

# 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  
g-index

33  
all docs

33  
docs citations

33  
times ranked

994  
citing authors

#	ARTICLE	IF	CITATIONS
1	Daily Mood and Stress Predict Pain, Health Care Use, and Work Activity in African American Adults With Sickle-Cell Disease.. Health Psychology, 2004, 23, 267-274.	1.6	151
2	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
3	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
4	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
5	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
6	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
7	Multiple Levels of Suffering. Clinical Journal of Pain, 2016, 32, 1076-1085.	1.9	52
8	Problematic Hospital Experiences among Adult Patients with Sickle Cell Disease. Journal of Health Care for the Poor and Underserved, 2010, 21, 1114-1123.	0.8	51
9	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
10	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
11	Race and social attitudes about sickle cell disease. Ethnicity and Health, 2011, 16, 423-429.	2.5	42
12	An unequal burden: Poor patientâ€™provider communication and sickle cell disease. Patient Education and Counseling, 2014, 96, 159-164.	2.2	37
13	Religious coping and hospital admissions among adults with sickle cell disease. Journal of Behavioral Medicine, 2011, 34, 120-127.	2.1	31
14	Illness-specific and general perceptions of social relationships in adjustment to Rheumatoid Arthritis: The Role of Interpersonal Expectations. Annals of Behavioral Medicine, 2004, 28, 203-210.	2.9	22
15	Predictors of Employment Status among African Americans with Sickle Cell Disease. Journal of Health Care for the Poor and Underserved, 2010, 21, 1124-1137.	0.8	18
16	Attitudes toward clinical trials among patients with sickle cell disease. Clinical Trials, 2014, 11, 275-283.	1.6	17
17	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
18	Racial Centrality and Health Care Use Among African American Adults With Sickle Cell Disease. Journal of Black Psychology, The, 2007, 33, 422-438.	1.7	14

#	ARTICLE	IF	CITATIONS
19	Depressive symptoms and sickle cell pain: The moderating role of internalized stigma.. Stigma and Health, 2017, 2, 271-280.	1.7	12
20	Optimism and Perceived Stress in Sickle-Cell Disease: The Role of an Afrocultural Social Ethos. Journal of Black Psychology, The, 2011, 37, 234-253.	1.7	11
21	Sickle Cell Disease in a "Postracial" America. Journal of the National Medical Association, 2009, 101, 1065-1066.	0.8	8
22	Metabolic syndrome among adults living with sickle cell disease. Blood Cells, Molecules, and Diseases, 2019, 74, 25-29.	1.4	8
23	Dietary behavior among African Americans: assessing cultural identity and health consciousness. Ethnicity and Disease, 2004, 14, 527-32.	2.3	8
24	Reducing and eliminating health disparities: a targeted approach. Journal of the National Medical Association, 2005, 97, 25-30.	0.8	7
25	Sickle Cell Disease, More Than Just Pain: The Mediating Role of Psychological Symptoms. Southern Medical Journal, 2019, 112, 253-258.	0.7	6
26	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
27	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
28	Public Support for Sickle-Cell Disease Funding: Does Race Matter?. Race and Social Problems, 2016, 8, 186-195.	2.2	3
29	Marijuana use and health behaviors in a US clinic sample of patients with sickle cell disease. PLoS ONE, 2020, 15, e0235192.	2.5	3
30	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
31	Perceived Discrimination In Health Care Is Associated With Daily Chronic Pain In Sickle Cell Disease. Blood, 2013, 122, 5577-5577.	1.4	1
32	Metabolic Syndrome Risk Among Adults Living with Sickle Cell Disease. Blood, 2015, 126, 3405-3405.	1.4	1
33	Interpersonal Contexts of Communication Between Sickle Cell Disease Patients and Providers. Southern Medical Journal, 2016, 109, 573-575.	0.7	1