Heterozygous \textit{RFX6} protein truncating variants are associated with MODY with reduced penetrance

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Finding new causes of monogenic diabetes helps understand glycaemic regulation in humans. To find novel genetic causes of maturity-onset diabetes of the young (MODY), we sequenced MODY cases with unknown aetiology and compared variant frequencies to large public databases. From 36 European patients, we identify two probands with novel \textit{RFX6} heterozygous nonsense variants. \textit{RFX6} protein truncating variants are enriched in the MODY discovery cohort compared to the European control population within ExAC (odds ratio $= 131$, $P = 1 \times 10^{-4}$). We find similar results in non-Finnish European ($n = 348$, odds ratio $= 43$, $P = 5 \times 10^{-5}$) and Finnish ($n = 80$, odds ratio $= 22$, $P = 1 \times 10^{-6}$) replication cohorts. \textit{RFX6} heterozygotes have reduced penetrance of diabetes compared to common \textit{HNF1A} and \textit{HNF4A}-MODY mutations ($27$, $70$ and $55\%$ at $25$ years of age, respectively). The hyperglycaemia results from beta-cell dysfunction and is associated with lower fasting and stimulated gastric inhibitory polypeptide (GIP) levels. Our study demonstrates that heterozygous \textit{RFX6} protein truncating variants are associated with MODY with reduced penetrance.
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inding the genetic cause of rare familial diabetes (monogenic diabetes) provides new biological insights into human pancreas development and function, as well as potentially novel therapeutic targets with important treatment implications. Maturity-onset diabetes of the young (MODY) is monogenic diabetes resulting from beta-cell dysfunction which usually present before the age of 25 years in non-obese patients who are non-insulin-dependent and have an autosomal dominant inheritance of diabetes. Mutations in HNF1A, HNF4A and GCK are the commonest causes of MODY responsible for ~60% of MODY aetiology.

There has been limited recent success in finding new MODY genes. WFS1 heterozygous variants and loss-of-function variants in the APPL1 gene were shown to be a rare cause of MODY. The reason for this limited success is the difficulty of distinguishing monogenic diabetes patients from those with type 1 diabetes, or from the increasing number of patients with early-onset type 2 diabetes due to rising rates of obesity. Another important reason is the lack of large pedigrees with an autosomal dominant pattern of inheritance of diabetes which would allow classical linkage analysis to be performed and which was used to discover the most common forms of MODY such as GCK, HNF1A and HNF4A.

Rare-variant association testing is an important step to confirm the pathogenicity of novel variants in monogenic disease. Rare-variant association testing particularly for comparing the frequency of novel protein-truncating variants (PTVs) in monogenic cases with unknown aetiology to the frequency in large control cohorts is now possible because of the availability of resources such as ExAC – a database of protein coding variants in large control populations. This allows burden testing of the frequency of novel or rare coding variants in diseases of interest and a comparison to rates in controls to identify new genetic causes of monogenic disease.

In this study, we have undertaken next-generation sequencing of MODY cases with unknown aetiology and compared the frequency of PTVs to large publicly available control cohorts to identify new MODY genes. Our study shows that heterozygous RFX6 PTVs are associated with MODY.

Results

Heterozygous RFX6 PTVs in MODY with unknown aetiology.

To identify patients with novel heterozygous PTVs, we first assessed 38 European (non-Finnish) probands with a strong MODY-like phenotype who did not have mutations in the common MODY genes (GCK, HNF1A, HNF4A) by Sanger sequencing (Supplementary Table 1). To exclude the other known/less common causes of monogenic diabetes, these patients underwent comprehensive-targeted-next generation sequencing (NGS) for all 29 known monogenic diabetes genes, including genes for neonatal diabetes, MODY and mitochondrial diabetes, lipodystrophy or other forms of syndromic diabetes (Supplementary Table 2). We identified two probands with mutations in the known MODY gene HNF1B.

RFX6 PTVs are enriched in a MODY discovery cohort.

