

Pier Mannuccio Mannucci

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3312698/publications.pdf>

Version: 2024-02-01

323
papers

18,066
citations

14655

66
h-index

15732

125
g-index

329
all docs

329
docs citations

329
times ranked

13763
citing authors

#	ARTICLE	IF	CITATIONS
1	Traffic-related air pollution and the coronavirus pandemia: shadows and lights. <i>European Journal of Preventive Cardiology</i> , 2022, 28, e3-e5.	1.8	7
2	Clinical factors associated with death in 3044 COVID-19 patients managed in internal medicine wards in Italy: comment. <i>Internal and Emergency Medicine</i> , 2022, 17, 299-302.	2.0	5
3	Clinical risk scores for the early prediction of severe outcomes in patients hospitalized for COVID-19: comment. <i>Internal and Emergency Medicine</i> , 2022, 17, 303-306.	2.0	2
4	Thrombotic thrombocytopenic purpura and other immune-mediated blood disorders following vaccination against SARS-CoV-2. <i>Haematologica</i> , 2022, 107, 785-786.	3.5	2
5	Integrated care: easy in theory, harder in practice?. <i>Internal and Emergency Medicine</i> , 2022, 17, 3-6.	2.0	6
6	Predictors of inhibitor eradication by primary immune tolerance induction in severe haemophilia A with high responding inhibitors. <i>Haemophilia</i> , 2022, 28, 55-64.	2.1	7
7	Cardiovascular health and ambient air pollution: lower is not enough. <i>European Journal of Preventive Cardiology</i> , 2022, 29, 1200-1201.	1.8	5
8	Impact of Diabetes Mellitus and Its Comorbidities on Elderly Patients Hospitalized in Internal Medicine Wards: Data from the RePoSi Registry. <i>Healthcare (Switzerland)</i> , 2022, 10, 86.	2.0	10
9	Vaccine-induced immune thrombotic thrombocytopenia with atypical vein thrombosis: Implications for clinical practice. <i>Phlebology</i> , 2022, , 026835552110689.	1.2	4
10	Histological and immunohistochemical findings in a fatal case of thrombotic thrombocytopenia after ChAdOx1 nCov-19 vaccination. <i>Pathology Research and Practice</i> , 2022, 231, 153796.	2.3	5
11	Managing prostatic illness in persons with haemophilia. <i>Haemophilia</i> , 2022, 28, 369-370.	2.1	0
12	Pharmaceutical pricing in Europe: time to take the right direction. <i>Internal and Emergency Medicine</i> , 2022, 17, 945-948.	2.0	5
13	An ecological alliance against air pollution and cardiovascular disease. , 2022, 1, 19-23.		0
14	Gene transfer in hemophilia A: not cogent yet. , 2022, 1, 7-8.		2
15	What changed in the Italian internal medicine and geriatric wards during the lockdown. <i>European Journal of Internal Medicine</i> , 2021, 84, 97-100.	2.2	0
16	Intelligent game engines for home exercises (exergames) in boys with haemophilia. <i>Haemophilia</i> , 2021, 27, e151-e154.	2.1	5
17	Pharmacist-driven medication recognition/ reconciliation in older medical patients. <i>European Journal of Internal Medicine</i> , 2021, 83, 39-44.	2.2	10
18	Pattern of comorbidities and 1-year mortality in elderly patients with COPD hospitalized in internal medicine wards: data from the RePoSi Registry. <i>Internal and Emergency Medicine</i> , 2021, 16, 389-400.	2.0	18

#	ARTICLE	IF	CITATIONS
19	The multifaceted spectrum of liver cirrhosis in older hospitalised patients: analysis of the REPOSI registry. <i>Age and Ageing</i> , 2021, 50, 498-504.	1.6	1
20	Community and hospital pharmacists in Europe: encroaching on medicine?. <i>Internal and Emergency Medicine</i> , 2021, 16, 7-10.	2.0	3
21	Improving primary care in Europe beyond COVID-19: from telemedicine to organizational reforms. <i>Internal and Emergency Medicine</i> , 2021, 16, 255-258.	2.0	26
22	Authors'™ Reply to Brunetti et al. "Potentially Inappropriate Medications, Drug-Drug Interactions, and Anticholinergic Burden in Elderly Hospitalized Patients: Does an Association Exist with Post-Discharge Health Outcomes?" <i>Drugs and Aging</i> , 2021, 38, 93-94.	2.7	1
23	IgG subclasses as biomarkers for persistence of factor VIII inhibitors in previously untreated patients with severe haemophilia A. <i>British Journal of Haematology</i> , 2021, 192, 621-625.	2.5	1
24	Von Willebrand disease type 2N: An update. <i>Journal of Thrombosis and Haemostasis</i> , 2021, 19, 909-916.	3.8	14
25	Noise and air pollution as triggers of hypertension. <i>European Heart Journal</i> , 2021, 42, 2085-2087.	2.2	23
26	Sustainable and personalized nutrition: From earth health to public health. <i>European Journal of Internal Medicine</i> , 2021, 86, 12-16.	2.2	10
27	Hyperglycemia at admission, comorbidities, and in-hospital mortality in elderly patients hospitalized in internal medicine wards: data from the RePoSI Registry. <i>Acta Diabetologica</i> , 2021, 58, 1225-1236.	2.5	6
28	Performance of a clinical risk prediction model for inhibitor formation in severe haemophilia A. <i>Haemophilia</i> , 2021, 27, e441-e449.	2.1	1
29	Perspective "The case for zero bleeds and drug bioequivalence in the treatment of congenital hemophilia A in 2021. <i>Blood Reviews</i> , 2021, 50, 100849.	5.7	4
30	Post-mortem findings in vaccine-induced thrombotic thrombocytopenia. <i>Haematologica</i> , 2021, 106, 2291-2293.	3.5	47
31	COVID-19 Vaccine and Death: Causality Algorithm According to the WHO Eligibility Diagnosis. <i>Diagnostics</i> , 2021, 11, 955.	2.6	49
32	Bleeding Disorders in Primary Fibrinolysis. <i>International Journal of Molecular Sciences</i> , 2021, 22, 7027.	4.1	8
33	Use of Lipid-Lowering Drugs and Associated Outcomes According to Health State Profiles in Hospitalized Older Patients. <i>Clinical Interventions in Aging</i> , 2021, Volume 16, 1251-1264.	2.9	7
34	Comparison of quality of life, and emotional and functional profiles in older people with and without severe haemophilia. <i>Haemophilia</i> , 2021, 27, e525-e529.	2.1	1
35	Association between air temperature, air pollution and hospital admissions for pulmonary embolism and venous thrombosis in Italy. <i>European Journal of Internal Medicine</i> , 2021, , .	2.2	5
36	Pharmaceutical patenting in the European Union: reform or riddance. <i>Internal and Emergency Medicine</i> , 2021, , 1.	2.0	2

