GENETIC TEST:
Facioscapulohumeral Muscular Dystrophy 2 (hypomethylation D4Z4 repeats)

FULL NAME:	Facioscapulohumeral Muscular Dystrophy 2 (hypomethylation D4Z4 repeats)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Methylation analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565390-565401
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	01 Aug 2018 - 11:55
CHANGED:	19 Apr 2022 - 12:18

RELATED CONTENT

Related Diseases

- Facioscapulohumeral dystrophy

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- FSHD region gene 1