

Joanne Yacobovich

List of Publications by Year in descending order

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Version: 2024-02-01

51
papers

1,001
citations

394421

19
h-index

434195

31
g-index

51
all docs

51
docs citations

51
times ranked

2007
citing authors

#	ARTICLE	IF	CITATIONS
1	A high-throughput sequencing test for diagnosing inherited bleeding, thrombotic, and platelet disorders. <i>Blood</i> , 2016, 127, 2791-2803.	1.4	157
2	Autoimmune lymphoproliferative syndrome-like disease in patients with LRBA mutation. <i>Clinical Immunology</i> , 2015, 159, 84-92.	3.2	96
3	Early-onset Evans syndrome, immunodeficiency, and premature immunosenescence associated with tripeptidyl-peptidase II deficiency. <i>Blood</i> , 2015, 125, 753-761.	1.4	66
4	Frequency and natural history of inherited bone marrow failure syndromes: the Israeli Inherited Bone Marrow Failure Registry. <i>Haematologica</i> , 2010, 95, 1300-1307.	3.5	57
5	Central Venous Line-Related Thrombosis in Children. <i>Acta Haematologica</i> , 2006, 115, 201-206.	1.4	54
6	Acquired Proximal Renal Tubular Dysfunction in β^2 -Thalassemia Patients Treated With Deferasirox. <i>Journal of Pediatric Hematology/Oncology</i> , 2010, 32, 564-567.	0.6	50
7	Further delineation of the phenotype of severe congenital neutropenia type 4 due to mutations in G6PC3. <i>European Journal of Human Genetics</i> , 2011, 19, 18-22.	2.8	50
8	An improved index for diagnosis and mortality prediction in malignancy-associated hemophagocytic lymphohistiocytosis. <i>Blood</i> , 2022, 139, 1098-1110.	1.4	46
9	Diagnosis and management of central-line-associated thrombosis in newborns and infants. <i>Seminars in Fetal and Neonatal Medicine</i> , 2011, 16, 340-344.	2.3	43
10	Thrombophilia in children with venous thromboembolic disease. <i>Thrombosis Research</i> , 2006, 118, 59-65.	1.7	33
11	Targeted next generation sequencing for the diagnosis of patients with rare congenital anemias. <i>European Journal of Haematology</i> , 2018, 101, 297-304.	2.2	27
12	Pediatric otogenic sigmoid sinus thrombosis: 12-Year experience. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2014, 78, 930-933.	1.0	26
13	Underdiagnosed Menorrhagia in Adolescents is Associated with Underdiagnosed Anemia. <i>Journal of Pediatrics</i> , 2012, 160, 468-472.	1.8	25
14	The conundrum of neonatal coagulopathy. <i>Hematology American Society of Hematology Education Program</i> , 2012, 2012, 450-4.	2.5	22
15	Infections caused by <i>Fusobacterium</i> in children: a 14-year single-center experience. <i>Infection</i> , 2015, 43, 663-670.	4.7	20
16	Patient and central venous catheter related risk factors for blood stream infections in children receiving chemotherapy. <i>Pediatric Blood and Cancer</i> , 2015, 62, 471-476.	1.5	20
17	Genetic analysis and clinical picture of severe congenital neutropenia in Israel. <i>Pediatric Blood and Cancer</i> , 2015, 62, 103-108.	1.5	20
18	Hematologic Adverse Events Associated With Prolonged Valganciclovir Treatment in Congenital Cytomegalovirus Infection. <i>Pediatric Infectious Disease Journal</i> , 2019, 38, 127-130.	2.0	20

