Paula Tanabe

List of Publications by Year
in descending order

83	1,911	279798 23	276875 41
papers	citations	h-index	g-index

84	84	84	1613
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Reliability and Validity of Scores on the Emergency Severity Index Version 3. Academic Emergency Medicine, 2004, 11, 59-65.	1.8	247
2	The Emergency Severity Index (version 3) 5-Level Triage System Scores Predict ED Resource Consumption. Journal of Emergency Nursing, 2004, 30, 22-29.	1.0	149
3	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
4	Stigma of Sickle Cell Disease: A Systematic Review. Issues in Mental Health Nursing, 2018, 39, 675-686.	1.2	97
5	Emergency Department Management of Acute Pain Episodes in Sickle Cell Disease. Academic Emergency Medicine, 2007, 14, 419-425.	1.8	92
6	Emergency Provider Analgesic Practices and Attitudes Toward Patients With Sickle Cell Disease. Annals of Emergency Medicine, 2013, 62, 293-302.e10.	0.6	71
7	Increased Blood Pressure in the Emergency Department: Pain, Anxiety, or Undiagnosed Hypertension?. Annals of Emergency Medicine, 2008, 51, 221-229.	0.6	66
8	Emergency Department Management of Acute Pain Episodes in Sickle Cell Disease. Academic Emergency Medicine, 2007, 14, 419-425.	1.8	56
9	A Comparison of Analgesic Management for Emergency Department Patients With Sickle Cell Disease and Renal Colic. Clinical Journal of Pain, 2010, 26, 199-205.	1.9	55
10	Refining Emergency Severity Index Triage Criteria. Academic Emergency Medicine, 2005, 12, 497-501.	1.8	51
11	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.	1.8	47
12	Sickle Cell Disease: A Review of Nonpharmacological Approaches for Pain. Journal of Pain and Symptom Management, 2016, 51, 163-177.	1.2	45
13	Transition to adult care in sickle cell disease: A longitudinal study of clinical characteristics and disease severity. Pediatric Blood and Cancer, 2019, 66, e27463.	1.5	42
14	A Qualitative Analysis of Best Self-management Practices: Sickle Cell Disease. Journal of the National Medical Association, 2010, 102, 1033-1041.	0.8	39
15	How Do Emergency Department Patients Store and Dispose of Opioids After Discharge? A Pilot Study. Journal of Emergency Nursing, 2012, 38, 273-279.	1.0	39
16	A randomized controlled trial comparing two vasoâ€occlusive episode (VOE) protocols in sickle cell disease (SCD). American Journal of Hematology, 2018, 93, 159-168.	4.1	37
17	Adult Emergency Department Patients with Sickle Cell Pain Crisis: A Learning Collaborative Model to Improve Analgesic Management. Academic Emergency Medicine, 2010, 17, 399-407.	1.8	36
18	CE: Understanding the Complications of Sickle Cell Disease. American Journal of Nursing, 2019, 119, 26-35.	0.4	32

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19	Knowledge Translation of the American College of Emergency Physicians Clinical Policy on Hypertension. Academic Emergency Medicine, 2007, 14, 1090-1096.	1.8	30
20	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.	0.8	29
21	Undiagnosed Hypertension in the ED Settingâ€"An Unrecognized Opportunity by Emergency Nurses. Journal of Emergency Nursing, 2004, 30, 225-229.	1.0	28
22	Barriers to Care for Persons With Sickle Cell Disease. Professional Case Management, 2018, 23, 213-219.	0.4	28
23	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.	2.5	27
24	Emergency Severity Index Version 4: Clarifying Common Questions. Journal of Emergency Nursing, 2007, 33, 182-185.	1.0	23
25	Sickle-Cell Disease Co-Management, Health Care Utilization, and Hydroxyurea Use. Journal of the American Board of Family Medicine, 2020, 33, 91-105.	1.5	23
26	Intentional and unintentional nonadherence to hydroxyurea among people with sickle cell disease: a qualitative study. Blood Advances, 2020, 4, 4463-4473.	5.2	23
27	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.	1.4	22
28	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.	0.6	22
29	Acceptability and Feasibility of a Mindfulness-Based Intervention for Pain Catastrophizing among Persons with Sickle Cell Disease. Pain Management Nursing, 2019, 20, 261-269.	0.9	20
30	Challenges in Shifting Management Responsibility From Parents to Adolescents With Sickle Cell Disease. Journal of Pediatric Nursing, 2016, 31, 678-690.	1.5	18
31	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.	1.9	18
32	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.	1.8	15
33	Impact of Medicaid expansion on access and healthcare among individuals with sickle cell disease. Pediatric Blood and Cancer, 2020, 67, e28152.	1.5	15
34	Factors Affecting the Risk of Blood Bank Specimen Hemolysis. Academic Emergency Medicine, 2003, 10, 897-900.	1.8	15
35	Factors Affecting Pain Scores during Female Urethral Catheterization. Academic Emergency Medicine, 2004, 11, 699-702.	1.8	14
36	Barriers to Screening and Intervention for ED Patients at Risk For Undiagnosed or Uncontrolled Hypertension. Journal of Emergency Nursing, 2011, 37, 17-23.	1.0	14

