Rare Variant Analysis of Human and Rodent Obesity Genes in Individuals with Severe Childhood Obesity

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Obesity is a genetically heterogeneous disorder. Using targeted and whole-exome sequencing, we studied 32 human and 87 rodent obesity genes in 2,548 severely obese children and 1,117 controls. We identified 52 variants contributing to obesity in 2% of cases including multiple novel variants in GNAS, which were sometimes found with accelerated growth rather than short stature as described previously. Nominally significant associations were found for rare functional variants in BBS1, BBS9, GNAS, MKKS, CLOCK and ANGPTL6. The p.S284X variant in ANGPTL6 drives the association signal (rs201622589, MAF=0.1%, odds ratio = 10.13, p-value = 0.042) and results in complete loss of secretion in cells. Further analysis including additional case-control studies and population controls (N = 260,642) did not support association of this variant with obesity (odds ratio = 2.34, p-value = 2.59 × 10⁻³), highlighting the challenges of testing rare variant associations and the need for very large sample sizes. Further validation in cohorts with severe obesity and engineering the variants in model organisms will be needed to explore whether human variants in ANGPTL6 and other genes that lead to obesity when deleted in mice, do contribute to obesity. Such studies may yield druggable targets for weight loss therapies.

Studies focused on severe early onset obesity alone, or obesity with developmental delay and/or dysmorphic features have identified a number of genes harbouring highly penetrant causal mutations1-3. The further characterisation of rare, highly penetrant variants identified in such individuals can provide insights into the cellular and

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physiological mechanisms involved in energy homeostasis and human obesity, and can identify and/or validate targets for therapeutic intervention. The aim of our study was to examine the prevalence of rare/novel variants in human and mouse obesity genes using high-throughput next-generation sequencing in a large cohort of individuals with severe early onset obesity. Here we describe results from an analysis of 119 candidate genes sequenced in 2,548 individuals with severe, early-onset obesity from the Severe Childhood Onset Obesity Project (SCOOP) (UK individuals of European ancestry recruited to the Genetics of Obesity Study, GOOS; BMI standard deviation score (SDS) > 3; onset of obesity before the age of 10 years; Methods) from the UK10K project. Our analysis includes 737 SCOOP individuals with whole-exome sequence, and 1,811 additional SCOOP individuals, not consented for whole-exome analysis, in whom we performed targeted sequencing. As this work was performed as part of a consortium (UK10K project), this data was compared to 1,117 individuals with other disorders (e.g. neurodevelopmental and rare disease) in whom exome sequencing and analysis were performed using the same methods. For the purposes of this analysis, these individuals are designated as “controls” (Methods), although we recognise that there are limitations with this design.

Study Design
SCOOP individuals likely to have congenital leptin deficiency, a treatable cause of severe obesity, were excluded by measurement of serum leptin, and individuals with mutations in the melanocortin 4 receptor gene (MC4R) (the most common genetic form of penetrant obesity) were excluded by prior Sanger sequencing.

We focused on six tiers of genes (Methods and Supplementary Table 1): (1) genes known to harbour variants causing human obesity alone (Obesity Alone; n = 6); (2) genes in which known variants cause human obesity combined with developmental delay and/or dysmorphology (Obesity and Delay; n = 26); (3) genes in which loss-of-function leads to obesity in mice (LoF Mice; n = 51); (4) genes in which gain-of-function leads to obesity in mice (GoF Mice; n = 5); (5) genes encoding anorectic molecules and their receptors (Anorectic Molecules; n = 7), and (6) genes in which loss-of-function is associated with other metabolic phenotypes in mice (Complex Metabolic Effects, n = 24). Our aim was to include genes where there was sufficient evidence to indicate disorders with Mendelian inheritance (obesity syndromes) or where complete deletion or overexpression (as opposed to conditional knockouts) causes an obesity phenotype in mice (Methods) (i.e. genes in which inherited loss/gain of function variants might exist). To identify variants more likely to be causally linked to obesity, we focused on rare (MAF < 1%) and novel (not seen in the data we used for filtering) variants predicted to be functional (i.e. nonsense variants, missense amino acid substitutions, alterations of conserved splice sites or small insertions/deletions (indels) that introduced a frameshift) (Methods). We filtered our data against approximately 8,000 publicly available sequenced samples, as well as 2,097 exomes (sequenced in parallel on the same platform as our samples) and 3,781 whole-genome sequenced samples also from the UK10K project (Methods).

