Genome-wide association study of ulcerative colitis identifies three new susceptibility loci, including the HNF4A region

The UK IBD Genetics Consortium and The Wellcome Trust Case Control Consortium 2

Abstract

Ulcerative colitis (UC) is a common form of inflammatory bowel disease with a complex aetiology. As part of the Wellcome Trust Case Control Consortium 2, we performed a genome-wide association scan for UC in 2361 cases and 5417 controls. Loci showing evidence of association at P < 1 × 10^{-5} were followed up by genotyping in an independent set of 2321 cases and 4818 controls. We find genome-wide significant evidence of association at three new loci, each containing at least one biologically relevant candidate gene, on chromosomes 20q13 (HNF4A; P = 3.2 × 10^{-17}), 16q22 (CDH1 and CDH3; P = 2.8 × 10^{-8}) and 7q31 (LAMB1; 3.0 × 10^{-8}). Of note, CDH1 has recently been associated with susceptibility to colorectal cancer, which is an established complication of longstanding UC. The new associations suggest that changes in the integrity of the intestinal epithelial barrier may contribute to the pathogenesis of UC.
scans in UC reported associations at IL23R, IL10 and loci on chromosomes 1p36 and 12q15 which meet accepted genome-wide significance thresholds.\textsuperscript{14, 15}

As part of the Wellcome Trust Case Control Consortium 2 (WTCCC2) study of 15 complex disorders and traits, we report here the results of the largest GWA scan in UC to date. All study subjects were UK residents of white, European ancestry; clinical data are presented in Table 1. Cases and controls were genotyped on the Affymetrix 6.0 array. After application of quality control filters (see Methods), we analysed GWA data from 2361 individuals with UC and 5417 controls (Figure 1). An initial analysis revealed 24 distinct loci (comprising 156 SNPs) which showed evidence of association at $P < 1 \times 10^{-5}$. Sixteen of these had not been previously reported, and were followed up by genotyping the most strongly associated SNP from each locus using the Sequenom iPlex platform in an independent panel of 2321 UC cases and 4818 controls. Three new loci showed evidence for association at $P < 5 \times 10^{-8}$ in the combined panel, with three further new loci showing nominal ($P < 0.05$) replication (Table 2 and Figure 2). We describe these loci below and highlight the most plausible candidate gene for each, recognizing that fine mapping and functional studies are required to define causal variants and identify the gene from which each signal arises. A list of all loci for which replication was attempted is shown in Supplementary Table 1.

The most significant new association was seen at rs6017342 (GWA scan $P = 3.2 \times 10^{-13}$; combined GWA and replication $P = 8.5 \times 10^{-17}$), which maps within a recombination hotspot on chromosome 20q13 containing the 3′ untranslated region (UTR) of just one gene, HNF4A. The SNP rs6017342 itself maps 5kb distal to the 3′UTR. Although within an expressed sequence tag DB076868, this has been detected in just a single testis cDNA library and does not encode a significant open reading frame. The region contains two small blocks of sequence that are conserved in mammals and may include regulatory sequences affecting the expression of surrounding genes. Since rs6017342 is located within a recombination hotspot, there are few known SNPs in strong linkage disequilibrium ($r^2 > 0.5$) with it; there are none on the Affymetrix chip used in this study or on the Illumina chips used in previous studies. As the evidence for this association rests on this single SNP, we subjected these data to careful scrutiny; genotype cluster plots for this SNP showed clear resolution of the 3 genotype classes (Supplementary Figure 1), with 99.3% completeness of genotypes within this dataset.

