Michael Makris

List of Publications by Citations

370 papers	10,163 citations	50 h-index	91 g-index
433 ext. papers	12,022 ext. citations	6.1 avg, IF	6.32 L-index

#	Paper	IF	Citations
370	Increased fetal loss in women with heritable thrombophilia. *Lancet, The*, **1996**, 348, 913-6	40	550
369	Acquired hemophilia A in the United Kingdom: a 2-year national surveillance study by the United Kingdom Haemophilia Centre Doctors' Organisation. *Blood*, **2007**, 109, 1870-7	2.2	501
368	Guidelines on oral anticoagulation with warfarin - fourth edition. *British Journal of Haematology*, **2011**, 154, 311-24	4.5	399
367	Clinical guidelines for testing for heritable thrombophilia. *British Journal of Haematology*, **2010**, 149, 209-20	4.5	349
366	Recurrent miscarriage: aetiology, management and prognosis. *Human Reproduction Update*, **2002**, 8, 463-81	15.8	334
365	Emergency Oral Anticoagulant Reversal: The Relative Efficacy of Infusions of Fresh Frozen Plasma and Clotting Factor Concentrate on Correction of the Coagulopathy. *Thrombosis and Haemostasis*, **1997**, 77, 477-480	7	284
364	Demonstration of viraemia patterns in haemophiliacs treated with hepatitis-C-virus-contaminated factor VIII concentrates. *Lancet, The*, **1990**, 336, 1022-5	40	209
363	Hepatitis C antibody and chronic liver disease in haemophilia. *Lancet, The*, **1990**, 335, 1117-9	40	204
362	Thrombin generation testing in routine clinical practice: are we there yet?. *British Journal of Haematology*, **2008**, 142, 889-903	4.5	164
361	Guideline on the selection and use of therapeutic products to treat haemophilia and other hereditary bleeding disorders. A United Kingdom Haemophilia Center Doctors' Organisation (UKHCDO) guideline approved by the British Committee for Standards in Haematology. *Haemophilia*, **2008**, 14, 671-84	3.3	159
360	Clinical Features of Vaccine-Induced Immune Thrombocytopenia and Thrombosis. *New England Journal of Medicine*, **2021**, 385, 1680-1689	59.2	156
359	Guideline on the management of bleeding in patients on antithrombotic agents. *British Journal of Haematology*, **2013**, 160, 35-46	4.5	155
358	The natural history of chronic hepatitis C in haemophiliacs. *British Journal of Haematology*, **1996**, 94, 746-53	4.5	152
357	A comparison of the efficacy and rate of response to oral and intravenous Vitamin K in reversal of over-anticoagulation with warfarin. *British Journal of Haematology*, **2001**, 115, 145-9	4.5	149
356	Enzymatically catalyzed disulfide exchange is required for platelet adhesion to collagen via integrin alpha2beta1. *Blood*, **2003**, 102, 2085-92	2.2	147
355	Risk of a first venous thrombotic event in carriers of a familial thrombophilic defect. The European Prospective Cohort on Thrombophilia (EPCOT). *Journal of Thrombosis and Haemostasis*, **2005**, 3, 459-64	15.4	147
354	The risk of spinal haematoma following neuraxial anaesthesia or lumbar puncture in thrombocytopenic individuals. *British Journal of Haematology*, **2010**, 148, 15-25	4.5	146

353	Enhanced thrombin generation in patients with cirrhosis-induced coagulopathy. *Journal of Thrombosis and Haemostasis*, **2010**, 8, 1994-2000	15.4	145
352	Home treatment of haemarthroses using a single dose regimen of recombinant activated factor VII in patients with haemophilia and inhibitors. *Thrombosis and Haemostasis*, **2006**, 95, 600-605	7	140
351	A spontaneous prothrombotic disorder resembling heparin-induced thrombocytopenia. *American Journal of Medicine*, **2008**, 121, 632-6	2.4	124