Human Obesity Syndrome Genes.
Firstly, we sought to identify rare and novel functional variants in the 32 genes known to cause human obesity, with or without, additional developmental delay and/or dysmorphology features (Obesity Alone, or Obesity and Delay) (Methods). We identified 11 rare potentially functional variants in ClinVar with pathogenic/likely pathogenic status and 321 novel functional variants, which we confirmed by Sanger sequencing (Fig. 1, Methods). Based on inheritance patterns (where available) and the functional properties of variants that have previously been characterised, variants in these genes may contribute to obesity, sometimes in a non-fully penetrant manner, in 52 (2%) individuals (Supplementary Figure 1 & Supplementary Table 2; Fig. 2, Methods). As variants in MC4R account for approximately 5% of severe obesity in this cohort, these findings indicate that >90% of patients within this cohort do not have their phenotype explained by variants in known human obesity genes. Further analysis exploring the whole-exome in an agnostic manner, with appropriately matched non-disease controls not available in the UK10K project, will be an aim of future investigations.

Notably, we identified thirteen clinically associated GNAS variants in fourteen SCOOP individuals (12 of these confirmed on Sanger sequencing; 0.5%) (Supplementary Table 3). GNAS is an imprinted gene in which heterozygous loss-of-function variants are associated with obesity, short stature and skeletal abnormalities, and, when maternally inherited, hormone resistance syndromes. As GNAS sequencing has traditionally only been performed in individuals with classical clinical features, our findings suggest that the true prevalence in childhood obesity may be underappreciated. Three of the variants found in our study have been described previously in patients with classical features (p.Y163X; p.R258W, and p.R265H). We identified a novel nonsense variant (p.Y169X) predicted to remove the entire Ras-like GTPase domain and several missense variants predicted to affect downstream signalling by affecting the interaction with G-protein coupled receptors (GPCRs), G-protein β- and γ-subunits, or downstream adenylyl cyclase when mapped onto the protein structure of GNAS (Fig. 3). We confirmed maternal transmission of variants in three out of the five families where parental samples were available for genotyping. Although four GNAS variant carriers exhibited endocrinopathies and nine had developmental delay as anticipated, unexpectedly four individuals had accelerated linear growth in childhood (height SDS > 2) rather than short stature (defined as height SDS < 2.0) (Supplementary Table 3). Further molecular and physiological studies will be needed to investigate potential genotype-phenotype correlations. As studies in rodents have shown that Gnas is imprinted in the paraventricular nucleus of the hypothalamus, the location of the majority of neurons expressing the G-protein coupled receptor MC4R, it is plausible to hypothesize that some GNAS variants may contribute to obesity and accelerated linear growth by reducing melanocortin signalling.

