Evidence for a causal relationship between low vitamin D, high BMI, and pediatric-onset MS

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ABSTRACT

Objective: To utilize Mendelian randomization to estimate the causal association between low serum vitamin D concentrations, increased body mass index (BMI), and pediatric-onset multiple sclerosis (MS) using genetic risk scores (GRS).

Methods: We constructed an instrumental variable for vitamin D (vitD GRS) by computing a GRS for 3 genetic variants associated with levels of 25(OH)D in serum using the estimated effect of each risk variant. A BMI GRS was also created that incorporates the cumulative effect of 97 variants associated with BMI. Participants included non-Hispanic white individuals recruited from over 15 sites across the United States (n = 394 cases, 10,875 controls) and Sweden (n = 175 cases, 5,376 controls; total n = 16,820).

Results: Meta-analysis findings demonstrated that a vitD GRS associated with increasing levels of 25(OH)D in serum decreased the odds of pediatric-onset MS (odds ratio [OR] 0.72, 95% confidence interval [CI] 0.55, 0.94; p=0.02) after controlling for sex, genetic ancestry, *HLA-DRB1*15:01*, and over 100 non-human leukocyte antigen MS risk variants. A significant association between BMI GRS and pediatric disease onset was also demonstrated (OR 1.17, 95% CI 1.05, 1.30; p=0.01) after adjusting for covariates. Estimates for each GRS were unchanged when considered together in a multivariable model.

Conclusions: We provide evidence supporting independent and causal effects of decreased vitamin D levels and increased BMI on susceptibility to pediatric-onset MS. **Neurology® 2017;88:1623-1629**

GLOSSARY

BMI = body mass index; **chBMI** = childhood body mass index; **CI** = confidence interval; **CIS** = clinically isolated syndrome; **EIMS** = Epidemiologic Investigation of Risk Factors for MS; **GEMS** = Genes and Environment in MS; **GRS** = genetic risk score; **HLA** = human leukocyte antigen; **GWAS** = genome-wide association studies; **HGDP** = Human Genome Diversity Project; **KPNC** = Kaiser Permanente Northern California; **MDS** = multidimensional scaling; **MS** = multiple sclerosis; **OR** = odds ratio; **RPGEH** = Research Program on Genes, Environment, and Health; **SNP** = single nucleotide polymorphism; **UCSF** = University of California San Francisco; **vitD GRS** = genetic instrumental variable for vitamin D; **wGRS** = weighted genetic risk score.

Disease onset in multiple sclerosis (MS) typically occurs between the ages of 20 and 40 years; however, approximately 5% of patients experience symptoms before the age of 18.^{1–3} The exact mechanisms involved in pediatric-onset MS pathogenesis have yet to be defined; however, similar to adult MS, infection with Epstein-Barr virus, exposure to

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cigarette smoking, and the genetic risk factor *HLA-DRB1* are associated with pediatric-onset MS.^{1,4–6}

While there is strong research demonstrating the association between low vitamin D status and increased risk of adult-onset MS,^{7–9} there is a paucity of studies examining this relationship in pediatric MS. In addition, an association between childhood obesity and risk of pediatric MS has been reported,¹⁰ though a causal relationship remains to be confirmed. Further, it remains unclear whether the association between increased obesity and MS risk is mediated by low circulating levels of vitamin D, which can be stored in adipose tissue.

Mendelian randomization uses genetic variants strongly associated with an exposure as an instrumental variable, rather than a direct measure of the exposure, to estimate the effect of the exposure on an outcome. Inherited alleles are not affected by most potential confounding variables or disease status, thus reverse causation is unlikely. Our goal was to estimate the causal association of vitamin D and body mass index (BMI) on pediatric-onset MS risk using instrumental variable analysis based on genetic variants from established large genome-wide association studies (GWAS) in 2 large datasets of cases and controls.

