

# Mendelian randomization shows a causal effect of low vitamin D on multiple sclerosis risk

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## ABSTRACT

**Objective:** We sought to estimate the causal effect of low serum 25(OH)D on multiple sclerosis (MS) susceptibility that is not confounded by environmental or lifestyle factors or subject to reverse causality.

**Methods:** We conducted mendelian randomization (MR) analyses using an instrumental variable (IV) comprising 3 single nucleotide polymorphisms found to be associated with serum 25(OH)D levels at genome-wide significance. We analyzed the effect of the IV on MS risk and both age at onset and disease severity in 2 separate populations using logistic regression models that controlled for sex, year of birth, smoking, education, genetic ancestry, body mass index at age 18–20 years or in 20s, a weighted genetic risk score for 110 known MS-associated variants, and the presence of one or more *HLA-DRB1\*15:01* alleles.

**Results:** Findings from MR analyses using the IV showed increasing levels of 25(OH)D are associated with a decreased risk of MS in both populations. In white, non-Hispanic members of Kaiser Permanente Northern California (1,056 MS cases and 9,015 controls), the odds ratio (OR) was 0.79 ( $p = 0.04$ , 95% confidence interval (CI): 0.64–0.99). In members of a Swedish population from the Epidemiological Investigation of Multiple Sclerosis and Genes and Environment in Multiple Sclerosis MS case-control studies (6,335 cases and 5,762 controls), the OR was 0.86 ( $p = 0.03$ , 95% CI: 0.76–0.98). A meta-analysis of the 2 populations gave a combined OR of 0.85 ( $p = 0.003$ , 95% CI: 0.76–0.94). No association was observed for age at onset or disease severity.

**Conclusions:** These results provide strong evidence that low serum 25(OH)D concentration is a cause of MS, independent of established risk factors. *Neurol Genet* 2016;2:e97; doi: 10.1212/NXG.000000000000097

## GLOSSARY

**CI** = confidence interval; **EHR** = electronic health record; **EIMS** = Epidemiological Investigation of Multiple Sclerosis; **GERA** = Genetic Epidemiology Research on Adult Health and Aging; **GEMS** = Genes and Environment in Multiple Sclerosis; **GWAS** = genome-wide association study; **HWE** = Hardy-Weinberg equilibrium; **ICD-9** = *International Classification of Diseases, 9th Revision*; **IV** = instrumental variable; **KPNC** = Kaiser Permanente in Northern California; **LD** = linkage disequilibrium; **MAF** = minor allele frequency; **MDS** = multidimensional scaling; **MR** = mendelian randomization; **MS** = multiple sclerosis; **MSSS** = Multiple Sclerosis Severity Scores; **SNP** = single nucleotide polymorphism; **VDRE** = vitamin D response element; **wGRS** = weighted genetic risk score.

Multiple sclerosis (MS) is an immune-mediated, demyelinating disease that leads to a wide variety of symptoms and disability. Both genetic and environmental factors have been implicated in its etiology, including vitamin D deficiency. Observational studies have consistently shown an association of low serum 25(OH)D and increased risk of MS, but it has not been shown that low 25(OH)D is actually a cause of MS.<sup>1</sup> The apparent beneficial effects of 25(OH)D on MS might

Supplemental data  
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alternately be explained by reverse causation (i.e., MS could be leading to low 25(OH)D) or by confounding by sun exposure, obesity, or some other unknown factors.

Mendelian randomization (MR), equivalently, instrumental variable (IV) analysis using a genetic instrument, is a technique that can overcome the problems of both reverse causation and confounding when assessing the causal relationship between an exposure and an outcome.<sup>2</sup> Single nucleotide polymorphisms (SNPs) known to be associated with 25(OH)D levels, rather than measured 25(OH)D, can be used as an IV to estimate the effect of low 25(OH)D on MS. Because SNP genotypes are determined at birth and are not likely to be influenced by potential confounding variables, the effect estimate from MR analysis should not be confounded, and reverse causation is unlikely because MS does not determine which 25(OH)D-associated SNPs are inherited (figure). We used MR analysis to estimate the causal relationship between serum 25(OH)D levels and MS susceptibility in 2 large case-control studies. We also investigated 2 clinical phenotypes for MS: age at onset and disease severity.

