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Genome-wide association meta-analysis in 269,867 individuals identifies new genetic and functional links to intelligence

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Genome-wide association meta-analysis in 269,867 individuals identifies new genetic and functional links to intelligence

Intelligence is highly heritable\(^1\) and a major determinant of human health and well-being\(^2\). Recent genome-wide meta-analyses have identified 24 genomic loci linked to variation in intelligence\(^3–7\), but much about its genetic underpinnings remains to be discovered. Here, we present the largest genetic association study of intelligence to date (N=269,867), identifying 205 associated genomic loci (190 novel) and 1,016 genes (939 novel) via positional mapping, expression quantitative trait locus (eQTL) mapping, chromatin interaction mapping, and gene-based association analysis. We find enrichment of genetic effects in conserved and coding regions and associations with 146 nonsynonymous exonic variants. Associated genes are strongly expressed in the brain, specifically in striatal medium spiny neurons and hippocampal pyramidal neurons. Gene-set analyses implicate pathways related to nervous system development and synaptic structure. We confirm previous strong genetic correlations with multiple health-related outcomes, and Mendelian randomization results suggest protective effects of intelligence for Alzheimer’s disease and ADHD, and bidirectional causation with pleiotropic effects for schizophrenia. These results are a major step forward in understanding the neurobiology of cognitive function as well as genetically related neurological and psychiatric disorders.

We performed a genome-wide association (GWAS) meta-analysis of 14 independent epidemiological cohorts of European ancestry and 9,295,118 genetic variants passing quality control. Additional details are provided in the Supplementary Methods.

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Competing Financial Interests Statement
PF Sullivan reports the following potentially competing financial interests: Lundbeck (advisory committee), Pfizer (Scientific Advisory Board member), and Roche (grant recipient, speaker reimbursement). O Breen reports consultancy and speaker fees from Eli Lilly and Illumina and grant funding from Eli Lilly. J Hjerling-Leffler reports interests from Cartana (Scientific Advisor) and Roche (grant recipient). TD Cannon is a consultant to Boehringer Ingelheim Pharmaceuticals and Lundbeck A/S. All other authors declare no financial interests or potential conflicts of interest.

Data Availability Statement
Summary statistics will be available for download upon publication from https://ctg.cncr.nl.
control (Table 1; Supplementary Table 1; Supplementary Figure 1). A flowchart of the study methodology is presented in Supplementary Figure 2 and additional details of the methods and results are presented in the Supplementary Note.

Intelligence was assessed using various neurocognitive tests, primarily gauging fluid domains of cognitive functioning (Supplementary Information 1.1–1.2). Despite variation in form and content, cognitive test scores display a positive manifold of correlations, a robust empirical phenomenon that is observed in multiple populations. Statistically, the variance common across cognitive tasks can be modeled as a latent factor denoted as \( g \) (the general factor of intelligence). In addition, twin- and family studies show strong genetic correlations across diverse cognitive domains, suggesting pleiotropy, and across levels of ability, substantiating the view of general intelligence as an aetiological continuum (with rare syndromic forms of severe intellectual disability being the exception). Additionally, \( g \)-factors extracted from different sets of cognitive tests correlate very strongly (>.98), supporting the universality of \( g \).

Despite sample and methodological variations, genetic correlations \( (r_g) \) between cohorts were considerable (mean=0.67), and there was no evidence of heterogeneity between cohorts in the single nucleotide polymorphism (SNP) associations (Supplementary Table 2; Supplementary Results 2.1). Age-stratified meta-analyses indicated high genetic correlations \( (r_g>0.62) \), and comparable heritability across age, as captured by the SNPs included in the analysis \( (h^2_{SNP}=0.19–0.22) \) (Supplementary Table 3; Supplementary Results 2.2). The full sample \( h^2_{SNP} \) was 0.19 (SE=0.01), in line with previous findings, and an LD score intercept of 1.08 (SE=0.02) indicated that most of the inflation \( (\lambda_{GC}=1.92) \) could be explained by polygenic signal.

In the meta-analysis, 12,110 variants indexed by 242 lead SNPs in approximate linkage equilibrium \( (r^2<0.1) \) reached genome-wide significance (GWS; \( P<5\times10^{-8} \)) (Figure 1a; Supplementary Tables 5–7; Supplementary Figures 4–5). These were located in 205 distinct genomic loci (Supplementary Results 2.3.1). We tested for replication using the proxy phenotype of educational attainment, which is correlated phenotypically \( (r=0.40) \) and genetically \( (r=0.70) \) with intelligence. We confirmed this high genetic correlation \( (r_g=0.73) \) and observed sign concordance with educational attainment for 93% of GWS SNPs \( (P<1\times10^{-300}) \), with replication for 48 loci (Supplementary Results 2.3.2; Supplementary Table 8). Using polygenic score prediction, the current results explain up to 5.2% of the variance in intelligence in four independent samples (Supplementary Table 9, Supplementary Results 2.3.3).

