

# catherine Lambert

## List of Publications by Year in descending order

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Version: 2024-02-01

78  
papers

1,748  
citations

430874

18  
h-index

302126

39  
g-index

81  
all docs

81  
docs citations

81  
times ranked

2351  
citing authors

#	ARTICLE	IF	CITATIONS
1	Primary immune thrombocytopenia in adults: Belgian recommendations for diagnosis and treatment anno 2021 made by the Belgian Hematology Society. <i>Acta Clinica Belgica</i> , 2022, 77, 470-483.	1.2	4
2	Apixaban and Dalteparin for the Treatment of Venous Thromboembolism in Patients with Different Sites of Cancer. <i>Thrombosis and Haemostasis</i> , 2022, 122, 796-807.	3.4	21
3	Impact of therapeutic plasma exchange on acquired vaccinal anti-SARS-CoV-2 antibodies. <i>European Journal of Internal Medicine</i> , 2022, , .	2.2	0
4	Gaining more insight into ankle pain in haemophilia: A study exploring pain, structural and functional evaluation of the ankle joint. <i>Haemophilia</i> , 2022, 28, 480-490.	2.1	9
5	Pain coping behaviour strategies in people with haemophilia: A systematic literature review. <i>Haemophilia</i> , 2022, 28, 902-916.	2.1	3
6	Management of COVID-19 Coagulopathy in a Patient with Severe Haemophilia A. <i>Acta Haematologica</i> , 2021, 144, 319-321.	1.4	5
7	Patient perspectives regarding gene therapy in haemophilia: Interviews from the PAVING study. <i>Haemophilia</i> , 2021, 27, 129-136.	2.1	27
8	Feasibility and outcomes of low-dose and low-frequency prophylaxis with recombinant extended half-life products (FcFVIII and FcFIX) in Ivorian children with hemophilia: Two-year experience in the setting of World Federation of Haemophilia humanitarian aid programme. <i>Haemophilia</i> , 2021, 27, 33-40.	2.1	11
9	Patient selection for hemophilia gene therapy: Real-life data from a single center. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, 390-394.	2.3	11
10	Peripheral embolism as first and only clinical symptom of a true aneurysmal degeneration of the radial artery after ligation of a radiocephalic fistula. <i>Journal of Vascular Access</i> , 2021, , 112972982110333.	0.9	1
11	Diagnosis of hepatitis C-related liver disease in patients with mild hemophilia. <i>European Journal of Internal Medicine</i> , 2021, 91, 102-103.	2.2	0
12	Patient preferences for gene therapy in haemophilia: Results from the PAVING threshold technique survey. <i>Haemophilia</i> , 2021, 27, 957-966.	2.1	14
13	Inhibitor epidemiology and genetic-related risk factors in people with haemophilia from Cote d'Ivoire. <i>Haemophilia</i> , 2020, 26, 79-85.	2.1	3
14	Can we compare haemophilia carriers with clotting factor deficiency to male patients with mild haemophilia?. <i>Haemophilia</i> , 2020, 26, 117-121.	2.1	20
15	EHLFIX in haemophilia B carriers with FIX deficiency. <i>Haemophilia</i> , 2020, 26, e38-e40.	2.1	4
16	First report of emicizumab use in a female patient with severe hemophilia A. <i>Blood Advances</i> , 2020, 4, 2950-2952.	5.2	3
17	In-hospital management of persons with haemophilia and COVID-19: Practical guidance. <i>Haemophilia</i> , 2020, 26, 768-772.	2.1	19
18	Cross-cultural adaptation and validation of HaemAQoL in Cote d'Ivoire. <i>Haemophilia</i> , 2020, 26, 459-466.	2.1	4

