Von Willebrand Disease, Type 1

National Cancer Institute

Source

National Cancer Institute. von Willebrand Disease, Type 1. NCI Thesaurus. Code C131685.

An autosomally inherited (generally dominant) coagulation disorder characterized by quantitative partial deficiency of circulating von Willebrand factor (VWF) which account for 60 to 80% of cases of von Willebrand disease. It is characterized by mild to moderate quantitative deficiencies of VWF and factor VIII, which are coordinately reduced from normal plasma levels.