We observed enrichment for heritability of SNPs in conserved regions \( (P=2.01\times10^{-12}) \), coding regions \( (P=1.67\times10^{-6}) \), and H3K9ac histone regions/peaks \( (P=6.26\times10^{-5}) \), and among common (minor allele frequency > 0.3) variants (Figure 1b; Supplementary Results 2.3.4; Supplementary Table 10; Supplementary Figures 6–7). Conserved and regulatory regions have previously been implicated in cognitive functioning but coding regions have not.
Functional annotation of all candidate SNPs in the associated loci (SNPs with an $r^2 \geq 0.6$ with one of the independent significant SNPs, a suggestive $P$-value ($<1 \times 10^{-5}$) and a MAF $>0.0001$; $n=21,368$) showed that these were mostly intronic/intergenic (Supplementary Table 6; Figure 1), yet 146 (81 GWS) SNPs were exonic non-synonymous (ExNS) (Supplementary Table 11, Supplementary Results 2.3.5). Convergent evidence of strong association ($Z=9.49$) and the highest observed probability of a deleterious protein effect (CADD$^{23}$ score=34) was found for rs13107325. This missense mutation (MAF=0.065, $P=2.23 \times 10^{-21}$) in SLC39A8 was the lead SNP in locus 71 and the ancestral allele C was associated with higher scores on intelligence measures. The effect sizes for ExNS were individually small, with each effect allele accounting for a difference of 0.01 to 0.08 standard deviations. Supplementary Tables 6 and 11 and Supplementary Results 2.3.5 present a detailed catalog of variants in the associated genomic loci.

To link the associated variants to genes, we applied three gene-mapping strategies implemented in FUMA$^{24}$. Positional gene-mapping aligned SNPs to 522 genes by genomic location, eQTL (expression quantitative trait loci) gene-mapping matched cis-eQTL SNPs to 684 genes whose expression levels they influence, and chromatin interaction mapping annotated SNPs to 227 genes based on three-dimensional DNA-DNA interactions (Figure 2; Supplementary Results 2.3.6; Supplementary Figures 8–9; Supplementary Tables 12–14). This resulted in 859 unique mapped genes, 435 of which were implicated by at least two mapping strategies and 139 by all three (Figure 3). Although not all of these genes are certain to have a role in intelligence, they point to potential functional links for the GWAS associated variants and give higher credibility to genes with convergent evidence of association from multiple sources. The FUMA-mapped genes were enriched for brain tissue expression and several regulatory biological gene-sets (Supplementary Results 2.3.6). Fifteen genes are particularly notable as they are implicated via chromatin interactions between two independent genomic risk loci (Figure 2; Supplementary Results 2.3.6). Cross-locus interactions implicated ELAVL2, PTCH1, ATF4, FBXL17, and MAN2A1 in left ventricle of the heart tissue, SATB2 in liver tissue, and MEF2C in 5 tissues. Multiple interactions in multiple tissue types were seen for a cluster of 8 genes on chromosome 6 encoding zinc finger proteins and histones.

We performed genome-wide gene-based association analysis (GWGAS) using MAGMA$^{25}$ to estimate aggregate associations based on all SNPs in a gene (whereas FUMA annotates individually significant SNPs to genes). GWGAS identified 507 associated genes (Figure 3a; Supplementary Results 2.4.1; Supplementary Table 15), of which 350 were also mapped by FUMA (Figure 3b). In total, 105 genes were implicated by all four strategies (Supplementary Table 16).

In gene-set analysis, six Gene Ontology$^{26}$ gene-sets were significantly associated with intelligence: neurogenesis ($P=4.78 \times 10^{-7}$), neuron differentiation ($P=4.82 \times 10^{-6}$), central nervous system neuron differentiation ($P=3.31 \times 10^{-6}$), regulation of nervous system development ($P=9.30 \times 10^{-7}$), positive regulation of nervous system development ($P=1.00 \times 10^{-6}$), and regulation of synapse structure or activity ($P=5.42 \times 10^{-6}$) (Supplementary Results 2.4.2; Supplementary Tables 17–18). Conditional analysis indicated that there were three independent associations, regulation of nervous system development,
central nervous system neuron differentiation, and regulation of synapse structure or activity, which together accounted for the associations of the other sets.

Linking gene-based P-values to tissue-specific gene-sets, we observed strong associations with gene expression across multiple brain areas (Figure 3c; Supplementary Results 2.4.2; Supplementary Table 19), particularly the frontal cortex ($P=3.10 \times 10^{-9}$). In brain single-cell expression gene-set analyses, we found significant associations of striatal medium spiny neurons ($P=2.02 \times 10^{-14}$) and pyramidal neurons in the CA1 hippocampal ($P=5.67 \times 10^{-11}$) and cortical somatosensory regions ($P=2.72 \times 10^{-9}$) (Figure 3d; Supplementary Results 2.4.2; Supplementary Table 20). Conditional analysis showed that the independent association signal in brain cells was driven by medium spiny neurons, neuroblasts, and pyramidal CA1 neurons.

Intelligence has been associated with a wide variety of human behaviors$^{15}$ and brain anatomy$^{27}$. Confirming previous reports$^{5,6}$, we observed negative genetic correlations with ADHD ($r_g=-0.36$, $P=4.58 \times 10^{-23}$), depressive symptoms ($r_g=-0.27$, $P=6.20 \times 10^{-10}$), Alzheimer’s disease ($r_g=-0.27$, $P=2.03 \times 10^{-5}$), and schizophrenia ($r_g=-0.21$, $P=3.82 \times 10^{-17}$), and positive correlations with longevity ($r_g=0.43$, $P=7.96 \times 10^{-8}$) and autism ($r_g=0.25$, $P=3.14 \times 10^{-7}$), among others (Supplementary Table 21; Supplementary Figure 10). Comparison with previous GWAS$^{28}$ supported these correlations, showing numerous shared genetic variants across phenotypes (Supplementary Results 2.5; Supplementary Tables 22–23). Low enrichment (87 of 1,518 genes, $P=0.05$) was found for genes previously linked to intellectual disability or developmental delay, indicating largely distinct biological processes. However, our results extend previous genetic research on normal variation in general intelligence, as catalogued in Supplementary Tables 24–25.

