GENETIC TEST:
Mc Ardle disease, glycogene storage disease type V

FULL NAME:	Mc Ardle disease, glycogene storage disease type V
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
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565471-565482
TURNAROUND TIME (MAXIMUM):	1 month
CREATED:	02 Sep 2019 - 07:49
CHANGED:	02 Jan 2023 - 17:00
URL:	https://www.chu.ulg.ac.be/jcms/c2_23256207/fr/glycogenose-par-deficit-en-glycog…

RELATED CONTENT

Related Diseases

- Glycogen storage disease due to muscle glycogen phosphorylase deficiency

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman

Related Analytes

- glycogen phosphorylase, muscle associated