#	ARTICLE	IF	CITATIONS
37	Prophylactic management of patients with von Willebrand disease. <i>Therapeutic Advances in Hematology</i> , 2021, 12, 204062072110640.	2.5	5
38	Thrombotic thrombocytopenic purpura and other immune mediated blood disorders following SARS-CoV-2 vaccination. <i>Haematologica</i> , 2021, , .	3.5	0
39	Comorbidity does not mean clinical complexity: evidence from the RePoSI register. <i>Internal and Emergency Medicine</i> , 2020, 15, 621-628.	2.0	18
40	Use and prescription appropriateness of drugs for peptic ulcer and gastrooesophageal reflux disease in hospitalized older people. <i>European Journal of Clinical Pharmacology</i> , 2020, 76, 459-465.	1.9	13
41	The dark age of Italian general practice research â€“ An Italian matter. <i>European Journal of Internal Medicine</i> , 2020, 73, 98-99.	2.2	4
42	Pain and Frailty in Hospitalized Older Adults. <i>Pain and Therapy</i> , 2020, 9, 727-740.	3.2	22
43	SARS-CoV2 vertical transmission with adverse effects on the newborn revealed through integrated immunohistochemical, electron microscopy and molecular analyses of Placenta. <i>EBioMedicine</i> , 2020, 59, 102951.	6.1	193
44	The never ending success story of tranexamic acid in acquired bleeding. <i>Haematologica</i> , 2020, 105, 1201-1205.	3.5	27
45	Potentially Inappropriate Medications, Drugâ€“Drug Interactions, and Anticholinergic Burden in Elderly Hospitalized Patients: Does an Association Exist with Post-Discharge Health Outcomes?. <i>Drugs and Aging</i> , 2020, 37, 585-593.	2.7	27
46	Acquired von Willebrand syndrome: focused for hematologists. <i>Haematologica</i> , 2020, 105, 2032-2037.	3.5	67
47	Witnessing the 1980s. <i>Haemophilia</i> , 2020, 26, 373-374.	2.1	0
48	Hemophilia therapy: the future has begun. <i>Haematologica</i> , 2020, 105, 545-553.	3.5	132
49	Conflicts of interest in medicine: a never-ending story. <i>Internal and Emergency Medicine</i> , 2020, 15, 357-359.	2.0	6
50	Benefits and limitations of extended plasma half-life factor VIII products in hemophilia A. <i>Expert Opinion on Investigational Drugs</i> , 2020, 29, 303-309.	4.1	16
51	Defensive medicine: Everything and its opposite. <i>European Journal of Internal Medicine</i> , 2020, 74, 117-118.	2.2	6
52	Direct oral anticoagulants and cirrhosis: More evidence still needed for efficacy and safety in portal vein thrombosis. <i>Vascular Pharmacology</i> , 2019, 113, 92-93.	2.1	5
53	Novel evidence for a greater burden of ambient air pollution on cardiovascular disease. <i>Haematologica</i> , 2019, 104, 2349-2357.	3.5	88
54	AGING WITH HEMOPHILIA: THE CHALLENGE OF APPROPRIATE DRUG PRESCRIPTION. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , 2019, 11, e2019056.	1.3	7

#	ARTICLE	IF	CITATIONS
55	Drug-drug interactions involving CYP3A4 and p-glycoprotein in hospitalized elderly patients. <i>European Journal of Internal Medicine</i> , 2019, 65, 51-57.	2.2	24
56	Need for Deprescribing in Hospital Elderly Patients Discharged with a Limited Life Expectancy: The REPOSI Study. <i>Medical Principles and Practice</i> , 2019, 28, 501-508.	2.4	5
57	New therapies for von Willebrand disease. <i>Hematology American Society of Hematology Education Program</i> , 2019, 2019, 590-595.	2.5	7
58	New therapies for von Willebrand disease. <i>Blood Advances</i> , 2019, 3, 3481-3487.	5.2	29
59	Sex-Differences in the Pattern of Comorbidities, Functional Independence, and Mortality in Elderly Inpatients: Evidence from the RePoSI Register. <i>Journal of Clinical Medicine</i> , 2019, 8, 81.	2.4	21
60	Factor VIII replacement is still the standard of care in haemophilia A. <i>Blood Transfusion</i> , 2019, 17, 479-486.	0.4	53
61	Telemedicine and telerehabilitation: current and forthcoming applications in haemophilia. <i>Blood Transfusion</i> , 2019, 17, 385-390.	0.4	18
62	Use of desmopressin in the treatment of hemophilia A: towards a golden jubilee. <i>Haematologica</i> , 2018, 103, 379-381.	3.5	2
63	Prognostic relevance of glomerular filtration rate estimation obtained through different equations in hospitalized elderly patients. <i>European Journal of Internal Medicine</i> , 2018, 54, 60-64.	2.2	4
64	Primary hyperfibrinolysis: Facts and fancies. <i>Thrombosis Research</i> , 2018, 166, 71-75.	1.7	18
65	Prediction of factor VIII inhibitor development in the SIPPET cohort by mutational analysis and factor VIII antigen measurement. <i>Journal of Thrombosis and Haemostasis</i> , 2018, 16, 778-790.	3.8	23
66	How to win space in medical journals: Bits and tips. <i>European Journal of Internal Medicine</i> , 2018, 50, 1-2.	2.2	0
67	Viral safety of coagulation factor concentrates: memoirs from an insider. <i>Journal of Thrombosis and Haemostasis</i> , 2018, 16, 630-633.	3.8	8
68	The importance of ABO blood group in pharmacokinetic studies in haemophilia A. <i>Haemophilia</i> , 2018, 24, e122-e123.	2.1	5
69	Risk factors for three-month mortality after discharge in a cohort of non-oncologic hospitalized elderly patients: Results from the REPOSI study. <i>Archives of Gerontology and Geriatrics</i> , 2018, 74, 169-173.	3.0	14
70	Comorbidities in persons with haemophilia aged 60 years or more compared with age-matched people from the general population. <i>Haemophilia</i> , 2018, 24, e6-e10.	2.1	15
71	A contemporary look at FVIII inhibitor development: still a great influence on the evolution of hemophilia therapies. <i>Expert Review of Hematology</i> , 2018, 11, 87-97.	2.2	14
72	Disability, and not diabetes, is a strong predictor of mortality in oldest old patients hospitalized with pneumonia. <i>European Journal of Internal Medicine</i> , 2018, 54, 53-59.	2.2	19

