Lori E Crosby

List of Publications by Year
in descending order

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	School Challenges and Services Related to Executive Functioning for Fully Included Middle Schoolers With Autism. Focus on Autism and Other Developmental Disabilities, 2023, 38, 90-100.	1.3	3
2	The influence of perceived racial bias and health-related stigma on quality of life among children with sickle cell disease. Ethnicity and Health, 2022, 27, 833-846.	2.5	21
3	Assessment of Competency-Based Behavioral Health Anticipatory Guidance Skills Among Pediatric Residents: the Role of Virtual Reality. Journal of Technology in Behavioral Science, 2022, 7, 115-124.	2.3	5
4	Perceptions of a self-management intervention for adolescents with sickle cell disease.. Clinical Practice in Pediatric Psychology, 2022, 10, 79-90.	0.3	4
5	Chronic pediatric diseases and risk for reading difficulties: a narrative review with recommendations. Pediatric Research, 2022, 92, 966-978.	2.3	4
6	COVID-19 Exposure and Family Impact Scales for Adolescents and Young Adults. Journal of Pediatric Psychology, 2022, 47, 631-640.	2.1	9
7	Development and Psychometric Evaluation of the PROMIS Pediatric Pain Intensity Measure in Children and Adolescents with Chronic Pain. Journal of Pain, 2021, 22, 48-56.	1.4	11
8	Weight status and health behaviors of adolescents and young adults with sickle cell disease: The emerging risk for obesity. Pediatric Hematology and Oncology, 2021, 38, 265-271.	0.8	4
9	Pain-Related Injustice Appraisals in Youth with Sickle Cell Disease: A Preliminary Investigation. Pain Medicine, 2021, 22, 2207-2217.	1.9	1
10	Attitudes About COVID-19 and Health (ATTACH): Online Survey and Mixed Methods Study. JMIR Mental Health, 2021, 8, e29963.	3.3	1
11	Mobile health use predicts self-efficacy and self-management in adolescents with sickle cell disease. Translational Behavioral Medicine, 2021, 11, 1823-1831.	2.4	10
12	Engaging Caregivers and Providers of Children With Sickle Cell Anemia in Shared Decision Making for Hydroxyurea: Protocol for a Multicenter Randomized Controlled Trial. JMIR Research Protocols, 2021, 10, e27650.	1.0	8
13	Neurodevelopmental and psychosocial interventions for individuals with CHD: a research agenda and recommendations from the Cardiac Neurodevelopmental Outcome Collaborative. Cardiology in the Young, 2021, 31, 888-899.	0.8	27
14	A Virtual Reality Resident Training Curriculum on Behavioral Health Anticipatory Guidance: Development and Usability Study. JMIR Pediatrics and Parenting, 2021, 4, e29518.	1.6	16
15	An Immersive Virtual Reality Curriculum for Pediatric Hematology Clinicians on Shared Decision-making for Hydroxyurea in Sickle Cell Anemia. Journal of Pediatric Hematology/Oncology, 2021, Publish Ahead of Print, e799-e803.	0.6	5
16	Effects of the COVID-19 Pandemic on Caregivers of Young Children with Sickle Cell Disease Enrolled in the Engage-HU Trial. Blood, 2021, 138, 1891-1891.	1.4	0
17	Disease Self-Efficacy and Health-Related Quality of Life in Adolescents With Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2020, 42, 141-144.	0.6	14
18	Academic Needs in Middle School: Perspectives of Parents and Youth with Autism. Journal of Autism and Developmental Disorders, 2020, 50, 3126-3139.	2.7	16

