DISEASE:
15q13.3 microdeletion syndrome

NAME:	15q13.3 microdeletion syndrome
DESCRIPTION:	15q13.3 microdeletion (microdel15q13.3) syndrome is characterized by a wide spectrum of neurodevelopmental disorders with no or subtle dysmorphic features.
ORPHACODE:	199318
SYNONYMS:	Del(15)(q13.3) Monosomy 15q13.3
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	CHRNA7
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/454

RELATED CONTENT

Related Analytes

- cholinergic receptor nicotinic alpha 7 subunit

Source URL: http://gentest.healthdata.be/disease/454