#	ARTICLE	IF	CITATIONS
73	Polypharmacy in older adults with severe haemophilia. <i>Haemophilia</i> , 2018, 24, e1-e3.	2.1	4
74	Ageing successfully with haemophilia: A multidisciplinary programme. <i>Haemophilia</i> , 2018, 24, 57-62.	2.1	21
75	Polypharmacy in older people: lessons from 10 years of experience with the REPOSIT register. <i>Internal and Emergency Medicine</i> , 2018, 13, 1191-1200.	2.0	45
76	Miracle of haemophilia drugs: Personal views about a few main players. <i>Haemophilia</i> , 2018, 24, 557-562.	2.1	21
77	Mitigation of air pollution by greenness: A narrative review. <i>European Journal of Internal Medicine</i> , 2018, 55, 1-5.	2.2	55
78	Rate and appropriateness of polypharmacy in older patients with hemophilia compared with age-matched controls. <i>Haemophilia</i> , 2018, 24, 726-732.	2.1	7
79	More on air pollution and venous thromboembolism. <i>European Journal of Internal Medicine</i> , 2017, 37, e11.	2.2	1
80	Efficacy and safety of a recombinant factor VIII produced from a human cell line (simoctocog alfa). <i>Expert Opinion on Drug Safety</i> , 2017, 16, 405-410.	2.4	4
81	Laboratory monitoring of replacement therapy for major surgery in von Willebrand disease. <i>Haemophilia</i> , 2017, 23, 182-187.	2.1	18
82	Oral anticoagulant use in octogenarian European patients with atrial fibrillation: A subanalysis of PREFER in AF. <i>International Journal of Cardiology</i> , 2017, 232, 98-104.	1.7	16
83	Reply to the letter by Iorio. <i>Haemophilia</i> , 2017, 23, e248-e249.	2.1	3
84	Antipsychotic prescription and mortality in hospitalized older persons. <i>Psychogeriatrics</i> , 2017, 17, 397-405.	1.2	9
85	Progress in the contemporary management of hemophilia: The new issue of patient aging. <i>European Journal of Internal Medicine</i> , 2017, 43, 16-21.	2.2	26
86	Involvement of the IgE-C1q system and mild complement activation in haemophilia B with anti-factor IX neutralizing antibodies and anaphylaxis. <i>Haemophilia</i> , 2017, 23, e348-e353.	2.1	8
87	Homeopathy provided by a national health service: Only in Italy?. <i>European Journal of Internal Medicine</i> , 2017, 41, 1-2.	2.2	2
88	SIPPET: methodology, analysis and generalizability. <i>Haemophilia</i> , 2017, 23, 353-361.	2.1	27
89	Nonneutralizing antibodies against factor VIII and risk of inhibitor development in severe hemophilia A. <i>Blood</i> , 2017, 129, 1245-1250.	1.4	41
90	Relationship between atrial fibrillation and cognitive decline in individuals aged 80 and older. <i>European Journal of Internal Medicine</i> , 2017, 46, 6-10.	2.2	16

#	ARTICLE	IF	CITATIONS
91	Genetic risk stratification to reduce inhibitor development in the early treatment of hemophilia A: a SIPPET analysis. <i>Blood</i> , 2017, 130, 1757-1759.	1.4	44
92	Management of Hemophilia in Older Patients. <i>Drugs and Aging</i> , 2017, 34, 881-889.	2.7	6
93	Use of non-steroidal anti-inflammatory drugs and analgesics in a cohort of hospitalized elderly patients: Results from the REPOSI study. <i>European Journal of Internal Medicine</i> , 2017, 38, e11-e12.	2.2	1
94	Prognostic value of degree and types of anaemia on clinical outcomes for hospitalised older patients. <i>Archives of Gerontology and Geriatrics</i> , 2017, 69, 21-30.	3.0	17
95	Air pollution levels and cardiovascular health: Low is not enough. <i>European Journal of Preventive Cardiology</i> , 2017, 24, 1851-1853.	1.8	15
96	Health Effects of Ambient Air Pollution in Developing Countries. <i>International Journal of Environmental Research and Public Health</i> , 2017, 14, 1048.	2.6	319
97	The increased demand for plasma-derived factor VIII in Italy. <i>Blood Transfusion</i> , 2017, 15, 279-280.	0.4	7
98	Tailoring hemostatic therapies to lower inhibitor development in previously untreated patients with severe hemophilia A. <i>Journal of Thrombosis and Haemostasis</i> , 2016, 14, 1330-1336.	3.8	17
99	Ndufc2 Gene Inhibition Is Associated With Mitochondrial Dysfunction and Increased Stroke Susceptibility in an Animal Model of Complex Human Disease. <i>Journal of the American Heart Association</i> , 2016, 5, .	3.7	43
100	The safety of pharmacologic options for the treatment of persons with hemophilia. <i>Expert Opinion on Drug Safety</i> , 2016, 15, 1391-1400.	2.4	12
101	Defining Aging Phenotypes and Related Outcomes: Clues to Recognize Frailty in Hospitalized Older Patients. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2016, 72, glw188.	3.6	41
102	Predictors of clinical events occurring during hospital stay among elderly patients admitted to medical wards in Italy. <i>European Journal of Internal Medicine</i> , 2016, 32, 38-42.	2.2	6
103	Innovative Pharmacological Therapies for the Hemophilias Not Based on Deficient Factor Replacement. <i>Seminars in Thrombosis and Hemostasis</i> , 2016, 42, 526-532.	2.7	32
104	Direct oral anticoagulants and venous thromboembolism. <i>European Respiratory Review</i> , 2016, 25, 295-302.	7.1	27
105	Therapeutic Duplicates in a Cohort of Hospitalized Elderly Patients: Results from the REPOSI Study. <i>Drugs and Aging</i> , 2016, 33, 647-654.	2.7	4
106	Von Willebrand factor (VonvendiÂ®): the first recombinant product licensed for the treatment of von Willebrand disease. <i>Expert Review of Hematology</i> , 2016, 9, 825-830.	2.2	32
107	Uncertain thrombophilia markers. <i>Thrombosis and Haemostasis</i> , 2016, 115, 25-30.	3.4	32
108	Association between particulate air pollution and venous thromboembolism: A systematic literature review. <i>European Journal of Internal Medicine</i> , 2016, 27, 10-13.	2.2	43

#	ARTICLE	IF	CITATIONS
109	Imatinib and polypharmacy in very old patients with chronic myeloid leukemia: effects on response rate, toxicity and outcome. <i>Oncotarget</i> , 2016, 7, 80083-80090.	1.8	24
110	Atypical hemolytic uremic syndrome (aHUS): essential aspects of an accurate diagnosis. <i>Clinical Advances in Hematology and Oncology</i> , 2016, 14 Suppl 11, 2-15.	0.3	23
111	Half-life extension technologies for haemostatic agents. <i>Thrombosis and Haemostasis</i> , 2015, 113, 165-176.	3.4	29
112	Factor VIII products in haemophilia A: one size fits all?. <i>Thrombosis and Haemostasis</i> , 2015, 113, 911-914.	3.4	14
113	Classic thrombophilic gene variants. <i>Thrombosis and Haemostasis</i> , 2015, 114, 885-889.	3.4	93
114	Management of Bleeding Associated with New Oral Anticoagulants. <i>Seminars in Thrombosis and Hemostasis</i> , 2015, 41, 788-801.	2.7	17
115	Gender difference in drug use in hospitalized elderly patients. <i>European Journal of Internal Medicine</i> , 2015, 26, 483-490.	2.2	25
116	Recombinant porcine factor <scp>VIII</scp>: a new instalment of a long story. <i>Haemophilia</i> , 2015, 21, 149-151.	2.1	3
117	Impact on human health of climate changes. <i>European Journal of Internal Medicine</i> , 2015, 26, 1-5.	2.2	107
118	The stigma of low opioid prescription in the hospitalized multimorbid elderly in Italy. <i>Internal and Emergency Medicine</i> , 2015, 10, 305-313.	2.0	7
119	Introduction and overview. <i>Blood Reviews</i> , 2015, 29, S1-S3.	5.7	0
120	Effects on health of air pollution: a narrative review. <i>Internal and Emergency Medicine</i> , 2015, 10, 657-662.	2.0	200
121	Brain and kidney, victims of atrial microembolism in elderly hospitalized patients? Data from the REPOSI study. <i>European Journal of Internal Medicine</i> , 2015, 26, 243-249.	2.2	14
122	Adherence to antibiotic treatment guidelines and outcomes in the hospitalized elderly with different types of pneumonia. <i>European Journal of Internal Medicine</i> , 2015, 26, 330-337.	2.2	25
123	Understanding organ dysfunction in thrombotic thrombocytopenic purpura. <i>Intensive Care Medicine</i> , 2015, 41, 715-718.	8.2	17
124	The complex differential diagnosis between thrombotic thrombocytopenic purpura and the atypical hemolytic uremic syndrome: Laboratory weapons and their impact on treatment choice and monitoring. <i>Thrombosis Research</i> , 2015, 136, 851-854.	1.7	27
125	Appropriateness of antithrombotic prophylaxis in the oldest old with non-valvular atrial fibrillation: ARAPACIS and REPOSI. <i>European Journal of Internal Medicine</i> , 2015, 26, e47-e48.	2.2	1
126	The Health and Economic Burden of Air Pollution. <i>American Journal of Medicine</i> , 2015, 128, 931-932.	1.5	24

