

# Raimondo De Cristofaro

## List of Publications by Year in descending order

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173  
papers

4,692  
citations

109321

35  
h-index

128289

60  
g-index

177  
all docs

177  
docs citations

177  
times ranked

5801  
citing authors

#	ARTICLE	IF	CITATIONS
1	Thrombotic risk factors in patients with liver cirrhosis: Correlation with MELD scoring system and portal vein thrombosis development. <i>Journal of Hepatology</i> , 2009, 51, 682-689.	3.7	409
2	Portal vein thrombosis: Insight into physiopathology, diagnosis, and treatment. <i>World Journal of Gastroenterology</i> , 2010, 16, 143.	3.3	248
3	Platelet Cyclooxygenase Inhibition by Low-Dose Aspirin Is Not Reflected Consistently by Platelet Function Assays. <i>Journal of the American College of Cardiology</i> , 2009, 53, 667-677.	2.8	234
4	Binding of Thrombin to Glycoprotein Ib Accelerates the Hydrolysis of Par-1 on Intact Platelets. <i>Journal of Biological Chemistry</i> , 2001, 276, 4692-4698.	3.4	193
5	The Bovine Basic Pancreatic Trypsin Inhibitor (Kunitz Inhibitor): A Milestone Protein. <i>Current Protein and Peptide Science</i> , 2003, 4, 231-251.	1.4	163
6	High-Mobility Group Box-1 Protein Promotes Angiogenesis After Peripheral Ischemia in Diabetic Mice Through a VEGF-Dependent Mechanism. <i>Diabetes</i> , 2010, 59, 1496-1505.	0.6	110
7	Crucial role of the protein C pathway in governing microvascular inflammation in inflammatory bowel disease. <i>Journal of Clinical Investigation</i> , 2007, 117, 1951-1960.	8.2	105
8	Indoleamine 2,3-dioxygenase-expressing leukemic dendritic cells impair a leukemia-specific immune response by inducing potent T regulatory cells. <i>Haematologica</i> , 2010, 95, 2022-2030.	3.5	95
9	The effect of shear stress on protein conformation. <i>Biophysical Chemistry</i> , 2010, 153, 1-8.	2.8	82
10	Cystic fibrosis transmembrane conductance regulator (CFTR) expression in human platelets: impact on mediators and mechanisms of the inflammatory response. <i>FASEB Journal</i> , 2010, 24, 3970-3980.	0.5	75
11	Congenital Prothrombin Deficiency: An Update. <i>Seminars in Thrombosis and Hemostasis</i> , 2013, 39, 596-606.	2.7	68
12	Lipid and protein oxidation contribute to a prothrombotic state in patients with type 2 diabetes mellitus. <i>Journal of Thrombosis and Haemostasis</i> , 2003, 1, 250-256.	3.8	67
13	The Asp272â€“Glu282 Region of Platelet Glycoprotein Ib $\alpha$ Interacts with the Heparin-binding Site of $\alpha$ -Thrombin and Protects the Enzyme from the Heparin-catalyzed Inhibition by Antithrombin III. <i>Journal of Biological Chemistry</i> , 2000, 275, 3887-3895.	3.4	61
14	Thrombin Domains: Structure, Function and Interaction with Platelet Receptors. <i>Journal of Thrombosis and Thrombolysis</i> , 2003, 15, 151-163.	2.1	56
15	Formation of methionine sulfoxide by peroxynitrite at position 1606 of von Willebrand factor inhibits its cleavage by ADAMTS-13: A new prothrombotic mechanism in diseases associated with oxidative stress. <i>Free Radical Biology and Medicine</i> , 2010, 48, 446-456.	2.9	56
16	Indoleamine 2,3-dioxygenase 1 (IDO1) activity correlates with immune system abnormalities in multiple myeloma. <i>Journal of Translational Medicine</i> , 2012, 10, 247.	4.4	56
17	Hyposialylation of neprilysin possibly affects its expression and enzymatic activity in hereditary inclusionâ€“body myopathy muscle. <i>Journal of Neurochemistry</i> , 2008, 105, 971-981.	3.9	53
18	Congenital Prothrombin Deficiency. <i>Seminars in Thrombosis and Hemostasis</i> , 2009, 35, 367-381.	2.7	53

