of this manuscript, and the views and opinions described in the publication reflect solely those of the authors.

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Supplemental Data

Supplementary material is available for this article online.

References

1. Compston A and Coles A. Multiple sclerosis. *Lancet* 2008; 372: 1502–1517.
2. Simpson S Jr, Blizzard L, Othahal P, et al. Latitude is significantly associated with the prevalence of multiple sclerosis: A meta-analysis. *J Neurol Neurosurg Psychiatry* 2011; 82: 1132–1141.
3. Ramagopalan SV, Handel AE, Giovannoni G, et al. Relationship of UV exposure to prevalence of multiple sclerosis in England. *Neurology* 2011; 76: 1410–1414.
4. Orton SM, Wald L, Confavreux C, et al. Association of UV radiation with multiple sclerosis prevalence and sex ratio in France. *Neurology* 2011; 76: 425–431.
5. Salzer J, Hallmans G, Nystrom M, et al. Vitamin D as a protective factor in multiple sclerosis. *Neurology* 2012; 79: 2140–2145.
6. Munger KL, Aivo J, Hongell K, et al. Vitamin D status during pregnancy and risk of multiple sclerosis in offspring of women in the finnish maternity cohort. *JAMA Neurol* 2016; 73: 515–519.
7. Dobson R, Giovannoni G and Ramagopalan S. The month of birth effect in multiple sclerosis: Systematic review, meta-analysis and effect of latitude. *J Neurol Neurosurg Psychiatry* 2013; 84: 427–432.
8. Simpson S Jr, Taylor B, Blizzard L, et al. Higher 25-hydroxyvitamin D is associated with lower relapse risk in multiple sclerosis. *Ann Neurol* 2010; 68: 193–203.
9. Mowry EM, Krupp LB, Milazzo M, et al. Vitamin D status is associated with relapse rate in pediatric-onset multiple sclerosis. *Ann Neurol* 2010; 67: 618–624.
10. Mowry EM, Waubant E, McCulloch CE, et al. Vitamin D status predicts new brain magnetic resonance imaging activity in multiple sclerosis. *Ann Neurol* 2012; 72: 234–240.
11. Correale J, Ysrraelit MC and Gaitan MI. Immunomodulatory effects of Vitamin D in multiple sclerosis. *Brain* 2009; 132: 1146–1160.
12. Adorini L and Penna G. Control of autoimmune diseases by the vitamin D endocrine system. *Nat Clin Pract Rheumatol* 2008; 4: 404–412.
13. Smolders J, Menheere P, Kessels A, et al. Association of vitamin D metabolite levels with relapse rate and

