Jose Maria Bastida

List of Publications by Year
in descending order

55	941	430874 18	477307 29
papers	citations	h-index	g-index

55	55	55	1391
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Introducing high-throughput sequencing into mainstream genetic diagnosis practice in inherited platelet disorders. Haematologica, 2018, 103, 148-162.	3.5	96
2	Novel mutations in RASGRP2, which encodes CalDAG-GEFI, abrogate Rap1 activation, causing platelet dysfunction. Blood, 2016, 128, 1282-1289.	1.4	68
3	Defects in memory B-cell and plasma cell subsets expressing different immunoglobulin-subclasses in patients with CVID and immunoglobulin subclass deficiencies. Journal of Allergy and Clinical Immunology, 2019, 144, 809-824.	2.9	55
4	Inherited Platelet Disorders: An Updated Overview. International Journal of Molecular Sciences, 2021, 22, 4521.	4.1	44
5	Design and application of a 23â€gene panel by nextâ€generation sequencing for inherited coagulation bleeding disorders. Haemophilia, 2016, 22, 590-597.	2.1	43
6	Influence of donor age in allogeneic stem cell transplant outcome in acute myeloid leukemia and myelodisplastic syndrome. Leukemia Research, 2015, 39, 828-834.	0.8	41
7	Eltrombopag in immune thrombocytopenia: efficacy review and update on drug safety. Therapeutic Advances in Drug Safety, 2018, 9, 263-285.	2.4	40
8	Thromboprophylaxis in multiple myeloma patients treated with lenalidomide â€“ A systematic review. Thrombosis Research, 2016, 141, 84-90.	1.7	36
9	Application of a molecular diagnostic algorithm for haemophilia A and B using next-generation sequencing of entire F8, F9 and VWF genes. Thrombosis and Haemostasis, 2017, 117, 66-74.	3.4	36
10	Two novel variants of the ABCG5 gene cause xanthelasmas and macrothrombocytopenia: a brief review of hematologic abnormalities of sitosterolemia. Journal of Thrombosis and Haemostasis, 2017, 15, 1859-1866.	3.8	34
11	Thrombopoietin Receptor Agonists for Severe Thrombocytopenia after Allogeneic Stem Cell Transplantation: Experience of the Spanish Group of Hematopoietic Stem Cell Transplant. Biology of Blood and Marrow Transplantation, 2019, 25, 1825-1831.	2.0	34
12	Molecular Diagnosis of Inherited Coagulation and Bleeding Disorders. Seminars in Thrombosis and Hemostasis, 2019, 45, 695-707.	2.7	32
13	Use of eltrombopag for secondary immune thrombocytopenia in clinical practice. British Journal of Haematology, 2017, 178, 959-970.	2.5	30
14	Usefulness of eltrombopag for treating thrombocytopenia after allogeneic stem cell transplantation. Bone Marrow Transplantation, 2019, 54, 757-761.	2.4	27
15	Identification of two novel mutations in RASGRP2 affecting platelet CalDAG-GEFI expression and function in patients with bleeding diathesis. Platelets, 2018, 29, 192-195.	2.3	26
16	CRISPR/Cas9-generated models uncover therapeutic vulnerabilities of del(11q) CLL cells to dual BCR and PARP inhibition. Leukemia, 2020, 34, 1599-1612.	7.2	21
17	GoldVariants, a resource for sharing rare genetic variants detected in bleeding, thrombotic, and platelet disorders: Communication from the ISTH SSC Subcommittee on Genomics in Thrombosis and Hemostasis. Journal of Thrombosis and Haemostasis, 2021, 19, 2612-2617.	3.8	21
18	Sitosterolemia: Diagnosis, Metabolic and Hematological Abnormalities, Cardiovascular Disease and Management. Current Medicinal Chemistry, 2019, 26, 6766-6775.	2.4	21

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19	Valores de referencia de parÃ¡metros de rigidez arterial y su relaciÃ³n con los factores de riesgo cardiovascular en poblaciÃ³n espaÃ±ola. Estudio EVA. Revista Espanola De Cardiologia, 2020, 73, 43-52.	1.2	20
20	Wiskottâ€“Aldrich syndrome in a child presenting with macrothrombocytopenia. Platelets, 2017, 28, 417-420.	2.3	19
21	Management of acquired hemophilia A: results from the Spanish registry. Blood Advances, 2021, 5, 3821-3829.	5.2	18
22	Management patterns and outcomes in symptomatic venous thromboembolism following allogeneic hematopoietic stem cell transplantation. A 15-years experience at a single center. Thrombosis Research, 2016, 142, 52-56.	1.7	12
23	RASGRP2 gene variations associated with platelet dysfunction and bleeding. Platelets, 2019, 30, 535-539.	2.3	12
24	Role of Thrombopoietin Receptor Agonists in Inherited Thrombocytopenia. International Journal of Molecular Sciences, 2021, 22, 4330.	4.1	12
25	Biological significance of monoallelic and biallelic BIRC3 loss in del(11q) chronic lymphocytic leukemia progression. Blood Cancer Journal, 2021, 11, 127.	6.2	12
26	Expanding the genetic spectrum of TUBB1-related thrombocytopenia. Blood Advances, 2021, 5, 5453-5467.	5.2	12
27	Identification of novel variants in ten patients with Hermansky-Pudlak syndrome by high-throughput sequencing. Annals of Medicine, 2019, 51, 141-148.	3.8	11
28	ABCG5 and ABCG8 gene variations associated with sitosterolemia and platelet dysfunction. Platelets, 2020, 32, 1-5.	2.3	11
29	Low molecular weight heparin is useful in adult COVID-19 inpatients. Experience during the first Spanish wave: observational study. Sao Paulo Medical Journal, 2021, , .	0.9	11
30	Transcriptomic analysis of patients with immune thrombocytopenia treated with eltrombopag. Platelets, 2020, 31, 993-1000.	2.3	10
31	Hemorrhagic pericardial effusion as the debut of acquired hemophilia in a chronic lymphocytic leukemia patient. Medicine (United States), 2017, 96, e8669.	1.0	8
32	Thrombocytopenia and Therapeutic Strategies after Allogeneic Hematopoietic Stem Cell Transplantation. Journal of Clinical Medicine, 2022, 11, 1364.	2.4	7
33	Novel Therapies to Address Unmet Needs in ITP. Pharmaceuticals, 2022, 15, 779.	3.8	7
34	Nuevos mÃ©todos diagnÃ³sticos en los trastornos plaquetarios hereditarios. Medicina ClÃ¬nica, 2017, 148, 71-77.	0.6	6
35	PTGS1 gene variations associated with bleeding and platelet dysfunction. Platelets, 2021, 32, 710-716.	2.3	6
36	Hidden myelodysplastic syndrome (MDS): A prospective study to confirm or exclude MDS in patients with anemia of uncertain etiology. International Journal of Laboratory Hematology, 2019, 41, 109-117.	1.3	5