To identify patients with novel heterozygous PTVs, we compared the frequency of RFX6 PTVs in our discovery cohort to a large control population with whole-exome data from ExAC. Neither of the RFX6 variants from the discovery cohort were present in the 60,706 individuals in ExAC. There were 15 individuals with RFX6 PTVs in the 33,346 ExAC non-Finnish European control population (Supplementary Table 3). The frequency of the RFX6 PTVs in the MODY discovery cohort was significantly higher (after accounting for the multiple testing of 29 genes) than the ExAC non-Finnish European control population (5.5 vs. 0.045%, odds ratio (OR) 131, 95% confidence interval (CI) 14-595, P = 1 × 10-8) (Table 1).

RFX6 PTVs are enriched in a MODY replication cohort.

To replicate the findings of our discovery cohort, we then examined 348 non-Finnish European probands who were routinely referred for MODY genetic testing to the Molecular Genetics Laboratory, Exeter, UK and in whom the common causes of MODY were excluded using targeted-NGS assay (Supplementary Table 1). The analysis of heterozygous PTVs identified four unrelated probands with two novel RFX6 nonsense variants (p.Gln25Ter, p.Arg377Ter) (Supplementary Table 1 and Supplementary Table 3). Similarly to the discovery cohort, the MODY replication cohort was enriched for RFX6 PTVs compared to the ExAC non-Finnish European control population (1.15 vs. 0.045%, OR = 26, 95% CI 6-82, P = 3 × 10-7) (Supplementary Table 4). This association was maintained when compared to an independent non-Finnish European control population with whole-genome sequence data from gnomAD (http://gnomad.broadinstitute.org) (Table 1 and Supplementary Table 4). The frequency of RFX6 PTVs in the gnomAD genome data set (0.027%) is not statistically different to that in ExAC (0.045%, P = 0.76).

Higher frequency of RFX6 PTVs in Finnish population.

Finnish individuals had ~10-fold higher frequency of RFX6 PTVs compared to non-Finnish Europeans. The ExAC database showed a relative abundance of RFX6 PTVs in Finnish Europeans (15/3305, 0.45%) compared to non-Finnish Europeans (15/33,346, 0.045%) (Supplementary Table 3). All of the Finnish individuals in ExAC with RFX6 PTVs had the same variant, p.His293Leufs. To further validate this finding in a larger Finnish control population, we analysed RFX6 PTVs in 7040 control individuals from the METSIM study in

Table 1 Frequency of heterozygous RFX6 protein-truncating variants in all study cohorts and control populations

| MODY cohorts, frequency of RFX6 PTV | Control population, frequency of RFX6 PTV | Odds ratio (95% CI) | P-value |
|------------------------------------|-----------------------------------------|---------------------|-------|
| Non-Finnish European Discovery cohort (n = 2/36), 5.5% | ExAC - exomes (n = 15/33,346), 0.045% | 131 (14-595) | 1 × 10-4 |
| Replication cohort (n = 4/348), 1.15% | gnomAD - genomes (n = 2/7508), 0.027% | 43 (6-483) | 5 × 10-5 |
| Finnish European Replication cohort (n = 6/80), 7.5% | METSIM-exomes (n = 26/7040), 0.37% | 22 (7-56) | 1 × 10-6 |
| Meta-analysis Heterogeneity (I² = 2.73 (d.f. = 2), P = 0.256) | | 34 (15-80) | 1 × 10-16 |

CI, confidence interval; MODY, maturity-onset diabetes of the young; PTV, protein-truncating variant.
Eastern Finland\textsuperscript{16}. There were 26 individuals with RFX6 PTVs in this cohort and all had the p.His293Leufs variant. The frequency of p.His293Leufs was not significantly different from the ExAC Finnish population frequency (0.37 vs. 0.45%, \( P = 0.63 \)) (Supplementary Table 3). The METSIM study has contributed to the ExAC Finnish cohort, so to prevent duplication we used the data from the larger METSIM study for further analysis\textsuperscript{12}.

