

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

116 papers	1,927 citations	24 h-index	40 g-index
118 ext. papers	2,192 ext. citations	2.8 avg, IF	4.8 L-index

#	Paper	IF	Citations
116	The impact of prophylactic treatment on children with severe haemophilia. <i>British Journal of Haematology</i> , 1996 , 92, 973-8	4.5	178
115	Prophylactic and therapeutic recombinant factor VIIa administration to patients with Glanzmann's thrombasthenia: results of an international survey. <i>Journal of Thrombosis and Haemostasis</i> , 2004 , 2, 1096-103	15.4	151
114	The use of recombinant factor VIIa in children with inherited platelet function disorders. <i>British Journal of Haematology</i> , 2003 , 121, 477-81	4.5	124
113	Purpura fulminans: recognition, diagnosis and management. <i>Archives of Disease in Childhood</i> , 2011 , 96, 1066-71	2.2	123
112	Health status and health-related quality of life of children with haemophilia from six West European countries. <i>Haemophilia</i> , 2004 , 10 Suppl 1, 26-33	3.3	88
111	Flow cytometric analysis of reticulated platelets: evidence for a large proportion of non-specific labelling of dense granules by fluorescent dyes. <i>British Journal of Haematology</i> , 1998 , 100, 351-7	4.5	68
110	Bruising and bleeding in infants and children--a practical approach. <i>British Journal of Haematology</i> , 2006 , 133, 221-31	4.5	61
109	Rituximab in the treatment of alloimmune factor VIII and IX antibodies in two children with severe haemophilia. <i>British Journal of Haematology</i> , 2004 , 125, 366-8	4.5	60
108	C1-inhibitor concentrate home therapy for hereditary angioedema: a viable, effective treatment option. <i>Clinical and Experimental Immunology</i> , 2007 , 147, 11-7	6.2	59
107	Central venous access devices in children with congenital coagulation disorders: complications and long-term outcome. <i>British Journal of Haematology</i> , 2000 , 110, 461-8	4.5	43
106	Factors affecting the Haemophilia Joint Health Score in children with severe haemophilia. <i>Haemophilia</i> , 2013 , 19, 626-31	3.3	42
105	The use of intermediate purity factor VIII concentrate BPL 8Y as prophylaxis and treatment in congenital thrombotic thrombocytopenic purpura. <i>British Journal of Haematology</i> , 2006 , 135, 101-4	4.5	36
104	Can early subclinical gait changes in children with haemophilia be identified using the GAITRite walkway. <i>Haemophilia</i> , 2007 , 13, 542-7	3.3	35
103	Assessment of treatment practice patterns for severe hemophilia A: a global nurse perspective. <i>Acta Haematologica</i> , 2008 , 119, 115-23	2.7	34
102	Increased burden on caregivers of having a child with haemophilia complicated by inhibitors. <i>Pediatric Blood and Cancer</i> , 2014 , 61, 706-11	3	33
101	Recombinant factor VIIa in the management of surgery and acute bleeding episodes in children with haemophilia and high responding inhibitors. <i>British Journal of Haematology</i> , 2002 , 116, 632-5	4.5	33
100	Inhibitors in nonsevere haemophilia A: outcome and eradication strategies. <i>Thrombosis and Haemostasis</i> , 2015 , 114, 46-55	7	29

