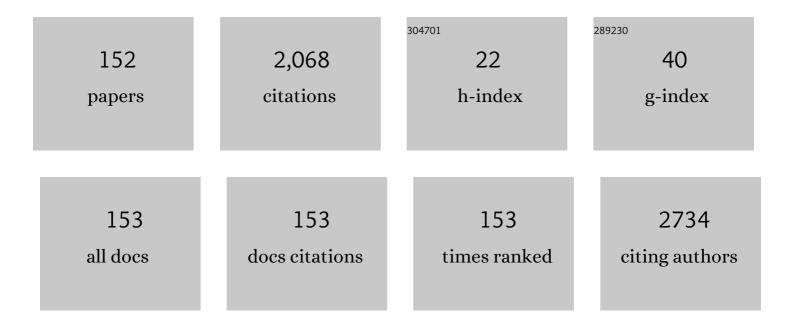
Santosh L Saraf

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Update on Meningiomas. Oncologist, 2011, 16, 1604-1613.	3.7	151
2	Nonmyeloablative Stem Cell Transplantation with Alemtuzumab/Low-Dose Irradiation to Cure and Improve theÂQuality of Life of Adults with Sickle Cell Disease. Biology of Blood and Marrow Transplantation, 2016, 22, 441-448.	2.0	111
3	Differences in the clinical and genotypic presentation of sickle cell disease around the world. Paediatric Respiratory Reviews, 2014, 15, 4-12.	1.8	97
4	Haemoglobinuria is associated with chronic kidney disease and its progression in patients with sickle cell anaemia. British Journal of Haematology, 2014, 164, 729-739.	2.5	91
5	COVIDâ€19 infection in patients with sickle cell disease. British Journal of Haematology, 2020, 189, 851-852.	2.5	90
6	Kidney Disease among Patients with Sickle Cell Disease, Hemoglobin SS and SC. Clinical Journal of the American Society of Nephrology: CJASN, 2016, 11, 207-215.	4.5	75
7	Feasibility of Implementing a Comprehensive Warfarin Pharmacogenetics Service. Pharmacotherapy, 2013, 33, 1156-1164.	2.6	70
8	Age-related differences in disease characteristics and clinical outcomes in polycythemia vera. Leukemia and Lymphoma, 2013, 54, 1989-1995.	1.3	65
9	Clinical effectiveness of decitabine in severe sickle cell disease. British Journal of Haematology, 2008, 141, 126-129.	2.5	64
10	Genetic variants and cell-free hemoglobin processing in sickle cell nephropathy. Haematologica, 2015, 100, 1275-1284.	3.5	60
11	Curative therapies: Allogeneic hematopoietic cell transplantation from matched related donors using myeloablative, reduced intensity, and nonmyeloablative conditioning in sickle cell disease. Seminars in Hematology, 2018, 55, 87-93.	3.4	51
12	Haploidentical Peripheral Blood Stem Cell Transplantation Demonstrates Stable Engraftment in Adults with Sickle Cell Disease. Biology of Blood and Marrow Transplantation, 2018, 24, 1759-1765.	2.0	50
13	APOL1 , α-thalassemia, and BCL11A variants as a genetic risk profile for progression of chronic kidney disease in sickle cell anemia. Haematologica, 2017, 102, e1-e6.	3.5	47
14	Metabolomic Markers of Kidney Function Decline in Patients With Diabetes: Evidence From the Chronic Renal Insufficiency Cohort (CRIC) Study. American Journal of Kidney Diseases, 2020, 76, 511-520.	1.9	45
15	Nonâ€myeloablative human leukocyte antigenâ€matched related donor transplantation in sickle cell disease: outcomes from three independent centres. British Journal of Haematology, 2021, 192, 761-768.	2.5	41
16	Combination of Linear Accelerator–Based Intensity-Modulated Total Marrow Irradiation and Myeloablative Fludarabine/Busulfan: A Phase I Study. Biology of Blood and Marrow Transplantation, 2014, 20, 2034-2041.	2.0	40
17	Characterization of opioid use in sickle cell disease. Pharmacoepidemiology and Drug Safety, 2018, 27, 479-486.	1.9	37
18	Losartan for the nephropathy of sickle cell anemia: A phaseâ€2, multicenter trial. American Journal of Hematology, 2017, 92, E520-E528.	4.1	36

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#	Article	IF	CITATIONS
19	Hyperfiltration is associated with the development of microalbuminuria in patients with sickle cell anemia. American Journal of Hematology, 2014, 89, 1156-1157.	4.1	35
