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ORIGINAL ARTICLE

Revealing the complex genetic architecture of obsessive–compulsive disorder using meta-analysis

International Obsessive Compulsive Disorder Foundation Genetics Collaborative (IOCDF-GC) and OCD Collaborative Genetics Association Studies (OCGAS)¹

Two obsessive–compulsive disorder (OCD) genome-wide association studies (GWASs) have been published by independent OCD consortia, the International Obsessive–Compulsive Disorder Foundation Genetics Collaborative (IOCDF-GC) and the OCD Collaborative Genetics Association Study (OCGAS), but many of the top-ranked signals were supported in only one study. We therefore conducted a meta-analysis from the two consortia, investigating a total of 2688 individuals of European ancestry with OCD and 7037 genomically matched controls. No single-nucleotide polymorphisms (SNPs) reached genome-wide significance. However, in comparison with the two individual GWASs, the distribution of *P*-values shifted toward significance. The top haplotypic blocks were tagged with rs4733767 ($P=7.1 \times 10^{-7}$; odds ratio (OR)=1.21; confidence interval (CI): 1.12–1.31, *CASC8/CASC11*), rs1030757 ($P=1.1 \times 10^{-6}$; OR=1.18; CI: 1.10–1.26, *GRID2*) and rs12504244 ($P=1.6 \times 10^{-6}$; OR=1.18; CI: 1.11–1.27, *KIT*). Variants located in or near the genes *ASB13*, *RSPO4*, *DLGAP1*, *PTPRD*, *GRIK2*, *FAIM2* and *CDH20*, identified in linkage peaks and the original GWASs, were among the top signals. Polygenic risk scores for each individual study predicted case–control status in the other by explaining 0.9% ($P=0.003$) and 0.3% ($P=0.0009$) of the phenotypic variance in OCGAS and the European IOCDF-GC target samples, respectively. The common SNP heritability in the combined OCGAS and IOCDF-GC sample was estimated to be 0.28 (s.e.=0.04). Strikingly, ~65% of the SNP-based heritability in the OCGAS sample was accounted for by SNPs with minor allele frequencies of $\geq 40\%$. This joint analysis constituting the largest single OCD genome-wide study to date represents a major integrative step in elucidating the genetic causes of OCD.

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INTRODUCTION

Obsessive–compulsive disorder (OCD) is a psychiatric condition characterized by persistent, intrusive thoughts and urges (obsessions) and repetitive, intentional behaviors (compulsions), typically, but not always, performed to reduce anxiety caused by obsessions.¹ The estimated lifetime prevalence of OCD is 1–3%, based on national surveys.² Individuals with OCD experience a chronic or episodic course with exacerbations that can substantially impair social and occupational functioning.¹

Since the early twentieth century, clinicians have suspected that heredity plays an important role in susceptibility to OCD. Consistent with this, several family studies have found a substantially greater prevalence of OCD (~10-fold increase) in the first-degree relatives of probands, compared with relatives of controls.^{3–6} Family studies of OCD in child and adolescent probands report even greater differences in the risk of OCD in relatives of cases compared with controls,^{7,8} consistent with previous reports of increased familial loading with an early age at onset.^{3,4}

The few existing studies that have examined twin concordance rates for OCD are insufficient in size to allow for accurate heritability estimates.⁹ However, population-based twin studies estimate the heritability of dimensional measures of obsessive–compulsive symptoms (OCS) to be 40–50%, with a similar contribution from nonshared environment and no significant contribution from shared environment.^{10–14} More recently, direct interrogation of the genome using genome-wide complex trait

analysis (GCTA) on data from the International OCD Foundation Genetics Consortium (IOCDF-GC) genome-wide association study (GWAS) provided heritability estimates of 0.37 (s.e.=0.07, $P=1.5 \times 10^{-07}$) for OCD. In the same sample, the estimate of heritability for childhood-onset OCD (symptoms before the age of 17 years¹⁵) was 0.43 (s.e.=0.10, $P=1 \times 10^{-05}$). Partitioning by minor allele frequency (MAF) suggested that the vast majority of the heritability was accounted for by single-nucleotide polymorphisms (SNPs) with MAF >0.30; little heritability was accounted for by SNPs with a MAF of <5%.¹⁵

To date, eight whole-genome studies of OCD or OCS have been published, including five linkage studies,^{16–22} two GWASs of OCD^{23,24} and one GWAS of OCS.²⁵ The five linkage studies identified several chromosomal regions with suggestive evidence for linkage,^{16–20} although there was little overlap between them and only one (1p36) met criteria for statistical significance for linkage.¹⁶ Consistent with sample size expectations for highly polygenic traits, no individual susceptibility variants have yet been identified for OCD using these methods.

The two published GWASs of OCD were conducted by independent OCD consortia: the IOCDF-GC²⁴ and the OCD Collaborative Genetics Association Study (OCGAS).²³ The IOCDF-GC published the first OCD GWAS, comprising 1465 cases and 5557 ancestry-matched controls, as well as 400 complete trios, from 22 sites worldwide.²⁶ The top signal from the combined trio–case–control sample was rs297941 on chromosome 19p13.2, near *FAIM2*

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($P=4.99 \times 10^{-7}$). Although no SNPs were found to be associated with OCD at a genome-wide significance level, a significant enrichment of methylation quantitative trait loci ($P < 0.001$) and frontal lobe expression quantitative trait loci (eQTLs; $P=0.001$) were observed within the top-ranked SNPs, providing evidence, consistent with other disease reports,^{27,28} that biologically relevant associations are present within subthreshold GWAS results. The OCGAS reported a second GWAS, conducted by six research centers in the United States.²³ In this study, 1065 families (containing 1406 patients with OCD), combined with population-based control samples (resulting in a total sample of 5061 individuals), were studied. The smallest P -value ($P=4.13 \times 10^{-7}$) was detected for a SNP on chromosome 9p23, in close proximity to the *protein tyrosine phosphate receptor D (PTPRD)* gene. The second smallest P -value was 1.76×10^{-6} near the *cadherin type 9* and *10 (CDH9 and CDH10)* genes on chromosome 5p15.

A third GWAS, this one examining quantitative OCS, was conducted in 6931 individuals from the Netherlands Twin Registry.²⁵ This study identified one gene that met criteria for genome-wide significance, the *myocyte enhancer factor 2B neighbor (MEF2BNB)* ($P=2.56 \times 10^{-8}$), on chromosome 19p13. The total SNP-based heritability for OCS in this sample was 0.14 (s.e.=0.05, $P=0.003$), and the polygenic risk score (PRS) derived from the IOCDF-GC GWAS was significantly associated with OCS, explaining 0.2% of the variance.

