Paula Tanabe

List of Publications by Year
in descending order

83 papers	1,911 citations	279701 23 h-index	276775 41 g-index
84 all docs	84 docs citations	84 times ranked	1613 citing authors

#	Article	IF	Citations
1	Stigma and quality of life in adults with sickle cell disease in Jamaica and the United States. Psychology, Health and Medicine, 2023, 28, 1133-1147.	1.3	4
2	Perspectives of individuals with sickle cell disease on barriers to care. PLoS ONE, 2022, 17, e0265342.	1.1	13
3	Effective Recruitment Strategies for a Sickle Cell Patient Registry Across Sites from the Sickle Cell Disease Implementation Consortium (SCDIC). Journal of Immigrant and Minority Health, 2021, 23, 725-732.	0.8	10
4	â€œPain is Subjectiveâ€• A Mixed-Methods Study of Provider Attitudes and Practices Regarding Pain Management in Sickle Cell Disease Across Three Countries. Journal of Pain and Symptom Management, 2021, 61, 474-487.	0.6	13
5	Trial design of comparing patient-specific versus weight-based protocols to treat vaso-occlusive episodes in sickle cell disease (COMPARE-VOE). Contemporary Clinical Trials, 2021, 101, 106252.	0.8	1
6	Pediatric Neurodevelopmental Delays in Children 0 to 5 Years of Age With Sickle Cell Disease: A Systematic Literature Review. Journal of Pediatric Hematology/Oncology, 2021, 43, 104-111.	0.3	6
7	Sickle cell disease is a global prototype for integrative research and healthcare. Genetics & Genomics Next, 2021, 2, e10037.	0.8	10
8	Patient Perspectives of Sickle Cell Management in the Emergency Department. Critical Care Nursing Quarterly, 2021, 44, 160-174.	0.4	12
9	Dissemination of Evidence-Based Recommendations for Sickle Cell Disease to Primary Care and Emergency Department Providers in North Carolina: A Cost Benefit Analysis. Journal of Health Economics and Outcomes Research, 2021, 8, 18-28.	0.6	2
10	Electronic Health Recordâ€“Embedded Individualized Pain Plans for Emergency Department Treatment of Vaso-occlusive Episodes in Adults With Sickle Cell Disease: Protocol for a Preimplementation and Postimplementation Study. JMIR Research Protocols, 2021, 10, e24818.	0.5	6
11	Veteransâ€™ Interpretation of Diabetes Distress in Diabetes Self-Management: Findings From Cognitive Interviews. Science of Diabetes Self-Management and Care, 2021, 47, 391-403.	0.9	4
12	A Needs Assessment of Persons With Sickle Cell Disease in a Major Medical Center in North Carolina. North Carolina Medical Journal, 2021, 82, 312-320.	0.1	0
13	Awareness and Use of the Sickle Cell Disease Toolbox by Primary Care Providers in North Carolina. Journal of Primary Care and Community Health, 2021, 12, 215013272110490.	1.0	1
14	It's Time to Provide Evidence-Based Care to Individuals with Sickle Cell Disease: A Call to Action. Journal of Emergency Nursing, 2021, 47, 684-688.	0.5	0
15	Sex-based differences in the manifestations and complications of sickle cell disease: Report from the Sickle Cell Disease Implementation Consortium. PLoS ONE, 2021, 16, e0258638.	1.1	13
16	Identifying barriers to evidence-based care for sickle cell disease: results from the Sickle Cell Disease Implementation Consortium cross-sectional survey of healthcare providers in the USA. BMJ Open, 2021, 11, e050880.	0.8	18
17	Prevalence of High BMI Status in Adults with Sickle Cell Disease. Blood, 2021, 138, 2039-2039.	0.6	0
18	Sickle-Cell Disease Co-Management, Health Care Utilization, and Hydroxyurea Use. Journal of the American Board of Family Medicine, 2020, 33, 91-105.	0.8	23

