Jerlym S Porter

List of Publications by Year
in descending order

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52 papers	957 citations	471509 17 h-index	477307 29 g-index
55 all docs	55 docs citations	55 times ranked	1148 citing authors

#	Article	IF	Citations
1	International and Interdisciplinary Identification of Health Care Transition Outcomes. JAMA Pediatrics, 2016, 170, 205.	6.2	193
2	Sickle Cell Clinical Research and Intervention Program (SCCRIP): A lifespan cohort study for sickle cell disease progression from the pediatric stage into adulthood. Pediatric Blood and Cancer, 2018, 65, e27228.	1.5	57
3	Gender, Ethnicity, Psychosocial Factors, and Quality of Life Among Severely Overweight, Treatment-Seeking Adolescents. Journal of Pediatric Psychology, 2007, 32, 90-94.	2.1	55
4	Psychosocial Factors and Perspectives on Weight Gain and Barriers to Weight Loss Among Adolescents Enrolled in Obesity Treatment. Journal of Clinical Psychology in Medical Settings, 2010, 17, 98-102.	1.4	40
5	A Qualitative Analysis of Best Self-management Practices: Sickle Cell Disease. Journal of the National Medical Association, 2010, 102, 1033-1041.	0.8	39
6	Development of the InCharge Health Mobile App to Improve Adherence to Hydroxyurea in Patients With Sickle Cell Disease: User-Centered Design Approach. JMIR MHealth and UHealth, 2020, 8, e14884.	3.7	38
7	A program of transition to adult care for sickle cell disease. Hematology American Society of Hematology Education Program, 2019, 2019, 496-504.	2.5	37
8	Transition From Pediatric to Adult Care in Sickle Cell Disease: Perspectives on the Family Role. Journal of Pediatric Nursing, 2014, 29, 158-167.	1.5	34
9	Hydroxyurea treatment and neurocognitive functioning in sickle cell disease from school age to young adulthood. British Journal of Haematology, 2021, 195, 256-266.	2.5	30
10	Pediatric to Adult Care Transition: Perspectives of Young Adults With Sickle Cell Disease. Journal of Pediatric Psychology, 2017, 42, 1016-1027.	2.1	27
11	Prejudice and Racism, Year 2008â€"Still Going Strong: Research on Reducing Prejudice With Recommended Methodological Advances. Journal of Counseling and Development, 2008, 86, 339-347.	2.4	26
12	APHON/ASPHO Policy Statement for the Transition of Patients With Sickle Cell Disease From Pediatric to Adult Health Care. Journal of Pediatric Oncology Nursing, 2015, 32, 355-359.	1.5	26
13	Caregiver Perspectives of Stigma Associated With Sickle Cell Disease in Adolescents. Journal of Pediatric Nursing, 2016, 31, 55-63.	1.5	24
14	Racial and ethnic disparities in neurocognitive, emotional, and qualityâ€ofâ€life outcomes in survivors of childhood cancer: A report from the Childhood Cancer Survivor Study. Cancer, 2019, 125, 3666-3677.	4.1	24
15	Intentional and unintentional nonadherence to hydroxyurea among people with sickle cell disease: a qualitative study. Blood Advances, 2020, 4, 4463-4473.	5.2	23
16	Association between hydroxycarbamide exposure and neurocognitive function in adolescents with sickle cell disease. British Journal of Haematology, 2020, 189, 1192-1203.	2.5	23
17	Paediatric to adult transition care for patients with sickle cell disease: a global perspective. Lancet Haematology,the, 2020, 7, e329-e341.	4.6	22
18	Sickle Cell Disease Patients' Perceptions of Emergency Department Pain Management. Journal of the National Medical Association, 2012, 104, 449-454.	0.8	18

