DISEASE:
17p13.3 microduplication syndrome

NAME:	17p13.3 microduplication syndrome
DESCRIPTION:	17p13.3 microduplication syndrome is characterized by variable psychomotor delay and dysmorphic features.
ORPHACODE:	217385
SYNONYMS:	17p13.3 duplication syndrome Dup(17)(p13.3) Trisomy 17p13.3
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	PAFAH1B1 YWHAE
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Analytes

- platelet activating factor acetylhydrolase 1b regulatory subunit 1
- tyrosine 3-monooxygenase/tryptophan 5-monooxygenase activation protein epsilon