Belgian Genetic Tests database

DISEASE:
Beckwith-Wiedemann syndrome due to imprinting defect of 11p15

NAME:	Beckwith-Wiedemann syndrome due to imprinting defect of 11p15
ORPHACODE:	231117
XREF(S):	Orphanet ICD-10
ANALYTE(S):	H19 KCNQ1OT1 IGF2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/1579

RELATED CONTENT

Related Genetic Tests

- Beckwith-Wiedemann syndrome
- Beckwith-Wiedemann syndrome (11p15 methylation)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- H19 imprinted maternally expressed transcript
- insulin like growth factor 2
- KCNQ1 opposite strand/antisense transcript 1