**Enrichment of RFX6 p.His293Leufs in Finnish MODY patients.** To assess whether the p.His293Leufs variant is associated with MODY in Finnish patients, we genotyped the RFX6 p.His293Leufs variant in 80 Finnish probands who were routinely referred for MODY genetic testing to Genome Center of Eastern Finland, University of Eastern Finland and did not have mutations in the most common MODY genes (GCK, HNF1A, HNF4A and HNF1B) (Supplementary Table 1). We identified six probands with the p.His293Leufs variant. The frequency of this variant was significantly higher in the Finnish MODY cohort compared to the METSIM controls (7.5 vs. 0.37%, OR = 22, 95% CI 7–56, \( P = 1 \times 10^{-6} \)) (Table 1). The meta-analysis of the three independent case–control analyses confirmed the strong
The penetrance of diabetes for RFX6 PTVs with MODY in the study cohorts (OR = 34, 95% CI 15–80, \( P = 1 \times 10^{-19} \)) (Table 1).

**Enrichment of RFX6 PTVs is not due to technical artefacts.** To ensure that the association we observed is not due to differences in sequencing technologies or analysis pipelines between cases and controls, we performed a series of sensitivity analyses. This included comparisons to additional whole exome, whole genome and in-house control cohorts and an analysis that removed exon 1 which was the least well covered exon in ExAC. These sensitivity analyses (Supplementary Table 4) show that results are consistent for all these analyses.

**RFX6 PTVs co-segregate with diabetes.** To further assess the causality of RFX6 PTVs, we conducted a co-segregation analysis in families with genetic data available on more than three affected individuals. We had only one family (family 1) with >3 affected individuals with genetic data (Fig. 1). The analysis showed that the RFX6 variant p.Leu292Ter co-segregated in 9 out of 10 individuals with diabetes (LOD score = 0.65, \( P = 0.04 \)). One individual without the RFX6 variant had diabetes which is likely to be a phenocopy of type 2 diabetes considering the large pedigree, age of diagnosis and obesity (51 years, body mass index (BMI) 30 kg/m\(^2\)). There were two family members with an RFX6 variant but with normal HbA1c level at the time of study (18 and 57 years) suggesting that RFX6 PTVs may have reduced penetrance.

**Reduced penetrance of diabetes with RFX6 PTVs.** To assess the penetrance of RFX6 PTVs for diabetes compared to common causes of MODY, we combined data for all six non-Finnish European proband families. There were 18 RFX6 heterozygotes of whom five had not developed diabetes at study entry. 27% (95% CI 11–58) developed diabetes by the age of 25 years and 78% (95% CI 55–95) by 51 years (Fig. 2). Two out of six probands did not have affected parents at study entry (Supplementary Fig. 1). The penetrance of diabetes for RFX6 heterozygotes was substantially lower compared to pathogenic variants of HNF1A (70%, 95% CI 67–72 by the age of 25 years and 97%, 95% CI 96–98 by 50 years) and moderately lower than pathogenic variants of HNF4A (55%, 95% CI 50–60 by the age of 25 years and 91%, 95% CI 88–94 by 50 years) (Fig. 2). Similar to non-Finnish European proband families, the Finnish RFX6 p.His293Leufs variant also showed reduced penetrance in Finnish families (Supplementary Fig. 2). In two previously reported families of neonatal diabetes children with homozygous p.Arg181Gln RFX6\(^{13,18} \) or RFX6 p. His293Leufs\(^{19} \), the genetic information available on RFX6 heterozygous family members was also suggestive of reduced penetrance of diabetes (Supplementary Figs 2 and 3).

**RFX6 PTVs are not enriched in type 2 diabetes.** The reduced penetrance and later age of onset of diabetes with RFX6 PTVs raised the possibility that these variants may be associated with type 2 diabetes. To assess this, we used freely available data from the Type 2 Diabetes Knowledge Portal which contains whole-exome data on type 2 diabetes patients\(^{20} \). Burden testing of RFX6 PTVs for exome sequencing data from 8373 type 2 diabetes cases and 8466 controls showed no significant association with type 2 diabetes (0.14 vs. 0.083%, OR = 1.79, 95% CI 0.7–4.57, \( P = 0.22 \)) (Supplementary Table 3).

