

Suvankar Majumdar

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

1,774
citations

687363

13
h-index

414414

32
g-index

40
all docs

40
docs citations

40
times ranked

2431
citing authors

#	ARTICLE	IF	CITATIONS
1	Complementary and Integrative Medicine Use in Pediatric Patients with Sickle Cell Disease. , 2022, 1, 24-28.		0
2	Effect of Poloxamer 188 vs Placebo on Painful Vaso-Occlusive Episodes in Children and Adults With Sickle Cell Disease. JAMA - Journal of the American Medical Association, 2021, 325, 1513.	7.4	24
3	Concordance with comprehensive iron assessment, hepatitis A vaccination, and hepatitis B vaccination recommendations among patients with sickle cell disease and thalassaemia receiving chronic transfusions: an analysis from the Centers for Disease Control haemoglobinopathy blood safety project. British Journal of Haematology, 2021, 195. e160-e164.	2.5	2
4	Low Rates of Cerebral Infarction after Hematopoietic Stem Cell Transplantation in Patients with Sickle Cell Disease at High Risk for Stroke. Transplantation and Cellular Therapy, 2021, 27, 1018.e1-1018.e9.	1.2	7
5	Multisystem Inflammatory Syndrome of Children: Subphenotypes, Risk Factors, Biomarkers, Cytokine Profiles, and Viral Sequencing. Journal of Pediatrics, 2021, 237, 125-135.e18.	1.8	40
6	Pediatric Sickle Cell Disease and the COVID-19 Pandemic: A Year in Review at Children's National Hospital. Blood, 2021, 138, 3036-3036.	1.4	1
7	Parvovirus B19 infection in sickle cell disease: An analysis from the Centers for Disease Control haemoglobinopathy blood surveillance project. Transfusion Medicine, 2020, 30, 226-230.	1.1	5
8	542. SARS CoV-2-Associated Multisystem Inflammatory Syndrome of Children (MIS-C) in the Washington DC Metropolitan Region. Open Forum Infectious Diseases, 2020, 7, S338-S338.	0.9	0
9	A phase 1 dose-finding study of intravenous L-citrulline in sickle cell disease: a potential novel therapy for sickle cell pain crisis. British Journal of Haematology, 2019, 184, 634-636.	2.5	10
10	Recurrent Acute Chest Syndrome in Pediatric Sickle Cell Disease: Clinical Features and Risk Factors. Journal of Pediatric Hematology/Oncology, 2018, 40, 51-55.	0.6	12
11	Inhaled corticosteroid use to prevent severe vaso-occlusive episode recurrence in children between 1 and 4 years of age with sickle cell disease: a multicenter feasibility trial. American Journal of Hematology, 2018, 93, E101-E103.	4.1	6
12	Children with sickle cell anemia with normal transcranial Doppler ultrasounds and without silent infarcts have a low incidence of new strokes. American Journal of Hematology, 2018, 93, 760-768.	4.1	8
13	Malaria, Epstein-Barr virus, vitamin A and Burkitt's lymphoma: Response to Joob and Wiwanitkit. International Journal of Cancer, 2018, 142, 864-864.	5.1	0
14	Prediction of Anti-FVIII Inhibitor Persistence By Anti-FVIII IgG Subclasses in Patients with Severe Hemophilia A in the Sippet Cohort Study. Blood, 2018, 132, 384-384.	1.4	0
15	Assessment of Iron Overload Impact on QTc Interval in Patients with Sickle Cell Disease. Blood, 2018, 132, 3673-3673.	1.4	0
16	Malaria, Epstein-Barr virus infection and the pathogenesis of Burkitt's lymphoma. International Journal of Cancer, 2017, 141, 1849-1855.	5.1	25
17	Hemophilia B Gene Therapy with a High-Specific-Activity Factor IX Variant. New England Journal of Medicine, 2017, 377, 2215-2227.	27.0	549
18	Platelet activation and erythrocyte lysis during brief exposure of blood to pathophysiological shear stress in vitro. Clinical Hemorheology and Microcirculation, 2017, 67, 159-172.	1.7	4

