DISEASE:
Congenital thrombotic thrombocytopenic purpura

NAME:	Congenital thrombotic thrombocytopenic purpura
DESCRIPTION:	A hereditary form of thrombotic thrombocytopenic purpura (TTP) characterized by profound peripheral thrombocytopenia, microangiopathic hemolytic anemia (MAHA) and single or multiple organ failure of variable severity.
ORPHACODE:	93583
SYNONYMS:	Congenital ADAMTS-13 deficiency Congenital TTP Familial TTP Upshaw-Schulman syndrome
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ADAMTS13
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RELATED CONTENT

Related Genetic Tests

- Atypical Hemolytic Uremic Syndrome (aHUS) (gene panel)
- Trombosis - Hemostasis (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ADAM metallopeptidase with thrombospondin type 1 motif 13

Related Gene Panels

- Atypical Hemolytic Uremic Syndrome (aHUS) and Complement disorders (17 genes) - IPG
- Trombosis - Hemostasis (107 genes) - KUL