**Phenotype of RFX6 heterozygotes with diabetes.** We assessed the diabetes phenotype in 27 RFX6 heterozygote individuals with diabetes. The clinical features are shown in Table 2. The median age at diagnosis of diabetes was 32 years (IQR 24–46, range 13–64 years) and median BMI of 25.1 kg/m\(^2\) (IQR 23–28). After a median 10 years (IQR 5–22) of diabetes 69% of patients were treated with insulin but there was significant endogenous insulin present in 24/25 patients at recruitment. There was no history of sulphonylurea sensitivity and they did not have islet auto-antibodies (GADA/IA2-Ab). All patients had isolated diabetes and there were no reports of the other features of homozygous RFX6 mutations, such as duodenal or gall bladder atresia.

**RFX6 haploinsufficiency is associated with reduced GIP.** RFX6 is a transcription factor and has been shown to increase expression and secretion of gastric inhibitory polypeptide (GIP) in mouse enterodendocrine K-cells\(^{21} \). We therefore measured the incretin hormone GIP in 17 RFX6 heterozygotes (eight with diabetes) and compared to 26 controls (two with diabetes). The fasting GIP was markedly lower in RFX6 heterozygotes compared to controls (16 (10–24) vs. 49 (28–65) pg ml\(^{-1}\), \( P = 1.2 \times 10^{-20} \)). Fasting glucagon-like peptide-1 (GLP-1) levels were not different in both groups (23 (12.5–32) vs. 24 (14–32) pg ml\(^{-1}\), \( P = 0.98 \)).

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**Table 2 Clinical characteristics of patients with RFX6-MODY**

| Characteristic                             | (n = 27) |
|-------------------------------------------|----------|
| Age at diagnosis (years), median (IQR)    | 32 (24–46) |
| Duration of diabetes, median (IQR)        | 10 (5–22) |
| Female, n (%)                             | 22 (81%) |
| BMI (kg/m\(^2\)), median (IQR)            | 25.1 (23–28) |
| **Initial treatment, n (%)**              |          |
| Diet                                      | 4 (15%)  |
| Oral hypoglycaemic agents                 | 14 (54%) |
| Insulin                                   | 5 (19%)  |
| Insulin + oral hypoglycaemic agents       | 3 (12%)  |
| **Current treatment, n (%)**              |          |
| Diet                                      | 2 (8%)   |
| Oral hypoglycaemic agents                 | 6 (23%)  |
| Insulin                                   | 12 (46%) |
| Insulin + oral hypoglycaemic agents       | 6 (23%)  |
| HbA1c at recruitment, mmol/mol, median (IQR) | 51 (45–70) |

Significant endogenous insulin at recruitment\(^4\), 24 (96%) n = 25

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BMI: body mass index; IQR: interquartile range; MODY: maturity-onset diabetes of the young.

\(^1\)Non-insulin treated or insulin treated with urine/blood random C-peptide > 200 pmol/l at recruitment

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**Fig. 2** Penetrance of diabetes in people with MODY. Heterozygous RFX6 PTV (n = 18), pathogenic HNF1A variant (n = 1265) or HNF4A variant (n = 427)

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remove potential confounding factors, we compared the OGTT data for the 11 Finnish RFX6 p.His293Leu fs heterozygotes without diabetes to five matched (age, sex and BMI) controls for each heterozygote from the PPP-Botnia Study (Fig. 3 and Supplementary Table 6). This confirmed that both fasting and 120 min stimulated GIP was reduced (18.3 vs. 48.9 pg ml$^{-1}$, $P=8 \times 10^{-3}$, 167 vs. 241 pg ml$^{-1}$, $P=0.029$, respectively). In addition, the non-diabetic heterozygotes had higher fasting glucose (5.5 vs. 5.1 mmol l$^{-1}$, $P=0.02$) with a similar fasting insulin level suggesting a beta-cell defect (Supplementary Table 6).

