

Identification of maturity onset diabetes of the young caused by mutation in *FOXM1* via whole-exome sequencing in Northern China

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Supplementary Table 1. Pairwise comparison between diabetic groups

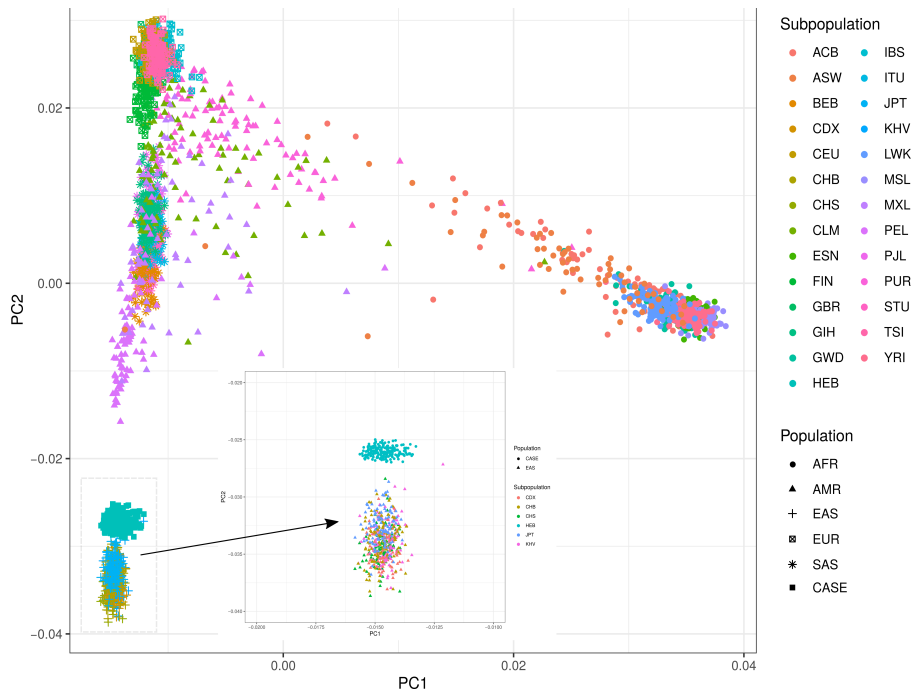
	<i>Group1</i>	<i>Group2</i>	<i>p-Value</i>	<i>Adjusted p value</i>	<i>Significance</i>
<i>Age</i>	MODY	T1DM	0.00725823	0.0073	**
<i>Age</i>	MODY	T2DM	1.49E-06	4.50E-06	****
<i>Age</i>	T1DM	T2DM	0.000194237	0.00039	***
<i>BMI</i>	MODY	T1DM	0.2	0.22	ns
<i>BMI</i>	MODY	T2DM	0.00756331	0.023	**
<i>BMI</i>	T1DM	T2DM	0.109961397	0.22	ns
<i>FPG</i>	MODY	T1DM	0.406344451	0.41	ns
<i>FPG</i>	MODY	T2DM	0.00868066	0.017	**
<i>FPG</i>	T1DM	T2DM	0.001240885	0.0037	**
<i>FPGC-peptide</i>	MODY	T1DM	0.574835652	0.57	ns
<i>FPGC-peptide</i>	MODY	T2DM	0.000220456	0.00044	****
<i>FPGC-peptide</i>	T1DM	T2DM	1.28E-05	3.90E-05	****
<i>HbA1c</i>	MODY	T1DM	0.06022889	0.12	ns
<i>HbA1c</i>	MODY	T2DM	0.000239988	0.00072	****
<i>HbA1c</i>	T1DM	T2DM	0.688947849	0.69	ns
<i>Insulin</i>	MODY	T1DM	0.969851098	0.97	ns
<i>Insulin</i>	MODY	T2DM	0.063225047	0.13	ns
<i>Insulin</i>	T1DM	T2DM	0.002002185	0.006	**
<i>Hepatic Fat Content</i>	T1DM	T2DM	0.203	0.2	ns
<i>Cholesterol</i>	MODY	T1DM	0.49214177	0.98	ns
<i>Cholesterol</i>	MODY	T2DM	0.216746626	0.65	ns
<i>Cholesterol</i>	T1DM	T2DM	0.981847741	0.98	ns
<i>Serum creatinine</i>	MODY	T1DM	0.53932819	1	ns
<i>Serum creatinine</i>	MODY	T2DM	0.918421945	1	ns
<i>Serum creatinine</i>	T1DM	T2DM	0.23998686	0.72	ns
<i>Triglyceride</i>	MODY	T1DM	0.197685565	0.4	ns
<i>Triglyceride</i>	MODY	T2DM	0.683405314	0.68	ns
<i>Triglyceride</i>	T1DM	T2DM	0.062083274	0.19	ns
<i>HDL</i>	MODY	T1DM	0.920108282	1	ns
<i>HDL</i>	MODY	T2DM	0.666186795	1	ns
<i>HDL</i>	T1DM	T2DM	0.897402908	1	ns
<i>LDL</i>	MODY	T1DM	0.569700822	1	ns
<i>LDL</i>	MODY	T2DM	0.304233578	0.91	ns
<i>LDL</i>	T1DM	T2DM	0.755838935	1	ns

¹ The pairwise comparison was performed by Wilcoxon test. The P value was adjusted by Benjamini & Hochberg(FDR) method.

