Belgian Genetic Tests database

DISEASE:
Angelman syndrome due to imprinting defect in 15q11-q13

NAME:	Angelman syndrome due to imprinting defect in 15q11-q13
ORPHACODE:	411515
XREF(S):	Orphanet
ANALYTE(S):	SNRPN UBE3A ATP10A
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/2324

RELATED CONTENT

Related Genetic Tests

- Angelman / Prader Willi Syndrome
- Angelman / Prader Willi Syndrome
- Angelman / Prader Willi Syndrome
- Angelman / Prader Willi Syndrome
- Angelman / Prader Willi Syndrome
- Angelman / Prader Willi Syndrome
- Angelman / Prader Willi Syndrome

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

- ATPase phospholipid transporting 10A (putative)
- small nuclear ribonucleoprotein polypeptide N
- ubiquitin protein ligase E3A

Source URL: http://gentest.healthdata.be/disease/2324