

Maaïke G J M Van Bergen

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2320739/publications.pdf>

Version: 2024-02-01

10
papers

143
citations

1478505

6
h-index

1588992

8
g-index

10
all docs

10
docs citations

10
times ranked

354
citing authors

#	ARTICLE	IF	CITATIONS
1	Neutrophil specific granule and NETosis defects in gray platelet syndrome. <i>Blood Advances</i> , 2021, 5, 549-564.	5.2	18
2	Specific proteome changes in platelets from individuals with GATA1-, GF11B-, and RUNX1-linked bleeding disorders. <i>Blood</i> , 2021, 138, 86-90.	1.4	7
3	Characterization of a genomic region 8Åkb downstream of GF11B associated with myeloproliferative neoplasms. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2021, 1867, 166259.	3.8	0
4	Platelet CD34 expression in a patient with a partial deletion of transcription factor subunit CBFβ. <i>American Journal of Hematology</i> , 2020, 95, E136-E139.	4.1	1
5	Targeting the GF11/1Bâ€”CoREST Complex in Acute Myeloid Leukemia. <i>Frontiers in Oncology</i> , 2019, 9, 1027.	2.8	21
6	CBFÎ²-MYH11 interferes with megakaryocyte differentiation via modulating a gene program that includes GATA2 and KLF1. <i>Blood Cancer Journal</i> , 2019, 9, 33.	6.2	7
7	Inherited missense variants that affect GF11B function do not necessarily cause bleeding diatheses. <i>Haematologica</i> , 2019, 104, e260-e264.	3.5	7
8	Molecular mechanisms of bleeding disorder associated GF11B ^{Q287*} mutation and its affected pathways in megakaryocytes and platelets. <i>Haematologica</i> , 2019, 104, 1460-1472.	3.5	21
9	Platelet CD34 Expression and a Congenital Collar Bone Malformation Associated with a Partial CBFβ Deletion in a Case with a Bleeding Disorder. <i>Blood</i> , 2019, 134, 1077-1077.	1.4	0
10	Association of a Haplotype in the <i>NR3C2</i> Gene, Encoding the Mineralocorticoid Receptor, With Chronic Central Serous Chorioretinopathy. <i>JAMA Ophthalmology</i> , 2017, 135, 446.	2.5	61