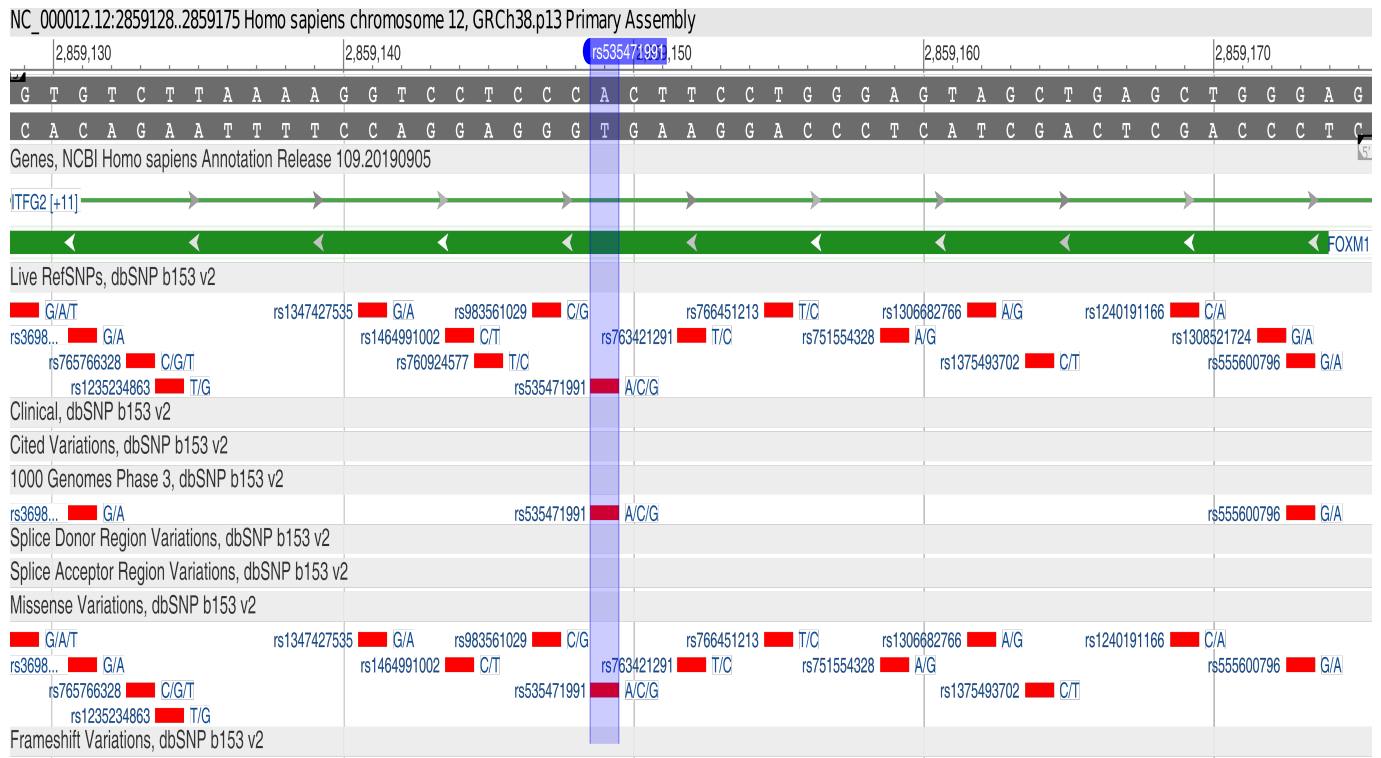
² 'ns': p > 0.05 ; '*': p <= 0.05 ; '**': p <= 0.01 ; '***': p <= 0.001 ; '****': p <= 0.0001

Supplementary Table 2. Variation count in 14 type MODY gene

	D620	D855	D884	D913	D1103	D1111	D1112	D1174	D1384	D1389
<i>Hnf4A</i>	9	16	18	3	8	12	2	2	16	3
<i>Gck</i>	12	10	10	5	1	5	5	10	10	5
<i>Hnf1A</i>	9	18	17	20	22	22	20	21	24	21
<i>Pdx1</i>	0	0	1	0	2	3	1	0	0	1
<i>Hnf1B</i>	0	0	0	1	0	3	2	0	2	0
<i>Neurod1</i>	1	1	1	1	1	1	1	1	1	1
<i>Klf11</i>	1	1	2	2	1	1	2	1	2	1
<i>Cel</i>	0	1	1	1	0	2	1	1	1	1
<i>Pax4</i>	5	0	0	8	0	5	0	4	7	5
<i>Ins</i>	5	5	3	3	6	6	6	5	6	6
<i>Blk</i>	20	16	20	17	27	18	18	33	23	21
<i>Abcc8</i>	17	19	25	14	18	16	13	14	21	20
<i>Kcjn11</i>	0	0	0	0	0	0	0	0	0	0
<i>Appl1</i>	3	2	6	2	6	6	5	2	4	5



Supplementary Figure 1. PCA plot 200 subjects and 1000 Genome Project. The combined PCA analysis of 200 subjects and all samples from 1000 Genome Project. The main population (AFR: African; AMR: American; EAS: East Asian; EUR: European; SAS: South Asian) and 200 diabetic subjects (CASE) were grouped by different shapes. 27 Sub-populations were grouped by different colors. The detailed PCA analysis of EAS samples and the 200 subjects was also displayed in the bottom of the figure.



Supplementary Figure 2. rs535471991 annotated in variation databases The genomic information and the variation annotation from various databases of rs535471991.