**METHODS KPNC participants.** Data were collected from members of Kaiser Permanente Medical Care Plan, Northern California Region (KPNC). KPNC is an integrated health service delivery system with a membership of 3.2 million that comprises

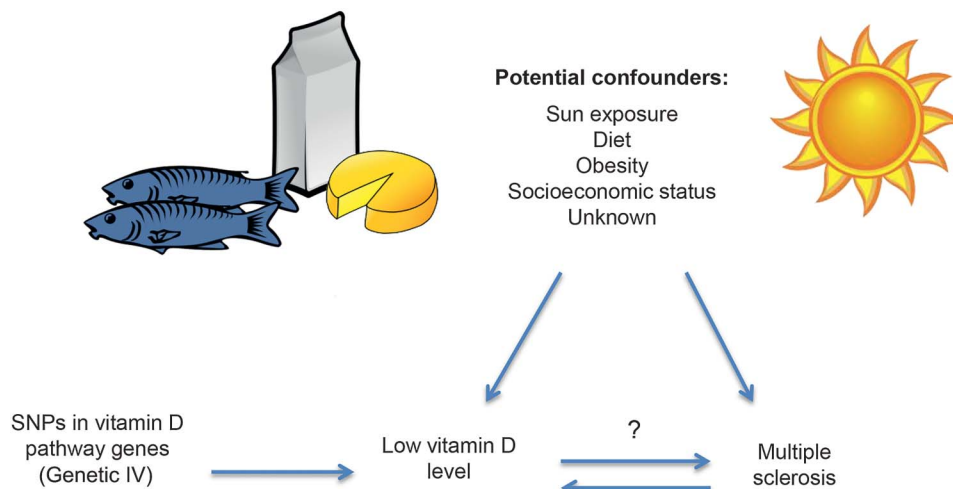
about 25–30% of the population of a 22-county service area and is the largest health care provider in northern California. Membership is largely representative of the general population in the service area; however, persons in impoverished neighborhoods are underrepresented.<sup>3</sup>

Eligible KPNC cases were defined as individuals with a diagnosis of MS by a neurologist (*ICD-9* code 340.xx), age 18–69 years, and membership in KPNC at initial contact. The study was restricted to self-identified white (non-Hispanic) race/ethnicity, the population with the highest prevalence of MS. The treating neurologist was contacted for approval to contact each case as a potential MS study participant. A total of 3,293 potential MS cases were reviewed by KPNC neurologists, who approved contact with 2,823 (86%) at the time of the data freeze (August 2014). Diagnoses were validated using electronic health record (EHR) review and according to published diagnostic criteria.<sup>4</sup> Multiple Sclerosis Severity Scores (MSSS) were calculated for each case at the time of study entry (mean disease duration = 17.7 years), as described,<sup>5</sup> and participants were asked to recall the age of first MS symptom onset which was validated using EHR data when possible.

Controls were white (non-Hispanic) current KPNC members without a diagnosis of MS or related condition (optic neuritis, transverse myelitis, or demyelinating disease; *ICD-9* codes: 340, 341.0, 341.1, 341.2, 341.20, 341.21, 341.22, 341.8, 341.9, 377.3, 377.30, 377.39, and 328.82) confirmed through EHR data. Potential study participants were contacted by email with a follow-up phone call. The participation rate was 80% for cases and 66% for controls. Genetic data were available for approximately 80% of study participants.

Additional controls were individuals of the Genetic Epidemiology Research on Adult Health and Aging (GERA) cohort participating in the KPNC Research Program on Genes, Environment, and Health, which is described elsewhere (dbGaP phs000674.v2.p2).<sup>6,7</sup> Respondents completed a written consent form and provided a saliva sample for DNA extraction. A total of 110,266 participant samples were initially collected. Approximately 103,000 samples were successfully genotyped, and 77% of participants subsequently returned new consent forms for placement in dbGaP (NIH), resulting in a final sample size of 78,486 participants. From these

**Figure** Relationship of exposure, outcome, confounding variables, and genetic instrumental variable used in mendelian randomization



IV = instrumental variable; SNP = single nucleotide polymorphism.

participants, we selected a subset of 12,605 self-reported non-Hispanic white individuals matched to MS cases for sex and age ( $\pm 2$  years) at a 10:1 ratio.

