Richard Ward

List of Publications by Year
in descending order

54	595	759055 12	642610 23
papers	citations	h-index	g-index

54	54	54	1072
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Thromboprophylaxis Reduced Venous Thromboembolism in Sickle Cell Patients with Central Venous Access Devices: A Retrospective Cohort Study. Journal of Clinical Medicine, 2022, 11, 1193.	1.0	4
2	Adverse outcome of acute splenic sequestration crisis in pregnancy. Obstetric Medicine, 2021, 14, 113-115.	0.5	0
3	Screening for Cognitive Dysfunction Using the Rowland Universal Dementia Assessment Scale in Adults With Sickle Cell Disease. JAMA Network Open, 2021, 4, e217039.	2.8	7
4	Comparison of Inline R2* MRI versus FerriScan for liver iron quantification in patients on chelation therapy for iron overload: preliminary results. European Radiology, 2021, 31, 9296-9305.	2.3	5
5	Distinct maternal and fetal pregnancy outcomes in women with sickle cell disease can be predicted using routine clinical and laboratory data. British Journal of Haematology, 2021, 194, 1063-1073.	1.2	10
6	Exploration of Barriers and Facilitators to Optimal Emergency Department Care of Sickle Cell Disease: Opportunities for Patientâ€“Physician Partnerships to Improve Care. Hemoglobin, 2021, 45, 13-19.	0.4	1
7	Glucose dysregulation in patients with iron overload: is there a relationship with quantitative pancreas and liver iron and fat content measured by MRI?. European Radiology, 2020, 30, 1616-1623.	2.3	11
8	Impact of MRI technique on clinical decision-making in patients with liver iron overload: comparison of FerriScan- versus R2*-derived liver iron concentration. European Radiology, 2020, 30, 1959-1968.	2.3	4
9	Impact of a transition program with navigator on loss to followâ€up, medication adherence, and appointment attendance in hemoglobinopathies. Pediatric Blood and Cancer, 2019, 66, e27781.	0.8	33
10	Deferiprone exerts a doseâ€dependent reduction of liver iron in adults with iron overload. European Journal of Haematology, 2019, 103, 80-87.	1.1	14
11	Development of a patientâ€reported outcomes symptom measure for patients with nontransfusionâ€dependent thalassemia (NTDTâ€PRO^{Â©}). American Journal of Hematology, 2019, 94, 171-176.	2.0	7
12	Prospective Evaluation of an R2* Method for Assessing Liver Iron Concentration (LIC) Against FerriScan: Derivation of the Calibration Curve and Characterization of the Nature and Source of Uncertainty in the Relationship. Journal of Magnetic Resonance Imaging, 2019, 49, 1467-1474.	1.9	25
13	An innovative short-stay health care model for treatment of uncomplicated vaso-occlusive crisis in adult sickle cell disease patients in Canada to reduce emergency department utilization. Canadian Journal of Emergency Medicine, 2019, 21, 55-62.	0.5	4
14	Cord gas parameters in infants born to women with sickle cell disease: a retrospective matched cohort study. British Journal of Haematology, 2019, 184, 653-657.	1.2	3
15	Patient Education Interventions for Improving Self-Management in Adults with Hemoglobinopathies: A Systematic Review and Meta-Analysis. Blood, 2019, 134, 5783-5783.	0.6	0
16	A proposed treatment algorithm for adults with Haemoglobin <scp>SC</scp> disease. British Journal of Haematology, 2018, 182, 607-609.	1.2	11
17	Design and implementation of a novel advanced training curriculum in hemoglobinopathies. American Journal of Hematology, 2018, 93, E75-E77.	2.0	0
18	Approach to transfusion in pregnant women with sickle cell disease: a survey of physicians. British Journal of Haematology, 2018, 183, 516-519.	1.2	3