METHODS US participants. Pediatric-onset MS cases were enrolled through pediatric MS centers established at University of California San Francisco (UCSF), Stony Brook, Children's Hospital of Philadelphia, Texas Children's Hospital, University of Colorado School of Medicine, University of Texas Southwestern, State University of New York Buffalo, Loma Linda University, Mayo Clinic, University of Alabama at Birmingham, Ann and Robert Lurie Children's Hospital of Chicago, University of Utah, Boston Children's Hospital, Massachusetts General Hospital, Washington University St. Louis, and Children's National Medical Center between January 2006 and December 2014. These centers are tertiary referral centers, but also serve local patients from all socioeconomic groups. Consecutive patients with onset of MS or clinically isolated syndrome (CIS) suggestive of early MS before age 18 years seen at some of these pediatric MS clinics were offered participation as previously described.¹¹ In addition, we utilized adult cases from Kaiser Permanente Northern California (KPNC) with reported age at onset <18 years. A separate cohort of distinct cases were enrolled in an NIHsupported case-control study (NS071463, PI Waubant) with MS or CIS onset before the age of 18, seen within 4 years of disease onset, with at least 2 silent T2-bright foci on brain and cord MRI, and ascertained by a panel of at least 2 pediatric MS experts.¹² Additional cases were provided from another MS

genetic study and only patients with a disease onset prior to age 18 based on medical records were included through UCSF. Established diagnostic criteria were used for all cases (total n=738).^{3,13}

Control individuals were derived from multiple sources to increase statistical power. These included (1) adult controls recruited from the KPNC membership without a diagnosis of MS or related condition (optic neuritis, transverse myelitis, or demyelination disease) confirmed through electronic medical records (n = 1,182)¹⁴; (2) adult controls from the Genetic Epidemiology Research on Adult Health and Aging cohort who participated in the KPNC Research Program on Genes, Environment, and Health (RPGEH) without evidence of MS in electronic medical records (dbGaP phs000674.v2.p2; N = 10,819); (3) pediatric controls (R01 NS071463, PI Waubant; n = 70); and (4) pediatric controls recruited as part of the Northern California Childhood Leukemia Study (n = 229).¹⁵

Whole blood was collected, processed, and extracted for DNA using Gentra Puregene protocol or received as Guthrie Card blood samples and extracted for DNA using the QIAamp DNA micro kit for dried blood spots (Qiagen, Venlo, Netherlands). Saliva was collected for DNA extraction using Oragene (Genotek, Murrieta, CA) kits. Genome-wide profiling was performed for all pediatric-onset MS cases and a subset of controls using the Illumina (San Diego, CA) Infinium platform. The Infinium 660K BeadChip or HumanOmniExpressExom BeadChip was used to genotype each study participant. RPGEH was genotyped using custom-designed Affymetrix (Santa Clara, CA) Axiom arrays. Genotype imputation was performed using IMPUTE2 and the 1,000 Genomes phase 3 reference haplotypes. Only single nucleotide polymorphisms (SNPs) with a high imputation quality score (info score ≥0.3) were retained for analysis. We eliminated SNPs that were associated with being genotyped on a particular array, were imputed in fewer than 99% of individuals, and deviated from Hardy-Weinberg equilibrium in controls (p < 0.000001).

Classical multidimensional scaling (MDS) was performed to visualize population substructure and provide quantitative measures of population genetic variation. MDS was completed for all genotyped data, which was merged with the Human Genome Diversity Project (HGDP) reference. Analysis of genome-wide average proportion of alleles shared identical by state was performed, and related/identical individuals were removed. Study samples were aligned with HGDP references, and the first 3 dimensions from MDS of the HGDP European population were used. We restricted our sample to white non-Hispanic participants (European ancestry), the largest group in our dataset, to ensure a genetically homogenous sample and avoid the possibility of confounding by genetic ancestry. After restriction and excluding population outliers (identified using MDS), the final dataset comprised 394 pediatric-onset MS cases and 10,875 controls.