**EIMS/GEMS participants.** Data were collected from 2 population-based case-control studies on incident MS patients (the Epidemiological Investigation of Multiple Sclerosis [EIMS] study) and prevalent MS patients (the Genes and Environment in Multiple Sclerosis [GEMS] study). In the EIMS study, the inclusion criteria were age 16–70 years, recently (within 2 years) diagnosed MS according to the McDonald criteria,<sup>4</sup> and ability to understand Swedish. For the GEMS study, the participants were identified from the Swedish National MS registry and recruited during 2009–2011. In both studies, controls (without MS) were randomly chosen from the population register and matched to the patients with MS by sex, age at inclusion in the study, and region of residence (2 controls per case in EIMS and 1 control per case in GEMS). Information on age at onset was self-reported. Data on MSSS<sup>5</sup> were retrieved from the Swedish National MS registry for all participants, in both EIMS and GEMS studies as per September 2014 (mean disease duration = 20.1 years). All participants in the EIMS study were distinct from those in the GEMS study. Details of the study design have been described elsewhere.<sup>8,9</sup> The participation rate was 92% for cases and 67% for controls in EIMS, and 82% for cases and 66% for controls in GEMS. Genotyping data were available for 75% of EIMS and 91% of GEMS participants.

**Standard protocol approvals, registrations, and patient consents.** Study protocols for KPNC participants were approved by the Institutional Review Boards of KPNC and the University of California, Berkeley. Ethical approval for both EIMS and GEMS was obtained from the Regional Ethical Review Board in Stockholm at Karolinska Institutet, and participants provided written informed consent.

**Exposure assessment in KPNC.** KPNC participants completed a computer-assisted telephone interview comprised questions related to various events and exposures, as described elsewhere.<sup>10,11</sup> GERA controls were mailed a survey consisting of questions related to health behaviors, sociodemographic information, and diagnoses (dbGaP phs000674.v2.p2).

Whole blood was collected and processed using the Gentra Puregene protocol. Saliva was collected using Oragene kits. Medium resolution *HLA-DRB1* and genome-wide SNP genotyping were performed as previously described<sup>12,13</sup> using Illumina Infinium 660K and Human Omni Express BeadChip arrays for KPNC cases and controls, and Axiom (Affymetrix) custom chips for the additional GERA controls. SNPs with a minor allele frequency (MAF) <1%, success rate <90%, or not in Hardy-Weinberg equilibrium (HWE) among controls ( $p < 0.000001$ ) were removed from analysis. Samples with >10% failed genotype

calls and related individuals were also removed. Imputation against reference haplotypes from Phase I of the 1000 Genomes Project was conducted using SHAPEIT and IMPUTE2. To ensure high-quality imputation results, only SNPs with info score >0.8 on all 3 genotyping platforms and with MAF in controls with SD <0.03 across all 3 platforms were retained for analysis. Cross-platform association tests were conducted to remove SNPs associated with the genotype array (FDR  $q < 0.05$ ). Population outliers were identified using multidimensional scaling (MDS) and removed from analysis. Two MDS components were used to adjust for population stratification due to genetic ancestry differences between cases and controls in the association analysis.

**Exposure assessment in EIMS and GEMS.** The exposure assessment was done through an extensive questionnaire that covered demographic and environmental factors; details are described elsewhere.<sup>8,9</sup> Participants were asked to provide blood samples, which were genotyped on an Illumina custom array (MS replication chip). *HLA-DRB1* information was imputed with a modified version of HLA\*IMP:02<sup>14</sup> with 400 Swedish controls added to the reference panel for MHC class II genes. SNPs with <2% MAF, success rate <98%, or not in HWE among controls ( $p < 0.0001$ ) were removed from analysis. Individuals with >2% failed genotype calls, with increased heterozygosity ( $> \text{mean} + 2\text{SD}$ ), related individuals, or whose recorded sex differed from genotype were removed from analysis. We removed population outliers identified using the smartpca program<sup>15</sup> with standard settings using 3,736 ancestry informative markers. Two principal component analysis vectors, those with  $p < 0.05$ , were used to adjust for population stratification in the association analysis, similar to KPNC comparisons described above.

