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Extended haplotype association study in Crohn's disease identifies a novel, Ashkenazi Jewish-specific missense mutation in the *NF- κ B* pathway gene, *HEATR3*

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Conflict of interest

The authors declare no conflict of interest.

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Abstract

The Ashkenazi Jewish population has a several-fold higher prevalence of Crohn's disease compared to non-Jewish European ancestry populations and has a unique genetic history. Haplotype association is critical to Crohn's disease etiology in this population, most notably at *NOD2*, in which three causal, uncommon, and conditionally independent *NOD2* variants reside on a shared background haplotype. We present an analysis of extended haplotypes which showed significantly greater association to Crohn's disease in the Ashkenazi Jewish population compared to a non-Jewish population (145 haplotypes and no haplotypes with P-value < 10⁻³, respectively). Two haplotype regions, one each on chromosomes 16 and 21, conferred increased disease risk within established Crohn's disease loci. We performed exome sequencing of 55 Ashkenazi Jewish individuals and follow-up genotyping focused on variants in these two regions. We observed Ashkenazi Jewish-specific nominal association at R755C in *TRPM2* on chromosome 21. Within the chromosome 16 region, R642S of *HEATR3* and rs9922362 of *BRD7* showed genome-wide significance. Expression studies of *HEATR3* demonstrated a positive role in *NOD2*-mediated NF- κ B signaling. The *BRD7* signal showed conditional dependence with only the downstream rare Crohn's disease-causal variants in *NOD2*, but not with the background haplotype; this elaborates *NOD2* as a key illustration of synthetic association.

Keywords

haplotype association; Ashkenazi Jewish; Crohn's disease; *NF- κ B* signaling; synthetic association

Introduction

Crohn's disease (CD) is a complex genetic disorder characterized by chronic intestinal inflammation resulting from a dysregulated host immune response to intestinal microbiota.¹ The *NOD2* gene, involved in innate immune responses to bacterial peptidoglycan, is strongly associated with Crohn's disease and was initially identified through genetic linkage.² Uncommon, loss-of-function coding mutations (Arg702Trp, Gly908Arg, Leu1007fsinsC) in *NOD2* confer a 17.1 fold (95% CI: 10.7 – 27.2) increased risk for disease with homozygous or compound heterozygote risk allele carriage.³ The advent of genome-wide association studies (GWAS) has resulted in the identification of 140 loci with genome-wide significance in CD, implicating numerous immune mechanisms in disease pathogenesis. However, most identified loci involve variants of modest effects, and the presently identified 140 genetic loci account for only 13.6% of the estimated heritability.⁴

Because genotyping platforms utilized thus far have focused on common variants, it is possible that untested rare mutations may contribute significantly to complex disorders and account for some portion of missing heritability. Furthermore, precise models of disease pathogenesis integrating multiple disease associations are largely lacking, reflecting in part the significant pathophysiologic heterogeneity underlying the myriad associations reported

thus far in Crohn's disease. Approaches to reduce genetic complexity, such as through focused studies in selected populations, may be of benefit.

The Ashkenazi Jewish population has a several-fold higher prevalence of Crohn's disease compared to non-Jewish European ancestry cohorts, with estimates of increased prevalence ranging between 4.3 to 7.7-fold.^{5,6} The Ashkenazim have a unique genetic history, characterized by population bottlenecks, expansions and endogamy.⁷ However, the associated polymorphisms identified thus far through genome-wide association studies, as well as at *NOD2*, do not account for the higher disease prevalence in the Ashkenazim. Although the basis for the increased frequency of CD in the Ashkenazim is not known, it is possible that unidentified, uncommon variants, unique to or at a higher frequency within Ashkenazi Jews, contribute to higher disease prevalence. An additional possibility, given the greater linkage disequilibrium observed within the Ashkenazim, is that multiple functional polymorphisms inherited in cis, contribute to disease more commonly within Ashkenazi Jewish compared to non-Jewish European ancestry populations. Such a configuration of mutations would be consistent with the well-documented presence of longer haplotype blocks with greater levels of linkage disequilibrium, relative to non-Jewish populations.^{8,9} As such, association testing to identify significant extended regions may point the way to understanding the unique genetic architecture underlying Crohn's disease in Ashkenazi Jews.

Studying haplotype structure in the context of disease association also informs our conception of synthetic association, which has been hypothesized to be a major contributor to genome-wide association signals. Under this framework, molecular evolution, which creates varying amounts of haplotype diversity, allows for significant disease association near clusters of rare causal variants.¹⁰ For example, in CD, *NOD2* is distinguished by three uncommon functional mutations whose carriers are a subset of individuals who have a common background haplotype.^{3,11} The multiple hierarchical levels in the tree-based genealogy originally proposed by Dickson, et al., are reflected in various stringencies for the clustering parameters in an analysis of contemporary haplotype diversity; for analyses with strict parameters, causal variants may not lie within the associated haplotypes themselves. Using haplotypes may be a particularly powerful approach for tagging functional variants; a recent study found that using a multi-marker method for predicting Crohn's disease risk showed improved power over a single-variant approach.¹² Toward this end, we present an analysis using extended shared haplotypes as a filter for prioritizing novel variants and detecting conditional dependence structure.