As is evident from the data above, although multiple regions of interest have been reported, there is currently little convergence of results to identify OCD susceptibility variants. This is likely because of genetic and phenotypic heterogeneity, and insufficient sample sizes. Thus, a logical next step is to use comparable data sets in combined analyses to increase power. Here, we report findings from combined analyses of the IOCDF-GC and OCGAS GWAS data aimed at further exploring the genetic underpinnings of OCD. We first used the genotypes of these two studies, after imputation to a common reference, to conduct a joint GWAS. We then used each individual study as a discovery sample for PRS analysis and predicted case-control status in the alternate data set to investigate the amount of phenotypic variation explained by the respective PRS. To replicate the finding that SNPs with high MAF account for the majority of the heritability in OCD, we next computed the common variation heritability of the OCGAS sample using GCTA and performed the same partitioning, as previously reported.¹⁵ Finally, we used linkage disequilibrium (LD) score regression²⁹ to estimate the heritability of OCD based on the combined meta-analysis cohort.

MATERIALS AND METHODS

Samples

For these analyses we used only individuals of European ancestry from the original GWAS samples, yielding 1429 cases, 5089 controls and 285 trios from IOCDF-GC and 344 cases and 1033 controls and 630 trios from OCGAS (Supplementary Table S1), after the addition of screened controls from the Genomic Psychiatry Cohort,³⁰ matched to the OCGAS cases. All cases met DSM-IV (Diagnostic and Statistical Manual of Mental Disorders, 4th edition), criteria for OCD.¹ Controls from the IOCDF-GC GWAS were unscreened. Additional information on the IOCDF-GC and OCGAS samples and methods has been previously published.^{23,26} This work was approved by the relevant institutional review boards at all participating sites, and all participants provided written informed consent.

GWAS analyses

We imputed genotype-level data from the IOCDF-GC (except the Dutch samples that were imputed separately, see below), OCGAS and Genomic Psychiatry Cohort samples using IMPUTE2 (ref. 31) and reference haplotypes from the 1000 Genomes Project (Phase I integrated variant set release); NCBI build 37 (hg19) was constructed with SHAPEIT2.³² We assessed genetic relatedness between samples through identity-by-descent estimation between all sample pairs using PLINK³³ and retained

only one member of each pair of samples with $\text{pi_hat} > 0.2$. Samples were excluded if they had a call rate of < 0.98 , an absolute value of $F_{\text{HET}} > 0.20$ or absence or unambiguous correct genotypic sex. SNPs were excluded from pre-imputation data set if the call rate was < 0.98 , MAF < 0.01 ,

case-control differential missingness was > 0.02 or the P -value of Hardy-Weinberg equilibrium was $< 1.0 \times 10^{-6}$ for controls and $< 1.0 \times 10^{-10}$ for cases. After imputation, SNPs were excluded if IMPUTE2 info was < 0.6 , IMPUTE2 certainty was < 0.9 or MAF was < 0.01 . We assessed population structure using multidimensional scaling and, as previously observed,²⁴ samples of Ashkenazi Jewish or Afrikaans (South African) ethnicity clustered as separate groups (Supplementary Figures S1-S5). We conducted separate association analyses for each case-control subpopulation (IOCDF-GC European (IOEU), IOCDF-GC Ashkenazi Jewish (IOAJ), IOCDF-GC South African (IOSA), OCGAS case-control (OCCC)) and trio sample (IOCDF-GC trios (IOTR) and OCGAS trios (OCTR)); as probands versus pseudo-controls. We defined 'pseudo-controls' as the non-transmitted haplotype pairs from parents to affected offspring in the trio samples.

Because of more stringent data sharing restrictions for Dutch cases, imputation and summary statistics for the Dutch cases and population-matched controls (IODU) were calculated separately by the site investigators following the same imputation and quality control (QC) procedures. We then performed meta-analysis using the summary statistics of all case-control subpopulations (including IODU) and trio samples using METAL³⁴ with the inverse variance method on SNPs that passed QC in at least 500 cases and 500 controls. Results were visualized with AssocPlots.³⁵ The top loci of meta-analysis were defined by the regions of LD pruned independent top SNPs passing predefined P -value threshold ($r^2 < 0.2$, 500 kb window, clump function in PLINK) and their tagged SNPs ($r^2 > 0.2$, 1000 kb window, show-tags function in PLINK) using 1KG samples of European independent founders (EUR, TSI and GBR, phase 1) as the reference panel.

PRS analysis

We conducted PRS analyses using PLINK, as previously described,³⁶ to test whether multiple variants of small effect jointly contribute to OCD. PRSs for subsets of the IOCDF-GC sample (IOEU) and the OCGAS sample (OCTR) (target samples) were calculated based on the SNP effect size estimated from the discovery samples, the OCGAS and European ancestry IOCDF-GC samples (excluding IOAJ), respectively. Imputed SNPs with high quality (IMPUTE2 info > 0.95 , MAF > 0.05) and GWAS P -values passing predetermined significance thresholds ($P < 0.0001, 0.001, 0.01, 0.1, 0.2, 0.3, 0.4$ and 0.5) in the discovery samples were extracted along with their risk alleles and odds ratios (ORs), and then LD pruned within 500 kb window at $r^2 > 0.2$ (clump function in PLINK) using 1KG samples of European independent founders (EUR, TSI and GBR, phase 1) as the reference panel.

For each significance threshold, a quantitative aggregate risk score was calculated for each individual in the target sample IOEU and OCTR, defined as the sum of the number of risk alleles present at each locus weighted by the log of the OR for that locus estimated from the discovery sample. We examined the relationship between aggregate risk score and case-control status in the target samples, IOEU and OCTR, at each significance threshold, using logistic regression controlling for population stratification. We then estimated the percentage of phenotypic variance explained by the aggregate risk score (Nagelkerke's pseudo- R^2).

