Will Lester

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1816373/publications.pdf

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30 papers	2,324 citations	471509 17 h-index	526287 27 g-index
30 all docs	30 docs citations	30 times ranked	2280 citing authors

#	Article	IF	Citations
1	Liver graft outcomes from donors with vaccine induced thrombosis and thrombocytopenia (VITT): United Kingdom multicenter experience. American Journal of Transplantation, 2022, 22, 996-998.	4.7	6
2	Pitfalls in laboratory monitoring of treatment in people with Haemophilia. Blood Reviews, 2022, 55, 100946.	5.7	0
3	Natural history of PF4 antibodies in vaccine-induced immune thrombocytopenia and thrombosis. Blood, 2022, 139, 2553-2560.	1.4	20
4	Laboratory coagulation tests and recombinant porcine factor VIII: A United Kingdom Haemophilia Centre Doctorsâ€™ Organisation guideline. Haemophilia, 2022, 28, 515-519.	2.1	7
5	Interindividual variability in transgene mRNA and protein production following adeno-associated virus gene therapy for hemophilia A. Nature Medicine, 2022, 28, 789-797.	30.7	48
6	Intracranial hemorrhage in immune thrombotic thrombocytopenic purpura treated with caplacizumab. Journal of Thrombosis and Haemostasis, 2021, 19, 1922-1925.	3.8	10
7	Coronary artery thrombus resulting in ST-elevation myocardial infarction in a patient with COVID-19. BMJ Case Reports, 2021, 14, e243811.	0.5	0
8	Endoscopy in patients on antiplatelet or anticoagulant therapy: British Society of Gastroenterology (BSG) and European Society of Gastrointestinal Endoscopy (ESGE) guideline update. Endoscopy, 2021, 53, 947-969.	1.8	47
9	Endoscopy in patients on antiplatelet or anticoagulant therapy: British Society of Gastroenterology (BSG) and European Society of Gastrointestinal Endoscopy (ESGE) guideline update. Gut, 2021, 70, 1611-1628.	12.1	61
10	Persistence of haemostatic response following gene therapy with valoctocogene roxaparvovec in severe haemophilia A. Haemophilia, 2021, 27, 947-956.	2.1	62
11	Laboratory measurement of factor replacement therapies in the treatment of congenital haemophilia: A United Kingdom Haemophilia Centre Doctorsâ€™ Organisation guideline. Haemophilia, 2020, 26, 6-16.	2.1	31
12	Multiyear Follow-up of AAV5-hFVIII-SQ Gene Therapy for Hemophilia A. New England Journal of Medicine, 2020, 382, 29-40.	27.0	316
13	Laboratory coagulation tests and emicizumab treatment A United Kingdom Haemophilia Centre Doctors' Organisation guideline. Haemophilia, 2020, 26, 151-155.	2.1	44
14	Guidelines on the use of liver biopsy in clinical practice from the British Society of Gastroenterology, the Royal College of Radiologists and the Royal College of Pathology. Gut, 2020, 69, 1382-1403.	12.1	172
15	Addendum to British Society for Haematology Guidelines on Investigation and Management of Antiphospholipid syndrome, 2012 (Br. J. Haematol. 2012; 157: 47â€“58): use of direct acting oral anticoagulants. British Journal of Haematology, 2020, 189, 212-215.	2.5	53
16	Are we assessing venous thromboembolism (VTE) risk appropriately for hospitalised medical patients? The National VTE Risk Assessment Tool versus Padua Prediction Score. British Journal of Haematology, 2020, 189, e16-e18.	2.5	4
17	Management of acute myocardial infarction in a patient with idiopathic thrombocytopenic purpura, the value of optical coherence tomography: a case report. European Heart Journal - Case Reports, 2020, 4, 1-5.	0.6	5
18	NICE NG89 recommendations for extended pharmacological thromboprophylaxis â€“ is it justified and is it cost effective: a rebuttal from the British Society for Haematology. British Journal of Haematology, 2019, 186, 790-791.	2.5	6

#	Article	IF	Citations
19	Investigation of the contribution of an underlying platelet defect in women with unexplained heavy menstrual bleeding. Platelets, 2019, 30, 56-65.	2.3	9
20	A United Kingdom Immune Thrombocytopenia (<scp>ITP</scp>) Forum review of practice: thrombopoietin receptor agonists. British Journal of Haematology, 2018, 180, 591-594.	2.5	7
21	Varicose Veins and Deep Venous Thrombosis. JAMA - Journal of the American Medical Association, 2018, 320, 509.	7.4	7
22	AAV5â€“Factor VIII Gene Transfer in Severe Hemophilia A. New England Journal of Medicine, 2017, 377, 2519-2530.	27.0	529
23	Outcomes of pregnancy in patients with known Budd-Chiari syndrome. World Journal of Hepatology, 2017, 9, 945.	2.0	21
24	The diagnosis and management of von <scp>W</scp>illebrand disease: a <scp>U</scp>nited <scp>K</scp>ingdom <scp>H</scp>aemophilia <scp>C</scp>entre <scp>D</scp>octors <scp>O</scp>rganization guideline approved by the <scp>B</scp>ritish <scp>C</scp>ommittee for <scp>S</scp>tandards in <scp>H</scp>aematology. British Journal of Haematology, 2014, 167, 453-465.	2.5	297
25	Fatal venous thromboembolism associated with hospital admission: a cohort study to assess the impact of a national risk assessment target. Heart, 2013, 99, 1734-1739.	2.9	46
26	Expression of 14 von Willebrand factor mutations identified in patients with typeÂ1 von Willebrand disease from the MCMDM-1VWD study. Journal of Thrombosis and Haemostasis, 2009, 7, 1304-1312.	3.8	57
27	Identification and characterization of a novel P2Y12 variant in a patient diagnosed with type 1 von Willebrand disease in the European MCMDM-1VWD study. Blood, 2009, 113, 4110-4113.	1.4	67
28	Phenotype and genotype of a cohort of families historically diagnosed with type 1 von Willebrand disease in the European study, Molecular and Clinical Markers for the Diagnosis and Management of Type 1 von Willebrand Disease (MCMDM-1VWD). Blood, 2007, 109, 112-121.	1.4	364
29	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
30	Thromboprophylaxis for the Obese Surgical Patient. , 0, , 181-185.		0