

Haematologica
HAEMATOL/2018/214239
Version 2

Glanzmann thrombasthenia: genetic basis and clinical correlates

Juliana Perez Botero, Kristy Lee, Brian R. Branchford, Paul F. Bray,
Kathleen Freson, Michele P. Lambert, Minjie Luo, Shruthi Mohan,
Justyne E. Ross, Wolfgang Bergmeier, Jorge Di Paola, and ClinGen
Platelet Disorder Variant Curation Expert Panel
Collaborative Groups: ClinGen Platelet Disorder Variant Curation
Expert Panel)

Disclosures: The ClinGen Expert Curation Panel on Platelet Disorders is supported by the American Society of Hematology (ASH). ClinGen is primarily funded by the National Human Genome Research Institute (NHGRI), through the following three grants: U41HG006834 (Rehm), U41HG009649 (Bustamante), and U41HG009650 (Berg). ClinGen also receives support for content curation from the Eunice Kennedy Shriver National Institute of Child Health and Human Development (NICHD), through the following three grants: U24HD093483, U24HD093486, and U24HD093487

Contributions: JPB, KL, BRB, PFB, KF, MPL, ML, SM, JER, WB, JDP wrote the manuscript