Heritability and genetic correlations

Genetic complex trait analysis. We used GCTA v1.24 (<http://cns.genomics.com/software/gcta/>) to estimate the proportion of phenotypic variance explained by directly genotyped and imputed SNPs in the OCGAS sample, as has previously been done in the IOCDF-GC sample.¹⁵ Because of the sensitivity of GCTA to low-quality SNPs and remotely related samples, we conducted more stringent QC for these analyses by removing SNPs with Hardy-Weinberg equilibrium $P < 0.05$ and platform effects with $P < 0.01$ (detected by GWAS comparison of platforms among population-matched controls) and removing one member of each sample pair with $\text{pi_hat} > 0.05$. For directly genotyped and imputed SNPs respectively, this resulted in 487 459 directly genotyped and 5 843 119 imputed SNPs on 999 cases and 1064 controls. After QC, we used the GCTA software to generate a genetic relationship matrix (GRM) file that included identity-by-descent relationships calculated from genotype data. Genomic restricted maximum likelihood (GREML) analysis was conducted using the respective

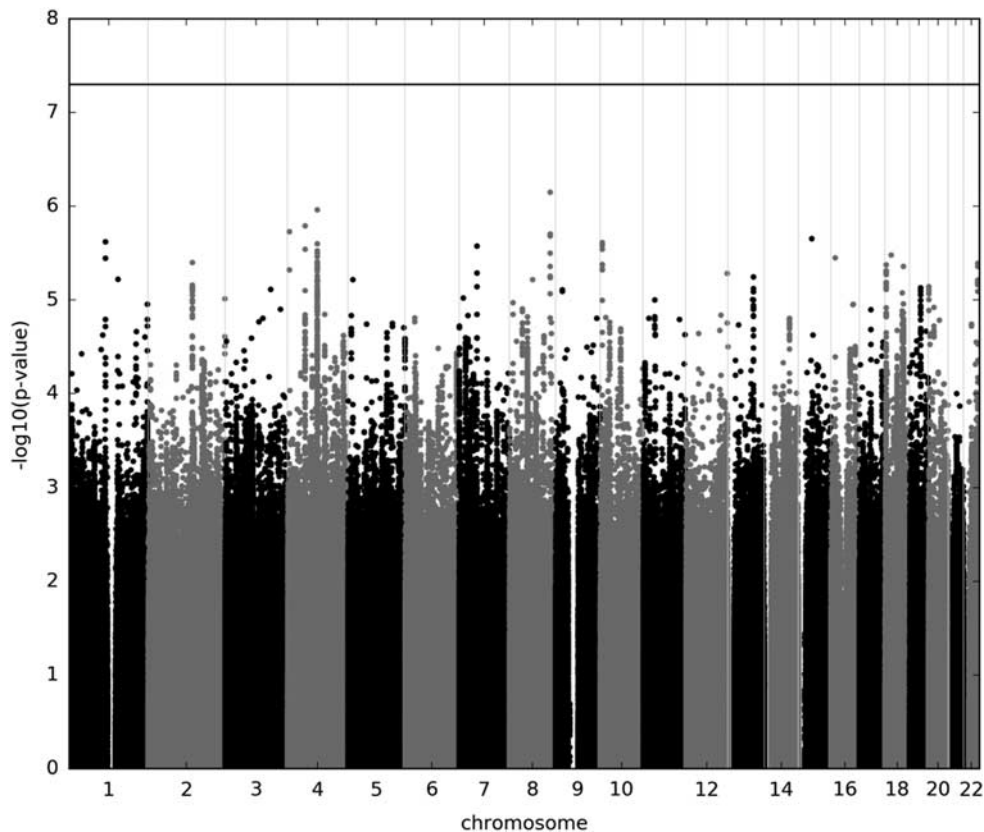


Figure 1. Manhattan plot of genotyped and imputed single-nucleotide polymorphisms (SNPs) for 2688 obsessive-compulsive disorder (OCD) cases and 7031 controls from the International Obsessive-Compulsive Disorder Foundation Genetics Collaborative (IOCDF-GC), OCD Collaborative Genetics Association Study (OCGAS) and Genomic Psychiatry Cohort (GPC) consortia. Black line near the top of the figure indicates the genome-wide significance threshold of $P = 5 \times 10^{-8}$.

GRM estimated from all the SNPs and 20 principal component quantitative covariates. In order to account for the oversampling of cases, we used the OCD population prevalence (2.5%) to transform the estimate of variance explained to the liability scale. Finally, we estimated the chromosome-specific heritability and heritability partitioned by MAF for five MAF bins (0.01–0.1, 0.1–0.2, 0.2–0.3, 0.3–0.4 and 0.4–0.5).

We then combined the two largest European IOCDF-GC and OCGAS data sets (not including trios, IOAJ, IOSA or Dutch), and performed a second round of QC to remove any samples that fell outside the European genetic cluster, and one sample of any pair with a π -hat > 0.05 , resulting in 1323 cases and 4938 controls. At the SNP level, we filtered the imputed SNPs based on the imputation info quality metric (info > 0.6), certainty ($< 80\%$) and MAF (< 0.05) that resulted in 5 235 858 SNPs. A prevalence of 2.5% and 20 principal components were used for the GREML analysis.

LD score regression analysis. We applied the LD score regression (LDSC) method²⁹ to 1 159 580 imputed and directly genotyped SNPs (that overlapped with a panel of high confidence HapMap3 SNPs) measured on all 9725 (2688 cases and 7037 controls) individuals included in the OCD meta-analysis. Regression weights were calculated using the HapMap European reference sample provided by Bulik-Sullivan *et al.*²⁹ To transform from the observed heritability scale to the liability scale we used a population prevalence of 2.5%. Using LDSC, we calculated heritability, checked for residual population stratification (based on the LDSC intercept), and calculated genetic correlation between the two consortium sample collections.

RESULTS

Genome-wide association study

After data cleaning to remove samples falling outside the European genetic cluster, we had a total sample size of 2688

OCD cases and 7037 genomically matched controls, comprising seven subsamples (Supplementary Table S1) that were analyzed individually and then combined by meta-analysis to provide overall P -values on 8 693 187 autosomal SNPs. We generated quantile–quantile plots of the observed versus expected $\log(P)$ values under the null hypothesis and calculated the genomic control lambda for the final sample. We observed no evidence for significant residual stratification effects (Supplementary Figure S6; $\lambda = 1.028$; $\lambda_{1000} = 1.007$). In addition, the LDSC analysis demonstrated no evidence of residual population stratification (LDSC intercept = 1.0005; s.e. = 0.0068).

