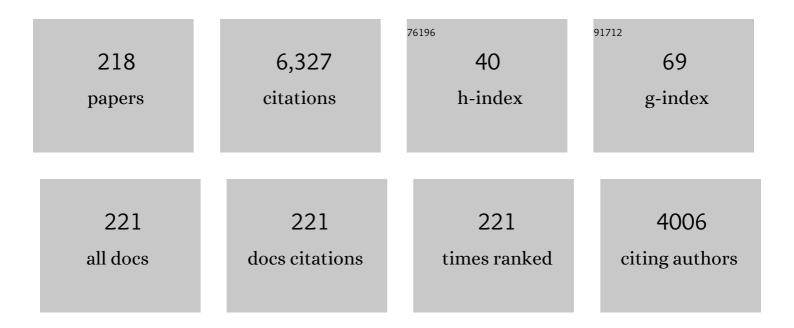
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	An international collaborative study to assign value for Total Factor XIIIâ€B Subunit Antigen to the WHO 1st International Standard for Factor XIII Plasma, (02/206): Communication from the ISTH SSC Subcommittee on Factor XIII and Fibrinogen. Journal of Thrombosis and Haemostasis, 2022, 20, 525-531.	1.9	2
2	Activation mechanism dependent surface exposure of cellular factor XIII on activated platelets and platelet microparticles. Journal of Thrombosis and Haemostasis, 2022, 20, 1223-1235.	1.9	14
3	The Effect of Activated FXIII, a Transglutaminase, on Vascular Smooth Muscle Cells. International Journal of Molecular Sciences, 2022, 23, 5845.	1.8	0
4	Genetic Background of Inherited Factor XIII-A Subunit Deficiency: Review of the Literature and Description of Two New Cases. Seminars in Thrombosis and Hemostasis, 2021, 47, 885-889.	1.5	1
5	Effect of α2-plasmin inhibitor heterogeneity on the risk of venous thromboembolism. Thrombosis Research, 2021, 203, 110-116.	0.8	2
6	Terminal Phase Components of the Clotting Cascade in Patients with End-Stage Renal Disease Undergoing Hemodiafiltration or Hemodialysis Treatment. International Journal of Molecular Sciences, 2020, 21, 8426.	1.8	5
7	Nâ€glycosylation of blood coagulation factor XIII subunit B and its functional consequence. Journal of Thrombosis and Haemostasis, 2020, 18, 1302-1309.	1.9	7
8	The factor XIIIâ€A Val34Leu polymorphism decreases whole blood clot mass at high fibrinogen concentrations. Journal of Thrombosis and Haemostasis, 2020, 18, 885-894.	1.9	8
9	Autoimmune factor XIII deficiency with unusual laboratory and clinical phenotype. Journal of Thrombosis and Haemostasis, 2020, 18, 1330-1334.	1.9	2
10	A new ELISA method for the measurement of total α2-plasmin inhibitor level in human body fluids. Journal of Immunological Methods, 2019, 471, 27-33.	0.6	6
11	FXIII levels in patients with systemic lupus erythematosus and antiphospholipid syndrome. Clinica Chimica Acta, 2019, 493, S416-S417.	0.5	Ο
12	First Step of the Transglutaminase Reaction Catalyzed by Activated Factor XIII Subunit A, Hybrid Quantum Chemistry/Molecular Mechanics Calculations. Journal of Physical Chemistry B, 2019, 123, 3887-3897.	1.2	2
13	Factor XIII: What does it look like?. Journal of Thrombosis and Haemostasis, 2019, 17, 714-716.	1.9	7
14	Cellular Factor XIII, a Transglutaminase in Human Corneal Keratocytes. International Journal of Molecular Sciences, 2019, 20, 5963.	1.8	6
15	PB2431 CAROTID INTIMAâ€MEDIA THICKNESS PROGRESSION AND BLOOD COAGULATION FACTOR XIII LEVELS IN TYPE 2 DIABETES: 8 YEARS OF FOLLOWâ€UP. HemaSphere, 2019, 3, 1077.	1.2	Ο
16	Auto―and alloantibodies against factor XIII: laboratory diagnosis and clinical consequences. Journal of Thrombosis and Haemostasis, 2018, 16, 822-832.	1.9	18
17	Effect of factor XIII levels and polymorphisms on the risk of myocardial infarction in young patients. Molecular and Cellular Biochemistry, 2018, 448, 199-209.	1.4	11
18	Evaluation of endogenous thrombin potential among patients with antithrombin deficiency. Thrombosis Research, 2018, 166, 50-53.	0.8	4

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19	Low factor XIII levels after intravenous thrombolysis predict short-term mortality in ischemic stroke patients. Scientific Reports, 2018, 8, 7662.	1.6	17
20	Early onset of abdominal venous thrombosis in a newborn with homozygous type II heparin-binding site antithrombin deficiency. Blood Coagulation and Fibrinolysis, 2017, 28, 264-266.	0.5	6
21	Evaluation of flow cytometric <scp>HIT</scp> assays in relation to an <scp>I</scp> g <scp>G</scp> â€5pecific immunoassay and clinical outcome. Cytometry Part B - Clinical Cytometry, 2017, 92, 389-397.	0.7	7
22	Severe bleeding diatheses in an elderly patient with combined type autoantibody against factor <scp>XIII</scp> A subunit; novel approach to the diagnosis and classification of antiâ€factor XIII antibodies. Haemophilia, 2017, 23, 590-597.	1.0	4
