DISEASE:
Isolated permanent neonatal diabetes mellitus

NAME:	Isolated permanent neonatal diabetes mellitus
DESCRIPTION:	Permanent neonatal diabetes mellitus (PNDM) is a monogenic form of neonatal diabetes (NDM, see this term) characterized by persistent hyperglycemia within the first 12 months of life in general, requiring continuous insulin treatment.
ORPHACODE:	99885
SYNONYMS:	Isolated PNDM Monogenic diabetes of infancy
XREF(S):	Orphanet OMIM OMIM ICD-10 OMIM OMIM

ANALYTE(S):	STAT3 ABCC8 GCK KCNJ11 PDX1 INS
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (gene panel)
- Hyperinsulinism (gene panel)
- Maturity onset Diabete of the Young (MODY), type 5 / Renal cysts and diabetes syndrome (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- ATP binding cassette subfamily C member 8
- glucokinase
- insulin
- potassium inwardly rectifying channel subfamily J member 11
- pancreatic and duodenal homeobox 1
- signal transducer and activator of transcription 3

Related Gene Panels

- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (genepanel) - UZA
- Hyperinsulinism (5 genes) - UZA
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