#	ARTICLE	IF	CITATIONS
127	Under-detection of delirium and impact of neurocognitive deficits on in-hospital mortality among acute geriatric and medical wards. <i>European Journal of Internal Medicine</i> , 2015, 26, 696-704.	2.2	65
128	Gastrointestinal angiodysplasia and bleeding in von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2014, 112, 427-431.	3.4	60
129	The real value of thrombophilia markers in identifying patients at high risk of venous thromboembolism. <i>Expert Review of Hematology</i> , 2014, 7, 757-765.	2.2	19
130	Reply to: The importance and challenge of pediatric trials of hemophilia drugs. <i>Nature Medicine</i> , 2014, 20, 466-466.	30.7	3
131	Pediatric requirements in Europe stymie help for hemophilia. <i>Nature Medicine</i> , 2014, 20, 117-117.	30.7	8
132	Making clinical decisions on the basis of RODIN. <i>Haemophilia</i> , 2014, 20, e174-e175.	2.1	0
133	Emerging drugs for hemophilia B. <i>Expert Opinion on Emerging Drugs</i> , 2014, 19, 407-414.	2.4	4
134	Integrated postural analysis in children with haemophilia. <i>Haemophilia</i> , 2014, 20, 263-267.	2.1	12
135	Multimorbidity and polypharmacy in the elderly: lessons from REPOSI. <i>Internal and Emergency Medicine</i> , 2014, 9, 723-734.	2.0	121
136	Haematology clinic: Haemophilia A. <i>Hematology</i> , 2014, 19, 181-182.	1.5	11
137	Prognostic value of estimated glomerular filtration rate in hospitalized elderly patients. <i>Internal and Emergency Medicine</i> , 2014, 9, 735-747.	2.0	16
138	Gout, allopurinol intake and clinical outcomes in the hospitalized multimorbid elderly. <i>European Journal of Internal Medicine</i> , 2014, 25, 847-852.	2.2	1
139	Novel investigations on the protective role of the <sc>FVIII</sc>/<sc>VWF</sc> complex in inhibitor development. <i>Haemophilia</i> , 2014, 20, 2-16.	2.1	17
140	Prevalence of potentially inappropriate medications and risk of adverse clinical outcome in a cohort of hospitalized elderly patients: results from the REPOSI Study. <i>Journal of Clinical Pharmacy and Therapeutics</i> , 2014, 39, 511-515.	1.5	59
141	Gender-differences in disease distribution and outcome in hospitalized elderly: Data from the REPOSI study. <i>European Journal of Internal Medicine</i> , 2014, 25, 617-623.	2.2	75
142	Inhibitor eradication with rituximab in haemophilia: where do we stand?. <i>British Journal of Haematology</i> , 2014, 165, 600-608.	2.5	51
143	Procoagulant imbalance in patients with non-alcoholic fatty liver disease. <i>Journal of Hepatology</i> , 2014, 61, 148-154.	3.7	149
144	Prescription Drug Use Among Older Adults in Italy: A Country-Wide Perspective. <i>Journal of the American Medical Directors Association</i> , 2014, 15, 531.e11-531.e15.	2.5	22

#	ARTICLE	IF	CITATIONS
145	The bleeding score predicts clinical outcomes and replacement therapy in adults with von Willebrand disease. <i>Blood</i> , 2014, 123, 4037-4044.	1.4	123
146	Dynamics of complement activation in aHUS and how to monitor eculizumab therapy. <i>Blood</i> , 2014, 124, 1715-1726.	1.4	288
147	Haemophilia Centre Certification Systems: optional or optimal choice for healthcare systems?. <i>Blood Transfusion</i> , 2014, 12 Suppl 3, s492-4.	0.4	0
148	The European standards of Haemophilia Centres. <i>Blood Transfusion</i> , 2014, 12 Suppl 3, s525-30.	0.4	12
149	The methodology for defining the European standards for the certification of Haemophilia Centres in Europe. <i>Blood Transfusion</i> , 2014, 12 Suppl 3, s519-24.	0.4	4
150	A certification/accreditation model for Haemophilia Centres in Italy. <i>Blood Transfusion</i> , 2014, 12 Suppl 3, s505-9.	0.4	6
151	Prophylaxis in severe forms of von Willebrand's disease: results from the von Willebrand Disease Prophylaxis Network (VWD PN). <i>Haemophilia</i> , 2013, 19, 76-81.	2.1	99
152	Association of Anticholinergic Burden with Cognitive and Functional Status in a Cohort of Hospitalized Elderly: Comparison of the Anticholinergic Cognitive Burden Scale and Anticholinergic Risk Scale. <i>Drugs and Aging</i> , 2013, 30, 103-112.	2.7	140
153	Is haemophilia B less severe than haemophilia A?. <i>Haemophilia</i> , 2013, 19, 499-502.	2.1	61
154	Thromboprophylaxis in the oldest old with atrial fibrillation: Between Scylla and Charybdis. <i>European Journal of Internal Medicine</i> , 2013, 24, 285-287.	2.2	12
155	Investigational drugs for coagulation disorders. <i>Expert Opinion on Investigational Drugs</i> , 2013, 22, 945-953.	4.1	12
156	Von Willebrand disease-associated angiodysplasia: a few answers, still many questions. <i>British Journal of Haematology</i> , 2013, 161, 177-182.	2.5	68
157	Evolution of the European guidelines for the clinical development of factor VIII products: little progress towards improved patient management. <i>Haemophilia</i> , 2013, 19, 344-348.	2.1	11
158	Drug-drug interactions in a cohort of hospitalized elderly patients. <i>Pharmacoepidemiology and Drug Safety</i> , 2013, 22, 1054-1060.	1.9	53
159	Pharmacokinetics and safety of a novel recombinant human von Willebrand factor manufactured with a plasma-free method: a prospective clinical trial. <i>Blood</i> , 2013, 122, 648-657.	1.4	120
160	Acquired haemophilia A: A 2013 update. <i>Thrombosis and Haemostasis</i> , 2013, 110, 1114-1120.	3.4	103
161	Liver disease, coagulopathies and transfusion therapy. <i>Blood Transfusion</i> , 2013, 11, 32-6.	0.4	26
162	Plasma-derived medicinal products: demand and clinical use. <i>Blood Transfusion</i> , 2013, 11 Suppl 4, s2-5.	0.4	4

