Thrombocytopenia 2

National Cancer Institute

Source

National Cancer Institute. Thrombocytopenia 2. NCI Thesaurus. Code C129035.

An autosomal dominant disorder caused by mutation(s) in the ANKRD26 gene, encoding ANKRD26 protein. Additionally, in one family, a mutation(s) has been identified in the MASTL gene, encoding serine/threonine-protein kinase greatwall. The condition is characterized by mild to moderate bruisability.