350	Co-inheritance of the 20210A Allele of the Prothrombin Gene Increases the Risk of Thrombosis in Subjects with Familial Thrombophilia. *Thrombosis and Haemostasis*, **1997**, 78, 1426-1429	7	117
349	Establishing the Prevalence and Prevalence at Birth of Hemophilia in Males: A Meta-analytic Approach Using National Registries. *Annals of Internal Medicine*, **2019**, 171, 540-546	8	115
348	Stomatocytic haemolysis and macrothrombocytopenia (Mediterranean stomatocytosis/macrothrombocytopenia) is the haematological presentation of phytosterolaemia. *British Journal of Haematology*, **2005**, 130, 297-309	4.5	113
347	Ligelizumab for Chronic Spontaneous Urticaria. *New England Journal of Medicine*, **2019**, 381, 1321-1332	59.2	102
346	The management of coumarin-induced over-anticoagulation Annotation. *British Journal of Haematology*, **2001**, 114, 271-80	4.5	98
345	Inhibitor development in haemophilia according to concentrate. Four-year results from the European HAemophilia Safety Surveillance (EUHASS) project. *Thrombosis and Haemostasis*, **2015**, 113, 968-75	7	89
344	Whole exome sequencing identifies genetic variants in inherited thrombocytopenia with secondary qualitative function defects. *Haematologica*, **2016**, 101, 1170-1179	6.6	89
343	Corn trypsin inhibitor in fluorogenic thrombin-generation measurements is only necessary at low tissue factor concentrations and influences the relationship between factor VIII coagulant activity and thrombogram parameters. *Blood Coagulation and Fibrinolysis*, **2008**, 19, 183-9	1	75
342	Recurrence rate after a first venous thrombosis in patients with familial thrombophilia. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **2005**, 25, 1992-7	9.4	75
341	Progression to end-stage liver disease in patients with inherited bleeding disorders and hepatitis C: an international, multicenter cohort study. *Blood*, **2007**, 109, 3667-71	2.2	72
340	Thrombotic adverse events to coagulation factor concentrates for treatment of patients with haemophilia and von Willebrand disease: a systematic review of prospective studies. *Haemophilia*, **2012**, 18, e173-87	3.3	71
339	Hereditary thrombophilia and fetal loss: a prospective follow-up study. *Journal of Thrombosis and Haemostasis*, **2004**, 2, 592-6	15.4	71
338	Familial thrombophilia and lifetime risk of venous thrombosis. *Journal of Thrombosis and Haemostasis*, **2004**, 2, 1526-32	15.4	71
337	A randomized controlled trial of recombinant interferon-alpha in chronic hepatitis C in hemophiliacs. *Blood*, **1991**, 78, 1672-1677	2.2	70
336	Heterogeneity of hepatitis C virus genotypes in hemophilia: relationship with chronic liver disease. *Blood*, **1995**, 85, 1259-1262	2.2	69

335	Emergency oral anticoagulant reversal: the relative efficacy of infusions of fresh frozen plasma and clotting factor concentrate on correction of the coagulopathy. *Thrombosis and Haemostasis*, **1997**, 77, 477-80	7	69
334	Hepatitis C viral RNA in clotting factor concentrates and the development of hepatitis in recipients. *Blood*, **1993**, 81, 1898-1902	2.2	67
333	Thrombin generation assays are superior to traditional tests in assessing anticoagulation reversal in vitro. *Thrombosis and Haemostasis*, **2008**, 100, 350-355	7	66
332	Definition, aims, and implementation of GA(2) LEN Urticaria Centers of Reference and Excellence. *Allergy: European Journal of Allergy and Clinical Immunology*, **2016**, 71, 1210-8	9.3	65
331	Biomarkers and clinical characteristics of autoimmune chronic spontaneous urticaria: Results of the PURIST Study. *Allergy: European Journal of Allergy and Clinical Immunology*, **2019**, 74, 2427-2436	9.3	61
