TEMPI syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. TEMPI syndrome. ORPHA:284227

TEMPI syndrome is a rare multi-systemic disease characterized by the presence of Telangiectasias, Erythrocytosis with elevated erythropoietin levels, Monoclonal gammopathy, Perinephric-fluid collections, and Intrapulmonary shunting.