DISEASE:
Rare autosomal recessive non-syndromic sensorineural deafness type DFNB

NAME:	Rare autosomal recessive non-syndromic sensorineural deafness type DFNB
ORPHACODE:	90636
SYNONYMS:	Autosomal recessive isolated neurosensory deafness type DFNB Autosomal recessive isolated neurosensory hearing loss type DFNB Autosomal recessive isolated sensorineural deafness type DFNB Autosomal recessive isolated sensorineural hearing loss type DFNB Autosomal recessive non-syndromic neurosensory deafness type DFNB Autosomal recessive non-syndromic neurosensory hearing loss type DFNB Autosomal recessive non-syndromic sensorineural hearing loss type DFNB

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ANALYTE(S):	PPIP5K2 TMEM132E GRAP BDP1 AFG2B CEACAM16 NARS2 ROR1 WBP2 EPS8L2 SLC26A4 BSND CDH23 TECTA TMC1 TMIE TMPRSS3 USH1C COL11A2 ELMOD3 GJB2 GJB3 GJB6 MET MYO15A MYO6 MYO7A OTOF PCDH15 WHRN STRC TRIOBP RDX LHFPL5 ESPN ESRRB MARVELD2 MYO3A SLC26A5

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RELATED CONTENT

Related Genetic Tests

- Corneal dystrophy (gene panel)
- Deafness, autosomal recessive 1A
- Frequent hearing deficiency (4 genes)
- Hearing loss, STRC gene
- Recessive nonsyndromic hearing loss and deafness (2 genes)
- Recessive nonsyndromic hearing loss and deafness DFNB (2 genes)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Humaine - Erasme ULB
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Gent

Related Analytes

- adenylate cyclase 1
- AFG2 AAA ATPase homolog B
- ATPase plasma membrane Ca2+ transporting 2
- B double prime 1, subunit of RNA polymerase III transcription initiation factor IIIB
- barttin CLCNK type accessory subunit beta
- calcium binding protein 2

- cell division cycle 14A
- cadherin related 23
- CEA cell adhesion molecule 16, tectorial membrane component
- calcium and integrin binding family member 2
- claudin 14
- chloride intracellular channel 5
- collagen type XI alpha 2 chain
- doublecortin domain containing 2
- ELMO domain containing 3
- epidermal growth factor receptor pathway substrate 8
- EPS8 like 2
- espin
- estrogen related receptor beta
- GIPC PDZ domain containing family member 3
- gap junction protein alpha 1
- gap junction protein beta 2
- gap junction protein beta 3
- gap junction protein beta 6
- G protein signaling modulator 2
- GRB2 related adaptor protein
- glutaredoxin and cysteine rich domain containing 1
- glutaredoxin and cysteine rich domain containing 2
- hepatocyte growth factor
- immunoglobulin like domain containing receptor 1
- lysyl-tRNA synthetase 1
- LHFPL tetraspan subfamily member 5
- lipoxygenase homology PLAT domains 1
- leucine rich transmembrane and O-methyltransferase domain containing
- MARVEL domain containing 2
- MET proto-oncogene, receptor tyrosine kinase
- membrane integral NOTCH2 associated receptor 2
- myelin protein zero like 2
- methionine sulfoxide reductase B3

- myosin XVA
- myosin IIIA
- myosin VI
- myosin VIIA
- asparaginyl-tRNA synthetase 2, mitochondrial
- otoancorin
- otoferlin
- otogelin
- otogelin like
- protocadherin related 15
- pejvakin
- polyribonucleotide nucleotidyltransferase 1
- diphosphoinositol pentakisphosphate kinase 2
- protein tyrosine phosphatase receptor type Q
- radixin
- RHO family interacting cell polarization regulator 2
- receptor tyrosine kinase like orphan receptor 1
- sphingosine-1-phosphate receptor 2
- serpin family B member 6
- solute carrier family 26 member 4
- solute carrier family 26 member 5
- SLIT and NTRK like family member 6
- stereocilin
- spectrin repeat containing nuclear envelope family member 4
- TBC1 domain family member 24
- tectorin alpha
- transmembrane channel like 1
- transmembrane protein 132E
- transmembrane inner ear
- transmembrane serine protease 3
- taperin
- TRIO and F-actin binding protein
- thrombospondin type laminin G domain and EAR repeats

- USH1 protein network component harmonin
- WW domain binding protein 2
- whirlin

Related Gene Panels

- Corneal dystrophy - UGent
- Non syndromic hearing loss and deafness (2 genes) - IPG - ULG