Swedish participants. Data were collected from 2 population-based case-control studies of incident (Epidemiologic Investigation of Risk Factors for MS [EIMS]) and prevalent (Genes and Environment in MS [GEMS]) MS. The EIMS study (2005–2014) inclusion criteria were age 16–70 years, diagnosed MS according to the to the McDonald criteria^{13,16} within 2 years, and ability to understand the Swedish language. GEMS study participants were identified from the Swedish National MS registry, fulfilled the McDonald criteria, and were recruited during 2009–2011. We included cases with a reported age at onset <18 years. For both studies, controls were randomly chosen from the population register and matched to cases by sex, age at inclusion in the study, and region of residence. Two controls were matched to

each case in the EIMS study and one control per case in the GEMS study. All participants in the EIMS study were distinct from those in the GEMS study. Details of the study design have been described elsewhere. The participation rate in the EIMS study was 92% for cases and 67% for controls, and in the GEMS study 82% for cases and 66% for controls. Genotyping data were available for 75% of EIMS and 91% of GEMS participants. Data for 175 cases and 5,376 controls were available for the current study.

All participants were asked to give blood samples, which were genotyped on an Illumina custom array and on OmniExpress-24. HLA-DRB1 information was imputed with HLA*IMP02 using genotypes in the major histocompatibility complex region from the custom array. SNPs with <2% minor allele frequency, genotyped in <98% of individuals, or not in Hardy-Weinberg equilibrium among controls (p < 0.0001) were removed from analysis. Individuals with >2% failed genotype calls, related individuals, or population outliers identified using the SmartPCA program were removed. Ninety of the 110 MS risk SNPs were present on the custom array. Proxy SNPs for an additional 18 MS risk SNPs were selected from the custom array ($R^2 > 0.8$, except rs28723576, which had $R^2 > 0.71$). All vitamin D SNPs were present on the custom array. Twelve BMI SNPs were taken from the custom array, and the remaining from the OmniExpress chip. Forty-three BMI SNPs were not present on the array and were imputed using MaCH 1.0 with standard settings and the Northern European 1,000 Genomes reference panel. Seventeen markers utilized the August 2009 reference panel, 25 markers the August 2010 panel, and 1 marker the July 2011 panel. The variant rs2245368 could not be imputed with high quality and was therefore omitted from the BMI GRS.

Standard protocol approvals, registrations, and patient consents. Study protocols were approved by the institutional review boards for human subjects at UCSF, Stony Brook, Children's Hospital of Philadelphia, Texas Children's Hospital, University of Colorado School of Medicine, University of Texas Southwestern, State University of New York Buffalo, Loma Linda University, Mayo Clinic, University of Alabama at Birmingham, Ann and Robert Lurie Children's Hospital of Chicago, University of Utah, Boston Children's Hospital, Massachusetts General Hospital, Washington University St. Louis, Children's National Medical Center, Kaiser Permanente Division of Research, University of California Berkeley, Regional Ethical Review Board in North Stockholm, and Karolinska Institutet. Informed consent or assent (children) was obtained for all study participants and their parents when appropriate.

Exposure assessment. A previous GWAS identified 3 SNPs strongly associated with serum vitamin D level: rs2282679, rs2060793, and rs3829251, which together explain approximately 2.8% of the variance in circulating vitamin D levels. ¹⁸ A genetic instrumental variable for vitamin D (vitD GRS) was constructed using the 3 risk variants, weighting each allele by the effect size reported in the GWAS and summing across the variants. The BMI instrumental variable was derived using β coefficients as weights for 97 variants associated with BMI ($R^2 = 2.7\%$) identified through the largest and most recent GWAS. ¹⁹ The sum of risk alleles multiplied by the estimated effect reported of each risk allele on the phenotype was calculated as each individual's BMI genetic risk score (GRS).

