

Federica Melazzini

List of Publications by Year in descending order

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Version: 2024-02-01

36
papers

1,654
citations

430442

18
h-index

395343

33
g-index

38
all docs

38
docs citations

38
times ranked

1696
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in ANKRD26 are responsible for a frequent form of inherited thrombocytopenia: analysis of 78 patients from 21 families. <i>Blood</i> , 2011, 117, 6673-6680.	0.6	263
2	New roles for mean platelet volume measurement in the clinical practice?. <i>Platelets</i> , 2016, 27, 607-612.	1.1	164
3	Eltrombopag for the treatment of the inherited thrombocytopenia deriving from MYH9 mutations. <i>Blood</i> , 2010, 116, 5832-5837.	0.6	141
4	Platelet diameters in inherited thrombocytopenias: analysis of 376 patients with all known disorders. <i>Blood</i> , 2014, 124, e4-e10.	0.6	112
5	Clinical and laboratory features of 103 patients from 42 Italian families with inherited thrombocytopenia derived from the monoallelic Ala156Val mutation of GPIbA (Bolzano mutation). <i>Haematologica</i> , 2012, 97, 82-88.	1.7	99
6	Clinical and pathogenic features of <i>ETV6</i> -related thrombocytopenia with predisposition to acute lymphoblastic leukemia. <i>Haematologica</i> , 2016, 101, 1333-1342.	1.7	92
7	Bleeding risk of surgery and its prevention in patients with inherited platelet disorders. <i>Haematologica</i> , 2017, 102, 1192-1203.	1.7	92
8	Clinical and genetic aspects of Bernard-Soulier syndrome: searching for genotype/phenotype correlations. <i>Haematologica</i> , 2011, 96, 417-423.	1.7	90
9	Platelet size distinguishes between inherited macrothrombocytopenias and immune thrombocytopenia. <i>Journal of Thrombosis and Haemostasis</i> , 2009, 7, 2131-2136.	1.9	86
10	Platelet size for distinguishing between inherited thrombocytopenias and immune thrombocytopenia: a multicentric, real life study. <i>British Journal of Haematology</i> , 2013, 162, 112-119.	1.2	86
11	Inherited thrombocytopenias—recent advances in clinical and molecular aspects. <i>Platelets</i> , 2017, 28, 3-13.	1.1	51
12	Thrombopoietin mutation in congenital amegakaryocytic thrombocytopenia treatable with romiplostim. <i>EMBO Molecular Medicine</i> , 2018, 10, 63-75.	3.3	47
13	Peptic Ulcer Disease as a Common Cause of Bleeding in Patients with Coronavirus Disease 2019. <i>American Journal of Gastroenterology</i> , 2020, 115, 1139-1140.	0.2	43
14	Impact of COVID-19 on liver function: results from an internal medicine unit in Northern Italy. <i>Internal and Emergency Medicine</i> , 2020, 15, 1399-1407.	1.0	37
15	Analysis of 65 pregnancies in 34 women with five different forms of inherited platelet function disorders. <i>British Journal of Haematology</i> , 2015, 170, 559-563.	1.2	35
16	5'UTR point substitutions and N-terminal truncating mutations of ANKRD26 in acute myeloid leukemia. <i>Journal of Hematology and Oncology</i> , 2017, 10, 18.	6.9	33
17	Depletion of circulating IgM memory B cells predicts unfavourable outcome in COVID-19. <i>Scientific Reports</i> , 2020, 10, 20836.	1.6	32
18	A new form of inherited thrombocytopenia due to monoallelic loss of function mutation in the thrombopoietin gene. <i>British Journal of Haematology</i> , 2018, 181, 698-701.	1.2	21

#	ARTICLE	IF	CITATIONS
19	Personalized reference intervals for platelet count reduce the number of subjects with unexplained thrombocytopenia. <i>Haematologica</i> , 2015, 100, e338-e340.	1.7	19
20	Bleeding is not the main clinical issue in many patients with inherited thrombocytopaenias. <i>Haemophilia</i> , 2017, 23, 673-681.	1.0	19
21	The ISTH bleeding assessment tool as predictor of bleeding events in inherited platelet disorders: Communication from the ISTH SSC Subcommittee on Platelet Physiology. <i>Journal of Thrombosis and Haemostasis</i> , 2021, 19, 1364-1371.	1.9	19
22	<i>ACTN1</i> mutations lead to a benign form of platelet macrocytosis not always associated with thrombocytopenia. <i>British Journal of Haematology</i> , 2018, 183, 276-288.	1.2	16
23	Mutations of <i>RUNX1</i> in families with inherited thrombocytopenia. <i>American Journal of Hematology</i> , 2017, 92, E86-E88.	2.0	15
24	Extramedullary hematopoiesis: a new feature of inherited thrombocytopenias?. <i>Journal of Thrombosis and Haemostasis</i> , 2017, 15, 2226-2229.	1.9	8
25	Venous thromboembolism and COVID-19: a single center experience from an academic tertiary referral hospital of Northern Italy. <i>Internal and Emergency Medicine</i> , 2021, 16, 1141-1152.	1.0	8
26	Antithrombotic prophylaxis for surgery-associated venous thromboembolism risk in patients with inherited platelet disorders. The SPATA-DVT Study. <i>Haematologica</i> , 2020, 105, 1948-1956.	1.7	7
27	Massive mediastinal enlargement due to extramedullary haematopoiesis in a patient with MYH9-related thrombocytopenia. <i>British Journal of Haematology</i> , 2017, 178, 10-10.	1.2	5
28	Diagnostic Delay of Pulmonary Embolism in COVID-19 Patients. <i>Frontiers in Medicine</i> , 2021, 8, 637375.	1.2	4
29	COVID-19-related symptom clustering in a primary care vs internal medicine setting. <i>Internal and Emergency Medicine</i> , 2021, , 1.	1.0	3
30	Venous thromboembolism in chronic gastrointestinal disorders. <i>Expert Review of Gastroenterology and Hepatology</i> , 2022, 16, 437-448.	1.4	3
31	Research at the heart of hematology: thrombocytopenias and platelet function disorders. <i>Haematologica</i> , 2017, 102, 203-205.	1.7	2
32	Dysregulation of oncogenic factors by GFI1B p32: investigation of a novel <i>GFI1B</i> germline mutation. <i>Haematologica</i> , 2022, 107, 260-267.	1.7	1
33	Eltrombopag for the Treatment of the Inherited Thrombocytopenia Deriving From MYH9 Mutations. <i>Blood</i> , 2010, 116, 2533-2533.	0.6	1
34	Letter. <i>Neurosurgery</i> , 2016, 78, E895-E896.	0.6	0
35	Inherited Thrombocytopenias. , 2017, , 727-747.		0
36	Clinical and Laboratory Features of 103 Patients From 42 Italian Families with Inherited Thrombocytopenia Derived From the Monoallelic Ala156Val Mutation of GPIb Alpha (Bolzano) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 57 T		