

Kate Khair

List of Publications by Year in descending order

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Version: 2024-02-01

116
papers

2,521
citations

201658

27
h-index

223791

46
g-index

118
all docs

118
docs citations

118
times ranked

1775
citing authors

#	ARTICLE	IF	CITATIONS
1	The impact of prophylactic treatment on children with severe haemophilia. <i>British Journal of Haematology</i> , 1996, 92, 973-978.	2.5	192
2	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, 1096-1103.	3.8	179
3	Purpura fulminans: recognition, diagnosis and management. <i>Archives of Disease in Childhood</i> , 2011, 96, 1066-1071.	1.9	166
4	The use of recombinant factor VIIa in children with inherited platelet function disorders. <i>British Journal of Haematology</i> , 2003, 121, 477-481.	2.5	154
5	Health status and health-related quality of life of children with haemophilia from six West European countries. <i>Haemophilia</i> , 2004, 10, 26-33.	2.1	95
6	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, 351-357.	2.5	91
7	Bruising and bleeding in infants and children - a practical approach. <i>British Journal of Haematology</i> , 2006, 133, 221-231.	2.5	74
8	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, 366-368.	2.5	73
9	C1-inhibitor concentrate home therapy for hereditary angioedema: a viable, effective treatment option. <i>Clinical and Experimental Immunology</i> , 2006, 147, 061124011224001-???	2.6	73
10	Factors affecting the Haemophilia Joint Health Score in children with severe haemophilia. <i>Haemophilia</i> , 2013, 19, 626-631.	2.1	57
11	Central venous access devices in children with congenital coagulation disorders: complications and long-term outcome. <i>British Journal of Haematology</i> , 2000, 110, 461-468.	2.5	50
12	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, 101-104.	2.5	47
13	Increased burden on caregivers of having a child with haemophilia complicated by inhibitors. <i>Pediatric Blood and Cancer</i> , 2014, 61, 706-711.	1.5	41
14	Assessment of Treatment Practice Patterns for Severe Hemophilia A: A Global Nurse Perspective. <i>Acta Haematologica</i> , 2008, 119, 115-123.	1.4	40
15	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, 632-635.	2.5	39
16	Can early subclinical gait changes in children with haemophilia be identified using the GAITRite [®] 1/2 walkway. <i>Haemophilia</i> , 2007, 13, 542-547.	2.1	39
17	The impact of sport on children with haemophilia. <i>Haemophilia</i> , 2012, 18, 898-905.	2.1	38
18	Obesity in the global haemophilia population: prevalence, implications and expert opinions for weight management. <i>Obesity Reviews</i> , 2018, 19, 1569-1584.	6.5	34

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19	Inhibitors in nonsevere haemophilia A: outcome and eradication strategies. <i>Thrombosis and Haemostasis</i> , 2015, 114, 46-55.	3.4	33
20	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.	2.1	33
21	Pattern of bleeding in a large prospective cohort of haemophilia A patients: A three-year follow-up of the <sc>AHEAD</sc> (Advate in HaEmophilia A outcome Database) study. <i>Haemophilia</i> , 2018, 24, 85-96.	2.1	33
22	The benefits of prophylaxis: views of adolescents with severe haemophilia. <i>Haemophilia</i> , 2012, 18, e286-9.	2.1	32
23	Self-management and skills acquisition in boys with haemophilia. <i>Health Expectations</i> , 2015, 18, 1105-1113.	2.6	32
24	The impact of sport on health status, psychological well-being and physical performance of adults with haemophilia. <i>Haemophilia</i> , 2016, 22, 521-530.	2.1	32
25	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.	1.7	30
26	Stem cell transplantation for children with Glanzmann thrombasthenia. <i>British Journal of Haematology</i> , 2008, 140, 568-571.	2.5	29
27	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, 830-836.	2.5	28
28	The experience of girls and young Women with inherited bleeding disorders. <i>Haemophilia</i> , 2013, 19, e276-81.	2.1	28
29	Hermansky-Pudlak syndrome: infrequent bleeding and first report of Turkish and Pakistani kindreds. <i>Archives of Disease in Childhood</i> , 2002, 86, 297-301.	1.9	27
30	Physiotherapy interventions for pain management in haemophilia: A systematic review. <i>Haemophilia</i> , 2020, 26, 667-684.	2.1	27
31	Compliance, concordance and adherence: what are we talking about?. <i>Haemophilia</i> , 2014, 20, 601-603.	2.1	23
32	European principles of care for women and girls with inherited bleeding disorders. <i>Haemophilia</i> , 2021, 27, 837-847.	2.1	23
33	Long-term consequences of intracranial haemorrhage in children with haemophilia. <i>Haemophilia</i> , 2009, 15, 184-192.	2.1	21
34	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.	2.5	21
35	Prophylaxis with prothrombin complex concentrate in four children with severe congenital factor X deficiency. <i>Haemophilia</i> , 2009, 15, 401-403.	2.1	18
36	Study of physical function in adolescents with haemophilia: The SOFIT study. <i>Haemophilia</i> , 2017, 23, 918-925.	2.1	18

