DISEASE:
Facioscapulohumeral dystrophy

NAME:	Facioscapulohumeral dystrophy
DESCRIPTION:	A rare neuromuscular disease characterized by progressive muscle weakness with focal involvement of the facial, shoulder and limb muscles.
ORPHACODE:	269
SYNONYMS:	FSH dystrophy FSHD Facioscapulohumeral muscular dystrophy Facioscapulohumeral myopathy Landouzy-Dejerine dystrophy Landouzy-Dejerine myopathy
XREF(S):	Orphanet OMIM OMIM MedDRA ICD-10 OMIM OMIM OMIM

ANALYTE(S):	DNMT3B DUX4 DUX4L1 FRG1 SMCHD1 SMCHD1
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/80

RELATED CONTENT

Related Genetic Tests

- Facioscapulohumeral Muscular Dystrophy 2 (hypomethylation D4Z4 repeats)
- Facioscapulohumeral muscular dystrophy 1A (D4Z4 repeat)
- Facioscapulohumeral muscular dystrophy 1A (D4Z4 repeat)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- DNA methyltransferase 3 beta
- double homeobox 4
- double homeobox 4 like 1 (pseudogene)
- FSHD region gene 1
- structural maintenance of chromosomes flexible hinge domain containing 1