We used Generalized Summary-statistic-based Mendelian Randomization$^{29}$ to test for potential credible causal associations between intelligence and genetically correlated traits (Supplementary Results 2.5.3; Supplementary Figures 11–12; Supplementary Table 26). We observed a strong bidirectional effect of cognitive ability on educational attainment ($b_{xy}=0.549$, $P=6.85 \times 10^{-82}$) and of educational attainment on intelligence ($b_{yx}=0.480$, $P=6.19 \times 10^{-52}$). Such findings are consistent with previous studies implicating bidirectional causal effects$^{30,31}$. There was also a bidirectional association showing a strong protective effect of intelligence on schizophrenia (OR=0.50, $b_{xy}=−0.685$, $P=2.02 \times 10^{-57}$) and a relatively smaller reverse effect ($b_{yx}=−0.214$, $P=4.19 \times 10^{-52}$), with additional evidence for pleiotropy (Supplementary Results 2.5.3). A number of previous reports support both a causal link and genetic overlap between these phenotypes$^{32,33}$. Our results also suggested that higher intelligence had a protective effect on ADHD (OR=0.48, $b_{xy}=−0.734$, $P=2.57 \times 10^{-46}$) and Alzheimer’s disease (OR=0.65, $b_{xy}=-0.435$, $P=3.59 \times 10^{-14}$), but was associated with higher risk of autism (OR=1.38, $b_{xy}=0.321$, $P=1.12 \times 10^{-3}$).

In the present study, we have affirmed and expanded existing knowledge of the genetics of general intelligence, identifying 190 novel loci and 939 novel associated genes and replications previous associations with 15 loci and 77 genes. The combined strategies of functional annotation and gene-mapping using biological data resources provide extensive information on the likely consequences of relevant genetic variants and put forward a rich
set of plausible gene targets and biological mechanisms for functional follow-up. Gene-set analyses contribute novel insight into underlying neurobiological pathways, confirming the importance of brain-expressed genes and neurodevelopmental processes in fluid domains of intelligence and pointing towards the involvement of specific cell types. Our results indicate overlap in the genetic processes involved in both cognitive functioning and neurological and psychiatric traits and provide suggestive evidence of causal associations that may drive these correlations. These results are important for understanding the biological underpinnings of cognitive functioning and contribute to our understanding of related neurological and psychiatric disorders.

Online Methods

Study Cohorts

The meta-analysis included new and previously reported GWAS summary statistics from 14 cohorts: UK Biobank (UKB), Cognitive Genomics Consortium (COGENT), Rotterdam Study (RS), Generation R Study (GENR), Swedish Twin Registry (STR), Spit for Science (S4S), High-IQ/Health and Retirement Study (HiQ/HRS), Twins Early Development Study (TEDS), Danish Twin Registry (DTR), IMAGEN, Brisbane Longitudinal Twin Study (BLTS), Netherlands Study of Cognition, Environment and Genes (NESCOG), Genes for Good (GfG), and the Swedish Twin Studies of Aging (STSA). All samples were obtained from epidemiological cohorts ascertained for research on a variety of physical and psychological outcomes. Participants ranged from children to older adults, with older samples being screened for cognitive decline to exclude the possibility of dementia affecting performance on cognitive tests.

Different measures of intelligence were assessed in each cohort but were all operationalized to index a common latent g factor underlying multiple dimensions of cognitive functioning. With the exception of HiQ/HRS, all cohorts extracted a single sum score, mean score, or factor score from a multidimensional set of cognitive performance tests and used this normally-distributed score as the phenotype in a covariate-adjusted (e.g. age, sex, ancestry principal components) GWAS using linear regression methods. For HiQ/HRS, a logistic regression GWAS was run with “case” status reflecting whether participants were drawn from an extreme-sampled population of very high intelligence (i.e. at the upper ~0.03% of the tail of the normal distribution) versus an epidemiological sample of unselected population “controls”. Detailed descriptions of the samples, measures, genotyping, quality control, and analysis procedures for each cohort are provided in the Supplementary Note (Supplementary Information 1.1–1.2), Supplementary Table 1, and in the Life Sciences Reporting Summary.

Meta-analysis

Stringent quality control measures were applied to the summary statistics for each GWAS cohort before combining. All files were checked for data integrity and accuracy. SNPs were filtered from further analysis if they met any of the following criteria: imputation quality (INFO/R^2) score < 0.6, Hardy-Weinberg equilibrium (HWE) P < 5×10^{-6}, study-specific minor allele frequency (MAF) corresponding to a minor allele count (MAC) < 100, and
mismatch of alleles or allele frequency difference greater than 20% from the Haplotype Reference Consortium (HRC) genome reference panel\textsuperscript{16}. Some cohorts used more stringent criteria (see Supplementary Information 1.1). Indels and SNPs that were duplicated, multi-allelic, monomorphic, or ambiguous (A/T or C/G with a MAF >0.4) were also excluded. Visual inspection of the distribution of the summary statistics was completed, and Manhattan plots and QQ plots were created for the cleaned summary statistics from each cohort (Supplementary Figure 1).