99	Stem cell transplantation for children with Glanzmann thrombasthenia. <i>British Journal of Haematology</i> , 2008 , 140, 568-71	4.5	28
98	The benefits of prophylaxis: views of adolescents with severe haemophilia. <i>Haemophilia</i> , 2012 , 18, e286-93	3.3	27
97	The prevalence of the cysteine1584 variant of von Willebrand factor is increased in type 1 von Willebrand disease: co-segregation with increased susceptibility to ADAMTS13 proteolysis but not clinical phenotype. <i>British Journal of Haematology</i> , 2005 , 128, 830-6	4.5	27
96	The impact of sport on children with haemophilia. <i>Haemophilia</i> , 2012 , 18, 898-905	3.3	26
95	Assessment and management of pain in children and adolescents with bleeding disorders: a cross-sectional study from three haemophilia centres. <i>Haemophilia</i> , 2016 , 22, 65-71	3.3	25
94	The impact of sport on health status, psychological well-being and physical performance of adults with haemophilia. <i>Haemophilia</i> , 2016 , 22, 521-30	3.3	24
93	The experience of girls and young women with inherited bleeding disorders. <i>Haemophilia</i> , 2013 , 19, e276-81	3.3	24
92	Self-management and skills acquisition in boys with haemophilia. <i>Health Expectations</i> , 2015 , 18, 1105-13	3.7	22
91	Hermansky-Pudlak syndrome: infrequent bleeding and first report of Turkish and Pakistani kindreds. <i>Archives of Disease in Childhood</i> , 2002 , 86, 297-301	2.2	22
90	Obesity in the global haemophilia population: prevalence, implications and expert opinions for weight management. <i>Obesity Reviews</i> , 2018 , 19, 1569-1584	10.6	21
89	Pattern of bleeding in a large prospective cohort of haemophilia A patients: A three-year follow-up of the AHEAD (Advate in HaEmophilia A outcome Database) study. <i>Haemophilia</i> , 2018 , 24, 85-96	3.3	20
88	Compliance, concordance and adherence: what are we talking about?. <i>Haemophilia</i> , 2014 , 20, 601-3	3.3	19
87	Long-term consequences of intracranial haemorrhage in children with haemophilia. <i>Haemophilia</i> , 2009 , 15, 184-92	3.3	17
86	Prophylaxis with prothrombin complex concentrate in four children with severe congenital factor X deficiency. <i>Haemophilia</i> , 2009 , 15, 401-3	3.3	17
85	Unusual presentation of factor XIII deficiency. <i>Haemophilia</i> , 2002 , 8, 703-5	3.3	15
84	Treatment burden, haemostatic strategies and real world inhibitor screening practice in non-severe haemophilia A. <i>British Journal of Haematology</i> , 2017 , 176, 796-804	4.5	14
83	Study of physical function in adolescents with haemophilia: The SO-FIT study. <i>Haemophilia</i> , 2017 , 23, 918-925	3.3	13
82	Factor XIII deficiency: new nonsense and deletion mutations in the human factor XIII A gene. <i>Haematologica</i> , 2005 , 90, 1718-20	6.6	13

81	Wilate use in 47 children with von Willebrand disease: the North London paediatric haemophilia network experience. <i>Haemophilia</i> , 2015 , 21, e44-50	3.3	12
80	Physiotherapy interventions for pain management in haemophilia: A systematic review. <i>Haemophilia</i> , 2020 , 26, 667-684	3.3	12
79	Multidisciplinary teams in the United Kingdom: problems and solutions. <i>Journal of Pediatric Nursing</i> , 2000 , 15, 131-4	2.2	12
78	HOW Patients view Extended half-life products: Impressions from real-world experience (The HOPE study). <i>Haemophilia</i> , 2019 , 25, 814-820	3.3	11
77	Social networking for adolescents with severe haemophilia. <i>Haemophilia</i> , 2012 , 18, e290-6	3.3	11
76	Minimizing joint damage: the role of nurses in promoting adherence to hemophilia treatment. <i>Orthopaedic Nursing</i> , 2010 , 29, 193-200; quiz 201-2	0.9	11
75	Health-Related Quality of Life and Psychological Aspects of Adults With Hemophilia in Iran. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2018 , 24, 1073-1081	3.3	10
74	An action research study to explore the nature of the nurse consultant role in the care of children and young people. <i>Journal of Clinical Nursing</i> , 2013 , 22, 201-10	3.2	10
73	The use of recombinant factor VIIa in a patient with severe Glanzmann's thrombasthenia to facilitate insertion of a Port-a-Cath. <i>Blood Coagulation and Fibrinolysis</i> , 1999 , 10, 447-8	1	10
72	The impact of psychosocial determinants on caregivers' burden of children with haemophilia (results of the BBC study). <i>Haemophilia</i> , 2019 , 25, 424-432	3.3	9
71	Management of surgical procedures in children with severe FV deficiency: experience of 13 surgeries. <i>Haemophilia</i> , 2013 , 19, 256-8	3.3	9
70	Intranasal desmopressin (Octim): a safe and efficacious treatment option for children with bleeding disorders. <i>Haemophilia</i> , 2007 , 13, 548-51	3.3	9
69	The incidence, risk and functional outcomes of intracranial haemorrhage in children with inherited bleeding disorders at one haemophilia center. <i>Haemophilia</i> , 2016 , 22, 556-63	3.3	9
68	Spirituality Experiences in Hemophilia Patients: A Phenomenological Study. <i>Journal of Religion and Health</i> , 2019 , 58, 992-1002	2.6	9
67	The burden of bleeds and other clinical determinants on caregivers of children with haemophilia (the BBC Study). <i>Haemophilia</i> , 2019 , 25, 416-423	3.3	8
66	Supporting adherence and improving quality of life in haemophilia care. <i>British Journal of Nursing</i> , 2013 , 22, 692	0.7	8
65	The impact of clinical practice on the outcome of central venous access devices in children with haemophilia. <i>Haemophilia</i> , 2017 , 23, e276-e281	3.3	7
64	Education needs of nurses in thrombosis and hemostasis: An international, mixed-methods study. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2019 , 3, 99-108	5.1	7