20	Hypoxic Response Contributes to Altered Gene Expression and Precapillary Pulmonary Hypertension in Patients With Sickle Cell Disease. Circulation, 2014, 129, 1650-1658.	1.6	32
21	Clinical, laboratory, and genetic risk factors for thrombosis in sickle cell disease. Blood Advances, 2020, 4, 1978-1986.	5.2	28
22	Progression of albuminuria in patients with sickle cell anemia: a multicenter, longitudinal study. Blood Advances, 2020, 4, 1501-1511.	5.2	28
23	The nephropathy of sickle cell trait and sickle cell disease. Nature Reviews Nephrology, 2022, 18, 361-377.	9.6	26
24	Comparison of Patients from Nigeria and the USA Highlights Modifiable Risk Factors for Sickle Cell Anemia Complications. Hemoglobin, 2014, 38, 236-243.	0.8	24
25	Patterns of opioid use in sickle cell disease. American Journal of Hematology, 2016, 91, 1102-1106.	4.1	24
26	Association of circulating transcriptomic profiles with mortality in sickle cell disease. Blood, 2017, 129, 3009-3016.	1.4	22
27	Iron status, fibroblast growth factor 23 and cardiovascular and kidney outcomes in chronic kidney disease. Kidney International, 2021, 100, 1292-1302.	5.2	22
28	Genetic polymorphism of APOB is associated with diabetes mellitus in sickle cell disease. Human Genetics, 2015, 134, 895-904.	3.8	20
29	Red blood cell alloimmunization in sickle cell disease: assessment of transfusion protocols during two time periods. Transfusion, 2018, 58, 1588-1596.	1.6	20
30	HMOX1 and acute kidney injury in sickle cell anemia. Blood, 2018, 132, 1621-1625.	1.4	20
31	Systematic Review of Crizanlizumab: A New Parenteral Option to Reduce Vasoâ€occlusive Pain Crises in Patients with Sickle Cell Disease. Pharmacotherapy, 2020, 40, 535-543.	2.6	19
32	Systematic Review of Voxelotor: A Firstâ€in lass Sickle Hemoglobin Polymerization Inhibitor for Management of Sickle Cell Disease. Pharmacotherapy, 2020, 40, 525-534.	2.6	17
33	Non-p53 Dependent, Leukemia Initiating-Cell Selective, Therapy Blood, 2009, 114, 2077-2077.	1.4	17
34	Identification of ceruloplasmin as a biomarker of chronic kidney disease in urine of sickle cell disease patients by proteomic analysis. American Journal of Hematology, 2018, 93, E45-E47.	4.1	16
35	Risk factors for vitamin D deficiency in sickle cell disease. British Journal of Haematology, 2018, 181, 828-835.	2.5	16
36	Hydroxycarbamide adherence and cumulative dose associated with hospital readmission in sickle cell disease: a 6â€year populationâ€based cohort study. British Journal of Haematology, 2018, 182, 259-270.	2.5	16

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37	Progressive glomerular and tubular damage in sickle cell trait and sickle cell anemia mouse models. Translational Research, 2018, 197, 1-11.	5.0	15
38	Urinary orosomucoid is associated with progressive chronic kidney disease stage in patients with sickle cell anemia. American Journal of Hematology, 2018, 93, E107-E109.	4.1	15
39	Ex vivo expansion of human mobilized peripheral blood stem cells using epigenetic modifiers. Transfusion, 2015, 55, 864-874.	1.6	14
40	Outcome Disparities in Caucasian andÂNon-Caucasian Patients With Myeloproliferative Neoplasms. Clinical Lymphoma, Myeloma and Leukemia, 2016, 16, 350-357.	0.4	14
41	Hemolysis and hemolysisâ€related complications in females vs. males with sickle cell disease. American Journal of Hematology, 2018, 93, E376-E380.	4.1	14
