DISEASE:
Prader-Willi syndrome due to paternal deletion of 15q11q13 type 2

NAME:	Prader-Willi syndrome due to paternal deletion of 15q11q13 type 2
ORPHACODE:	177904
XREF(S):	Orphanet
ANALYTE(S):	SNORD115@ SNRPN MAGEL2 NDN OCA2 SNORD116@
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Angelman / Prader Willi Syndrome
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Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Humaine - Erasme ULB
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- MAGE family member L2
- necdin, MAGE family member
- OCA2 melanosomal transmembrane protein
- small nucleolar RNA, C/D box 115 cluster
- small nucleolar RNA, C/D box 116 cluster
- small nuclear ribonucleoprotein polypeptide N