We also explored whether variants specifically associated with childhood BMI (chBMI) were associated with pediatric-onset MS susceptibility. A chBMI GRS was constructed using 28 independent variants associated with pediatric/childhood BMI in the

literature. $^{20-24}$ The score was unweighted, and calculated by summing the number of risk alleles across each locus. Of the 28 variants in the chBMI GRS, 11 overlapped or were highly correlated with ($R^2 > 0.6$) variants in the BMI GRS.

A weighted GRS (wGRS) that combines the weighted odds ratio (OR) from each of 110 non-MHC MS susceptibility loci identified through recent GWAS²⁵ was calculated for each pediatric-onset case and control by multiplying the number of risk alleles for each locus by the weight for that variant and then taking the sum across the 110 loci.²⁶ The weight for each locus is the natural log of the OR for each allele. Three SNPs were missing in the US study (rs201202118, rs201847125, and imm_5_141486748), and 1 SNP failed Hardy-Weinberg equilibrium, rs2744148. Five SNPs were missing from the Swedish data and adequate proxies were not found (rs4772201, rs2028597, rs4679081, imm_5_141486748, and rs533646). Characterization of *HLA-DRB1*15:01* (0, 1, or 2 allele copies) for US participants was based on the rs3135388 tag SNP, which is highly correlated with *DRB1*15:01* alleles ($r^2 = 0.97$).²⁷

Statistical analyses. Logistic regression was used to estimate the effect of BMI and vitD GRS on pediatric-onset MS case/control status, controlling for sex, wGRS, presence of any *HLA-DRB1*15:01* alleles, and genetic ancestry. Ninety-five percent confidence intervals (CIs), OR, and p values were reported. Interaction was assessed on the multiplicative scale. Meta-analysis assuming fixed effects was performed if χ^2 tests of heterogeneity demonstrated p > 0.05. All analyses were conducted in PLINK, STATA, or R.

RESULTS Clinical and genetic characteristics of cases and controls are shown in table 1. Mean age at onset for pediatric cases was 14.05 years (± 3.30) in the US study and 14.91 (± 2.67) in the Swedish study. There was a significant difference in wGRS and *HLA-DRB1*15:01* status between cases and controls in both studies (p < 0.001). Average GRS for each instrumental variable in cases and controls are also reported in table 1. There was no association between age at onset and any of the instrumental variables in either study population (data not shown).

In the meta-analysis of both US and Swedish studies, vitD GRS was significantly associated with a decreased risk of MS (p = 0.02) after adjusting for sex, HLA-DRB1*15:01, wGRS, and genetic ancestry (table 2). The instrumental variable analysis for the BMI GRS also demonstrated a causal association between BMI and pediatric-onset MS as represented by the BMI GRS after adjusting for covariates (p = 0.01). There was no significant association between chBMI GRS and pediatric-onset MS. These results were consistent with results from analyses that also included pediatric-onset cases with age at onset 18 years (total n = 415 US cases and n = 262Swedish cases; table e-1 at Neurology.org), with meta-analysis estimates demonstrating a significant association between MS and BMI GRS (OR 1.15, 95% CI 1.04, 1.27; p = 0.01), vitD GRS (OR 0.75, 95% CI 0.59, 0.97; p = 0.03), but not chBMI GRS (OR 1.01, 95% CI 0.99, 1.04; p = 0.31).

Table 1 Clinical and genetic characteristics of patients with pediatric-onset multiple sclerosis and controls

	United States			Sweden		
	Pediatric-onset MS (n = 394)	Controls (n = 10,875)	p Value	Pediatric-onset MS (n = 175)	Controls (n = 5,376)	p Value
Age at onset, y	14.05 ± 3.30	_	_	14.91 ± 2.67	_	_
Female:male	3.0:1	1.6:1	0.02	2.5:1	3.2:1	0.15
wGRS	12.78 ± 0.67	12.31 ± 0.69	<0.001	12.41 ± 0.66	11.91 ± 0.69	< 0.001
HLA-DRB1*15:01			<0.001			< 0.001
0	193 (49)	7,973 (72)		57 (33)	3,806 (71)	
1-2	201 (51)	2,902 (28)		118 (67)	1,570 (29)	
vitD GRS	1.02 ± 0.28	1.04 ± 0.31	0.11	1.01 ± 0.32	1.06 ± 0.32	0.07
BMI GRS	11.51 ± 0.85	11.38 ± 0.81	0.002	11.54 ± 0.79	11.50 ± 0.82	0.49
chBMI GRS	25.45 ± 3.15	25.32 ± 3.26	0.44	26.06 ± 3.50	26.09 ± 3.32	0.93

