

Amy D Shapiro

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4009689/publications.pdf>

Version: 2024-02-01

18
papers

2,082
citations

1163117

8
h-index

1125743

13
g-index

18
all docs

18
docs citations

18
times ranked

1471
citing authors

#	ARTICLE	IF	CITATIONS
1	Prophylaxis versus Episodic Treatment to Prevent Joint Disease in Boys with Severe Hemophilia. <i>New England Journal of Medicine</i> , 2007, 357, 535-544.	27.0	1,681
2	Haemophilia. <i>Nature Reviews Disease Primers</i> , 2021, 7, 45.	30.5	103
3	Cardiomyopathy With Restrictive Physiology in Sickle Cell Disease. <i>JACC: Cardiovascular Imaging</i> , 2016, 9, 243-252.	5.3	97
4	Sickle cell anemia mice develop a unique cardiomyopathy with restrictive physiology. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E5182-91.	7.1	65
5	Management of people with haemophilia A undergoing surgery while receiving emicizumab prophylaxis: Real-world experience from a large comprehensive treatment centre in the US. <i>Haemophilia</i> , 2021, 27, 90-99.	2.1	37
6	The Hemophilia Joint Health Score version 2.1 Validation in Adult Patients Study: A multicenter international study. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022, 6, e12690.	2.3	37
7	Musculoskeletal ultrasound in hemophilia: Results and recommendations from a global survey and consensus meeting. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12531.	2.3	18
8	Merging into the mainstream: the evolution of the role of point-of-care musculoskeletal ultrasound in hemophilia. <i>F1000Research</i> , 2019, 8, 1029.	1.6	13
9	Factor VIII prophylaxis effects outweigh other hemostasis contributors in predicting severe haemophilia A joint outcomes. <i>Haemophilia</i> , 2019, 25, 867-875.	2.1	8
10	Therapeutic and technological advancements in haemophilia care: Quantum leaps forward. <i>Haemophilia</i> , 2022, 28, 77-92.	2.1	8
11	Iron-Refractory Microcytic Anemia as the Presenting Feature of Unicentric Castleman Disease in Children. <i>Journal of Pediatrics</i> , 2014, 164, 928-930.	1.8	7
12	Clinical features of children, adolescents, and adults with coexisting hypermobility syndromes and von Willebrand disease. <i>Pediatric Blood and Cancer</i> , 2018, 65, e27370.	1.5	6
13	A new wave in the evaluation of haemophilic arthropathy. <i>Haemophilia</i> , 2017, 23, 491-493.	2.1	2
14	Genomic Characterization Of Histiocytic Lesions Following Pediatric T-Cell Acute Lymphoblastic Leukemia. <i>Blood</i> , 2013, 122, 4940-4940.	1.4	0
15	Reactive Oxygen Species Produced by NADPH Oxidase Contribute to Cardiac Pathology in a Mouse Model of Sickle Cell Disease. <i>Blood</i> , 2016, 128, 853-853.	1.4	0
16	Risk Factors for Hospital-Acquired Venous Thromboembolism in Children: Findings from the Children's Hospital-Acquired Thrombosis (CHAT) Registry. <i>Blood</i> , 2018, 132, 142-142.	1.4	0
17	The Children's Hospital-Acquired Thrombosis (CHAT) Consortium Admission Risk-Assessment Models from Traditional Biostatistics and Machine Learning. <i>Blood</i> , 2019, 134, 635-635.	1.4	0
18	Assessing Venous Thromboembolism Risk in Critically Ill Children: A Report from the Children's Hospital-Acquired Thrombosis (CHAT) Consortium. <i>Blood</i> , 2019, 134, 1150-1150.	1.4	0