42	Similar burden of type 2 diabetes among adult patients with sickle cell disease relative to African Americans in the U.S. population: a sixâ€year populationâ€based cohort analysis. British Journal of Haematology, 2019, 185, 116-127.	2.5	14
43	Improved health care utilization and costs in transplanted versus non-transplanted adults with sickle cell disease. PLoS ONE, 2020, 15, e0229710.	2.5	14
44	Comparing the Effectiveness of Education Versus Digital Cognitive Behavioral Therapy for Adults With Sickle Cell Disease: Protocol for the Cognitive Behavioral Therapy and Real-time Pain Management Intervention for Sickle Cell via Mobile Applications (CaRISMA) Study. JMIR Research Protocols, 2021, 10, e29014.	1.0	14
45	Changes in Conjunctival Hemodynamics Predict Albuminuria in Sickle Cell Nephropathy. American Journal of Nephrology, 2015, 41, 487-493.	3.1	12
46	Associations of \hat{I}_{\pm} -thalassemia and BCL11A with stroke in Nigerian, United States, and United Kingdom sickle cell anemia cohorts. Blood Advances, 2017, 1, 693-698.	5.2	12
47	Allogeneic Hematopoietic Stem Cell Transplantation for Adults with Sickle Cell Disease. Journal of Clinical Medicine, 2019, 8, 1565.	2.4	12
48	FT-4202, an Allosteric Activator of Pyruvate Kinase-R, Demonstrates Proof of Mechanism and Proof of Concept after a Single Dose and after Multiple Daily Doses in a Phase 1 Study of Patients with Sickle Cell Disease. Blood, 2020, 136, 19-20.	1.4	12
49	Role of Ethnicity in Clinical Outcomes of Patients with Ph-Negative Myeloproliferative Neoplasms. Blood, 2012, 120, 2076-2076.	1.4	12
50	Use of anti-inflammatory analgesics in sickle-cell disease. Journal of Clinical Pharmacy and Therapeutics, 2017, 42, 656-660.	1.5	11
51	Fixed lowâ€dose hydroxyurea for the treatment of adults with sickle cell anemia in <scp>N</scp> igeria. American Journal of Hematology, 2018, 93, E193.	4.1	11
52	The morbidity and mortality of end stage renal disease in sickle cell disease. American Journal of Hematology, 2019, 94, E138-E141.	4.1	11
53	Rapid decline in estimated glomerular filtration rate in sickle cell anemia: results of a multicenter pooled analysis. Haematologica, 2021, 106, 1749-1753.	3.5	11
54	Impact of a Clinical Pharmacy Service on the Management of Patients in a Sickle Cell Disease Outpatient Center. Pharmacotherapy, 2016, 36, 1166-1172.	2.6	10

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55	Health-Related Quality of Life and Personal Life Goals of Adults With Sickle Cell Disease After Hematopoietic Stem Cell Transplantation. Western Journal of Nursing Research, 2019, 41, 555-575.	1.4	10
56	Anemia and Incident End-Stage Kidney Disease. Kidney360, 2020, 1, 623-630.	2.1	10
57	A Multi-Center, Phase-2 Trial of Losartan for the Nephropathy of Sickle Cell Anemia. Blood, 2016, 128, 265-265.	1.4	10
58	Standard clinical practice underestimates the role and significance of erythropoietin deficiency in sickle cell disease. British Journal of Haematology, 2011, 153, 386-392.	2.5	9
59	A prospective study of intravenous pentamidine for PJP prophylaxis in adult patients undergoing intensive chemotherapy or hematopoietic stem cell transplant. Bone Marrow Transplantation, 2018, 53, 300-306.	2.4	9
60	Erythropoiesisâ€stimulating agents in sickle cell anaemia. British Journal of Haematology, 2018, 182, 602-605.	2.5	9
61	Safety of chronic transdermal fentanyl use in patients receiving hemodialysis. American Journal of Health-System Pharmacy, 2016, 73, 947-948.	1.0	8