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37	The experiences of people with haemophilia and their families of gene therapy in a clinical trial setting: regaining control, the Exigency study. Orphanet Journal of Rare Diseases, 2022, 17, 155.	2.7	18
38	Multidisciplinary teams in the United Kingdom: Problems and solutions. Journal of Pediatric Nursing, 2000, 15, 131-134.	1.5	17
39	Wilate use in 47 children with von Willebrand disease: the North London paediatric haemophilia network experience. Haemophilia, 2015, 21, e44-50.	2.1	17
40	Unusual presentation of factor XIII deficiency. Haemophilia, 2002, 8, 703-705.	2.1	16
41	The incidence, risk and functional outcomes of intracranial haemorrhage in children with inherited bleeding disorders at one haemophilia center. Haemophilia, 2016, 22, 556-563.	2.1	16
42	The burden of bleeds and other clinical determinants on caregivers of children with haemophilia (the Tj ETQq0 0 0 rgBT /Overlock 10 T	2.1	16
43	Prophylaxis for hemophilia A without inhibitors: treatment options and considerations. Expert Review of Hematology, 2020, 13, 731-743.	2.2	16
44	Social networking for adolescents with severe haemophilia. Haemophilia, 2012, 18, e290-6.	2.1	15
45	The use of recombinant factor Vila in a patient with severe Glanzmann??s thrombasthenia to facilitate insertion of a Port-a-Cath??. Blood Coagulation and Fibrinolysis, 1999, 10, 447-448.	1.0	13
46	HOw Patients view Extended halfâ€life products: Impressions from realâ€world experience (The HOPE) Tj ETQq0 0 0 rgBT /Overlock 10 T	2.1	13
47	The impact of psychosocial determinants on caregiversâ€™ burden of children with haemophilia (results) Tj ETQq1 1 0.784314 rgBT /Ov	2.1	13
48	Spirituality Experiences in Hemophilia Patients: A Phenomenological Study. Journal of Religion and Health, 2019, 58, 992-1002.	1.7	13
49	Minimizing Joint Damage. Orthopaedic Nursing, 2010, 29, 193-200.	0.4	13
50	Factor XIII deficiency: new nonsense and deletion mutations in the human factor XIII gene. Haematologica, 2005, 90, 1718-20.	3.5	13
51	Burden on parents of children with haemophilia: The impact of sociodemographic and child's medical condition. Journal of Clinical Nursing, 2019, 28, 4077-4086.	3.0	12
52	Parental perspectives on gene therapy for children with haemophilia: The Exigency study. Haemophilia, 2021, 27, 120-128.	2.1	12
53	An exploration of why men with severe haemophilia might not want gene therapy: The exigency study. Haemophilia, 2021, 27, 760-768.	2.1	12
54	An action research study to explore the nature of the nurse consultant role in the care of children and young people. Journal of Clinical Nursing, 2013, 22, 201-210.	3.0	11