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19	High mobility group box 1 is a novel substrate of dipeptidyl peptidase-IV. <i>Diabetologia</i> , 2012, 55, 236-244.	6.3	51
20	Haemostatic system in inflammatory bowel diseases: New players in gut inflammation. <i>World Journal of Gastroenterology</i> , 2011, 17, 594.	3.3	50
21	Qualitative and quantitative modifications of von Willebrand factor in patients with essential thrombocythemia and controlled platelet count. <i>Journal of Thrombosis and Haemostasis</i> , 2015, 13, 1226-1237.	3.8	48
22	Structural and Functional Mapping of the Thrombin Domain Involved in the Binding to the Platelet Glycoprotein Ib. <i>Biochemistry</i> , 2001, 40, 13268-13273.	2.5	47
23	Low-grade exercise enhances platelet aggregability in patients with obstructive coronary disease independently of myocardial ischemia. <i>American Journal of Cardiology</i> , 2001, 87, 16-20.	1.6	46
24	Prostaglandin E2 Differentially Modulates Human Platelet Function through the Prostanoid EP2 and EP3 Receptors. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2011, 336, 391-402.	2.5	45
25	Italian intersociety consensus on DOAC use in internal medicine. <i>Internal and Emergency Medicine</i> , 2017, 12, 387-406.	2.0	44
26	Management and 1-Year Outcomes of Patients With Newly Diagnosed Atrial Fibrillation and Chronic Kidney Disease: Results From the Prospective GARFIELD-AF Registry. <i>Journal of the American Heart Association</i> , 2019, 8, e010510.	3.7	44
27	Role of Chloride Ions in Modulation of the Interaction between von Willebrand Factor and ADAMTS-13. <i>Journal of Biological Chemistry</i> , 2005, 280, 23295-23302.	3.4	43
28	Linkage between proton binding and amidase activity in human .alpha.-thrombin: effect of ions and temperature. <i>Biochemistry</i> , 1991, 30, 7913-7924.	2.5	40
29	A Novel Venombin B from <i>Agkistrodon contortrix contortrix</i> : Evidence for Recognition Properties in the Surface around the Primary Specificity Pocket Different from Thrombin. <i>Biochemistry</i> , 2000, 39, 10294-10308.	2.5	40
30	Impaired primary hemostasis with normal platelet function in Duchenne muscular dystrophy during highly-invasive spinal surgery. <i>Neuromuscular Disorders</i> , 2005, 15, 532-540.	0.6	40
31	Presence of portal vein thrombosis in liver cirrhosis is strongly associated with low levels of ADAMTS-13: a pilot study. <i>Internal and Emergency Medicine</i> , 2016, 11, 959-967.	2.0	40
32	Thrombin-Induced Platelet Activation Is Inhibited by High- and Low-Molecular-Weight Heparin. <i>Circulation</i> , 1999, 99, 3308-3314.	1.6	39
33	Mechanisms of the interaction between two ADAMTS13 gene mutations leading to severe deficiency of enzymatic activity. <i>Human Mutation</i> , 2006, 27, 330-336.	2.5	39
34	Fluorinated Benzyloxyphenyl Piperidine-4-carboxamides with Dual Function against Thrombosis: Inhibitors of Factor Xa and Platelet Aggregation. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 1018-1028.	6.4	38
35	Cyclooxygenase-2 (COX-2) Inhibition Constrains Indoleamine 2,3-Dioxygenase 1 (IDO1) Activity in Acute Myeloid Leukaemia Cells. <i>Molecules</i> , 2013, 18, 10132-10145.	3.8	38
36	Carbon monoxide and oxygen binding to human hemoglobin F0. <i>Biochemistry</i> , 1989, 28, 2631-2638.	2.5	36