Next we formally tested for enrichment of rare, or novel, functional variants in case-control analysis (Methods). Although no gene attained experiment-wide threshold of p-value < 2 × 10^{-4} (Methods and Supplementary Tables 4–9), four genes (BBS1, BBS9, GNAS, and MKKS) known to cause obesity and developmental delay and/or dysmorphology (Obesity and Delay) had nominally significant burden test p-values (p-value < 0.05) and a higher burden of variants in cases than controls (Table 1).
Mouse Obesity Genes. We found a higher burden of rare variants in cases than controls for two genes known to cause obesity when disrupted in mice - ANGPTL6 and CLOCK (Table 1). Testing the functional consequences of CLOCK variants in cells is not straightforward as the molecular mechanisms that lead to obesity are not fully understood. As such, we focused on verifying the accuracy of the computationally predicted function of all twelve rare ANGPTL6 variants (Methods, Fig. 4, Supplementary Table 10). All variants found in cases (but not the one variant found exclusively in controls, V143L) were predicted to affect the fibrinogen-like domain of the protein (Fig. 4a). ANGPTL6 is predominantly expressed in the liver but is also expressed in white adipose tissue and kidney. To evaluate the effect of these variants, HEK293 cells were transiently transfected with constructs encoding wild type and variant forms of ANGPTL6. Whereas wild-type ANGPTL6 was readily detected in the medium, several mutants reduced, and the S286X mutant abolished, secretion of ANGPTL6 (Fig. 4b). Incorporation of this functional information did not qualitatively change the results of the ANGPTL6 gene-based test (Supplementary Table 10). However, limiting the analysis to the single variant that completely abolished protein secretion (p.S286X, rs201622589) led to an adjusted OR = 10.13 and p-value = 0.028 (Fisher’s p-value = 0.041) suggesting that the association signal at this gene is primarily driven by this variant (Supplementary Table 10). To increase power, and given that there were no additional cases of European descent from our obesity cohort, we analysed existing exome-chip data from an additional 253,587 unrelated European, non-Finnish population controls (Methods, Supplementary Table 11). In an updated analysis with the 2,548 cases and 253,587 controls, we observed an odds ratio (OR) = 2.90 and a Fisher’s p-value = 0.0022, which did not reach experiment or exome-wide significance. Further analysis of 1,436 non-overlapping obesity cases and 1,954 non-overlapping controls from two studies did not lend additional support, and when combined with the original data, yielded an overall OR = 2.34 and chi-squared p-value = 0.0060 (total 3,984 cases vs 256,658 controls, Supplementary Table 11). The results are similar when limiting to non-obese controls (Supplementary Table 12). These findings highlight the difficulty in studying very rare variants in complex diseases, and potentially the effect of winner’s curse, reinforcing the need for extremely large sample sizes\[^{12}\]. Of interest, mice with targeted deletion of Angptl6 that survive to birth (20%) develop marked obesity, have increased food intake, reduced energy expenditure, exhibit lipid accumulation in liver and muscle, and develop insulin resistance\[^{13}\]. Conversely, mice with targeted overexpression of Angptl6 are lean, insulin sensitive and are protected from diet-induced obesity\[^{13}\]. As ANGPTL6 is a liver-derived circulating peptide, and thus could potentially be manipulated for therapeutic purposes\[^{14}\], further genetic studies in larger cohorts and experimental studies in mice and humans are necessary to explore its role in obesity and potential utility as an anti-obesity drug target.

Power and Gene Set Analysis. Assuming an experiment-wide significance level of $2 \times 10^{-4}$ and that 30% of rare (MAF < 1%) variants within a given gene are causal we have very limited power (~20%) to detect an association to a gene region using our sample of 2,548 cases and 1,117 controls (Supplementary Figure 2, Online Methods, Supplementary Note 4)\[^{12}\]. As there is evidence that looking across group of genes with stronger priors may increase power\[^{15,17}\], we tested for association of rare or novel functional variants within each of the six
candidate gene sets. We further filtered variants with respect to their likelihood of being deleterious and performed six nested tests for each gene set (Methods). Although none of the analyses of tiered gene sets reached experiment-wide significance (Supplementary Table 13, Fig. 5), we see an increase in OR as we focus on rarer variants with more evidence of predicted deleterious effect within the Obesity Alone gene set. This suggests that focusing on extremely rare variants with strong evidence of being deleterious may be a good strategy for diseases with a complex genetic architecture, which likely includes rare, highly penetrant alleles. Our power analyses also suggests that restricting the MAF threshold to be near the very rare allele frequency of that expected for the causal variants, rather than a more lenient MAF threshold of 1%, increases power (Supplementary Figure 3). Finally, there are nominally significant signals (p-value < 0.05) within the LoF Mice candidate gene set for the novel, functional variant tests suggesting that within this rather large gene set (n = 51) there may be additional genes associated with human early onset obesity (Fig. 5).

**Discussion**

In summary, analysis of sequence data in 119 genes across 2,548 severely obese children and 1,117 other disease controls from the UK10K project identified variants in known obesity genes, contributing to obesity in 2% of children within this cohort, in whom mutations in MC4R and leptin had been excluded. It also highlighted that GNAS mutations may be more prevalent in human obesity than previously thought, and that the spectrum of phenotypic consequences may be broader and more heterogeneous than previously described (6). Additional detailed genotype-phenotype studies will be required to further elucidate the molecular underpinnings of the physiological consequences of these variants.