**Discussion**

Heterozygous RFX6 PTVs are associated with MODY. We identified RFX6 PTVs predicted to be pathogenic$^{11, 19}$ in unrelated MODY patients in whom the known causes of monogenic diabetes had been excluded. These variants were enriched in Finnish and non-Finnish European MODY probands but were rare in the control cohorts and in patients with type 2 diabetes. We observed co-segregation within pedigrees albeit with reduced penetrance. Finally, these variants are likely to have a functional effect due to nonsense-mediated decay causing haploinsufficiency$^{12, 22}$. This is further supported by the studies that showed that the homozygous RFX6 PTVs cause neonatal diabetes$^{15, 23, 24}$. Among unknown MODY cases, RFX6 PTVs were responsible for 7.5% Finnish cases compared to only ~1% of non-Finnish European cases.

Large-scale control cohorts such as ExAC in combination with next-generation sequencing of well-characterised cases is a useful strategy for identifying new causes of monogenic disease. The large ExAC database provides sufficient power for reliable burden testing of rare variants in monogenic disease$^{12}$ without the need for large pedigrees or linkage analysis. RFX6 PTVs were highly enriched in both discovery and replication cohorts compared to control cohorts, supporting their pathogenicity. The ExAC database has been very useful in identifying benign variants because of an unusually high frequency in the population compared to frequency of the disease in question$^{11, 12, 25}$. However, caution is required for reduced penetrance variants as their frequency can be higher than estimated disease frequency in the general population. RFX6 PTVs are an example where the reduced penetrance explains the higher frequency in control cohorts compared to the estimated frequency of MODY (0.01%) in the general population$^{26}$. In addition, our study highlights the importance of population specific control and disease cohorts. The frequency of RFX6 PTVs is ~10-fold higher in the Finnish population compared to non-Finnish populations due to the well-documented bottleneck in population genetics$^{27, 28}$. This observation is not
restricted to RFX6 alone. It has been shown that the Finnish population has an overall higher burden of genome-wide PTVs (0.5–5%) in many genes compared to non-Finnish Europeans, and some of these have been associated with disease. RFX6 PTVs are associated with reduced penetrance in MODY. This reduced penetrance explains the lack of complete co-segregation in the RFX6 pedigrees. It also clarifies why diabetes was only reported in the parents or grandparents (obligate heterozygous for functional RFX6 variants) in seven out of the 12 published recessive RFX6 neonatal diabetes pedigrees. The lack of enrichment of RFX6 PTVs in type 2 diabetes patients compared to controls further supports their association with reduced penetrance MODY rather than type 2 diabetes. Further studies are needed to understand the mechanism of reduced penetrance of diabetes in RFX6 heterozygotes. It could be due to a combination of factors, such as expression patterns of normal alleles, epigenetic modifications and rare or common genetic variant modifiers.

There are differences as well as similarities between RFX6-MODY and HNF1A/HNF4A-MODY. In contrast to HNF1A and HNF4A-MODY patients, RFX6-MODY patients do not show enhanced sensitivity to sulphonylureas. All patients with HNF1A/HNF4A-MODY have significant endogenous insulin 3–5 years post diagnosis. RFX6-MODY patients showed a similar pattern, except for one patient who did not have detectable endogenous insulin. Similar to HNF1A/HNF4A-MODY, RFX6-MODY patients lack islet autoantibodies and have isolated diabetes. This suggests that persistent C-peptide, lack of islet autoantibodies and parental history of diabetes, which are currently used to distinguish common forms of MODY from type 1 diabetes, can also be used to identify RFX6-MODY. However, a similar strategy will not help to distinguish late onset RFX6-MODY from type 2 diabetes.

