DISEASE:
Sensorineural deafness with dilated cardiomyopathy

NAME:	Sensorineural deafness with dilated cardiomyopathy
DESCRIPTION:	Sensorineural deafness with dilated cardiomyopathy is an extremely rare autosomal dominant syndrome described in two families to date and characterized by moderate to severe sensorineural hearing loss manifesting during childhood, and associated with late-onset dilated cardiomyopathy that generally progresses to heart failure.
ORPHACODE:	217622
SYNONYMS:	Neurosensory deafness with dilated cardiomyopathy Neurosensory hearing loss with dilated cardiomyopathy Sensorineural hearing loss with dilated cardiomyopathy
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	EYA4
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- EYA transcriptional coactivator and phosphatase 4

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB