

Juvenal Paiva

List of Publications by Year in descending order

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Version: 2024-02-01

8
papers

47
citations

2258059

3
h-index

1720034

7
g-index

8
all docs

8
docs citations

8
times ranked

67
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of p.W246L As a Novel Mutation in the GP1BA Gene Responsible for Platelet-Type von Willebrand Disease. <i>Seminars in Thrombosis and Hemostasis</i> , 2014, 40, 151-160.	2.7	18
2	Type 2A and 2M von Willebrand Disease: Differences in Phenotypic Parameters According to the Affected Domain by Disease-Causing Variants and Assessment of Pathophysiological Mechanisms. <i>Seminars in Thrombosis and Hemostasis</i> , 2021, 47, 862-874.	2.7	10
3	Phenotypic Parameters in Genotypically Selected Type 2B von Willebrand Disease Patients: A Large, Single-Center Experience Including a New Novel Mutation. <i>Seminars in Thrombosis and Hemostasis</i> , 2017, 43, 092-100.	2.7	7
4	Diagnosis of von Willebrand disease in Argentina: a single institution experience. <i>Annals of Blood</i> , 0, 2, 22-22.	0.4	4
5	Thrombotic microangiopathies: First report of 294 cases from a single institution experience in Argentina. <i>EJHaem</i> , 2021, 2, 149-156.	1.0	3
6	Combined effects of two mutations in von Willebrand disease 2M phenotype. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2018, 2, 162-167.	2.3	2
7	Type 2N von Willebrand disease: Is it always a recessive trait?. <i>Thrombosis Research</i> , 2021, 198, 49-51.	1.7	2
8	Von Willebrand disease type 2M: Correlation between genotype and phenotype: Comment from Woods et al.. <i>Journal of Thrombosis and Haemostasis</i> , 2022, 20, 1022-1023.	3.8	1