

**DISEASE:**  
**Pitt-Hopkins-like syndrome**

<b>NAME:</b>	Pitt-Hopkins-like syndrome
<b>DESCRIPTION:</b>	Pitt-Hopkins-like syndrome is a rare, genetic, syndromic intellectual disability disorder characterized by severe intellectual disability, lack of speech with normal, or mildly delayed, motor development, episodic breathing abnormalities, early-onset seizures and facial dysmorphism which only includes a wide mouth. Abnormal sleep-wake cycles, autistic behavior and stereotypic movements are commonly associated.
<b>ORPHACODE:</b>	221150
<b>XREF(S):</b>	<u>Orphanet</u> <u>OMIM</u> <u>OMIM</u>
<b>ANALYTE(S):</b>	<u>CNTNAP2</u> <u>NRXN1</u>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	01 Feb 2020 - 12:07

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