GENETIC TEST:
Homocystinuria (hot spot mutation - c.1298A>C)

FULL NAME:	Homocystinuria (hot spot mutation - c.1298A>C)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565331-565342
EQA:	• MTHFR (Set 01-D)
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	30 Aug 2019 - 11:14
CHANGED:	13 Dec 2022 - 11:53

Source URL: http://gentest.healthdata.be/genetic_test/847

RELATED CONTENT

Related Diseases

- Homocystinuria due to methylene tetrahydrofolate reductase deficiency

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman

Related Analytes

- methylenetetrahydrofolate reductase