DISEASE:
Early-onset non-syndromic cataract

NAME:	Early-onset non-syndromic cataract
DESCRIPTION:	A rare, genetic, non-syndromic developmental defect of the eye disorder, with high clinical and genetic heterogeneity, most frequently characterized by bilateral, symmetrical, non-progressive cataracts which present at birth or in early-childhood. Additional ocular manifestations (e.g. anterior segment dysgenesis, colobomas, nystagmus, microcornea, microphthalmia, myopia) may be associated, however other organs/systems are usually not affected.
ORPHACODE:	91492
XREF(S):	Orphanet ICD-10
CREATED:	29 Jul 2019 - 12:21
CHANGED:	29 Jul 2019 - 12:23

RELATED CONTENT

Related Genetic Tests

- Cataract (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Gene Panels

- Cataract - UGent
- test