63	Prophylaxis for hemophilia A without inhibitors: treatment options and considerations. <i>Expert Review of Hematology</i> , 2020 , 13, 731-743	2.8	7
62	Burden on parents of children with haemophilia: The impact of sociodemographic and child's medical condition. <i>Journal of Clinical Nursing</i> , 2019 , 28, 4077-4086	3.2	7
61	What is a nurse-led service? A discussion paper. <i>The Journal of Haemophilia Practice</i> , 2017 , 4, 4-13	0.2	7
60	Potential for development of haemophilia link nurse role within UK hospitals. <i>Haemophilia</i> , 2013 , 19, 578-82	3.3	6
59	'Just an unfortunate coincidence': children's understanding of haemophilia genetics and inheritance. <i>Haemophilia</i> , 2011 , 17, 470-5	3.3	6
58	Children's preferences of transfer devices for reconstitution of factors VIII and IX for the treatment of haemophilia. <i>Haemophilia</i> , 2009 , 15, 159-67	3.3	6
57	Evaluating a self infusion device for children with haemophilia. <i>Paediatric Nursing</i> , 2006 , 18, 19-20		6
56	Immune tolerance in children with factors VIII and IX inhibitors: a single centre experience. <i>Haemophilia</i> , 2005 , 11, 340-5	3.3	6
55	I can always rely on them—the importance of social support for boys with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2013 , 1, 10-16	0.2	6
54	A core competency framework for haemophilia nurses in the UK. <i>The Journal of Haemophilia Practice</i> , 2013 , 1, 32-36	0.2	6
53	Successful use of BPL Factor X concentrate in a child with severe factor X deficiency. <i>The Journal of Haemophilia Practice</i> , 2013 , 1, 8-10	0.2	6
52	Physical function and quality of life in adolescents with haemophilia (SO-FIT study). <i>The Journal of Haemophilia Practice</i> , 2013 , 1, 11-14	0.2	6
51	HaemophiliaLIVE: an ethnographic study on the impact of haemophilia on daily life. <i>The Journal of Haemophilia Practice</i> , 2014 , 1, 14-20	0.2	6
50	Caregiver burden in haemophilia: results from a single UK centre. <i>The Journal of Haemophilia Practice</i> , 2017 , 4, 40-48	0.2	6
49	Pain Experience in Hemophilia Patients: A Hermeneutic Phenomenological Study. <i>International Journal of Community Based Nursing and Midwifery</i> , 2016 , 4, 309-319	1.6	6
48	European principles of care for women and girls with inherited bleeding disorders. <i>Haemophilia</i> , 2021 , 27, 837-847	3.3	6
47	Validation of the Haemophilia & Exercise Project-Test-Questionnaire (HEP-Test-Q)-An instrument for the assessment of subjective physical functioning in children with haemophilia. <i>Haemophilia</i> , 2018 , 24, 888-895	3.3	5
46	Rituximab Is Effective and Well Tolerated in Children with Hemophila and Factor VIII Inhibitors.. <i>Blood</i> , 2006 , 108, 1031-1031	2.2	5

