GENETIC TEST:
Deafness, X-linked

FULL NAME:	Deafness, X-linked
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:	Bi-directional Sanger Sequence analysis MLPA based techniques
RIZIV CODE:	565471-565482
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	06 Aug 2019 - 13:16
CHANGED:	19 Apr 2022 - 11:52

RELATED CONTENT

Related Diseases

- Rare mitochondrial non-syndromic sensorineural deafness
- Xq21 microdeletion syndrome

Related Laboratories

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

Related Analytes

- POU class 3 homeobox 4