Our study supports the role of RFX6 in the adult human pancreas. RFX6 is from a family of transcription factors that contains winged-helix DNA-binding domains. RFX6 is expressed almost exclusively in pancreatic islets, small intestine and colon. It acts downstream of NGN3, regulates islet cell differentiation and the development of the endocrine pancreas. The homozygous RFX6 missense and PTVs cause syndromic neonatal diabetes (gall bladder aplasia, gut atresia and diabetes). RFX6 whole-body null mice show phenotypes consistent with human disease and die soon after birth, but the heterozygous whole-body RFX6 mouse has not been reported to develop diabetes. This is not surprising considering the lack of phenotype in heterozygous null mice of HNF1A and HNF4B. Interestingly, the defect in glucose-induced insulin secretion was present in models that are more akin to haploinsufficiency of RFX6 in adult humans. 80% depletion of RFX6 protein in the adult mouse pancreas in vivo as well as in human beta cells in vitro showed that this defect was due to reduced expression of ABCCC8, GCK and Ca2+ channels in beta cells and disruption of Ca2+-mediated insulin secretion. These data support the role of RFX6 in the physiology of adult beta cells. This along with evidence of impaired of beta-cell function (requirement of insulin to maintain euglycaemia, one patient with C-peptide <200 pmol l−1, lower C-peptide in heterozygotes with diabetes compared to without diabetes) suggests that insulin deficiency is the cause of diabetes in these patients.

Incretin analysis: We completed physiological studies on 25/47 RFX6 heterozygotes in whom 10 had diabetes. 7/25 had fasting blood sample analysis and 18/25 had 75 g oral glucose tolerance test (OGTT). We used 27 (12 family and 15 unrelated) controls that included two individuals with diabetes for an initial comparison of fasting GIP/GLP-1. RFX6 heterozygotes had decreased GIP secretion.

Methods

Study populations. Discovery MODY cohort: The discovery cohort comprises 38 European probands with strong MODY-like phenotype who did not have mutations in the three most common MODY genes (GCK, HNF1A and HNF4A) and 7040). They were all males aged 45–70 years, randomly selected from the population register of the Kuopio town, Eastern Finland. Non-Finnish European replication MODY cohort: The replication cohort was derived from 469 non-Finnish European routine MODY diagnostic referrals to the Molecular Genetic Laboratory, Exeter, UK. A monogenic aetiology in a known monogenic diabetes gene was identified in 121 patients and the remaining 348 patients with unknown aetiology comprised the replication cohort (Supplementary Table 1).

Finnish-European replication MODY cohort: This cohort consisted of 80 patients who were routinely referred for MODY diagnostic testing to the Genome Center of Eastern Finland, University of eastern Finland in whom no mutation was found in the common MODY genes (GCK, HNF1A, HNF4B and HNF4A) when tested by Sanger sequencing (Supplementary Table 1). These 80 patients comprise 78% of the total MODY X Finnish cohort.

Finnish-European control cohort: Individuals of this cohort were part of the METSIM study (n = 7040). They were all males aged 45–70 years, randomly selected from the population register of the Kuopio town, Eastern Finland. From a previously reported family from Belgium (Supplementary Fig. 3).

Incretin analysis: We completed physiological studies on 25/47 RFX6 heterozygotes in whom 10 had diabetes. 7/25 had fasting blood sample analysis and 18/25 had 75 g oral glucose tolerance test (OGTT). We used 27 (12 family and 15 unrelated) controls that included two individuals with diabetes for an initial comparison of fasting GIP/GLP-1. RFX6 heterozygotes had decreased GIP secretion.
Plasma glucose was analysed using the Hemocue Glucose System (HemoCue AB, Angelholm, Sweden). Serum insulin was measured by an AutoDelfia fluoroenzymometric assay (PerkinElmer, Waltham, Massachusetts, US) and serum C-peptide by Cobas e411 electrochemiluminometric immunoanalysis (Roche, Mannheim, Germany). Serum GIP was analysed using Millipore’s Human GIP Total ELISA (Merck, Darmstadt, Germany; catalogue # EZHGP-54K) which has 100% cross-reactivity to both human GIP (1–42) and GIP (3–42). Serum total PLA-1 concentrations, which detects both intact PLA-1 and PLA-1 (9–36 amide), were determined using Millipore’s radioimmunoassay (Merck, Darmstadt, Germany; catalogue # GLPIT-36HK). Serum total cholesterol, high-density lipoprotein and triglyceride concentrations were measured first on a Cobas Mira analyser (Hoffmann-La Roche, UK), and low-density lipoprotein was calculated with the Friedewald formula, and since January 2006 with an enzymatic method (Konelab 60 analyser; Thermo Electron Oy, Vantaa, Finland).

