

Full name:	MODY (7 genes) - UZA
Description:	Last update: 24/12/2020
Version number:	MODY_v1
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Related Diseases

- Autosomal recessive hyperinsulinism due to SUR1 deficiency
- Congenital hyperinsulinism due to HNF4A deficiency
- Hyperinsulinism due to HNF1A deficiency
- Hyperinsulinism due to INSR deficiency
- Isolated permanent neonatal diabetes mellitus
- MODY
- Transient neonatal diabetes mellitus

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
<u>ABCC8</u>			
<u>GCK</u>			
<u>HNF1A</u>			
<u>HNF1B</u>			
<u>HNF4A</u>			
<u>INS</u>			
<u>KCNJ11</u>			