Results

Extended haplotype analysis

To address the presence and relevance to disease of extended linkage disequilibrium stretches, we performed a haplotype-based analysis of a cohort comprising 397 Ashkenazi Crohn's disease cases and 431 controls, and 547 non-Ashkenazi Crohn's disease cases and 549 controls.^{13,14} Our analysis of associated extended haplotypes showed that there were markedly distinct patterns of haplotype association signals in the Ashkenazim compared to the non-Jewish cohort (Figure 1). We observed 145 haplotypes demonstrating nominal

evidence for association with P-values $< 10^{-3}$, including three distinct regions with P-values $< 10^{-4}$ (Supplementary Table 2). At this threshold, no regions were found to be significant in the non-Jewish cohort. Comparing the 145 haplotypes with 71 previously identified Crohn's disease loci in a GWAS-based meta-analysis,¹⁵ we found that there were four haplotypes overlapping three established loci (Table 1); no haplotypes overlapped the 69 additional CD-associated loci in a larger ImmunoChip-based study.⁴ Among all loci, the most significant haplotype association signal was found at chromosome 16q12 in a region that includes the *NOD2* gene. Three of these four haplotypes showed higher allele frequencies in Crohn's disease cases, corresponding to increased disease risk, while the remaining haplotype on chromosome 2 appeared to confer a protective function.

Chromosome 16 haplotypes—Two of the significant haplotypes in established CD loci (chr16_hap6721 and chr16_hap6830) were located contiguously on chromosome 16, spanning 48.88 – 49.18 and 49.18 – 49.48 Mb, respectively (Table 1). Haplotype chr16_hap6830 was carried by 25 Crohn's disease Ashkenazi cases and 4 Ashkenazi controls, and chr16_hap6721 was carried by 15 Crohn's disease cases and no controls. The 15 Crohn's disease cases that carried chr16_hap6721 were completely a subset of the 25 cases that carried chr16_hap6721. Importantly, of these 15 Crohn's disease extended haplotype carriers, 12 also carried the Gly908Arg polymorphism within the *NOD2* gene. Of the three major disease-associated polymorphisms, only the Gly908Arg variant is present at a significantly higher frequency in Ashkenazi compared to non-Ashkenazi Crohn's disease cases.^{16–22} *NOD2* encodes an intracellular pathogen recognition receptor that functions as a sensor for peptidoglycan found in the cell wall of most bacteria. In response to stimulation by a component of bacterial peptidoglycan, muramyl dipeptide (MDP), *NOD2* signaling leads to the activation of the *NF-κB* (nuclear factor of kappa light polypeptide gene enhancer in B-cells) family of transcription factors.^{23,24} In addition to the established association signal at *NOD2*, the associated region on chromosome 16 also contained the *BRD7* (bromodomain containing 7) gene, which has demonstrated genome-wide significant association at rs9922362 (P-value = 3.26×10^{-16}) in a GWAS dataset, separate from the one used in the haplotype analysis, comprising 907 Ashkenazi CD and 2345 Ashkenazi controls (Figure 2).²⁵ *BRD7* lies within 350Kb of *NOD2* and encodes a protein involved in chromatin remodeling.²⁶ Notably, the GWAS dataset did not show any evidence for independent association in *CYLD* (cylindromatosis), which is immediately adjacent to *NOD2*, as has previously been reported.²⁷

Chromosomes 2 and 21 haplotypes—Like the chromosome 16 region, the haplotype on chromosome 21, chr21_hap8015 (chr21: 44.47 – 44.65 Mb) also spans multiple immune-diseases related genes, such as *ICOSLG* (inducible T-cell co-stimulator ligand) and *AIRE* (autoimmune regulator). The chromosome 2 haplotype, chr2_hap10940 (chr2: 61.18 – 61.87 Mb), which had a higher frequency in controls compared to cases, was within 125Kb of *COMMD1* (copper metabolism [Murr1] domain containing 1), a regulator of the *NF-κB* signaling pathway.

Exome sequencing

To further explore the haplotype regions of interest, we performed exome sequencing on 55 Ashkenazi Jewish samples, including 50 Crohn's disease cases and 5 healthy controls. The samples' Jewish ancestry was validated genetically using GWAS data, and the case samples included two carriers each of the risk haplotypes on chromosomes 16 and 21. Within the associated haplotype regions chromosomes 16 and 21, we identified 40 previously unreported missense or nonsense polymorphisms among all 55 Ashkenazi Jewish individuals sequenced (Supplementary Table 3). None of the coding variants identified in the chromosome 2 haplotype were predicted by Polyphen-2 to be probably damaging in function, and no further investigation of these SNPs was conducted.²⁸

