# Kate Khair

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8139322/publications.pdf

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| 116 | 2,521 | 201674 | 223800 |
|---|---|---|---|
| papers | citations | 27 h-index | 46 g-index |

| 118 | 118 | 118 | 1775 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | The impact of prophylactic treatment on children with severe haemophilia. British Journal of Haematology, 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. Journal of Thrombosis and Haemostasis, 2004, 2, 1096-1103. | 3.8 | 179 |
| 3 | Purpura fulminans: recognition, diagnosis and management. Archives of Disease in Childhood, 2011, 96, 1066-1071. | 1.9 | 166 |
| 4 | The use of recombinant factor VIIa in children with inherited platelet function disorders. British Journal of Haematology, 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. Haemophilia, 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. British Journal of Haematology, 1998, 100, 351-357. | 2.5 | 91 |
| 7 | Bruising and bleeding in infants and children - a practical approach. British Journal of Haematology, 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. British Journal of Haematology, 2004, 125, 366-368. | 2.5 | 73 |
| 9 | C1-inhibitor concentrate home therapy for hereditary angioedema: a viable, effective treatment option. Clinical and Experimental Immunology, 2006, 147, 061124011224001-???. | 2.6 | 73 |
| 10 | Factors affecting the Haemophilia Joint Health Score in children with severe haemophilia. Haemophilia, 2013, 19, 626-631. | 2.1 | 57 |
| 11 | Central venous access devices in children with congenital coagulation disorders: complications and long-term outcome. British Journal of Haematology, 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. British Journal of Haematology, 2006, 135, 101-104. | 2.5 | 47 |
| 13 | Increased burden on caregivers of having a child with haemophilia complicated by inhibitors. Pediatric Blood and Cancer, 2014, 61, 706-711. | 1.5 | 41 |
| 14 | Assessment of Treatment Practice Patterns for Severe Hemophilia A: A Global Nurse Perspective. Acta Haematologica, 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. British Journal of Haematology, 2002, 116, 632-635. | 2.5 | 39 |
| 16 | Can early subclinical gait changes in children with haemophilia be identified using the GAITRiteï¿½walkway. Haemophilia, 2007, 13, 542-547. | 2.1 | 39 |
| 17 | The impact of sport on children with haemophilia. Haemophilia, 2012, 18, 898-905. | 2.1 | 38 |
| 18 | Obesity in the global haemophilia population: prevalence, implications and expert opinions for weight management. Obesity Reviews, 2018, 19, 1569-1584. | 6.5 | 34 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Inhibitors in nonsevere haemophilia A: outcome and eradication strategies. Thrombosis and Haemostasis, 2015, 114, 46-55. | 3.4 | 33 |
| 20 | Assessment and management of pain in children and adolescents with bleeding disorders: a crossâ€sectional study from three haemophilia centres. Haemophilia, 2016, 22, 65-71. | 2.1 | 33 |
| 21 | Pattern of bleeding in a large prospective cohort of haemophilia A patients: A threeâ€year followâ€up of the <scp>AHEAD</scp> (Advate in HaEmophilia A outcome Database) study. Haemophilia, 2018, 24, 85-96. | 2.1 | 33 |
| 22 | The benefits of prophylaxis: views of adolescents with severe haemophilia. Haemophilia, 2012, 18, e286-9. | 2.1 | 32 |
| 23 | Selfâ€management and skills acquisition in boys with haemophilia. Health Expectations, 2015, 18, 1105-1113. | 2.6 | 32 |
| 24 | The impact of sport on health status, psychological wellâ€being and physical performance of adults with haemophilia. Haemophilia, 2016, 22, 521-530. | 2.1 | 32 |
| 25 | Health-Related Quality of Life and Psychological Aspects of Adults With Hemophilia in Iran. Clinical and Applied Thrombosis/Hemostasis, 2018, 24, 1073-1081. | 1.7 | 30 |
| 26 | Stem cell transplantation for children with Glanzmann thrombasthenia. British Journal of Haematology, 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: coâ€segregation with increased susceptibility to ADAMTS13 proteolysis but not clinical phenotype. British Journal of Haematology, 2005, 128, 830-836. | 2.5 | 28 |
| 28 | The experience of girls and young Women with inherited bleeding disorders. Haemophilia, 2013, 19, e276-81. | 2.1 | 28 |
| 29 | Hermansky-Pudlak syndrome: infrequent bleeding and first report of Turkish and Pakistani kindreds. Archives of Disease in Childhood, 2002, 86, 297-301. | 1.9 | 27 |
| 30 | Physiotherapy interventions for pain management in haemophilia: A systematic review. Haemophilia, 2020, 26, 667-684. | 2.1 | 27 |
| 31 | Compliance, concordance and adherence: what are we talking about?. Haemophilia, 2014, 20, 601-603. | 2.1 | 23 |
| 32 | European principles of care for women and girls with inherited bleeding disorders. Haemophilia, 2021, 27, 837-847. | 2.1 | 23 |
| 33 | Longâ€term consequences of intracranial haemorrhage in children with haemophilia. Haemophilia, 2009, 15, 184-192. | 2.1 | 21 |
| 34 | Treatment burden, haemostatic strategies and real world inhibitor screening practice in nonâ€severe haemophilia A. British Journal of Haematology, 2017, 176, 796-804. | 2.5 | 21 |
| 35 | Prophylaxis with prothrombin complex concentrate in four children with severe congenital factor X deficiency. Haemophilia, 2009, 15, 401-403. | 2.1 | 18 |
| 36 | Study of physical function in adolescents with haemophilia: The SOâ€FIT study. Haemophilia, 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 Tf | 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 / | 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 XIIIA 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. Haemophilia, 2013, 19, 256-258. | 2.1 | 11 |
| 56 | Education needs of nurses in thrombosis and hemostasis: An international, mixedâ€methods study. Research and Practice in Thrombosis and Haemostasis, 2019, 3, 99-108. | 2.3 | 11 |
| 57 | What is a nurse-led service? A discussion paper. The Journal of Haemophilia Practice, 2017, 4, 4-13. | 0.4 | 11 |
| 58 | Intranasal desmopressin (Octim<sup>TM</sup>): a safe and efficacious treatment option for children with bleeding disorders. Haemophilia, 2007, 13, 548-551. | 2.1 | 10 |
| 59 | Supporting adherence and improving quality of life in haemophilia care. British Journal of Nursing, 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. Haemophilia, 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. Haemophilia, 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. Haemophilia, 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. Disability and Rehabilitation, 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. Haemophilia, 2018, 24, 888-895. | 2.1 | 8 |
| 65 | The top 10 research priorities in bleeding disorders: a James Lind Alliance Priority Setting Partnership. British Journal of Haematology, 2019, 186, e98-e100. | 2.5 | 8 |
| 66 | The UK haemophilia specialist nurse: Competencies fit for practice in the 21st century. Haemophilia, 2020, 26, 622-630. | 2.1 | 8 |
| 67 | A core competency framework for haemophilia nurses in the UK. The Journal of Haemophilia Practice, 2013, 1, 32-36. | 0.4 | 8 |
| 68 | Pain Experience in Hemophilia Patients: A Hermeneutic Phenomenological Study. International Journal of Community Based Nursing and Midwifery, 2016, 4, 309-319. | 0.2 | 8 |
| 69 | The lived experience of women with a bleeding disorder: AÂsystematic review. Research and Practice in Thrombosis and Haemostasis, 2022, 6, e12652. | 2.3 | 8 |
| 70 | Immune tolerance in children with factors VIII and IX inhibitors: a single centre experience. Haemophilia, 2005, 11, 340-345. | 2.1 | 7 |
| 71 | â€˜Just an unfortunate coincidenceâ€™: childrenâ€™s understanding of haemophilia genetics and inheritance. Haemophilia, 2011, 17, 470-475. | 2.1 | 7 |
| 72 | Potential for development of haemophilia link nurse role within UK hospitals. Haemophilia, 2013, 19, 578-582. | 2.1 | 7 |