Abbreviations: BMI = body mass index; chBMI GRS = childhood body mass index genetic risk score; vitD GRS = genetic instrumental variable for vitamin D; wGRS = weighted genetic risk score.

Table values are mean \pm SD for continuous variables and n (column %) for categorical variables; p value is for t test or χ^2 test between cases and controls.

When modeled together, both the vitD and BMI GRS independently contributed to pediatric-onset MS susceptibility. Joint estimates were consistent with those found when each GRS was modeled alone in both populations. For example, in the US study, estimates from the multivariate analysis reflected OR 1.21 (95% CI 1.06, 1.37) for the BMI GRS and OR 0.74 (95% CI 0.53, 1.03) for the vitD GRS.

There was no evidence of interaction between vitD GRS or chBMI GRS and *HLA-DRB1*15:01* (data not shown); however, a significant interaction was present between BMI GRS and *HLA-DRB1*15:01* in the US study (*p* interaction 0.04). Individuals carrying 1–2 *DRB1*15:01* risk alleles demonstrated a stronger association (OR 1.39) compared to noncarriers (OR 1.05). However, this interaction was not detected in the Swedish dataset (*p* interaction 0.66).

Table 2 Meta-analysis results support a causal association between both low vitamin D and high body mass index (BMI) and pediatric-onset multiple sclerosis

Instrumental variable ^a	Odds ratio ^b (95% CI)	p Value
Vitamin D GRS	0.72 (0.55-0.94)	0.02
BMI GRS	1.17 (1.05-1.30)	0.01
Childhood BMI GRS	1.02 (0.79-1.33)	0.88

Abbreviation: GRS = genetic risk score.

Analyses adjusted for sex, *HLA-DRB1*15:01*, weighted GRS, and genetic ancestry derived from principal components.

and independent association between low serum concentrations of vitamin D and increased BMI and risk of pediatric-onset MS after adjusting for sex, ancestry, *HLA-DRB1*15:01*, and over 100 non–human leukocyte antigen (HLA) MS risk variants. While previous studies have attributed an increased risk of MS associated with BMI to lower vitamin D levels seen in obese individuals, our findings support a role for independent contributions to disease susceptibility.

Several studies have shown a protective effect of vitamin D levels on MS risk. Two prospective studies showed a significantly reduced risk of MS in those with higher 25(OH)D.^{28,29} Maternal vitamin D deficiency (25[OH]D levels <12.02 ng/mL) during early pregnancy may be associated with a 2-fold risk of MS in offspring.30 Further, recent MR studies suggest a causal effect for low 25(OH)D on adult MS risk.8,9 Findings from our meta-analysis reveal that vitamin D represented as a GRS is significantly associated with pediatric-onset MS, with a stronger effect than in comparable adult-onset populations (OR 0.85).9 A larger sample size or instrumental variable constructed to include additional variants may help to demonstrate an even larger magnitude of effect, as each SNP used in the vitD GRS explained ≤1.2% of the variance in serum 25(OH)D concentrations.

Our findings also suggest that increased BMI is indeed a causal risk factor for pediatric-onset MS, similar to adults. Analogous to vitD GRS results, the BMI GRS demonstrates a stronger effect than that seen in adults (OR 1.10). Interestingly, previous studies have shown that specific HLA genotypes interact with self-reported BMI during adolescence to increase the risk of MS, an interaction that was also present in our study based on a BMI GRS; however,

^a GRS used as instrumental variables to capture low vitamin D and high BMI (see Methods).