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55	Management of surgical procedures in children with severe FV deficiency: experience of 13 surgeries. <i>Haemophilia</i> , 2013, 19, 256-258.	2.1	11
56	Education needs of nurses in thrombosis and hemostasis: An international, mixedâ€methods study. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2019, 3, 99-108.	2.3	11
57	What is a nurse-led service? A discussion paper. <i>The Journal of Haemophilia Practice</i> , 2017, 4, 4-13.	0.4	11
58	Intranasal desmopressin (OctimTM): a safe and efficacious treatment option for children with bleeding disorders. <i>Haemophilia</i> , 2007, 13, 548-551.	2.1	10
59	Supporting adherence and improving quality of life in haemophilia care. <i>British Journal of Nursing</i> , 2013, 22, 692-692.	0.7	9
60	The impact of clinical practice on the outcome of central venous access devices in children with haemophilia. <i>Haemophilia</i> , 2017, 23, e276-e281.	2.1	9
61	Prophylactic bypassing agent use before and during immune tolerance induction in patients with haemophilia A and inhibitors toFVIII. <i>Haemophilia</i> , 2018, 24, 570-577.	2.1	9
62	Assessing the supportive care needs of parents with a child with a bleeding disorder using the Parental Needs Scale for Rare Diseases (PNSâ€RD): A singleâ€centre pilot study. <i>Haemophilia</i> , 2019, 25, 831-837.	2.1	9
63	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> , 2022, 44, 8420-8428.	1.8	9
64	Validation of the Haemophilia & Exercise Projectâ€Testâ€Questionnaire (HEPâ€Testâ€Q)â€”An instrument for the assessment of subjective physical functioning in children with haemophilia. <i>Haemophilia</i> , 2018, 24, 888-895.	2.1	8
65	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.	2.5	8
66	The UK haemophilia specialist nurse: Competencies fit for practice in the 21st century. <i>Haemophilia</i> , 2020, 26, 622-630.	2.1	8
67	A core competency framework for haemophilia nurses in the UK. <i>The Journal of Haemophilia Practice</i> , 2013, 1, 32-36.	0.4	8
68	Pain Experience in Hemophilia Patients: A Hermeneutic Phenomenological Study. <i>International Journal of Community Based Nursing and Midwifery</i> , 2016, 4, 309-319.	0.2	8
69	The lived experience of women with a bleeding disorder: Aâ€systematic review. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2022, 6, e12652.	2.3	8
70	Immune tolerance in children with factors VIII and IX inhibitors: a single centre experience. <i>Haemophilia</i> , 2005, 11, 340-345.	2.1	7
71	â€Just an unfortunate coincidenceâ€™: childrenâ€™s understanding of haemophilia genetics and inheritance. <i>Haemophilia</i> , 2011, 17, 470-475.	2.1	7
72	Potential for development of haemophilia link nurse role within UK hospitals. <i>Haemophilia</i> , 2013, 19, 578-582.	2.1	7

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73	Pain: the voiceless scream in every haemophilia patient's life. The Journal of Haemophilia Practice, 2016, 3, 8-13.	0.4	7
74	Caregiver burden in haemophilia: results from a single UK centre. The Journal of Haemophilia Practice, 2017, 4, 40-48.	0.4	7
75	How does a lifetime of painful experiences influence sensations and beliefs about pain in adults with severe haemophilia? A qualitative study. Disability and Rehabilitation, 2022, 44, 8412-8419.	1.8	7
76	Evaluating a self infusion device for children with haemophilia. Paediatric Nursing, 2006, 18, 19-20.	0.1	6
77	Children's preferences of transfer devices for reconstitution of factors VIII and IX for the treatment of haemophilia. Haemophilia, 2009, 15, 159-167.	2.1	6
78	"I can always rely on them": the importance of social support for boys with haemophilia. The Journal of Haemophilia Practice, 2013, 1, 10-16.	0.4	6
79	Successful use of BPL Factor X concentrate in a child with severe factor X deficiency. The Journal of Haemophilia Practice, 2013, 1, 8-10.	0.4	6
80	Physical function and quality of life in adolescents with haemophilia (SO-FIT study). The Journal of Haemophilia Practice, 2013, 1, 11-14.	0.4	6
81	HaemophiliaLIVE: an ethnographic study on the impact of haemophilia on daily life. The Journal of Haemophilia Practice, 2014, 1, 14-20.	0.4	6
82	Multimethodology research with boys with severe haemophilia. Nurse Researcher, 2013, 20, 40-44.	0.5	6
83	Research Engagement in Pediatric Nursing Practice: Career Pathways are the Biggest Barrier. Journal of Pediatrics, 2020, 221, S62-S63.	1.8	5
84	Rituximab Is Effective and Well Tolerated in Children with Hemophilia and Factor VIII Inhibitors.. Blood, 2006, 108, 1031-1031.	1.4	5
85	Fitness enhances psychosocial well-being and self-confidence in young men with hemophilia: Results from Project GYM. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12622.	2.3	5
86	The views of women with bleeding disorders: Results from the Cinderella study. Haemophilia, 2022, 28, 316-325.	2.1	5
87	Haemophilia A: meeting the needs of individual patients. British Journal of Nursing, 2007, 16, 987-993.	0.7	4
88	Why don't haemophilia nurses do research?. Haemophilia, 2012, 18, 540-543.	2.1	4
89	Pharmacokinetics, Safety, and Efficacy of Recombinant Factor VIII Fc Fusion Protein. Journal of Infusion Nursing, 2017, 40, 65-75.	2.3	4
90	The impact on parents of having a child with haemophilia. The Journal of Haemophilia Practice, 2016, 3, 4-14.	0.4	4