Rare HNF4A mutations account for approximately 4% of UK cases of maturity-onset diabetes of the young (MODY),\textsuperscript{16} a monogenic form of diabetes mellitus characterized by autosomal dominant inheritance, young age of onset, pancreatic b-cell dysfunction and sensitivity to sulphonylureas. Common variants of HNF4A influence predisposition to Type II diabetes (rs2144908)\textsuperscript{17} and dyslipidaemia (rs1800961).\textsuperscript{18} The UC associated SNP, rs6017342 is not in LD with either of these 2 common variants, nor did it not show association in our study of CD ($P = 0.92$).\textsuperscript{3}

HNF4A encodes the transcription factor hepatocyte nuclear factor 4 α which regulates the expression of multiple components within all three key compartments of the cell-cell junction, namely the adherens junction, the tight junction and the desmosome.\textsuperscript{19} Such cell-cell junctions are fundamental to epithelial organization and barrier function. HNF4α also plays a key role in the development of the embryonic mammalian gastrointestinal tract. Previous studies demonstrated that mice with targeted deletion of HNF4α in epithelial cells of the foetal colon die perinatally. Histological analysis of colonic tissue recovered during late development (E18.5) demonstrated absent crypt formation, reduced epithelial cell proliferation and defective goblet cell maturation.\textsuperscript{20} In order to explore the role of HNF4α in murine intestinal inflammation, Ahn and colleagues circumnavigated the embryonic lethality of Hnf4α\textsuperscript{-/-} mice by generating a conditional model of intestinal Hnf4α deletion.\textsuperscript{21} These Hnf4α\textsuperscript{IEPC} mice (floxed Hnf4α driven by the villin promoter) developed increased epithelial permeability and
a markedly more severe colitis following dextran sodium sulphate (DSS) challenge, than their wild-type littermates.21 The same investigators provided preliminary evidence for dysregulated HNF4A gene expression in the intestinal epithelium in Crohn’s disease and in ulcerative colitis,21 a finding which now merits detailed re-exploration.

Significant association was also seen for a locus on chromosome 1q22, with the strongest signal at rs1728785 (GWA scan \(P = 1.8 \times 10^{-5}\); combined GWA and replication \(P = 2.8 \times 10^{-8}\)). The interval bounded by recombination hotspots spans 411 kb and encodes several genes. Among the strongest candidates for UC susceptibility is CDH1 which encodes E-cadherin. This transmembrane glycoprotein is one of the main components of the adherens junction and a key mediator of intercellular adhesion in the intestinal epithelium. It also plays a key role in epithelial restitution and repair following mucosal damage and expression of CDH1 is known to be significantly reduced in areas of active UC.22

Given the well-recognised association between UC and colorectal cancer,2 the observation of correlated association signals at the CDH1 locus in both diseases is striking. Thus variants in LD \((r^2 = 0.5)\) with the most strongly UC associated SNP in our study were recently identified in a GWA scan meta-analysis to be associated with colorectal cancer susceptibility23; conversely, we find that a perfect proxy for the most associated SNP in the colorectal cancer study is also associated with UC \((P = 8 \times 10^{-8})\). This locus did not show association with CD in a large international GWA meta-analysis of CD \((P = 0.549)\) (Supplementary Table 2). However, evidence for association of CDH1 with CD was reported recently in the Canadian population using a candidate gene approach,24 and the CD associated SNPs resulted in a truncated E-cadherin protein in vitro which accumulated in the cytoplasm and led to disorganized epithelial architecture.24

Of great potential relevance is the evidence that HNF4A and E-cadherin co-operate to maintain epithelial barrier integrity in the intestine. In experiments focused on the liver, HNF4a knockout mice failed to express E-cadherin,19 while in the gut E-cadherin dependent cell-cell contact was found to be critical in determining the amount and binding activity of nuclear HNF4a. This in turn affected the expression of several genes including ApoA-IV,25 an anti-inflammatory protein known to inhibit experimental colitis.26

The third newly confirmed UC susceptibility locus was a region on chromosome 7q31, previously suggested by a recent North American GWA scan.14 In the current study the peak association was seen at rs886774 (GWA scan \(P = 4.8 \times 10^{-7}\); combined GWA and replication \(P = 3.0 \times 10^{-8}\)). A strong positional candidate gene at this locus is LAMB1, encoding the laminin beta 1 subunit. Laminins are heterotrimers; the beta-1 light chain is present in laminins-1 -2 and -10. Laminins are expressed in the intestinal basement membrane, and play a key role in anchoring the single-layered epithelium; expression is known to be down-regulated in UC.27 rs886774 was not associated with CD in the meta-analysis.3 (Supplementary Table 2)

