GENETIC TEST:
Bethlem myopathy / Ullrich congenital muscular dystrophy / Myosclerosis Myopathy

FULL NAME:	Bethlem myopathy / Ullrich congenital muscular dystrophy / Myosclerosis Myopathy
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
TURNAROUND TIME (MAXIMUM):	8 weeks
CREATED:	29 Jul 2019 - 08:05
CHANGED:	13 Dec 2022 - 22:11

Source URL: http://gentest.healthdata.be/genetic_test/244

RELATED CONTENT

Related Diseases

- Bethlem myopathy
- Congenital muscular dystrophy, Ullrich type
- Myosclerosis
- Primary dystonia, DYT27 type

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- collagen type XII alpha 1 chain
- collagen type VI alpha 1 chain
- collagen type VI alpha 2 chain
- collagen type VI alpha 3 chain

Related Gene Panels

- Bethlem myopathy / Ullrich / Myosclerosis Myopathy - UGent