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37	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.	0.9	14
38	Identifying Social-Behavioral Health Needs of Adults with Sickle Cell Disease in the Emergency Department. Journal of Emergency Nursing, 2017, 43, 444-450.	1.0	14
39	Safety of an ED High-Dose Opioid Protocol for Sickle Cell Disease Pain. Journal of Emergency Nursing, 2015, 41, 227-235.	1.0	13
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.	1.1	13
41	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.	1.2	13
42	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.	2.5	13
43	Perspectives of individuals with sickle cell disease on barriers to care. PLoS ONE, 2022, 17, e0265342.	2.5	13
44	Factors Affecting the Risk of Blood Bank Specimen Hemolysis. Academic Emergency Medicine, 2003, 10, 897-900.	1.8	12
45	Patient Perspectives of Sickle Cell Management in the Emergency Department. Critical Care Nursing Quarterly, 2021, 44, 160-174.	0.8	12
46	Evaluation of a Train-the-Trainer Workshop on Sickle Cell Disease for ED Providers. Journal of Emergency Nursing, 2013, 39, 539-546.	1.0	10
47	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.	1.6	10
48	Sickle cell disease is a global prototype for integrative research and healthcare. Genetics & Genomics Next, 2021, 2, e10037.	1.5	10
49	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.	0.7	7
50	The Emergency Department Sickle Cell Assessment of Needs and Strengths (ED-SCANS). Advanced Emergency Nursing Journal, 2013, 35, 143-153.	0.5	6
51	Application of a Proactive Risk Analysis to Emergency Department Sickle Cell Care. Western Journal of Emergency Medicine, 2014, 15, 446-458.	1.1	6
52	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.	0.6	6
53	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.	1.0	6
54	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.	1.2	5

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55	Implementation of an Emergency Department Screening and Care Management Referral Process for Patients With Sickle Cell Disease. Professional Case Management, 2019, 24, 240-248.	0.4	5
56	Evaluation of a Sickle Cell Disease Educational Website for Emergency Providers. Advanced Emergency Nursing Journal, 2016, 38, 123-132.	0.5	4
57	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.	0.5	4
58	Veteransâ€™ Interpretation of Diabetes Distress in Diabetes Self-Management: Findings From Cognitive Interviews. Science of Diabetes Self-Management and Care, 2021, 47, 391-403.	1.6	4
59	Factors affecting pain scores during female urethral catheterization. Academic Emergency Medicine, 2004, 11, 699-702.	1.8	4
60	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.	2.4	4
61	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.	0.9	3
62	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.	1.0	2
63	Promoting Equity. American Journal of Medical Quality, 2012, 27, 80-82.	0.5	2
64	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.	0.5	2
65	Twelve tips for teaching a comprehensive disease-focused course with a global perspective: A sickle cell disease example. Medical Teacher, 2019, 41, 275-281.	1.8	2
66	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.	1.2	2
67	Intravenous Fluid Boluses Are Commonly Administered to Adults with Sickle Cell Disease and Vaso-Occlusive Pain. Blood, 2019, 134, 4839-4839.	1.4	2
68	Emergency Department Encounters, Hospitalizations and ED Reliance Among Medicaid Eligible Patients with Sickle Cell Disease in North Carolina. Blood, 2019, 134, 2113-2113.	1.4	2
69	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.	1.8	1
70	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.	2.1	1
71	Health Related Stigma and Quality of Life in Adults with Sickle Cell Disease in Jamaica. Blood, 2018, 132, 2285-2285.	1.4	1
72	Pathways and Protocols for the Triage Patient with Acute Pain. , 2008, , 67-74.		0

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73	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.	0.2	0
74	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.	1.0	0
75	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.	1.4	0
76	Emergency Department Follow-up for Adults with Sickle Cell Disease.. Blood, 2009, 114, 241-241.	1.4	0
77	Sickle Cell Crisis: Safety Of a High-Dose Opioid Protocol In The Emergency Department. Blood, 2013, 122, 5579-5579.	1.4	0
78	Trajectories of Sickle Cell Disease Severity during Transition to Adult Care. Blood, 2018, 132, 318-318.	1.4	0
79	Outpatient Healthcare Utilization and Rates of Co-Management Among Medicaid Patients with Sickle Cell Disease in North Carolina. Blood, 2018, 132, 4725-4725.	1.4	0
80	Hydroxyurea Prescription Fills and Adherence, Among Pediatric and Adult Medicaid Eligible Patients with Sickle Cell Disease in North Carolina. Blood, 2019, 134, 3391-3391.	1.4	0
81	Prevalence of High BMI Status in Adults with Sickle Cell Disease. Blood, 2021, 138, 2039-2039.	1.4	0
82	Predictors of Maternal Morbidity Among Participants Enrolled in the Sickle Cell Disease Implementation Consortium Registry. Blood, 2020, 136, 3-3.	1.4	0
83	Sex Based Differences in Sickle Cell Disease. Blood, 2020, 136, 37-37.	1.4	0