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|---|---|---|---|
| 73 | Pain: the voiceless scream in every haemophilia patientâ€™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 | Childrenâ€™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 | â€˜I can always rely on themâ€™: 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 Hemophila and Factor VIII Inhibitors.. Blood, 2006, 108, 1031-1031. | 1.4 | 5 |
| 85 | Fitness enhances psychosocial wellâ€being and selfâ€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. The Journal of Haemophilia Practice, 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. Health Expectations, 2022, 25, 443-454. | 2.6 | 4 |
| 93 | Shortening the paediatric Haemophilia Activities List (pedHAL) based on pooled data from international studies. Haemophilia, 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. TH Open, 2021, 05, e264-e272. | 1.4 | 3 |
| 95 | Evaluation of a pre-filled diluent syringe (MixPro<sup>Â®</sup>) among patient/carer users and nurses. The Journal of Haemophilia Practice, 2018, 5, 12-23. | 0.4 | 3 |
| 96 | Switching factor products: nursesâ€™ experience with NovoEight. The Journal of Haemophilia Practice, 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. Haemophilia, 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 3â€“5, 2021; virtual congress). Expert Review of Hematology, 2021, 14, 1-18. | 2.2 | 2 |
| 99 | Patient and caregiver perceptions of a pre-filled diluent syringe (MixProÂ®). The Journal of Haemophilia Practice, 2016, 3, 33-38. | 0.4 | 2 |
| 100 | Haemophilia nursing practice: A global survey of roles and responsibilities. The Journal of Haemophilia Practice, 2016, 3, 29-33. | 0.4 | 2 |
| 101 | A multi-country snapshot study of pain in people with haemophilia. The Journal of Haemophilia Practice, 2020, 7, 143-149. | 0.4 | 2 |
| 102 | Project GYM: A randomized feasibility study investigating effect on motivation of personal trainerâ€led exercise in young men with hemophilia. Research and Practice in Thrombosis and Haemostasis, 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. Blood, 2020, 136, 1-1. | 1.4 | 2 |
| 104 | Experience of recombinant-activated factor VII to control bleeding in children with Glanzmannâ€™s thrombasthenia. Comparative Clinical Pathology, 2007, 16, 167-172. | 0.7 | 1 |
| 105 | Pain self-management experiences in haemophilia patients: a qualitative study. The Journal of Haemophilia Practice, 2018, 5, 76-82. | 0.4 | 1 |
| 106 | Experience of switching to NovoEight: views of people with haemophilia. The Journal of Haemophilia Practice, 2020, 7, 70-77. | 0.4 | 1 |
| 107 | â€œYouâ€™re only a carrierâ€•â€“ women and the language of haemophilia. The Journal of Haemophilia Practice, 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/VWF Concentrate (wilateÂ®). Blood, 2012, 120, 4633-4633. | 1.4 | 0 |

| # | Article | IF | Citations |
|---|---|---|---|
| 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 outside-in?. 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 ADVATEÂ® 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 |