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37	Inhibitors of indoleamine 2,3-dioxygenase: a review of novel patented lead compounds. <i>Expert Opinion on Therapeutic Patents</i> , 2010, 20, 229-250.	5.0	35
38	Phenomenological analysis of the clotting curve. <i>The Protein Journal</i> , 1991, 10, 455-468.	1.1	34
39	Fibrinogen-elongated $\hat{1}^3$ Chain Inhibits Thrombin-induced Platelet Response, Hindering the Interaction with Different Receptors. <i>Journal of Biological Chemistry</i> , 2008, 283, 30193-30204.	3.4	34
40	The linkage between binding of the C-terminal domain of hirudin and amidase activity in human $\hat{1}^{\pm}$ -thrombin. <i>Biochemical Journal</i> , 1993, 289, 475-480.	3.7	33
41	Glycaemic variability affects ischaemia-induced angiogenesis in diabetic mice. <i>Clinical Science</i> , 2011, 121, 555-564.	4.3	32
42	Mechanochemistry of von Willebrand factor. <i>Biomolecular Concepts</i> , 2019, 10, 194-208.	2.2	32
43	Interaction of the 268-282 region of glycoprotein Ibalpha with the heparin-binding site of thrombin inhibits the enzyme activation of factor VIII. <i>Biochemical Journal</i> , 2003, 373, 593-601.	3.7	30
44	The natural mutation by deletion of Lys9 in the thrombin A-chain affects the pKa value of catalytic residues, the overall enzyme's stability and conformational transitions linked to Na <sup>+</sup> binding. <i>FEBS Journal</i> , 2006, 273, 159-169.	4.7	30
45	Ristocetin-induced self-aggregation of von Willebrand factor. <i>European Biophysics Journal</i> , 2010, 39, 1597-1603.	2.2	30
46	Thrombin Interaction with Platelet GPIB: Role of the Heparin Binding Domain. <i>Thrombosis and Haemostasis</i> , 1997, 77, 735-740.	3.4	30
47	The Oxidative Modification of Von Willebrand Factor Is Associated with Thrombotic Angiopathies in Diabetes Mellitus. <i>PLoS ONE</i> , 2013, 8, e55396.	2.5	30
48	Effect of High- and Low-Molecular-Weight Heparins on Thrombin-Thrombomodulin Interaction and Protein C Activation. <i>Circulation</i> , 1998, 98, 1297-1301.	1.6	29
49	Circulating endothelial progenitor cells and residual in vivo thromboxane biosynthesis in low-dose aspirin-treated polycythemia vera patients. <i>Blood</i> , 2008, 112, 1085-1090.	1.4	29
50	Major adverse cardiovascular events in non-valvular atrial fibrillation with chronic obstructive pulmonary disease: the ARAPACIS study. <i>Internal and Emergency Medicine</i> , 2018, 13, 651-660.	2.0	29
51	Binding of the bovine basic pancreatic trypsin inhibitor (Kunitz) to human $\hat{1}^{\pm}$ -, $\hat{1}^2$ - and $\hat{1}^3$ -thrombin; a kinetic and thermodynamic study. <i>BBA - Proteins and Proteomics</i> , 1988, 956, 156-161.	2.1	28
52	Molecular and functional characterization of a natural homozygous Arg67His mutation in the prothrombin gene of a patient with a severe procoagulant defect contrasting with a mild hemorrhagic phenotype. <i>Blood</i> , 2002, 100, 1347-1353.	1.4	28
53	Defective platelet responsiveness to thrombin and protease-activated receptors agonists in a novel case of gray platelet syndrome: correlation between the platelet defect and the $\hat{1}^{\pm}$ -granule content in the patient and four relatives. <i>Journal of Thrombosis and Haemostasis</i> , 2007, 5, 551-559.	3.8	28
54	Evaluation of assay methods to measure plasma ADAMTS13 activity in thrombotic microangiopathies. <i>Thrombosis and Haemostasis</i> , 2011, 105, 381-385.	3.4	27