Chromosome 21 haplotype—We identified a number of previously unreported missense or nonsense uncommon mutations within the *TRPM2* (transient receptor potential cation channel, subfamily M, member 2) gene, including three polymorphisms, Arg755Cys, Gln953Stop and Thr1347Met, predicted by PolyPhen-2 to be probably damaging in function. Genotyping of these three SNPs was performed using Taqman assays on a follow-up Ashkenazi cohort of 1220 cases and 1167 healthy controls (Supplementary Table 1). Table 2 summarizes the association evidence for the three *TRPM2* variants. We observed nominal evidence for association (P-value = 0.0015) for the most common of these variants, Arg755Cys in the Ashkenazi Jewish Crohn's disease cohort. At this SNP, in a comparably-sized non-Jewish cohort (915 non-Jewish European ancestry Crohn's disease cases, and 818 matched healthy controls), we observed no evidence for association (P-value = 0.725). In contrast to Arg755Cys, we observed no evidence for association in the Jewish cohort for either Gln953stop or Thr1347Met in *TRPM2*. Of interest, the Gln953stop variant is carried by an individual carrying the chromosome 21 risk haplotype (Table 1) and is specific to the Ashkenazi Jewish population, not being observed in a subset of the non-Jewish European ancestry samples comprising 192 Crohn's disease cases and 192 healthy controls. Additional studies with larger cohorts will be required to definitively determine a role for *TRPM2* in Ashkenazi Jewish Crohn's disease.

Chromosome 16 haplotypes—Within the region encompassing the chromosome 16 risk haplotypes (Table 1), we identified 21 missense or nonsense mutations among the 55 individuals undergoing exome sequencing (Supplementary Table 3). While most of the carriers of the chromosome 16 risk haplotypes also carry the *NOD2* Gly908Arg variant, the converse is not true; in the initial GWAS cohort, only 15 of the 68 Gly908Arg carriers (22%) also carried the extended chromosome 16 risk haplotypes. We then asked whether carriers of the extended risk haplotype might carry additional risk alleles. Importantly, two extended chromosome 16 risk haplotype carriers also carried a previously unidentified missense mutation, Arg642Ser, in *HEATR3*. The *HEATR3* gene encodes a 680 amino acid protein, and the Arg642Ser polymorphism is highly conserved between species and predicted to be probably damaging to gene function by PolyPhen-2.

Fine mapping of chromosome 16 haplotypes

To more precisely define the nature of the various association signals in the *NOD2* haplotype, we performed single-point and logistic regression analyses on the region's key

markers in the 1167 Ashkenazi cases and 1220 Ashkenazi controls described above (Table 2). Consistent with prior reports, the Gly908Arg variant is the most common *NOD2* risk allele (10.4% minor allele frequency in cases) in the Ashkenazi Jewish cohort; this is in contrast to non-Jewish European ancestry Crohn's disease cohorts, where the frameshift mutation, Leu1007fsinsC, is the most common.^{19,29} The Arg642Ser variant in *HEATR3* (3.2% minor allele frequency in cases) is highly associated with Crohn's disease (P-value = 3.53×10^{-7}). We genotyped the Arg642Ser variant using Sanger sequencing in 384 non-Jewish European ancestry Crohn's disease cases and identified no carriers, indicating an Ashkenazi Jewish predominance for this novel variant. Because the *HEATR3* Arg642Ser variant is present in Ashkenazi Jews on an extended risk haplotype also containing the more common *NOD2* Gly908Arg allele (D' between Arg642Ser in *HEATR3* and Gly908Arg in *NOD2* = 0.828), we next performed conditional logistic regression analysis to test for additional conditionally dependent relationships. Several models were compared using the Akaike Information Criterion (AIC) to determine the best subset of SNPs. Additive, dominant and factor coding for the SNP genotypes were examined during the modeling stage. The final model with minimum AIC contained the associated marker in *BRD7*, rs9922362, as well as all four SNPs in *NOD2*, but not Arg642Ser in *HEATR3*. Under this regression model, Gly908Arg in *NOD2* demonstrated the most significant evidence for association (P-value = 1.66×10^{-13}). Additionally, in a full model containing all six variants across the three genes, a conditional relationship to *NOD2* accounted for much of the association evidence observed at *HEATR3* Arg642Ser variant (P-value = 0.333).

HEATR3 knockdown and over-expression

Although the genetic association signal at Arg642Ser in *HEATR3* on chromosome 16 did not appear to be independent, we noted that this gene is highly expressed in innate immune cells,³⁰ and we thus investigated its function in regulating *NOD2* signaling. To test this, we transiently knocked down *HEATR3* using siRNA in cells stably expressing *NOD2* and stimulated them with muramyl dipeptide in order to activate *NOD2* signaling. We monitored *NOD2*-dependent signaling using both *NF-κB* activation, as measured by a luciferase reporter assay (Promega, Madison, WI), and IL-8 secretion, as measured by ELISA assay. Cells transfected with siRNA targeting *HEATR3* showed no significant loss in cell viability but resulted in reduced MDP-induced *NF-κB* activation and greater than 50% reduction in IL-8 secretion compared to non-targeting siRNA controls. These results suggest that *HEATR3* is a positive regulator of *NOD2* signaling. To further establish this relationship, we also over-expressed *HEATR3* in HEK293 cells stable expressing *NOD2*. Cells over-expressing *HEATR3* exhibited increased luciferase activity and increased IL-8 secretion, thereby establishing a role for *HEATR3* in positively regulating *NOD2* signaling (Figure 3).

