Belgian Genetic Tests database

DISEASE:
Pendred syndrome

NAME:	Pendred syndrome
DESCRIPTION:	A syndromic genetic deafness clinically variable characterized by bilateral sensorineural hearing loss and euthyroid goiter.
ORPHACODE:	705
SYNONYMS:	Goiter-deafness syndrome Goiter-hearing loss syndrome
XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	SLC26A4 KCNJ10 FOXI1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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