Two other loci previously implicated in UC-related phenotypes showed strong (but not genome-wide significant) association with UC. These comprise a SNP previously associated with osteoporosis28 (rs7524102 on chromosome 1p36, combined GWA and replication \(P = 3.1 \times 10^{-7}\)) and a SNP nearby (though not in LD with) a marker known to be associated with psoriasis29 (rs9548988 on 13q.13, combined GWA and replication \(P = 2.7 \times 10^{-7}\)).

In addition to the novel loci described above, our GWAS detected strong association at established UC loci such as the MHC, IL23R, 3p21/MST1 and NKX2-3 (one tailed \(P\) values in the direction of the previously reported association in Table 3). We also provide robust confirmation of two UC loci reported recently in genome-wide scans, the IL10 locus11 and the OTUD3/PLA2G2E locus12 on chromosome 1q31 and 1p36 respectively. Also of interest is our
finding that the PSMG1 locus on chromosome 21, which has previously been associated with pediatric-onset IBD, is likely to contribute specifically to disease susceptibility in UC. Variable degrees of support were obtained for some previously reported UC loci, including ECM1, CARD9, KIF21B/chromosome 1q32, and JAK2/chromosome 9p24, but weaker support for other loci such as IL2/IL21, IL12B and 12q15 (Table 3). Some of the UC loci are clearly associated with CD, while others are not, or have not been tested (Supplementary Table 2). We also tested for epistatic interaction among all pairwise combinations of these loci (both previously described and new) but found none.

This is the first report of a new series of GWA scans undertaken by the WTCCC2 consortium. We have identified three new susceptibility loci for UC, and provide the first genetic link between UC and colorectal cancer. Each of the strongest new association intervals that we have identified contains respectively HNF4A, CDH1 and LAMB1 as the most plausible positional candidate genes, thus providing further evidence for the re-emerging concept that altered epithelial barrier function may be a key factor in UC pathogenesis. Indeed, this is the first time that variants within genetic loci encoding such epithelial barrier genes have shown association with IBD at stringent genome-wide significant thresholds. Fine mapping and functional studies are clearly required to investigate this connection further, but our study provides strong scientific justification for the exploration of new therapeutic targets relevant to epithelial barrier function.

METHODS

Subjects

Cases—A total of 5319 unrelated patients of white, European, non-Jewish ancestry with a diagnosis of ulcerative colitis established using standard endoscopic, radiological and histological criteria, were recruited from ten centres within the United Kingdom (Cambridge, Oxford, London, Newcastle, Sheffield, Edinburgh, Dundee, Manchester, Torbay and Exeter, Supplementary Table 4). All patients provided written consent and either a sample of blood or saliva, from which DNA was extracted according to standard protocols. Research Ethics Committee approval was obtained prior to sample collection (Cambridge, Oxford, London, Newcastle, Sheffield, Edinburgh, Dundee, Manchester, Torbay and Exeter Local Research Ethics Committees). After QC (see below), we analyzed a total of 4682 samples, which were divided between the discovery panel (2361 samples) and replication panel (2321 samples).

Controls—A total of 10,235 control DNA samples from 3 sources passed our QC filters (see below). 5417 samples of the WTCCC2 common control set were used for the GWA experiment. This comprised 2675 healthy blood donors recruited from the United Kingdom Blood Service (UKBS), and 2742 samples from the 1958 Birth Cohort (1958BC) obtained from EBV-transformed cell lines from individuals born in England, Wales and Scotland during one week in 1958. The 4818 samples used as controls for the replication cohort were recruited from the Wellcome Trust-funded People of the British Isles (PoBI) DNA collection, obtained from rural populations throughout the British Isles, and from a further independent set of DNA samples obtained from 1958BC. All of the control samples used were from individuals with self-reported Caucasian ethnicity.

