

Shawn M Bediako

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

992
citations

516710

16
h-index

434195

31
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33
all docs

33
docs citations

33
times ranked

994
citing authors

#	ARTICLE	IF	CITATIONS
1	Marijuana use and health behaviors in a US clinic sample of patients with sickle cell disease. PLoS ONE, 2020, 15, e0235192.	2.5	3
2	Psychosocial and Clinical Risk Factors Associated with Substance Use in Observational Cohort of Patients with Sickle Cell Disease. Substance Use and Misuse, 2020, 55, 2205-2212.	1.4	5
3	Metabolic syndrome among adults living with sickle cell disease. Blood Cells, Molecules, and Diseases, 2019, 74, 25-29.	1.4	8
4	Sickle Cell Disease, More Than Just Pain: The Mediating Role of Psychological Symptoms. Southern Medical Journal, 2019, 112, 253-258.	0.7	6
5	Communalism Moderates the Association Between Racial Centrality and Emergency Department Use for Sickle Cell Disease Pain. Journal of Black Psychology, The, 2017, 43, 659-668.	1.7	4
6	Depressive symptoms and sickle cell pain: The moderating role of internalized stigma.. Stigma and Health, 2017, 2, 271-280.	1.7	12
7	Public Support for Sickle-Cell Disease Funding: Does Race Matter?. Race and Social Problems, 2016, 8, 186-195.	2.2	3
8	Multiple Levels of Suffering. Clinical Journal of Pain, 2016, 32, 1076-1085.	1.9	52
9	The Association between Educational Attainment and Patterns of Emergency Department Utilization among Adults with Sickle Cell Disease. International Journal of Behavioral Medicine, 2016, 23, 300-309.	1.7	16
10	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
11	Interpersonal Contexts of Communication Between Sickle Cell Disease Patients and Providers. Southern Medical Journal, 2016, 109, 573-575.	0.7	1
12	Metabolic Syndrome Risk Among Adults Living with Sickle Cell Disease. Blood, 2015, 126, 3405-3405.	1.4	1
13	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
14	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
15	Attitudes toward clinical trials among patients with sickle cell disease. Clinical Trials, 2014, 11, 275-283.	1.6	17
16	An unequal burden: Poor patientâ€“provider communication and sickle cell disease. Patient Education and Counseling, 2014, 96, 159-164.	2.2	37
17	Perceived Discrimination In Health Care Is Associated With Daily Chronic Pain In Sickle Cell Disease. Blood, 2013, 122, 5577-5577.	1.4	1
18	Religious coping and hospital admissions among adults with sickle cell disease. Journal of Behavioral Medicine, 2011, 34, 120-127.	2.1	31

#	ARTICLE	IF	CITATIONS
19	Examining the characteristics and beliefs of hydroxyurea users and nonusers among adults with sickle cell disease. <i>American Journal of Hematology</i> , 2011, 86, 85-87.	4.1	50
20	Race and social attitudes about sickle cell disease. <i>Ethnicity and Health</i> , 2011, 16, 423-429.	2.5	42
21	Optimism and Perceived Stress in Sickle-Cell Disease: The Role of an Afrocultural Social Ethos. <i>Journal of Black Psychology</i> , The, 2011, 37, 234-253.	1.7	11
22	Trends In the Hospital Treatment of Sickle Cell Disease-Related Priapism In the United States: 1998 to 2007,. <i>Blood</i> , 2011, 118, 4195-4195.	1.4	1
23	The Association of Provider Communication with Trust among Adults with Sickle Cell Disease. <i>Journal of General Internal Medicine</i> , 2010, 25, 543-548.	2.6	71
24	Hospital self-discharge among adults with sickle-cell disease (SCD): Associations with trust and interpersonal experiences with care. <i>Journal of Hospital Medicine</i> , 2010, 5, 289-294.	1.4	46
25	Problematic Hospital Experiences among Adult Patients with Sickle Cell Disease. <i>Journal of Health Care for the Poor and Underserved</i> , 2010, 21, 1114-1123.	0.8	51
26	Predictors of Employment Status among African Americans with Sickle Cell Disease. <i>Journal of Health Care for the Poor and Underserved</i> , 2010, 21, 1124-1137.	0.8	18
27	Health care provider attitudes toward patients with acute vaso-occlusive crisis due to sickle cell disease: Development of a scale. <i>Patient Education and Counseling</i> , 2009, 76, 272-278.	2.2	55
28	Sickle Cell Disease in a "Postracial" America. <i>Journal of the National Medical Association</i> , 2009, 101, 1065-1066.	0.8	8
29	Racial Centrality and Health Care Use Among African American Adults With Sickle Cell Disease. <i>Journal of Black Psychology</i> , The, 2007, 33, 422-438.	1.7	14
30	Reducing and eliminating health disparities: a targeted approach. <i>Journal of the National Medical Association</i> , 2005, 97, 25-30.	0.8	7
31	Illness-specific and general perceptions of social relationships in adjustment to Rheumatoid Arthritis: The Role of Interpersonal Expectations. <i>Annals of Behavioral Medicine</i> , 2004, 28, 203-210.	2.9	22
32	Daily Mood and Stress Predict Pain, Health Care Use, and Work Activity in African American Adults With Sickle-Cell Disease.. <i>Health Psychology</i> , 2004, 23, 267-274.	1.6	151
33	Dietary behavior among African Americans: assessing cultural identity and health consciousness. <i>Ethnicity and Disease</i> , 2004, 14, 527-32.	2.3	8