Familial thrombomodulin anomalies

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial thrombomodulin anomalies. ORPHA:3324

Familial thrombomodulin anomalies is a rare, life-threatening, genetic coagulation disorder characterized by an increased risk of blood clot formation in several members of a family due to a thrombomodulin gene mutation. Patients may manifest with venous thromboembolic disease, premature myocardial infarction and/or arterial thrombosis.