DNA sample preparation: Genomic DNA for all cases was shipped to the Sanger Institute, Cambridge. DNA quality plus subject identity were validated using the Sequenom iPLEX assay designed to genotype 4 gender SNPs and 26 SNPs present on the Affymetrix array. DNA concentrations were quantified using a PicoGreen assay (Invitrogen) and an aliquot assayed by agarose gel electrophoresis. A DNA sample was considered to pass quality control if the
original DNA concentration was ≥50 ng/µl, the DNA was not degraded, the gender assignment from the iPLEX assay matched that provided in the patient data manifest and genotypes were obtained for over 65% of the SNPs on the iPLEX.

**GWA Genotyping**

Samples were genotyped at Affymetrix’s service laboratory on the Genome-Wide Human SNP Array 6.0. For all samples passing Affymetrix’s laboratory QC, raw intensities (from the .CEL files) were renormalized within collections using CelQuantileNorm (see http://outmodedbonsai.sourceforge.net/). These normalized intensities were used to call genotypes with an updated version of the Chiamo software (see www.stats.ox.ac.uk/~marchini/software/gwas/chiamo.html), adapted for Affymetrix 6.0 SNP data. The Chiamo algorithm simultaneously calls genotypes for individuals in several collections; here it was applied to 15,068 individuals from five collections genotyped as part of the WTCCC2. Chiamo generates posterior probabilities for each of the three possible genotypes plus a fourth class of outliers. Our analyses use thresholded genotypes: for each individual, if one genotype had posterior probability greater than 0.9, this was set as the genotype for that individual, otherwise the genotype was set to be missing. After applying the QC filters described below, this threshold led to a study-wide level of missing data of 0.20%.

An overlapping set of 4830 controls were also genotyped on the Illumina 1.2M chip as part of a separate WTCCC2 project, and the 50,000 SNPs which are shared between that platform and the Affymetrix 6.0 (used in this study) were used to evaluate genotype accuracy. For the same QC thresholds and similar levels of missing data, discordance between Chiamo and Illuminus, which we regard as an upper bound on genotyping error rate, was 0.05857% for 1958BC and 0.07476% for UKBS.

We compared Chiamo to Birdsuite (the default Affymetrix calling algorithm applied on a plate-by-plate basis as recommended in 33) by making genotype calls at different confidence thresholds, and then plotting the fraction of calls made against concordance with the Illumina genotypes (Supplementary Figure 2). The general trend is that, when matched for the proportion of missing data, Chiamo has slightly higher concordance than Birdsuite. We are therefore confident that Chiamo is an acceptable alternative to Birdsuite.

**Replication Genotyping**

In the replication stage, genotyping was carried out at the Sanger Institute using the Sequenom iPLEX Gold assay. For one locus, the most associated SNP could not be genotyped with this technology, so a perfect (r² = 1 in all HapMap populations) proxy was used instead. 19 SNPs (including 3 gender markers) were typed in a multiplex reaction; 15 passed experimental QC (one SNP with Hardy-Weinberg P value < 1 × 10⁻⁶ was discarded). Samples with > 20% missing genotypes (n=300) were excluded; these samples are not included in the tallies in Table 1.

**Quality Control**

**Samples**—As is now standard practice for GWAS studies, we excluded sets of individuals whose genome-wide patterns of diversity are outliers compared to the bulk of those in the study, and SNPs where there is evidence that genotype calls do not provide precise estimates of genotype frequencies. Ignoring individuals and SNPs in this way throws away data gained at some expense, but because they typically violate assumptions underpinning standard tests for association, the payback in terms of increased accuracy for these tests can be substantial.