**Statistical analyses.** A 25(OH)D IV was constructed for each individual using the  $\beta$  coefficients for 3 published genetic variants associated at genome-wide significance with higher serum 25(OH)D: rs2282679-A, in an intron of *GC*; rs2060793-A, upstream of *CYP2R1*; and rs3829251-G, in an intron of *NADSYN1* and upstream of *DHCR7*.<sup>16</sup> The IV was calculated by multiplying the number of alleles that correspond to an increase in 25(OH)D for each locus by the  $\beta$  coefficient from the genome-wide association study (GWAS) for that variant and then taking the sum across the 3 loci (table 1). For KPNC, either the genotyped alleles or imputed allele probabilities were used to calculate the IV. For EIMS/GEMS, genotyped alleles were used to calculate the IV, and missing genotypes (2 for rs2282679 and 243 for rs3822951) were replaced with 2 times the MAF. In addition, rs10741657, which is in perfect linkage disequilibrium (LD) with rs2060793 ( $r^2 = 1$  in 1,000 Genomes Phase I European population), was used in the IV calculation for EIMS/GEMS, as the latter SNP was not genotyped. A weighted genetic risk score (wGRS) for 110 non-HLA MS susceptibility loci identified through the most recent MS GWAS

**Table 1** Characteristics of SNPs used to construct the 25(OH)D instrumental variable

SNP	Allele associated with increasing 25(OH)D	Allele frequency	Chromosome	Gene	Weight ( $\beta$ coefficient from GWAS of serum 25(OH)D)
rs2282679	A	0.74	4	GC	0.38
rs2060793; rs10741657	A; A	0.41	11	CYP2R1	0.25
rs3829251	G	0.81	11	NADSYN1/DHCR7	0.18

Abbreviations: EIMS = Epidemiological Investigation of Multiple Sclerosis; GEMS = Genes and Environment in Multiple Sclerosis; GWAS = genome-wide association study; KPNC = Kaiser Permanente in Northern California; SNP = single nucleotide polymorphism. The 2 SNPs near the *CYP2R1* gene are in linkage disequilibrium in European populations; rs2060793 was used in KPNC, and rs10741657 was used in EIMS/GEMS. Allele frequencies are those reported in the 25(OH)D GWAS.

was calculated for each individual.<sup>17</sup> The wGRS was calculated by multiplying the number of risk alleles for each locus by the logarithm of the odds ratio (OR) for that variant, then taking the sum across the 110 loci. One SNP was missing for KPNC (rs201202118), and 2 SNPs were missing for EIMS/GEMS (rs2028597 and rs6874308).

After quality control and removal of population outliers, 1,655 individuals from KPNC (1,056 cases and 599 controls) with genetic data were available as well as an additional 8,416 controls from the GERA study, for a total of 1,056 cases and 9,015 controls. Data for 6,335 cases and 5,762 controls were available from the Swedish studies (EIMS and GEMS).

Demographic differences between cases and controls were compared using  $\chi^2$  and independent sample *t* tests where appropriate. Linear regression was used to test the assumption that the IV is not associated with confounding factors. MR analysis, in this case a separate-sample IV analysis, was performed by regressing MS case status on the 25(OH)D IV in logistic models, and regressing MSSS and age at onset on the 25(OH)D IV in linear regression models. All analyses were controlled for sex, year of birth, ever smoking, college education, *HLA-DRB1\*15:01* carrier status, wGRS of non-HLA MS risk variants, and genetic ancestry (as derived from the first 2 components from MDS for KPNC or smartpca for Sweden). Swedish analyses were additionally controlled for region of residency and study type (EIMS vs GEMS). Matching variables were included as covariates because the final study populations included all participants with genotyping data available, and a matched analysis was not performed. ORs with 95% confidence intervals (CIs) were estimated. Meta-analysis assuming random effects was performed.

All analyses were conducted using PLINK,<sup>18</sup> STATA (StataCorp, College Station, TX), SAS (SAS Institute, Cary, NC), or R.<sup>19</sup>

**RESULTS** Demographic and disease characteristics of MS cases and controls are found in table 2 for both KPNC and Sweden studies. There were significant differences between cases and controls with respect to

smoking, college graduation, obesity at age 18–20 years or in 20s, and *HLA-DRB1\*15:01* status for both KPNC and EIMS/GEMS, as expected, and for age and sex in EIMS/GEMS. The 25(OH)D IV was lower among cases than that in controls in KPNC (1.026 vs 1.045, respectively; *p* = 0.049) and in EIMS/GEMS (1.044 vs 1.056, respectively; *p* = 0.043). In both KPNC and EIMS/GEMS, the IV was not associated with year of birth, sex, smoking, college education, obesity at age 18–20 years or in 20s, *HLA-DRB1\*15:01* status, or the wGRS of 110 non-HLA MS risk variants, or with study type in the Swedish population. In EIMS/GEMS, it was associated with 2 of the 6 geographic regions.