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19	Patient-Reported Outcomes for Pediatric Adherence and Self-Management: A Systematic Review. Journal of Pediatric Psychology, 2020, 45, 340-357.	2.1	24
20	Commentary: Reflections on the COVID-19 Pandemic and Health Disparities in Pediatric Psychology. Journal of Pediatric Psychology, 2020, 45, 839-841.	2.1	29
21	Improving selfâ€management in adolescents with sickle cell disease. Pediatric Blood and Cancer, 2020, 67, e28492.	1.5	11
22	Feasibility and acceptability of an innovative adherence intervention for young adults with childhood-onset systemic Lupus Erythematosus. Pediatric Rheumatology, 2020, 18, 36.	2.1	11
23	Vitamin D supplementation and pain-related emergency department visits in children with sickle cell disease. Complementary Therapies in Medicine, 2020, 49, 102342.	2.7	5
24	Pediatric sickle cell disease. , 2020, , 185-206.		0
25	Addressing Recruitment Challenges in the Engage-HU Trial in Young Children with Sickle Cell Disease. Blood, 2020, 136, 26-27.	1.4	0
26	Development of a Hydroxyurea Decision Aid for Parents of Children With Sickle Cell Anemia. Journal of Pediatric Hematology/Oncology, 2019, 41, 56-63.	0.6	11
27	Allocation of Treatment Responsibility and Adherence to Hydroxyurea Among Adolescents With Sickle Cell Disease. Journal of Pediatric Psychology, 2019, 44, 1196-1204.	2.1	5
28	Halo or horn? A qualitative study of mothersâ€™ experiences with feeding children during the first year following bariatric surgery. Appetite, 2019, 142, 104366.	3.7	5
29	Quality Improvement Initiative to Reduce Nighttime Noise in a Transplantation and Cellular Therapy Unit. Biology of Blood and Marrow Transplantation, 2019, 25, 1844-1850.	2.0	5
30	AAAPT Diagnostic Criteria for Acute Sickle Cell Disease Pain. Journal of Pain, 2019, 20, 746-759.	1.4	37
31	A preliminary investigation of the psychometric properties of PROMISÂ® scales in emerging adults with sickle cell disease.. Health Psychology, 2019, 38, 386-390.	1.6	10
32	An Immersive Virtual Reality Curriculum for Pediatric Providers on Shared Decision Making for Hydroxyurea. Blood, 2019, 134, 3402-3402.	1.4	2
33	Clinical Practice Patterns for Hydroxyurea Initiation in Young Children with Sickle Cell Disease. Blood, 2019, 134, 4713-4713.	1.4	0
34	Optimizing Digital Integrated Care via Microâ€Randomized Trials. Clinical Pharmacology and Therapeutics, 2018, 104, 53-58.	4.7	50
35	Sleep disruption in caregivers of pediatric stem cell recipients. Pediatric Blood and Cancer, 2018, 65, e26965.	1.5	11
36	Patient-Centered eHealth Interventions for Children, Adolescents, and Adults With Sickle Cell Disease: Systematic Review. Journal of Medical Internet Research, 2018, 20, e10940.	4.3	119