45	Pain: the voiceless scream in every haemophilia patient's life. <i>The Journal of Haemophilia Practice</i> , 2016 , 3, 8-13	0.2	5
44	Multimethodology research with boys with severe haemophilia. <i>Nurse Researcher</i> , 2013 , 20, 40-4	1	5
43	The UK haemophilia specialist nurse: Competencies fit for practice in the 21st century. <i>Haemophilia</i> , 2020 , 26, 622-630	3.3	5
42	Parental perspectives on gene therapy for children with haemophilia: The Exigency study. <i>Haemophilia</i> , 2021 , 27, 120-128	3.3	5
41	Prophylactic bypassing agent use before and during immune tolerance induction in patients with haemophilia A and inhibitors to FVIII. <i>Haemophilia</i> , 2018 , 24, 570-577	3.3	4
40	Assessing the supportive care needs of parents with a child with a bleeding disorder using the Parental Needs Scale for Rare Diseases (PNS-RD): A single-centre pilot study. <i>Haemophilia</i> , 2019 , 25, 831-837	3.3	4
39	Why don't haemophilia nurses do research?. <i>Haemophilia</i> , 2012 , 18, 540-3	3.3	4
38	Haemophilia A: meeting the needs of individual patients. <i>British Journal of Nursing</i> , 2007 , 16, 987-93	0.7	4
37	The top 10 research priorities in bleeding disorders: a James Lind Alliance Priority Setting Partnership. <i>British Journal of Haematology</i> , 2019 , 186, e98-e100	4.5	3
36	Evaluation of a pre-filled diluent syringe (MixPro [®]) among patient/carer users and nurses. <i>The Journal of Haemophilia Practice</i> , 2018 , 5, 12-23	0.2	3
35	The impact of factor infusion frequency on health-related quality of life in people with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2020 , 7, 102-109	0.2	3
34	The experiences and beliefs of people with severe haemophilia and healthcare professionals on pain management, and their views of using exercise as an aspect of intervention: a qualitative study.. <i>Disability and Rehabilitation</i> , 2021 , 1-9	2.4	3
33	The lived experience of a novel disruptive therapy in a group of men and boys with haemophilia A with inhibitors: Emi & Me. <i>Health Expectations</i> , 2021 ,	3.7	2
32	Patient and caregiver perceptions of a pre-filled diluent syringe (MixPro [®]). <i>The Journal of Haemophilia Practice</i> , 2016 , 3, 33-38	0.2	2
31	The impact on parents of having a child with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2016 , 3, 4-14	0.2	2
30	Switching factor products: nurses' experience with NovoEight. <i>The Journal of Haemophilia Practice</i> , 2020 , 7, 59-69	0.2	2
29	An exploration of why men with severe haemophilia might not want gene therapy: The exigency study. <i>Haemophilia</i> , 2021 , 27, 760-768	3.3	2
28	How does a lifetime of painful experiences influence sensations and beliefs about pain in adults with severe haemophilia? A qualitative study.. <i>Disability and Rehabilitation</i> , 2021 , 1-8	2.4	2

