ANALYTE:
CFI

NAME:	complement factor I
SYMBOL:	CFI
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	C3b-INA C3b-inactivator FI KAF Konglutinogen-activating factor
XREF(S):	Orphanet OMIM Reactome SwissProt Ensembl Genatlas HGNC
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/1129

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)

Related Diseases

- Atypical hemolytic uremic syndrome with complement gene abnormality
- Atypical hemolytic-uremic syndrome with I factor anomaly
- De novo thrombotic microangiopathy after kidney transplantation
- Familial drusen
- HELLP syndrome
- Immunodeficiency with factor I 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
