Belgian Genetic Tests database

DISEASE:
Prader-Willi syndrome due to imprinting mutation

NAME:	Prader-Willi syndrome due to imprinting mutation
ORPHACODE:	177910
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SNORD115@ SNRPN MAGEL2 NDN SNORD116@
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/391

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 Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- MAGE family member L2
- necdin, MAGE family member
- small nucleolar RNA, C/D box 115 cluster
- small nucleolar RNA, C/D box 116 cluster

- small nuclear ribonucleoprotein polypeptide N
