Carlton Haywood Jr

List of Publications by Year
in descending order

66 papers	2,977 citations	159585 30 h-index	168389 53 g-index
67 all docs	67 docs citations	67 times ranked	2575 citing authors

#	Article	IF	Citations
1	Mortality and Access to Kidney Transplantation in Patients with Sickle Cell Diseaseâ€“Associated Kidney Failure. Clinical Journal of the American Society of Nephrology: CJASN, 2021, 16, 407-414.	4.5	14
2	Treatment of Acute Pain in Adults With Sickle Cell Disease in an Infusion Center Versus the Emergency Department. Annals of Internal Medicine, 2021, 174, 1207-1213.	3.9	16
3	Comparison of US Federal and Foundation Funding of Research for Sickle Cell Disease and Cystic Fibrosis and Factors Associated With Research Productivity. JAMA Network Open, 2020, 3, e201737.	5.9	102
4	Metabolic syndrome among adults living with sickle cell disease. Blood Cells, Molecules, and Diseases, 2019, 74, 25-29.	1.4	8
5	Do Words Matter? Stigmatizing Language and the Transmission of Bias in the Medical Record. Journal of General Internal Medicine, 2018, 33, 685-691.	2.6	217
6	Clinical Outcomes Associated With Sickle Cell Trait. Annals of Internal Medicine, 2018, 169, 619.	3.9	78
7	Increased acute care utilization in a prospective cohort of adults with sickle cell disease. Blood Advances, 2018, 2, 2412-2417.	5.2	31
8	Effect of Free Dental Services on Individuals with Sickle Cell Disease. Southern Medical Journal, 2016, 109, 576-578.	0.7	5
9	Psychometric Validation of the Insomnia Severity Index in Adults with Sickle Cell Disease. Journal of Health Care for the Poor and Underserved, 2016, 27, 209-218.	0.8	15
10	Improving Emergency Providersâ€™ Attitudes Toward Sickle Cell Patients in Pain. Journal of Pain and Symptom Management, 2016, 51, 628-632.e3.	1.2	25
11	Racial differences in acute kidney injury of hospitalized adults with diabetes. Journal of Diabetes and Its Complications, 2016, 30, 1129-1136.	2.3	18
12	Chronic Opioid Therapy and Central Sensitization in Sickle Cell Disease. American Journal of Preventive Medicine, 2016, 51, S69-S77.	3.0	65
13	The Measure of Sickle Cell Stigma: Initial findings from the Improving Patient Outcomes through Respect and Trust study. Journal of Health Psychology, 2016, 21, 808-820.	2.3	53
14	Acute Pain and Depressive Symptoms: Independent Predictors of Insomnia Symptoms among Adults with Sickle Cell Disease. Pain Management Nursing, 2016, 17, 38-46.	0.9	16
15	An Evaluation of Central Sensitization in Patients With Sickle Cell Disease. Journal of Pain, 2016, 17, 617-627.	1.4	79
16	Acute Care Utilization Is More Common in Patients with Sickle Cell Disease Who Have Chronic Complications and Chronic Pain: A Preliminary Report from the Escaped Trial. Blood, 2016, 128, 2490-2490.	1.4	1
17	Utility of the Montreal Cognitive Assessment as a Screening Test for Neurocognitive Dysfunction in Adults with Sickle Cell Disease. Southern Medical Journal, 2016, 109, 560-565.	0.7	10
18	Examination of the Patient and Hospitalization Characteristics of 30-Day SCD Readmissions. Southern Medical Journal, 2016, 109, 583-587.	0.7	9

