GENETIC TEST:
Carnitine Palmitoyltransferase type II

FULL NAME:	Carnitine Palmitoyltransferase type II
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) Bi-directional Sanger Sequence analysis
RIZIV CODE:	565471-565482
TURNAROUND TIME (MAXIMUM):	unknown
CREATED:	06 Aug 2019 - 11:27
CHANGED:	19 Apr 2022 - 11:32

RELATED CONTENT

Related Diseases

- Carnitine palmitoyl transferase II deficiency, myopathic form
- Carnitine palmitoyl transferase II deficiency, neonatal form
- Carnitine palmitoyl transferase II deficiency, severe infantile form

Related Laboratories

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

Related Analytes

- carnitine palmitoyltransferase 2