In order to try to obtain the maximally powerful set of samples and SNPs we attempted to refine some standard QC practices. For all individuals we explicitly model the data as a mixture
of “normal” and “outlier” individuals for each of ancestry, missing data and heterozygosity, and sex assignment. We fit each model in a Bayesian framework, and exclude individuals whose posterior probability of belonging to the outlier class was above 0.5. This approach replaces (and we believe improves upon) the traditional concept of fixed exclusion thresholds for parameters such as call rate, heterozygosity and ancestry. In total 413 case individuals and 567 control individuals were excluded from the analyses (Supplementary Table 3).

To assess relatedness amongst study individuals we compared each individual with the 100 individuals they were most closely related to (on the basis of genome-wide levels of allele sharing) and used a hidden Markov Model (HMM) to decide, at each position in their genome, whether the two individuals shared 0, 1, or 2 chromosomes identical by descent. This allows more refined assessment of the relatedness between individuals than do genome-wide sharing statistics (for example, parent-child relationships can be distinguished from siblings). We obtained a set of individuals with IBD < 5% by iteratively removing the member of each pair of putatively related individuals with more missing genotypes.

SNPs—For each SNP we considered a measure of the (Fisher) information carried by the genotype calls for the underlying allele frequency. Informally, this will decrease as the number of individuals with low posterior probabilities for the most likely call increases, and it can be thought of as a more refined measure of both missing data levels and minor allele frequency (Supplementary Figure 3). The measure is calculated automatically by the program SNPtest (www.stats.ox.ac.uk/~marchini/software/gwas/snptest.html). SNPs were removed if this information measure was below 0.98, or if the estimated MAF was below 0.01% (both calculated on the combined case-control data). 14.7% of SNPs were removed by these criteria. Again, this approach appears to offer advantages over conventional SNP filters, in excluding fewer SNPs for the same level of improved data quality. Because associated SNPs are expected to be enriched in the tiny fraction of poorly performing markers on these chips, we subsequently examined 155 cluster plots for SNPs with p < 1×10^{-5}, and excluded 16 from further analysis as likely genotyping errors.

Supplementary Figure 4 provides QQ plots for the post-QC comparison of our two control collections, and for association statistics based on the post-QC trend test comparing cases and the combined control set. Both visual inspection, and the inflation statistic for each (\(\lambda = 1.037\) and \(\lambda = 1.079\) respectively), suggest that the QC filtered data provides a good basis for association analyses.

**Statistical Methods**

We report p-values from 1-d.f. Cochran-Armitage tests for trend as implemented in the software SNPTTEST and PLINK. We also performed 2-d.f. genotypic tests to verify that none of our associations show significant deviation from a multiplicative model, and two marker logistic regressions to test for epistasis between associated markers. Effect size estimates are based on replication samples only, and represent per-allele increase of risk in a multiplicative model.

**URLs**

Supplementary Material
Refer to Web version on PubMed Central for supplementary material.

Complete List of Authors
†The complete list of authors who contributed to this study is as follows:

: The UK IBD Genetics Consortium

Jeffrey C Barrett†1, James Lee2, Charlie Lees3, Natalie Prescott3, Carl A Anderson1,5, Anne Phillips3, Emma Wesley6, Kirstie Parnell6, Hu Zhang2, Hazel Drummond3, Elaine R Nimmo3, Duncan Massey2, Kasia Blaszczyk4, Timothy Elliott7, Lynn Cotterill8, Helen Dallal9, Alan Lobo10, Craig Mowat11, Jeremy Sanderson7, Derek P Jewell12, William Newman8, Cathryn Edwards13, Tariq Ahmad9, John C Mansfield14, Jack Satsangi3, Miles Parkes2, Christopher G Mathew4

The Wellcome Trust Case Control Consortium 2

Management Committee

Peter Donnelly (Chair)1,2, Leena Peltonen (Deputy Chair)3, Elvira Bramon4, Matthew Brown5, Juan Casas6, Aiden Corvin7 Nicholas Craddock8, Panos Deloukas3, Janus Jankowski9, Hugh Markus10, Christopher G Mathew11, Mark McCarthy12, Colin Palmer13,

