

Lori E Crosby

List of Publications by Year in descending order

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Version: 2024-02-01

68
papers

1,442
citations

331670

21
h-index

377865

34
g-index

74
all docs

74
docs citations

74
times ranked

1946
citing authors

#	ARTICLE	IF	CITATIONS
1	Patient-Centered eHealth Interventions for Children, Adolescents, and Adults With Sickle Cell Disease: Systematic Review. <i>Journal of Medical Internet Research</i> , 2018, 20, e10940.	4.3	119
2	Medication Adherence Among Pediatric Patients With Sickle Cell Disease: A Systematic Review. <i>Pediatrics</i> , 2014, 134, 1175-1183.	2.1	103
3	Psychopathology of Adolescent Social Phobia. <i>Journal of Psychopathology and Behavioral Assessment</i> , 2007, 29, 46-53.	1.2	89
4	Development and evaluation of iManage: A self-management app co-designed by adolescents with sickle cell disease. <i>Pediatric Blood and Cancer</i> , 2017, 64, 139-145.	1.5	84
5	Mediators and Moderators of Outcome in the Behavioral Treatment of Childhood Social Phobia. <i>Journal of the American Academy of Child and Adolescent Psychiatry</i> , 2009, 48, 945-953.	0.5	79
6	Perceived Barriers to Clinic Appointments for Adolescents With Sickle Cell Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2009, 31, 571-576.	0.6	58
7	Development and validation of the self-reported PROMIS pediatric pain behavior item bank and short form scale. <i>Pain</i> , 2017, 158, 1323-1331.	4.2	55
8	Barriers to Treatment Adherence for Pediatric Patients With Sickle Cell Disease and Their Families. <i>Children's Health Care</i> , 2009, 38, 107-122.	0.9	51
9	Optimizing Digital Integrated Care via Micro-Randomized Trials. <i>Clinical Pharmacology and Therapeutics</i> , 2018, 104, 53-58.	4.7	50
10	Qualitative Evaluation of Pediatric Pain Behavior, Quality, and Intensity Item Candidates and the PROMIS Pain Domain Framework in Children With Chronic Pain. <i>Journal of Pain</i> , 2015, 16, 1243-1255.	1.4	37
11	AAAPT Diagnostic Criteria for Acute Sickle Cell Disease Pain. <i>Journal of Pain</i> , 2019, 20, 746-759.	1.4	37
12	Applicability of the SMART Model of Transition Readiness for Sickle-Cell Disease. <i>Journal of Pediatric Psychology</i> , 2016, 41, 543-554.	2.1	36
13	Integrating Interactive Web-Based Technology to Assess Adherence and Clinical Outcomes in Pediatric Sickle Cell Disease. <i>Anemia</i> , 2012, 2012, 1-8.	1.7	33
14	Eating dinner away from home: Perspectives of middle-to high-income parents. <i>Appetite</i> , 2016, 96, 147-153.	3.7	32
15	A Biopsychosocial Model for the Management of Patients With Sickle-Cell Disease Transitioning to Adult Medical Care. <i>Advances in Therapy</i> , 2015, 32, 293-305.	2.9	31
16	Pilot of the Chronic Disease Self-Management Program for Adolescents and Young Adults With Sickle Cell Disease. <i>Journal of Adolescent Health</i> , 2017, 60, 120-123.	2.5	30
17	Commentary: Reflections on the COVID-19 Pandemic and Health Disparities in Pediatric Psychology. <i>Journal of Pediatric Psychology</i> , 2020, 45, 839-841.	2.1	29
18	Neurodevelopmental and psychosocial interventions for individuals with CHD: a research agenda and recommendations from the Cardiac Neurodevelopmental Outcome Collaborative. <i>Cardiology in the Young</i> , 2021, 31, 888-899.	0.8	27