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37	Characterization of the Platelet Phenotype Caused by a Germline RUNX1 Variant in a CRISPR/Cas9-Generated Murine Model. Thrombosis and Haemostasis, 2021, 121, 1193-1205.	3.4	5
38	GENYOi005-A: An induced pluripotent stem cells (iPSCs) line generated from a patient with Familial Platelet Disorder with associated Myeloid Malignancy (FPDMM) carrying a p.Thr196Ala variant. Stem Cell Research, 2019, 41, 101603.	0.7	4
39	HLA specificities are associated with prognosis in IGHV-mutated CLL-like high-count monoclonal B cell lymphocytosis. PLoS ONE, 2017, 12, e0172978.	2.5	4
40	Novel approaches for diagnosing inherited platelet disorders. Medicina ClÃnica (English Edition), 2017, 148, 71-77.	0.2	3
41	A novel nonsense variant in TPM4 caused dominant macrothrombocytopenia, mild bleeding tendency and disrupted cytoskeleton remodeling. Journal of Thrombosis and Haemostasis, 2022, 20, 1248-1255.	3.8	3
42	Src-related thrombocytopenia: a fine line between a megakaryocyte dysfunction and an immune-mediated disease. Blood Advances, 2022, 6, 5244-5255.	5.2	3
43	Functional Alterations Involved in Increased Bleeding in Hereditary Hemorrhagic Telangiectasia Mouse Models. Frontiers in Medicine, 2022, 9, .	2.6	3
44	Topâ€level sport in athletes with severe haemophilia A. Haemophilia, 2017, 23, e391-e393.	2.1	2
45	A Modern Approach to the Molecular Diagnosis of Inherited Bleeding Disorders. Journal of Molecular and Genetic Medicine: an International Journal of Biomedical Research, 2018, 12, .	0.1	2
46	A novel genetic variant in <scp>PTGS1</scp> affects Nâ€glycosylation of cyclooxygenaseâ€1 causing a dominantâ€negative effect on platelet function and bleeding diathesis. American Journal of Hematology, 2021, 96, E83-E88.	4.1	2
47	Design and Validate of Next-Generation Sequencing Panel for Inherited Platelet Disorders. Blood, 2014, 124, 4210-4210.	1.4	2
48	Acquired Haemophilia A in Association with Influenza A and Urinary Tract Infection. European Journal of Case Reports in Internal Medicine, 2019, 7, 001678.	0.4	2
49	Bleeding Complications Associated with Pregnancy with Primary Immune Thrombocytopenia: A Meta-Analysis. TH Open, 2022, 06, e230-e237.	1.4	1
50	Effect of sarpogrelate treatment on 5-HT modulation of vascular sympathetic innervation and platelet activity in diabetic rats. Biomedicine and Pharmacotherapy, 2022, 153, 113276.	5.6	1
51	Protocolo de prÃ¡ctica asistencial de las linfocitosis crÃ³nicas. Medicine, 2016, 12, 1245-1248.	0.0	0
52	Protocolo de prÃ¡ctica asistencial de las poliadenopatÃas crÃ³nicas. Medicine, 2016, 12, 1249-1252.	0.0	0
53	PD-1 and PD-L1 Are Overexpressed in the "Intermediate CD14+CD16+" and "Non Classical CD14lowCD16+" but Not in the "Classical CD14+CD16-" Monocytes in the Peripheral Blood of Chronic Myelomonocytic Leukemia. Blood, 2015, 126, 1694-1694.	1.4	0
54	Impact of Treatment on Overall Survival (OS) in Higher-Risk Myelodysplastic Syndromes (MDS): A Report from the Erasme Study. Blood, 2015, 126, 5240-5240.	1.4	0

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55	Identification By Whole Exome Sequencing of the Molecular Defect in a Novel Gene Related to Glycosylation in Two Unrelated Families with Syndromic Macrothrombocytopenia. Blood, 2021, 138, 588-588.	1.4	0