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19	Experience of a 40-day (6-week) LMWH treatment for isolated distal deep vein thrombosis. <i>Journal of Thrombosis and Thrombolysis</i> , 2020, 50, 837-843.	2.1	1
20	Cross-cultural adaptation and validation of the Canadian Haemophilia Outcomes-Kids™ Life Assessment Tool (CHO-KLAT) in Côte d'Ivoire (the Ivory Coast). <i>Health and Quality of Life Outcomes</i> , 2020, 18, 76.	2.4	4
21	Apixaban for the Treatment of Venous Thromboembolism Associated with Cancer. <i>New England Journal of Medicine</i> , 2020, 382, 1599-1607.	27.0	658
22	Impact of the COVID-19 pandemic on therapeutic choices in thrombosis and hemostasis. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 1794-1795.	3.8	26
23	Low rate of subclinical venous thrombosis in patients with haemophilia undergoing major orthopaedic surgery in the absence of pharmacological thromboprophylaxis. <i>Haemophilia</i> , 2020, 26, 1064-1071.	2.1	9
24	Implementation and assessment of a self- and community-based rehabilitation programme in patients with haemophilia from Côte d'Ivoire. <i>Haemophilia</i> , 2019, 25, 859-866.	2.1	12
25	Development and evaluation of appropriate, culturally adapted educational tools for Ivoirian patients with haemophilia, haemophilia carriers and their families. <i>Haemophilia</i> , 2019, 25, 838-844.	2.1	4
26	Incidental finding of unreported large duplication in F8 gene during prenatal analysis: Which management for genetic counselling?. <i>Thrombosis Research</i> , 2019, 182, 39-42.	1.7	4
27	VWF concentrates in the treatment of inherited von Willebrand disease: A single-centre retrospective study. <i>Haemophilia</i> , 2019, 25, e300-e303.	2.1	2
28	Haemophilia in Côte d'Ivoire (the Ivory Coast) in 2017: Extensive data collection as part of the World Federation of Hemophilia's twinning programme. <i>Haemophilia</i> , 2019, 25, 236-243.	2.1	13
29	Haemophilia carrier's awareness, diagnosis, and management in emerging countries: a cross-sectional study in Côte d'Ivoire (Ivory Coast). <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 26.	2.7	11
30	Compartment syndrome of the forearm with life-threatening bleeding after fasciotomy as the presenting sign of postpartum acquired hemophilia A. <i>Blood Coagulation and Fibrinolysis</i> , 2019, 30, 120-126.	1.0	4
31	Does haemophilia slow down the development of liver fibrosis?. <i>Haemophilia</i> , 2019, 25, e32-e35.	2.1	3
32	Screening of haemophilia carriers in moderate and severe haemophilia A and B: Prevalence and determinants. <i>Haemophilia</i> , 2018, 24, e142-e144.	2.1	5
33	Management of immune thrombocytopenia in multiple sclerosis patients treated with alemtuzumab: a Belgian consensus. <i>Acta Neurologica Belgica</i> , 2018, 118, 7-11.	1.1	10
34	Apixaban versus Dalteparin for the Treatment of Acute Venous Thromboembolism in Patients with Cancer: The Caravaggio Study. <i>Thrombosis and Haemostasis</i> , 2018, 118, 1668-1678.	3.4	121
35	Dabigatran etexilate in the treatment of localized intravascular coagulopathy associated with venous malformations. <i>Thrombosis Research</i> , 2018, 168, 114-120.	1.7	14
36	The value and impact of anti-Xa activity monitoring for prophylactic dose adjustment of low-molecular-weight heparin during pregnancy. <i>Blood Coagulation and Fibrinolysis</i> , 2017, 28, 199-204.	1.0	20