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19	GROWTH STATUS IN CHILDREN AND ADOLESCENTS WITH SICKLE CELL DISEASE. <i>Pediatric Hematology and Oncology</i> , 2009, 26, 202-215.	0.8	25
20	Patient-Reported Outcomes for Pediatric Adherence and Self-Management: A Systematic Review. <i>Journal of Pediatric Psychology</i> , 2020, 45, 340-357.	2.1	24
21	Feasibility of Web-based Technology to Assess Adherence to Clinic Appointments in Youth With Sickle Cell Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2012, 34, e93-e96.	0.6	23
22	Translating sickle cell guidelines into practice for primary care providers with Project ECHO. <i>Medical Education Online</i> , 2016, 21, 33616.	2.6	23
23	Patient Perspectives on Gene Transfer Therapy for Sickle Cell Disease. <i>Advances in Therapy</i> , 2017, 34, 2007-2021.	2.9	22
24	Improving Sickle Cell Transitions of Care Through Health Information Technology. <i>American Journal of Preventive Medicine</i> , 2016, 51, S17-S23.	3.0	21
25	The influence of perceived racial bias and health-related stigma on quality of life among children with sickle cell disease. <i>Ethnicity and Health</i> , 2022, 27, 833-846.	2.5	21
26	The Community Leaders Institute. <i>Academic Medicine</i> , 2013, 88, 335-342.	1.6	18
27	School Performance and Disease Interference in Adolescents with Sickle Cell Disease. <i>Physical Disabilities: Education and Related Services</i> , 2015, 34, 14-30.	0.3	17
28	Shared decision making for hydroxyurea treatment initiation in children with sickle cell anemia. <i>Pediatric Blood and Cancer</i> , 2015, 62, 184-185.	1.5	16
29	Psychometric Properties of the Psychosocial Assessment Tool-General in Adolescents and Young Adults With Sickle Cell Disease. <i>Journal of Pediatric Psychology</i> , 2016, 41, 397-405.	2.1	16
30	Academic Needs in Middle School: Perspectives of Parents and Youth with Autism. <i>Journal of Autism and Developmental Disorders</i> , 2020, 50, 3126-3139.	2.7	16
31	A Virtual Reality Resident Training Curriculum on Behavioral Health Anticipatory Guidance: Development and Usability Study. <i>JMIR Pediatrics and Parenting</i> , 2021, 4, e29518.	1.6	16
32	Implementation of a Process for Initial Transcranial Doppler Ultrasonography in Children With Sickle Cell Anemia. <i>American Journal of Preventive Medicine</i> , 2016, 51, S10-S16.	3.0	14
33	Disease Self-Efficacy and Health-Related Quality of Life in Adolescents With Sickle Cell Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2020, 42, 141-144.	0.6	14
34	Understanding the Social Networks of Parents of Children with Sickle Cell Disease. <i>Journal of Health Care for the Poor and Underserved</i> , 2011, 22, 1014-1029.	0.8	13
35	Using Quality Improvement Methods to Implement an Electronic Medical Record (EMR) Supported Individualized Home Pain Management Plan for Children with Sickle Cell Disease. <i>Journal of Clinical Outcomes Management</i> , 2014, 21, 210-217.	1.7	13
36	Sleep disruption in caregivers of pediatric stem cell recipients. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26965.	1.5	11