#	ARTICLE	IF	CITATIONS
163	The demand for factor VIII and for factor IX and the toll fractionation product surplus management. Blood Transfusion, 2013, 11 Suppl 4, s64-76.	0.4	12
164	Present and future challenges in the treatment of haemophilia: a clinician's perspective. Blood Transfusion, 2013, 11 Suppl 4, s77-81.	0.4	20
165	Recombinant factor VIIa as haemostatic therapy in advanced liver disease. Blood Transfusion, 2013, 11, 487-90.	0.4	4
166	Prophylactic efficacy and pharmacokinetically guided dosing of a von Willebrand factor/factor VIII concentrate in adults and children with von Willebrand's disease undergoing elective surgery: a pooled and comparative analysis of data from USA and European Union clinical trials. Blood Transfusion, 2013, 11, 533-40.	0.4	14
167	Management of antithrombotic therapy for acute coronary syndromes and atrial fibrillation in patients with hemophilia. Expert Opinion on Pharmacotherapy, 2012, 13, 505-510.	1.8	20
168	Getting rid of refractory hemophilia. Blood, 2012, 119, 1326-1327.	1.4	1
169	How we choose factor VIII to treat hemophilia. Blood, 2012, 119, 4108-4114.	1.4	74
170	Air pollution and cardiovascular disease. Thrombosis Research, 2012, 129, 230-234.	1.7	91
171	Health-related quality of life and psychological well-being in elderly patients with haemophilia. Haemophilia, 2012, 18, 345-352.	2.1	29
172	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.	2.1	88
173	The role of natural VWF/FVIII complex concentrates in contemporary haemophilia care: a guideline for the next decade. Haemophilia, 2012, 18, 2-7.	2.1	10
174	Clinical governance: many fancies, very few facts. Internal and Emergency Medicine, 2012, 7, 3-4.	2.0	3
175	Hemostatic defects in liver and renal dysfunction. Hematology American Society of Hematology Education Program, 2012, 2012, 168-73.	2.5	15
176	Hemostatic defects in liver and renal dysfunction. Hematology American Society of Hematology Education Program, 2012, 2012, 168-173.	2.5	48
177	ADAMTS13 Content of Plasma-Derived Factor VIII-Von Willebrand Factor Concentrates. Blood, 2012, 120, 3383-3383.	1.4	1
178	Old and new anticoagulant drugs: A minireview. Annals of Medicine, 2011, 43, 116-123.	3.8	48
179	The Coagulopathy of Chronic Liver Disease. New England Journal of Medicine, 2011, 365, 147-156.	27.0	1,171
180	Multiple Diseases and Polypharmacy in the Elderly: Challenges for the Internist of the Third Millennium. Journal of Comorbidity, 2011, 1, 28-44.	3.9	192

#	ARTICLE	IF	CITATIONS
181	Thrombogenicity and cardiovascular effects of ambient air pollution. <i>Blood</i> , 2011, 118, 2405-2412.	1.4	167
182	Treatment of haemophilia: building on strength in the third millennium. <i>Haemophilia</i> , 2011, 17, 1-24.	2.1	16
183	A randomized clinical trial of prophylaxis in children with hemophilia A (the ESPRIT Study). <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 700-710.	3.8	390
184	Polypharmacy, length of hospital stay, and in-hospital mortality among elderly patients in internal medicine wards. The REPOSI study. <i>European Journal of Clinical Pharmacology</i> , 2011, 67, 507-519.	1.9	255
185	Fine particulate: it matters. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 659-661.	3.8	10
186	Red cells playing as activated platelets in thalassemia intermedia. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 2149-2151.	3.8	23
187	Efficacy and safety of highly purified, doubly virus-inactivated VWF/FVIII concentrates in inherited von Willebrand's disease: results of an Italian cohort study on 120 patients characterized by bleeding severity score. <i>Haemophilia</i> , 2010, 16, 101-110.	2.1	39
188	Comorbidities and quality of life in elderly persons with haemophilia. <i>British Journal of Haematology</i> , 2010, 148, 522-533.	2.5	90
189	Aspirin as antiplatelet agent in diabetes: Cons. <i>European Journal of Internal Medicine</i> , 2010, 21, 154-156.	2.2	1
190	Mechanism of hemostasis defects and management of bleeding in patients with acute coronary syndromes. <i>European Journal of Internal Medicine</i> , 2010, 21, 254-259.	2.2	9
191	Autoimmune hemophilia at rescue. <i>Haematologica</i> , 2009, 94, 459-461.	3.5	9
192	Activated Partial Thromboplastin Time. <i>Scandinavian Journal of Haematology</i> , 2009, 25, 308-317.	0.0	8
193	von Willebrand disease in the 21st century: current approaches and new challenges. <i>Haemophilia</i> , 2009, 15, 1154-1158.	2.1	11
194	Health status and quality of life of elderly persons with severe hemophilia born before the advent of modern replacement therapy. <i>Journal of Thrombosis and Haemostasis</i> , 2009, 7, 780-786.	3.8	122
195	Factor VIII gene (F8) mutations as predictors of outcome in immune tolerance induction of hemophilia A patients with high-responding inhibitors. <i>Journal of Thrombosis and Haemostasis</i> , 2009, 7, 1809-1815.	3.8	103
196	Rate of Inhibitor Development in Hemophilia A Patients Treated with Plasma Derived or Recombinant Factor VIII Concentrates. A Systematic Review of the Literature. <i>Blood</i> , 2009, 114, 3154-3154.	1.4	0
197	Evidence-based recommendations on the treatment of von Willebrand disease in Italy. <i>Blood Transfusion</i> , 2009, 7, 117-26.	0.4	63
198	Desmopressin: an historical introduction. <i>Haemophilia</i> , 2008, 14, 1-4.	2.1	12