The RXf6 p.His293Leufs variant was genotyped in 5187 individuals by the Kompetitive Allele Specific PCR genotyping system (KASPTM) on Demand (KOD) assay according to the manufacturer’s testing conditions including six positive control samples identified by direct sequencing (LGc Hoddesdon, Herts, UK). Two out of 5180 participants had RXf6 p.His293Leufs (the genotyping failed in 7), which was confirmed by direct sequencing.

DNA analysis. Targeted next-generation sequencing: The analysis of all known monogenic diabetes genes in European cohorts was conducted using targeted-NGS[9]. The panel included 29 genes in which variants are known to cause monogenic neonatal diabetes, MODY, mitochondrial diabetes, lipodystrophy or other forms of syndromic diabetes[10] (Supplementary Table 2). The RXf6 PTVs identified by targeted-NGS were identified using Sanger sequencing. The essential splice site, nonsense and frameshift variants excluding the last exon were considered PTVs in this study[11,12]. The targeted-NGS assay covered 100% bases of the RXf6 coding region with >10x read depth for all the samples.

Sanger sequencing: Genomic DNA was extracted from whole blood using standard procedures and the coding region and intron/exon boundaries of the RXf6 gene were amplified by PCR. Amplicons were sequenced using the Big Dye Terminator Cycle Sequencing Kit v3.1 (Applied Biosystems, Warrington, UK) according to manufacturer’s instructions and reactions were analysed on an ABI 3730 Capillary sequencer (Applied Biosystems, Warrington, UK). Sequences were compared with the reference sequences (NM_173580.3) using Mutation Surveyor v3.24 software (Soft Genetics, State College, PA, USA).

The Finnish-European MODY cohort was analysed for p.His293Leufs variant using Sanger sequencing as described above. Family co-segregation analysis was performed in available family members using a Sanger sequencing assay for the specific RXf6 variant identified in that family. DNA analysis of the METSIM study has been described before[12].

Statistical analysis. Fisher’s exact test was used to compare the frequency of RXf6 PTVs. The threshold P-value for association was 1 × 10^-5 as there were 29 genes on the panel (0.05/29). The penetrance of diabetes was assessed using survival time analysis method. The statistical analysis was conducted using Stata 14 (StataCorp, Texas, USA). The comparison of RXf6 heterozygotes to PPP-Botnia controls were conducted using R (3.3.2) with packages for the data manipulation (dplyr) and visualization (ggplot2). Continuous variables were compared with Mann–Whitney U-test for categorical variables with chi-squared test. Single point non-parametric linkage analyses were performed using MERLIN 1.1.2[17]. The Z-score was converted into a LOD score by use of the Kong and Cox exponential model[18].

Ethics. Informed consent was obtained from all subjects. The UK study is approved by the North Wales Research Ethics Committee. The FINNMODY/PPP-Botnia study is approved by the Research Ethics committee for Medicine of the Helsinki University Hospital.

Data availability. The majority of data used in this study are publically available and can be accessed via the studies cited in the text. Considering issues of patient confidentiality and restrictions in IBR permissions, other original data are available through specific request.

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Author contributions
K.A.P., J.K., M.C. and M.N.W. researched data and performed statistical analyses. K.A.P., J.K., M.C. and T.T. wrote the first draft of the manuscript, which was modified by all authors. All authors contributed to the discussion and reviewed or edited the manuscript. M.N.W., M.C. and T.T. are the guarantors of this work and, as such, had full access to all the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

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