

DISEASE:
Autosomal thrombocytopenia with normal platelets

NAME:	Autosomal thrombocytopenia with normal platelets
ORPHACODE:	168629
XREF(S):	<u>Orphanet</u> <u>ICD-10</u> <u>OMIM</u> <u>OMIM</u> <u>OMIM</u>
ANALYTE(S):	<u>IKZF5</u> <u>MASTL</u> <u>CYCS</u> <u>ANKRD26</u>
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RELATED CONTENT

Related Genetic Tests

- [Myeloid neoplasms with germline predisposition \(Hereditary MDS/Acute Leukemia\) \(gene panel\)](#)
- [Myeloid/lymphoid neoplasms with germline predisposition \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
- [Trombosis - Hemostasis \(gene panel\)](#)
- [« Inherited bone marrow failures syndromes » with or without organ dysfunction](#)

Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)
- [Centrum Menselijke Erfelijkheid - KUL](#)

Related Analytes

- [ankyrin repeat domain containing 26](#)
- [cytochrome c, somatic](#)
- [IKAROS family zinc finger 5](#)
- [microtubule associated serine/threonine kinase like](#)

Related Gene Panels

- [Hematologic Familiar Forms - ULG](#)
- [Hereditary Myelodysplastic /Acute Leukemia Predisposition Syndromes \(gene panel\)](#)

- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Primary immune deficiencies (444 genes) - KUL
- Trombosis - Hemostasis (108 genes) - KUL

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