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55	Î²2-GlycoproteinÂI binds to thrombin and selectively inhibits the enzyme procoagulant functions. <i>Journal of Thrombosis and Haemostasis</i> , 2013, 11, 1093-1102.	3.8	27
56	Oxidation of Human Î±-Thrombin by the Myeloperoxidase-H2O2-chloride System: Structural and Functional Effects. <i>Thrombosis and Haemostasis</i> , 2000, 83, 253-261.	3.4	26
57	Pain assessment and management in haemophilia: A survey among Italian patients and specialist physicians. <i>Haemophilia</i> , 2018, 24, 766-773.	2.1	26
58	Onâ€pump Cardiac Surgery Enhances Platelet Renewal and Impairs Aspirin Pharmacodynamics: Effects of Improved Dosing Regimens. <i>Clinical Pharmacology and Therapeutics</i> , 2017, 102, 849-858.	4.7	24
59	Meningeal hematopoiesis causing exophthalmus and hemiparesis in myelofibrosis: Effect of radiotherapy: A case report. <i>Cancer</i> , 1988, 62, 2346-2349.	4.1	23
60	Nucleotide-Derived Thrombin Inhibitors: A New Tool for an Old Issue. <i>Cardiovascular and Hematological Agents in Medicinal Chemistry</i> , 2009, 7, 19-28.	1.0	23
61	Molecular characterization of in-frame and out-of-frame alternative splicings in coagulation factor XI pre-mRNA. <i>Blood</i> , 2010, 115, 2065-2072.	1.4	23
62	The dominant-negative von Willebrand factor gene deletion p.P1127_C1948delinsR: molecular mechanism and modulation. <i>Blood</i> , 2010, 116, 5371-5376.	1.4	23
63	Modulation of thrombin-fibrinogen interaction by specific ion effects. <i>Biochemistry</i> , 1992, 31, 257-265.	2.5	22
64	Binding of human Î±-thrombin to platelet Gplb: energetics and functional effects. <i>Biochemical Journal</i> , 1998, 332, 643-650.	3.7	22
65	In vivo and in vitro effects of different anaesthetics on platelet function. <i>British Journal of Haematology</i> , 2004, 125, 79-82.	2.5	22
66	Mechanistic Studies on ADAMTS13 Catalysis. <i>Biophysical Journal</i> , 2008, 95, 2450-2461.	0.5	22
67	The first deletion mutation in the TSP1-6 repeat domain of ADAMTS13 in a family with inherited thrombotic thrombocytopenic purpura. <i>Haematologica</i> , 2009, 94, 289-293.	3.5	22
68	Platelet reactive conformation and multimeric pattern of von Willebrand factor in acquired thrombotic thrombocytopenic purpura during acute disease and remission. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 1744-1751.	3.8	22
69	The expanding spectrum of PRPS1-associated phenotypes: three novel mutations segregating with X-linked hearing loss and mild peripheral neuropathy. <i>European Journal of Human Genetics</i> , 2015, 23, 766-773.	2.8	22
70	Emergency management in patients with haemophilia A and inhibitors on prophylaxis with emicizumab: AICE practical guidance in collaboration with SIBioC, SIMEU, SIMEUP, SIPMeL and Siset. <i>Blood Transfusion</i> , 2020, 18, 143-151.	0.4	22
71	Effect of protons on the amidase activity of human Î±-thrombin. <i>Journal of Molecular Biology</i> , 1990, 216, 1077-1085.	4.2	21
72	Thermodynamics of Substrates and Reversible Inhibitors Binding to the Active Site Cleft of Human Î±-thrombin. <i>Journal of Molecular Biology</i> , 1994, 239, 569-577.	4.2	21

