

Sarina Levy-Mendelovich

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

281
citations

1039880

9
h-index

940416

16
g-index

28
all docs

28
docs citations

28
times ranked

450
citing authors

#	ARTICLE	IF	CITATIONS
1	Pediatric severe factor XI deficiency: A multicenter study. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29545.	0.8	2
2	Anti-TFPI for hemostasis induction in patients with rare bleeding disorders, an ex vivo thrombin generation (TG) guided pilot study. <i>Blood Cells, Molecules, and Diseases</i> , 2022, 95, 102663.	0.6	3
3	Severe Protein C Deficiency due to Novel Biallelic Variants in <i>PROC</i> and Their Phenotype Correlation. <i>Acta Haematologica</i> , 2021, 144, 327-331.	0.7	1
4	Pediatric immune thrombocytopenia: apoptotic markers may help in predicting the disease course. <i>Pediatric Research</i> , 2021, 90, 93-98.	1.1	2
5	Activated Protein C (APC) and 3K3A-APC-Induced Regression of Choroidal Neovascularization (CNV) Is Accompanied by Vascular Endothelial Growth Factor (VEGF) Reduction. <i>Biomolecules</i> , 2021, 11, 358.	1.8	5
6	Pediatric literature trends: high-level analysis using text-mining. <i>Pediatric Research</i> , 2021, 90, 212-215.	1.1	2
7	Molecular Mechanisms of Skewed X-Chromosome Inactivation in Female Hemophilia Patients—Lessons from Wide Genome Analyses. <i>International Journal of Molecular Sciences</i> , 2021, 22, 9074.	1.8	8
8	Real-World Data on Bleeding Patterns of Hemophilia A Patients Treated with Emicizumab. <i>Journal of Clinical Medicine</i> , 2021, 10, 4303.	1.0	15
9	Inherited SLP76 deficiency in humans causes severe combined immunodeficiency, neutrophil and platelet defects. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	20
10	Allergy and inhibitors in hemophilia - a rare complication with potential novel solutions. <i>Blood Cells, Molecules, and Diseases</i> , 2020, 80, 102370.	0.6	2
11	Emicizumab for infants and children with haemophilia A. <i>British Journal of Haematology</i> , 2020, 191, 145-146.	1.2	3
12	Rivaroxaban for the treatment of venous thromboembolism in pediatric patients. <i>Expert Review of Cardiovascular Therapy</i> , 2020, 18, 733-741.	0.6	8
13	Emicizumab prophylaxis among infants and toddlers with severe hemophilia A and inhibitors—a single-center cohort. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27886.	0.8	65
14	Low Concentrations of Recombinant Factor VIIa May Improve the Impaired Thrombin Generation of Glanzmann Thrombasthenia Patients. <i>Thrombosis and Haemostasis</i> , 2019, 119, 117-127.	1.8	9
15	Treatment tailoring for factor V deficient patients and perioperative management using global hemostatic coagulation assays. <i>Blood Cells, Molecules, and Diseases</i> , 2018, 71, 5-10.	0.6	6
16	Case 2: Is It Neonatal Purpura Fulminans?. <i>NeoReviews</i> , 2018, 19, e301-e303.	0.4	0
17	Thrombosis in pediatric patients with leukemia. <i>Thrombosis Research</i> , 2018, 164, S94-S97.	0.8	24
18	Cytoreductive surgery (CRS) and hyperthermic intraperitoneal chemotherapy (HIPEC) for disseminated intra-abdominal malignancies in children—a single-institution experience. <i>Journal of Pediatric Surgery</i> , 2018, 53, 1381-1386.	0.8	15

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19	Alternative treatment options for pediatric hemophilia B patients with highâ€responding inhibitors: A thrombin generationâ€guided study. <i>Pediatric Blood and Cancer</i> , 2018, 65, e27381.	0.8	4
20	From thrombasthenia to next generation thrombocytopenia: Neonatal alloimmune thrombocytopenia induced by maternal Glanzmann thrombasthenia. <i>Pediatric Blood and Cancer</i> , 2018, 65, e27376.	0.8	8
21	Rare bleeding disorders-old diseases in the era of novel options for therapy. <i>Blood Cells, Molecules, and Diseases</i> , 2017, 67, 63-68.	0.6	9
22	Quantification of specific T and B cells immunological markers in children with chronic and transient ITP. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26646.	0.8	10
23	Combination of hemostatic therapies for treatment of patients with hemophilia A and inhibitors. <i>Blood Cells, Molecules, and Diseases</i> , 2017, 66, 1-5.	0.6	8
24	Combined immunodeficiency in a patient with mosaic monosomy 21. <i>Immunologic Research</i> , 2016, 64, 841-847.	1.3	5
25	Highlighting the problematic reliance on CD18 for diagnosing leukocyte adhesion deficiency type 1. <i>Immunologic Research</i> , 2016, 64, 476-482.	1.3	23
26	Long-term follow-up of distal intestinal obstruction syndrome in cystic fibrosis. <i>World Journal of Gastroenterology</i> , 2015, 21, 318.	1.4	24