23	Factor XIII levels and factor XIII B subunit polymorphisms in patients with venous thromboembolism. Thrombosis Research, 2017, 158, 93-97.	0.8	11
24	Minimal factor XIII activity level to prevent major spontaneous bleeds. Journal of Thrombosis and Haemostasis, 2017, 15, 1728-1736.	1.9	34
25	Assessment of Factor XIII. Methods in Molecular Biology, 2017, 1646, 277-293.	0.4	10
26	Clinical and laboratory characteristics of antithrombin deficiencies: A large cohort study from a single diagnostic center. Thrombosis Research, 2017, 160, 119-128.	0.8	16
27	Intracardiac Hemostasis and Fibrinolysis Parameters in Patients with Atrial Fibrillation. BioMed Research International, 2017, 2017, 1-10.	0.9	17
28	Antithrombin Debrecen (p.Leu205Pro) – Clinical and molecular characterization of a novel mutation associated with severe thrombotic tendency. Thrombosis Research, 2017, 158, 1-7.	0.8	4
29	Alloantibody developed in a factor XIII A subunit deficient patient during substitution therapy; characterization of the antibody. Haemophilia, 2016, 22, 268-275.	1.0	15
30	Founder effect is responsible for the p.Leu131Phe heparinâ€bindingâ€site antithrombin mutation common in Hungary: phenotype analysis in a large cohort. Journal of Thrombosis and Haemostasis, 2016, 14, 704-715.	1.9	38
31	Diagnosis and Management of Congenital and Acquired FXIII Deficiencies. Seminars in Thrombosis and Hemostasis, 2016, 42, 429-439.	1.5	66
32	Protein cross-linking by chlorinated polyamines and transglutamylation stabilizes neutrophil extracellular traps. Cell Death and Disease, 2016, 7, e2332-e2332.	2.7	24
33	Regulation of plasma factor XIII levels in healthy individuals; a major impact by subunit B intron K c.1952+144 C>G polymorphism. Thrombosis Research, 2016, 148, 101-106.	0.8	8
34	The lack of aspirin resistance in patients with coronary artery disease. Journal of Translational Medicine, 2016, 14, 74.	1.8	9
35	Neutralizing autoantibody against factor XIII A subunit resulted in severe bleeding diathesis with a fatal outcome – characterization of the antibody. Journal of Thrombosis and Haemostasis, 2016, 14, 1517-1520.	1.9	5
36	Pregnancy related stroke in the setting of homozygous type-II HBS antithrombin deficiency. Thrombosis Research, 2016, 139, 111-113.	0.8	6

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37	How to test the effect of aspirin and clopidogrel in patients on dual antiplatelet therapy?. Platelets, 2016, 27, 59-65.	1.1	16
38	Plasma clot properties in patients with a mild-to-moderate bleeding tendency of unknown cause. Annals of Hematology, 2015, 94, 1301-1310.	0.8	11
39	The use of recombinant factor <scp>XIII</scp> in a major bleeding episode of a patient with congenital factor <scp>XIII</scp> deficiency – the first experience. Haemophilia, 2015, 21, e118-21.	1.0	5
40	Factor XIII B Subunit Polymorphisms and the Risk of Coronary Artery Disease. International Journal of Molecular Sciences, 2015, 16, 1143-1159.	1.8	17
41	Molecular characterization of p.Asp77Gly and the novel p.Ala163Val and p.Ala163Glu mutations causing protein C deficiency. Thrombosis Research, 2015, 135, 718-726.	0.8	13
42	Interaction of factor XIII subunits. Blood, 2014, 123, 1757-1763.	0.6	55
43	Factor XIII deficiency: complete phenotypic characterization of two cases with novel causative mutations. Haemophilia, 2014, 20, 114-120.	1.0	5
44	Progressive chromogenic anti-factor Xa assay and its use in the classification of antithrombin deficiencies. Clinical Chemistry and Laboratory Medicine, 2014, 52, 1797-806.	1.4	8
45	Evaluation of laboratory methods routinely used to detect the effect of aspirin against new reference methods. Thrombosis Research, 2014, 133, 811-816.	0.8	19
46	Poor pregnancy outcome in women with homozygous type-II HBS antithrombin deficiency. Thrombosis Research, 2014, 133, 1158-1160.	0.8	11