Taking the information generated in this study together with the published literature, we suggest that molecular genetic investigations, including testing known obesity genes, or agnostic exome-wide or genome-wide approaches (as these become more established at the point of care), should become part of the assessment for a child presenting with severe obesity (BMI SDS > 3) in the absence of other syndromic features. Genetic studies can allow the identification of congenital leptin deficiency which is entirely treatable with recombinant leptin therapy18. Also, early reports suggest that patients with mutations that disrupt POMC signalling may be effectively
treated with Setmelanotide (a MC4R agonist)\textsuperscript{19}. Finally, our data suggest that focusing on very rare, deleterious variants may increase power to find genetic association. Further detailed investigation of genes that lead to obesity when deleted in mice may identify new pathophysiological mechanisms involved in human obesity that can be targeted for drug discovery.

**Methods**

**Sample Sets.** All studies were approved by the Cambridge Local Research Ethics Committee and all participants and their parents gave written informed consent. All methods were performed in accordance with the relevant laboratory/clinical guidelines and regulations.

The SCOOP cohort\textsuperscript{3} was studied as part of the UK10K consortium\textsuperscript{4}. Data was compared to subsets within the neurodevelopmental and rare disease groups that were consented for use as controls. Details and further
Table 1. Summary of case-control results. Summary of case-control results in genes with nominally significant (p-value < 0.05, bold) burden of Rare (top) or Novel (bottom) functional variants in SCOOP cases (N = 1,811 with targeted sequence and N = 737 with whole-exome sequence), compared to 1,117 controls with whole-exome sequence data from the UK10K project. The number of variants (# variants) per gene are shown, as well as the number of alleles in cases and controls, odds ratios (OR) and p-values from SKAT-O (p-value SKATO) and from burden tests (p-value BURDEN). For variants not detected in controls an adjusted odds ratio (Adjusted OR) was calculated by adding 0.5 to the number of alleles in each cell of the two-by-two table.

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<th>Gene</th>
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<th># Variants</th>
<th>Rare alleles in SCOOP TS (N=1,811)</th>
<th>Rare alleles in SCOOP WES (N=737)</th>
<th>Rare alleles in Control WES (N=1,117)</th>
<th>OR</th>
<th>Adjusted OR</th>
<th>p-value SKATO</th>
<th>p-value BURDEN</th>
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Novel Functional

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information about the UK10K project can be found at http://www.uk10k.org/ and in the UK10K consortium paper from 2015.4 We gathered the genotypes for the variant S284X (rs201622589) in 202,981 unrelated population controls with European non-Finnish ancestry from existing cohorts (UKHLS22, Fenland, EPIC Norfolk,23, CCHS24-26, CGPS24-26, CHOIDS24-26, EPIC-CVD37, UK Biobank28 and ExAC29) and in two additional studies of obese children and healthy or thin controls30,31, including one from the Leipzig Childhood Obesity Cohort32. (Supplementary Note 1)

Sequencing, Variant Calling, and Quality Control. Targeted Sequencing (TS) and Whole Exome Sequencing (WES) was performed as described elsewhere (refs 33 and 4 respectively). All UK10K WES samples from all arms of the exome study available at the time of the variant calling and SCOOP TS samples (N = 5,233 and 2,819 respectively) were called together on the non-redundant targets +/− 100bp using multi-sample calling. Using SAMtools34, a BCF file was created and the site genotype likelihoods were calculated. Variants (SNPs and Indels) were called using BCFTools. Indels were left-aligned using ‘vcf norm’ from the htshlib package. Variants were filtered for sequencing and genotype quality at the site and genotype level using vcf-annotate35. Additional details in Supplementary Note 2.