Discussion

The chromosome 16 associations at *BRD7* and *NOD2* represent an empirical paradigm of synthetic association that illustrates and extends the proposals of Dickson, et al.¹⁰ Using computer simulation, the researchers demonstrated that a cluster of low frequency, highly penetrant mutations often occurs stochastically more frequently with one allele than the other at a common SNP, resulting in "synthetic" association signals at these common

variants. While it has been argued that synthetic associations are not common in general, the interactions between and within *BRD7* and *NOD2* provide an important illustration of the complexities inherent in regression analyses to identify independent alleles. The minor allele (minor allele frequency 25.5% in healthy controls) at the common *NOD2* polymorphism rs2076756 tags a shared haplotype on which four independent, causal, and uncommon *NOD2* variants (Arg702Trp, Asn852Ser, Gly908Arg, Leu1007fsinsC) completely reside. This has been widely acknowledged as an example of synthetic association, as the *NOD2* GWAS signal is created by this cluster of rare variants with high effect, and the signal is mappable using linkage, which was originally predicted by Dickson et al.^{10,31}

In our analysis, common variants at *BRD7* (rs9922362) and *NOD2* (rs2076756) appeared to confer independent evidence for association, with P-values = 1.42×10^{-13} and 3.25×10^{-13} , respectively, in a logistic regression model consisting of the two SNPs. After conditioning the *BRD7* SNP on four uncommon causal *NOD2* variants, however, the *BRD7* association signal largely diminishes (conditional P-value = 0.0292). Taken together, these statistical results imply that the *BRD7* SNP functions as a partial background for the directly causal *NOD2* mutations, but not for the entire *NOD2* IBD-associated haplotype. This is visualized more directly in Figure 4, which shows that the *BRD7* SNP indeed does not appear to simply be associated through linkage disequilibrium with the *NOD2* background SNP, but rather that its carriers are enriched for the rare causal variants within the *NOD2* background haplotype. Importantly, the 0.4 Mb distance between the *BRD7* variant and *NOD2* is consistent with the findings by Dickson, et al. which showed that causal relationships can extend over long regions surrounding synthetic association, in the range of several megabases. However, we note that the absence of conditional dependence between the two common background SNPs in *BRD7* and *NOD2* conflicts with the hierarchical genealogy model supported by Dickson, et al. Overall, the example of Crohn's disease illustrates the variability of conditioning on highly associated, non-causal background variants such as the *NOD2* common haplotype background variant, rs2076756, as opposed to functional uncommon variants.

In this study, we report associations of uncommon extended haplotypes in Ashkenazi Jewish, but not non-Jewish European ancestry populations, thereby illustrating the greater power of this approach in endogamous populations. The haplotype associations on chromosome 16 tag the Gly908Arg variant within *NOD2*, the most prevalent causal mutation in Ashkenazi Jewish populations. The extended haplotype carriers in this region tag a less common, Ashkenazi Jewish-specific variant, Arg642Ser in *HEATR3*, which we demonstrated to function in the *NOD2* pathway. Given the extensive linkage disequilibrium between Gly908Arg in *NOD2* and Arg642Ser in *HEATR3*, however, we did not observe independent evidence for association for the *HEATR3* SNP. This marker, along with others in *TRPM2* on chromosome 21, demonstrated the capability of our haplotype-based analysis to direct the discovery of new disease-associated and population-specific polymorphisms. Furthermore, the extended length of the Jewish-specific haplotypes may indicate the presence of long-distance conditional relationships, as in the case of rs9922362 of *BRD7* and the *NOD2* causal variants on chromosome 16, an example of synthetic association. As additional fine-mapping studies of GWAS loci are reported, an increasing number of

population-specific polymorphisms and haplotypes will be identified. The etiology for the higher prevalence of Crohn's disease in the Ashkenazi Jewish population remains largely undefined; however, our study indicates that extended haplotypes containing multiple functional polymorphisms inherited in cis may contribute in a population-specific manner.

Materials and Methods

Subjects

Three hundred and ninety-seven Ashkenazi Crohn's disease cases and 431 controls, and 547 non-Ashkenazi Crohn's disease cases and 549 controls (Supplementary Table 1) were ascertained through Genetics Research Centers in Baltimore, Chicago, Montreal, Pittsburgh, Los Angeles, and Toronto, and through the New York Health Project. In all cases, informed consent was obtained following protocols approved by each local institutional review board. Diagnostic inclusion and exclusion criteria are described elsewhere.¹³ Ashkenazi Jewish ancestry was validated using principal components analysis.

Genotyping and haplotype association analysis

Genomic DNA was isolated from whole blood from the case-control cohort. Genotyping was performed on the Illumina HumanHap300 platform, with an average inter-marker distance less than 10Kb. All samples had genotype yields greater than 94%, and missing genotypes were imputed using BEAGLE 3.1.0 with default parameters.³² Analysis was performed using GERMLINE 1.5.0 (Genetic Error-tolerant Regional Matching with LINEar-time Extension), a computationally efficient program for identifying shared identical-by-descent segments between pairs of individuals in a large population, and DASH 1.1.0 (DASH Associates Shared Haplotypes), a program for clustering these segments into haplotype regions that are shared by multiple individuals.^{33,34} Our analysis utilized default parameters, with a minimum haplotype length of 3 Mb and a maximum of 4 homozygous mismatches between shared segments. In the Ashkenazi samples, 473,621 haplotype clusters across the autosomes were tested for association, compared to 8,986 clusters in the non-Jewish samples. Due to correlation between the haplotype clusters, precise multiple testing corrections could not be calculated, but given that all clusters in the Jewish and non-Jewish analyses had P-values greater than 10^{-5} and 10^{-3} , respectively, it was clear that none were genome-wide significant.

