## J Michael Soucie

List of Publications by Year in descending order

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#	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	Vascular access and increased risk of death among hemodialysis patients. Kidney International, 2002, 62, 620-626.	5.2	395
3	Occurrence of hemophilia in the United States. , 1998, 59, 288-294.		288
4	Joint range-of-motion limitations among young males with hemophilia: prevalence and risk factors. Blood, 2004, 103, 2467-2473.	1.4	225
5	A study of variations in the reported haemophilia A prevalence around the world. Haemophilia, 2010, 16, 20-32.	2.1	224
6	Impact of inhibitors on hemophilia a mortality in the <scp>U</scp> nited <scp>S</scp> tates. American Journal of Hematology, 2015, 90, 400-405.	4.1	127
7	Prophylaxis usage, bleeding rates, and joint outcomes of hemophilia, 1999 to 2010: a surveillance project. Blood, 2017, 129, 2368-2374.	1.4	121
8	Prevalence and risk factors for heart disease among males with hemophilia. American Journal of Hematology, 2005, 79, 36-42.	4.1	115
9	High oneÂyear mortality in adults with sickle cell disease and endâ€stage renal disease. British Journal of Haematology, 2012, 159, 360-367.	2.5	100
10	Associations between intracranial haemorrhage and prescribed prophylaxis in a large cohort of haemophilia patients in the United States. British Journal of Haematology, 2011, 152, 211-216.	2.5	98
11	A study of variations in the reported haemophilia B prevalence around the world. Haemophilia, 2012, 18, e91-4.	2.1	93
12	Men with severe hemophilia in the United States: birth cohort analysis of a large national database. Blood, 2016, 127, 3073-3081.	1.4	93
13	Changes in the occurrence of and risk factors for hemophilia-associated intracranial hemorrhage. American Journal of Hematology, 2001, 68, 37-42.	4.1	89
14	Renal disease among males with haemophilia. Haemophilia, 2003, 9, 703-710.	2.1	84
15	The frequency of joint hemorrhages and procedures in nonsevere hemophilia A vs B. Blood Advances, 2018, 2, 2136-2144.	5.2	69
16	Epidemiology of Device-Associated Infections Related to a Long-Term Implantable Vascular Access Device. Infection Control and Hospital Epidemiology, 1999, 20, 187-191.	1.8	57
17	Pediatric Hemophilia: A Review. Seminars in Thrombosis and Hemostasis, 2011, 37, 737-744.	2.7	56
18	The Universal Data Collection Surveillance System for Rare Bleeding Disorders. American Journal of Preventive Medicine, 2010, 38, S475-S481.	3.0	54