No SNP exceeded the genome-wide threshold for significance (Figure 1 and Table 1). A total of 130 SNPs (Supplementary Table S2) from 29 LD independent loci (Table 1) were observed with $P < 1.0 \times 10^{-5}$. The SNP with the lowest P -value was rs4733767 ($P = 7.1 \times 10^{-7}$; OR = 1.21; confidence interval (CI): 1.12–1.31) that tagged a haplotypic block of 53.7 kb (chr8: 128 568 359–128 622 083) on chromosome 8q24.21 and is 87.2 kb 5' to *CASC8* (*Cancer Susceptibility Candidate 8*, also known as LOC727677) (Table 1, Supplementary Table S2 and Supplementary Figure S7). The SNP with the second lowest P -value, rs1030757 ($P = 1.1 \times 10^{-6}$; OR = 1.18; CI: 1.10–1.26), on chromosome 4q22.1 tagged the second best haplotype block (chr4: 93 479 275–94 230 511; 751.2 kb; Supplementary Figure S8) that lies wholly within *GRID2* (*Glutamate Ionotropic Receptor Delta Type Subunit 2*). Within this haplotype there were SNPs with heterogeneity P -values of < 0.05 ; we conducted random-effects meta-analysis using PLINK and the findings were different. We identified that the heterogeneity came from the isolate subpopulation South African. Excluding this sample (*post hoc*) from the meta analysis, the

Table 1. LD-independent genomic regions with $P < \times 10^{-5}$ in the combined OCD genome-wide association study (GWAS)

SNP	Chr	BP	A1/A2	A1 FRQ	INFO	Odd ratio (95% CI)	P	LD block (hg19)	Genes
rs4733767	8	128 581 578	A/G	0.274	0.867	1.21 (1.12–1.31)	7.1E–07	128568359–128622083	
rs1030757	4	93 697 153	C/A	0.488	0.979	1.18 (1.10–1.26)	1.1E–06	93479275–94230511	GRID2
rs12504244	4	55 485 188	G/C	0.393	0.974	1.18 (1.11–1.27)	1.6E–06	55476381–55580596	KIT
rs13141765	4	6 243 646	C/T	0.393	0.713	1.31 (1.17–1.46)	1.9E–06	6239015–6249840	LOC285484
rs116347760	1	114 201 251	A/T	0.020	0.793	1.88 (1.44–2.44)	2.4E–06	113751726–114621340	AP4B1, AP4B1-AS1, BCL2L15, DCLRE1B, HIPK1, LOC643441, LOC101928846, MAGI3, OLFML3, PHTF1, PTPN22, RSNB1, SYT6
rs72781967	10	5 622 426	C/T	0.344	0.995	1.18 (1.10–1.27)	2.4E–06	5607539–5659916	
rs55687617	7	56 775 429	A/G	0.119	0.881	0.76 (0.68–0.85)	2.7E–06	56314381–57203788	DKFZp434L192, LOC650226, LOC100130849, LOC100240728, LOC101928401, MIR4283-1, MIR4283-2, ZNF479
rs117310268	18	19 675 267	T/C	0.038	0.780	1.57 (1.30–1.89)	3.3E–06	19505213–19683750	
rs5019028	4	94 222 825	G/T	0.279	0.995	1.19 (1.11–1.28)	3.4E–06	93665778–94232270	GRID2
rs72783425	16	14 148 431	A/C	0.053	0.960	1.40 (1.22–1.62)	3.5E–06	13982844–14318912	ERCC4, LOC101927311, LOC101927348, MKL2
rs56343802	2	139 823 241	T/A	0.281	0.999	1.19 (1.10–1.27)	4.0E–06	139821308–139882668	
rs909701	22	44 973 368	G/C	0.500	0.999	1.17 (1.09–1.25)	4.1E–06	44971548–44988209	LINC00207, LINC00229
rs9952159	18	3 660 801	T/C	0.220	1.026	1.20 (1.11–1.30)	4.2E–06	3638103–3710355	DLGAP1
rs77885126	18	58 420 429	C/T	0.015	0.932	1.83 (1.41–2.36)	4.4E–06	58250265–59372219	CDH20
rs10773765	12	130 767 334	T/C	0.252	0.907	1.20 (1.11–1.30)	5.2E–06	130739141–130857497	PIWIL1
rs9544927	13	79 505 864	G/A	0.208	0.996	0.82 (0.76–0.90)	5.7E–06	79496764–79551012	
rs28599745	1	153 396 665	A/G	0.160	0.650	0.70 (0.59–0.81)	6.0E–06	153349194–153396665	S100A7A, S100A7L2, S100A8, S100A9, S100A12
rs1652783	8	73 279 728	G/A	0.225	0.764	1.31 (1.16–1.47)	6.1E–06	73279414–73312929	
rs190543171	5	15 840 912	T/C	0.012	0.787	2.32 (1.61–3.35)	6.1E–06	15726922–16139424	FBXL7, MARCH11
rs56025909	20	955 893	T/C	0.032	0.960	1.51 (1.26–1.81)	7.2E–06	931170–991579	RSP04
rs3097331	19	34 648 956	C/T	0.368	1.000	0.85 (0.80–0.92)	7.5E–06	34632485–34727202	KIAA0355, LSM14A
rs138445568	3	143 922 936	T/A	0.013	0.630	2.53 (1.68–3.80)	7.7E–06	142971745–144868438	C3orf58, SLC9A9, SLC9A9-AS1
rs139286049	9	20 688 387	G/A	0.019	0.760	1.82 (1.40–2.37)	7.8E–06	20688387–21100533	FOCAD, IFNB1, MIR491, PTPLAD2
rs146238482	9	20 856 226	C/G	0.010	0.823	2.68 (1.74–4.13)	8.1E–06	20836624–21016372	FOCAD, PTPLAD2
rs35894340	18	54 307 062	G/A	0.252	0.948	1.19 (1.10–1.29)	8.4E–06	54246712–54518866	TXNL1, WDR7
rs149952789	7	12 716 170	T/A	0.010	0.603	3.33 (1.96–5.68)	9.6E–06	12716170–12716170	ARL4A
rs4444795	4	93 355 172	T/C	0.183	0.912	1.22 (1.12–1.33)	9.7E–06	93195208–94077524	GRID2
rs75740353	2	242 741 686	A/G	0.103	0.869	0.65 (0.54–0.79)	9.7E–06	242685298–242764651	D2HGDH, GAL3ST2, INGS5, NEU4
rs116969557	13	77 337 736	A/G	0.016	0.957	1.77 (1.38–2.29)	9.9E–06	77082597–77566923	BTF3P11, CLN5, FBXL3, IRG1, KCTD12

Abbreviations: A1, minor allele; A1 FRQ, frequency of A1 allele; A2, major allele; BP, chromosomal position, in base pairs; Chr, chromosome; CI, confidence interval; INFO, INFO score; LD, linkage disequilibrium; OCD, obsessive-compulsive disorder; SNP, single-nucleotide polymorphism LD block indicates tagged region by the index SNP at $r^2 > 0.2$; Genes indicate genes and their 20 kb flanking regions on each side overlapped with the LD block.

association with OCD for the SNPs with heterogeneity became more significant (for example, rs7683744: $P = 8.0 \times 10^{-7}$; OR = 0.84). The SNP with the third lowest P -value, rs12504244 ($P = 1.6 \times 10^{-6}$; OR = 1.18; CI: 1.11–1.27), on chromosome 4q12 tagged the third best haplotypic block (chr4: 55 476 381–55 580 596; 104.2 kb; Supplementary Figure S9) that overlies the promoter and much of the gene body of *KIT* (*KIT Proto-Oncogene Receptor Tyrosine Kinase*).

