Wiskott-Aldrich syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Wiskott-Aldrich syndrome. ORPHA:906

Wiskott-Aldrich syndrome (WAS) is a primary immunodeficiency disease characterized by microthrombocytopenia, eczema, infections and an increased risk for autoimmune manifestations and malignancies.