# 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 | 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 |
| 2 | Src-related thrombocytopenia: a fine line between a megakaryocyte dysfunction and an immune-mediated disease. Blood Advances, 2022, 6, 5244-5255. | 5.2 | 3 |
| 3 | Thrombocytopenia and Therapeutic Strategies after Allogeneic Hematopoietic Stem Cell Transplantation. Journal of Clinical Medicine, 2022, 11, 1364. | 2.4 | 7 |
| 4 | Bleeding Complications Associated with Pregnancy with Primary Immune Thrombocytopenia: A Meta-Analysis. TH Open, 2022, 06, e230-e237. | 1.4 | 1 |
| 5 | Functional Alterations Involved in Increased Bleeding in Hereditary Hemorrhagic Telangiectasia Mouse Models. Frontiers in Medicine, 2022, 9, . | 2.6 | 3 |
| 6 | Novel Therapies to Address Unmet Needs in ITP. Pharmaceuticals, 2022, 15, 779. | 3.8 | 7 |
| 7 | 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 |
| 8 | PTGS1 gene variations associated with bleeding and platelet dysfunction. Platelets, 2021, 32, 710-716. | 2.3 | 6 |
| 9 | 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 |
| 10 | Role of Thrombopoietin Receptor Agonists in Inherited Thrombocytopenia. International Journal of Molecular Sciences, 2021, 22, 4330. | 4.1 | 12 |
| 11 | Inherited Platelet Disorders: An Updated Overview. International Journal of Molecular Sciences, 2021, 22, 4521. | 4.1 | 44 |
| 12 | Biological significance of monoallelic and biallelic BIRC3 loss in del(11q) chronic lymphocytic leukemia progression. Blood Cancer Journal, 2021, 11, 127. | 6.2 | 12 |
| 13 | 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 |
| 14 | 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 |
| 15 | Management of acquired hemophilia A: results from the Spanish registry. Blood Advances, 2021, 5, 3821-3829. | 5.2 | 18 |
| 16 | Expanding the genetic spectrum of TUBB1-related thrombocytopenia. Blood Advances, 2021, 5, 5453-5467. | 5.2 | 12 |
| 17 | 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 |
| 18 | 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 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 | Transcriptomic analysis of patients with immune thrombocytopenia treated with eltrombopag. Platelets, 2020, 31, 993-1000. | 2.3 | 10 |
| 21 | ABCG5 and ABCG8 gene variations associated with sitosterolemia and platelet dysfunction. Platelets, 2020, 32, 1-5. | 2.3 | 11 |
| 22 | 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 |
| 23 | 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 |
| 24 | 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 |
| 25 | 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 |
| 26 | Molecular Diagnosis of Inherited Coagulation and Bleeding Disorders. Seminars in Thrombosis and Hemostasis, 2019, 45, 695-707. | 2.7 | 32 |
| 27 | RASGRP2 gene variations associated with platelet dysfunction and bleeding. Platelets, 2019, 30, 535-539. | 2.3 | 12 |
| 28 | 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 |
| 29 | Usefulness of eltrombopag for treating thrombocytopenia after allogeneic stem cell transplantation. Bone Marrow Transplantation, 2019, 54, 757-761. | 2.4 | 27 |
| 30 | 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 |
| 31 | Sitosterolemia: Diagnosis, Metabolic and Hematological Abnormalities, Cardiovascular Disease and Management. Current Medicinal Chemistry, 2019, 26, 6766-6775. | 2.4 | 21 |
| 32 | 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 |
| 33 | Introducing high-throughput sequencing into mainstream genetic diagnosis practice in inherited platelet disorders. Haematologica, 2018, 103, 148-162. | 3.5 | 96 |
| 34 | 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 |
| 35 | 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 |
| 36 | Eltrombopag in immune thrombocytopenia: efficacy review and update on drug safety. Therapeutic Advances in Drug Safety, 2018, 9, 263-285. | 2.4 | 40 |

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|---|---|---|---|
| 37 | Nuevos mÃ©todos diagnÃ³sticos en los trastornos plaquetarios hereditarios. Medicina ClÃ¡nica, 2017, 148, 71-77. | 0.6 | 6 |
| 38 | Novel approaches for diagnosing inherited platelet disorders. Medicina ClÃ¡nica (English Edition), 2017, 148, 71-77. | 0.2 | 3 |
| 39 | Topâ€level sport in athletes with severe haemophilia A. Haemophilia, 2017, 23, e391-e393. | 2.1 | 2 |
| 40 | Use of eltrombopag for secondary immune thrombocytopenia in clinical practice. British Journal of Haematology, 2017, 178, 959-970. | 2.5 | 30 |
| 41 | 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 |
| 42 | Hemorrhagic pericardial effusion as the debut of acquired hemophilia in a chronic lymphocytic leukemia patient. Medicine (United States), 2017, 96, e8669. | 1.0 | 8 |
| 43 | Wiskottâ€“Aldrich syndrome in a child presenting with macrothrombocytopenia. Platelets, 2017, 28, 417-420. | 2.3 | 19 |
| 44 | 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 |
| 45 | 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 |
| 46 | 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 |
| 47 | Protocolo de prÃ¡ctica asistencial de las linfocitosis crÃ³nicas. Medicine, 2016, 12, 1245-1248. | 0.0 | 0 |
| 48 | Protocolo de prÃ¡ctica asistencial de las poliadenopatÃas crÃ³nicas. Medicine, 2016, 12, 1249-1252. | 0.0 | 0 |
| 49 | Thromboprophylaxis in multiple myeloma patients treated with lenalidomide â€“ A systematic review. Thrombosis Research, 2016, 141, 84-90. | 1.7 | 36 |
| 50 | Novel mutations in RASGRP2, which encodes CalDAG-GEFI, abrogate Rap1 activation, causing platelet dysfunction. Blood, 2016, 128, 1282-1289. | 1.4 | 68 |
| 51 | 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 |
| 52 | 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 |
| 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 | Design and Validate of Next-Generation Sequencing Panel for Inherited Platelet Disorders. Blood, 2014, 124, 4210-4210. | 1.4 | 2 |