PRS analysis

We used SNP effect sizes derived from the individual OCGAS and IOCDF-GC meta-analyses to calculate PRS and predict OCD status in individuals of European ancestry from the IOCDF-GC sample (IOEU) and in trios of European ancestry from the OCGAS sample (OCTR), respectively. As shown in Figure 2 and Supplementary Table S3, the PRS derived from meta-analysis of the European IOCDF-GC samples (excluding the IOAJ samples to avoid heterogeneity in the discovery sample) reasonably predicted case-control status in the OCGAS trio target sample ($P = 0.003$), explaining ~0.9% of the phenotypic variance. Conversely, risk

scores derived using the OCGAS as a discovery sample explained 0.3% of the phenotypic variance in the IOEU samples ($P = 0.0009$).

Heritability analyses

GCTA-based heritability in the OCGAS sample alone (999 cases and 1064 controls; Table 2) was 0.25 (s.e. = 0.11, $P = 0.0096$). The GCTA heritability estimate of OCD in the combined OCGAS and IOCDF-GC European sample was also 0.25 (s.e. = 0.05; $P = 0.0096$). LDSC analysis yielded a heritability estimate of 0.28 (s.e. = 0.04) for the combined OCD sample (Table 2). When the sample was then split into its constituent parts (OCGAS and IOCDF-GC), we observed a significant genetic correlation between the two ($r_g = 0.83$; s.e. = 0.28; $P = 0.003$).

In parallel with the univariate and bivariate analyses, we partitioned heritability by allele frequency in the OCGAS sample using five MAF bins as 0.01–0.1, 0.1–0.2, 0.2–0.3, 0.3–0.4 and 0.4–0.5 in the GRMEL joint analysis; results are presented in Supplementary Table S4. As has been previously reported for the IOCDF-GC sample,¹⁵ the largest proportion of heritability was observed in the highest frequency allele bins (MAF > 0.4).

DISCUSSION

We report the results of a meta-analysis of GWAS from the two published genome-wide association studies of OCD, with a sample of 2688 individuals with OCD that include both family-based and singleton cases, and 7037 controls. PRS and LDSC analyses confirm that the two samples share genetic risk factors for OCD, and are thus appropriate for combined GWAS analyses. With LDSC we observed a strong genetic correlation between the IOCDF-GC and OCGAS samples ($r_g = 0.83$, s.e. = 0.28; $P = 0.003$). PRS derived from each sample significantly predicted case-control status in the other sample. Although the phenotypic variance explained was relatively small ($R^2 = 0.9\%$ for the OCGAS trios and 0.3% for the IOCDF-GC European cases and controls), they are comparable to those found for schizophrenia, using similar discovery sample sizes³⁷ (PGC2 SCZ GWAS, 2014).

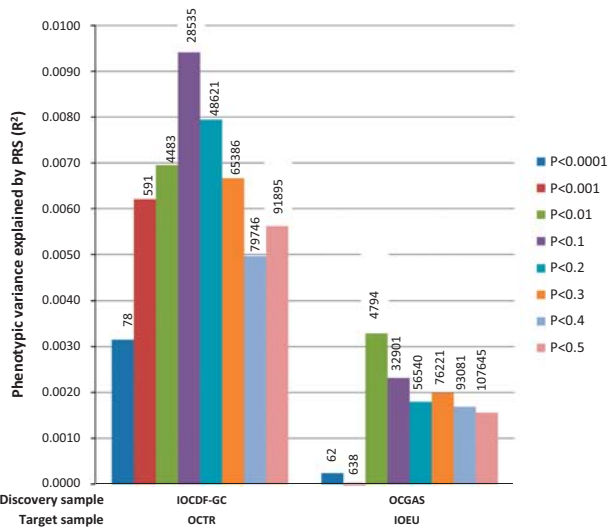


Figure 2. Polygenic risk score (PRS) analysis in obsessive-compulsive disorder (OCD). The variance explained in two target samples (OCTR, consisting of 630 cases and 630 pseudo-controls; IOEU, consisting of 1032 cases and 4100 controls) is based on risk scores derived from an aggregated sum of weighted single-nucleotide polymorphism (SNP) risk allele effect sizes estimated from the discovery samples (IOCDF-GC without IOAJ, consisting of 1623 cases and 5113 controls; OCGAS, consisting of 974 cases and 663 controls) at eight significance thresholds. The numbers of SNPs used at each significance thresholds for PRS were listed on the top of the corresponding bars. The y axis indicates Nagelkerke's pseudo- R^2 . IOCDF-GC, International Obsessive-Compulsive Disorder Foundation Genetics Collaborative; OCGAS, OCD Collaborative Genetics Association Study.

Using GCTA, the heritability tagged by the SNPs in the OCGAS sample was slightly lower (25%) than previously observed for the IOCDF-GC sample (32%).¹⁵ The ascertainment strategies differed in IOCDF-GC and OCGAS studies, with the former recruiting individuals, and the latter primarily families (trios) that may underestimate the heritability tagged by SNPs, as the polygenic load in family members of affected individuals is elevated in comparison with controls.³⁸ Joint heritability analyses of the two samples, using GCTA and LDSC, resulted in similar estimates (0.25 and 0.28, respectively), suggesting that the common variation heritability of OCD is between 25 and 30% (that is, $\geq 50\%$ of the total heritability than estimated by twin studies).

We also examined the allele frequency distribution of the common variation heritability of OCD. Although the confidence intervals of each allele frequency bin are large because of the limited sample sizes, the majority of the heritability ($\sim 65\%$) was accounted for by SNPs with high MAF (for example, $> 40\%$) in both the OCGAS sample alone and combined sample (Supplementary Table S4).

Although there were no genome-wide significant findings, the 53.7 kb (chr8: 128 568 359–128 622 083) haplotype block encompassing the top SNP, rs4733767 (Supplementary Figure S7), contains 25 H3K27Ac peaks in the ENCODE/ROADMAP data, suggesting it has regulatory potential,³⁹ although the current release of the Genotype-Tissue Expression project (GTEx Release V6 (dbGaP Accession phs000424.v6.p1))⁴⁰ has no eQTLs in the block. The closest genes on either side of rs4733767 (CASC8 and CASC11) are long noncoding RNAs that are thought as a class to have regulatory functions.⁴¹ Both are only expressed at low levels in the brain in the GTEx database. The potential transcriptional consequences of genomic risk for OCD in this region are unclear, at present.