#	Article	IF	Citations
19	The five key things you need to know to manage adult patients with sickle cell disease. Hematology American Society of Hematology Education Program, 2015, 2015, 420-425.	2.5	9
20	Sickle cell trait diagnosis: clinical and social implications. Hematology American Society of Hematology Education Program, 2015, 2015, 160-167.	2.5	89
21	The Association of Clinician Characteristics with their Attitudes Toward Patients with Sickle Cell Disease: Secondary Analyses of a Randomized Controlled Trial. Journal of the National Medical Association, 2015, 107, 89-96.	0.8	13
22	Quality Improvement Process in a Sickle Cell Infusion Center. American Journal of Medicine, 2015, 128, 541-544.	1.5	9
23	Impact of a dedicated infusion clinic for acute management of adults with sickle cell pain crisis. American Journal of Hematology, 2015, 90, 376-380.	4.1	43
24	Improving Clinician Attitudes of Respect and Trust for Persons With Sickle Cell Disease. Hospital Pediatrics, 2015, 5, 377-384.	1.3	22
25	Metabolic Syndrome Risk Among Adults Living with Sickle Cell Disease. Blood, 2015, 126, 3405-3405.	1.4	1
26	Primary Care Providersâ€™ Comfort Levels in Caring for Patients with Sickle Cell Disease. Southern Medical Journal, 2015, 108, 531-536.	0.7	28
27	Social and Behavioral Implications of National Collegiate Athletic Association Sickle Cell Trait Screening: The Athletesâ€™ Perspective. Journal of the Georgia Public Health Association, 2015, 5, .	0.1	0
28	Perceived Discrimination, Patient Trust, and Adherence to Medical Recommendations Among Persons with Sickle Cell Disease. Journal of General Internal Medicine, 2014, 29, 1657-1662.	2.6	96
29	Attitudes Toward Patients With Sickle Cell Disease in a Multicenter Sample of Emergency Department Providers. Advanced Emergency Nursing Journal, 2014, 36, 335-347.	0.5	48
30	Perceived Discrimination in Health Care Is Associated With a Greater Burden of Pain in Sickle Cell Disease. Journal of Pain and Symptom Management, 2014, 48, 934-943.	1.2	91
31	Attitudes toward clinical trials among patients with sickle cell disease. Clinical Trials, 2014, 11, 275-283.	1.6	17
32	An unequal burden: Poor patientâ€“provider communication and sickle cell disease. Patient Education and Counseling, 2014, 96, 159-164.	2.2	37
33	Screening for Neurocognitive Dysfunction in an Adult Population with Sickle Cell Disease. Blood, 2014, 124, 2717-2717.	1.4	5
34	Venous Thromboembolism in Adults with Sickle Cell Disease: A Serious and Under-recognized Complication. American Journal of Medicine, 2013, 126, 443-449.	1.5	156
35	The impact of depressive symptoms on patientâ€“provider communication in HIV care. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2013, 25, 1185-1192.	1.2	16
36	The impact of race and disease on sickle cell patient wait times in the emergency department. American Journal of Emergency Medicine, 2013, 31, 651-656.	1.6	98