Variant Annotation. All variant annotation was applied using the GRCh37 human reference. Variants were annotated with rsIDs from dbSNP137, and allele frequencies from the final 1000Genomes Phase 1 integrated (v3) callset36, the NHLBI Exome Sequencing Project (ESP) v237, and the UK10K WGS sample set4. The Ensembl Variant Effect Predictor (http://www.ensembl.org/info/docs/variation/vep/index.html) v2.8 with Ensembl 66 was used to add variant consequence annotations including the predicted deleteriousness of each missense variant as predicted by SIFT38,40, PolyPhen41,42, and Condel43. For the 32 known human obesity genes, only the variant consequences on the most clinically relevant transcript were considered for further analysis. For the other 87 candidate obesity genes, the most severe consequence on any transcript was retained for each variant (Supplementary Table 1). Using Sequence Ontology terms44,45, variant consequences were defined as functional (i.e. essential splice site, stop gained, stop lost, complex indel, frameshift coding, non synonymous coding, within mature miRNA, partial codon).

To identify rare variants, we used the thirteen UK10K WES sample sets that were not obese sample sets and were not included as controls in this study (Supplementary Note 1). These sample sets were processed in parallel with our cases and controls and thus provide a similar sequencing coverage and depth across the called regions. We removed variants that had a MAF > 1% across all or a MAF > 10% in any of the thirteen sample sets. We then removed variants with a MAF > 1% in any of seven additional sample sets: 1. UK10K WGS sample set4, 2–3. European and African American NHLBI ESP v2 sample sets37, and 4–7, the four continent sample sets (AFR, AMR, ASN, EUR) from 1000 Genomes Phase 1 integrated v3 call set36, Novel variants were identified as sites not seen in any of the internal or external datasets used for MAF annotation, i.e. novel at the time of this study.

Sample quality control. Samples were identified as contaminated using a combination of two methods: VerifyBamID v1.046 and “fraction skewed hets”47. For the TS samples, the FREEMIX value was estimated using 11,250 high quality, autosomal, biallelic SNPs with an alternate AF ≥ 0.01, and a call rate ≥ 0.5 in both the TS sample set and the 1000Genomes Project Phase 1 v336. Of the 2,819 TS SCOOP samples, the 784 WES SCOOP samples, and the 1427 WES samples used as controls 37, one, and nine were excluded due to contamination.
respectively. Sixteen non-contaminated TS samples were excluded based on a mean sample read-depth 3 SD below the average for all samples (i.e. mean sample read-depth < 12.09). Genotype concordance for 436 WES SCOOP samples and 1,035 TS SCOOP samples for which we had both sequence and GWAS data was also calculated. Four WES samples and six TS samples with a concordance rate below 90% were identified and excluded. A set of highly polymorphic markers (MAF > 0.3) was genotyped and compared to the sequencing calls from each sample. Non-concordant WES samples were removed prior to variant calling and are not included in the original sequencing numbers. Eighty-one TS samples with low concordance were removed after variant calling. Three TS samples were removed due to having an extremely high genotype missing rate of > 50%.

To identify non-European samples, we calculated principal components (PCs) from the 1000Genomes Phase I integrated call set using either EIGENSTRAT v4.2 or LASER 2.0 for the WES and TS samples respectively (Supplementary Note 3). Of the 2,676 TS samples, 837 were classified as non-European due to genetic ancestry and three were excluded due to a reported non-European ancestry resulting in 1,836 good quality TS samples of European ancestry. Of the 779 WES SCOOP samples, 37 samples were classified as non-European due to genetic ancestry resulting in 742 good quality WES SCOOP samples of European ancestry. Of the 1,418 control samples, 146 were identified as being non-European leaving 1272 high quality WES controls.

Known and cryptic relatedness was identified by estimating pairwise identity by descent using PLINK v1.07. Genetic relationships within the WES sample set is described in detail elsewhere. We removed individuals sequentially by: (1) largest number of relationships, (2) diseased controls, non-diseased controls, cases, (3) lower mean depth of sequencing. Five SCOOP cases and 155 controls were removed resulting in 737 SCOOP WES cases and 1,117 WES controls all of high quality, unrelated, and European ancestry. A similar process was used to identify genetic relationships within the TS sample set and between the TS and WES sample sets (Supplementary Note 3). This resulted in 25 additional exclusions (18 TS cases related to WES cases and seven related within the TS sample set) for a total of 1,811 high quality, unrelated, TS samples of European ancestry.