#	ARTICLE	IF	CITATIONS
199	Dosing anticoagulant therapy with coumarin drugs: is genotyping clinically useful? No. Journal of Thrombosis and Haemostasis, 2008, 6, 1450-1452.	3.8	8
200	Italian Registry of Haemophilia and Allied Disorders. Objectives, methodology and data analysis. Haemophilia, 2008, 14, 444-453.	2.1	88
201	Back to the future: a recent history of haemophilia treatment. Haemophilia, 2008, 14, 10-18.	2.1	137
202	Understanding hemostasis and thrombosis : The lesson learnt from natural models. Japanese Journal of Thrombosis and Hemostasis, 2008, 19, 85-91.	0.1	0
203	Clinical use of Haemate(R) P in inherited von Willebrand's disease: a cohort study on 100 Italian patients. Haematologica, 2007, 92, 944-951.	3.5	85
204	Prevention and Treatment of Major Blood Loss. New England Journal of Medicine, 2007, 356, 2301-2311.	27.0	445
205	Reply:. Hepatology, 2007, 45, 832-833.	7.3	1
206	Immune tolerance induction with a high purity von Willebrand factor/VIII complex concentrate in haemophilia A patients with inhibitors at high risk of a poor response. Haemophilia, 2007, 13, 373-379.	2.1	81
207	Factor VIII products and inhibitor development: the SIPPET study (survey of inhibitors in) Tj ETQq1 1 0.784314 rgBT/Overlock 10 Tf 50	2.1	90
208	Thrombosis and bleeding disorders outside Western countries. Journal of Thrombosis and Haemostasis, 2007, 5, 68-72.	3.8	8
209	Treatment of severe von Willebrand disease with a high purity von Willebrand factor concentrate (Wilfactin®): a prospective study of 50 patients. Journal of Thrombosis and Haemostasis, 2007, 5, 1115-1124.	3.8	111
210	von Willebrand factor/factor VIII concentrate (Haemate® P) dosing based on pharmacokinetics: a prospective multicenter trial in elective surgery. Journal of Thrombosis and Haemostasis, 2007, 5, 1420-1430.	3.8	118
211	Short-term effects of air pollution on cardiovascular diseases: outcomes and mechanisms. Journal of Thrombosis and Haemostasis, 2007, 5, 2169-2174.	3.8	115
212	Editorial. Journal of Thrombosis and Haemostasis, 2006, 4, 1-2.	3.8	10
213	Need for randomized trials in hemophilia. Journal of Thrombosis and Haemostasis, 2006, 4, 501-502.	3.8	12
214	Abnormal hemostasis tests and bleeding in chronic liver disease: are they related? No. Journal of Thrombosis and Haemostasis, 2006, 4, 721-723.	3.8	82
215	Update on the pathophysiology and classification of von Willebrand disease: a report of the Subcommittee on von Willebrand Factor. Journal of Thrombosis and Haemostasis, 2006, 4, 2103-2114.	3.8	1,061
216	Non-Sense-Mediated mRNA Decay in ADAMTS13 Gene Caused by 29 Nucleotide Deletion.. Blood, 2006, 108, 1061-1061.	1.4	0

#	ARTICLE	IF	CITATIONS
217	ADAMTS13 autoantibodies in patients with thrombotic microangiopathies and other immunomediated diseases. <i>Blood</i> , 2005, 106, 1262-1267.	1.4	275
218	Thrombolytic therapy in acute myocardial infarction. <i>Journal of Thrombosis and Haemostasis</i> , 2005, 3, 2806-2807.	3.8	0
219	Localization and Function of Platelet ADAMTS-13.. <i>Blood</i> , 2005, 106, 3967-3967.	1.4	0
220	Plasma levels of von Willebrand factor regulate ADAMTS-13, its major cleaving protease. <i>British Journal of Haematology</i> , 2004, 126, 213-218.	2.5	99
221	llsley Ingram. <i>Journal of Thrombosis and Haemostasis</i> , 2004, 2, 1219-1219.	3.8	0
222	Treatment of von Willebrand's Disease. <i>New England Journal of Medicine</i> , 2004, 351, 683-694.	27.0	464
223	Recessively inherited coagulation disorders. <i>Blood</i> , 2004, 104, 1243-1252.	1.4	479
224	The International Society for Thrombosis and Haemostasis owns its official journal: the future has begun!. <i>Journal of Thrombosis and Haemostasis</i> , 2003, 1, 1-2.	3.8	33
225	Consistency of ADAMTS-13 activity assays: a moderately optimistic view. <i>Journal of Thrombosis and Haemostasis</i> , 2003, 1, 1880-1881.	3.8	8
226	AIDS, hepatitis and hemophilia in the 1980s: memoirs from an insider. <i>Journal of Thrombosis and Haemostasis</i> , 2003, 1, 2065-2069.	3.8	24
227	Cost of care and quality of life for patients with hemophilia complicated by inhibitors: the COCIS Study Group. <i>Blood</i> , 2003, 102, 2358-2363.	1.4	351
228	Overview of Bleeding in Cancer Patients. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 2003, 33, 44-45.	0.3	6
229	Patients with localized and disseminated tumors have reduced but measurable levels of ADAMTS-13 (von Willebrand factor cleaving protease). <i>Haematologica</i> , 2003, 88, 454-8.	3.5	37
230	Von Willebrand factor cleaving protease (ADAMTS-13) in 123 patients with connective tissue diseases (systemic lupus erythematosus and systemic sclerosis). <i>Haematologica</i> , 2003, 88, 914-8.	3.5	85
231	Venous thrombosis: the history of knowledge. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 2002, 32, 209-212.	0.3	24
232	Venous thromboembolism in von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2002, 88, 378-9.	3.4	41
233	How I treat patients with von Willebrand disease. <i>Blood</i> , 2001, 97, 1915-1919.	1.4	157
234	Changes in health and disease of the metalloprotease that cleaves von Willebrand factor. <i>Blood</i> , 2001, 98, 2730-2735.	1.4	488

#	ARTICLE	IF	CITATIONS
235	Venous thrombosis and anticoagulant therapy. <i>British Journal of Haematology</i> , 2001, 114, 258-270.	2.5	31
236	Coexistence of a novel homozygous nonsense mutation in exon 13 of the factor V gene with the homozygous Leiden mutation in two unrelated patients with severe factor V deficiency. <i>British Journal of Haematology</i> , 2001, 114, 871-874.	2.5	20
237	The Hemophilias " From Royal Genes to Gene Therapy. <i>New England Journal of Medicine</i> , 2001, 344, 1773-1779.	27.0	936
238	Use of Prophylaxis to Prevent Complications of Hemophilia. <i>Advances in Experimental Medicine and Biology</i> , 2001, 489, 59-64.	1.6	13
239	Treatment of von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2001, 86, 149-53.	3.4	10
240	The search for the causes of myocardial infarction. Introduction. <i>Italian Heart Journal: Official Journal of the Italian Federation of Cardiology</i> , 2001, 2, 489.	0.1	0
241	Autoimmune bleeding disorders in cancer patients. <i>Haemostasis</i> , 2001, 31 Suppl 1, 45-6.	0.0	0
242	Desmopressin (DDAVP) in the treatment of bleeding disorders: the first twenty years. <i>Haemophilia</i> , 2000, 6, 60-67.	2.1	104
243	A novel two base pair deletion in the factor V gene associated with severe factor V deficiency. <i>British Journal of Haematology</i> , 2000, 111, 1240-1246.	2.5	1
244	The risk of venous thromboembolism in family members with mutations in the genes of factor V or prothrombin or both. <i>British Journal of Haematology</i> , 2000, 111, 1223-1229.	2.5	6
245	Clinical manifestations and complications of childbirth and replacement therapy in 385 Iranian patients with type 3 von Willebrand disease. <i>British Journal of Haematology</i> , 2000, 111, 1236-1239.	2.5	91
246	Aspects of the Clinical Management of Hereditary Thrombophilia: A Personal Perspective. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 2000, 30, 11-15.	0.3	3
247	Choice of replacement therapy for hemophilia: recombinant products only?. <i>The Hematology Journal</i> , 2000, 1, 72-76.	1.4	6
248	Gly319"Arg substitution in the dysfunctional prothrombin Segovia. <i>British Journal of Haematology</i> , 1999, 105, 667-669.	2.5	12
249	Thrombotic thrombocytopenic purpura: a simpler diagnosis at last?. <i>Thrombosis and Haemostasis</i> , 1999, 82, 1380-1.	3.4	3
250	Treatment of von Willebrand disease. <i>International Journal of Clinical and Laboratory Research</i> , 1998, 28, 211-214.	1.0	7
251	Treatment of von Willebrand disease. <i>Haemophilia</i> , 1998, 4, 661-664.	2.1	30
252	Hemostatic Drugs. <i>New England Journal of Medicine</i> , 1998, 339, 245-253.	27.0	581

