

Marzia Menegatti

List of Publications by Year in descending order

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Version: 2024-02-01

55
papers

2,029
citations

279701

23
h-index

243529

44
g-index

60
all docs

60
docs citations

60
times ranked

1806
citing authors

#	ARTICLE	IF	CITATIONS
1	Coagulation factor activity and clinical bleeding severity in rare bleeding disorders: results from the European Network of Rare Bleeding Disorders. <i>Journal of Thrombosis and Haemostasis</i> , 2012, 10, 615-621.	1.9	362
2	No Evidence of Association Between Prothrombotic Gene Polymorphisms and the Development of Acute Myocardial Infarction at a Young Age. <i>Circulation</i> , 2003, 107, 1117-1122.	1.6	191
3	Factor X Deficiency. <i>Seminars in Thrombosis and Hemostasis</i> , 2009, 35, 407-415.	1.5	125
4	Genetic diagnosis of haemophilia and other inherited bleeding disorders. <i>Haemophilia</i> , 2006, 12, 82-89.	1.0	123
5	Introduction: Rare Bleeding Disorders: General Aspects of Clinical Features, Diagnosis, and Management. <i>Seminars in Thrombosis and Hemostasis</i> , 2009, 35, 349-355.	1.5	123
6	Treatment of rare factor deficiencies other than hemophilia. <i>Blood</i> , 2019, 133, 415-424.	0.6	92
7	Rare bleeding disorders. <i>Haemophilia</i> , 2006, 12, 137-142.	1.0	76
8	Gynecological and obstetrical manifestations of inherited bleeding disorders in women. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 236-245.	1.9	64
9	Gene mutations and three-dimensional structural analysis in 13 families with severe factor X deficiency. <i>British Journal of Haematology</i> , 2002, 117, 685-692.	1.2	61
10	Rare Bleeding Disorders: Worldwide Efforts for Classification, Diagnosis, and Management. <i>Seminars in Thrombosis and Hemostasis</i> , 2013, 39, 579-584.	1.5	58
11	Treatment of rare factor deficiencies in 2016. <i>Hematology American Society of Hematology Education Program</i> , 2016, 2016, 663-669.	0.9	53
12	Phenotype and genotype report on homozygous and heterozygous patients with congenital factor X deficiency. <i>Haematologica</i> , 2008, 93, 934-938.	1.7	47
13	Phenotype-genotype characterization of 10 families with severe a subunit factor XIII deficiency. <i>Human Mutation</i> , 2004, 23, 98-98.	1.1	45
14	Relatively Poor Performance of Clinical Laboratories for DNA Analyses in the Detection of Two Thrombophilic Mutations – A Cause for Concern. <i>Thrombosis and Haemostasis</i> , 2002, 88, 690-691.	1.8	41
15	Exploring the global landscape of genetic variation in coagulation factor XI deficiency. <i>Blood</i> , 2017, 130, e1-e6.	0.6	41
16	Oxidative stress is increased in primary and post-polycythemia vera myelofibrosis. <i>Experimental Hematology</i> , 2010, 38, 1058-1065.	0.2	39
17	Efficacy of prophylaxis and genotype-phenotype correlation in patients with severe Factor X deficiency in Iran. <i>Haemophilia</i> , 2012, 18, 211-215.	1.0	34
18	Minimal factor XIII activity level to prevent major spontaneous bleeds. <i>Journal of Thrombosis and Haemostasis</i> , 2017, 15, 1728-1736.	1.9	34