Figure 4. Functional characterisation of ANGPTL6 variants location and functional effect (A) Human ANGPTL6 protein and location of the genetic variants. Schematic of the human ANGPTL6 protein with the three recognisable domains: signal peptide (SP) in green, coiled-coil domain (CCD) in purple, and fibrinogen-like domain (FLD) in yellow. As all genetic variants identified are located predominantly in the FLD, the partial tertiary structure of the C-terminal protein portion (amino acids 135–470), based on previously solved fibrinogen structure (1lwu) is presented. The single variant seen in controls only is shown in blue, variants identified in cases are in purple, the S284X variant is starred. (B) Functional characterisation of ANGPTL6 variants. Cells were transiently transfected with constructs encoding wild-type (WT) or mutant ANGPTL6; levels of protein secretion into the media were measured by ELISA; means ± standard deviation (SD) for experiments performed in triplicate are shown (results were confirmed by Western blotting; data not shown). Statistical significance was measured using unpaired T-test with Welch’s correction using the GraphPad Prism software. *p < 0.05; ***p < 0.001.
Candidate Gene Sets. We performed database searches using keywords ‘obesity’, ‘growth’, ‘size’, ‘adipose tissue’, (details below) and manually curated the results to arrive at a set of six tiers of candidate gene with definite or likely links to obesity. The Online Mendelian Inheritance of Man (OMIM database, http://www.omim.org/) accessed in February, 2013 was used to identify 32 genes (Supplementary Table 1) which directly lead to human obesity. These were further categorised into two groups based on consistent clinical features: genes characterised by “Obesity alone” (Obesity Alone; n = 6), and “Obesity and developmental delay and/or dysmorphology” (Obesity and Delay; n = 26). Any putative list of candidate genes has limitations and other genes that contribute to energy homeostasis/obesity in animal models deserve interrogation in the future.

Genes that cause obesity when disrupted in rodents were identified using the Mouse Genome Informatics Database (MGI, http://www.informatics.jax.org/) and the Rat Genome Database (RGD, http://rgd.mcw.edu) both accessed in February 2013, as well as published research (www.ncbi.nlm.gov/Pubmed). Eighty-seven genes were associated with a growth/size phenotype when perturbed in rodent animal models. We evaluated the strength of evidence for a role of each gene in obesity phenotype and divided them into four further categories, specifically: Loss of function associated with obesity in mice (LoF Mice; n = 51); gain of function associated with obesity in mice (GoF Mice, n = 5); anorectic peptides and receptors (Anorectic Molecules, n = 7); loss of function associated with other metabolic phenotypes in mice (Complex Metabolic Effects, n = 24) (Supplementary Table 1).

Sequence validation of rare functional variants. Novel, functional variants in all 119 candidate genes were taken forward for validation by conventional Sanger sequencing (90% validation rate). Briefly, customized PCR primers were designed +/− 250 bp surrounding the variant; and sequencing was performed using BigDye Terminator v3 kit (Applied Biosystems) and analysed by capillary electrophoresis on an ABI3730 DNA Analyzer platform (Applied Biosystems), according to the manufacturers’ instructions. Familial segregation analysis of variants was performed where family samples were available.

Identification of clinically-associated variants in known human obesity genes. The strategy for identification of clinically-associated variants in known human obesity genes is outlined in Fig. 5. Briefly, novel and rare variants in the known human obesity genes were filtered using the ClinVar database (http://www.ncbi.nlm.nih.gov/clinvar/). We retained novel variants and those with a ClinVar status of Pathogenic/Likely pathogenic (N = 332).

Structural Analysis of GNAS Variants. For structural analysis of the genetic variants and prediction of their impact on protein function, we modelled their location in relation to previously solved protein crystal structures of GNAS in complex with either G-protein coupled receptor (ADRB2) or downstream effector (adenylyl cyclase). Specifically, the first model represented in Fig. 3a is based on crystal structure of the active state ternary protein complex of GPCR beta-2-adrenergic receptor (ADRB2), and the nucleotide-free Gs heterotrimer, composed of Gs-α(GNAS), Gs-β, and Gs-γ subunits (ref. 20, PDB number: 3SN6, www.rcsb.org). The second model, represented in Fig. 3b, is based on complex of GNAS with the catalytic domains of mammalian adenylyl cyclase (ref. 21, PDB number: 3SN6, www.rcsb.org).