The 25(OH)D IV was associated with MS susceptibility after controlling for sex, year of birth, ancestry, smoking, wGRS of 110 non-HLA MS risk variants, and number of *HLA-DRB1\*15:01* alleles, as well as study type and region of residency in the Swedish study. An increasing IV score corresponding to increasing 25(OH)D levels was protective against MS (OR 0.79, 95% CI: 0.64–0.99; *p* = 0.04) in the KPNC study. An association was also found for the Swedish study (OR 0.86, 95% CI: 0.76–0.98; *p* = 0.03). Adjustment did not alter the findings substantially; however, for both KPNC and EIMS/GEMS data sets, *p* values and ORs were both lower in the full models compared with those in the unadjusted models (data not shown). Expanded models are shown for KPNC (table e-1 at [Neurology.org/ng](http://Neurology.org/ng)) and EIMS/GEMS (table e-2). Both studies were combined into a meta-analysis: the association remained (OR 0.85, 95% CI: 0.76–0.94; *p* = 0.003). No evidence of

**Table 2** Population characteristics for the KPNC and EIMS/GEMS study participants

Characteristic	KPNC MS cases (N = 1,056)	KPNC controls (N = 9,015)	<i>p</i> Value	EIMS/GEMS MS cases (N = 6,335)	EIMS/GEMS controls (N = 5,762)	<i>p</i> Value
Females:males	841:215	7,328:1,687	0.196	4,162:1,723	4,378:1,384	<0.001
Year of birth	1958 ± 8.8	1958 ± 8.9	0.246	1960 ± 13.1	1961 ± 13.3	0.021
Body mass index <sup>a</sup>	23.0 ± 4.4	21.5 ± 3.3	<0.001	22.0 ± 3.5	21.7 ± 3.2	<0.001
College graduate			<0.001			0.002
Yes	468 (44)	3,141 (35)		4,521 (71)	3,963 (69)	
No	588 (56)	5,874 (65)		1,814 (29)	1,799 (31)	
Smoker			<0.001			<0.001
Ever	525 (50)	2,876 (32)		3,578 (56)	2,742 (48)	
Never	531 (50)	6,139 (68)		2,757 (44)	3,020 (52)	
<i>HLA-DRB1*15:01</i>			<0.001			<0.001
0 copies	498 (47)	6,613 (73)		2,615 (41)	4,095 (71)	
1–2 copies	558 (53)	2,402 (27)		3,720 (59)	1,667 (29)	

Abbreviations: EIMS = Epidemiological Investigation of Multiple Sclerosis; GEMS = Genes and Environment in Multiple Sclerosis; KPNC = Kaiser Permanente in Northern California; MS = multiple sclerosis.

Values are mean ± SD or N (%). *p* Values for differences between cases and controls are from Student *t* tests or  $\chi^2$  tests.

<sup>a</sup>At age 18 years or in 20s (KPNC), or at age 20 years (EIMS/GEMS).



heterogeneity between populations was observed ( $I^2 = 0.0\%$ , heterogeneity  $\chi^2 p = 0.51$ ). Results are summarized in table 3. The 25(OH)D IV was not associated with MSSS or age at onset in either study (data not shown).

**Validating additional assumptions.** To test the MR model assumptions for the 25(OH)D IV, we conducted overidentification tests to evaluate the null hypothesis that effect estimates from multiple IVs are identical.<sup>20</sup> Each variant was split into a separate instrument. Estimates suggested the same direction of causal effect, with ORs ranging from 0.92 to 0.94 in KPNC and 0.84 to 0.99 in EIMS/GEMS (table 4).

**DISCUSSION** This study, using data from 2 different populations, provides strong evidence that low 25(OH)D is causally associated with MS susceptibility. Numerous observational studies have shown an association between low serum 25(OH)D level and increased risk of MS, and some have provided evidence supporting a causal association. In a study that examined patients with clinically isolated syndrome, a first event suggestive of MS, low 25(OH)D levels early in the disease course predicted higher MS activity.<sup>21</sup> Two prospective, nested case-control studies, which collected serum prior to MS onset, showed a significantly reduced risk of MS in those with high 25(OH)D.<sup>22,23</sup> Results from the current study rule out the most plausible alternative (noncausal) interpretations of these observational studies, thus strengthening evidence of a causal link between 25(OH)D and MS, even after accounting for other risk factors and potential confounders.

