

DISEASE:**Atypical hemolytic-uremic syndrome with H factor anomaly**

NAME:	Atypical hemolytic-uremic syndrome with H factor anomaly
ORPHACODE:	93579
SYNONYMS:	Atypical HUS with H factor anomaly D-HUS with H factor anomaly Hemolytic-uremic syndrome without diarrhea with H factor anomaly aHUS with H factor anomaly
XREF(S):	<u>Orphanet</u> <u>OMIM</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>CFH</u>
CREATED:	13 May 2019 - 01:02
CHANGED:	01 Jul 2019 - 06:57

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RELATED CONTENT

Related Genetic Tests

- Atypical Hemolytic Uremic Syndrome (aHUS) (gene panel)
- Cardiomyopathy, hereditary (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- complement factor H

Related Gene Panels

- Atypical Hemolytic Uremic Syndrome (aHUS) and Complement disorders (17 genes) - IPG
- Cardiomyopathy, hereditary (208 genes) - VUB
- Neuromuscular disorders (166 genes) - VUB

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