Pascual Marco

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4029762/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Demographic and clinical data in acquired hemophilia A: results from the European Acquired Haemophilia Registry (EACH2). Journal of Thrombosis and Haemostasis, 2012, 10, 622-631.	1.9	395
2	Management of bleeding in acquired hemophilia A: results from the European Acquired Haemophilia (EACH2) Registry. Blood, 2012, 120, 39-46.	0.6	326
3	Immunosuppression for acquired hemophilia A: results from the European Acquired Haemophilia Registry (EACH2). Blood, 2012, 120, 47-55.	0.6	284
4	Pregnancyâ€associated acquired haemophilia A: results from the European Acquired Haemophilia (EACH2) registry. BJOG: an International Journal of Obstetrics and Gynaecology, 2012, 119, 1529-1537.	1.1	96
5	Premature myocardial infarction: Clinical profile and angiographic findings. International Journal of Cardiology, 2008, 126, 127-129.	0.8	67
6	Hypofibrinolysis in atrial fibrillation. American Heart Journal, 1998, 136, 956-960.	1.2	59
7	Premature coronary artery disease in young (age < 45) subjects: Interactions of lipid profile, thrombophilic and haemostatic markers. International Journal of Cardiology, 2009, 136, 222-225.	0.8	56
8	Molecular and clinical profile of von Willebrand disease in Spain (PCM–EVW–ES): Proposal for a new diagnostic paradigm. Thrombosis and Haemostasis, 2016, 115, 40-50.	1.8	36
9	Molecular and clinical profile of von Willebrand disease in Spain (PCM-EVW-ES): comprehensive genetic analysis by next-generation sequencing of 480 patients. Haematologica, 2017, 102, 2005-2014.	1.7	35
10	Von Willebrand factor and ADAMTS13 activity as clinical severity markers in patients with COVID-19. Journal of Thrombosis and Thrombolysis, 2021, 52, 497-503.	1.0	20
11	Effects of Body Mass Index on the Lipid Profile and Biomarkers of Inflammation and a Fibrinolytic and Prothrombotic State. Journal of Atherosclerosis and Thrombosis, 2015, 22, 610-617.	0.9	19
12	Management of acquired hemophilia A: results from the Spanish registry. Blood Advances, 2021, 5, 3821-3829.	2.5	18
13	The Unravelling of the Genetic Architecture of Plasminogen Deficiency and its Relation to Thrombotic Disease. Scientific Reports, 2016, 6, 39255.	1.6	15
14	Unraveling the effect of silent, intronic and missense mutations on <i>VWF</i> splicing: contribution of next generation sequencing in the study of mRNA. Haematologica, 2019, 104, 587-598.	1.7	13
15	Identification of 58 Mutations (26 Novel) in 94 of 109 Symptomatic Spanish Probands with Protein C Deficiency. Thrombosis and Haemostasis, 2019, 119, 1409-1418.	1.8	8
16	Multiparameter Flow Cytometry Identification of Neoplastic Subclones: A New Biomarker in Monoclonal Gammopathy of Undetermined Significance and Multiple Myeloma. Acta Haematologica, 2019, 141, 1-6.	0.7	8
17	Acquired Haemophilia: Clinical and Demographic Data.Results of European Acquired Haemophilia Registry (EACH2) Blood, 2010, 116, 1398-1398.	0.6	8
18	Role of multimeric analysis of von Willebrand factor (VWF) in von Willebrand disease (VWD) diagnosis: Lessons from the PCM-EVW-ES Spanish project. PLoS ONE, 2018, 13, e0197876.	1.1	6

PASCUAL MARCO

#	Article	IF	CITATIONS
19	Diurnal Variation in the Intensity of Anticoagulation in Atrial Fibrillation. Stroke, 2002, 33, 322-324.	1.0	6
20	Measurement of procoagulant activity of microparticles in plasma: Feasibility of new functional assays. Thrombosis Research, 2014, 134, 1363-1364.	0.8	5
21	Management of Bleeding In Acquired Hemophilia: Results of the European Acquired Hemophilia Registry (EACH2). Blood, 2010, 116, 716-716.	0.6	5
22	Acquired Haemophilia A and Pregnancy/Postpartum – a Report From a European Registry. Blood, 2010, 116, 717-717.	0.6	5
23	Identification of new markers of recurrence in patients with unprovoked deep vein thrombosis by gene expression profiling: the retro study. European Journal of Haematology, 2016, 97, 128-136.	1.1	2
24	Unraveling the Influence of Common von Willebrand factor variants on von Willebrand Disease Phenotype: An Exploratory Study on the Molecular and Clinical Profile of von Willebrand Disease in Spain Cohort. Thrombosis and Haemostasis, 2020, 120, 437-448.	1.8	2
25	Dealing with refractoriness in obstetric primary antiphospholipid syndrome—often not a matter of success. Lupus, 2014, 23, 964-965.	0.8	1
26	Involvement of antifactor VIII autoantibodies specificity in the outcome of inhibitor eradication therapies in acquired hemophilia a patients. Blood Coagulation and Fibrinolysis, 2019, 30, 127-132.	0.5	1
27	Autoimmune Acquired Factor XIII Deficiency: A Case Report. Journal of Blood Medicine, 2021, Volume 12, 63-68.	0.7	1
28	Upshaw-Schulman Syndrome: Novel homozygous missense mutation. Thrombosis Research, 2017, 158, 83-85.	0.8	0
29	Improvement in the cardiovascular profile of patients with morbid obesity following bariatric surgery. Medicine (United States), 2021, 100, e25280.	0.4	0