DISEASE:
2q32q33 microdeletion syndrome

NAME:	2q32q33 microdeletion syndrome
DESCRIPTION:	A rare autosomal monosomy characterized by a variable phenotype with moderate to severe intellectual disability, behavioral problems, short stature, microcephaly, dysplastic nails, sparse hair, cleft palate and dysmorphic craniofacial features.
ORPHACODE:	251019
SYNONYMS:	Del(2)(q32) Del(2)(q32q33) Monosomy 2q32 Monosomy 2q32q33
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	SATB2
CREATED:	13 May 2019 - 01:02
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