GENETIC TEST:
CHARGE syndrome

FULL NAME:	CHARGE syndrome
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
TURNAROUND TIME (MAXIMUM):	20-60 days
CREATED:	12 Dec 2019 - 16:53

CHANGED:	25 Mar 2022 - 11:52

Source URL: http://gentest.healthdata.be/genetic_test/968

RELATED CONTENT

Related Diseases

- CHARGE syndrome

Related Laboratories

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

Related Analytes

- chromodomain helicase DNA binding protein 7