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19	Genetic therapies for the first molecular disease. Journal of Clinical Investigation, 2021, 131, .	8.2	17
20	Pediatric to adult care coâ€location transitional model for youth with sickle cell disease. American Journal of Hematology, 2018, 93, E30-E32.	4.1	16
21	Self-Esteem, Teasing and Quality of Life: African American Adolescent Girls Participating in a Family-Based Pediatric Overweight Intervention. Journal of Clinical Psychology in Medical Settings, 2006, 13, 217-228.	1.4	15
22	Relations Among Teasing, Body Satisfaction, Self-Esteem, and Depression in Treatment-Seeking Obese African American Adolescents. Journal of Black Psychology, The, 2013, 39, 375-395.	1.7	15
23	Patientâ€reported neurocognitive symptoms influence instrumental activities of daily living in sickle cell disease. American Journal of Hematology, 2021, 96, 1396-1406.	4.1	15
24	Exploring Family Communication About Sickle Cell Disease in Adolescence. Journal of Pediatric Oncology Nursing, 2012, 29, 323-336.	1.5	13
25	Genetic Education and Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2014, 36, 572-577.	0.6	13
26	Cognitive performance as a predictor of healthcare transition in sickle cell disease. British Journal of Haematology, 2021, 192, 1082-1091.	2.5	13
27	Web-Based Technology to Improve Disease Knowledge Among Adolescents With Sickle Cell Disease: Pilot Study. JMIR Pediatrics and Parenting, 2020, 3, e15093.	1.6	13
28	Prematurity stereotyping and perceived vulnerability at 5â€months: Relations with mothers and their premature and fullâ€term infants at 9â€months. Journal of Reproductive and Infant Psychology, 2009, 27, 168-181.	1.8	9
29	A scoping review of transition interventions for young adults with sickle cell disease. Pediatric Blood and Cancer, 2021, 68, e29135.	1.5	9
30	Empirically Derived Profiles of Health-Related Quality of Life in Youth and Young Adults with Sickle Cell Disease. Journal of Pediatric Psychology, 2021, 46, 293-303.	2.1	9
31	Using qualitative perspectives of adolescents with sickle cell disease and caregivers to develop healthcare transition programming.. Clinical Practice in Pediatric Psychology, 2017, 5, 319-329.	0.3	7
32	Neurocognitive risk in sickle cell disease: Utilizing neuropsychology services to manage cognitive symptoms and functional limitations. British Journal of Haematology, 2022, 197, 260-270.	2.5	7
33	Neurocognitive functioning in preschool children with sickle cell disease. Pediatric Blood and Cancer, 2022, 69, e29531.	1.5	7
34	Characterizing Body Image in Youth with HIV. AIDS and Behavior, 2016, 20, 1585-1590.	2.7	6
35	The Adolescent and Caregiver Sickle Cell Disease Self-management Skills Checklist: Preliminary Reliability and Validity. Journal of Pediatric Hematology/Oncology, 2020, 42, 12-19.	0.6	5
36	Transition care continuity promotes longâ€term retention in adult care among young adults with sickle cell disease. Pediatric Blood and Cancer, 2021, 68, e29209.	1.5	5

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37	Clinic Attendance of Youth With Sickle Cell Disease on Hydroxyurea Treatment. Journal of Pediatric Hematology/Oncology, 2017, 39, 345-349.	0.6	4
38	Attention difficulties are associated with lower engagement in adult care amongst youth with sickle cell disease. British Journal of Haematology, 2020, 189, e27-e30.	2.5	4
39	Understanding barriers to transition from pediatric to adult care among young adults with sickle cell disease to develop a transition mentor program.. Clinical Practice in Pediatric Psychology, 2021, 9, 68-81.	0.3	4
40	Use of Wise Device Technology to Measure Adherence to Hydroxyurea Therapy in Youth With Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2021, 43, e19-e25.	0.6	3
41	Body Image and Risk Behaviors in Youth with HIV. AIDS Patient Care and STDs, 2017, 31, 176-181.	2.5	2
42	Feasibility of Medical Student Mentors to Improve Transition in Sickle Cell Disease. Journal of Pediatric Psychology, 2021, 46, 650-661.	2.1	2
43	Nocturnal Enuresis in Sickle Cell: Sociodemographic, Medical, and Quality of Life Factors. Journal of Pediatric Psychology, 2022, 47, 75-85.	2.1	2
44	Efficacy of a Health Care Transition Program for Patients with Sickle Cell Disease. Blood, 2018, 132, 5820-5820.	1.4	2
45	Interruption in Care Continuity during Healthcare Transition from Pediatric to Adult Care Increases Acute Care Utilization. Blood, 2018, 132, 2226-2226.	1.4	1
46	Transition Continuity Promotes Long-Term Retention in Adult Care Among Young Adults with Sickle Cell Disease. Blood, 2019, 134, 4676-4676.	1.4	1
47	Utilizing Medical Students As Patient Mentors: An Intervention to Improve Transition Readiness in Young Adults with Sickle Cell Disease. Blood, 2020, 136, 34-35.	1.4	1
48	Racial/ethnic differences in neurocognitive, emotional and quality of life outcomes in adult survivors of childhood cancer: A report from the Childhood Cancer Survivor Study (CCSS).. Journal of Clinical Oncology, 2018, 36, 10567-10567.	1.6	0
49	Neurocognitive Impairment Predicts Poor Transition Outcomes Among Patients with Sickle Cell Disease. Blood, 2019, 134, 519-519.	1.4	0
50	Social Determinants of Health and Neurocognitive Functioning in Sickle Cell Disease. Blood, 2021, 138, 2030-2030.	1.4	0
51	Impact of Gaps in Care during Adult Care Transfer in Sickle Cell Disease. Blood, 2021, 138, 2992-2992.	1.4	0
52	Food Deserts Are Associated with Acute Care Utilization Among Preschool Children with Sickle Cell Disease. Blood, 2020, 136, 19-19.	1.4	0