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37	Usual and unusual mutations in a cohort of Belgian patients with hemophilia B. <i>Thrombosis Research</i> , 2017, 149, 25-28.	1.7	5
38	Successful management of a severe anti-M alloimmunization during pregnancy. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2017, 217, 175-176.	1.1	6
39	Comparative study of the prevalence of clotting factor deficiency in carriers of haemophilia A and haemophilia B. <i>Haemophilia</i> , 2017, 23, e471-e473.	2.1	8
40	Biodistribution of Liver-Derived Mesenchymal Stem Cells After Peripheral Injection in a Hemophilia A Patient. <i>Transplantation</i> , 2017, 101, 1845-1851.	1.0	29
41	Successful Management of Acquired Hemophilia A Associated with Bullous Pemphigoid: A Case Report and Review of the Literature. <i>Case Reports in Hematology</i> , 2017, 2017, 1-7.	0.4	16
42	Successful Treatment and Secondary Prevention of Venous Thrombosis Secondary to Behçet Disease with Rivaroxaban. <i>Case Reports in Hematology</i> , 2016, 2016, 1-3.	0.4	5
43	Dabigatran etexilate versus low-molecular weight heparin to control consumptive coagulopathy secondary to diffuse venous vascular malformations. <i>Blood Coagulation and Fibrinolysis</i> , 2016, 27, 216-219.	1.0	18
44	Is the cardiovascular toxicity of NSAIDs and COX-2 selective inhibitors underestimated in patients with haemophilia?. <i>Critical Reviews in Oncology/Hematology</i> , 2016, 100, 25-31.	4.4	6
45	Stop only advising physical activity in adults with haemophilia – prescribe it now! The role of exercise therapy and nutrition in chronic musculoskeletal diseases. <i>Haemophilia</i> , 2016, 22, e554-e556.	2.1	16
46	Complement activation and effect of eculizumab in scleroderma renal crisis. <i>Medicine (United States)</i> , 2016, 95, e4459.	1.0	57
47	Continuous infusion of factor VIII concentrates in obese patients with severe haemophilia A: is weight-based dose adjustment required?. <i>Haemophilia</i> , 2016, 22, e62-4.	2.1	5
48	Plerixafor prescription modalities in autologous haematopoietic stem cell mobilization in Belgium. <i>Acta Clinica Belgica</i> , 2015, 70, 16-22.	1.2	1
49	Management of pregnancy in paroxysmal nocturnal hemoglobinuria on long-term eculizumab. <i>Blood Coagulation and Fibrinolysis</i> , 2015, 26, 464-466.	1.0	11
50	The use of short-term central venous catheters for optimizing continuous infusion of coagulation factor concentrate in haemophilia patients undergoing major surgical procedures. <i>Haemophilia</i> , 2015, 21, e364-8.	2.1	3
51	A large polyp in the rectum: not always an epithelial lesion. <i>International Journal of Colorectal Disease</i> , 2015, 30, 147-148.	2.2	1
52	Overrepresentation of missense mutations in mild hemophilia A patients from Belgium: founder effect or independent occurrence?. <i>Thrombosis Research</i> , 2015, 135, 1057-1063.	1.7	6
53	Rivaroxaban for arterial thrombosis related to heparin-induced thrombocytopenia. <i>Blood Coagulation and Fibrinolysis</i> , 2015, 26, 205-206.	1.0	26
54	Does the site of platelet sequestration predict the response to splenectomy in adult patients with immune thrombocytopenic purpura?. <i>Platelets</i> , 2015, 26, 573-576.	2.3	23

