

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):	<u>Orphanet</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>PAFAH1B1</u> <u>YWHAE</u>
CREATED:	13 May 2019 - 01:02
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Source URL: <http://gentest.healthdata.be/disease/1434>

RELATED CONTENT

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

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