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19	Intentional and unintentional nonadherence to hydroxyurea among people with sickle cell disease: a qualitative study. Blood Advances, 2020, 4, 4463-4473.	2.5	23
20	A Survey-Based Needs Assessment of Barriers to Optimal Sickle Cell Disease Care in the Emergency Department. Annals of Emergency Medicine, 2020, 76, S64-S72.	0.3	22
21	Impact of Medicaid expansion on access and healthcare among individuals with sickle cell disease. Pediatric Blood and Cancer, 2020, 67, e28152.	0.8	15
22	Predictors of Maternal Morbidity Among Participants Enrolled in the Sickle Cell Disease Implementation Consortium Registry. Blood, 2020, 136, 3-3.	0.6	0
23	Sex Based Differences in Sickle Cell Disease. Blood, 2020, 136, 37-37.	0.6	0
24	A pilot test of the Adult Sickle Cell Quality of Life Measurement Information System (ASCQ-Me) and the Jenerette Self-Care Assessment (J-SAT) Tools in adults with sickle cell disease. Pilot and Feasibility Studies, 2019, 5, 85.	0.5	5
25	Acceptability and Feasibility of a Mindfulness-Based Intervention for Pain Catastrophizing among Persons with Sickle Cell Disease. Pain Management Nursing, 2019, 20, 261-269.	0.4	20
26	Barriers and facilitators to care for individuals with sickle cell disease in central North Carolina: The emergency department providersâ€™ perspective. PLoS ONE, 2019, 14, e0216414.	1.1	27
27	Improving the Care of Individuals With Sickle Cell Disease in the Emergency Department Using a Quality Improvement Framework. Advanced Emergency Nursing Journal, 2019, 41, 261-270.	0.2	4
28	CE: Understanding the Complications of Sickle Cell Disease. American Journal of Nursing, 2019, 119, 26-35.	0.2	32
29	Exploring Emergency Department Provider Experiences With and Perceptions of Weight-Based Versus Individualized Vaso-Occlusive Treatment Protocols in Sickle Cell Disease. Advanced Emergency Nursing Journal, 2019, 41, 86-97.	0.2	2
30	Implementation of an Emergency Department Screening and Care Management Referral Process for Patients With Sickle Cell Disease. Professional Case Management, 2019, 24, 240-248.	0.2	5
31	Transition to adult care in sickle cell disease: A longitudinal study of clinical characteristics and disease severity. Pediatric Blood and Cancer, 2019, 66, e27463.	0.8	42
32	Twelve tips for teaching a comprehensive disease-focused course with a global perspective: A sickle cell disease example. Medical Teacher, 2019, 41, 275-281.	1.0	2
33	Intravenous Fluid Boluses Are Commonly Administered to Adults with Sickle Cell Disease and Vaso-Occlusive Pain. Blood, 2019, 134, 4839-4839.	0.6	2
34	Emergency Department Encounters, Hospitalizations and ED Reliance Among Medicaid Eligible Patients with Sickle Cell Disease in North Carolina. Blood, 2019, 134, 2113-2113.	0.6	2
35	Hydroxyurea Prescription Fills and Adherence, Among Pediatric and Adult Medicaid Eligible Patients with Sickle Cell Disease in North Carolina. Blood, 2019, 134, 3391-3391.	0.6	0
36	Stigma of Sickle Cell Disease: A Systematic Review. Issues in Mental Health Nursing, 2018, 39, 675-686.	0.6	97