Exome sequencing and variant screening

Genomic DNA was extracted from whole blood and whole-exome captured with the NimbleGen 2.1 M human exome array following the manufacturer's protocol (Roche/NimbleGen, Madison, WI). Captured libraries were sequenced on the Illumina genome analyzer as paired-end 75-bp reads, following the manufacturer's protocol. Sequence reads were mapped to the reference genome (hg18) using the BWA program using default parameters.³⁵ The Genome Analysis Toolkit (GATK) v2 was used to call alleles at variant sites.³⁶ Sample-level realignment and multi-sample SNP calling were performed using default parameters and the GATKStandard variant filter.

HEATR3 knockdown and over-expression

HEK293 cells stably expressing *NOD2* were cultured in Dulbecco's modified Eagle's medium with 10% fetal bovine serum, 2 mM glutamine, 1 mM sodium pyruvate, and 1x penicillin-streptomycin (Gibco, Grand Island, NY). All experiments were performed in duplicate. In the silencing experiments, these cells were then reverse-transfected with siGENOME siRNA pools (Thermo Scientific, Hudson, NH) targeting *HEATR3* (or non-targeting siRNA as a negative control) at a concentration of 20 nM with the lipid transfection agent Lipofectamine 2000 (Invitrogen, Grand Island, NY) diluted in Opti-MEM (Invitrogen) for 48 hours. Cells were either left unstimulated or subsequently stimulated with MDP (20 ng/ml; Bachem, Torrance, CA) for 18 hours. In the over-expression experiment, HEK293 cells were transiently transfected using Lipofectamine 2000 with a pCMV-SPORT6 based expression plasmid encoding murine *HEATR3* (Thermo Scientific), followed by MDP stimulation as described above. Transfection with the empty vector was used as a negative control.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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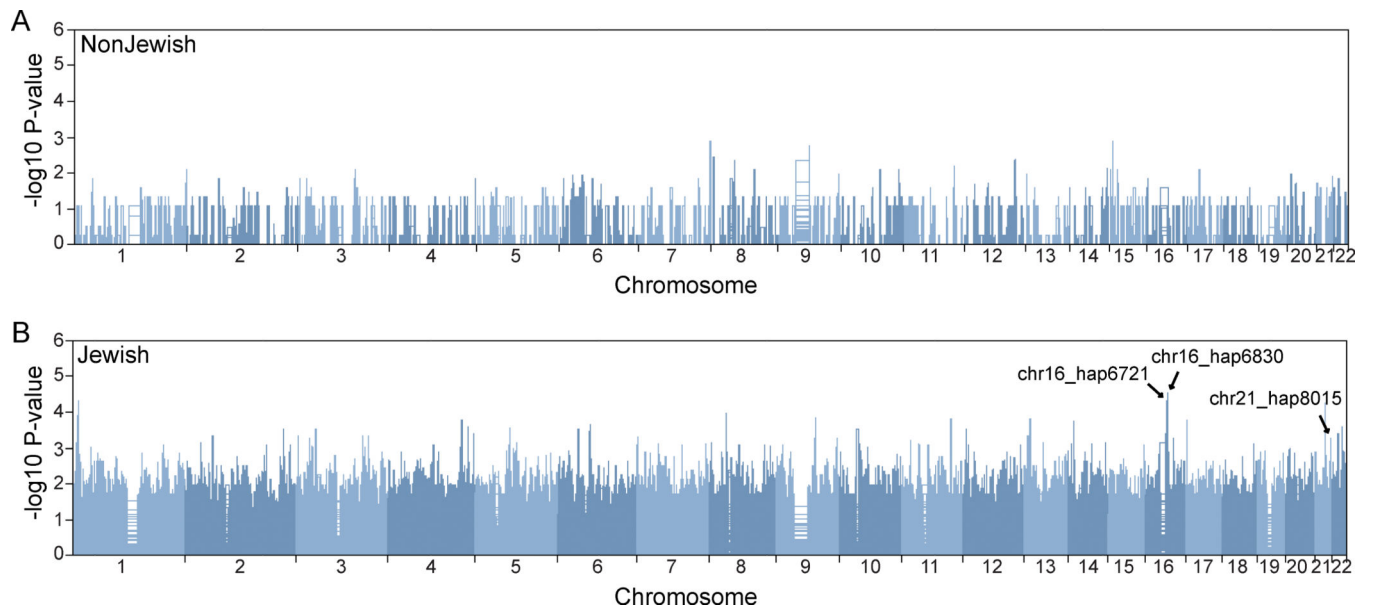


Figure 1. Manhattan plot of haplotype associations in non-Ashkenazi Jewish (A) and Ashkenazi Jewish (B) European Ancestry cohorts. In the non-Jewish analysis, no regions demonstrated evidence for association. In contrast, in the Ashkenazim, we observed 145 haplotypes demonstrating nominal evidence for association with P-values $< 10^{-3}$, including three distinct regions with P-values $< 10^{-4}$. Associated haplotypes that overlapped with established CD loci, and which are detailed in Table 1, are labeled.