330	EUHASS: The European Haemophilia Safety Surveillance system. *Thrombosis Research*, **2011**, 127 Suppl 2, S22-5	8.2	61
329	Identification and characterization of a novel P2Y 12 variant in a patient diagnosed with type 1 von Willebrand disease in the European MCMDM-1VWD study. *Blood*, **2009**, 113, 4110-3	2.2	59
328	Mobile technology offers novel insights into the control and treatment of allergic rhinitis: The MASK study. *Journal of Allergy and Clinical Immunology*, **2019**, 144, 135-143.e6	11.5	57
327	Prophylaxis in haemophilia should be life-long. *Blood Transfusion*, **2012**, 10, 165-8	3.6	56
326	Treatment of bleeding episodes in haemophilia A complicated by a factor VIII inhibitor in patients receiving Emicizumab. Interim guidance from UKHCDO Inhibitor Working Party and Executive Committee. *Haemophilia*, **2018**, 24, 344-347	3.3	55
325	Evaluation of laboratory assays for anti-platelet factor 4 antibodies after ChAdOx1 nCOV-19 vaccination. *Journal of Thrombosis and Haemostasis*, **2021**, 19, 2007-2013	15.4	53
324	Unfavourable cardiovascular disease risk profiles in a cohort of Dutch and British haemophilia patients. *Thrombosis and Haemostasis*, **2013**, 109, 16-23	7	52
323	Warfarin anticoagulation reversal: management of the asymptomatic and bleeding patient. *Journal of Thrombosis and Thrombolysis*, **2010**, 29, 171-81	5.1	52
322	The international EAACI/GA□LEN/EuroGuiDerm/APAAACI guideline for the definition, classification, diagnosis, and management of urticaria. *Allergy: European Journal of Allergy and Clinical Immunology*, **2021**,	9.3	50
321	Inhibitor development in previously treated hemophilia A patients: a systematic review, meta-analysis, and meta-regression. *Journal of Thrombosis and Haemostasis*, **2013**, 11, 1655-62	15.4	47
320	Hyperhomocysteinemia and thrombosis. *International Journal of Laboratory Hematology*, **2000**, 22, 133-43		47
319	Gastrointestinal bleeding in von Willebrand disease. *Thrombosis Research*, **2006**, 118 Suppl 1, S13-7	8.2	46
318	The responsiveness of different APTT reagents to mild factor VIII, IX and XI deficiencies. *International Journal of Laboratory Hematology*, **2011**, 33, 154-8	2.5	44

317	The phenotypic and genetic assessment of antithrombin deficiency. *International Journal of Laboratory Hematology*, **2011**, 33, 227-37	2.5	44
316	Calibrated automated thrombin generation and modified thromboelastometry in haemophilia A. *Thrombosis Research*, **2009**, 123, 895-901	8.2	44
315	Long-term follow-up of hepatitis C infection in a large cohort of patients with inherited bleeding disorders. *Journal of Hepatology*, **2014**, 60, 39-45	13.4	42
314	The development of antiphospholipid antibodies in haemophilia is linked to infection with hepatitis C. *British Journal of Haematology*, **1994**, 88, 845-8	4.5	41
313	Thrombosis in Inherited Fibrinogen Disorders. *Transfusion Medicine and Hemotherapy*, **2017**, 44, 70-76	4.2	39
312	Vaccine-induced Immune Thrombocytopenia and Thrombosis (VITT). *Research and Practice in Thrombosis and Haemostasis*, **2021**, 5, e12529	5.1	39
311	Management of peri-operative anti-thrombotic therapy. *Anaesthesia*, **2015**, 70 Suppl 1, 58-67, e21-3	6.6	38
310	Phenotypic approaches to gene mapping in platelet function disorders. *Hamostaseologie*, **2010**, 30, 29-38	1.9	38
309	Clotting factor concentrate switching and inhibitor development in hemophilia A. *Blood*, **2012**, 120, 720-7	7.2	37