---

1Wellcome Trust Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, UK
2Gastroenterology Research Unit, Addenbrooke's Hospital, Hills Road, Cambridge CB2 2QZ, UK
3Gastrointestinal Unit, Molecular Medicine Centre, University of Edinburgh, Western General Hospital, Crewe Road, Edinburgh EH4 2XU
4Department of Medical and Molecular Genetics, King's College London School of Medicine, Floor 8 Tower Wing, Guy's Hospital, London SE1 9RT, UK
5Wellcome Trust Centre for Human Genetics, Roosevelt Drive, Oxford OX3 7BN, UK
6Peninsula College of Medicine and Dentistry, Barrack Road, Exeter EX2 5DW, UK
7Dept Gastroenterology, Guy's & St Thomas' NHS Foundation Trust, St Thomas' Hospital, London SE1 9RT, UK
8Department of Medical Genetics, Manchester Academic Health Science Centre (MAHSC), University of Manchester and NIHR Biomedical Research Centre, Central Manchester NHS Foundation Trust, Manchester M13 0JH, UK
9Department of Gastroenterology, James Cook University Hospital, South Tees Hospitals NHS Trust, Marton Road, Middlesbrough TS4 3BW, UK
10Division of Molecular and Genetic Medicine, University of Sheffield Medical School, Royal Hallamshire Hospital, Sheffield S10 2JF, UK
11Department of General Internal Medicine, Ninewells Hospital and Medical School, Ninewells Avenue, Dundee DD1 9SY, UK
12Gastroenterology Unit, Gibson Laboratories, Radcliffe Infirmary, Woodstock Road, Oxford OX2 6HE, UK
13Endoscopy Regional Training Unit, Torbay Hospital, Torbay TQ2 7AA, UK
14Institute of Human Genetics, Newcastle University, Newcastle upon Tyne NE1 3BZ, UK
1Wellcome Trust Centre for Human Genetics, Roosevelt Drive, Oxford OX3 7LI, UK
2Dept Statistics, University of Oxford, Oxford OX1 3TG, UK
3Wellcome Trust Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, UK
4Dept Psychological Medicine, King's College London Institute of Psychiatry Denmark Hill, London SE5 8AF, UK
5Nuffield Department of Orthopaedics, Rheumatology and Musculoskeletal Sciences, University of Oxford, Nuffield Orthopaedic Centre, Oxford OX3 7LD, UK
6Dept Epidemiology and Population Health, London School of Hygiene and Tropical Medicine, London WC1E 7HT, UK
7Neuropsychiatric Genetics Research Group, Institute of Molecular Medicine, Trinity College Dublin, Dublin 2, Eire
8Dept Psychological Medicine, Cardiff University School of Medicine, Heath Park, Cardiff CF14 4XN, UK
9Centre for Gastroenterology, Bart's and The London School of Medicine and Dentistry, London E1 2AT, UK
10Division of Cardiac and Vascular Sciences, Dept Clinical Neurosciences, St George's Hospital, London SW17 0RE, UK
11Dept Medical and Molecular Genetics, King's College London School of Medicine, Guy's Hospital, London SE1 9RT, UK
12Oxford Centre for Diabetes, Endocrinology and Metabolism (ICDEM), Churchill Hospital, Oxford OX3 7LJ, UK
13Biomedical Research Centre, Ninewells Hospital and Medical School, Dundee DD1 9SY, UK