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19	Phase 3 Study of L-Glutamine in Sickle Cell Disease: Analyses of Time to First and Second Crisis and Average Cumulative Recurrent Events. <i>Blood</i> , 2017, 130, 685-685.	1.4	3
20	A Randomized Trial of Factor VIII and Neutralizing Antibodies in Hemophilia A. <i>New England Journal of Medicine</i> , 2016, 374, 2054-2064.	27.0	414
21	Daily Pain, Physical Activity, and Home Fluid Intake in Pediatric Sickle Cell Disease. <i>Journal of Pediatric Psychology</i> , 2016, 42, jsw061.	2.1	10
22	Spk-9001: Adeno-Associated Virus Mediated Gene Transfer for Hemophilia B Achieves Sustained Mean Factor IX Activity Levels of >30% without Immunosuppression. <i>Blood</i> , 2016, 128, 3-3.	1.4	24
23	Quality of life among pediatric patients with cancer: Contributions of time since diagnosis and parental chronic stress. <i>Pediatric Blood and Cancer</i> , 2015, 62, 1232-1236.	1.5	24
24	Source of Factor VIII Replacement (PLASMATIC OR RECOMBINANT) and Incidence of Inhibitory Alloantibodies in Previously Untreated Patients with Severe Hemophilia a: The Multicenter Randomized Sippet Study. <i>Blood</i> , 2015, 126, 5-5.	1.4	16
25	An Update on Sickle Cell Anemia in Children and Adolescents. <i>Journal of the Mississippi State Medical Association</i> , 2015, 56, 268-71.	0.1	1
26	Outcome of overt stroke in sickle cell anaemia, a single institution's experience. <i>British Journal of Haematology</i> , 2014, 165, 707-713.	2.5	15
27	Associations among emergency room visits, parenting styles, and psychopathology among pediatric patients with sickle cell. <i>Pediatric Blood and Cancer</i> , 2014, 61, 1822-1827.	1.5	5
28	Controlled Trial of Transfusions for Silent Cerebral Infarcts in Sickle Cell Anemia. <i>New England Journal of Medicine</i> , 2014, 371, 699-710.	27.0	421
29	1225. <i>Critical Care Medicine</i> , 2014, 42, A1646.	0.9	0
30	The use and effectiveness of complementary and alternative medicine for pain in sickle cell anemia. <i>Complementary Therapies in Clinical Practice</i> , 2013, 19, 184-187.	1.7	30
31	Stroke with intracranial stenosis is associated with increased platelet activation in sickle cell anemia. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1192-1197.	1.5	9
32	The Adolescent with Sickle Cell Disease. , 2013, , 295-306.		1
33	The adolescent with sickle cell disease. <i>Adolescent Medicine: State of the Art Reviews</i> , 2013, 24, 295-306, xv.	0.2	1
34	A Rare Case of Peripheral Neuropathy From Relapse of Acute Lymphoblastic Leukemia to the Brachial Plexus. <i>Journal of Pediatric Hematology/Oncology</i> , 2012, 34, e77-e79.	0.6	5
35	Overweight and Obesity in Hemophilia. <i>American Journal of Preventive Medicine</i> , 2011, 41, S369-S375.	3.0	58
36	Compound heterozygous mutation with a novel splice donor region DNA sequence variant in the succinate dehydrogenase subunit B gene in malignant paraganglioma. <i>Pediatric Blood and Cancer</i> , 2010, 54, 473-475.	1.5	17

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37	What are the mechanisms of tumor formation in patients with paraganglioma and <i>SDHB</i> germline mutations?. <i>Pediatric Blood and Cancer</i> , 2010, 55, 212-212.	1.5	0
38	Gender Differences in Incidence Rates of Childhood B-Precursor Acute Lymphocytic Leukemia in Mississippi. <i>Journal of Pediatric Oncology Nursing</i> , 2010, 27, 164-167.	1.5	11
39	Outcome of Hematopoietic Cell Transplantation in Children with Sickle Cell Disease, a Single Center's Experience.. <i>Blood</i> , 2007, 110, 1112-1112.	1.4	6
40	Successful Prophylactic Treatment for Bleeding in a Girl With Severe Hereditary Prothrombin Deficiency Using a Prothrombin Complex Concentrate (Bebulin® VH). <i>Journal of Pediatric Hematology/Oncology</i> , 2004, 26, 480-483.	0.6	10