62	Trans-ethnic genome-wide association study of blood metabolites in the Chronic Renal Insufficiency Cohort (CRIC) study. Kidney International, 2022, 101, 814-823.	5.2	8
63	Program expansion of a day hospital dedicated to manage sickle cell pain. American Journal of Hematology, 2018, 93, E20-E21.	4.1	7
64	"Maximum tolerated dose―vs "fixed lowâ€dose―hydroxyurea for treatment of adults with sickle cell anemia. American Journal of Hematology, 2019, 94, E112-E115.	4.1	7
65	ARTS: automated randomization of multiple traits for study design. Bioinformatics, 2014, 30, 1637-1639.	4.1	6
66	Conjunctival microvascular hemodynamics following vaso-occlusive crisis in sickle cell disease. Clinical Hemorheology and Microcirculation, 2016, 62, 359-367.	1.7	6
67	The experience of adults with sickle cell disease and their HLAâ€matched adult sibling donors after allogeneic hematopoietic stem cell transplantation. Journal of Advanced Nursing, 2019, 75, 2943-2951.	3.3	6
68	Biomarkers of clinical severity in treated and untreated sickle cell disease: a comparison by genotypes of a single center cohort and African Americans in the NHANES study. British Journal of Haematology, 2021, 194, 767-778.	2.5	6
69	Phase 1 Single (SAD) and Multiple Ascending Dose (MAD) Studies of the Safety, Tolerability, Pharmacokinetics (PK) and Pharmacodynamics (PD) of FT-4202, an Allosteric Activator of Pyruvate Kinase-R, in Healthy and Sickle Cell Disease Subjects. Blood, 2019, 134, 616-616.	1.4	6
70	Use of metformin in patients with sickle cell disease. American Journal of Hematology, 2019, 94, E13-E15.	4.1	5
71	Voxelotor and albuminuria in adults with sickle cell anaemia. British Journal of Haematology, 2022, , .	2.5	5
72	Longitudinal study of glomerular hyperfiltration in adults with sickle cell anemia: a multicenter pooled analysis. Blood Advances, 2022, 6, 4461-4470.	5.2	5

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73	A genetic variation associated with plasma erythropoietin and a non-coding transcript ofPRKAR1Ain sickle cell disease. Human Molecular Genetics, 2016, 25, ddw299.	2.9	4
74	Increased vancomycin dosing requirements in sickle cell disease due to hyperfiltration-dependent and independent pathways. Haematologica, 2017, 102, e282-e284.	3.5	4
75	Kidney ultrasound findings according to kidney function in sickle cell anemia. American Journal of Hematology, 2019, 94, E288-E291.	4.1	4
76	Engulfment and cell motility 1 (ELMO1) and apolipoprotein A1 (APOA1) as candidate genes for sickle cell nephropathy. British Journal of Haematology, 2021, 193, 628-632.	2.5	4
77	Urinary Kringle Domain-Containing Protein HGFL: A Validated Biomarker of Early Sickle Cell Anemia-Associated Kidney Disease. American Journal of Nephrology, 2021, 52, 582-587.	3.1	4
78	Urinary Ceruloplasmin Concentration Predicts Development of Kidney Disease in Sickle Cell Disease Patients. Blood, 2016, 128, 4865-4865.	1.4	4
79	Chronic opioid use can be reduced or discontinued after haematopoietic stem cell transplantation for sickle cell disease. British Journal of Haematology, 2020, 191, e70-e72.	2.5	3
80	Heme A1M'ed at the kidney in sickle cell disease. Blood, 2020, 135, 979-981.	1.4	3
81	Association of Blood Pressure Genetic Risk Score with Cardiovascular Disease and CKD Progression: Findings from the CRIC Study. Kidney360, 2021, 2, 1251-1260.	2.1	3