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37	Defining Sickle Cell Disease Acute Painful Episodes: The Pisces Project. Blood, 2018, 132, 3510-3510.	1.4	0
38	Development and validation of the self-reported PROMIS pediatric pain behavior item bank and short form scale. Pain, 2017, 158, 1323-1331.	4.2	55
39	Reply to iManage: A novel selfâ€management app for sickle cell disease. Pediatric Blood and Cancer, 2017, 64, e26358.	1.5	1
40	Patient Perspectives on Gene Transfer Therapy for Sickle Cell Disease. Advances in Therapy, 2017, 34, 2007-2021.	2.9	22
41	Development and evaluation of iManage: A selfâ€management app coâ€designed by adolescents with sickle cell disease. Pediatric Blood and Cancer, 2017, 64, 139-145.	1.5	84
42	Pilot of the Chronic Disease Self-Management Program for Adolescents and Young Adults With Sickle Cell Disease. Journal of Adolescent Health, 2017, 60, 120-123.	2.5	30
43	Translating sickle cell guidelines into practice for primary care providers with Project ECHO. Medical Education Online, 2016, 21, 33616.	2.6	23
44	Applicability of the SMART Model of Transition Readiness for Sickle-Cell Disease. Journal of Pediatric Psychology, 2016, 41, 543-554.	2.1	36
45	Implementation of a Process for Initial Transcranial Doppler Ultrasonography in Children With Sickle Cell Anemia. American Journal of Preventive Medicine, 2016, 51, S10-S16.	3.0	14
46	Improving Sickle Cell Transitions of Care Through Health Information Technology. American Journal of Preventive Medicine, 2016, 51, S17-S23.	3.0	21
47	Eating dinner away from home: Perspectives of middle-to high-income parents. Appetite, 2016, 96, 147-153.	3.7	32
48	Psychometric Properties of the Psychosocial Assessment Tool-General in Adolescents and Young Adults With Sickle Cell Disease. Journal of Pediatric Psychology, 2016, 41, 397-405.	2.1	16
49	Society of Pediatric Psychology Diversity Award: Training underrepresented minority students in psychology.. Clinical Practice in Pediatric Psychology, 2016, 4, 349-357.	0.3	0
50	Using Project Echo Telementoring to Improve Sickle Cell Disease Care in the Midwest. Blood, 2016, 128, 5923-5923.	1.4	0
51	Shared decision making for hydroxyurea treatment initiation in children with sickle cell anemia. Pediatric Blood and Cancer, 2015, 62, 184-185.	1.5	16
52	A Biopsychosocial Model for the Management of Patients With Sickle-Cell Disease Transitioning to Adult Medical Care. Advances in Therapy, 2015, 32, 293-305.	2.9	31
53	Qualitative Evaluation of Pediatric Pain Behavior, Quality, andÂIntensity Item Candidates and the PROMIS Pain Domain Framework in Children With Chronic Pain. Journal of Pain, 2015, 16, 1243-1255.	1.4	37
54	School Performance and Disease Interference in Adolescents with Sickle Cell Disease. Physical Disabilities: Education and Related Services, 2015, 34, 14-30.	0.3	17

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55	Medication Adherence Among Pediatric Patients With Sickle Cell Disease: A Systematic Review. Pediatrics, 2014, 134, 1175-1183.	2.1	103
56	Using Quality Improvement Methods to Implement an Electronic Medical Record (EMR) Supported Individualized Home Pain Management Plan for Children with Sickle Cell Disease. Journal of Clinical Outcomes Management, 2014, 21, 210-217.	1.7	13
57	The Community Leaders Institute. Academic Medicine, 2013, 88, 335-342.	1.6	18
58	Impact Of Use Of a Disease-Specific Patient Portal On Transition Readiness and Quality Of Life In Adolescents With Sickle Cell Disease. Blood, 2013, 122, 2982-2982.	1.4	3
59	The Community Engagement and Translational Research Speaker Series: An Innovative Model of Health Education. , 2013, 03, .		0
60	Integrating Interactive Web-Based Technology to Assess Adherence and Clinical Outcomes in Pediatric Sickle Cell Disease. Anemia, 2012, 2012, 1-8.	1.7	33
61	Feasibility of Web-based Technology to Assess Adherence to Clinic Appointments in Youth With Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2012, 34, e93-e96.	0.6	23
62	Understanding the Social Networks of Parents of Children with Sickle Cell Disease. Journal of Health Care for the Poor and Underserved, 2011, 22, 1014-1029.	0.8	13
63	Barriers to Treatment Adherence for Pediatric Patients With Sickle Cell Disease and Their Families. Children's Health Care, 2009, 38, 107-122.	0.9	51
64	GROWTH STATUS IN CHILDREN AND ADOLESCENTS WITH SICKLE CELL DISEASE. Pediatric Hematology and Oncology, 2009, 26, 202-215.	0.8	25
65	Mediators and Moderators of Outcome in the Behavioral Treatment of Childhood Social Phobia. Journal of the American Academy of Child and Adolescent Psychiatry, 2009, 48, 945-953.	0.5	79
66	Perceived Barriers to Clinic Appointments for Adolescents With Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2009, 31, 571-576.	0.6	58
67	Psychopathology of Adolescent Social Phobia. Journal of Psychopathology and Behavioral Assessment, 2007, 29, 46-53.	1.2	89
68	Considerations for Selecting Cognitive Endpoints and Psychological Patient-Reported Outcomes for Clinical Trials in Pediatric Patients With Sickle Cell Disease. Frontiers in Neurology, 0, 13, .	2.4	4