- disability in multiple sclerosis. *Mult Scler* 2008; 14: 1220–1224.
14. Bhalla AK, Amento EP, Serog B, et al. 1,25-Dihydroxyvitamin D3 inhibits antigen-induced T cell activation. *J Immunol* 1984; 133: 1748–1754.
 15. Chen S, Sims GP, Chen XX, et al. Modulatory effects of 1,25-dihydroxyvitamin D3 on human B cell differentiation. *J Immunol* 2007; 179: 1634–1647.
 16. Boonstra A, Barrat FJ, Crain C, et al. 1 α ,25-dihydroxyvitamin d3 has a direct effect on naive CD4(+) T cells to enhance the development of Th2 cells. *J Immunol* 2001; 167: 4974–4980.
 17. Sotirchos ES, Bhargava P, Eckstein C, et al. Safety and immunologic effects of high- vs low-dose cholecalciferol in multiple sclerosis. *Neurology* 2016; 86: 382–390.
 18. Kimball S, Vieth R, Dosch HM, et al. Cholecalciferol plus calcium suppresses abnormal PBMC reactivity in patients with multiple sclerosis. *J Clin Endocrinol Metab* 2011; 96: 2826–2834.
 19. Smolders J, Peelen E, Thewissen M, et al. Safety and T cell modulating effects of high dose vitamin D3 supplementation in multiple sclerosis. *PLoS One* 2010; 5: e15235.
 20. Ashtari F, Toghianifar N, Zarkesh-Esfahani SH, et al. Short-term effect of high-dose vitamin D on the level of interleukin 10 in patients with multiple sclerosis: A randomized, double-blind, placebo-controlled clinical trial. *Neuroimmunomodulation* 2015; 22: 400–404.
 21. Toghianifar N, Ashtari F, Zarkesh-Esfahani SH, et al. Effect of high dose vitamin D intake on interleukin-17 levels in multiple sclerosis: A randomized, double-blind, placebo-controlled clinical trial. *J Neuroimmunol* 2015; 285: 125–128.
 22. Muris AH, Smolders J, Rolf L, et al. Immune regulatory effects of high dose vitamin D3 supplementation in a randomized controlled trial in relapsing remitting multiple sclerosis patients receiving IFN β ; the SOLARIUM study. *J Neuroimmunol* 2016; 300: 47–56.
 23. Aivo J, Hanninen A, Ilonen J, et al. Vitamin D3 administration to MS patients leads to increased serum levels of latency activated peptide (LAP) of TGF- β . *J Neuroimmunol* 2015; 280: 12–15.
 24. Polman CH, Reingold SC, Banwell B, et al. Diagnostic criteria for multiple sclerosis: 2010 Revisions to the McDonald criteria. *Ann Neurol* 2011; 69: 292–302.
 25. Miller DH, Chard DT and Ciccarelli O. Clinically isolated syndromes. *Lancet Neurol* 2012; 11: 157–169.
 26. Lonergan R, Kinsella K, Fitzpatrick P, et al. Multiple sclerosis prevalence in Ireland: Relationship to vitamin D status and HLA genotype. *J Neurol Neurosurg Psychiatry* 2011; 82: 317–322.
 27. O'Connell K, Kelly S, Kinsella K, et al. Dose-related effects of vitamin D on immune responses in patients with clinically isolated syndrome and healthy control participants: Study protocol for an exploratory randomized double-blind placebo-controlled trial. *Trials* 2013; 14: 272.
 28. Allen AC, Kelly S, Basdeo SA, et al. A pilot study of the immunological effects of high-dose vitamin D in healthy volunteers. *Mult Scler* 2012; 18: 1797–1800.
 29. Munger KL, Levin LI, Hollis BW, et al. Serum 25-hydroxyvitamin D levels and risk of multiple sclerosis. *JAMA* 2006; 296: 2832–2838.
 30. Bhargava P, Steele SU, Waubant E, et al. Multiple sclerosis patients have a diminished serologic response to vitamin D supplementation compared to healthy controls. *Mult Scler* 2016; 22: 753–760.
 31. Fletcher JM, Lalor SJ, Sweeney CM, et al. T cells in multiple sclerosis and experimental autoimmune encephalomyelitis. *Clin Exp Immunol* 2010; 162: 1–11.
 32. Rolf L, Muris AH, Hupperts R, et al. Illuminating vitamin D effects on B-cells – the multiple sclerosis perspective. *Immunology* 2016; 147: 275–284.
 33. Prietl B, Treiber G, Mader JK, et al. High-dose cholecalciferol supplementation significantly increases peripheral CD4(+) Tregs in healthy adults without negatively affecting the frequency of other immune cells. *Eur J Nutr* 2014; 53: 751–759.
 34. Bscheider M and Butcher EC. Vitamin D immunoregulation through dendritic cells. *Immunology* 2016; 148: 227–236.
 35. Ghashut RA, Talwar D, Kinsella J, et al. The effect of the systemic inflammatory response on plasma vitamin 25 (OH) D concentrations adjusted for albumin. *PLoS One* 2014; 9: e92614.
 36. Maverakis E, Miyamura Y, Bowen MP, et al. Light, including ultraviolet. *J Autoimmun* 2010; 34: J247–J257.
 37. Fitzgerald KC, Munger KL, Kochert K, et al. Association of vitamin D levels with multiple sclerosis activity and progression in patients receiving interferon beta-1b. *JAMA Neurol* 2015; 72: 1458–1465.
 38. Ascherio A, Munger KL, White R, et al. Vitamin D as an early predictor of multiple sclerosis activity and progression. *JAMA Neurol* 2014; 71: 306–314.
 39. Smolders J, Vieth R, Holmøy T, et al. High dose cholecalciferol (vitamin D3) oil as add-on therapy in subjects with relapsing–remitting multiple sclerosis receiving subcutaneous interferonB-1a. *Mult Scler* 2016; 22.
 40. Bhargava P, Cassard S, Steele SU, et al. The vitamin D to ameliorate multiple sclerosis (VIDAMS) trial: Study design for a multicenter, randomized, double-blind controlled trial of vitamin D in multiple sclerosis. *Contemp Clin Trials* 2014; 39: 288–293.
 41. Dorr J, Ohlraun S, Skarabis H, et al. Efficacy of vitamin D supplementation in multiple sclerosis (EVIDIMS Trial): Study protocol for a randomized controlled trial. *Trials* 2012; 13: 15.