The second best haplotypic block (chr4: 93 479 275–94 230 511) lies entirely within *GRID2*, a gene expressing a subunit of an ionotropic glutamate receptor, and contains ~ 300 H3K27Ac peaks. The region between $\sim 94 120 000$ and $94 230 000$ kb contains multiple SNPs that regulate *GRID2* in both brain (www.BRAINEAC.org, intralobular white matter ($n = 131$))⁴² and testis (GTEx Release V6). In the latter, two of these SNPs overlap with those observed in this study (rs7684707 and rs5019028), and the OCD risk allele is predicted to increase expression. These eQTLs were not detected in brain in the GTEx study, most likely as a consequence of small sample size. *GRID2* is highly expressed in the cerebellum, but is also expressed in other regions of the brain throughout the lifespan (www.BrainSpan.org), with detectable expression in the caudate, putamen, nucleus accumbens and the anterior cingulate cortex, all regions that have been implicated in OCD,⁴³ and is part of the glutamatergic signaling system that is thought to be important in OCD.⁴⁴ Deletions of portions of *GRID2* in humans are responsible for spinocerebellar ataxia, autosomal recessive 18 (SCAR18; <http://omim.org/entry/602368>) that are severe when homozygous and milder when heterozygous.⁴⁵

Table 2. Heritability estimates for OCD in the IOCDF-GC and OCGAS samples

Sample characteristics	Method	Cases	Controls	Number of SNPs	Reference	V(g)/V(p)_Liability (s.e.)
IOCDF-GC case-control	GCTA	1061	4236	7 657 106	Davis et al. ¹⁵	0.32
OCGAS case-control and trio controls	GCTA	999	1064	5 843 119	This study	0.25
OCGAS and IOCDF-GC cases-control EU only	GCTA	1323	4398	5 235 858	This study	0.25
OCGAS and IOCDF-GC case-control and pseudo-control	LDSC	2936	7279	1 159 580	This study	0.28

Abbreviations: GCTA, genome-wide complex trait analysis; IOCDF-GC, International Obsessive-Compulsive Disorder Foundation Genetics Collaborative; LDSC, linkage disequilibrium score regression; OCD, obsessive-compulsive disorder; OCGAS, OCD Collaborative Genetics Association Study; SNP, single-nucleotide polymorphism.

These observations suggest that lower *GRID2*, particularly in the cerebellum causes ataxia, whereas higher *GRID2*, especially in the non-cerebellar brain, may increase risk for OCD.

The third best haplotypic block observed contains the promoter and much of the gene body of *KIT*. No eQTLs for *KIT* are found in GTEx (V6), but the haplotypic block is likely to regulate the gene, as it contains 47.7 kb 5' to the transcription start site, and has 76 H3K27Ac peaks. *KIT* is expressed in multiple brain regions (BrainSpan and GTEx) and across the human lifespan, with highest expression during fetal development (BrainSpan). Allelic variants of *KIT* in humans have been observed in individuals with piebaldism, various leukemias and gastrointestinal stromal tumors (<http://omim.org/entry/164920?search=kit&highlight=kit>). Of note, both *KIT* and *GRID2* are regulated by transforming growth factor- β 1 in rodents.^{46–48}

Comparison of findings in prior linkage or GWAS studies of OCD Of the signals with P -values of $< 1 \times 10^{-5}$ in this meta-analysis, two were in genomic regions that have been previously identified in genome-wide linkage studies. These include 7 SNPs on chromosome 10p15,¹⁸ all of which are eQTLs of *ASB13* (*ankyrin repeat and SOCS box containing 13*), in Epstein–Barr virus-transformed lymphocytes, and predict high expression (GTEx Release V6). A 60.4 kb haplotypic block was seen in a linkage peak on 20p13 (ref.¹⁶) that encompasses the gene for *RSPO4* and part of its promoter. The block contains SNPs that are eQTLs for *RSPO4* in about a dozen tissues and an eQTL for *SRXN1* in thyroid. As mentioned above, the number of brain samples in the V6 release is ≤ 100 , limiting the power to detect brain eQTLs. Overall, most of the eQTLs being detected at the present sample sizes in the GTEx project affect multiple tissues, and hence it is plausible that these SNPs may also be regulating the same genes in brain. Final determination of this will require more data.

Of the signals identified in the three prior OCD GWASs,^{23–25} SNPs within *DLGAP1*, which was identified in the IOCDF-GC GWAS, represented the signal most strongly supported in this meta-analysis (best was rs9952159, $P = 4.2 \times 10^{-6}$, OR = 1.20). Among the other genes of interest, signals in or near *PTPRD*, which was previously identified in the OCGAS GWAS ($P = 2.4 \times 10^{-4}$, OR = 1.45), *GRIK2* (rs116966225, $P = 5.4 \times 10^{-4}$, -158.5 kb and rs78014006, $P = 7.2 \times 10^{-3}$, intronic) and *FAIM2* (rs297941, $P = 6.1 \times 10^{-5}$, 21.3 kb), were also identified in this meta-analysis, although not among the top hits. Although no signal was identified in this meta-analysis for either *CDH9* or *CDH10*, we did identify a strong signal for a related cadherin gene *CDH20* (rs77885126, $P = 4.4 \times 10^{-6}$, OR = 1.83). It should be noted that according to a power analysis, the sample had low power to detect genome-wide significant association with a common SNP conferring an OR of ≤ 1.2 . The OCS GWAS was omitted from this study because it employed a different phenotype; that is, it used a self-report assessment tool that measured the presence of symptoms and not the diagnosis of OCD, as opposed to a clinical assessment that was based on OCD diagnostic criteria. We omitted it because of these differences in phenotypes, and the desire to not introduce additional heterogeneity into the study. The OCS GWAS identified *RFXANK* as a top signal; it was also identified among the top SNPs of this meta-analysis (rs11666960, $P = 6.3 \times 10^{-4}$, OR = 0.87).

Overall, the results from this meta-analysis support some of the preexisting findings generated from two previous GWASs of OCD. Among the best findings in this study are several glutamatergic system genes (that is, *GRID2*, *DLGAP1*). Evidence has implicated abnormalities in this system as part of the etiology of OCD and the most robust candidate gene study results have consistently identified genes involved in this neurotransmitter system (*GRIN2B*⁴⁹ and *SLC1A1* (ref. 50)). Therefore, further dissection of glutamatergic system genes along with increasing sample sizes

will improve our understanding of the underlying mechanism of OCD.