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19	Clinical and molecular characterisation of 21 patients affected by quantitative fibrinogen deficiency. <i>Thrombosis and Haemostasis</i> , 2015, 113, 567-576.	1.8	33
20	Identification of genetic risk variants for deep vein thrombosis by multiplexed next-generation sequencing of 186 hemostatic/pro-inflammatory genes. <i>BMC Medical Genomics</i> , 2012, 5, 7.	0.7	32
21	Reduced fibrinolytic resistance in patients with factor XI deficiency. Evidence of a thrombin-independent impairment of the thrombin-activatable fibrinolysis inhibitor pathway. <i>Journal of Thrombosis and Haemostasis</i> , 2016, 14, 1603-1614.	1.9	27
22	A rare inherited coagulation disorder: Combined homozygous factor VII and factor X deficiency. <i>American Journal of Hematology</i> , 2004, 77, 90-91.	2.0	26
23	Retrospective evaluation of bleeding tendency and simultaneous thrombin and plasmin generation in patients with rare bleeding disorders. <i>Haemophilia</i> , 2012, 18, 630-638.	1.0	26
24	Establishment of a bleeding score as a diagnostic tool for patients with rare bleeding disorders. <i>Thrombosis Research</i> , 2016, 148, 128-134.	0.8	22
25	Genome editing of factor X in zebrafish reveals unexpected tolerance of severe defects in the common pathway. <i>Blood</i> , 2017, 130, 666-676.	0.6	22
26	Analysis of factor V in zebrafish demonstrates minimal levels needed for early hemostasis. <i>Blood Advances</i> , 2019, 3, 1670-1680.	2.5	18
27	Management of rare acquired bleeding disorders. <i>Hematology American Society of Hematology Education Program</i> , 2019, 2019, 80-87.	0.9	15
28	Genotype and phenotype report on patients with combined deficiency of factor V and factor VIII in Iran. <i>Blood Coagulation and Fibrinolysis</i> , 2014, 25, 360-363.	0.5	14
29	Performance of Clinical Laboratories for DNA Analyses to Detect Thrombophilia Mutations. <i>Clinical Chemistry</i> , 2005, 51, 1310-1311.	1.5	13
30	Potential misdiagnosis of dysfibrinogenemia: Data from multicentre studies amongst UK NEQAS and PRO-BDD project laboratories. <i>International Journal of Laboratory Hematology</i> , 2017, 39, 653-662.	0.7	13
31	An international registry of patients with plasminogen deficiency (HISTORY). <i>Haematologica</i> , 2020, 105, 554-561.	1.7	13
32	Molecular characterization of an Italian patient with plasminogen deficiency and ligneous conjunctivitis. <i>Blood Coagulation and Fibrinolysis</i> , 2007, 18, 81-84.	0.5	12
33	Post-partum hemorrhage in women with rare bleeding disorders. <i>Thrombosis Research</i> , 2011, 127, S116-S119.	0.8	12
34	Arg77His and Trp187Arg are the Most Common Mutations Causing FXIII Deficiency in Iran. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2012, 18, 100-103.	0.7	12
35	Identification of Four Novel Polymorphisms in the α_1 and β Fibrinogen Genes and Analysis of Association with Plasma Levels of the Protein. <i>Thrombosis Research</i> , 2001, 103, 299-307.	0.8	11
36	Profiling the mutational landscape of coagulation factor V deficiency. <i>Haematologica</i> , 2020, 105, e180-e185.	1.7	10

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37	Thrombin generation in patients with idiopathic sudden sensorineural hearing loss. <i>Thrombosis Research</i> , 2014, 133, 1130-1134.	0.8	9
38	Clinical and laboratory diagnosis of rare coagulation disorders (RCDs). <i>Thrombosis Research</i> , 2020, 196, 603-608.	0.8	9
39	Rare variants lowering the levels of coagulation factor X are protective against ischemic heart disease. <i>Haematologica</i> , 2020, 105, e365-e369.	1.7	9
40	A recurrent Gly43Asp substitution in coagulation Factor X rigidifies its catalytic pocket and impairs catalytic activity and intracellular trafficking. <i>Thrombosis Research</i> , 2014, 133, 481-487.	0.8	8
41	Frequency of the p.Gly262Asp mutation in congenital Factor X deficiency. <i>European Journal of Clinical Investigation</i> , 2015, 45, 1087-1091.	1.7	7
42	Detection of Factor XIII deficiency: data from multicentre exercises amongst UK NEQAS and PRO-RBDD project laboratories. <i>International Journal of Laboratory Hematology</i> , 2017, 39, 350-358.	0.7	6
43	Molecular investigation of 41 patients affected by coagulation factor XI deficiency. <i>Haemophilia</i> , 2018, 24, e50-e55.	1.0	6
44	Understanding the Impact of Aberrant Splicing in Coagulation Factor V Deficiency. <i>International Journal of Molecular Sciences</i> , 2019, 20, 910.	1.8	5
45	Effect of prothrombin 19911 A>G polymorphism on the risk of cerebral sinus venous thrombosis. <i>European Journal of Neurology</i> , 2010, 17, 1482-1485.	1.7	4
46	Neonatal onset of congenital factor X deficiency. <i>Blood Coagulation and Fibrinolysis</i> , 2015, 26, 679-681.	0.5	4
47	Disseminated intravascular coagulation with positive D-dimer. <i>Blood Coagulation and Fibrinolysis</i> , 2016, 27, 933-935.	0.5	4
48	A homozygous duplication of the <i>FGG</i> exon 8-intron 8 junction causes congenital afibrinogenemia. Lessons learned from the study of a large consanguineous Turkish family. <i>Haematologica</i> , 2022, 107, 1064-1071.	1.7	3
49	Effect of adenosine derivatives on in vitro thrombus formation induced by shear stress. <i>Haematologica</i> , 1999, 84, 721-5.	1.7	3
50	Safety and effectiveness of recombinant factor XIII in congenital factor XIII deficiency: Real world evidence. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022, 6, e12628.	1.0	3
51	Minimal factor XIII activity level to prevent major spontaneous bleeds: reply. <i>Journal of Thrombosis and Haemostasis</i> , 2017, 15, 2280-2282.	1.9	2
52	Recombinant factor XIII subunit in a patient with factor XIII deficiency and recurrent pregnancy loss. <i>Journal of Thrombosis and Haemostasis</i> , 2018, 16, 1052-1054.	1.9	2
53	Inherited Bleeding Disorders in Pregnancy: Rare Coagulation Factor Defects. , 2015, , 209-221.		0
54	Inherited Bleeding Disorders in Pregnancy: Rare Coagulation Factor Defects. , 2012, , 131-141.		0

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55	Prospective Evaluation of Bleeding Incidence in Fibrinogen Deficiency (PRO-RBDD Study). Blood, 2016, 128, 207-207.	0.6	0