The SNP association \textit{P}-values from the GWAS cohorts were meta-analyzed with METAL\textsuperscript{34} (see URLs) in two phases. First, we meta-analyzed all cohorts with quantitative phenotypes (all except HiQ/HRS) using a sample-size weighted scheme. In the second phase, we added the HiQ/HRS study results to the first phase results, weighting each set of summary statistics by their respective non-centrality parameter (NCP). This method improves power when using an extreme case sampling design such as HiQ\textsuperscript{35} and provides a comparable metric with which to combine information from different analytic designs while accounting for their differences in power/effective sample size. NCPs were estimated using the Genetic Power Calculator\textsuperscript{36}, as described by Coleman et al.\textsuperscript{37}. After combining all data, meta-analysis results were further filtered to exclude any variants with N < 50,000. We additionally included a random-effects meta-analysis for each phase, as implemented in METAL, to evaluate potential heterogeneity in the SNP association statistics between cohorts.

The X chromosome was treated separately in the meta-analysis because imputed genotypes were not available for the X chromosome in the largest cohort (UKB), and there was little overlap between the UKB called genotypes and imputed data from other cohorts (N\textsubscript{SNPs} < 500). We therefore included only the called X chromosome variants in UKB for these analyses after performing X-specific quality control steps\textsuperscript{38}.

We conducted a series of meta-analyses on subsets of the full sample using the same methods as above. Age group-specific meta-analyses were run in the cohorts of children (age < 17; GENR, TEDS, IMAGEN, BLTS; N=9,814), young adults (age ~17–18; S4S, STR; N=6,033), adults (age > 18, primarily middle-aged or older: UKB, RS, DTR, NESCOG, STSA; N=204,228), and older adults (mean age > 60, RS, DTR, STSA; N=204,228).

URLs:

- UK Biobank website: http://ukbiobank.ac.uk
- UK Biobank genotyping: http://www.biorxiv.org/content/early/2017/07/20/166298
- Health and Retirement study: http://hrsonline.isr.umich.edu
- Genes for Good study: http://genesforgood.org
- International Cognitive Ability Resource measure (Genes for Good): https://icar-project.com/
- Functional Mapping and Annotation (FUMA) software: http://fuma.ctglab.nl
- Multi-marker Analysis of GenoMic Annotation (MAGMA) software: http://ctg.cncr.nl/software/magma
- METAL software: http://genome.sph.umich.edu/wiki/METAL_Program
- LD Score Regression software: https://github.com/bulik/ldsc
- LD Hub (GWAS summary statistics): http://ldsc.broadinstitute.org/
- LD scores: https://data.broadinstitute.org/alkesgroup/LDSCORE/
- GeneCards: http://www.genecards.org
- MSigDB curated gene-set database: http://software.broadinstitute.org/gsea/msigdb/collections.jsp
- NHGRI GWAS catalog: https://www.ebi.ac.uk/gwas/
- RegionAnnotator: https://github.com/ivankosmos/RegionAnnotator
N=8,323), excluding studies whose samples overlapped child/young adult and adult groups (COGENT, HiQ/HRS, GfG; N=49,792). To create independent discovery samples for use in polygenic score validation, we also conducted meta-analyses with a “leave-one-out” strategy in which summary statistics from four validation datasets were, respectively, excluded from the meta-analysis (see Polygenic Scoring, below).

Cohort Heritability and Genetic Correlation

LD score regression\(^{17}\) was used to estimate genomic inflation and heritability of the intelligence phenotypes in each of the 14 cohorts using their post-quality control summary statistics, and to estimate the cross-cohort genetic correlations\(^{39}\). Pre-calculated LD scores from the 1000 Genomes European reference population were obtained online (see URLs). Genetic correlations were calculated on HapMap3 SNPs only. LD score regression was also used on the age subgroup meta-analyses to estimate heritability and cross-age genetic correlations.

Genomic Risk Loci Definition

Independently associated loci from the meta-analysis were defined using FUMA\(^{24}\) (see URLs), an online platform for functional mapping of genetic variants. We first identified independent significant SNPs which had a Bonferroni-corrected genome-wide significant two-tailed \(P\)-value \((P<5 \times 10^{-8})\) and represented signals that were independent from each other at \(r^2<0.6\). These SNPs were further represented by lead SNPs, which are a subset of the independent significant SNPs that are in approximate linkage equilibrium with each other at \(r^2<0.1\). We then defined associated genomic loci by merging any physically overlapping lead SNPs (linkage disequilibrium [LD] blocks <250kb apart). Borders of the associated genomic loci were defined by identifying all SNPs in LD \((r^2 \geq 0.6)\) with one of the independent significant SNPs in the locus, and the region containing all of these candidate SNPs was considered to be a single independent genomic locus. All LD information was calculated from UK Biobank genotype data.

