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	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
2	Hydroxyurea for Sickle Cell Disease: A Systematic Review for Efficacy and Toxicity in Children. Pediatrics, 2008, 122, 1332-1342.	2.1	189
3	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
4	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
5	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
6	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
7	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
8	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
9	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
10	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
11	Sickle cell trait diagnosis: clinical and social implications. Hematology American Society of Hematology Education Program, 2015, 2015, 160-167.	2.5	89
12	An Evaluation of Central Sensitization in Patients With Sickle Cell Disease. Journal of Pain, 2016, 17, 617-627.	1.4	79
13	Clinical Outcomes Associated With Sickle Cell Trait. Annals of Internal Medicine, 2018, 169, 619.	3.9	78
14	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
15	Emergency Provider Analgesic Practices and Attitudes Toward Patients With Sickle Cell Disease. Annals of Emergency Medicine, 2013, 62, 293-302.e10.	0.6	71
16	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
17	Sickle Cell Disease Related Pain: Crisis and Conflict. Journal of Pain, 2006, 7, 453-458.	1.4	66
18	Chronic Opioid Therapy and Central Sensitization in Sickle Cell Disease. American Journal of Preventive Medicine, 2016, 51, S69-S77.	3.0	65

#	Article	IF	Citations
19	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
20	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
21	A Rose by Any Other Name: Pain Contracts/Agreements. American Journal of Bioethics, 2010, 10, 5-12.	0.9	50
22	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
23	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
24	Prediction of onset and course of high hospital utilization in sickle cell disease. Journal of Hospital Medicine, 2011, 6, 248-255.	1.4	48
25	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
26	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
27	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
28	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
29	An unequal burden: Poor patientâ€“provider communication and sickle cell disease. Patient Education and Counseling, 2014, 96, 159-164.	2.2	37
30	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
31	Religious coping and hospital admissions among adults with sickle cell disease. Journal of Behavioral Medicine, 2011, 34, 120-127.	2.1	31
32	Increased acute care utilization in a prospective cohort of adults with sickle cell disease. Blood Advances, 2018, 2, 2412-2417.	5.2	31
33	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
34	Primary Care Providersâ€™ Comfort Levels in Caring for Patients with Sickle Cell Disease. Southern Medical Journal, 2015, 108, 531-536.	0.7	28
35	A â€œNarcotics Contractâ€•for a Patient With Sickle Cell Disease and Chronic Pain. Pediatrics, 2011, 128, 127-131.	2.1	27
36	Improving Emergency Providersâ€™ Attitudes Toward Sickle Cell Patients in Pain. Journal of Pain and Symptom Management, 2016, 51, 628-632.e3.	1.2	25

#	Article	IF	Citations
37	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
38	Improving Clinician Attitudes of Respect and Trust for Persons With Sickle Cell Disease. Hospital Pediatrics, 2015, 5, 377-384.	1.3	22
39	Patient reactions to personalized medicine vignettes: An experimental design. Genetics in Medicine, 2011, 13, 421-428.	2.4	19
40	Racial differences in acute kidney injury of hospitalized adults with diabetes. Journal of Diabetes and Its Complications, 2016, 30, 1129-1136.	2.3	18
41	Attitudes toward clinical trials among patients with sickle cell disease. Clinical Trials, 2014, 11, 275-283.	1.6	17
42	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
43	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
44	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
45	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
46	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
47	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
48	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
49	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
50	Quality Improvement Process in a Sickle Cell Infusion Center. American Journal of Medicine, 2015, 128, 541-544.	1.5	9
51	Examination of the Patient and Hospitalization Characteristics of 30-Day SCD Readmissions. Southern Medical Journal, 2016, 109, 583-587.	0.7	9
52	Metabolic syndrome among adults living with sickle cell disease. Blood Cells, Molecules, and Diseases, 2019, 74, 25-29.	1.4	8
53	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
54	Effect of Free Dental Services on Individuals with Sickle Cell Disease. Southern Medical Journal, 2016, 109, 576-578.	0.7	5

#	Article	IF	Citations
55	Screening for Neurocognitive Dysfunction in an Adult Population with Sickle Cell Disease. Blood, 2014, 124, 2717-2717.	1.4	5
56	Do Sickle Cell Patients Wait Longer to See Physicians in the Emergency Department?. Blood, 2011, 118, 2070-2070.	1.4	2
57	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
58	Perceived Discrimination In Health Care Is Associated With Daily Chronic Pain In Sickle Cell Disease. Blood, 2013, 122, 5577-5577.	1.4	1
59	Metabolic Syndrome Risk Among Adults Living with Sickle Cell Disease. Blood, 2015, 126, 3405-3405.	1.4	1
60	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
61	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
62	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
63	The First National Level Estimate of Emergency Department Charges for Sickle Cell Disease.. Blood, 2009, 114, 553-553.	1.4	0
64	Trends In Mortality Rates and Age of Death In Sickle Cell Disease (SCD): 1979â€“2005. Blood, 2010, 116, 736-736.	1.4	0
65	Describing Adherence to Recommended Preventative Care Behaviors Among Adults with Sickle Cell Disease. Blood, 2012, 120, 2058-2058.	1.4	0
66	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