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19	Diagnostic accuracy and yield of screening tests for atrial fibrillation in the family practice setting: a multicentre cohort study. CMAJ Open, 2018, 6, E308-E315.	1.1	21
20	Exploring the Needs of Adolescents With Sickle Cell Disease to Inform a Digital Self-Management and Transitional Care Program: Qualitative Study. JMIR Pediatrics and Parenting, 2018, 1, e11058.	0.8	19
21	Should Women with Sickle Cell Disease be Administered Prophylactic Transfusions to Avoid Complications during Pregnancy?. Blood, 2018, 132, 3679-3679.	0.6	0
22	Validation of Sickle Cell Disease Severity Score in a Cohort of Hemoglobin SC Disease Patients. Blood, 2018, 132, 2287-2287.	0.6	0
23	Is There a Role for Thromboprophylaxis in Sickle Cell Patients with Central Venous Access Devices?. Blood, 2018, 132, 3518-3518.	0.6	1
24	Examining the Role of Sickle Cell Disease Patients As Teachers in the Emergency Department: Patient Directed Physician Education. Blood, 2018, 132, 4722-4722.	0.6	0
25	Real World Experience with Sublingual Fentanyl for the Treatment of Sickle Cell Vaso-Occlusive Episodes in Adults in a Tertiary Canadian Emergency Department. Blood, 2018, 132, 5879-5879.	0.6	0
26	Comprehensive Structured Transition Program With Dedicated Transition Navigator Reduced Lost to Follow-Up and Improved Medication Adherence in Adolescents and Young Adults With Sickle Cell Disease and Thalassemia. Journal of Adolescent Health, 2017, 60, S40-S41.	1.2	5
27	Quantification of global myocardial function by cine MRI deformable registration-based analysis: Comparison with MR feature tracking and speckle-tracking echocardiography. European Radiology, 2017, 27, 1404-1415.	2.3	31
28	Safety and feasibility of red cell exchange for sickle cell disease across Canada. Transfusion and Apheresis Science, 2016, 55, 129-130.	0.5	0
29	Patient Involvement as Experts in the Development and Assessment of a Smartphone App as a Patient Education Tool for the Management of Thalassemia and Iron Overload Syndromes. Hemoglobin, 2016, 40, 323-329.	0.4	11
30	Characterization of Chronic Pain and Opioid Usage in Adult Sickle Cell Disease Patients Referred to a Comprehensive Pain Clinic. Pain Medicine, 2016, 17, 2145-2146.	0.9	0
31	Quantification of Myocardial Extracellular Volume Fraction with Cardiac MR Imaging in Thalassemia Major. Radiology, 2016, 279, 720-730.	3.6	44
32	A 19-year-old woman with sickle cell disease and pain. Cmaj, 2016, 188, 745-746.	0.9	1
33	Comprehensive Structured Transition Program with Dedicated Transition Navigator Reduced Lost to Follow-up and Improved Medication Adherence in Sickle Cell Disease and Thalassemia Adolescents and Young Adults. Blood, 2016, 128, 317-317.	0.6	2
34	Factors Impacting Quality of Life in Thalassemia Patients; Results from the Intercontinenthal Collaborative Study. Blood, 2016, 128, 3633-3633.	0.6	15
35	Pilot Study of Online Learning Modules for Hemoglobinopathy Education in Canadian Hematology Training Programs. Blood, 2016, 128, 314-314.	0.6	1
36	Prophylactic transfusion for pregnant women with sickle cell disease: a systematic review and meta-analysis. Blood, 2015, 126, 2424-2435.	0.6	81

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37	A comparison of chronic manual and automated red blood cell exchange transfusion in sickle cell disease patients. British Journal of Haematology, 2015, 170, 425-428.	1.2	38
38	Optimal Manual Exchange Transfusion Protocol for Sickle Cell Disease: A Retrospective Comparison of Two Comprehensive Care Centers in the United Kingdom and Canada. Hemoglobin, 2015, 39, 310-315.	0.4	5
39	An Unusual Case of Acquired Hemophilia a and Factor XIII Consumption. Blood, 2015, 126, 4701-4701.	0.6	0
40	Aligning Health Care Policy With Evidence-Based Medicine: The Case for Funding Direct Oral Anticoagulants in Atrial Fibrillation. Canadian Journal of Cardiology, 2014, 30, 1245-1248.	0.8	11
41	Neglected conditions. Cmaj, 2014, 186, 452.2-452.	0.9	0
42	A doubleâ€blind, placeboâ€controlled phase II study of the efficacy and safety of 2,2â€dimethylbutyrate (HQKâ€1001), an oral fetal globin inducer, in sickle cell disease. American Journal of Hematology, 2014, 89, 709-713.	2.0	54
43	Hemoglobinopathy Education in Canadian Hematology Training Programs: How Much Are Residents Learning?. Blood, 2014, 124, 2168-2168.	0.6	0
44	Heterogeneity of myocardial iron distribution in response to chelation therapy in patients with transfusion-dependent anemias. International Journal of Cardiovascular Imaging, 2013, 29, 1517-1526.	0.7	5
45	A North American Experience Of Hemoglobin SC Disease, Its Complications, and Management. Blood, 2013, 122, 2221-2221.	0.6	2
46	Characterization Of Pulmonary Compliance In Sickle Cell Patients Revealed Wide Variability. Blood, 2013, 122, 989-989.	0.6	0
47	Treatment Patterns and Outcomes Of Sickle Cell Patients With Frequent ER Visits: A Single Center Experience. Blood, 2013, 122, 1010-1010.	0.6	1
48	Effectiveness Of An Analgesia Protocol For The Treatment Of Painful Vaso-Occlusive Crisis Of Sickle Cell Disease Patients In Emergency Room: A Retrospective Cohort Study. Blood, 2013, 122, 2222-2222.	0.6	1
49	A Comparison of Chronic Manual and Automated Red Blood Cell Exchange Transfusion in Sickle Cell Disease Patients From Two Comprehensive Care Centres in the United Kingdom. Blood, 2012, 120, 3430-3430.	0.6	1
50	Comparing Patterns for Transitioning the Care of Young Adults with Sickle Cell Disease Versus Hemophilia: The Toronto Experience. Blood, 2011, 118, 2072-2072.	0.6	1
51	Effectiveness, Time Utilization and Clinical Outcome of Partial Manual Red Cell Exchange in Patients with Sickle Cell Disease. Blood, 2011, 118, 2326-2326.	0.6	0
52	Application of Self-Efficacy Theory in Adherence to Iron Chelation Therapy: A Single-Center Cross-Sectional Study. Blood, 2011, 118, 5284-5284.	0.6	1
53	The Effect of Comprehensive Care on Maternal and Fetal Outcomes in Sickle Cell Disease Pregnancies. Blood, 2011, 118, 4842-4842.	0.6	0
54	The antiparasitic agent ivermectin induces chloride-dependent membrane hyperpolarization and cell death in leukemia cells. Blood, 2010, 116, 3593-3603.	0.6	101