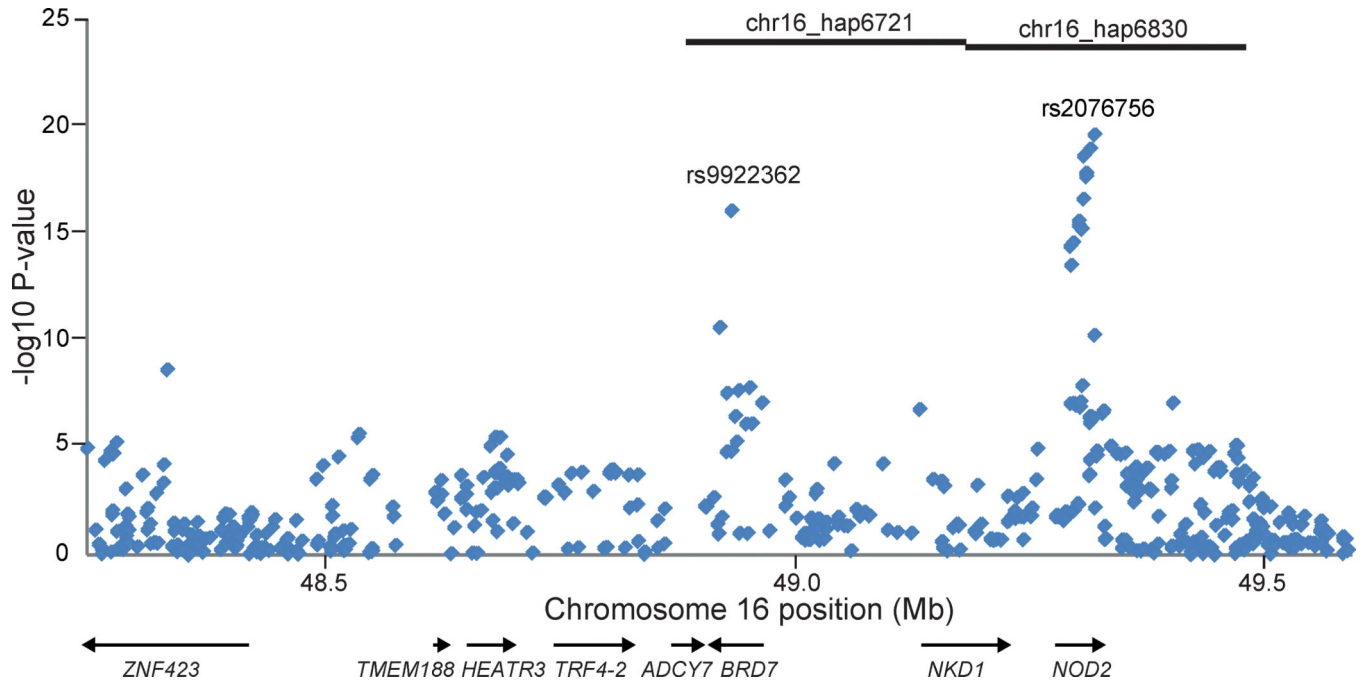


Figure 2.

Single-marker associations in the chromosome 16 haplotype region. We observed genome-wide significant associations at rs2076756 ($P\text{-value} = 2.32 \times 10^{-20}$) in *NOD2* and rs9922362 ($P\text{-value} = 3.26 \times 10^{-16}$) in *BRD7*. Positions of two nominally associated Ashkenazi Jewish haplotypes are shown by black bars at top.

