Joanne Yacobovich

List of Publications by Year
in descending order

51	1,001	394421	434195
papers	citations	19 h-index	31 g-index

51	51	51	2007
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	An improved index for diagnosis and mortality prediction in malignancy-associated hemophagocytic lymphohistiocytosis. Blood, 2022, 139, 1098-1110.	1.4	46
2	Congenital Thrombocytopenia Associated with a Heterozygous Variant in the MEIS1 Gene Encoding a Transcription Factor Essential for Megakaryopoiesis. Platelets, 2022, , 1-4.	2.3	0
3	Syndromes predisposing to leukemia are a major cause of inherited cytopenias in children. Haematologica, 2022, 107, 2081-2095.	3.5	5
4	Pediatric immune thrombocytopenia: apoptotic markers may help in predicting the disease course. Pediatric Research, 2021, 90, 93-98.	2.3	2
5	Bone pain at leukemia diagnosis and other risk factors for symptomatic osteonecrosis in children with acute lymphoblastic leukemia. Pediatric Blood and Cancer, 2021, 68, e29033.	1.5	3
6	Improved transplant outcomes with myeloablative conditioning for hemophagocytic lymphohistiocytosis in HLA-matched and mismatched donors: a national multicenter retrospective study. Bone Marrow Transplantation, 2021, 56, 2088-2096.	2.4	5
7	Pediatric myelodysplastic syndrome with inflammatory manifestations: Diagnosis, genetics, treatment, and outcome. Pediatric Blood and Cancer, 2021, 68, e29138.	1.5	4
8	A novel index using inflammatory markers improves the diagnosis of hemophagocytic lymphohistiocytosis in patients with hematologic malignancies.. Journal of Clinical Oncology, 2021, 39, 7563-7563.	1.6	1
9	Mutations in RASGRP2 gene identified in patients misdiagnosed as Glanzmann thrombasthenia patients. Blood Cells, Molecules, and Diseases, 2021, 89, 102560.	1.4	4
10	Incorporation of somatic panels for the detection of haematopoietic transformation in children and young adults with leukaemia predisposition syndromes and with acquired cytopenias. British Journal of Haematology, 2021, 193, 570-580.	2.5	9
11	Primary autoimmune myelofibrosis: A case report in a child. EJHaem, 2020, 1, 304-308.	1.0	1
12	Alpha-Thalassemia Carrier due to â€“Î±^{3.7} Deletion: Not So Silent. Acta Haematologica, 2020, 143, 432-437.	1.4	4
13	Splenectomy in childhood for nonâ€malignant haematologic disorders â€“ longâ€term followâ€up shows minimal adverse effects. British Journal of Haematology, 2020, 190, 909-915.	2.5	10
14	Serum soluble CD25 and ferritin in distinguishing patients with uncomplicated hematologic malignancies from patients with hematologic malignancies complicated by hemophagocytic lymphohistiocytosis.. Journal of Clinical Oncology, 2020, 38, e20072-e20072.	1.6	0
15	A Novel Inflammatory Index Is Sufficient to Identify Hemophagocytic Lymphohistiocytosis in Adult Patients with Hematologic Malignancies. Blood, 2020, 136, 1-2.	1.4	0
16	Vox Sanguinis International Forum on paediatric indications for blood component transfusion. Vox Sanguinis, 2019, 114, e36-e90.	1.5	0
17	Splenectomy and emerging novel treatments in rare inherited hemolytic anemias. HemaSphere, 2019, 3, 160-162.	2.7	0
18	Thrombophilia Testing in High Pediatric Migraine Risk Children With Migraine. Journal of Child Neurology, 2019, 34, 121-127.	1.4	0

