Open Peer Review on Qeios

Bleeding Disorder, Platelet-Type 17

National Cancer Institute

Source

National Cancer Institute. <u>Bleeding Disorder, Platelet-Type 17</u>. NCI Thesaurus. Code C142084.

An autosomal dominant condition caused by mutation(s) in the GFI1B gene, encoding zinc finger protein Gfi-1b. It is characterized by a tendency for increased bleeding due to abnormal platelet function.