308	Wide variation in thrombin generation in patients with atrial fibrillation and therapeutic International Normalized Ratio is not due to inflammation. *British Journal of Haematology*, **2008**, 142, 946-52	4.5	36
307	Specific and global coagulation assays in the diagnosis of discrepant mild hemophilia A. *Haematologica*, **2013**, 98, 1980-7	6.6	35
306	Familial thrombophilia: genetic risk factors and management. *Journal of Internal Medicine*, **1997**, 242, 9-15	10.8	35
305	Antiviral therapy for chronic hepatitis C in patients with inherited bleeding disorders: an international, multicenter cohort study. *Journal of Thrombosis and Haemostasis*, **2007**, 5, 1624-9	15.4	35
304	Poor reversal of low molecular weight heparin by protamine. *British Journal of Haematology*, **2000**, 108, 884-5	4.5	35
303	Detection by PCR of hepatitis C virus in factor VIII concentrates. *Lancet, The*, **1990**, 335, 1473	40	35
302	Rupatadine inhibits proinflammatory mediator secretion from human mast cells triggered by different stimuli. *International Archives of Allergy and Immunology*, **2010**, 151, 38-45	3.7	34
301	Haemoperitoneum associated with ovulation in women with bleeding disorders: the case for conservative management and the role of the contraceptive pill. *Haemophilia*, **2007**, 13, 93-7	3.3	34
300	A rapid method for haemophilia B mutation detection using conformation sensitive gel electrophoresis. *British Journal of Haematology*, **1999**, 104, 915-8	4.5	34

299	Further evidence that activated protein C resistance can be misdiagnosed as inherited functional protein S deficiency. *British Journal of Haematology*, **1994**, 88, 201-3	4.5	34
298	Hyperhomocysteinemia and venous thrombosis. *Seminars in Hematology*, **2007**, 44, 70-6	4	33
297	Activation of coagulation in diabetes mellitus in relation to the presence of vascular complications. *Diabetic Medicine*, **1991**, 8, 322-9	3.5	33
296	Acquired Glanzmann's thrombasthenia without thrombocytopenia: a severe acquired autoimmune bleeding disorder. *British Journal of Haematology*, **2004**, 127, 209-13	4.5	32
295	Emicizumab and thrombosis: The story so far. *Journal of Thrombosis and Haemostasis*, **2019**, 17, 1269-1272	15.4	31
294	Immunization of patients with bleeding disorders. *Haemophilia*, **2003**, 9, 541-6	3.3	31
293	The natural history of occult or angiodysplastic gastrointestinal bleeding in von Willebrand disease. *Haemophilia*, **2015**, 21, 338-42	3.3	30
292	Hyperhomocysteinemia in women with advanced breast cancer. *International Journal of Laboratory Hematology*, **2007**, 29, 421-5	2.5	30
291	Performing and interpreting individual pharmacokinetic profiles in patients with Hemophilia A or B: Rationale and general considerations. *Research and Practice in Thrombosis and Haemostasis*, **2018**, 2, 535-548	5.1	30
290	Venous thrombosis following the use of intermediate purity FVIII concentrate to treat patients with von Willebrand's disease. *Thrombosis and Haemostasis*, **2002**, 88, 387-8	7	30
289	How to discuss gene therapy for haemophilia? A patient and physician perspective. *Haemophilia*, **2019**, 25, 545-557	3.3	29
288	Parvovirus transmission by blood products - a cause for concern?. *British Journal of Haematology*, **2012**, 159, 385-93	4.5	29
287	Disease activity only moderately correlates with quality of life impairment in patients with chronic spontaneous urticaria. *Dermatology*, **2013**, 226, 371-9	4.4	29
286	Recombinant Factor VIIa concentrate versus plasma derived concentrates for the treatment of acute bleeding episodes in people with haemophilia and inhibitors. *Cochrane Database of Systematic Reviews*, **2010**, CD004449		29