**Association Analyses.** We implemented the optimal Sequence Kernel Association Test SKAT-O\(^{11,53}\) with the SKAT R package v1.1.2\(^{14}\) using options method = "optimal.adj" for SKAT-O and r.corr = 1 for burden. For each gene and candidate gene set, we performed two primary tests using the burden test: 1. rare functional (Rare Func) or (2) novel functional (Novel Func). We performed secondary tests restricting either to variants that were predicted to be deleterious by at least one of the three algorithms (i.e. SIFT, PolyPhen, and Condel) \((1 + del)\) or to variants that were predicted to be deleterious by all three algorithms \((all\ del)\). For SIFT and Condel, variants were classified as deleterious if they were labelled as “deleterious”. For PolyPhen, variants were classified as deleterious if they were labelled as “probably damaging” or “damaging”. Our conservative Bonferroni adjusted significance level of the primary analyses for 119 genes and 6 gene sets was \(0.05/(2 \times 125) = 2.0E-4\). Once considering our additional ten secondary tests, our Bonferroni adjusted significance level was \(0.05/(12 \times 125) = 3.3E-5\). We repeated all primary and secondary tests using SKAT-O (Table 1; Supplementary Table 4–9, 13).

**Functional studies of variants in ANGPTL6.** To predict the impact of variants on the ANGPTL6 protein, we modelled their location using the crystal structure of a related protein fibrinogen in a complex with a peptide Gly-His-Pro-amide (described in Yang et al.\(^{20}\), PDB number: ILWU, www.rcsb.org). The structural representations using ribbon-depicted models were generated using the Open-Source PyMOL Molecular Graphics System, Version 1.7.x Schrödinger, LLC (http://pymol.org). N-terminal triple Flag tag was added to human ANGPTL6 cDNA cloned into a PEZ-M14 mammalian expression vector (Capital Bioscience). Mutations were introduced into this construct using QuikChange (Agilent technologies), and confirmed by Sanger sequencing. ANGPTL6 protein expression was studied by transient transfection in HEK293 cells. Media was collected 48 h post-transfection, the cell medium was centrifuged for 5 min \((5,000 \times g; \text{at } 4^\circ C)\) and the supernatants were collected. Cells were harvested prior to centrifugation for 15 min \((15,000 \times g; \text{at } 4^\circ C)\). Aliquots from the medium and cells were subjected to SDS-PAGE and immunoblot analysis. Primary antibodies (monoclonal Flag M2 antibody (Sigma) and a polyclonal antibody to Calnexin (Cell signaling)) were used at 1:1000, and secondary Horseradish peroxidase–conjugated anti-mouse or anti-rabbit IgG (Dako) antibody at 1:2000 dilution. After staining with ECL West Dura Substrate kit (Thermo Scientific), visualisation was performed on Chemidoc Digital Image (Bio-Rad) Software. The relative protein secretion of ANGPTL6 for each variant was standardized to wild-type protein secretion. Experiments were performed in triplicate and analysed using an unpaired \(t\)-test with Welch’s correction.

**ANGPTL6 statistical follow-up analyses.** We ran gene-based analyses using SKAT-O as described above on two subsets of ANGPTL6 variants (Supplementary Table 10): (1) the one variant that resulted in a complete loss of protein secretion \((S284X; \text{rs201622589})\), and (2) excluding \(S284X\). For (1), we also used a Fisher’s Exact Test. We repeated the Fisher’s Exact Test of the \(S284X\) variant including additional population controls with pre-existing genotype data and replication samples (Online Methods, Sample Sets; Supplementary Tables 11 and 12).

**Power Analysis.** We calculated the power to detect association to a gene region using the burden test with the \(\text{Power}_{\text{Logistic}}\ R\) function and the haplotype dataset within the SKAT R-package\(^{16}\). We performed power calculations using 500 simulations on a random 2kb sub region. The effect sizes of the causal variants are equal to \(\log_{10}(\text{MAF})\) with a maximum effect size of 1.6 \((\text{MAF} = 0.0001)\) and all have the same direction of effect. We limited the MAF threshold for causal variants to \(\leq 0.01, 0.001, 0.0005\) and varied the percentage of causal variants in the region to be between 10–90%. (More details in Supplementary Note 4).

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