Belgian Genetic Tests database

DISEASE:
Beckwith-Wiedemann syndrome due to NSD1 mutation

NAME:	Beckwith-Wiedemann syndrome due to NSD1 mutation
ORPHACODE:	238613
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	NSD1
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/1615

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