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73	A Natural Prothrombin Mutant Reveals an Unexpected Influence of A-chain Structure on the Activity of Human $\alpha_2$ -Thrombin. <i>Journal of Biological Chemistry</i> , 2004, 279, 13035-13043.	3.4	21
74	Carbamazepine interaction with direct oral anticoagulants: help from the laboratory for the personalized management of oral anticoagulant therapy. <i>Journal of Thrombosis and Thrombolysis</i> , 2019, 48, 528-531.	2.1	21
75	Conformational Transitions Linked to Active Site Ligation in Human Thrombin: Effect on the Interaction with Fibrinogen and the Cleavable Platelet Receptor. <i>Journal of Molecular Biology</i> , 1995, 245, 447-458.	4.2	19
76	Plasma Protein Oxidation Is Associated with an Increase of Procoagulant Markers Causing an Imbalance between Pro- and Anticoagulant Pathways in Healthy Subjects. <i>Thrombosis and Haemostasis</i> , 2002, 87, 58-67.	3.4	19
77	Therapeutic management and costs of severe haemophilia A patients with inhibitors in Italy. <i>Haemophilia</i> , 2014, 20, e243-50.	2.1	19
78	Oxidation of Met1606 in von Willebrand factor is a risk factor for thrombotic and septic complications in chronic renal failure. <i>Biochemical Journal</i> , 2012, 442, 423-432.	3.7	18
79	A phase III study comparing secondary long-term prophylaxis versus on-demand treatment with vWF/FVIII concentrates in severe inherited von Willebrand disease. <i>Blood Transfusion</i> , 2019, 17, 391-398.	0.4	18
80	Molecular Mapping of the Chloride-binding Site in von Willebrand Factor (VWF). <i>Journal of Biological Chemistry</i> , 2006, 281, 30400-30411.	3.4	17
81	Kinetic study of von Willebrand factor self-aggregation induced by ristocetin. <i>Biophysical Chemistry</i> , 2009, 144, 101-107.	2.8	17
82	Oxidized von Willebrand factor is efficiently cleaved by serine proteases from primary granules of leukocytes: divergence from ADAMTS-13. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 1620-1627.	3.8	17
83	Biochemical Properties of Indoleamine 2,3-dioxygenase: From Structure to Optimized Design of Inhibitors. <i>Current Medicinal Chemistry</i> , 2011, 18, 2205-2214.	2.4	17
84	Management of patients with severe haemophilia a without inhibitors on prophylaxis with emicizumab: AICE recommendations with focus on emergency in collaboration with SIBioC, SIMEU, SIMEUP, SIPMeL and Siset. <i>Haemophilia</i> , 2020, 26, 937-945.	2.1	17
85	Function and dysfunction of dendritic cells in autoimmune rheumatic diseases. <i>Human Immunology</i> , 2009, 70, 360-373.	2.4	16
86	Heparin versus prostacyclin in continuous hemodiafiltration for acute renal failure: Effects on platelet function in the systemic circulation and across the filter. <i>Thrombosis Research</i> , 2010, 126, 24-31.	1.7	16
87	Risk profiles and one-year outcomes of patients with newly diagnosed atrial fibrillation in India: Insights from the GARFIELD-AF Registry. <i>Indian Heart Journal</i> , 2018, 70, 828-835.	0.5	16
88	Marked von Willebrand factor and factor VIII elevations in severe acute respiratory syndrome coronavirus-2-positive, but not severe acute respiratory syndrome coronavirus-2-negative, pneumonia: a case-control study. <i>Blood Coagulation and Fibrinolysis</i> , 2021, 32, 285-289.	1.0	16
89	Modulation of thrombin-hirudin interaction by specific ion effects. <i>Journal of Molecular Biology</i> , 1992, 226, 263-269.	4.2	15
90	Effect of Sodium on the Energetics of Thrombin $\alpha_2$ Thrombomodulin Interaction and its Relevance for Protein C Hydrolysis. <i>Journal of Molecular Biology</i> , 1996, 258, 190-200.	4.2	15

