DISEASE:
Severe hereditary thrombophilia due to congenital protein C deficiency

NAME:	Severe hereditary thrombophilia due to congenital protein C deficiency
DESCRIPTION:	A rare inherited coagulation disorder characterized by deep venous thrombosis symptoms due to reduced synthesis and/or activity levels of protein C.
ORPHACODE:	745
SYNONYMS:	Autosomal recessive thrombophilia due to PC deficiency Autosomal recessive thrombophilia due to congenital protein C deficiency
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	PROC
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Thrombophilia due to protein C deficiency (PROC gene)
- Trombosis - Hemostasis (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- protein C, inactivator of coagulation factors Va and VIIIa

Related Gene Panels

- Trombosis - Hemostasis (107 genes) - KUL