DISEASE:
Wolfram syndrome

NAME:	Wolfram syndrome
DESCRIPTION:	A rare, genetic, endocrine disorder characterized by type I diabetes mellitus (DM), diabetes insipidus (DI), sensorineural deafness (D), bilateral optical atrophy (OA) and neurological signs.
ORPHACODE:	3463
SYNONYMS:	DIDMOAD syndrome Diabetes insipidus-diabetes mellitus-optic atrophy-deafness syndrome Diabetes insipidus-diabetes mellitus-optic atrophy-hearing loss syndrome
XREF(S):	Orphanet ICD-10 MeSH OMIM OMIM OMIM
ANALYTE(S):	WFS1 CISD2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/446

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