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91	Enhanced platelet adhesion induces angiogenesis in intestinal inflammation and inflammatory bowel disease microvasculature. <i>Journal of Cellular and Molecular Medicine</i> , 2011, 15, 625-634.	3.6	15
92	The type 2B p.R1306W natural mutation of von Willebrand factor dramatically enhances the multimer sensitivity to shear stress. <i>Journal of Thrombosis and Haemostasis</i> , 2013, 11, 1688-1698.	3.8	15
93	PROTEOLYTIC PROCESSING OF VON WILLEBRAND FACTOR BY ADAMTS13 AND LEUKOCYTE PROTEASES. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , 2013, 5, e2013058.	1.3	15
94	Perceived challenges and attitudes to regimen and product selection from Italian haemophilia treaters: the 2013 AICE survey. <i>Haemophilia</i> , 2014, 20, e128-35.	2.1	15
95	Molecular Dynamics Characterization of Five Pathogenic Factor X Mutants Associated with Decreased Catalytic Activity. <i>Biochemistry</i> , 2014, 53, 6992-7001.	2.5	15
96	Temperature- and pH-dependence of the oxygen-binding reaction of human fetal haemoglobin. <i>Biochemical Journal</i> , 1989, 260, 617-619.	3.7	14
97	Effect of fibrinogen concentration and platelet count on the inhibitory effect of abciximab and tirofiban. <i>Thrombosis and Haemostasis</i> , 2003, 89, 348-354.	3.4	14
98	Homocysteinemia is inversely correlated with platelet count and directly correlated with sE- and sP-selectin levels in females homozygous for C677T methylenetetrahydrofolate reductase. <i>Platelets</i> , 2005, 16, 185-190.	2.3	14
99	Effects of pegylated G-CSF on immune cell number and function in patients with gynecological malignancies. <i>Journal of Translational Medicine</i> , 2010, 8, 114.	4.4	14
100	The D173G mutation in ADAMTS-13 causes a severe form of congenital thrombotic thrombocytopenic purpura. <i>Thrombosis and Haemostasis</i> , 2016, 115, 51-62.	3.4	14
101	Human platelet glycoprotein IIb/IIIa purification by phenyl boronate affinity chromatography coupled to anion-exchange high-performance liquid chromatography. <i>Biomedical Applications</i> , 1988, 426, 376-380.	1.7	13
102	Allosteric equilibria in the binding of fibrinogen to platelets. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1988, 85, 8473-8476.	7.1	13
103	Allosteric modulation of BPTI interaction with human alpha- and zeta-thrombin. <i>FEBS Journal</i> , 1999, 260, 97-102.	0.2	13
104	Comparison of international normalized ratio audit parameters in patients enrolled in GARFIELD and treated with vitamin K antagonists. <i>British Journal of Haematology</i> , 2016, 174, 610-623.	2.5	13
105	Effect of temperature on the association step in thrombin-fibrinogen interaction. <i>Biochemical Journal</i> , 1993, 294, 563-567.	3.7	12
106	Structure and Proteolytic Properties of ADAMTS13, A Metalloprotease Involved in the Pathogenesis of Thrombotic Microangiopathies. <i>Progress in Molecular Biology and Translational Science</i> , 2011, 99, 105-144.	1.7	12
107	Effects of protons on the thrombin-fibrinogen interaction. <i>FEBS Journal</i> , 1994, 219, 1013-1021.	0.2	11
108	Inherited Macrothrombocytopenia with Distinctive Platelet Ultrastructural and Functional Features. <i>Thrombosis and Haemostasis</i> , 2000, 83, 35-41.	3.4	11

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109	ADAMTS-13/von Willebrand factor ratio: A prognostic biomarker for portal vein thrombosis in compensated cirrhosis. A prospective observational study. <i>Digestive and Liver Disease</i> , 2022, 54, 1672-1680.	0.9	11
110	High-performance liquid chromatography in protein sequence determinations. <i>Journal of Chromatography A</i> , 1988, 440, 231-251.	3.7	10
111	Linkage between proton binding and amidase activity in human $\gamma$ -thrombin. <i>Biochemistry</i> , 1992, 31, 1147-1153.	2.5	10
112	Duodenal and gastric Dieulafoy's lesions in a patient with type 2A von Willebrand's disease. <i>Gastrointestinal Endoscopy</i> , 2005, 61, 322-325.	1.0	10
113	Inhibition of fibrinogen binding to human platelets by blockage of Na <sup>+</sup> /H <sup>+</sup> exchange. <i>Biochemical and Biophysical Research Communications</i> , 1989, 161, 1228-1232.	2.1	9
114	Increased von Willebrand factor levels in polycythemia vera and phenotypic differences with essential thrombocythemia. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2020, 4, 413-421.	2.3	9
115	The linkage between adenosine nucleotide binding and amidase activity in human $\alpha$ -thrombin. <i>Biophysical Chemistry</i> , 1990, 36, 77-84.	2.8	8
116	Prothrombotic response to coronary angioplasty in patients with unstable angina and raised C-reactive protein. <i>Journal of Thrombosis and Thrombolysis</i> , 2002, 14, 131-138.	2.1	8
117	A recurrent Gly43Asp substitution in coagulation Factor X rigidifies its catalytic pocket and impairs catalytic activity and intracellular trafficking. <i>Thrombosis Research</i> , 2014, 133, 481-487.	1.7	8
118	Polypharmacy in the elderly: A population based cross-sectional study in Lazio, Italy. <i>European Geriatric Medicine</i> , 2016, 7, 484-487.	2.8	8
119	Diagnostic testing for differential diagnosis in Thrombotic Microangiopathies. <i>Turkish Journal of Haematology</i> , 2019, 36, 222-229.	0.5	8
120	Modeling ADAMTS13-von Willebrand Factor interaction: Implications for oxidative stress-related cardiovascular diseases and type 2A von Willebrand Disease. <i>Biophysical Chemistry</i> , 2012, 160, 1-11.	2.8	7
121	Diagnosis and management of cerebral venous sinus thrombosis in children: a single-center retrospective analysis. <i>Child's Nervous System</i> , 2021, 37, 153-160.	1.1	7
122	Does chronic oral anticoagulation reduce in-hospital mortality among COVID-19 older patients?. <i>Aging Clinical and Experimental Research</i> , 2021, 33, 2335-2343.	2.9	7
123	Thrombin-thrombomodulin interaction: energetics and potential role of water as an allosteric effector. <i>Biochemical Journal</i> , 1995, 310, 49-53.	3.7	6
124	Portal vein thrombosis occurrence in a cirrhotic patient during treatment with rivaroxaban. <i>Liver International</i> , 2017, 37, 1251-1251.	3.9	6
125	Noncanonical type 2B von Willebrand disease associated with mutations in the VWF D $\alpha$ 2D3 and D4 domains. <i>Blood Advances</i> , 2020, 4, 3405-3415.	5.2	6
126	Thrombin interaction with platelet Gplb: structural mapping and effects on platelet activation (review).. <i>International Journal of Molecular Medicine</i> , 1999, 3, 363-71.	4.0	5