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55	Successful emergency resection of a massive intra-abdominal hemophilic pseudotumor. <i>World Journal of Gastrointestinal Surgery</i> , 2015, 7, 43.	1.5	9
56	Optimal management of hemophilic arthropathy and hematomas. <i>Journal of Blood Medicine</i> , 2014, 5, 207.	1.7	88
57	Budd-Chiari syndrome in a patient with acute promyelocytic leukaemia. <i>British Journal of Haematology</i> , 2014, 166, 1-1.	2.5	2
58	Physical and mental quality of life in adult patients with haemophilia in Belgium: the impact of financial issues. <i>Haemophilia</i> , 2014, 20, 479-485.	2.1	19
59	Multicentric Study Evaluating Venous Thrombosis Among Patients with Haemophilia Undergoing Major Orthopaedic Surgery. <i>Blood</i> , 2014, 124, 1496-1496.	1.4	1
60	Intron 22 homologous regions are implicated in exons 1-22 duplications of the F8 gene. <i>European Journal of Human Genetics</i> , 2013, 21, 970-976.	2.8	20
61	Polymyalgia rheumatica. <i>Blood Coagulation and Fibrinolysis</i> , 2013, 24, 211-212.	1.0	0
62	Successful management of hereditary angioedema during tonsillectomy. <i>Blood Coagulation and Fibrinolysis</i> , 2012, 23, 155-157.	1.0	5
63	Dabigatran etexilate (Pradaxa®) for preventing warfarin-induced skin necrosis in a patient with severe protein C deficiency. <i>Thrombosis and Haemostasis</i> , 2012, 107, 1189-1191.	3.4	27
64	Computational and molecular approaches for predicting unreported causal missense mutations in Belgian patients with haemophilia A. <i>Haemophilia</i> , 2012, 18, e331-9.	2.1	11
65	Acquired Antithrombin Type IIb Deficiency After Liver Transplantation: A Case Report. <i>American Journal of Transplantation</i> , 2012, 12, 1329-1332.	4.7	6
66	Novel head-to-head gene fusion of MLL with ZC3H13 in a JAK2 V617F-positive patient with essential thrombocythemia without blast cells. <i>Leukemia Research</i> , 2012, 36, e27-e30.	0.8	3
67	Percutaneous transcatheter closure of interatrial septal defect in adults: Procedural outcome and long-term results. <i>Catheterization and Cardiovascular Interventions</i> , 2012, 79, 322-330.	1.7	29
68	Deep vein thrombosis induced by thalidomide to control epistaxis secondary to hereditary haemorrhagic telangiectasia. <i>Blood Coagulation and Fibrinolysis</i> , 2011, 22, 616-618.	1.0	20
69	Plasma extraction rate and collection efficiency during therapeutic plasma exchange with Spectra Optia in comparison with Haemonetics MCS+. <i>Journal of Clinical Apheresis</i> , 2011, 26, 17-22.	1.3	23
70	Subclinical deep venous thrombosis observed in 10% of hemophilic patients undergoing major orthopedic surgery. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 1138-1140.	3.8	43
71	The role of physiotherapy after total knee arthroplasty in patients with haemophilia. <i>Haemophilia</i> , 2008, 14, 989-998.	2.1	40
72	ACUTE ADRENAL INSUFFICIENCY ASSOCIATED WITH HEPARIN-INDUCED THROMBOCYTOPENIA. <i>Acta Clinica Belgica</i> , 2008, 63, 112-115.	1.2	6

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73	Continuous intravenous infusion of a low-molecular-weight heparin during allogeneic haematopoietic stem-cell transplantation. <i>Blood Coagulation and Fibrinolysis</i> , 2008, 19, 735-737.	1.0	3
74	Safety of bevacizumab in mild haemophilia B. <i>Thrombosis and Haemostasis</i> , 2008, 99, 963-964.	3.4	10
75	The JAK2 V617F mutation is not a cause of central retinal vein occlusion. <i>Thrombosis and Haemostasis</i> , 2008, 100, 515-516.	3.4	3
76	Adjuvant treatment in resected pancreatic adenocarcinoma: A retrospective analysis of survival and prognostic factors in 141 patients. <i>Journal of Clinical Oncology</i> , 2007, 25, 15104-15104.	1.6	0
77	Inadvertent anticoagulation of a haemophiliac child with routine line flushing. <i>Haemophilia</i> , 2006, 12, 548-550.	2.1	3
78	High incidence of complications after 2-chloro-2'-deoxyadenosine combined with cyclophosphamide in patients with advanced lymphoproliferative malignancies. <i>Annals of Hematology</i> , 2004, 83, 356-363.	1.8	15