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37	A randomized controlled trial comparing two vasoâ€•occlusive episode (VOE) protocols in sickle cell disease (SCD). American Journal of Hematology, 2018, 93, 159-168.	2.0	37
38	Bereaved Parentsâ€™ Health Status During the First 6 Months After Their Childâ€™s Death. American Journal of Hospice and Palliative Medicine, 2018, 35, 829-839.	0.8	22
39	Social and Behavioral Factors in Sickle Cell Disease: Employment Predicts Decreased Health Care Utilization. Journal of Health Care for the Poor and Underserved, 2018, 29, 814-829.	0.4	29
40	Emergency Department (ED), ED Observation, Day Hospital, and Hospital Admissions for Adults with Sickle Cell Disease. Western Journal of Emergency Medicine, 2018, 19, 311-318.	0.6	13
41	Barriers to Care for Persons With Sickle Cell Disease. Professional Case Management, 2018, 23, 213-219.	0.2	28
42	Health Related Stigma and Quality of Life in Adults with Sickle Cell Disease in Jamaica. Blood, 2018, 132, 2285-2285.	0.6	1
43	Trajectories of Sickle Cell Disease Severity during Transition to Adult Care. Blood, 2018, 132, 318-318.	0.6	0
44	Outpatient Healthcare Utilization and Rates of Co-Management Among Medicaid Patients with Sickle Cell Disease in North Carolina. Blood, 2018, 132, 4725-4725.	0.6	0
45	A Prospective Emergency Department Quality Improvement Project to Improve the Treatment of Vaso-Occlusive Crisis in Sickle Cell Disease: Lessons Learned. Joint Commission Journal on Quality and Patient Safety, 2017, 43, 116-126.	0.4	7
46	Identifying Social-Behavioral Health Needs of Adults with Sickle Cell Disease in the Emergency Department. Journal of Emergency Nursing, 2017, 43, 444-450.	0.5	14
47	Implementation of a Schedule II patient agreement for opioids and stimulants in an adult primary care practice. Journal of Family Medicine and Primary Care, 2017, 6, 52.	0.3	3
48	Evaluation of a Sickle Cell Disease Educational Website for Emergency Providers. Advanced Emergency Nursing Journal, 2016, 38, 123-132.	0.2	4
49	Challenges in Shifting Management Responsibility From Parents to Adolescents With Sickle Cell Disease. Journal of Pediatric Nursing, 2016, 31, 678-690.	0.7	18
50	Does Attendance at a Sickle Cell Educational Conference Improve Clinician Knowledge and Attitude Toward Patients with Sickle Cell Disease?. Pain Management Nursing, 2016, 17, 226-234.	0.4	14
51	Sickle Cell Disease: A Review of Nonpharmacological Approaches for Pain. Journal of Pain and Symptom Management, 2016, 51, 163-177.	0.6	45
52	Safety of an ED High-Dose Opioid Protocol for Sickle Cell Disease Pain. Journal of Emergency Nursing, 2015, 41, 227-235.	0.5	13
53	Application of a Proactive Risk Analysis to Emergency Department Sickle Cell Care. Western Journal of Emergency Medicine, 2014, 15, 446-458.	0.6	6
54	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.	0.7	98