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14Social, Genetic and Developmental Psychiatry Centre, King’s College London Institute of Psychiatry, Denmark Hill, London SE5 8AF, UK
15Dept Clinical Neurosciences, University of Cambridge, Addenbrooke’s Hospital, Cambridge CB2 2QQ, UK
16Glaucoma Research Unit, Moorfields Eye Hospital NHS Foundation Trust, London EC1V 2PD, UK
17Dept Molecular Neuroscience, Institute of Neurology, Queen Square, London WC1N 3BG, UK
18Genetics and Infection Laboratory, Cambridge Institute of Medical Research, Addenbrooke’s Hospital, Cambridge CB2 0XY, UK
19Dept Haematology, University of Cambridge and National Health Service Blood and Transplant, Long Road, Cambridge CB2 2PT, UK
20ALSPAC DNA Bank, Dept Social Medicine, University of Bristol, 24 Tyndall Avenue, Bristol BS8 1TQ, UK
21ALSPAC Laboratory, Dept Social Medicine, Clifton, Bristol BS8 2BN, UK
22Division of Community Health Sciences, St George’s Hospital, London SW17 0RE, UK.
Reference List


34. Spencer CCA. A simple clustering approach to pre-analysis exclusion of individuals from GWAS. In preparation. 2009

Figure 1.
$-\log_{10}(P)$ values from the 1 d.f. trend test. Alternating chromosomes shown in shades of blue. SNPs with $P < 1 \times 10^{-5}$ which had not been previously reported are highlighted in green. The three new loci identified in this study are noted.
Figure 2.
$-\log_{10}(P)$ values from the 1 d.f. trend test from three new loci, along with local recombination rate estimated from HapMap data. Combined $P$ values for replicated SNPs are indicated with a red diamond.
### Table 1
Clinical details of cases and controls

<table>
<thead>
<tr>
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<th>GWAS</th>
<th>Replication cohort</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>CASES</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Age at diagnosis</strong>&lt;sup&gt;a&lt;/sup&gt;</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Early onset (&lt;18 years)</td>
<td>5.9% (112)</td>
<td>6.5% (130)</td>
</tr>
<tr>
<td>Not early onset (&gt;18 years)</td>
<td>94.1% (1783)</td>
<td>93.5% (1861)</td>
</tr>
<tr>
<td>Median</td>
<td>33.4</td>
<td>35.2</td>
</tr>
<tr>
<td>Mean</td>
<td>36.3</td>
<td>38.2</td>
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<tr>
<td><strong>Disease Extent</strong>&lt;sup&gt;b&lt;/sup&gt;</td>
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<tr>
<td>Proctitis</td>
<td>17.9% (357)</td>
<td>15.1% (285)</td>
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<tr>
<td>Left sided</td>
<td>38.8% (774)</td>
<td>47.5% (897)</td>
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<tr>
<td>Extensive</td>
<td>43.3% (864)</td>
<td>37.4% (707)</td>
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<td><strong>Smoking at diagnosis</strong>&lt;sup&gt;c&lt;/sup&gt;</td>
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<tr>
<td>Ex-smoker</td>
<td>36.7% (556)</td>
<td>30.3% (553)</td>
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<tr>
<td>Current smoker</td>
<td>10.8% (163)</td>
<td>16.4% (300)</td>
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<tr>
<td>Never smoked</td>
<td>52.5% (794)</td>
<td>53.2% (971)</td>
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<tr>
<td><strong>Colectomy</strong>&lt;sup&gt;d&lt;/sup&gt;</td>
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<td>15.7% (266)</td>
<td>12.0% (226)</td>
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<tr>
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<td>84.3% (1432)</td>
<td>88.0% (1660)</td>
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<td></td>
<td>1.00% (23)</td>
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<tr>
<td>POBI</td>
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<sup>a</sup>Data available for 80% (GWAS) 86% (Replication),

<sup>b</sup>Data available for 84% (GWAS) 81% (Replication),

<sup>c</sup>Data available for 64% (GWAS) 79% (Replication),

<sup>d</sup>Data available for 72% (GWAS) 81% (Replication),

<sup>e</sup>Data available for 93% (GWAS) 99% (Replication).
New hits from the GWAS

Top tier reaches $5 \times 10^{-8}$ in combined analysis, second tier hits have replication $P < 0.05$ and require further study to completely verify.

