
GENETIC TEST:
Glycogen storage disease type 9

FULL NAME:	Glycogen storage disease type 9
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:	30 Aug 2019 - 10:00
CHANGED:	02 Jan 2023 - 17:02
URL:	https://www.chu.ulg.ac.be/jcms/c2_23256219/fr/glycogenose-par-deficit-en-phosph...

RELATED CONTENT

Related Diseases

- [Glycogen storage disease due to liver phosphorylase kinase deficiency](#)

Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)

Related Analytes

- [phosphorylase kinase regulatory subunit alpha 2](#)

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