82	Race/ethnicity and underlying disease influences hematopoietic stem/progenitor cell mobilization response: A single center experience. Journal of Clinical Apheresis, 2021, 36, 634-644.	1.3	3
83	Chronic Opioid Use Pattern in Adult Patients with Sickle Cell Disease. Blood, 2015, 126, 3400-3400.	1.4	3
84	Low Fixed Dose Hydroxyurea for the Treatment of Adults with Sickle Cell Disease in Nigeria. Blood, 2017, 130, 981-981.	1.4	3
85	Clinical, Laboratory, and Genetic Risk Factors for Thrombosis in Sickle Cell Disease. Blood, 2018, 132, 9-9.	1.4	3
86	Antimicrobial resistance is a risk factor for mortality in adults with sickle cell disease. Haematologica, 2021, 106, 1745-1748.	3.5	3
87	The Effect of Crizanlizumab on the Number of Days Requiring Opioid Use for Management of Pain Associated with Vaso-Occlusive Crises in Patients with Sickle Cell Disease: Results from the Sustain Trial. Blood, 2020, 136, 32-33.	1.4	3
88	Hyperkalemia and Metabolic Acidosis Occur at Higher Estimated Glomerular Filtration Rates in Sickle Cell Disease. Kidney360, 0, , 10.34067/KID.0006802021.	2.1	3
89	Potential Contribution of Pulmonary Thromboembolic Disease in Pulmonary Hypertension in Sickle Cell Disease. Annals of the American Thoracic Society, 2020, 17, 899-901.	3.2	2
90	Effects of reninâ€angiotensin blockade and APOL1 on kidney function in sickle cell disease. EJHaem, 2021, 2, 483-484.	1.0	2

#	Article	IF	CITATIONS
91	Type 2 Diabetes Mellitus in Patients with Sickle Cell Disease: A Population-Based Longitudinal Analysis of Three Cohorts. Blood, 2018, 132, 4817-4817.	1.4	2
92	Thrombomodulin and Endothelial Dysfunction in Sickle Cell Anemia. Blood, 2019, 134, 3558-3558.	1.4	2
93	Phase 1 Evaluation of Oral Tetrahydrouridine-Decitabine As Non-Cytotoxic Epigenetic Disease Modification for Sickle Cell Disease. Blood, 2016, 128, 124-124.	1.4	2
94	Hydroxyurea for Treatment of Sickle Cell Disease in Adults in Africa. Blood, 2016, 128, 1305-1305.	1.4	2
95	Manifestations of Reduced Kidney Function Occur at a Higher Estimated Glomerular Filtration Rate in Sickle Cell Anemia. Blood, 2019, 134, 2268-2268.	1.4	2
96	Utility of the revised cardiac risk index for predicting postsurgical morbidity in Hb SC and Hb Sβ+â€thalassemia sickle cell disease. American Journal of Hematology, 2016, 91, E316-7.	4.1	1
97	Reply to <scp>R</scp> uan <scp>X</scp> et al: "A comment on pattern of opioid use in sickle cell disease― American Journal of Hematology, 2017, 92, E43.	4.1	1
98	High inpatient dose of opioid at discharge compared to home dose predicts readmission risk in sickle cell disease. American Journal of Hematology, 2019, 94, E5-E7.	4.1	1
99	<i>S100B</i> has pleiotropic effects on vasoâ€occlusive manifestations in sickle cell disease. American Journal of Hematology, 2020, 95, E62-E65.	4.1	1
100	Using machine learning to predict rapid decline of kidney function in sickle cell anemia. EJHaem, 2021, 2, 257-260.	1.0	1
101	The vasculopathic cord between preâ€eclampsia and kidney function in sickle cell disease. British Journal of Haematology, 2021, 194, 947-949.	2.5	1
102	Health Care Utilization in Transplanted Versus Non-Transplanted Sickle Cell Disease Patients. Blood, 2018, 132, 313-313.	1.4	1
103	Regulatory Genetic Variation at the S100B Gene Associates with Vaso-Occlusive Manifestations in Sickle Cell Disease. Blood, 2018, 132, 1063-1063.	1.4	1
104	Assessment of Bone Marrow Function in Sickle Cell Anaemia Patients Using Corrected Reticulocyte Counts. Blood, 2015, 126, 4581-4581.	1.4	1