Proxy-replication with Educational Attainment

We conducted GWAS of educational attainment, an outcome with a high genetic correlation with intelligence\(^{5}\), in a non-overlapping European subset of the UKB sample (N=188,435) who did not complete the intelligence measure. Educational attainment was coded as maximum years of education completed, using the same methods as earlier analyses\(^{40}\) and GWAS was conducted using the same quality control and analytic procedures as described for the UKB intelligence phenotype (Supplementary Information 1.1.1). To test replication of the SNPs with this proxy phenotype, we performed a sign concordance test for all GWS SNPs from the meta-analysis using the two-tailed exact binomial test. For each independent genomic locus, we considered it to be evidence for replication if the lead SNP or another correlated SNP in the region was sign concordant with the corresponding SNP in the intelligence meta-analysis and had a two-tailed \(P\)-value of association with educational attainment smaller than 0.05/242 independent tests=0.0002.
Polygenic Scoring

We calculated polygenic scores (PGS) based on the SNP effect sizes of the leave-one-out meta-analyses, from which four cohorts were (separately) excluded and reserved for score validation. These included a child (GENR), young adult (S4S), and adult sample (RS). We also included the UKB-wb sample to test for validation in a very large (N = 53,576) cohort with the greatest phenotypic similarity to the largest contributor to the meta-analysis statistics (UKB-ts), in order to maximize potential predictive power. PGS were calculated on the genotype data using LDpred\textsuperscript{21}, a Bayesian PGS method that utilizes a prior on effect size distribution to remodel the SNP effect size and account for LD, and PRSice\textsuperscript{20}, a PLINK\textsuperscript{41}-based program that automates optimization of the set of SNPs included in the PGS based on a high-resolution filtering of the GWAS $p$-value threshold. LDpred PGS were applied to the called, cleaned, genotyped variants in each of the validation cohorts with UK Biobank as the LD reference panel. PRSice PGS were calculated on hard-called imputed genotypes using $P$-value thresholds from 0.0 to 0.5 in steps of 0.001. The explained variance ($\Delta R^2$) was derived from a linear model in which the GWAS intelligence phenotype was regressed on each PGS while controlling for the same covariates as in each cohort-specific GWAS, compared to a linear model with GWAS covariates only.

Stratified Heritability

We partitioned SNP heritability using stratified LD Score regression\textsuperscript{42} in three ways: 1) by functional annotation category, 2) by minor allele frequency (MAF) in six percentile bins, and 3) by chromosome. Annotations for 28 binary categories of putative functional genomic characteristics (e.g. coding or regulatory regions) were obtained from the LD score website (see URLs). With this method, enrichment/depletion of heritability in each category is calculated as the proportion of heritability attributable to SNPs in the specified category divided by the proportion of total SNPs annotated to that category. The Bonferroni-corrected significance threshold was .05/56 annotations=.0009.

Functional Annotation of SNPs

Functional annotation of SNPs implicated in the meta-analysis was performed using FUMA\textsuperscript{24} (see URLs). We selected all candidate SNPs in associated genomic loci having an $r^2 \geq 0.6$ with one of the independent significant SNPs (see above), a suggestive $P$-value ($P<1e-5$) and a MAF $> 0.0001$ for annotations. Predicted functional consequences for these SNPs were obtained by matching SNPs’ chromosome, base-pair position, and reference and alternate alleles to databases containing known functional annotations, including ANNOVAR\textsuperscript{43} categories, Combined Annotation Dependent Depletion (CADD) scores\textsuperscript{23}, RegulomeDB\textsuperscript{44} (RDB) scores, and chromatin states\textsuperscript{45,46}. ANNOVAR categories identify the SNP’s genic position (e.g. intron, exon, intergenic) and associated function. CADD scores predict how deleterious the effect of a SNP is likely to be for a protein structure/function, with higher scores referring to higher deleteriousness. A CADD score above 12.37 is the threshold to be potentially pathogenic\textsuperscript{23}. The RegulomeDB score is a categorical score based on information from expression quantitative trait loci (eQTLs) and chromatin marks, ranging from 1a to 7 with lower scores indicating an increased likelihood of having a regulatory function. Scores are as follows: 1a=eQTL + Transcription Factor (TF) binding + matched TF
motif + matched DNase Footprint + DNase peak; 1b=eQTL + TF binding + any motif + DNase Footprint + DNase peak; 1c=eQTL + TF binding + matched TF motif + DNase peak; 1d=eQTL + TF binding + any motif + DNase peak; 1e=eQTL + TF binding + matched TF motif; 1f=eQTL + TF binding / DNase peak; 2a=TF binding + matched TF motif + matched DNase Footprint + DNase peak; 2b=TF binding + any motif + DNase Footprint + DNase peak; 2c=TF binding + matched TF motif + DNase peak; 3a=TF binding + any motif + DNase peak; 3b=TF binding + matched TF motif; 4=TF binding + DNase peak; 5=TF binding or DNase peak; 6=other; 7=Not available. The chromatin state represents the accessibility of genomic regions (every 200bp) with 15 categorical states predicted by a hidden Markov model based on 5 chromatin marks for 127 epigenomes in the Roadmap Epigenomics Project. A lower state indicates higher accessibility, with states 1–7 referring to open chromatin states. We annotated the minimum chromatin state across tissues to SNPs. The 15-core chromatin states as suggested by Roadmap are as follows: 1=Active Transcription Start Site (TSS); 2=Flanking Active TSS; 3=Transcription at gene 5' and 3'; 4=Strong transcription; 5=Weak Transcription; 6=Genic enhancers; 7=Enhancers; 8=Zinc finger genes & repeats; 9=Heterochromatic; 10=Bivalent/Poised TSS; 11=Flanking Bivalent/Poised TSS/Enhancer; 12=Bivalent Enhancer; 13=Repressed PolyComb; 14=Weak Repressed PolyComb; 15=Quiescent/Low. Standardized SNP effect sizes were calculated for the most impactful SNPs by transforming the sample size-weighted meta-analysis Z score, as described by Zhu et al.47.