<table>
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<tr>
<th>SNP</th>
<th>Chr</th>
<th>LD region(Mb)$^a$</th>
<th>Gene of interest (#)$^b$</th>
<th>$P_{\text{comb}}$</th>
<th>$P_{\text{repl}}$</th>
<th>$P_{\text{comb}}$</th>
<th>Risk allele</th>
<th>RAF$^c$</th>
<th>OR (95% CI)</th>
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<tbody>
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<td>rs886774</td>
<td>7q31.1</td>
<td>107.25-107.39</td>
<td>LAMB1(2)</td>
<td>$4.8 \times 10^{-7}$</td>
<td>0.005</td>
<td>$3 \times 10^{-8}$</td>
<td>G</td>
<td>0.4136</td>
<td>1.11 (1.03-1.19)</td>
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<td>rs1728785</td>
<td>16q22.1</td>
<td>66.98-67.40</td>
<td>CDH1(5)</td>
<td>$1.8 \times 10^{-5}$</td>
<td>0.0004</td>
<td>$2.8 \times 10^{-8}$</td>
<td>G</td>
<td>0.7641</td>
<td>1.17 (1.07-1.27)</td>
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<td>rs6017342</td>
<td>20q13.12</td>
<td>42.49-42.52</td>
<td>HNF4A(7)</td>
<td>$3.2 \times 10^{-13}$</td>
<td>$7.1 \times 10^{-6}$</td>
<td>$8.5 \times 10^{-17}$</td>
<td>C</td>
<td>0.5168</td>
<td>1.17 (1.09-1.26)</td>
</tr>
<tr>
<td>rs7524102*</td>
<td>1p36.12</td>
<td>22.54-22.61</td>
<td>(0)</td>
<td>$1.4 \times 10^{-7}$</td>
<td>0.05</td>
<td>$3.1 \times 10^{-7}$</td>
<td>A</td>
<td>0.8264</td>
<td>1.10 (1.00-1.21)</td>
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<td>rs9548988</td>
<td>13q13.3</td>
<td>39.36-39.56</td>
<td>(0)</td>
<td>$5.0 \times 10^{-6}$</td>
<td>0.0061</td>
<td>$2.7 \times 10^{-7}$</td>
<td>T</td>
<td>0.4594</td>
<td>1.10 (1.03-1.19)</td>
</tr>
</tbody>
</table>

*replication genotyping at this locus is for SNP rs12568930, which is an $r^2 = 1$ proxy for rs7524102.

$^a$LD region of 0.2 cM centered on focal SNP, in NCBI Build 36 coordinates.

$^b$Number of genes in LD region.

$^c$Risk allele frequency.
Table 3

GWAS signals from previously reported UC loci

Top tier were previously reported at genome-wide significance \((5 \times 10^{-8})\), bottom tier were previously reported with weaker evidence. \(P\) values are one-tailed in the direction of the previously reported association.

<table>
<thead>
<tr>
<th>SNP</th>
<th>Chrom</th>
<th>Pos</th>
<th>Gene</th>
<th>Scan (P)</th>
<th>Ref</th>
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<tbody>
<tr>
<td>rs6426833</td>
<td>1p36.13</td>
<td>20044447</td>
<td>OTUD3/PLA2GE</td>
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<td>rs1209026</td>
<td>1p31.3</td>
<td>67478546</td>
<td>IL23R</td>
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<tr>
<td>rs302493</td>
<td>1q32.1</td>
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<td>(8.0 \times 10^{-8})</td>
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</tr>
<tr>
<td>rs10021288</td>
<td>4q27</td>
<td>123224984</td>
<td>IL2/21</td>
<td>0.0033</td>
<td>32</td>
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<tr>
<td>rs9268877</td>
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<td>32539125</td>
<td>MHC</td>
<td>(3.9 \times 10^{-23})</td>
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<td>rs12815372</td>
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<td>0.00070</td>
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<td>rs2094871</td>
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<td>PSMG1</td>
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<td>KIF21B</td>
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<td>CARD9</td>
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