DISEASE:
Adult-onset foveomacular vitelliform dystrophy

NAME:	Adult-onset foveomacular vitelliform dystrophy
DESCRIPTION:	A rare, genetic, macular dystrophy characterized by blurred vision, metamorphopsia and mild visual impairment secondary to a slightly elevated, yellow, egg yolk-like lesion located in the foveal or parafoveal region.
ORPHACODE:	99000
SYNONYMS:	AOFMD AVMD Adult-onset foveomacular dystrophy Adult-onset foveomacular dystrophy with choroidal neovascularization Adult-onset vitelliform macular dystrophy Gass disease Pseudo-Best disease Pseudo-vitelliform macular dystrophy

XREF(S):	Orphanet ICD-10 OMIM OMIM OMIM OMIM
ANALYTE(S):	PRPH2 BEST1 IMPG2 IMPG1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Vitelliform Macular Dystrophy

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- bestrophin 1
- interphotoreceptor matrix proteoglycan 1
- interphotoreceptor matrix proteoglycan 2
- peripherin 2

Related Gene Panels

- Vitelliform Macular Dystrophy (4 genes)