Marijke Bryckaert

List of Publications by Year
in descending order

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11 papers	475 citations	1040018 9 h-index	1281846 11 g-index
11 all docs	11 docs citations	11 times ranked	661 citing authors

#	Article	IF	Citations
1	New insights into regulation of Î±IIbÎ²3 integrin signaling by filamin A. Research and Practice in Thrombosis and Haemostasis, 2022, 6, e12672.	2.3	2
2	Filamin A: key actor in platelet biology. Blood, 2019, 134, 1279-1288.	1.4	62
3	Disrupted filamin A/Î±IIbÎ²3 interaction induces macrothrombocytopenia by increasing RhoA activity. Blood, 2019, 133, 1778-1788.	1.4	27
4	A mutation of the human EPHB2 gene leads to a major platelet functional defect. Blood, 2018, 132, 2067-2077.	1.4	17
5	Protein kinase C signaling dysfunction in von Willebrand disease (p.V1316M) type 2B platelets. Blood Advances, 2018, 2, 1417-1428.	5.2	9
6	Gain-of-Function Mutation in Filamin A Potentiates Platelet Integrin Î± _{IIb} Î² ₃ Activation. Arteriosclerosis, Thrombosis, and Vascular Biology, 2017, 37, 1087-1097.	2.4	28
7	A genetically-engineered von Willebrand disease type 2B mouse model displays defects in hemostasis and inflammation. Scientific Reports, 2016, 6, 26306.	3.3	19
8	A new form of macrothrombocytopenia induced by a germ-line mutation in the PRKACG gene. Blood, 2014, 124, 2554-2563.	1.4	69
9	Heterogeneity of Platelet Functional Alterations in Patients With Filamin A Mutations. Arteriosclerosis, Thrombosis, and Vascular Biology, 2013, 33, e11-8.	2.4	52
10	von Willebrand factor mutation promotes thrombocytopathy by inhibiting integrin Î±IIbÎ²3. Journal of Clinical Investigation, 2013, 123, 5071-5081.	8.2	42
11	Thrombocytopenia resulting from mutations in filamin A can be expressed as an isolated syndrome. Blood, 2011, 118, 5928-5937.	1.4	148