Belgian Genetic Tests database

DISEASE:
Angelman syndrome due to maternal 15q11q13 deletion

NAME:	Angelman syndrome due to maternal 15q11q13 deletion
ORPHACODE:	98794
SYNONYMS:	Angelman syndrome due to maternal monosomy 15q11q13
XREF(S):	Orphanet ICD-10
ANALYTE(S):	UBE3A OCA2
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/3649

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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

- OCA2 melanosomal transmembrane protein
- ubiquitin protein ligase E3A

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