DISEASE:
19p13.3 microduplication syndrome

NAME:	19p13.3 microduplication syndrome
DESCRIPTION:	A rare, genetic, syndromic intellectual disability characterized by intrauterine growth retardation, microcephaly, hypotonia, motor and neurodevelopmental delay, speech delay, intellectual disability, and mild dysmorphic features.
ORPHACODE:	447980
SYNONYMS:	Dup(19)(p13.13)
XREF(S):	Orphanet
ANALYTE(S):	NFIX
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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