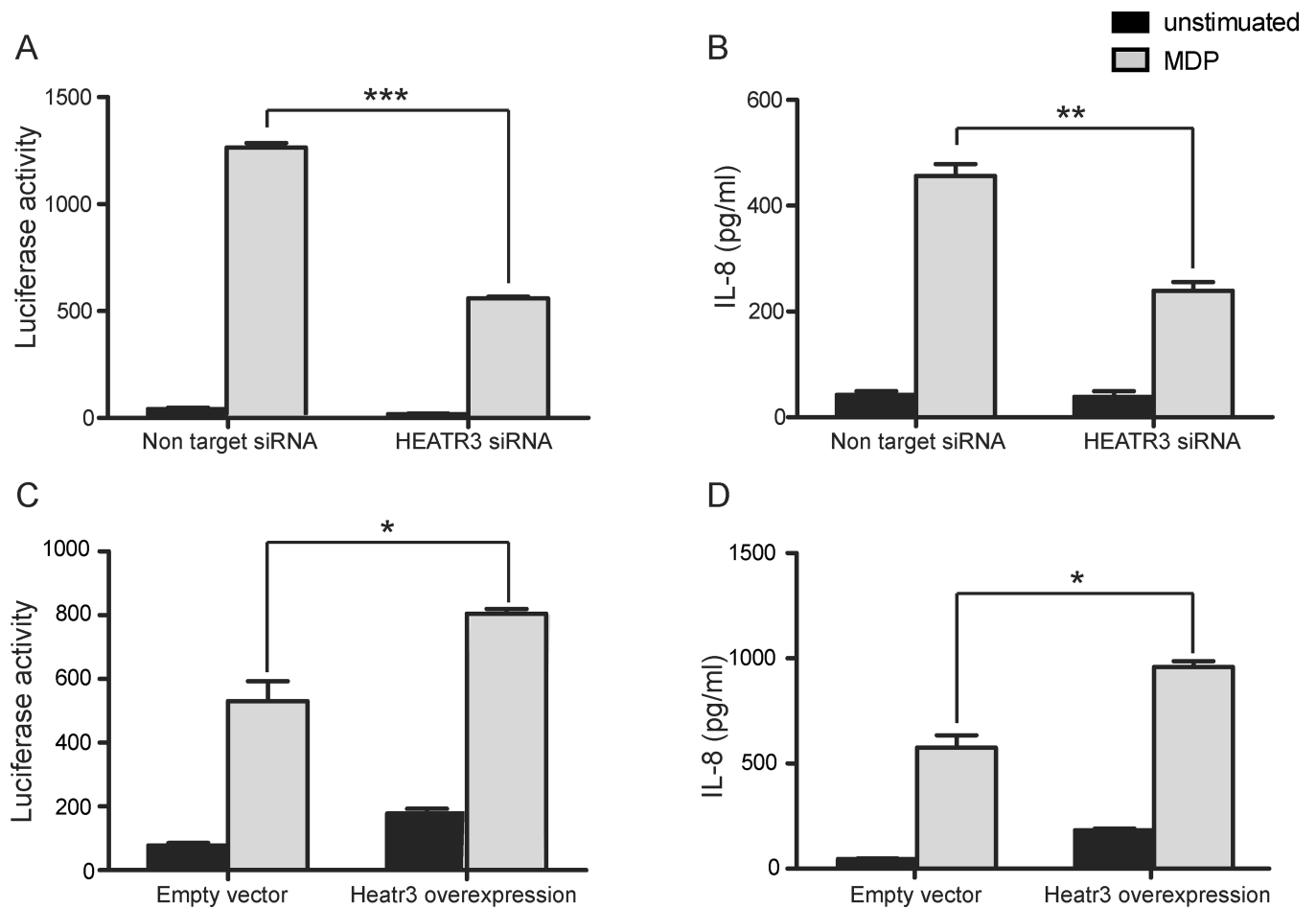


Figure 3. *HEATR3* is a positive component of the NOD2 signaling pathway. siRNA mediated knock down of *HEATR3* diminished NOD2-dependent NF- κ B induction (A) and IL-8 secretion (B) relative to non-targeting siRNA controls. Furthermore, over-expression of *HEATR3* increased NF- κ B signaling (C) and IL-8 production (D). Cells were left unstimulated (black bars) or stimulated overnight with 20 ng/mL of MDP (grey bars). Together these results support a positive role for *HEATR3* in NOD2 signaling. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

