Export to PDF

Full name:	Atypical Hemolytic Uremic Syndrome (aHUS) and Complement disorders (17 genes) - IPG
Abbreviation:	aHUS
Type of panel:	Custom panel
Version number:	V7
Laboratory:	Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
Created:	04 Jul 2019 - 08:43
Changed:	11 Dec 2023 - 11:23

Related Diseases

- Atypical hemolytic uremic syndrome with complement gene abnormality
- Atypical hemolytic-uremic syndrome with C3 anomaly
- Atypical hemolytic-uremic syndrome with H factor anomaly
- Complement component 3 deficiency
- Congenital thrombotic thrombocytopenic purpura
- Dense deposit disease
- Immunodeficiency due to a late component of complement deficiency

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ADAMTS13	100.00	1	NM_139027.6
C3	100.00	1	NM_000064.4
C5	100.00	1	NM_001735.3
CD46	100.00	1	NM_172351.3
CFB	100.00	1	NM_001710.6

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
CFH	100.00	1	NM_000186.4
CFHR1	93.00	1	NM_002113.3
CFHR2	91.00	1	NM_005666.4
CFHR3	100.00	1	NM_021023.6
CFHR4	100.00	1	NM_001201550.3
CFHR5	100.00	1	NM_030787.4
CFI	100.00	1	NM_000204.5
DGKE	100.00	1	NM_003647.3
MMACHC	100.00	1	NM_015506.3
PLG	100.00	1	NM_000301.5
THBD	100.00	1	NM_000361.3