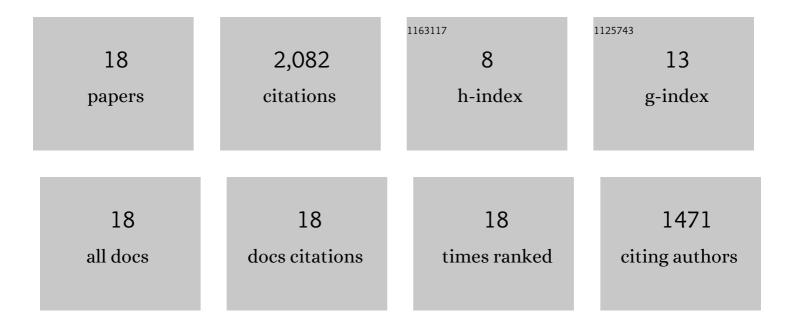
Amy D Shapiro

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4009689/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Prophylaxis versus Episodic Treatment to Prevent Joint Disease in Boys with Severe Hemophilia. New England Journal of Medicine, 2007, 357, 535-544.	27.0	1,681
2	Haemophilia. Nature Reviews Disease Primers, 2021, 7, 45.	30.5	103
3	Cardiomyopathy With Restrictive Physiology in Sickle CellÂDisease. JACC: Cardiovascular Imaging, 2016, 9, 243-252.	5.3	97
4	Sickle cell anemia mice develop a unique cardiomyopathy with restrictive physiology. Proceedings of the United States of America, 2016, 113, E5182-91.	7.1	65
5	Management of people with haemophilia A undergoing surgery while receiving emicizumab prophylaxis: Realâ€world experience from a large comprehensive treatment centre in the US. Haemophilia, 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. Research and Practice in Thrombosis and Haemostasis, 2022, 6, e12690.	2.3	37
7	Musculoskeletal ultrasound in hemophilia: Results and recommendations from a global survey and consensus meeting. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12531.	2.3	18
8	Merging into the mainstream: the evolution of the role of point-of-care musculoskeletal ultrasound in hemophilia. F1000Research, 2019, 8, 1029.	1.6	13
9	Factor VIII prophylaxis effects outweigh other hemostasis contributors in predicting severe haemophilia A joint outcomes. Haemophilia, 2019, 25, 867-875.	2.1	8
10	Therapeutic and technological advancements in haemophilia care: Quantum leaps forward. Haemophilia, 2022, 28, 77-92.	2.1	8
11	Iron-Refractory Microcytic Anemia as the Presenting Feature of Unicentric Castleman Disease in Children. Journal of Pediatrics, 2014, 164, 928-930.	1.8	7
12	Clinical features of children, adolescents, and adults with coexisting hypermobility syndromes and von Willebrand disease. Pediatric Blood and Cancer, 2018, 65, e27370.	1.5	6
13	A new wave in the evaluation of haemophilic arthropathy. Haemophilia, 2017, 23, 491-493.	2.1	2
14	Genomic Characterization Of Histiocytic Lesions Following Pediatric T-Cell Acute Lymphoblastic Leukemia. Blood, 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. Blood, 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. Blood, 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. Blood, 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. Blood, 2019, 134, 1150-1150.	1.4	0