GENETIC TEST:
Medium chain acyl-CoA dehydrogenase deficiency (MCAD hot spot mutation - p.Lys329Glu)

FULL NAME:	Medium chain acyl-CoA dehydrogenase deficiency (MCAD hot spot mutation - p.Lys329Glu)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Mutation screening and sequence analysis of selected exons
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565390-565401
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	07 Aug 2019 - 13:54
CHANGED:	21 Apr 2022 - 10:17

RELATED CONTENT

Related Diseases

- Medium chain acyl-CoA dehydrogenase deficiency

Related Laboratories

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

Related Analytes

- acyl-CoA dehydrogenase medium chain