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19	Hematologic Adverse Events Associated With Prolonged Valganciclovir Treatment in Congenital Cytomegalovirus Infection. Pediatric Infectious Disease Journal, 2019, 38, 127-130.	2.0	20
20	Labile plasma iron as an indicator of patient adherence to iron chelation treatment. Blood Cells, Molecules, and Diseases, 2018, 71, 1-4.	1.4	1
21	Targeted next generation sequencing for the diagnosis of patients with rare congenital anemias. European Journal of Haematology, 2018, 101, 297-304.	2.2	27
22	Evaluating platelet function disorders in children with bleeding tendency â€“ A single center study. Platelets, 2017, 28, 676-681.	2.3	7
23	Molecular diagnosis of Î±â€thalassemia in a multiethnic population. European Journal of Haematology, 2017, 98, 553-562.	2.2	19
24	Quantitation of bleeding symptoms in a national registry of patients with inherited platelet disorders. Blood Cells, Molecules, and Diseases, 2017, 67, 59-62.	1.4	3
25	Distal limb anomalies in patients with congenital dyserythropoietic anemia. American Journal of Medical Genetics, Part A, 2017, 173, 487-490.	1.2	5
26	Neonatal haemostasis. Hamostaseologie, 2016, 36, 261-264.	1.9	2
27	Refractory autoimmune disease: an overview of when first-line therapy is not enough. Seminars in Hematology, 2016, 53, S35-S38.	3.4	1
28	A high-throughput sequencing test for diagnosing inherited bleeding, thrombotic, and platelet disorders. Blood, 2016, 127, 2791-2803.	1.4	157
29	Anti-D treatment for pediatric immune thrombocytopenia: Is the bad reputation justified?. Seminars in Hematology, 2016, 53, S64-S66.	3.4	3
30	Association between renal cystic lesions and bilateral Wilmsâ€™ tumours. European Radiology, 2016, 26, 1665-1669.	4.5	2
31	Early-onset Evans syndrome, immunodeficiency, and premature immunosenescence associated with tripeptidyl-peptidase II deficiency. Blood, 2015, 125, 753-761.	1.4	66
32	Inferior vena cava (IVC) filters in children: A 10â€year single center experience. Pediatric Blood and Cancer, 2015, 62, 1974-1978.	1.5	12
33	Infections caused by Fusobacterium in children: a 14-year single-center experience. Infection, 2015, 43, 663-670.	4.7	20
34	Patient and central venous catheter related risk factors for blood stream infections in children receiving chemotherapy. Pediatric Blood and Cancer, 2015, 62, 471-476.	1.5	20
35	Autoimmune lymphoproliferative syndrome-like disease in patients with LRBA mutation. Clinical Immunology, 2015, 159, 84-92.	3.2	96
36	Genetic analysis and clinical picture of severe congenital neutropenia in Israel. Pediatric Blood and Cancer, 2015, 62, 103-108.	1.5	20

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37	Thalassemia Major and Sickle Cell Disease in Adolescents and Young Adults. Acta Haematologica, 2014, 132, 340-347.	1.4	11
38	Pediatric otogenic sigmoid sinus thrombosis: 12-Year experience. International Journal of Pediatric Otorhinolaryngology, 2014, 78, 930-933.	1.0	26
39	Extremity and caval deep venous thrombosis. , 2014, , 1-15.		1
40	The use of DDAVP in children with bleeding disorders. Pediatric Blood and Cancer, 2013, 60, S41-3.	1.5	14
41	Underdiagnosed Menorrhagia in Adolescents is Associated with Underdiagnosed Anemia. Journal of Pediatrics, 2012, 160, 468-472.	1.8	25
42	The conundrum of neonatal coagulopathy. Hematology American Society of Hematology Education Program, 2012, 2012, 450-4.	2.5	22
43	Diagnosis and management of central-line-associated thrombosis in newborns and infants. Seminars in Fetal and Neonatal Medicine, 2011, 16, 340-344.	2.3	43
44	Postoperative thrombotic thrombocytopenic purpura in an infant: case report and literature review. Journal of Pediatric Surgery, 2011, 46, 764-766.	1.6	8
45	Further delineation of the phenotype of severe congenital neutropenia type 4 due to mutations in G6PC3. European Journal of Human Genetics, 2011, 19, 18-22.	2.8	50
46	Clinical and Laboratory Assessment of the Bleeding Pediatric Patient. Seminars in Thrombosis and Hemostasis, 2011, 37, 756-762.	2.7	13
47	Acquired Proximal Renal Tubular Dysfunction in Î²-Thalassemia Patients Treated With Deferasirox. Journal of Pediatric Hematology/Oncology, 2010, 32, 564-567.	0.6	50
48	Frequency and natural history of inherited bone marrow failure syndromes: the Israeli Inherited Bone Marrow Failure Registry. Haematologica, 2010, 95, 1300-1307.	3.5	57
49	Postâ€thrombotic syndrome after central venous catheter removal in childhood cancer survivors is associated with a history of obstruction. Pediatric Blood and Cancer, 2010, 55, 153-156.	1.5	19
50	Thrombophilia in children with venous thromboembolic disease. Thrombosis Research, 2006, 118, 59-65.	1.7	33
51	Central Venous Line-Related Thrombosis in Children. Acta Haematologica, 2006, 115, 201-206.	1.4	54