

Jason R Hodges

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5945675/publications.pdf>

Version: 2024-02-01

26
papers

437
citations

933410

10
h-index

752679

20
g-index

28
all docs

28
docs citations

28
times ranked

660
citing authors

#	ARTICLE	IF	CITATIONS
1	Fertility as a priority among at-risk adolescent males newly diagnosed with cancer and their parents. Supportive Care in Cancer, 2015, 23, 333-341.	2.2	102
2	Pre-existing humoral immunity to human common cold coronaviruses negatively impacts the protective SARS-CoV-2 antibody response. Cell Host and Microbe, 2022, 30, 83-96.e4.	11.0	64
3	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
4	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
5	Risk Factors for Non-initiation of the Human Papillomavirus Vaccine among Adolescent Survivors of Childhood Cancer. Cancer Prevention Research, 2013, 6, 1101-1110.	1.5	27
6	Intentional and unintentional nonadherence to hydroxyurea among people with sickle cell disease: a qualitative study. Blood Advances, 2020, 4, 4463-4473.	5.2	23
7	Association between hydroxycarbamide exposure and neurocognitive function in adolescents with sickle cell disease. British Journal of Haematology, 2020, 189, 1192-1203.	2.5	23
8	A polygenic score for acute vaso-occlusive pain in pediatric sickle cell disease. Blood Advances, 2021, 5, 2839-2851.	5.2	14
9	Cognitive performance as a predictor of healthcare transition in sickle cell disease. British Journal of Haematology, 2021, 192, 1082-1091.	2.5	13
10	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
11	Neighborhood effect and obesity in adult survivors of pediatric cancer: A report from the St. Jude lifetime cohort study. International Journal of Cancer, 2020, 147, 338-349.	5.1	12
12	Cross-reactive Antibody Response to mRNA SARS-CoV-2 Vaccine After Recent COVID-19-Specific Monoclonal Antibody Therapy. Open Forum Infectious Diseases, 2021, 8, ofab420.	0.9	12
13	Height-corrected low bone density associates with severe outcomes in sickle cell disease: SCCRIP cohort study results. Blood Advances, 2019, 3, 1476-1488.	5.2	10
14	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
15	Transition care continuity promotes long-term retention in adult care among young adults with sickle cell disease. Pediatric Blood and Cancer, 2021, 68, e29209.	1.5	5
16	Pain in Youth With Sickle Cell Disease. Clinical Journal of Pain, 2021, 37, 43-50.	1.9	5
17	Precision Medicine for Sickle Cell Disease through Whole Genome Sequencing. Blood, 2018, 132, 3641-3641.	1.4	3
18	Efficacy of a Health Care Transition Program for Patients with Sickle Cell Disease. Blood, 2018, 132, 5820-5820.	1.4	2

#	ARTICLE	IF	CITATIONS
19	Data Access and Interactive Visualization of Whole Genome Sequence of Sickle Cell Patients within the St. Jude Cloud. <i>Blood</i> , 2018, 132, 723-723.	1.4	2
20	Interruption in Care Continuity during Healthcare Transition from Pediatric to Adult Care Increases Acute Care Utilization. <i>Blood</i> , 2018, 132, 2226-2226.	1.4	1
21	Transition Continuity Promotes Long-Term Retention in Adult Care Among Young Adults with Sickle Cell Disease. <i>Blood</i> , 2019, 134, 4676-4676.	1.4	1
22	Evaluation of Factors Influencing Health Literacy in Adolescents and Adults with Sickle Cell Disease. <i>Blood</i> , 2019, 134, 2110-2110.	1.4	1
23	Fetal Hemoglobin Level during Hydroxyurea Therapy Varies By Neighborhood. <i>Blood</i> , 2018, 132, 2221-2221.	1.4	0
24	Neighborhood effects and obesity in adult survivors of pediatric cancer: A report from the St. Jude Lifetime Cohort Study.. <i>Journal of Clinical Oncology</i> , 2019, 37, e23051-e23051.	1.6	0
25	Neurocognitive Impairment Predicts Poor Transition Outcomes Among Patients with Sickle Cell Disease. <i>Blood</i> , 2019, 134, 519-519.	1.4	0
26	Food Deserts Are Associated with Acute Care Utilization Among Preschool Children with Sickle Cell Disease. <i>Blood</i> , 2020, 136, 19-19.	1.4	0