DISEASE:
8p23.1 microdeletion syndrome

NAME:	8p23.1 microdeletion syndrome
DESCRIPTION:	8p23.1 deletion involves a partial deletion of the short arm of chromosome 8 characterized by low birth weight, postnatal growth deficiency, mild intellectual deficit, hyperactivity, craniofacial abnormalities, and congenital heart defects.
ORPHACODE:	251071
SYNONYMS:	Del(8)(p23.1) Monosomy 8p23.1
XREF(S):	Orphanet ICD-10
ANALYTE(S):	GATA4
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Analytes

- GATA binding protein 4