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19	Post-thrombotic syndrome after central venous catheter removal in childhood cancer survivors is associated with a history of obstruction. <i>Pediatric Blood and Cancer</i> , 2010, 55, 153-156.	1.5	19
20	Molecular diagnosis of α -thalassemia in a multiethnic population. <i>European Journal of Haematology</i> , 2017, 98, 553-562.	2.2	19
21	The use of DDAVP in children with bleeding disorders. <i>Pediatric Blood and Cancer</i> , 2013, 60, S41-3.	1.5	14
22	Clinical and Laboratory Assessment of the Bleeding Pediatric Patient. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 756-762.	2.7	13
23	Inferior vena cava (IVC) filters in children: A 10-year single center experience. <i>Pediatric Blood and Cancer</i> , 2015, 62, 1974-1978.	1.5	12
24	Thalassemia Major and Sickle Cell Disease in Adolescents and Young Adults. <i>Acta Haematologica</i> , 2014, 132, 340-347.	1.4	11
25	Splenectomy in childhood for non-malignant haematologic disorders – long-term follow-up shows minimal adverse effects. <i>British Journal of Haematology</i> , 2020, 190, 909-915.	2.5	10
26	Incorporation of somatic panels for the detection of haematopoietic transformation in children and young adults with leukaemia predisposition syndromes and with acquired cytopenias. <i>British Journal of Haematology</i> , 2021, 193, 570-580.	2.5	9
27	Postoperative thrombotic thrombocytopenic purpura in an infant: case report and literature review. <i>Journal of Pediatric Surgery</i> , 2011, 46, 764-766.	1.6	8
28	Evaluating platelet function disorders in children with bleeding tendency – A single center study. <i>Platelets</i> , 2017, 28, 676-681.	2.3	7
29	Distal limb anomalies in patients with congenital dyserythropoietic anemia. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 487-490.	1.2	5
30	Improved transplant outcomes with myeloablative conditioning for hemophagocytic lymphohistiocytosis in HLA-matched and mismatched donors: a national multicenter retrospective study. <i>Bone Marrow Transplantation</i> , 2021, 56, 2088-2096.	2.4	5
31	Syndromes predisposing to leukemia are a major cause of inherited cytopenias in children. <i>Haematologica</i> , 2022, 107, 2081-2095.	3.5	5
32	Alpha-Thalassemia Carrier due to α^0 ;3.7; Deletion: Not So Silent. <i>Acta Haematologica</i> , 2020, 143, 432-437.	1.4	4
33	Pediatric myelodysplastic syndrome with inflammatory manifestations: Diagnosis, genetics, treatment, and outcome. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29138.	1.5	4
34	Mutations in RASGRP2 gene identified in patients misdiagnosed as Glanzmann thrombasthenia patients. <i>Blood Cells, Molecules, and Diseases</i> , 2021, 89, 102560.	1.4	4
35	Anti-D treatment for pediatric immune thrombocytopenia: Is the bad reputation justified?. <i>Seminars in Hematology</i> , 2016, 53, S64-S66.	3.4	3
36	Quantitation of bleeding symptoms in a national registry of patients with inherited platelet disorders. <i>Blood Cells, Molecules, and Diseases</i> , 2017, 67, 59-62.	1.4	3

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37	Bone pain at leukemia diagnosis and other risk factors for symptomatic osteonecrosis in children with acute lymphoblastic leukemia. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29033.	1.5	3
38	Neonatal haemostasis. <i>Hamostaseologie</i> , 2016, 36, 261-264.	1.9	2
39	Association between renal cystic lesions and bilateral Wilms's™ tumours. <i>European Radiology</i> , 2016, 26, 1665-1669.	4.5	2
40	Pediatric immune thrombocytopenia: apoptotic markers may help in predicting the disease course. <i>Pediatric Research</i> , 2021, 90, 93-98.	2.3	2
41	Extremity and caval deep venous thrombosis. , 2014, , 1-15.		1
42	Refractory autoimmune disease: an overview of when first-line therapy is not enough. <i>Seminars in Hematology</i> , 2016, 53, S35-S38.	3.4	1
43	Labile plasma iron as an indicator of patient adherence to iron chelation treatment. <i>Blood Cells, Molecules, and Diseases</i> , 2018, 71, 1-4.	1.4	1
44	Primary autoimmune myelofibrosis: A case report in a child. <i>EJHaem</i> , 2020, 1, 304-308.	1.0	1
45	A novel index using inflammatory markers improves the diagnosis of hemophagocytic lymphohistiocytosis in patients with hematologic malignancies.. <i>Journal of Clinical Oncology</i> , 2021, 39, 7563-7563.	1.6	1
46	Vox Sanguinis International Forum on paediatric indications for blood component transfusion. <i>Vox Sanguinis</i> , 2019, 114, e36-e90.	1.5	0
47	Splenectomy and emerging novel treatments in rare inherited hemolytic anemias. <i>HemaSphere</i> , 2019, 3, 160-162.	2.7	0
48	Thrombophilia Testing in High Pediatric Migraine Risk Children With Migraine. <i>Journal of Child Neurology</i> , 2019, 34, 121-127.	1.4	0
49	Serum soluble CD25 and ferritin in distinguishing patients with uncomplicated hematologic malignancies from patients with hematologic malignancies complicated by hemophagocytic lymphohistiocytosis.. <i>Journal of Clinical Oncology</i> , 2020, 38, e20072-e20072.	1.6	0
50	A Novel Inflammatory Index Is Sufficient to Identify Hemophagocytic Lymphohistiocytosis in Adult Patients with Hematologic Malignancies. <i>Blood</i> , 2020, 136, 1-2.	1.4	0
51	Congenital Thrombocytopenia Associated with a Heterozygous Variant in the MEIS1 Gene Encoding a Transcription Factor Essential for Megakaryopoiesis. <i>Platelets</i> , 2022, , 1-4.	2.3	0