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37	Development of a Hydroxyurea Decision Aid for Parents of Children With Sickle Cell Anemia. <i>Journal of Pediatric Hematology/Oncology</i> , 2019, 41, 56-63.	0.6	11
38	Improving self-management in adolescents with sickle cell disease. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28492.	1.5	11
39	Feasibility and acceptability of an innovative adherence intervention for young adults with childhood-onset systemic Lupus Erythematosus. <i>Pediatric Rheumatology</i> , 2020, 18, 36.	2.1	11
40	Development and Psychometric Evaluation of the PROMIS Pediatric Pain Intensity Measure in Children and Adolescents with Chronic Pain. <i>Journal of Pain</i> , 2021, 22, 48-56.	1.4	11
41	Mobile health use predicts self-efficacy and self-management in adolescents with sickle cell disease. <i>Translational Behavioral Medicine</i> , 2021, 11, 1823-1831.	2.4	10
42	A preliminary investigation of the psychometric properties of PROMIS® scales in emerging adults with sickle cell disease.. <i>Health Psychology</i> , 2019, 38, 386-390.	1.6	10
43	COVID-19 Exposure and Family Impact Scales for Adolescents and Young Adults. <i>Journal of Pediatric Psychology</i> , 2022, 47, 631-640.	2.1	9
44	Engaging Caregivers and Providers of Children With Sickle Cell Anemia in Shared Decision Making for Hydroxyurea: Protocol for a Multicenter Randomized Controlled Trial. <i>JMIR Research Protocols</i> , 2021, 10, e27650.	1.0	8
45	Allocation of Treatment Responsibility and Adherence to Hydroxyurea Among Adolescents With Sickle Cell Disease. <i>Journal of Pediatric Psychology</i> , 2019, 44, 1196-1204.	2.1	5
46	Halo or horn? A qualitative study of mothers' experiences with feeding children during the first year following bariatric surgery. <i>Appetite</i> , 2019, 142, 104366.	3.7	5
47	Quality Improvement Initiative to Reduce Nighttime Noise in a Transplantation and Cellular Therapy Unit. <i>Biology of Blood and Marrow Transplantation</i> , 2019, 25, 1844-1850.	2.0	5
48	Vitamin D supplementation and pain-related emergency department visits in children with sickle cell disease. <i>Complementary Therapies in Medicine</i> , 2020, 49, 102342.	2.7	5
49	Assessment of Competency-Based Behavioral Health Anticipatory Guidance Skills Among Pediatric Residents: the Role of Virtual Reality. <i>Journal of Technology in Behavioral Science</i> , 2022, 7, 115-124.	2.3	5
50	An Immersive Virtual Reality Curriculum for Pediatric Hematology Clinicians on Shared Decision-making for Hydroxyurea in Sickle Cell Anemia. <i>Journal of Pediatric Hematology/Oncology</i> , 2021, Publish Ahead of Print, e799-e803.	0.6	5
51	Weight status and health behaviors of adolescents and young adults with sickle cell disease: The emerging risk for obesity. <i>Pediatric Hematology and Oncology</i> , 2021, 38, 265-271.	0.8	4
52	Perceptions of a self-management intervention for adolescents with sickle cell disease.. <i>Clinical Practice in Pediatric Psychology</i> , 2022, 10, 79-90.	0.3	4
53	Chronic pediatric diseases and risk for reading difficulties: a narrative review with recommendations. <i>Pediatric Research</i> , 2022, 92, 966-978.	2.3	4
54	Considerations for Selecting Cognitive Endpoints and Psychological Patient-Reported Outcomes for Clinical Trials in Pediatric Patients With Sickle Cell Disease. <i>Frontiers in Neurology</i> , 0, 13, .	2.4	4

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55	Impact Of Use Of a Disease-Specific Patient Portal On Transition Readiness and Quality Of Life In Adolescents With Sickle Cell Disease. <i>Blood</i> , 2013, 122, 2982-2982.	1.4	3
56	School Challenges and Services Related to Executive Functioning for Fully Included Middle Schoolers With Autism. <i>Focus on Autism and Other Developmental Disabilities</i> , 2023, 38, 90-100.	1.3	3
57	An Immersive Virtual Reality Curriculum for Pediatric Providers on Shared Decision Making for Hydroxyurea. <i>Blood</i> , 2019, 134, 3402-3402.	1.4	2
58	Reply to iManage: A novel self-management app for sickle cell disease. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26358.	1.5	1
59	Pain-Related Injustice Appraisals in Youth with Sickle Cell Disease: A Preliminary Investigation. <i>Pain Medicine</i> , 2021, 22, 2207-2217.	1.9	1
60	Attitudes About COVID-19 and Health (ATTACH): Online Survey and Mixed Methods Study. <i>JMIR Mental Health</i> , 2021, 8, e29963.	3.3	1
61	Pediatric sickle cell disease. , 2020, , 185-206.		0
62	The Community Engagement and Translational Research Speaker Series: An Innovative Model of Health Education. , 2013, 03, .		0
63	Society of Pediatric Psychology Diversity Award: Training underrepresented minority students in psychology.. <i>Clinical Practice in Pediatric Psychology</i> , 2016, 4, 349-357.	0.3	0
64	Using Project Echo Telementoring to Improve Sickle Cell Disease Care in the Midwest. <i>Blood</i> , 2016, 128, 5923-5923.	1.4	0
65	Defining Sickle Cell Disease Acute Painful Episodes: The Pisces Project. <i>Blood</i> , 2018, 132, 3510-3510.	1.4	0
66	Clinical Practice Patterns for Hydroxyurea Initiation in Young Children with Sickle Cell Disease. <i>Blood</i> , 2019, 134, 4713-4713.	1.4	0
67	Effects of the COVID-19 Pandemic on Caregivers of Young Children with Sickle Cell Disease Enrolled in the Engage-HU Trial. <i>Blood</i> , 2021, 138, 1891-1891.	1.4	0
68	Addressing Recruitment Challenges in the Engage-HU Trial in Young Children with Sickle Cell Disease. <i>Blood</i> , 2020, 136, 26-27.	1.4	0