#	ARTICLE	IF	CITATIONS
253	Treatment of von Willebrand Disease. Hematology, 1998, 3, 339-346.	1.5	15
254	Different Risks of Thrombosis in Four Coagulation Defects Associated With Inherited Thrombophilia: A Study of 150 Families. Blood, 1998, 92, 2353-2358.	1.4	378
255	A Common Mutation in the Methylenetetrahydrofolate Reductase Gene (C677T) Increases the Risk for Deep-Vein Thrombosis in Patients With Mutant Factor V (Factor V:Q 506). Arteriosclerosis, Thrombosis, and Vascular Biology, 1997, 17, 1662-1666.	2.4	161
256	Desmopressin (DDAVP) in the Treatment of Bleeding Disorders: The First 20 Years. Blood, 1997, 90, 2515-2521.	1.4	466
257	Partial inhibition of platelet aggregation by nebulized pentamidine in severe haemophiliacs. Haemophilia, 1997, 3, 31-34.	2.1	1
258	Clinical manifestations in 28 Italian and Iranian patients with severe factor VII deficiency. Haemophilia, 1997, 3, 242-246.	2.1	92
259	Resistance to activated protein C in unselected patients with arterial and venous thrombosis. American Journal of Hematology, 1997, 55, 59-64.	4.1	19
260	Deficiency of (33P)2MeS-ADP Binding Sites on Platelets with Secretion Defect, Normal Granule Stores and Normal Thromboxane A2 Production. Thrombosis and Haemostasis, 1997, 77, 0986-0990.	3.4	76
261	Transfusion Requirements Are Correlated with the Degree of Proteolysis of von Willebrand Factor during Orthotopic Liver Transplantation. Thrombosis and Haemostasis, 1997, 78, 813-819.	3.4	6
262	High Frequency of the C677T Mutation in the Methylenetetrahydrofolate Reductase (MTHFR) Gene in Northern Italy. Thrombosis and Haemostasis, 1997, 78, 963-964.	3.4	43
263	Desmopressin (DDAVP) in the treatment of bleeding disorders: the first 20 years. Blood, 1997, 90, 2515-21.	1.4	88
264	Treatment of von Willebrand's disease. Journal of Internal Medicine Supplement, 1997, 740, 129-32.	0.0	1
265	Proteolysis of von Willebrand factor is decreased in acute promyelocytic leukaemia by treatment with all-trans-retinoic acid. British Journal of Haematology, 1996, 92, 733-739.	2.5	27
266	William Hewson and the blood which issued last but clotted first: the beginning of the story of desmopressin in haemophilia and vWD. Haemophilia, 1996, 2, 180-183.	2.1	1
267	Hemostasis Abnormalities in Patients with Vascular Dementia and Alzheimer's Disease. Thrombosis and Haemostasis, 1996, 75, 216-218.	3.4	60
268	Frequency of Factor V Arg506 Gin in Italians. Thrombosis and Haemostasis, 1996, 75, 694-694.	3.4	4
269	Oral Contraceptives Are a Risk Factor for Cerebral Vein Thrombosis. Thrombosis and Haemostasis, 1996, 76, 477-478.	3.4	20
270	B19 Parvovirus Withstands "Super Heating" in Antihemophilic Concentrates. Thrombosis and Haemostasis, 1996, 76, 821-821.	3.4	7

#	ARTICLE	IF	CITATIONS
271	Purinoceptors on blood platelets: further pharmacological and clinical evidence to suggest the presence of two ADP receptors. <i>British Journal of Haematology</i> , 1995, 91, 434-444.	2.5	134
272	The knowledge and perception of factor concentrate in persons with haemophilia A. <i>Haemophilia</i> , 1995, 1, 232-235.	2.1	1
273	Viral safety of plasma-derived and recombinant products used in the management of haemophilia A and B. <i>Haemophilia</i> , 1995, 1, 14-20.	2.1	6
274	Effects of Factor VIII Concentrates on the Immune System of Patients with Hemophilia. <i>Thrombosis and Haemostasis</i> , 1995, 74, 437-439.	3.4	7
275	The Effect of Desmopressin on Reducing Blood Loss in Cardiac Surgery – A Meta-Analysis of Double-Blind, Placebo-Controlled Trials. <i>Thrombosis and Haemostasis</i> , 1995, 74, 1064-1070.	3.4	89
276	Very Low Frequency of “Normandy Type” Mutations among Type 1 von Willebrand Disease Families. <i>Thrombosis and Haemostasis</i> , 1995, 73, 324-324.	3.4	4
277	Preliminary In Vivo Evaluation of a Nanofiltered Factor IX Concentrate. <i>Thrombosis and Haemostasis</i> , 1995, 73, 737-738.	3.4	2
278	Risk of Inhibitor Development after Intermediate Purity Coagulation Factor Concentrates: Is It Truly Low?. <i>Thrombosis and Haemostasis</i> , 1995, 74, 997-997.	3.4	3
279	Patients with severe von Willebrand disease are insensitive to the releasing effect of DDAVP: evidence that the DDAVP-induced increase in plasma factor VIII is not secondary to the increase in plasma von Willebrand factor. <i>British Journal of Haematology</i> , 1994, 86, 333-337.	2.5	18
280	Repeated Release of the Tissue Factor Pathway Inhibitor. <i>Thrombosis and Haemostasis</i> , 1994, 72, 327-328.	3.4	6
281	Clinical Evaluation of Viral Safety of Coagulation Factor VIII and IX Concentrates. <i>Vox Sanguinis</i> , 1993, 64, 197-203.	1.5	73
282	Type I von Willebrand disease, subtype “platelet low”: decreased platelet adhesion can be explained by low synthesis of von Willebrand factor in endothelial cells. <i>British Journal of Haematology</i> , 1993, 83, 88-93.	2.5	26
283	Antiplatelet Agents in Acute Myocardial Infarction and in the Post-infarction Period. <i>Platelets</i> , 1993, 4, 24-25.	2.3	0
284	Pharmacokinetics of Monoclonally-Purified and Recombinant Factor VIII in Patients with Severe von Willebrand Disease. <i>Thrombosis and Haemostasis</i> , 1993, 70, 270-272.	3.4	60
285	Modern Treatment of Hemophilia: From the Shadows Towards the Light. <i>Thrombosis and Haemostasis</i> , 1993, 70, 017-023.	3.4	31
286	Comparison of Functional Assays for Protein S: European Collaborative Study of Patients with Congenital and Acquired Deficiency. <i>Thrombosis and Haemostasis</i> , 1993, 70, 0946-0950.	3.4	34
287	Risk of thromboembolic complications in patients with inflammatory bowel disease. <i>International Journal of Clinical and Laboratory Research</i> , 1992, 21, 165-170.	1.0	48
288	Patterns of development of tachyphylaxis in patients with haemophilia and von Willebrand disease after repeated doses of desmopressin (DDAVP). <i>British Journal of Haematology</i> , 1992, 82, 87-93.	2.5	169