105	Effect of Angiotensin Converting Enzyme Inhibitors and Angiotensin Receptor Blockers on Kidney Function in Patients with Sickle Cell Disease. Blood, 2016, 128, 3666-3666.	1.4	1
106	Genetic Modifiers Identify a High Risk Group for Stroke in Three Independent Cohorts of Sickle Cell Anemia Patients. Blood, 2016, 128, 1015-1015.	1.4	1
107	Biomarkers of Cardiopulmonary, Renal, and Liver Dysfunction in an Adult Sickle Cell Disease Cohort. Blood, 2019, 134, 3574-3574.	1.4	1
108	Biomarker Association with Hypertension in Mild Versus Severe Sickle Cell Disease Genotypes of a Single Center Cohort, in Comparison with African Americans from the Nhanes Study. Blood, 2021, 138, 2051-2051.	1.4	1

#	Article	IF	CITATIONS
109	Use of Multiple Urinary Biomarkers for Early Detection of Chronic Kidney Disease in Sickle Cell Anemia Patients. Blood, 2020, 136, 30-30.	1.4	1
110	Black and White Adults With CKD Hospitalized With Acute Kidney Injury: Findings From the Chronic Renal Insufficiency Cohort (CRIC) Study. American Journal of Kidney Diseases, 2022, , .	1.9	1
111	Thrombomodulin and <scp>multiorgan</scp> failure in sickle cell anemia. American Journal of Hematology, 2022, 97, .	4.1	1
112	Improvement of Hemolytic Anemia with GBT1118 is Reno-protective in Transgenic Sickle Mice. Blood Advances, 0, , .	5.2	1
113	Type 2 diabetes in adults with sickle cell disease: can we dive deeper? Response to Skinner <i>etÂal</i> . British Journal of Haematology, 2019, 186, 782-783.	2.5	0
114	Laparoscopic Sleeve Gastrectomy in Sickle Cell Disease: a Case Series. Obesity Surgery, 2019, 29, 3762-3764.	2.1	0
115	Evaluation of pointâ€ofâ€care International Normalized Ratio in sickle cell disease. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12533.	2.3	0
116	African American Patients with Multiple Myeloma Have Prolonged Responses after Autologous Stem Cell Transplantation Blood, 2005, 106, 3131-3131.	1.4	0
117	Significance of, and Difficulty in Diagnosing, Erythropoietin Deficiency in Sickle Cell Anemia. Blood, 2008, 112, 2479-2479.	1.4	0
118	Favorable Responses to Novel Agents for Multiple Myeloma in African American Patients,. Blood, 2011, 118, 4213-4213.	1.4	0
119	Conjunctival Biopsy to Guide Treatment of Chronic Ocular Gvhd. Blood, 2012, 120, 4491-4491.	1.4	0
120	Clinical and Laboratory Predictors for Renal Damage in Sickle Cell Disease. Blood, 2012, 120, 3252-3252.	1.4	0
121	Hemoglobinuria Is a Risk Factor For Kidney Disease Progression In Sickle Cell Anemia. Blood, 2013, 122, 996-996.	1.4	0
122	Myeloablative Fludarabine/ IV Busulfan Combined With Linac Based Intentsity Modulated Total Marrow Irradiation (IM-TMI) In Allogeneic Stem Cell Transplant For High Risk Hematologic Malignancies: A Phase I Study. Blood, 2013, 122, 3285-3285.	1.4	0
123	LINAC-based intensity modulated total marrow irradiation (TMI) in addition to myeloablative fludarabine/IV busulfan conditioning prior to allogeneic stem cell transplant for high-risk hematologic malignancies: A phase I study Journal of Clinical Oncology, 2014, 32, 7045-7045.	1.6	0
124	Utility of the Revised Cardiac Index Score for Predicting Post-Surgical Outcome in Hb SC or SBeta+ -Thalassemia Sickle Cell Disease. Blood, 2015, 126, 3413-3413.	1.4	0
125	CCN2 - Exploring a New Biomarker in Myelofibrosis. Blood, 2015, 126, 4063-4063.	1.4	0

Allogeneic Hematopoietic Cell Transplant in Sickle Cell Disease. , 2016, , 89-96.