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37	Emergency Provider Analgesic Practices and Attitudes Toward Patients With Sickle Cell Disease. Annals of Emergency Medicine, 2013, 62, 293-302.e10.	0.6	71
38	Disrespectful Care in the Treatment of Sickle Cell Disease Requires More Than Ethics Consultation. American Journal of Bioethics, 2013, 13, 12-14.	0.9	5
39	Measuring and explaining racial and ethnic differences in willingness to donate live kidneys in the United States. Clinical Transplantation, 2013, 27, 673-683.	1.6	30
40	A Preliminary Study of Psychiatric, Familial, and Medical Characteristics of High-utilizing Sickle Cell Disease Patients. Clinical Journal of Pain, 2013, 29, 317-323.	1.9	33
41	Perceived Discrimination In Health Care Is Associated With Daily Chronic Pain In Sickle Cell Disease. Blood, 2013, 122, 5577-5577.	1.4	1
42	Describing Adherence to Recommended Preventative Care Behaviors Among Adults with Sickle Cell Disease. Blood, 2012, 120, 2058-2058.	1.4	0
43	Respect, trust, and the management of sickle cell disease pain in hospital: comparative analysis of concern-raising behaviors, preliminary model, and agenda for international collaborative research to inform practice. Ethnicity and Health, 2011, 16, 405-421.	2.5	42
44	Religious coping and hospital admissions among adults with sickle cell disease. Journal of Behavioral Medicine, 2011, 34, 120-127.	2.1	31
45	A Video-Intervention to Improve Clinician Attitudes Toward Patients with Sickle Cell Disease: The Results of a Randomized Experiment. Journal of General Internal Medicine, 2011, 26, 518-523.	2.6	68
46	Prediction of onset and course of high hospital utilization in sickle cell disease. Journal of Hospital Medicine, 2011, 6, 248-255.	1.4	48
47	Examining the characteristics and beliefs of hydroxyurea users and nonusers among adults with sickle cell disease. American Journal of Hematology, 2011, 86, 85-87.	4.1	50
48	Patient reactions to personalized medicine vignettes: An experimental design. Genetics in Medicine, 2011, 13, 421-428.	2.4	19
49	A â€œNarcotics Contractâ€•for a Patient With Sickle Cell Disease and Chronic Pain. Pediatrics, 2011, 128, 127-131.	2.1	27
50	Do Sickle Cell Patients Wait Longer to See Physicians in the Emergency Department?. Blood, 2011, 118, 2070-2070.	1.4	2
51	Trends In the Hospital Treatment of Sickle Cell Disease-Related Priapism In the United States: 1998 to 2007,. Blood, 2011, 118, 4195-4195.	1.4	1
52	The Association of Provider Communication with Trust among Adults with Sickle Cell Disease. Journal of General Internal Medicine, 2010, 25, 543-548.	2.6	71
53	Sickle cell disease in the United States: Looking back and forward at 100 years of progress in management and survival. American Journal of Hematology, 2010, 85, 346-353.	4.1	48
54	The burden of emergency department use for sickleâ€cell disease: An analysis of the national emergency department sample database. American Journal of Hematology, 2010, 85, 797-799.	4.1	176

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55	Hospital self-discharge among adults with sickle-cell disease (SCD): Associations with trust and interpersonal experiences with care. Journal of Hospital Medicine, 2010, 5, 289-294.	1.4	46
56	Examining the Effectiveness of Hydroxyurea in People with Sickle Cell Disease. Journal of Health Care for the Poor and Underserved, 2010, 21, 277-286.	0.8	24
57	A Rose by Any Other Name: Pain Contracts/Agreements. American Journal of Bioethics, 2010, 10, 5-12.	0.9	50
58	Trends In Mortality Rates and Age of Death In Sickle Cell Disease (SCD): 1979â€“2005. Blood, 2010, 116, 736-736.	1.4	0
59	A Systematic Review of Barriers and Interventions to Improve Appropriate Use of Therapies for Sickle Cell Disease. Journal of the National Medical Association, 2009, 101, 1022-1033.	0.8	135
60	Health care provider attitudes toward patients with acute vaso-occlusive crisis due to sickle cell disease: Development of a scale. Patient Education and Counseling, 2009, 76, 272-278.	2.2	55
61	The First National Level Estimate of Emergency Department Charges for Sickle Cell Disease.. Blood, 2009, 114, 553-553.	1.4	0
62	The Burden of Hospital Readmissions for Patients with Sickle Cell Disease in California and North Carolina: 2004 to 2007.. Blood, 2009, 114, 554-554.	1.4	1
63	Hydroxyurea for Sickle Cell Disease: A Systematic Review for Efficacy and Toxicity in Children. Pediatrics, 2008, 122, 1332-1342.	2.1	189
64	Sickle Cell Disease Related Pain: Crisis and Conflict. Journal of Pain, 2006, 7, 453-458.	1.4	66
65	Hospitalization rates and costs of care of patients with sickle-cell anemia in the state of Maryland in the era of hydroxyurea. American Journal of Hematology, 2006, 81, 927-932.	4.1	140
66	Gender and Regional Differences in In-Hospital Mortality in Patients with Sickle Cell Anemia (SCA) in the United States: Results from the Nationwide Inpatient Sample (NIS) 1993-2003.. Blood, 2006, 108, 461-461.	1.4	0