

**DISEASE:**  
**19p13.3 microduplication syndrome**

<b>NAME:</b>	19p13.3 microduplication syndrome
<b>DESCRIPTION:</b>	A rare, genetic, syndromic intellectual disability characterized by intrauterine growth retardation, microcephaly, hypotonia, motor and neurodevelopmental delay, speech delay, intellectual disability, and mild dysmorphic features.
<b>ORPHACODE:</b>	447980
<b>SYNOMYS:</b>	Dup(19)(p13.13)
<b>XREF(S):</b>	<a href="#">Orphanet</a>
<b>ANALYTE(S):</b>	<a href="#">NFX1</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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