GENETIC TEST:
Familial hemiplegic Migraine (gene panel)

FULL NAME:	Familial hemiplegic Migraine (gene panel)
DESCRIPTION:	core panel: CACNA1A, ATP1A2, SCN1A full panel: CACNA1A, ATP1A2, SCN1A + ATP1A3, KCNK18, PRRT2, SLC1A3, SLC2A1
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:	565493-565504
TURNAROUND TIME (MAXIMUM):	4 months
CREATED:	19 Jul 2019 - 10:42

CHANGED:	15 Oct 2021 - 13:59
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/15164

Source URL: http://gentest.healthdata.be/genetic_test/83

RELATED CONTENT

Related Diseases

- Familial or sporadic hemiplegic migraine

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ATPase Na+/K+ transporting subunit alpha 2
- ATPase Na+/K+ transporting subunit alpha 3
- calcium voltage-gated channel subunit alpha1 A
- potassium two pore domain channel subfamily K member 18
- proline rich transmembrane protein 2
- sodium voltage-gated channel alpha subunit 1
- solute carrier family 1 member 3
- solute carrier family 2 member 1

Related Gene Panels

- Familial hemiplegic Migraine (8 genes) - KUL