DISEASE:
16q24.3 microdeletion syndrome

NAME:	16q24.3 microdeletion syndrome
DESCRIPTION:	16q24.3 microdeletion syndrome is a recently described syndrome associated with variable developmental delay, facial dysmorphism, seizures and autistic spectrum disorder.
ORPHACODE:	261250
SYNONYMS:	Del(16)(q24.3) Monosomy 16q24.3
XREF(S):	Orphanet ICD-10
ANALYTE(S):	ANKRD11
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Short Stature (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

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- ankyrin repeat domain containing 11

Related Gene Panels

- Short Stature (46 genes) - IPG