^b Fixed effects meta-analysis; all χ^2 tests of heterogeneity p > 0.05.

we were unable to replicate this finding in the Swedish dataset, likely due to lower power resulting from a smaller sample size.

There are several hypotheses linking low vitamin D and high BMI to autoimmune diseases, including MS. Increased exposure to vitamin D leads 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 risk by shifting the balance of the immune response toward a more proinflammatory state.34-36 Obesity may also induce a Th17 response via an interleukin-6-dependent process leading to exacerbation of inflammatory diseases such as MS.37 In addition, a Th17/Treg imbalance may lead to alteration of intestinal microbiome in obese individuals, altering the intestinal immune response, which may explain the association between increased BMI and MS.38,39

Our study had several strengths, including clinically well-characterized pediatric-onset MS cases with average symptom onset 20 years earlier than most adult MS cases, rigorous diagnostic criteria, and focus on a single racial group. Comprehensive methods were used for quality control of genotypes and to identify and remove population outliers prior to analysis. We also analyzed BMI and vitamin D risk scores together in multivariate modeling, and used 2 study samples to conduct a large, well-powered meta-analysis. In addition, evaluation of our genetic risk scores in a subset of participants demonstrated a significant association with childhood body size (BMI GRS: Sweden n = 1,359, R^2 = 1.3%, p < 0.001; US n = 723, $R^2 = 0.7\%$, p = 0.03) and serum vitamin D levels adjusted for month of sample draw (vitD GRS: Sweden n = 881, R^2 = 2.5%, p < 0.001; US results previously demonstrated).40 Further research involving larger sample sizes is needed to examine whether serum vitamin D levels and childhood obesity independently and directly increase the risk, severity, and progression of pediatric-onset MS.

Limitations include identification of pediatriconset MS cases through tertiary clinics in the United States, which may not represent all MS cases with pediatric onset. While many of our cases had confirmed physician-based reports of pediatric MS (n = 130), a fraction self-reported the onset of their first disease symptoms in childhood. When the 2 patient groups were examined separately, similar effect sizes for each instrumental variable were observed (data not shown); therefore, self-report of pediatric symptoms was not likely to lead to misclassification. It is possible that individuals with a more benign disease or without access to health care may have been missed. In addition, instrumental variable analysis involves certain assumptions that we have

addressed. We met required model assumptions by utilizing a vitD and BMI GRS constructed from weights derived from independent populations through large published GWAS. However, associations of genetic variants with unmeasured or unknown confounders cannot be ruled out, and our models assumed linearity of the relationship between instrumental variable and outcome. Our study included non-Hispanic white participants and small numbers of male patients, which may limit the generalizability of our findings. Larger studies are needed to determine whether sex-specific effects contribute to pediatric-onset MS.

Our results add to the growing evidence supporting an important role of genetic and environmental factors in pediatric-onset MS. A causal relationship between low vitamin D and increased BMI has implications for prevention strategies in children. The effect of both low vitamin D and obesity may involve independent predisposing genetic factors and biological pathways mediating disease onset that future studies will unravel.

AUTHOR CONTRIBUTIONS

Dr. Gianfrancesco: analysis and interpretation of the data, drafting the article, final approval of the version to be published. Dr. Stridh: analysis and interpretation of the data, drafting the article, final approval of the version to be published. B. Rhead: analysis and interpretation of the data, reviewing and revising of the article, final approval of the version to be published. X. Shao: analysis and interpretation of the data, reviewing and revising of the article, final approval of the version to be published. E. Xu: analysis and interpretation of the data, reviewing and revising of the article, final approval of the version to be published. Dr. Graves: acquisition of the data, reviewing and revising of the article, final approval of the version to be published. Dr. Chitnis: acquisition of the data, reviewing and revising of the article, final approval of the version to be published. 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