#	ARTICLE	IF	CITATIONS
289	Sustained correction of the bleeding time in an afibrinogenaemic patient after infusion of fresh frozen plasma. <i>British Journal of Haematology</i> , 1992, 82, 388-390.	2.5	28
290	Intramuscular anti- ϵ D immunoglobulins for home treatment of chronic immune thrombocytopenic purpura. <i>British Journal of Haematology</i> , 1992, 80, 337-340.	2.5	27
291	Immune Status of Asymptomatic HIV-Infected Hemophiliacs: Randomized, Prospective, Two-Year Comparison of Treatment with a High-Purity or an Intermediate-Purity Factor VIII Concentrate. <i>Thrombosis and Haemostasis</i> , 1992, 67, 310-313.	3.4	32
292	The Effect of Instrumentation on Thromboplastin Calibration. <i>Thrombosis and Haemostasis</i> , 1992, 67, 588-589.	3.4	19
293	Comparison of four virus-inactivated plasma concentrates for treatment of severe von Willebrand disease: a cross-over randomized trial. <i>Blood</i> , 1992, 79, 3130-7.	1.4	20
294	Low Levels of the Anticoagulant Activity of Protein C in Patients with Chronic Renal Insufficiency: an Inhibitor of Protein C Is Present in Uremic Plasma. <i>Thrombosis and Haemostasis</i> , 1991, 66, 420-425.	3.4	28
295	Adverse Effects of Treatment with Porcine Factor VIII. <i>Thrombosis and Haemostasis</i> , 1991, 65, 245-247.	3.4	47
296	Spontaneous platelet aggregation during pregnancy in a patient with von Willebrand disease type IIB can be blocked by monoclonal antibodies to both platelet glycoproteins Ib and IIb/IIIa. <i>British Journal of Haematology</i> , 1990, 75, 86-91.	2.5	2
297	Restriction of Polymerase Chain Reaction Products for Carrier. <i>Thrombosis and Haemostasis</i> , 1990, 63, 527-528.	3.4	2
298	Antibody to Hepatitis G Virus after a Vapour-Heated Factor VIII Concentrate. <i>Thrombosis and Haemostasis</i> , 1990, 64, 232-234.	3.4	26
299	Subcutaneous Desmopressin (DDAVP) Shortens the Prolonged Bleeding Time in Patients with Liver Cirrhosis. <i>Thrombosis and Haemostasis</i> , 1990, 64, 358-360.	3.4	45
300	Dissociated antibody responses to the s and pre-s2 regions of the hepatitis b virus after vaccination in hemophiliacs. <i>Journal of Medical Virology</i> , 1989, 28, 156-158.	5.0	5
301	Subcutaneous desmopressin (DDAVP) shortens the bleeding time in uremia. <i>American Journal of Hematology</i> , 1989, 31, 32-35.	4.1	59
302	Type II H von willebrand disease: New structural abnormality of plasma and platelet von willebrand factor in a patient with prolonged bleeding time and borderline levels of ristocetin cofactor activity. <i>American Journal of Hematology</i> , 1989, 32, 287-293.	4.1	18
303	Monoclonal Antibodies Directed to the Calcium-Free Conformation of Human Protein S. <i>Thrombosis and Haemostasis</i> , 1989, 62, 708-714.	3.4	12
304	Familial Dysfunction of Protein S. <i>Thrombosis and Haemostasis</i> , 1989, 62, 763-766.	3.4	34
305	Fibrin(ogen) Peptides in Early Breast Cancer. <i>Thrombosis and Haemostasis</i> , 1989, 62, 819.	3.4	2
306	Antithrombin Milano, Single Amino Acid Substitution at the Reactive Site, Arg393 to Cys. <i>Thrombosis and Haemostasis</i> , 1988, 60, 471-475.	3.4	29

#	ARTICLE	IF	CITATIONS
307	EVALUATION OF A FULLY AUTOMATED CENTRIFUGAL ANALYSER FOR ITS PERFORMANCE OF HEMOSTASIS TESTS. , 1987, 58, 0457.		0
308	Combined Use of DNA Probes in First-Trimester Prenatal Diagnosis of Hemophilia A. Thrombosis and Haemostasis, 1987, 58, 988-992.	3.4	4
309	Intravenous and Subcutaneous Administration of Desmopressin (DDAVP) to Hemophiliacs: Pharmacokinetics and Factor VIII Responses. Thrombosis and Haemostasis, 1987, 58, 1037-1039.	3.4	69
310	Multicenter Comparison of Five Functional and Two Immunological Assays for Protein C. Thrombosis and Haemostasis, 1987, 57, 044-048.	3.4	26
311	Laboratory Screening of Inherited Thrombotic Syndromes. Thrombosis and Haemostasis, 1987, 57, 247-251.	3.4	67
312	Fibrinogens "Milano II" and "Naples". Thrombosis and Haemostasis, 1987, 57, 375-375.	3.4	2
313	Von Willebrand factor contaminating porcine factor VIII concentrate (Hyate: C) causes platelet aggregation. British Journal of Haematology, 1986, 63, 703-711.	2.5	31
314	Fibrinogen Milano II: A Congenital Dysfibrinogenaemia Associated with Juvenile Arterial and Venous Thrombosis. Thrombosis and Haemostasis, 1986, 55, 131-135.	3.4	34
315	Electroblot and Immunoperoxidase Staining for Rapid Screening of the Abnormalities of the Multimeric Structure of von Willebrand Factor in von Willebrand's Disease. Thrombosis and Haemostasis, 1986, 55, 246-249.	3.4	13
316	Characterization of an Abnormal Antithrombin (Milano 2) with Defective Thrombin Binding. Thrombosis and Haemostasis, 1986, 56, 349-352.	3.4	6
317	Charge heterogeneity of human protein C revealed by isoelectric focusing in immobilized pH gradients. Electrophoresis, 1985, 6, 373-376.	2.4	14
318	ABSENCE OF ANTIBODIES TO AIDS VIRUS IN HAEMOPHILIACS TREATED WITH HEAT-TREATED FACTOR VIII CONCENTRATE. Lancet, The, 1985, 325, 271-272.	13.7	91
319	TRANSMISSION OF NON-A, NON-B HEPATITIS BY HEAT-TREATED FACTOR VIII CONCENTRATE. Lancet, The, 1985, 326, 1-4.	13.7	188
320	Human Platelet Aggregation and Release Reaction Induced by Platelet Activating Factor (PAF-Acether) " Effects of Acetylsalicylic Acid and External Ionized Calcium. Thrombosis and Haemostasis, 1985, 53, 221-224.	3.4	16
321	Hemostatic Functions in Hypoalpha and Hyperbetalipoproteinemias. Thrombosis and Haemostasis, 1983, 50, 624-625.	3.4	5
322	Biochemical and Metabolic Aspects of Platelet Dysfunction in Chronic Myeloproliferative Disorders. Thrombosis and Haemostasis, 1982, 47, 084-089.	3.4	91
323	Health technology assessment for pharmaceuticals in the European Union: what lessons after two decades?. Internal and Emergency Medicine, 0, , .	2.0	0