**Gene-mapping**

Genome-wide significant loci obtained by the GWAS meta-analysis were mapped to genes in FUMA using three strategies:

1. **Positional mapping** maps SNPs to genes based on physical distance (within a 10kb window) from known protein coding genes in the human reference assembly (GRCh37/hg19).

2. **eQTL mapping** maps SNPs to genes with which they show a significant eQTL association (i.e. allelic variation at the SNP is associated with the expression level of that gene). eQTL mapping uses information from 45 tissue types in 3 data repositories (GTEx, Blood eQTL browser, BIOS QTL browser), and is based on cis-eQTLs which can map SNPs to genes up to 1Mb apart. We used a false discovery rate (FDR) of 0.05 to define significant eQTL associations.

3. **Chromatin interaction mapping** was performed to map SNPs to genes when there is a three-dimensional DNA-DNA interaction between the SNP region and a gene region. Chromatin interaction mapping can involve long-range interactions as it does not have a distance boundary. FUMA currently contains Hi-C data of 14 tissue types from the study of Schmitt et al. Since chromatin interactions are often defined in a certain resolution, such as 40kb, an interacting region can span multiple genes. If a SNP is located in a region that interacts with a region containing multiple genes, it will be mapped to each of those genes. To further prioritize candidate genes, we selected only interaction-mapped genes in which one region involved in the interaction overlaps with a predicted enhancer region in any of the 111 tissue/cell types from the Roadmap Epigenomics Project and
the other region is located in a gene promoter region (250bp up and 500bp downstream of the transcription start site and also predicted by Roadmap to be a promoter region). This reduces the number of genes mapped but increases the likelihood that those identified will have a plausible biological function. We used a FDR of $1 \times 10^{-5}$ to define significant interactions, based on previous recommendations\textsuperscript{51} modified to account for the differences in cell lines used here.

**Functional annotation of mapped genes**

Genes implicated by mapping of significant GWAS SNPs were further investigated using the GENE2FUNC procedure in FUMA\textsuperscript{24}, which provides hypergeometric tests of enrichment of the list of mapped genes in 53 GTEx\textsuperscript{48} tissue-specific gene expression sets, 7,246 MSigDB gene-sets\textsuperscript{52}, and 2,195 GWAS catalog gene-sets\textsuperscript{28}. The Bonferroni-corrected significance threshold was $0.05/9,494$ gene-sets=$5.27 \times 10^{-6}$.

**Gene-based analysis**

SNP-based $P$-values from the meta-analysis were used as input for the gene-based genome-wide association analysis (GWGAS). 18,128 protein-coding genes (each containing at least 1 GWAS SNP) from the NCBI 37.3 gene definitions were used as basis for GWGAS in MAGMA\textsuperscript{25} (see URLs). The Bonferroni-corrected genome-wide significance threshold was $.05/18,128$ genes=$2.76 \times 10^{-6}$.

**Gene-set analysis**

Results from the GWGAS analyses were used to test for association in three types of predefined gene-sets:

1. 7,246 curated gene-sets representing known biological and metabolic pathways were derived from 9 data resources, catalogued by and obtained from the MsigDB version 5.2\textsuperscript{29} (see URLs)

2. gene expression values from 53 tissues obtained from GTEx\textsuperscript{48}, log2 transformed with pseudocount 1 after winsorization at 50 and averaged per tissue

3. cell-type specific gene expression in 24 types of brain cells, which were calculated following the method described in Skene et al.\textsuperscript{53} and Coleman et al.\textsuperscript{37} Briefly, brain cell-type expression data was drawn from single-cell RNA sequencing data from mouse brains. For each gene, the value for each cell-type was calculated by dividing the mean Unique Molecular Identifier (UMI) counts for the given cell type by the summed mean UMI counts across all cell types. Single-cell gene-sets were derived by grouping genes into 40 equal bins by specificity of expression.

These gene-sets were tested for association with the GWGAS gene-based test statistics using MAGMA. We computed competitive $P$-values, which represent the test of association for a specific gene-set compared to other gene-sets. This method is more robust to Type I error than self-contained tests that only test for association of a gene-set against the null hypothesis of no association\textsuperscript{25}. The Bonferroni-corrected significance threshold was...
0.05/7,323 gene-sets=6.83×10^{-6}. Conditional analyses were performed as a follow-up using MAGMA to test whether each significant association observed was independent of all others. The association between each gene-set was tested conditional on the most strongly associated set, and then - if any substantial (p<.05/number of gene-sets) associations remained - by conditioning on the first and second most strongly associated set, and so on until no associations remained. Gene-sets that retained their association after correcting for other sets were considered to be independent signals. We note that this is not a test of association per se, but rather a strategy to identify, among gene-sets with known significant associations whose defining genes may overlap, which set(s) are responsible for driving the observed association.

**Cross-Trait Genetic Correlation**

Genetic correlations \( r_g \) between intelligence and 38 phenotypes were computed using LD score regression\(^{39} \), as described above, based on GWAS summary statistics obtained from publicly available databases (see URLs; Supplementary Table 18). The Bonferroni-corrected significance threshold was 0.05/38 traits=1.32×10^{-3}.

