GENETIC TEST:
Mitochondrial non-syndromic sensorineural deafness with susceptibility to aminoglycoside exposure (hot spot mutation - 1555A-G in MT-RNR1)

FULL NAME:	Mitochondrial non-syndromic sensorineural deafness with susceptibility to aminoglycoside exposure (hot spot mutation - 1555A-G in MT-RNR1)
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:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565390-565401
ACCREDITATION (ISO 15189):	2023-11-09 / 2024-05-08
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	06 Aug 2019 - 13:11
CHANGED:	22 Jan 2024 - 14:01

URL:	https://labogidsmedgen.uza.be/analyses/slechthorendheid-mitochondriaal

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- Mitochondrial non-syndromic sensorineural deafness with susceptibility to aminoglycoside exposure

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- mitochondrially encoded 12S rRNA