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55	Emergency Provider Analgesic Practices and Attitudes Toward Patients With Sickle Cell Disease. Annals of Emergency Medicine, 2013, 62, 293-302.e10.	0.3	71
56	Evaluation of a Train-the-Trainer Workshop on Sickle Cell Disease for ED Providers. Journal of Emergency Nursing, 2013, 39, 539-546.	0.5	10
57	The Emergency Department Sickle Cell Assessment of Needs and Strengths (ED-SCANS). Advanced Emergency Nursing Journal, 2013, 35, 143-153.	0.2	6
58	Sickle Cell Crisis: Safety Of a High-Dose Opioid Protocol In The Emergency Department. Blood, 2013, 122, 5579-5579.	0.6	0
59	Promoting Equity. American Journal of Medical Quality, 2012, 27, 80-82.	0.2	2
60	Adult Emergency Department Patients With Sickle Cell Pain Crisis: Results From a Quality Improvement Learning Collaborative Model to Improve Analgesic Management. Academic Emergency Medicine, 2012, 19, 430-438.	0.8	47
61	How Do Emergency Department Patients Store and Dispose of Opioids After Discharge? A Pilot Study. Journal of Emergency Nursing, 2012, 38, 273-279.	0.5	39
62	Barriers to Screening and Intervention for ED Patients at Risk For Undiagnosed or Uncontrolled Hypertension. Journal of Emergency Nursing, 2011, 37, 17-23.	0.5	14
63	A Qualitative Analysis of Best Self-management Practices: Sickle Cell Disease. Journal of the National Medical Association, 2010, 102, 1033-1041.	0.6	39
64	A Comparison of Analgesic Management for Emergency Department Patients With Sickle Cell Disease and Renal Colic. Clinical Journal of Pain, 2010, 26, 199-205.	0.8	55
65	Adult Emergency Department Patients with Sickle Cell Pain Crisis: A Learning Collaborative Model to Improve Analgesic Management. Academic Emergency Medicine, 2010, 17, 399-407.	0.8	36
66	Emergency Department Sickle Cell Assessment of Needs and Strengths (EDâ€SCANS), a Focus Group and Decision Support Tool Development Project. Academic Emergency Medicine, 2010, 17, 848-858.	0.8	15
67	Can Education and Staff-based Participatory Research Change Nursing Practice in an Era of ED Overcrowding? A Focus Group Study. Journal of Emergency Nursing, 2009, 35, 290-298.	0.5	2
68	Development of a Decision Support Tool to Guide Management of Adults with Sickle Cell Disease: The Emergency Department Sickle Cell Assessment of Strengths and Needs (ED-SCANS).. Blood, 2009, 114, 1413-1413.	0.6	0
69	Emergency Department Follow-up for Adults with Sickle Cell Disease.. Blood, 2009, 114, 241-241.	0.6	0
70	Increased Blood Pressure in the Emergency Department: Pain, Anxiety, or Undiagnosed Hypertension?. Annals of Emergency Medicine, 2008, 51, 221-229.	0.3	66
71	Pathways and Protocols for the Triage Patient with Acute Pain. , 2008, , 67-74.		0
72	Emergency Department Management of Acute Pain Episodes in Sickle Cell Disease. Academic Emergency Medicine, 2007, 14, 419-425.	0.8	92

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73	Knowledge Translation of the American College of Emergency Physicians Clinical Policy on Hypertension. Academic Emergency Medicine, 2007, 14, 1090-1096.	0.8	30
74	Emergency Severity Index Version 4: Clarifying Common Questions. Journal of Emergency Nursing, 2007, 33, 182-185.	0.5	23
75	Emergency Department Management of Acute Pain Episodes in Sickle Cell Disease. Academic Emergency Medicine, 2007, 14, 419-425.	0.8	56
76	Refining Emergency Severity Index Triage Criteria. Academic Emergency Medicine, 2005, 12, 497-501.	0.8	51
77	The Emergency Severity Index (version 3) 5-Level Triage System Scores Predict ED Resource Consumption. Journal of Emergency Nursing, 2004, 30, 22-29.	0.5	149
78	Undiagnosed Hypertension in the ED Settingâ€"An Unrecognized Opportunity by Emergency Nurses. Journal of Emergency Nursing, 2004, 30, 225-229.	0.5	28
79	Reliability and Validity of Scores on the Emergency Severity Index Version 3. Academic Emergency Medicine, 2004, 11, 59-65.	0.8	247
80	Factors Affecting Pain Scores during Female Urethral Catheterization. Academic Emergency Medicine, 2004, 11, 699-702.	0.8	14
81	Factors affecting pain scores during female urethral catheterization. Academic Emergency Medicine, 2004, 11, 699-702.	0.8	4
82	Factors Affecting the Risk of Blood Bank Specimen Hemolysis. Academic Emergency Medicine, 2003, 10, 897-900.	0.8	12
83	Factors Affecting the Risk of Blood Bank Specimen Hemolysis. Academic Emergency Medicine, 2003, 10, 897-900.	0.8	15