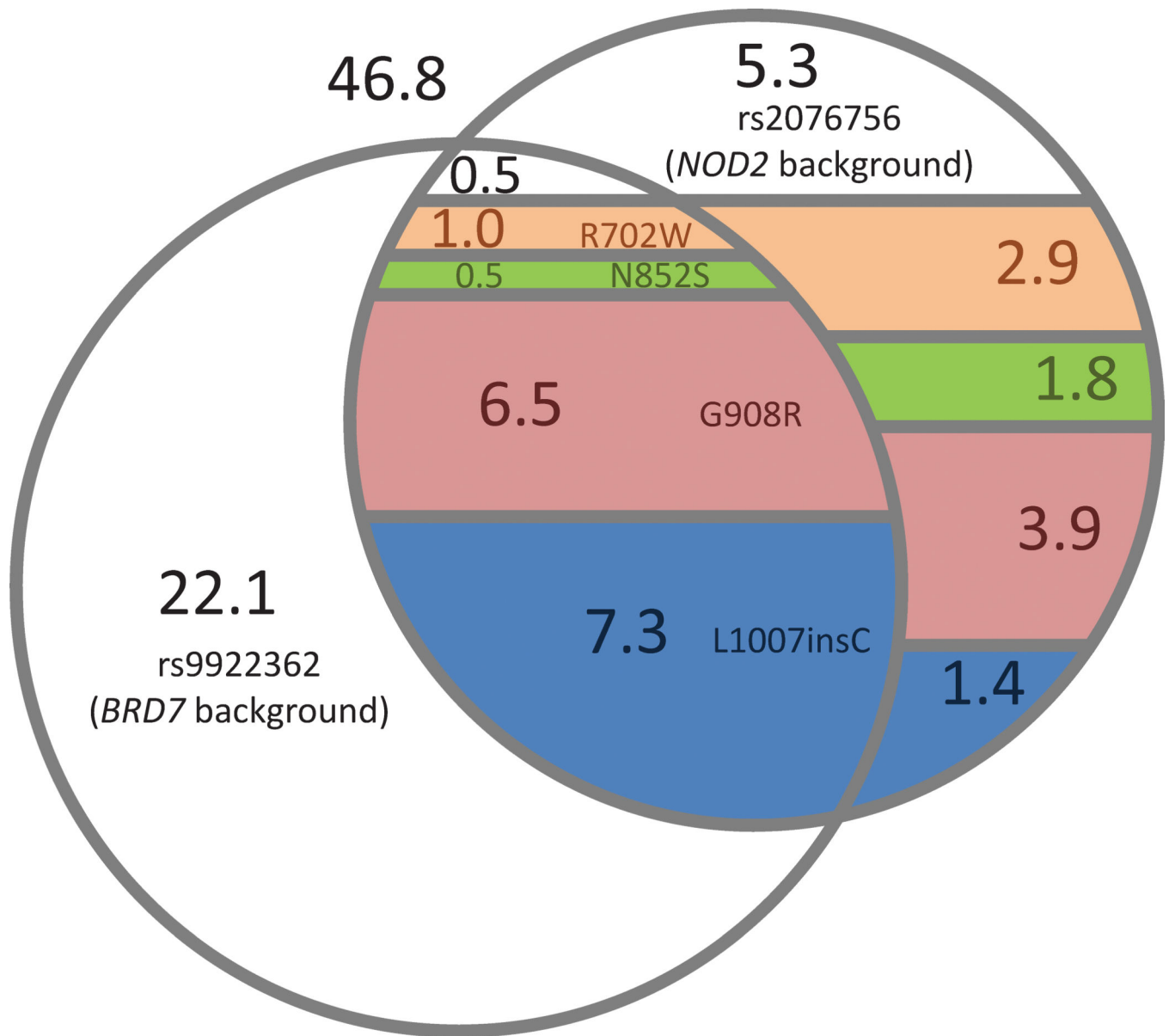


Figure 4.

Venn diagram of carriers for chromosome 16 haplotype variants among healthy controls. Numbers indicate the percentage of haploid chromosomes ($n = 2440$) that carry each combination of mutations. Colors indicate causal mutations within *NOD2*: peach = R702W; green = N852S; pink = G908R; blue = L1007insC. Missing genotypes and phasing of haploid carriers were inferred using BEAGLE with default parameters. Regions representing less than 0.5% of haploid carriers are not shown. These results indicate that both the common *BRD7* and *NOD2* SNPs serve as backgrounds for the *NOD2* uncommon causal variants, which in turn serve as the basis for association at the these two common background variants.

Associated haplotypes overlapping with established Crohn's disease loci in Ashkenazi Jewish samples

Table 1

Haplotype association							
Haplotype ID	Chr ¹	Start position ¹	End position ¹	P-value	MAF_Case ² (N =397)	MAF_Ctrl ² (N =431)	Established CD loci (Mb)
chr2_hap10940	2	61 184 852	61 865 369	4.63×10^{-04}	0.0038	0.024	60.76–61.87
chr16_hap6721	16	48 885 095	49 184 879	5.04×10^{-05}	0.019	0	49.02–49.41
chr16_hap6830	16	49 184 879	49 479 621	3.17×10^{-05}	0.032	0.0046	49.02–49.41
chr21_hap8015	21	44 471 849	44 657 340	5.26×10^{-04}	0.014	0	44.41–44.52

¹Chromosomal coordinates in hg 18.

²MAF = minor allele frequency.

Table 2

Single nucleotide polymorphism association and logistic regression analysis in Ashkenazi Jewish samples

SNP	Position ¹	Gene	Alleles	MAF_Case (N = 1167)	MAF_Ctrl (N = 1220)	P-value	P-value with model selection ²	P-value, full model ³
Chr16 (16q12.1)-HEATR 3, BRD7, and NOD2								
HEATR3_R642S	48 696 354	HEATR3	C/A	0.032	0.010	3.53×10^{-07}		0.33
BRD7_rs9922362	48 913 242	BRD7	G/A	0.39	0.32	2.36×10^{-07}	0.0292	0.041
NOD2_R702W	49 303 427	NOD2	C/T	0.040	0.022	5.00×10^{-04}	5.55×10^{-05}	5.27×10^{-05}
NOD2_N852S	49 311 345	NOD2	A/G	0.024	0.012	0.00140	0.0027	0.0028
NOD2_G908R	49 314 041	NOD2	G/C	0.10	0.040	1.93×10^{-17}	1.66×10^{-13}	3.57×10^{-10}
NOD2_L1007fsinsC	49 321 279	NOD2	G/C	0.087	0.028	1.65×10^{-18}	3.10×10^{-8}	3.90×10^{-08}
Chr21(21q22.3) - TRPM2								
TRPM2_R755C	44 644 624	TRPM2	C/T	0.043	0.026	0.0015		
TRPM2_Q953X	44 650 971	TRPM2	C/T	0.0080	0.0080	0.76		
TRPM2_T1347M	44 679 507	TRPM2	C/T	0.030	0.025	0.40		

¹ Chromosomal coordinates in hg 18.

² Logistic regression model with minimum AIC, comprising all four NOD2 variants and rs9922362 in BRD7.

³ Full logistic regression model comprising all six variants on chromosome 16.