27	Pharmacokinetics, Safety, and Efficacy of Recombinant Factor VIII Fc Fusion Protein: A Practical Review. <i>Journal of Infusion Nursing</i> , 2017 , 40, 65-75	1	1
26	Experience of recombinant-activated factor VII to control bleeding in children with Glanzmann's thrombasthenia. <i>Comparative Clinical Pathology</i> , 2007 , 16, 167-172	0.9	1
25	Effectiveness and Safety Outcomes in Patients with Hemophilia A Receiving Antihemophilic Factor (Recombinant) for at Least 5 Years in a Real-World Setting: 6-Year Interim Analysis of the Ahead International and German Studies. <i>Blood</i> , 2020 , 136, 1-1	2.2	1
24	The lived experience of women with a bleeding disorder: A systematic review.. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022 , 6, e12652	5.1	1
23	Experience of switching to NovoEight: views of people with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2020 , 7, 70-77	0.2	1
22	Project GYM: A randomized feasibility study investigating effect on motivation of personal trainer-led exercise in young men with hemophilia. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021 , 5, e12613	5.1	1
21	Pain self-management experiences in haemophilia patients: a qualitative study. <i>The Journal of Haemophilia Practice</i> , 2018 , 5, 76-82	0.2	1
20	A multi-country snapshot study of pain in people with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2020 , 7, 143-149	0.2	1
19	Nurse-Led Effort in Developing, Implementing and Evaluating Healthcare Transition and Promoting Developmentally Appropriate Healthcare for Young People with Haemophilia 2020 , 257-280		1
18	Research Engagement in Pediatric Nursing Practice: Career Pathways are the Biggest Barrier. <i>Journal of Pediatrics</i> , 2020 , 221S, S62-S63	3.6	1
17	Shortening the paediatric Haemophilia Activities List (pedHAL) based on pooled data from international studies. <i>Haemophilia</i> , 2021 , 27, 305-313	3.3	1
16	Real-World Data on the Effectiveness and Safety of wilate for the Treatment of von Willebrand Disease. <i>TH Open</i> , 2021 , 5, e264-e272	2.7	1
15	The experiences of people with haemophilia and their families of gene therapy in a clinical trial setting: regaining control, the Exigency study.. <i>Orphanet Journal of Rare Diseases</i> , 2022 , 17, 155	4.2	1
14	Fitness enhances psychosocial well-being and self-confidence in young men with hemophilia: Results from Project GYM. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021 , 5, e12622	5.1	0
13	You're only a carrier... Women and the language of haemophilia. <i>The Journal of Haemophilia Practice</i> , 2021 , 8, 128-132	0.2	0
12	Haemophilia nursing practice: A global survey of roles and responsibilities. <i>The Journal of Haemophilia Practice</i> , 2016 , 3, 29-33	0.2	0
11	The Kids' immune thrombocytopenia Tool is not suitable for assessing quality of life in children with platelet function disorders. <i>Haemophilia</i> , 2018 , 24, e259-e261	3.3	0
10	Addressing unmet needs in rare bleeding disorders: selected poster extracts of recent research in hemophilia A and von Willebrand disease presented at the 14th Annual Congress of the European Association for Haemophilia and Allied Disorders (EAHAD) (Feb 3-5, 2021; virtual congress). <i>Expert Review of Hematology</i> , 2021 , 14, 1-18	2.8	0

- 9 Final Results of the Prospective ADVATE Immune Tolerance Induction Registry (PAIR) Study with Plasma- and Albumin-Free Recombinant Factor VIII. *Journal of Blood Medicine*, **2021**, 12, 991-1001 2.3
- 8 Living with an inhibitor: Results from the Study of Haemophilia Experiences, Results and Opportunities in Children and young adults with long-standing inhibitors (the SO-HEROIC study). *The Journal of Haemophilia Practice*, **2018**, 5, 24-34 0.2
- 7 You think you know the patient inside-out. But do you know the outside-in?. *The Journal of Haemophilia Practice*, **2018**, 5, 130-137 0.2
- 6 Now and then, and half a world away. *The Journal of Haemophilia Practice*, **2019**, 6, s36-s38 0.2
- 5 Ahead Study: A 3 Year Follow-up of 522 Severe and Moderate Hemophilia a Patients. *Blood*, **2016**, 128, 3786-3786 2.2
- 4 Observational Study on Safety and Efficacy of Factor Replacement Therapy in the Management of Von Willebrand Disease Patients - Results from an Ongoing Study with a VWF/FVIII Concentrate. *Blood*, **2016**, 128, 4968-4968 2.2
- 3 A Neonate with Type 3 Von Willebrands Disease and Life Threatening Bleeding Treated Successfully with a FVIII/VWF Concentrate (wilate[®]). *Blood*, **2012**, 120, 4633-4633 2.2
- 2 Haemophilia. *Professional Nurse (london, England)*, **2002**, 17, 463-4
- 1 It's a way of life—Results from the Perceptions of Pain in Haemophilia study. *The Journal of Haemophilia Practice*, **2021**, 8, 145-154 0.2