DISEASE:
Familial platelet disorder with associated myeloid malignancy

NAME:	Familial platelet disorder with associated myeloid malignancy
DESCRIPTION:	A rare, genetic, constitutional thrombocytopenia disease characterized by mild to moderate thrombocytopenia, abnormal platelet function and a propensity to develop hematological malignancies, mainly of myeloid origin.
ORPHACODE:	71290
SYNONYMS:	FPD/AML FPDMM FPS/AML Familial platelet disorder with predisposition to acute myelogenous leukemia Familial platelet disorder with predisposition to myeloid malignancy Familial platelet disorder with propensity to acute myeloid leukemia Familial thrombocytopenia with propensity to acute myelogenous leukemia
XREF(S):	Orphanet OMIM ICD-10 OMIM

ANALYTE(S):	ETV6 ANKRD26 RUNX1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest-acc.healthdata.be/disease/2907

RELATED CONTENT

Related Genetic Tests

- Myeloid neoplasms with germline predisposition (Hereditary MDS/Acute Leukemia) (gene panel)
- Trombosis - Hemostasis (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ankyrin repeat domain containing 26
- ETS variant transcription factor 6
- RUNX family transcription factor 1

Related Gene Panels

- Hereditary Myelodysplastic /Acute Leukemia Predisposition Syndromes (gene panel)
- Trombosis - Hemostasis (108 genes) - KUL