**GWAS catalog lookup**

We used FUMA to identify SNPs with previously reported \( (P < 5×10^{-5}) \) phenotypic associations in published GWAS listed in the NHGRI-EBI catalog\(^{28} \) which overlapped with the genomic risk loci identified in the meta-analysis. As an additional relevant phenotype of interest, we examined whether the genes associated with intelligence in this study (by FUMA mapping or GWGAS) were overrepresented in a set of 1,518 genes linked to intellectual disability and/or developmental delay, as compiled by RegionAnnotater (see URLs). Many of these have been identified by non-GWAS sources and are not represented in the NHGRI catalog. We tested for enrichment using a hypergeometric test with a background set of 19,283 genomic protein-coding genes, as in FUMA. Manual lookups were also performed to identify overlapping loci_genes with known previous GWAS of intelligence.

**Mendelian Randomization**

To infer credible causal associations between intelligence and traits that are genetically correlated with intelligence, we performed Generalized Summary-data based Mendelian Randomization\(^{29} \) (GSMR; see URLs). This method utilizes summary-level data to test for causal associations between a putative risk factor (exposure) and an outcome by using independent genome-wide significant SNPs as instrumental variables. HEIDI-outlier detection was used to filter genetic instruments that show clear pleiotropic effects on both the exposure phenotype and the outcome phenotype. We used a threshold p-value of 0.01 for the outlier detection analysis in HEIDI which removes 1% of SNPs by chance if there is no pleiotropic effect. To test for a potential causal effect of intelligence on various outcomes, we selected traits in non-overlapping samples that showed significant genetic correlations \( r_g \) with intelligence. We tested for bi-directional causation by repeating the analyses while switching the role of each correlated phenotype as an exposure and intelligence as the outcome. For each trait, we selected independent \( (r^2<=0.1) \), GWS lead SNPs as instrumental variables in the analyses. For traits with less than 10 GWS lead SNPs (i.e. the minimum
number of SNPs on which GSMR can perform a reliable analysis), the GWS threshold was lowered to 1×10^{-5}, allowing a sufficient number of SNPs to conduct the reverse GSMR analysis for former smoker status, autism, and intracranial volume.

The method estimates a putative causal effect of the exposure on the outcome \( (b_{xy}) \) as a function of the relationship between the SNPs' effects on the exposure \( (b_{zx}) \) and the SNPs' effects on the outcome \( (b_{zy}) \), given the assumption that the effect of non-pleiotropic SNPs on an exposure (\( x \)) should be related to their effect on the outcome (\( y \)) in an independent sample only via mediation through the phenotypic causal pathway \( (b_{xy}) \). The estimated causal effect coefficients \( (b_{xy}) \) are approximately equal to the natural log odds ratio (OR) for a case-control trait\(^{29}\). An OR of 2 can be interpreted as a doubled risk compared to the population prevalence of a binary trait for every SD increase in the exposure trait. For quantitative traits the \( b_{xy} \) can be interpreted as a one standard deviation increase explained in the outcome trait for every SD increase in the exposure trait. This method can help differentiate the likely causal direction of association between two traits but cannot make any statement about the intermediate mechanisms involved in any potential causal process.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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Methods-Only References


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Figure 1. SNP-based associations with intelligence in the GWAS meta-analysis of N=269,867 independent individuals.

(a) Manhattan plot showing the $-\log_{10}$ transformed two-tailed $P$-value of each SNP from the GWAS meta-analysis (of linear and logistic regression statistics) on the y-axis and base pair positions along the chromosomes on the x-axis. The dotted red line indicates Bonferroni-corrected genome-wide significance ($P<5\times10^{-8}$); the blue line the threshold for suggestive associations ($P<1\times10^{-5}$). Independent lead SNPs are indicated by a diamond.

(b) Heritability enrichment of 28 functional annotation categories for SNPs in the meta-analysis.
calculated with stratified LD score regression. Error bars show 95% confidence intervals around the enrichment estimate. The dashed horizontal line indicates no enrichment of the annotation category. Red dots indicate significant Bonferroni-corrected two-tailed P-values and beige dots indicate suggestive (P<.05) values. UTR=untranslated region; TSS=transcription start site; CTCF=CCCTC-binding factor; DHS=DNaseI Hypersensitive Site; TFBS=transcription factor binding site; DGF=DNaseI digital genomic footprint. (c) Distribution of functional consequences of SNPs in genomic risk loci in the meta-analysis. (d) Distribution of RegulomeDB score for SNPs in genomic risk loci, with a low score indicating a higher likelihood of having a regulatory function (Online methods). (e) The minimum chromatin state across 127 tissue and cell types for SNPs in genomic risk loci, with lower states indicating higher accessibility and states 1–7 referring to open chromatin states (Online Methods).
Figure 2. Cross-locus interactions for genomic regions associated with intelligence in 269,867 independent individuals.