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91	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.	0.4	4
92	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> , 2022, 25, 443-454.	2.6	4
93	Shortening the paediatric Haemophilia Activities List (pedHAL) based on pooled data from international studies. <i>Haemophilia</i> , 2021, 27, 305-313.	2.1	3
94	Real-World Data on the Effectiveness and Safety of wilate for the Treatment of von Willebrand Disease. <i>TH Open</i> , 2021, 05, e264-e272.	1.4	3
95	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.	0.4	3
96	Switching factor products: nurses' experience with NovoEight. <i>The Journal of Haemophilia Practice</i> , 2020, 7, 59-69.	0.4	3
97	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.	2.1	2
98	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 rd , 2021; virtual congress). <i>Expert Review of Hematology</i> , 2021, 14, 1-18.	2.2	2
99	Patient and caregiver perceptions of a pre-filled diluent syringe (MixPro [®]). <i>The Journal of Haemophilia Practice</i> , 2016, 3, 33-38.	0.4	2
100	Haemophilia nursing practice: A global survey of roles and responsibilities. <i>The Journal of Haemophilia Practice</i> , 2016, 3, 29-33.	0.4	2
101	A multi-country snapshot study of pain in people with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2020, 7, 143-149.	0.4	2
102	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.	2.3	2
103	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.	1.4	2
104	Experience of recombinant-activated factor VII to control bleeding in children with Glanzmann's thrombasthenia. <i>Comparative Clinical Pathology</i> , 2007, 16, 167-172.	0.7	1
105	Pain self-management experiences in haemophilia patients: a qualitative study. <i>The Journal of Haemophilia Practice</i> , 2018, 5, 76-82.	0.4	1
106	Experience of switching to NovoEight: views of people with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2020, 7, 70-77.	0.4	1
107	"You're only a carrier" women and the language of haemophilia. <i>The Journal of Haemophilia Practice</i> , 2021, 8, 128-132.	0.4	1
108	A Neonate with Type 3 Von Willebrands Disease and Life Threatening Bleeding Treated Successfully with a FVIII/WWF Concentrate (wilate [®]). <i>Blood</i> , 2012, 120, 4633-4633.	1.4	0

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109	Ahead Study: A 3 Year Follow-up of 522 Severe and Moderate Hemophilia a Patients. Blood, 2016, 128, 3786-3786.	1.4	0
110	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.	1.4	0
111	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.	0.4	0
112	You think you know the patient inside-out. But do you know the <i>outside-in</i>?. The Journal of Haemophilia Practice, 2018, 5, 130-137.	0.4	0
113	Now and then, and half a world away. The Journal of Haemophilia Practice, 2019, 6, s36-s38.	0.4	0
114	Final Results of the Prospective ADVATEA® Immune Tolerance Induction Registry (PAIR) Study with Plasma- and Albumin-Free Recombinant Factor VIII. Journal of Blood Medicine, 2021, Volume 12, 991-1001.	1.7	0
115	Haemophilia. Professional Nurse (london, England), 2002, 17, 463-4.	0.1	0
116	â€œIt's a way of lifeâ€: Results from the Perceptions of Pain in Haemophilia study. The Journal of Haemophilia Practice, 2021, 8, 145-154.	0.4	0