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127	The P303T mutation in the human factor VII (FVII) gene alters the conformational state of the enzyme and causes a severe functional deficiency. <i>British Journal of Haematology</i> , 2004, 127, 576-584.	2.5	5
128	Sustained safe and effective anticoagulation using Edoxaban via percutaneous endoscopic gastrostomy. <i>ESC Heart Failure</i> , 2019, 6, 884-888.	3.1	5
129	Indoleamine 2,3-Dioxygenase (IDO) Is Expressed by Multiple Myeloma Plasma Cells and Promotes the Differentiation of Regulatory T Cells: Investigations into the Role of Hepatocyte Growth Factor.. <i>Blood</i> , 2008, 112, 1680-1680.	1.4	5
130	COX-2 Inhibition Suppresses the Interferon- $\gamma$ -Induced Expression of Indoleamine 2,3-Dioxygenase (IDO) in Human Leukemia Cell Lines.. <i>Blood</i> , 2008, 112, 1623-1623.	1.4	5
131	Effect of fibrinogen concentration and platelet count on the inhibitory effect of abciximab and tirofiban. <i>Thrombosis and Haemostasis</i> , 2003, 89, 348-54.	3.4	5
132	Two Naturally Occurring Mutations on FVII Gene (S363I-W364C) Altering Intrinsic Catalytic Activity. <i>Thrombosis and Haemostasis</i> , 2002, 88, 750-755.	3.4	4
133	Thrombin-mediated impairment of fibroblast growth factor- $\beta$ activity. <i>FEBS Journal</i> , 2009, 276, 3277-3289.	4.7	4
134	Apixaban Interacts with Haemoglobin: Effects on Its Plasma Levels. <i>Thrombosis and Haemostasis</i> , 2018, 118, 1701-1712.	3.4	4
135	Molecular Aggregation of Marketed Recombinant FVIII Products: Biochemical Evidence and Functional Effects. <i>TH Open</i> , 2019, 03, e123-e131.	1.4	4
136	A novel factor- $\beta$ missense mutation (Val371Ile) in the activation loop is responsible for a case of mild type- $\beta$ factor- $\beta$ deficiency. <i>FEBS Journal</i> , 2007, 274, 6128-6138.	4.7	3
137	Response: Further thoughts on the "phantom" $\beta$ 6/7 FXI isoform. <i>Blood</i> , 2010, 116, 1186-1187.	1.4	3
138	ORal anticoagulants In fraGile patients with percutaneous endoscopic gastrostomy and atrial fibrillation: the (ORIGAMI) study. <i>Journal of Cardiovascular Medicine</i> , 2021, 22, 175-179.	1.5	3
139	Current Choices and Management of Treatment in Persons with Severe Hemophilia A without Inhibitors: A Mini-Delphi Consensus. <i>Journal of Clinical Medicine</i> , 2022, 11, 801.	2.4	3
140	Detection of Platelet-Activating Antibodies Associated with Vaccine-Induced Thrombotic Thrombocytopenia by Flow Cytometry: An Italian Experience. <i>Viruses</i> , 2022, 14, 1133.	3.3	3
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