GENETIC TEST:
Amyloidosis hereditary / Dystransthyretinemic hyperthyroxinemia

FULL NAME:	Amyloidosis hereditary / Dystransthyretinemic hyperthyroxinemia
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:	565456-565460
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	06 Aug 2019 - 08:56
CHANGED:	25 Mar 2022 - 08:04

RELATED CONTENT

Related Diseases

- ATTRV122I amyloidosis
- ATTRV30M amyloidosis
- Hereditary ATTR amyloidosis

Related Laboratories

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

Related Analytes

- transthyretin