DISEASE:
Uveal melanoma

NAME:	Uveal melanoma
DESCRIPTION:	Uveal melanoma is a rare tumor of the eye, arising from the choroid in 90% of cases and from the iris and ciliary body in the other 10% of cases, which clinically presents with visual symptoms (including blurred vision, photopsia, floaters, and visual field reduction), a visible mass and pain. Fatal metastatic disease is seen in about half of all patients, with the liver being the most frequent site of metastasis.
ORPHACODE:	39044
SYNONYMS:	Choroidal melanoma Iris melanoma
XREF(S):	Orphanet MeSH MedDRA OMIM OMIM OMIM ICD-10

ANALYTE(S):	CYSLTR2 BAP1 BAP1 GNAQ GNA11 SF3B1
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RELATED CONTENT

Related Genetic Tests

- Congenital hemangioma (2 genes)
- Hereditary cancer (gene panel)
- Melanoma (6 genes)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- BRCA1 associated protein 1
- cysteinyl leukotriene receptor 2
- G protein subunit alpha 11
- G protein subunit alpha q
- splicing factor 3b subunit 1

Related Gene Panels

- Hereditary predisposition to cancer (47 genes) - IPG
- Melanoma (6 genes) - UCL
