Federica Melazzini

List of Publications by Year
in descending order

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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	Dysregulation of oncogenic factors by GFI1B p32: investigation of a novel <I>GFI1B</I> germline mutation. Haematologica, 2022, 107, 260-267.	1.7	1
2	Venous thromboembolism in chronic gastrointestinal disorders. Expert Review of Gastroenterology and Hepatology, 2022, 16, 437-448.	1.4	3
3	Venous thromboembolism and COVID-19: a single center experience from an academic tertiary referral hospital of Northern Italy. Internal and Emergency Medicine, 2021, 16, 1141-1152.	1.0	8
4	The ISTH bleeding assessment tool as predictor of bleeding events in inherited platelet disorders: Communication from the ISTH SSC Subcommittee on Platelet Physiology. Journal of Thrombosis and Haemostasis, 2021, 19, 1364-1371.	1.9	19
5	Diagnostic Delay of Pulmonary Embolism in COVID-19 Patients. Frontiers in Medicine, 2021, 8, 637375.	1.2	4
6	COVID-19-related symptom clustering in a primary care vs internal medicine setting. Internal and Emergency Medicine, 2021, , 1.	1.0	3
7	Impact of COVID-19 on liver function: results from an internal medicine unit in Northern Italy. Internal and Emergency Medicine, 2020, 15, 1399-1407.	1.0	37
8	Depletion of circulating IgM memory B cells predicts unfavourable outcome in COVID-19. Scientific Reports, 2020, 10, 20836.	1.6	32
9	Antithrombotic prophylaxis for surgery-associated venous thromboembolism risk in patients with inherited platelet disorders. The SPATA-DVT Study. Haematologica, 2020, 105, 1948-1956.	1.7	7
10	Peptic Ulcer Disease as a Common Cause of Bleeding in Patients with Coronavirus Disease 2019. American Journal of Gastroenterology, 2020, 115, 1139-1140.	0.2	43
11	A new form of inherited thrombocytopenia due to monoallelic loss of function mutation in the thrombopoietin gene. British Journal of Haematology, 2018, 181, 698-701.	1.2	21
12	Thrombopoietin mutation in congenital amegakaryocytic thrombocytopenia treatable withÂromiplostim. EMBO Molecular Medicine, 2018, 10, 63-75.	3.3	47
13	<scp>ACTN</scp>1 mutations lead to a benign form of platelet macrocytosis not always associated with thrombocytopenia. British Journal of Haematology, 2018, 183, 276-288.	1.2	16
14	Inherited thrombocytopeniasâ€"recent advances in clinical and molecular aspects. Platelets, 2017, 28, 3-13.	1.1	51
15	Research at the heart of hematology: thrombocytopenias and platelet function disorders. Haematologica, 2017, 102, 203-205.	1.7	2
16	Mutations of RUNX1 in families with inherited thrombocytopenia. American Journal of Hematology, 2017, 92, E86-E88.	2.0	15
17	Bleeding risk of surgery and its prevention in patients with inherited platelet disorders. Haematologica, 2017, 102, 1192-1203.	1.7	92
18	Inherited Thrombocytopenias. , 2017, , 727-747.		0

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19	Massive mediastinal enlargement due to extramedullary haematopoiesis in a patient with MYH9 -related thrombocytopenia. British Journal of Haematology, 2017, 178, 10-10.	1.2	5
20	Bleeding is not the main clinical issue in many patients with inherited thrombocytopaenias. Haemophilia, 2017, 23, 673-681.	1.0	19
21	Extramedullary hematopoiesis: a new feature of inherited thrombocytopenias?. Journal of Thrombosis and Haemostasis, 2017, 15, 2226-2229.	1.9	8
22	5â€™UTR point substitutions and N-terminal truncating mutations of ANKRD26 in acute myeloid leukemia. Journal of Hematology and Oncology, 2017, 10, 18.	6.9	33
23	Letter. Neurosurgery, 2016, 78, E895-E896.	0.6	0
24	Clinical and pathogenic features of ETV6 -related thrombocytopenia with predisposition to acute lymphoblastic leukemia. Haematologica, 2016, 101, 1333-1342.	1.7	92
25	New roles for mean platelet volume measurement in the clinical practice?. Platelets, 2016, 27, 607-612.	1.1	164
26	Analysis of 65 pregnancies in 34 women with five different forms of inherited platelet function disorders. British Journal of Haematology, 2015, 170, 559-563.	1.2	35
27	Personalized reference intervals for platelet count reduce the number of subjects with unexplained thrombocytopenia. Haematologica, 2015, 100, e338-e340.	1.7	19
28	Platelet diameters in inherited thrombocytopenias: analysis of 376 patients with all known disorders. Blood, 2014, 124, e4-e10.	0.6	112
29	Platelet size for distinguishing between inherited thrombocytopenias and immune thrombocytopenia: a multicentric, real life study. British Journal of Haematology, 2013, 162, 112-119.	1.2	86
30	Clinical and laboratory features of 103 patients from 42 Italian families with inherited thrombocytopenia derived from the monoallelic Ala156Val mutation of GPIbÂ (Bolzano mutation). Haematologica, 2012, 97, 82-88.	1.7	99
31	Mutations in ANKRD26 are responsible for a frequent form of inherited thrombocytopenia: analysis of 78 patients from 21 families. Blood, 2011, 117, 6673-6680.	0.6	263
32	Clinical and genetic aspects of Bernard-Soulier syndrome: searching for genotype/phenotype correlations. Haematologica, 2011, 96, 417-423.	1.7	90
33	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 ETQq1 1 0.784314 rgBT /Overlock 10	0.6	0
34	Eltrombopag for the treatment of the inherited thrombocytopenia deriving from MYH9 mutations. Blood, 2010, 116, 5832-5837.	0.6	141
35	Eltrombopag for the Treatment of the Inherited Thrombocytopenia Deriving From MYH9 Mutations. Blood, 2010, 116, 2533-2533.	0.6	1
36	Platelet size distinguishes between inherited macrothrombocytopenias and immune thrombocytopenia. Journal of Thrombosis and Haemostasis, 2009, 7, 2131-2136.	1.9	86