285	SSC/ISTH classification of hemophilia A: can hemophilia center laboratories achieve the new criteria?. *Journal of Thrombosis and Haemostasis*, **2004**, 2, 271-4	15.4	29
284	The definition, diagnosis and management of mild hemophilia A: communication from the SSC of the ISTH. *Journal of Thrombosis and Haemostasis*, **2018**, 16, 2530-2533	15.4	29
283	The impact of nail disorders on quality of life. *European Journal of Dermatology*, **2013**, 23, 366-71	0.8	28
282	Is VIII worse than IX?. *Blood*, **2009**, 114, 750-1	2.2	28

281	Acute generalized exanthematous pustulosis (AGEP) triggered by a spider bite. *Allergology International*, **2009**, 58, 301-3	4.4	28
280	Rupatadine inhibits inflammatory mediator release from human laboratory of allergic diseases 2 cultured mast cells stimulated by platelet-activating factor. *Annals of Allergy, Asthma and Immunology*, **2013**, 111, 542-7	3.2	27
279	Contact factor deficiencies and cardiopulmonary bypass surgery: detection of the defect and monitoring of heparin. *European Journal of Haematology*, **2009**, 82, 208-12	3.8	27
278	Characterisation of Type 2N von Willebrand Disease Using Phenotypic and Molecular Techniques. *Thrombosis and Haemostasis*, **1996**, 75, 959-964	7	26
277	Concentrate-related inhibitor risk: is a difference always real?. *Journal of Thrombosis and Haemostasis*, **2011**, 9, 2176-9	15.4	25
276	Guidelines on the diagnosis, management and prevention of hepatitis in haemophilia. *Haemophilia*, **2001**, 7, 339-45	3.3	25
275	Optimizing warfarin reversal--an ex vivo study. *Journal of Thrombosis and Haemostasis*, **2009**, 7, 1123-7	15.4	24
274	Kilt syndrome?. *British Journal of Haematology*, **2002**, 118, 1199-200	4.5	24
273	The management of von Willebrand's disease-associated gastrointestinal angiodysplasia. *Blood Coagulation and Fibrinolysis*, **2001**, 12, 143-8	1	24
272	Treatment of chronic hepatitis C in patients with haemophilia: a review of the literature. *Haemophilia*, **2006**, 12, 473-8	3.3	23
271	Major Structural Defects in the Antithrombin Gene in Four Families with Type I Antithrombin Deficiency. *Thrombosis and Haemostasis*, **2000**, 83, 715-721	7	23
270	Chronic hepatitis in haemophilia. *Blood Reviews*, **1993**, 7, 243-50	11.1	23
269	Vaccine-induced immune thrombotic thrombocytopenia. *Lancet Haematology,the*, **2021**,	14.6	23
268	Fundamentals for a Systematic Approach to Mild and Moderate Inherited Bleeding Disorders: An EHA Consensus Report. *HemaSphere*, **2019**, 3, e286	0.3	23
267	The effect of tissue factor concentration on calibrated automated thrombography in the presence of inhibitor bypass agents. *International Journal of Laboratory Hematology*, **2009**, 31, 189-98	2.5	22
266	First-line immune tolerance induction for children with severe haemophilia A: A protocol from the UK Haemophilia Centre Doctors' Organisation Inhibitor and Paediatric Working Parties. *Haemophilia*, **2017**, 23, 654-659	3.3	21
265	Pharmacokinetic properties of IB1001, an investigational recombinant factor IX, in patients with haemophilia B: repeat pharmacokinetic evaluation and sialylation analysis. *Haemophilia*, **2012**, 18, 881-7	3.3	21
264	Thrombophilia: grading the risk. *Blood*, **2009**, 113, 5038-9	2.2	21

263	Src family kinases are essential for primary aggregation by G(i) -coupled receptors. *Journal of Thrombosis and Haemostasis*, **2010**, 8, 2273-82	15.4	21
262	The successful use of protein C concentrate during pregnancy in a patient with type 1 protein C deficiency, previous thrombosis and recurrent fetal loss. *British Journal of Haematology*, **1997**, 98, 660-1	4.5	21