As sample sizes grow and sequencing costs reduce further, we anticipate that genetic associations with OCD will become increasingly robust, and that a proportion of these currently suggestive findings will reach genome-wide significance.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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REFERENCES

- 1 American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*, 5th edn. American Psychiatric Association: Arlington, VA, 2013.
- 2 Karno M, Golding JM. Obsessive compulsive disorder. In: Robins LN, Regier DA (eds). *Psychiatric Disorders in America: The Epidemiologic Catchment Area Study*. Free Press; Collier Macmillan Canada; Maxwell Macmillan International: New York, Toronto, 1991, pp 204–219.
- 3 Pauls DL, Alsobrook JP 2nd, Goodman W, Rasmussen S, Leckman JF. A family study of obsessive-compulsive disorder. *Am J Psychiatry* 1995; **152**: 76–84.
- 4 Nestadt G, Samuels J, Riddle M, Bienvenu OJ 3rd, Liang KY, LaBuda M *et al*. A family study of obsessive-compulsive disorder. *Arch Gen Psychiatry* 2000; **57**: 358–363.
- 5 Grabe HJ, Ruhrmann S, Ettl S, Buhtz F, Hochrein A, Schulze-Rauschenbach S *et al*. Familiality of obsessive-compulsive disorder in nonclinical and clinical subjects. *Am J Psychiatry* 2006; **163**: 1986–1992.
- 6 Fyer AJ, Lipsitz JD, Mannuzza S, Aronowitz B, Chapman TF. A direct interview family study of obsessive-compulsive disorder. I. *Psychol Med* 2005; **35**: 1611–1621.
- 7 Hanna GL, Himle JA, Curtis GC, Gillespie BW. A family study of obsessive-compulsive disorder with pediatric probands. *Am J Med Genet B Neuropsychiatr Genet* 2005; **134**: 13–19.
- 8 do Rosario-Campos MC, Leckman JF, Curi M, Quatrano S, Katsovtch L, Miguel EC *et al*. A family study of early-onset obsessive-compulsive disorder. *Am J Med Genet B Neuropsychiatr Genet* 2005; **136**: 92–97.
- 9 van Grootheest DS, Cath DC, Beekman AT, Boomsma DI. Twin studies on obsessive-compulsive disorder: a review. *Twin Res Hum Genet* 2005; **8**: 450–458.
- 10 Clifford CA, Murray RM, Fulker DW. Genetic and environmental influences on obsessional traits and symptoms. *Psychol Med* 1984; **14**: 791–800.
- 11 Jonnal AH, Gardner CO, Prescott CA, Kendler KS. Obsessive and compulsive symptoms in a general population sample of female twins. *Am J Med Genet* 2000; **96**: 791–796.
- 12 Eley TC, Bolton D, O'Connor TG, Perrin S, Smith P, Plomin R. A twin study of anxiety-related behaviours in pre-school children. *J Child Psychol Psychiatry* 2003; **44**: 945–960.
- 13 Hudziak JJ, Van Beijsterveldt CE, Althoff RR, Stanger C, Rettew DC, Nelson EC *et al*. Genetic and environmental contributions to the Child Behavior Checklist Obsessive-Compulsive Scale: a cross-cultural twin study. *Arch Gen Psychiatry* 2004; **61**: 608–616.
- 14 Taylor S. Etiology of obsessions and compulsions: a meta-analysis and narrative review of twin studies. *Clin Psychol Rev* 2011; **31**: 1361–1372.
- 15 Davis LK, Yu D, Keenan CL, Gamazon ER, Konkashbaev AI, Derks EM *et al*. Partitioning the heritability of tourette syndrome and obsessive compulsive disorder reveals differences in genetic architecture. *PLoS Genet* 2013; **9**: e1003864.
- 16 Mathews CA, Badner JA, Andresen JM, Sheppard B, Himle JA, Grant JE *et al*. Genome-wide Linkage Analysis of Obsessive-Compulsive Disorder Implicates Chromosome 1p36. *Biol Psychiatry* 2012; **72**: 629–636.
- 17 Ross J, Badner J, Garrido H, Sheppard B, Chavira DA, Grados M *et al*. Genomewide linkage analysis in Costa Rican families implicates chromosome 15q14 as a candidate region for OCD. *Hum Genet* 2011; **130**: 795–805.
- 18 Hanna GL, Veenstra-Vanderweele J, Cox NJ, Van Etten M, Fischer DJ, Himle JA *et al*. Evidence for a susceptibility locus on chromosome 10p15 in early-onset obsessive-compulsive disorder. *Biol Psychiatry* 2007; **62**: 856–862.
- 19 Shugart YY, Samuels J, Willour VL, Grados MA, Greenberg BD, Knowles JA *et al*. Genomewide linkage scan for obsessive-compulsive disorder: evidence for susceptibility loci on chromosomes 3q, 7p, 1q, 15q, and 6q. *Mol Psychiatry* 2006; **11**: 763–770.
- 20 Hanna GL, Veenstra-Vanderweele J, Cox NJ, Boehnke M, Himle JA, Curtis GC *et al*. Genome-wide linkage analysis of families with obsessive-compulsive disorder ascertained through pediatric probands. *Am J Med Genet* 2002; **114**: 541–552.
- 21 Wang Y, Samuels JF, Chang YC, Grados MA, Greenberg BD, Knowles JA *et al*. Gender differences in genetic linkage and association on 11p15 in obsessive-compulsive disorder families. *Am J Med Genet B Neuropsychiatr Genet* 2009; **150B**: 33–40.
- 22 Samuels J, Shugart YY, Grados MA, Willour VL, Bienvenu OJ, Greenberg BD *et al*. Significant linkage to compulsive hoarding on chromosome 14 in families with obsessive-compulsive disorder: results from the OCD Collaborative Genetics Study. *Am J Psychiatry* 2007; **164**: 493–499.
- 23 Mattheisen M, Samuels JF, Wang Y, Greenberg BD, Fyer AJ, McCracken JT *et al*. Genome-wide association study in obsessive-compulsive disorder: results from the OCGAS. *Mol Psychiatry* 2015; **20**: 337–344.
- 24 Stewart SE, Yu D, Scharf JM, Neale BM, Fagerness JA, Mathews CA *et al*. Genome-wide association study of obsessive-compulsive disorder. *Mol Psychiatry* 2013; **18**: 788–798.
- 25 den Braber A, Zilhao NR, Fedko IO, Hottenga JJ, Pool R, Smit DJ *et al*. Obsessive-compulsive symptoms in a large population-based twin-family sample are predicted by clinically based polygenic scores and by genome-wide SNPs. *Transl Psychiatry* 2016; **6**: e731.
- 26 Stewart SE, Yu D, Scharf JM, Neale BM, Fagerness JA, Mathews CA *et al*. Genome-wide association study of obsessive-compulsive disorder. *Mol Psychiatry* 2013; **18**: 788–798.
- 27 Davis LK, Gamazon ER, Kistner-Griffin E, Badner JA, Liu C, Cook EH *et al*. Loci nominally associated with autism from genome-wide analysis show enrichment of brain expression quantitative trait loci but not lymphoblastoid cell line expression quantitative trait loci. *Mol Autism* 2012; **3**: 3.
- 28 Below JE, Parra EJ, Gamazon ER, Torres J, Krithika S, Candille S *et al*. Meta-analysis of lipid-traits in Hispanics identifies novel loci, population-specific effects, and tissue-specific enrichment of eQTLs. *Sci Rep* 2016; **6**: 19429.
- 29 Bulik-Sullivan BK, Loh PR, Finucane HK, Ripke S, Yang J. Schizophrenia Working Group of the Psychiatric Genomics Consortium. LD Score regression distinguishes confounding from polygenicity in genome-wide association studies. *Nat Genet* 2015; **47**: 291–295.
- 30 Pato MT, Sobell JL, Medeiros H, Abbott C, Sklar BM, Buckley PF *et al*. The genomic psychiatry cohort: partners in discovery. *Am J Med Genet B Neuropsychiatr Genet* 2013; **162B**: 306–312.
- 31 Howie B, Marchini J, Stephens M. Genotype imputation with thousands of genomes. *G3 (Bethesda)* 2011; **1**: 457–470.
- 32 Delaneau O, Howie B, Cox AJ, Zagury JF, Marchini J. Haplotype estimation using sequencing reads. *Am J Hum Genet* 2013; **93**: 687–696.
- 33 Purcell S, Neale B, Todd-Brown K, Thomas L, Ferreira MA, Bender D *et al*. PLINK: a tool set for whole-genome association and population-based linkage analyses. *Am J Hum Genet* 2007; **81**: 559–575.
- 34 Willer CJ, Li Y, Abecasis GR. METAL: fast and efficient meta-analysis of genome-wide association scans. *Bioinformatics* 2010; **26**: 2190–2191.
- 35 Khrantsova EA, Stranger BE. Assocplots: a python package for static and interactive visualization of multiple-group gwas results. *Bioinformatics* 2017; **33**: 432–434.
- 36 Yu D, Mathews CA, Scharf JM, Neale BM, Davis LK, Gamazon ER *et al*. Cross-disorder genome-wide analyses suggest a complex genetic relationship between Tourette's syndrome and OCD. *Am J Psychiatry* 2015; **172**: 82–93.
- 37 Schizophrenia Working Group of the Psychiatric Genomics Consortium. Biological insights from 108 schizophrenia-associated genetic loci. *Nature* 2014; **511**: 421–427.
- 38 Klei L, Sanders SJ, Murtha MT, Hus V, Lowe JK, Willsey AJ *et al*. Common genetic variants, acting additively, are a major source of risk for autism. *Mol Autism* 2012; **3**: 9.
- 39 Malik AN, Vierbuchen T, Hemberg M, AA Rubin, Ling E, Couch CH *et al*. Genome-wide identification and characterization of functional neuronal activity-dependent enhancers. *Nat Neurosci* 2014; **17**: 1330–1339.
- 40 GTEx Consortium. The Genotype-Tissue Expression (GTEx) pilot analysis: multitissue gene regulation in humans. *Science* 2015; **348**: 648–660, 40.
- 41 Ng SY, Lin L, Soh BS, Stanton LW. Long noncoding RNAs in development and disease of the central nervous system. *Trends Genet* 2013; **29**: 461–468.
- 42 Ramasamy A, Trabzuni D, Guelfi S, Varghese V, Smith C, Walker R *et al*. Genetic variability in the regulation of gene expression in ten regions of the human brain. *Nat Neurosci* 2014; **17**: 1418–28.
- 43 Graybiel AM, Rauch SL. Toward a neurobiology of obsessive-compulsive disorder. *Neuron* 2000; **28**: 343–7.
- 44 Pittenger C, Bloch MH, Williams K. Glutamate abnormalities in obsessive compulsive disorder: neurobiology, pathophysiology, and treatment. *Pharmacol Ther* 2011; **132**: 314–332.
- 45 Coutelier M, Burglen L, Mundwiler E, Abada-Bendib M, Rodriguez D, Chantot-Bastarud S *et al*. GRID2 mutations span from congenital to mild adult-onset cerebellar ataxia. *Neurology* 2015; **28**: 1751–1759.
- 46 Fernando J, Faber TW, Pullen NA, Falanga YT, Kolawole EM, Oskertizian CA *et al*. Genotype-dependent effects of TGF- β 1 on mast cell function: targeting the Stat5 pathway. *J Immunol* 2013; **191**: 4505–4513.

