DISEASE:
6q16 microdeletion syndrome

NAME:	6q16 microdeletion syndrome
DESCRIPTION:	A rare Prader-Willi like syndrome due to an interstitial deletion located at 6q16.1q16.2 and characterized by obesity, hyperphagia, hypotonia, small hands and feet, eye/vision anomalies, and global developmental delay.
ORPHACODE:	171829
SYNONYMS:	Del(6)(q16) Monosomy 6q16 Prader-Willi-like syndrome due to microdeletion 6q16
XREF(S):	Orphanet
ANALYTE(S):	SIM1
CREATED:	13 May 2019 - 01:02
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- SIM bHLH transcription factor 1

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