261	ARIA digital anamorphosis: Digital transformation of health and care in airway diseases from research to practice. *Allergy: European Journal of Allergy and Clinical Immunology*, **2021**, 76, 168-190	9.3	21
260	Genetic analysis, phenotypic diagnosis, and risk of venous thrombosis in families with inherited deficiencies of protein S. *Blood*, **2000**, 95, 1935-41	2.2	21
259	Real-life experience in switching to new extended half-life products at European haemophilia centres. *Haemophilia*, **2019**, 25, 946-952	3.3	20
258	Molecular basis of protein S deficiency in three families also showing independent inheritance of factor V leiden. *Blood*, **1996**, 88, 1700-1707	2.2	20
257	Teicoplanin administration in patients experiencing reactions to vancomycin. *Journal of Antimicrobial Chemotherapy*, **1989**, 23, 810-2	5.1	20
256	The World Federation of Hemophilia Annual Global Survey 1999-2018. *Haemophilia*, **2020**, 26, 591-600	3.3	18
255	Cognitive appraisals and psychological distress following venous thromboembolic disease: an application of the theory of cognitive adaptation. *Social Science and Medicine*, **2006**, 63, 2395-406	5.1	18
254	Sensitivity of functional protein S assays to protein S deficiency: a comparative study of three commercial kits. *Journal of Thrombosis and Haemostasis*, **2003**, 1, 1112-4	15.4	18
253	Molecular basis of protein S deficiency in three families also showing independent inheritance of factor V leiden. *Blood*, **1996**, 88, 1700-1707	2.2	18
252	The global impact of the COVID-19 pandemic on the management and course of chronic urticaria. *Allergy: European Journal of Allergy and Clinical Immunology*, **2021**, 76, 816-830	9.3	18
251	Co-inheritance of the 20210A allele of the prothrombin gene increases the risk of thrombosis in subjects with familial thrombophilia. *Thrombosis and Haemostasis*, **1997**, 78, 1426-9	7	18
250	Prophylaxis in von Willebrand Disease: Coming of Age?. *Seminars in Thrombosis and Hemostasis*, **2016**, 42, 498-506	5.3	17
249	Risk reduction strategies for variant Creutzfeldt-Jakob disease transmission by UK plasma products and their impact on patients with inherited bleeding disorders. *Haemophilia*, **2010**, 16, 305-15	3.3	17
248	The reversal of anticoagulation in clinical practice . *Clinical Medicine*, **2018**, 18, 314-319	1.9	16
247	Pregnancy outcome in women with factor V Leiden and recurrent miscarriage. *BJOG: an International Journal of Obstetrics and Gynaecology*, **2009**, 116, 995-8	3.7	16
246	Real world usage of PCC to "rapidly" correct warfarin induced coagulopathy. *Blood Transfusion*, **2013**, 11, 500-5	3.6	16

245	New findings on inhibitor development: from registries to clinical studies. *Haemophilia*, **2017**, 23 Suppl 1, 4-13	3.3	15
244	Switching clotting factor concentrates: considerations in estimating the risk of immunogenicity. *Haemophilia*, **2014**, 20, 200-6	3.3	15
243	A Novel Mutation in Intron K of the PROS1 Gene Causes Aberrant RNA Splicing and Is a Common Cause of Protein S Deficiency in a UK Thrombophilia Cohort. *Thrombosis and Haemostasis*, **1998**, 79, 1086-1091	7	15
242	Spinal epidural haematoma in haemophilia A with inhibitors--efficacy of recombinant factor VIIa concentrate. *Haemophilia*, **1999**, 5, 209-12	3.3	15
241	Warfarin Induced Skin Necrosis Associated with Activated Protein C Resistance. *Thrombosis and Haemostasis*, **1996**, 75, 523-524	7	15
240	Comparable profiles of serum histamine and IgG4 levels in allergic beekeepers. *Allergy: European Journal of Allergy and Clinical Immunology*, **2015**, 70, 457-60	9.3	14