- 47 Kawakami T, Soma Y, Kawa Y, Ito M, Yamasaki E, Watabe H *et al*. Transforming growth factor beta1 regulates melanocyte proliferation and differentiation in mouse neural crest cells via stem cell factor/KIT signaling. *J Invest Dermatol* 2002; **118**: 471–478.
- 48 Cacheaux LP, Ivens S, David Y, Lakhter AJ, Bar-Klein G, Shapira M *et al*. Transcriptome profiling reveals TGF-beta signaling involvement in epileptogenesis. *J Neurosci* 2009; **29**: 8927–8935.
- 49 Arnold PD, Rosenberg DR, Mundo E, Tharmalingam S, Kennedy JL, Richter MA. Association of a glutamate (NMDA) subunit receptor gene (GRIN2B) with obsessive-compulsive disorder: a preliminary study. *Psychopharmacology (Berl)* 2004; **174**: 530–538.
- 50 Stewart SE, Mayerfeld C, Arnold PD, Crane JR, O'Dushlaine C, Fagerness JA *et al*. Meta-analysis of association between obsessive-compulsive disorder and the 3' region of neuronal glutamate transporter gene SLC1A1. *Am J Med Genet B Neuropsychiatr Genet* 2013; **162B**: 367–379.

Supplementary Information accompanies the paper on the Molecular Psychiatry website (<http://www.nature.com/mp>)