Belgian Genetic Tests database

DISEASE:
Neurofibromatosis type 1 due to NF1 mutation or intragenic deletion

NAME:	Neurofibromatosis type 1 due to NF1 mutation or intragenic deletion
ORPHACODE:	363700
SYNONYMS:	Von Recklinghausen disease due to NF1 mutation or intragenic deletion
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	NF1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/1846

RELATED CONTENT

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- Neurofibromatosis type 1 / Legius syndrome

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- Centrum Medische Genetica - UZ Antwerpen
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