239	Inherited Thrombophilia and Pregnancy Complications: Should We Test?. *Seminars in Thrombosis and Hemostasis*, **2019**, 45, 50-60	5.3	14
238	How I treat inhibitors in haemophilia. *Haemophilia*, **2012**, 18 Suppl 4, 48-53	3.3	14
237	Long-term management after splenectomy. Lifelong penicillin unproved in trials. *BMJ: British Medical Journal*, **1994**, 308, 131-2		14
236	Choosing and using non-steroidal anti-inflammatory drugs in haemophilia. *Haemophilia*, **2016**, 22, 179-187	3.3	14
235	Haemophilia and ageing. *British Journal of Haematology*, **2019**, 184, 712-720	4.5	14
234	Treatment of allergic rhinitis during and outside the pollen season using mobile technology. A MASK study. *Clinical and Translational Allergy*, **2020**, 10, 62	5.2	13
233	The coagulation laboratory monitoring of Afstyla single-chain FVIII concentrate. *Haemophilia*, **2017**, 23, e469-e470	3.3	13
232	Enhancing haemophilia care through registries. *Haemophilia*, **2014**, 20 Suppl 4, 121-9	3.3	13
231	Management of excessive anticoagulation or bleeding. *Seminars in Vascular Medicine*, **2003**, 3, 279-84		13
230	Influence of the -675 4G/5G dimorphism of the plasminogen activator inhibitor 1 promoter on thrombotic risk in patients with factor V Leiden. *British Journal of Haematology*, **2000**, 110, 135-8	4.5	13
229	Increased soluble IL-2 receptor levels in HCV-infected haemophiliacs: a possible indicator of liver disease severity. *British Journal of Haematology*, **1994**, 87, 419-21	4.5	13
228	Alternaria and Cladosporium calendar of Western Thrace: Relationship with allergic rhinitis symptoms. *Laryngoscope*, **2016**, 126, E51-6	3.6	13

227	The World Federation of Hemophilia guideline on management of haemophilia. *Haemophilia*, **2013**, 19, 1	3.3	12
226	A rapid, automated VWF ristocetin cofactor activity assay improves reliability in the diagnosis of Von Willebrand disease. *Thrombosis Research*, **2011**, 127, 341-4	8.2	12
225	Evaluation of the Roche LightCycler. *Blood Coagulation and Fibrinolysis*, **2003**, 14, 499-503	1	12
224	Recombinant VIIa concentrate in the management of bleeding following prothrombin complex concentrate-related myocardial infarction in patients with haemophilia and inhibitors. *British Journal of Haematology*, **2000**, 111, 974-979	4.5	12
223	The role of IL-17, IL-23 and IL-31, IL-33 in allergic skin diseases. *Current Opinion in Allergy and Clinical Immunology*, **2020**, 20, 367-373	3.3	12
222	Using pharmacokinetics for tailoring prophylaxis in people with hemophilia switching between clotting factor products: A scoping review. *Research and Practice in Thrombosis and Haemostasis*, **2019**, 3, 528-541	5.1	11
221	Kreuth V initiative: European consensus proposals for treatment of hemophilia using standard products, extended half-life coagulation factor concentrates and non-replacement therapies. *Haematologica*, **2020**, 105, 2038-2043	6.6	11
220	The clinical significance of the pollen calendar of the Western Thrace/northeast Greece region in allergic rhinitis. *International Forum of Allergy and Rhinology*, **2015**, 5, 1156-63	6.3	11
219	Inhibitor development in non-severe haemophilia across Europe. *Thrombosis and Haemostasis*, **2015** , 114, 670-5	7	11
218	Reduced cardiovascular mortality in hemophilia despite normal atherosclerotic load. *Journal of Thrombosis and Haemostasis*, **2012**, 10, 20-2	15.4	11
217	Peripheral mononuclear cells of haemophiliacs with chronic liver disease are infected with replicating hepatitis C virus. *British Journal of Haematology*, **1994**, 87, 215-7	4.5	11