Molecular studies also suggest that low serum 25(OH)D may be a causal risk factor for MS. Studies examining the location of vitamin D receptor-binding sites (also known as vitamin D response elements or VDREs) in the genome have shown that MS-associated loci are substantially enriched for VDREs, including in the promoter region of *HLA-DRB1*,<sup>24,25</sup>

**Table 4** Results of overidentification tests using each 25(OH)D SNP as a separate IV

SNP used in IV	Odds ratio	95% CI	p Value
<b>KPNC</b>			
rs2282679	0.94	0.84-1.05	0.26
rs2060793	0.92	0.83-1.02	0.11
rs3829251	0.93	0.81-1.06	0.27
<b>EIMS/GEMS</b>			
rs2282679	0.99	0.93-1.06	0.80
rs10741657	0.92	0.87-0.97	0.002
rs3829251	0.84	0.79-0.90	<0.0001

Abbreviations: CI = confidence interval; EIMS = Epidemiological Investigation of Multiple Sclerosis; GEMS = Genes and Environment in Multiple Sclerosis; IV = instrumental variable; KPNC = Kaiser Permanente in Northern California; SNP = single nucleotide polymorphism.

Values adjusted for matching variables (and study status in Swedish population), principal component analysis/multidimensional scaling variables, presence of *HLA-DRB1\*15:01* alleles, a non-HLA multiple sclerosis genetic risk score, ever smoking, college education, and BMI at age 18-20 years or in 20s.

and that VDREs are more often exposed in open chromatin regions in immune cells compared with non-immune cells.<sup>26</sup> These findings reinforce the plausibility that vitamin D regulates genes that play important roles in the development or progression of MS. Furthermore, increased exposure to vitamin D leads to changes in immune cells that lead to decreased production of inflammatory cytokines, a decrease in Th1 and Th17 cell differentiation, and an increase in T regulatory cells, suggesting that low vitamin D is acting on MS by shifting the balance of the immune system toward a more proinflammatory state.<sup>27-29</sup>

The results of this study are in agreement with another recent MR study that used summary-level data only from the International Multiple Sclerosis Genetics Consortium and showed a causal effect for low 25(OH)D on MS risk.<sup>30</sup> There is some data overlap between previous and current studies; specifically, 2,812 EIMS/GEMS participants included here were also in the summary data analysis.<sup>30</sup> However, because individual-level data were not available for the prior study, the authors were unable to control for established genetic and environmental risk factors associated with MS. Importantly, the current study shows that measured variables did not contribute substantially to confounding of the causal relationship between low 25(OH)D and MS risk. Furthermore, the previous analysis relied on stronger assumptions<sup>30</sup> compared with a traditional MR analysis as presented in the current study. Finally, a *CYP24A1* SNP described in the previous report<sup>30</sup> was not included in our analysis, as it did not demonstrate association

**Table 3** Results of mendelian randomization analysis for causal effect of vitamin D in KPNC and EIMS/GEMS and the combined meta-analysis result

Data set	No. cases	No. controls	Causal odds ratio	95% CI	p Value
KPNC	1,056	9,015	0.79	0.64-0.99	0.04
EIMS/GEMS	6,335	5,762	0.86	0.76-0.98	0.03
Meta-analysis	7,391	14,777	0.85	0.76-0.94	0.003

Abbreviations: CI = confidence interval; EIMS = Epidemiological Investigation of Multiple Sclerosis; GEMS = Genes and Environment in Multiple Sclerosis; KPNC = Kaiser Permanente in Northern California.

Values adjusted for matching variables (and study status in Swedish population), principal component analysis/multidimensional scaling variables, presence of *HLA-DRB1\*15:01* alleles, a non-HLA multiple sclerosis genetic risk score, ever smoking, college education, and BMI at age 18-20 years or in 20s. Full expanded models are shown in tables e-1 and e-2.

with 25(OH)D at genome-wide significance,<sup>16</sup> which was among the criteria for SNPs to be included in our IV for MR analysis. It is possible that the association between a new IV that included the *CYP24A1* SNP and risk of MS might show a stronger magnitude of effect. The number of SNPs that capture variation in serum 25(OH)D levels is likely to grow as GWAS become larger.

