

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):	<u>ETV6</u> <u>ANKRD26</u> <u>RUNX1</u>
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