GENETIC TEST:
Frequent hearing deficiency (4 genes)

FULL NAME:	Frequent hearing deficiency (4 genes)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi
METHOD CATEGORY:	Targeted variant analysis Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	PCR based technique Next Generation Sequencing (NGS)
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2021-03-25 / 2024-09-09

EQA:	• Hereditary deafness, • Hereditary deafness, • Hereditary deafness, • Hereditary deafness , • Hereditary deafness, • Hereditary deafness, • Hereditary deafness
TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	29 Jul 2019 - 17:06
CHANGED:	19 Jan 2024 - 09:20
URL:	https://ulbgenetics.be/compendium-des-analyses/#GM23

RELATED CONTENT

Related Diseases

- Rare autosomal recessive non-syndromic sensorineural deafness type DFNB

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- Centre de Génétique Humaine - Erasme ULB

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- gap junction protein beta 6
- otoancorin
- stereocilin