An MR study relies on the following assumptions regarding the validity of the IV: (1) the genotype (IV) is associated with the phenotype (25(OH)D levels), (2) it is independent of measured or unmeasured confounders, and (3) it can only influence the outcome via the causal effect of the exposure.<sup>20</sup> We were able to meet most model assumptions by using a 25(OH)D IV established to be strongly associated with serum 25(OH)D in an independent population through a large GWAS and testing whether the 25(OH)D IV is independent of measured confounders. However, associations with unmeasured or unknown confounders cannot be ruled out, leaving one assumption not fully testable.

Common reasons for violations of MR assumptions are population stratification, LD (between the loci under study and other polymorphisms associated with the outcome), pleiotropy, and developmental canalization.<sup>2</sup> The potential for population stratification has been handled rigorously in the data analysis stage, and it is unlikely that it has caused confounding in our study. We have evaluated possible loci that might be associated with the 25(OH)D SNPs and with MS and have not found any basis for LD being the explanation for our findings. Pleiotropy concerns the possibility of polymorphisms having different biological effects that may influence the outcome through other pathways than the one studied. To our knowledge, the SNPs do not have other biological effects than in vitamin D metabolism, with the exception of rs3892951, upstream of *DHCR7*, which plays a role in cholesterol synthesis. Cholesterol metabolism has been associated with MS, although the exact mechanisms are not yet understood; a role for pleiotropy cannot be excluded.<sup>31</sup> On the other hand, the SNPs are near genes that have a clear, biologically plausible role in determining 25(OH)D levels. *GC* codes for the vitamin D-binding protein, which transports vitamin D to target tissues; *CYP2R1* converts vitamin D to its main circulating form, 25(OH)D; and *DHCR7* converts 7-dehydrocholesterol, the compound that is converted to vitamin D<sub>3</sub> in the skin in the presence of UV radiation, to cholesterol. In addition, a 2012 study evaluated SNPs in these 3 genes to look for evidence of pleiotropic effects and found that there were no associations with several biomarkers.<sup>32</sup> A study of 1,500 Danish MS patients found that SNPs in *GC* and *CYP2R1* have significant effects on 25

(OH)D levels, further indicating that variants in those 2 genes are affecting MS via serum vitamin D.<sup>33</sup> Canalization refers to the buffering of the effects of genetic variants against changing 25(OH)D levels; if this effect is present, the association between the IV and MS risk would be biased toward the null.

**Strengths and limitations.** Our study included non-Hispanic whites, which limits the generalizability of our findings. This limitation is particularly relevant in light of recent findings indicating that 25(OH)D levels may not be associated with MS in Hispanic populations.<sup>34</sup> We assume that the 3 alleles associated with 25(OH)D levels in the previous GWAS on Europeans are also associated with 25(OH)D levels in our populations. However, we were able to test this assumption in a subset of 2,077 EIMS participants for whom serum samples were collected at the time of inclusion in the study: a linear regression model adjusted for the same confounders as the main analyses showed that the IV was associated with 25(OH)D levels ( $p < 0.0001$ ) among MS cases and healthy controls. Other limitations include the assumption of linearity and the assumption that the 3 alleles used to construct the IV have additive effects on 25(OH)D levels. Furthermore, our analysis of severity was restricted to a cross-sectional measure, the MSSS, and models could not account for the use of disease-modifying therapies. Additional studies should aim to replicate findings, specifically in populations of other races/ethnicities, and better examine how specific 25(OH)D-related variants might influence MS susceptibility and progressive disease. The strengths of this MR study are that results are very unlikely to be due to reverse causation or confounded by unmeasured variables, that we were able to adjust for genetic ancestry, which could feasibly still confound the relationship between 25(OH)D SNPs and MS, and that similar results were found in 2 independent populations from 2 areas with different sunlight exposure levels.

#### AUTHOR CONTRIBUTIONS

Brooke Rhead and Maria Bäärnhielm: analysis and interpretation and manuscript drafting. Milena Gianfrancesco, Amanda Mok, and Ingrid Kockum: interpretation of data and revision of manuscript. Xiaorong Shao: analysis and interpretation of data. Hong Quach: acquisition of data. Ling Shen, Catherine Schaefer, Jenny Link, Alexandra Gyllenberg, Tomas Olsson, and Jan Hillert: acquisition of data and revision of manuscript. Anna Karin Hedström and M. Maria Glymour: revision of manuscript. Lars Alfredsson and Lisa F. Barcellos: acquisition of data, revision of manuscript, and study supervision.

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## DISCLOSURE

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