DISEASE:
Nanophthalmos

NAME:	Nanophthalmos
DESCRIPTION:	A rare ophthalmic disease and a severe form of microphthalmia (small eye phenotype) characterized by a small eye with a short axial length, severe hyperopia, an elevated lens/eye ratio, and a high incidence of angle-closure glaucoma.
ORPHACODE:	35612
SYNONYMS:	Nanophthalmia
XREF(S):	Orphanet OMIM ICD-10 OMIM OMIM OMIM OMIM

ANALYTE(S):	MFRP TMEM98 OTX2 SIX6 RAX ALDH1A3 PRSS56 SOX2 BEST1 CRB1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Glaucoma (gene panel)
- Nanophthalmos

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- aldehyde dehydrogenase 1 family member A3
- bestrophin 1
- crumbs cell polarity complex component 1
- membrane frizzled-related protein
- orthodenticle homeobox 2
- serine protease 56
- retina and anterior neural fold homeobox
- SIX homeobox 6
- SRY-box transcription factor 2
- transmembrane protein 98

Related Gene Panels

- Glaucoma - UGent
- Leber - Retinal, early onset
