DISEASE:
17q23.1q23.2 microdeletion syndrome

NAME:	17q23.1q23.2 microdeletion syndrome
DESCRIPTION:	17q23.1q23.2 microdeletion syndrome is a recently described syndrome characterized by developmental delay, microcephaly, short stature, heart defects and limb abnormalities.
ORPHACODE:	261279
SYNONYMS:	Del(17)(q23.1q23.2) Monosomy 17q23.1q23.2
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	TBX4
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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