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127	Elevated Levels of Hgfl Protein in Sickle Cell Disease Urine Samples That Induce Glomerular Permeability. Blood, 2016, 128, 4841-4841.	1.4	0
128	Progressive Glomerular Damage in Sickle Cell Trait and Sickle Cell Anemia Mouse Models. Blood, 2016, 128, 3637-3637.	1.4	0
129	Quantitative Proteomics Identify Urinary Hgfl Protein As a Potential Marker for the Development of Chronic Kidney Disease in Sickle Cell Disease Patients. Blood, 2017, 130, 967-967.	1.4	0
130	Pulmonary Function Abnormalities in Adults with Sickle Cell Anemia. Blood, 2018, 132, 3664-3664.	1.4	0
131	Role of Automated Red Cell Exchange in Acute and Chronic Complications of Sickle Cell Disease. Blood, 2018, 132, 3674-3674.	1.4	Ο
132	Cancer Incidence in Sickle Cell Disease:an Institutional Experience. Blood, 2018, 132, 1087-1087.	1.4	0
133	Kidney Ultrasound Findings in Sickle Cell Anemia According to Kidney Disease and the APOL1 G1/G2 Risk Variants. Blood, 2018, 132, 3663-3663.	1.4	0
134	Association of Inpatient Opioid Utilization and Readmission Risk in Sickle Cell Disease. Blood, 2018, 132, 4699-4699.	1.4	0
135	Maximum Tolerated Dose Versus Fixed Low-Dose Hydroxyurea for Treatment of Adults with Sickle Cell Anemia - Retrospective Comparison of Two Studies. Blood, 2018, 132, 3656-3656.	1.4	0
136	A Safety Study of the Addition of Omacetaxine to the Standard-of-Care Induction Regimen of Cytarabine and Idarubicin in Newly-Diagnosed AML Patients. Blood, 2018, 132, 5218-5218.	1.4	0
137	Progression of Albuminuria in Sickle Cell Anemia: A Multicenter, Longitudinal Study. Blood, 2019, 134, 1004-1004.	1.4	Ο
138	Risk Factors for Kidney Disease in Hb SC and Hb Sβ+-Thalassemia Sickle Cell Disease. Blood, 2019, 134, 2299-2299.	1.4	0
139	Impact of Intravenous Opioid Shortage on Managing Pain Crisis in Sickle Cell Disease. Blood, 2019, 134, 3390-3390.	1.4	0
140	The Burden of Atrial Fibrillation in Sickle Cell Disease. Blood, 2021, 138, 3119-3119.	1.4	0
141	HIF-Mediated and Non-HIF-Mediated Differential Gene Expressions in Sickle Cell Reticulocyte and Their Impact on Clinical Manifestations. Blood, 2021, 138, 950-950.	1.4	0
142	Naloxone Use for Opioid Reversal in Patients with Sickle Cell Disease. Blood, 2021, 138, 2038-2038.	1.4	0
143	Cancer Incidence and Chemotherapy Tolerance in Patients with Sickle Cell Disease. Blood, 2020, 136, 24-25.	1.4	Ο
144	Effects of Hydroxyurea and Renin-Angiotensin Blockade on Kidney Function in Sickle Cell Disease. Blood, 2020, 136, 21-22.	1.4	0

#	Article	IF	CITATIONS
145	Mass-Spectrometry Analysis of Urinary Biomarkers of Endothelial Injury in Sickle Cell Anemia Patients. Blood, 2020, 136, 28-29.	1.4	0
146	Lower Apache II Score and Exchange Transfusions Predict Better Outcomes in the Intensive Care Unit for Patients with Sickle Cell Disease. Blood, 2020, 136, 18-19.	1.4	0
147	Correction of Point-of-Care International Normalized Ratio (INR) Values in Patients with Sickle Cell Disease. Blood, 2020, 136, 34-35.	1.4	0
148	<scp>COVID</scp> â€19 thromboembolism incidence, risk factors, and anticoagulation practices from a Chicago metropolitan <scp>US</scp> population. American Journal of Hematology, 2022, 97, .	4.1	0
149	Title is missing!. , 2020, 15, e0229710.		0
150	Title is missing!. , 2020, 15, e0229710.		0
151	Title is missing!. , 2020, 15, e0229710.		0
152	Title is missing!. , 2020, 15, e0229710.		0