# DISEASE:
# Wiskott-Aldrich syndrome

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| **NAME:** | Wiskott-Aldrich syndrome |
| **DESCRIPTION:** | A primary immunodeficiency disease characterized by microthrombocytopenia, eczema, infections and an increased risk for autoimmune manifestations and malignancies. |
| **ORPHACODE:** | 906 |
| **SYNONYMS:** | Eczema-thrombocytopenia-immunodeficiency syndrome<br>WAS |
| **XREF(S):** | Orphanet<br>MeSH<br>MedDRA<br>ICD-10<br>OMIM<br>OMIM<br>OMIM |
| **ANALYTE(S):** | WAS<br>WIPF1 |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Primary immune deficiencies (gene panel)
- Trombosis - Hemostasis (gene panel)
- Wiskott-Aldrich syndrome
- « 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

- WASP actin nucleation promoting factor
- WAS/WASL interacting protein family member 1

## Related Gene Panels

- Hematologic Familiar Forms - ULG
- Immunogenetics (21 genes)
- Primary immune deficiencies (444 genes) - KUL
- Trombosis - Hemostasis (108 genes) - KUL

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