Belgian Genetic Tests database

ANALYTE:
CFB

NAME:	complement factor B
SYMBOL:	CFB
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	H2-Bf
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/3908

RELATED CONTENT

Related Genetic Tests

- Atypical Hemolytic Uremic Syndrome (aHUS) (gene panel)
- Hereditary Hemolytic Anemias due to unknown or doubtful origin (gene panel - 52 genes)
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- Atypical hemolytic uremic syndrome with complement gene abnormality
- Atypical hemolytic-uremic syndrome with B factor anomaly
- NON RARE IN EUROPE: Age-related macular degeneration

Related Gene Panels

- Atypical Hemolytic Uremic Syndrome (aHUS) and Complement disorders (17 genes) - IPG
- Hereditary Hemolytic Anemias (52 genes) - ULB
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent

- Retinal dystrophy - UGent