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19	The CDC Hemophilia A Mutation Project (CHAMP) Mutation List: A New Online Resource. Human Mutation, 2013, 34, E2382-E2392.	2.5	52
20	Increased prevalence of inhibitors in Hispanic patients with severe haemophilia A enrolled in the Universal Data Collection database. Haemophilia, 2012, 18, e260-5.	2.1	45
21	Evidence for the transmission of parvovirus B19 in patients with bleeding disorders treated with plasmaâ€derived factor concentrates in the era of nucleic acid test screening. Transfusion, 2013, 53, 1217-1225.	1.6	40
22	Physical Functioning in Boys with Hemophilia in the U.S American Journal of Preventive Medicine, 2011, 41, S360-S368.	3.0	35
23	A Prospective Comparison of Methods for Determining if Cardiovascular Disease is a Predictor of Mortality in Dialysis Patients. American Journal of Kidney Diseases, 1994, 23, 382-388.	1.9	28
24	Public health surveillance and data collection: general principles and impact on hemophilia care. Hematology, 2012, 17, s144-s146.	1.5	28
25	Evaluation of CDC's Hemophilia Surveillance Program — Universal Data Collection (1998–2011) and Community Counts (2011–2019), United States. MMWR Surveillance Summaries, 2020, 69, 1-18.	34.6	26
26	Burden of Disease Resulting from Hemophilia in the U.S American Journal of Preventive Medicine, 2010, 38, S482-S488.	3.0	25
27	Facility Mortality Rates for New End-Stage Renal Disease Patients: Implications for Quality Improvement. American Journal of Kidney Diseases, 1994, 24, 280-289.	1.9	24
28	Human parvovirus B19 in young male patients with hemophilia A: associations with treatment product exposure and joint range-of-motion limitation. Transfusion, 2004, 44, 1179-1185.	1.6	24
29	Septic arthritis in males with haemophilia. Haemophilia, 2008, 14, 494-503.	2.1	24
30	The effect of secondary prophylaxis <i>versus</i> episodic treatment on the range of motion of target joints in patients with haemophilia. British Journal of Haematology, 2013, 161, 424-433.	2.5	21
31	Prevalence of malignancies among U.S. male patients with haemophilia: a review of the Haemophilia Surveillance System. Haemophilia, 2012, 18, 532-539.	2.1	20
32	High School Completion Rates Among Men with Hemophilia. American Journal of Preventive Medicine, 2010, 38, S489-S494.	3.0	16
33	Relevance of Abusive Head Trauma to Intracranial Hemorrhages and Bleeding Disorders. Pediatrics, 2018, 141, e20173485.	2.1	15
34	Assessing Emerging Infectious Threats to Blood Safety for the Blood Disorders Community. American Journal of Preventive Medicine, 2010, 38, S468-S474.	3.0	14
35	Women and girls with haemophilia receiving care at specialized haemophilia treatment centres in the United States. Haemophilia, 2021, 27, 1037-1044.	2.1	14
36	Knowledge and Therapeutic Gaps. American Journal of Preventive Medicine, 2011, 41, S324-S331.	3.0	13

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#	Article	IF	CITATIONS
37	Characteristics of hemophilia patients with factor <scp>VIII</scp> inhibitors detected by prospective screening. American Journal of Hematology, 2015, 90, 871-876.	4.1	11
38	Potential of the Community Counts registry to characterize rare bleeding disorders. Haemophilia, 2019, 25, 1045-1050.	2.1	11
39	A Public Health Approach to the Prevention of Inhibitors in Hemophilia. American Journal of Preventive Medicine, 2014, 47, 669-673.	3.0	10
40	Executive summary of the NHLBI State of the Science (SOS) Workshop: Overview and next steps in generating a national blueprint for future research on factor VIII inhibitors. Haemophilia, 2019, 25, 610-615.	2.1	8
41	Factor VIII prophylaxis effects outweigh other hemostasis contributors in predicting severe haemophilia A joint outcomes. Haemophilia, 2019, 25, 867-875.	2.1	8
42	Origins and organization of the NHLBI State of the Science Workshop: Generating a national blueprint for future research on factor VIII inhibitors. Haemophilia, 2019, 25, 575-580.	2.1	6
43	Global Hemophilia Care: Data for Action. Annals of Internal Medicine, 2019, 171, 585.	3.9	5
44	Characteristics, complications, and sites of bleeding among infants and toddlers less than 2 years of age with VWD. Blood Advances, 2021, 5, 2079-2086.	5.2	4
45	Hemophilia without prophylaxis: Assessment of joint range of motion and factor activity. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 1035-1045.	2.3	3
46	The national haemophilia program standards, evaluation and oversight systems in the United States of America. Blood Transfusion, 2014, 12 Suppl 3, e542-8.	0.4	3
47	Occurrence rates of inherited bleeding disorders other than haemophilia and von Willebrand disease among people receiving care in specialized treatment centres in the United States. Haemophilia, 2022, 28, .	2.1	3
48	Increased Prevalence of Inhibitors in Mexican-Hispanic Patients with Severe Hemophilia A Enrolled in the Universal Data Collection Project Blood, 2009, 114, 3488-3488.	1.4	2