216	A randomized controlled trial of recombinant interferon-alpha in chronic hepatitis C in hemophiliacs. *Blood*, **1991**, 78, 1672-7	2.2	11
215	Autoimmune Diseases Are Linked to Type IIb Autoimmune Chronic Spontaneous Urticaria. *Allergy, Asthma and Immunology Research*, **2021**, 13, 545-559	5.3	11
214	Definition, aims, and implementation of GA LEN/HAEi Angioedema Centers of Reference and Excellence. *Allergy: European Journal of Allergy and Clinical Immunology*, **2020**, 75, 2115-2123	9.3	10
213	Pharmacotherapy of chronic spontaneous urticaria. *Expert Opinion on Pharmacotherapy*, **2013**, 14, 2511-9	4	10
212	Clinical assessment of Optivate , a high-purity concentrate of factor VIII with von Willebrand factor, in the management of patients with haemophilia A. *Haemophilia*, **2011**, 17, 456-62	3.3	10
211	Maculopapular eruption to rivastigmine's transdermal patch application and successful oral desensitization. *Allergy: European Journal of Allergy and Clinical Immunology*, **2010**, 65, 925-6	9.3	10
210	Evaluation of a new venom-based clotting assay of protein C. *International Journal of Laboratory Hematology*, **2008**, 30, 437-43	2.5	10

209	Acute renal failure in acquired haemophilia following the use of high dose intravenous immunoglobulin. *Haemophilia*, **1999**, 5, 270-2	3.3	10
208	Differentiation of COVID-19 signs and symptoms from allergic rhinitis and common cold: An ARIA-EAACI-GA LEN consensus. *Allergy: European Journal of Allergy and Clinical Immunology*, **2021**, 76, 2354-2366	9.3	10
207	FVIII inhibitor development according to concentrate: data from the EUHASS registry excluding overlap with other studies. *Haemophilia*, **2016**, 22, e36-8	3.3	10
206	Pharmacokinetics, safety and efficacy of a recombinant factor IX product, trenonacog alfa in previously treated haemophilia B patients. *Haemophilia*, **2018**, 24, 104-112	3.3	9
205	Minimal dataset for post-registration surveillance of new drugs in hemophilia: communication from the SSC of the ISTH. *Journal of Thrombosis and Haemostasis*, **2017**, 15, 1878-1881	15.4	9
204	Potential misdiagnosis of dysfibrinogenaemia: Data from multicentre studies amongst UK NEQAS and PRO-RBDD project laboratories. *International Journal of Laboratory Hematology*, **2017**, 39, 653-662	2.5	9
203	Food protein-induced enterocolitis syndrome: pitfalls in the diagnosis. *Pediatric Allergy and Immunology*, **2014**, 25, 622-9	4.2	9
202	The use of biologicals in cutaneous allergies - present and future. *Current Opinion in Allergy and Clinical Immunology*, **2014**, 14, 409-16	3.3	9
201	Biovigilance and pharmacovigilance for haemophilia. *Haemophilia*, **2010**, 16 Suppl 5, 17-21	3.3	9
200	Temporal relationship of allergic rhinitis with asthma and other co-morbidities in a Mediterranean country: a retrospective study in a tertiary reference allergy clinic. *Allergologia Et Immunopathologia*, **2010**, 38, 246-53	1.9	9
199	An interpretative phenomenological analysis of adaptation to recurrent venous thrombosis and heritable thrombophilia: the importance of multi-causal models and perceptions of primary and secondary control. *Journal of Health Psychology*, **2008**, 13, 776-84	3.1	9
198	Altitude and coagulation activation: does going high provoke thrombosis?. *Acta Haematologica*, **2008**, 119, 156-7	2.7	9
197	Axillary vein thrombosis in a healthy donor following platelet apheresis. *British Journal of Haematology*, **2002**, 116, 390-1	4.5	9
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