Circos plots showing genes on chromosomes 2 (a), 5 (b), 6 (c), 9 (d), and 22 (e) that were linked to genomic risk loci in the GWAS meta-analysis (blue regions) by eQTL mapping (green lines connecting an eQTL SNP to its associated gene), and/or chromatin interactions (orange lines connecting two interacting regions) and showed evidence of interaction across two independent genomic risk loci. Genes implicated by eQTL are in green, by chromatin interactions in orange, and by both eQTL and chromatin interactions mapping in red. The
The outer layer shows a Manhattan plot containing the −log10 transformed two-tailed $P$-value of each SNP from the GWAS meta-analysis (of linear and logistic regression statistics), with genome-wide significant SNPs colored according to linkage disequilibrium patterns with the lead SNP. Circos plots for all chromosomes are provided in Supplementary Fig. 8.
Figure 3. Implicated genes, pathways, and tissue- and cell-expression profiles for intelligence in 269,867 independent individuals.

(a) Manhattan plot of the genome-wide gene-based association analysis (GWGAS). The y-axis shows the $-\log_{10}$ transformed two-tailed $P$-value of each gene from a linear model, and the chromosomal position on the x-axis. The red dotted line indicates the Bonferroni-corrected threshold for genome-wide significance of the gene-based test ($P<2.76\times10^{-6}$, 0.05/18,128 genes), and the blue line indicates the suggestive threshold ($P<2.76\times10^{-5}$, 0.5/18,128 genes).

(b) Venn diagram showing overlap of genes implicated by positional mapping, eQTL mapping, chromatin interaction mapping, and GWGAS.

(c) Gene expression profiles of associated genes in 53 tissue types. The y-axis shows the $-\log_{10}$ transformed two-tailed $P$-value of association of GWGAS test statistics with tissue-specific gene expression levels in a linear model. Expression data were extracted from the Genotype-Tissue Expression (GTEx) database. Expression values (RPKM) were log2 transformed with pseudocount 1 after winsorization at 50 and averaged per tissue. The dotted blue line indicates the Bonferroni-corrected significance threshold ($P=0.05/7,323$ gene-sets=$6.83\times10^{-6}$).

(d) Single-cell gene-expression analysis of genes related to intelligence in 24 cell-types. The x-axis shows the $-\log_{10}$ transformed two-tailed $P$-value of association of GWGAS test statistics with cell-specific gene expression levels in a linear model. The dotted blue line indicates the Bonferroni-corrected significance threshold ($P=0.05/7,323$ gene-sets=$6.83\times10^{-6}$).
### Table 1.
Overview of cohorts included in a GWAS meta-analysis of general intelligence.

<table>
<thead>
<tr>
<th>Cohort</th>
<th>N</th>
<th>Age</th>
<th>Phenotype</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. UKB</td>
<td>195,653</td>
<td>39–72</td>
<td>Verbal and mathematical reasoning</td>
</tr>
<tr>
<td>2. COGENT</td>
<td>35,289</td>
<td>8–96</td>
<td>One or more neuropsychological tests from three or more domains of cognitive performance</td>
</tr>
<tr>
<td>3. RS</td>
<td>6,182</td>
<td>45–98</td>
<td>Letter-digit substitution, Stroop, verbal fluency, delayed recall</td>
</tr>
<tr>
<td>4. GENR</td>
<td>1,929</td>
<td>5–9</td>
<td>SON-R (spatial visualization and abstract reasoning subsets)</td>
</tr>
<tr>
<td>5. STR</td>
<td>3,215</td>
<td>18</td>
<td>Logical, verbal, spatial, and technical ability subtests</td>
</tr>
<tr>
<td>6. S4S</td>
<td>2,818</td>
<td>17–18</td>
<td>SAT test scores</td>
</tr>
<tr>
<td>7. HiQ / HRS</td>
<td>9,410</td>
<td>*</td>
<td>High IQ cases / unselected population controls</td>
</tr>
<tr>
<td>8. TEDS</td>
<td>3,414</td>
<td>12</td>
<td>WISC-III verbal and nonverbal reasoning; Raven's progressive matrices</td>
</tr>
<tr>
<td>9a. DTR - MADT</td>
<td>737</td>
<td>55–80</td>
<td>Verbal fluency, digit span, immediate and delayed recall tests</td>
</tr>
<tr>
<td>9b. DTR - LSADT</td>
<td>253</td>
<td>73–94</td>
<td>Verbal fluency, digit span, immediate and delayed recall tests</td>
</tr>
<tr>
<td>10. IMAGEN</td>
<td>1,343</td>
<td>14</td>
<td>WISC-IV, CANTAB factor score</td>
</tr>
<tr>
<td>11a. BLTS - Children</td>
<td>530</td>
<td>12–13</td>
<td>VSRT-C factor score</td>
</tr>
<tr>
<td>11b. BLTS - Adolescents</td>
<td>2,598</td>
<td>15–30</td>
<td>MAB-II IQ score</td>
</tr>
<tr>
<td>12. NESCOG</td>
<td>252</td>
<td>18–79</td>
<td>WAIS IQ score</td>
</tr>
<tr>
<td>13. GfG</td>
<td>5,084</td>
<td>15–91</td>
<td>ICAR verbal reasoning test</td>
</tr>
<tr>
<td>14a. STSA - SATSA+GENDER</td>
<td>703</td>
<td>50–94</td>
<td>Verbal, spatial, episodic memory, and processing speed tests</td>
</tr>
<tr>
<td>14b. STSA - HARMONY</td>
<td>448</td>
<td>65–96</td>
<td>Verbal, spatial, episodic memory, and processing speed tests</td>
</tr>
</tbody>
</table>

* HiQ/HRS sample used a case-control design rather than a cognitive test score ascertained at a specific age; see Online Methods and Supplementary information 1.1.