

DISEASE:
15q14 microdeletion syndrome

NAME:	15q14 microdeletion syndrome
DESCRIPTION:	15q14 microdeletion syndrome is a recently described syndrome characterized by developmental delay, short stature and facial dysmorphism.
ORPHACODE:	261190
SYNONYMS:	Del(15)(q14) Monosomy 15q14
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	MEIS2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/index.php/index.php/disease/1055>

RELATED CONTENT

Related Analytes

- [Meis homeobox 2](#)

Source URL: <http://gentest.healthdata.be/index.php/index.php/disease/1055>