47	The Use of Recombinant FXIII in a Major Acute Bleeding Episode of a Patient with Congenital FXIII Deficiency – the First Experience. Blood, 2014, 124, 5065-5065.	0.6	Ο
48	New direct and indirect methods for the detection of cyclooxygenase 1 acetylation by aspirin; the lack of aspirin resistance among healthy individuals. Thrombosis Research, 2013, 131, 320-324.	0.8	25
49	Mechanism of the irreversible inhibition of human cyclooxygenase-1 by aspirin as predicted by QM/MM calculations. Journal of Molecular Graphics and Modelling, 2013, 40, 99-109.	1.3	48
50	Validation of Reference Genes for the Determination of Platelet Transcript Level in Healthy Individuals and in Patients with the History of Myocardial Infarction. International Journal of Molecular Sciences, 2013, 14, 3456-3466.	1.8	19
51	The Superiority of Anti-FXa Assay Over Anti-Flla Assay in Detecting Heparin-Binding Site Antithrombin Deficiency. American Journal of Clinical Pathology, 2013, 140, 675-679.	0.4	26
52	Comparison of a New P2Y12 Receptor Specific Platelet Aggregation Test with Other Laboratory Methods in Stroke Patients on Clopidogrel Monotherapy. PLoS ONE, 2013, 8, e69417.	1.1	19
53	Measurement of factor XIII activity in plasma. Clinical Chemistry and Laboratory Medicine, 2012, 50, 1191-1202.	1.4	33
54	Thrombomodulin-dependent effect of factor VLeiden mutation on factor XIII activation. Thrombosis Research, 2012, 129, 508-513.	0.8	4

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55	Interaction between homocysteine and lipoprotein(a) increases the prevalence of coronary artery disease/myocardial infarction in women: A case-control study. Thrombosis Research, 2012, 129, 133-138.	0.8	21
56	Factor XIII, clot structure, thrombosis. Thrombosis Research, 2012, 129, 382-387.	0.8	107
57	Factor XIII and inflammatory cells. Thrombosis Research, 2012, 129, S77-S81.	0.8	42
58	Thrombomodulin-dependent effect of factor V Leiden mutation on the cross-linking of $\hat{1}\pm 2$ -plasmin inhibitor to fibrin and its consequences on fibrinolysis. Thrombosis Research, 2012, 130, 528-534.	0.8	8
59	Expression of coagulation factor XIII subunit A in acute promyelocytic leukemia. Cytometry Part B - Clinical Cytometry, 2012, 82B, 209-216.	0.7	14
60	Factor XIII: A Coagulation Factor With Multiple Plasmatic and Cellular Functions. Physiological Reviews, 2011, 91, 931-972.	13.1	384
61	Factor XIII subunits in human tears; their highly elevated levels following penetrating keratoplasty. Clinica Chimica Acta, 2011, 412, 271-276.	0.5	10
62	Novel aspects of factor XIII deficiency. Current Opinion in Hematology, 2011, 18, 366-372.	1.2	67
63	Factor XIII: novel structural and functional aspects. Journal of Thrombosis and Haemostasis, 2011, 9, 9-20.	1.9	146
64	Diagnosis and classification of factorÂXIII deficiencies. Journal of Thrombosis and Haemostasis, 2011, 9, 1404-1406.	1.9	157
65	Factor XIII and Venous Thromboembolism. Seminars in Thrombosis and Hemostasis, 2011, 37, 305-314.	1.5	34
66	A highly sensitive chemiluminescence immunoassay for the measurement of coagulation factor XIII subunits and their complex in tears. Journal of Immunological Methods, 2010, 353, 87-92.	0.6	9
67	Antithrombin deficiency and its laboratory diagnosis. Clinical Chemistry and Laboratory Medicine, 2010, 48, S67-78.	1.4	29
68	Clinical and immunoserological characteristics of the transition from primary to overlap antiphospholipid syndrome. Lupus, 2010, 19, 1520-1526.	0.8	9
69	Factor XIII and Atherothrombotic Diseases. Seminars in Thrombosis and Hemostasis, 2010, 36, 018-033.	1.5	36
70	Variant type Glanzmann thrombasthenia caused by homozygous p.724R>X mutation in β3 integrin. Thrombosis Research, 2010, 125, 427-431.	0.8	0
71	Factor XIII A subunit Val34Leu polymorphism in patients suffering atherothrombotic ischemic stroke. Thrombosis Research, 2010, 126, 159-162.	0.8	14
72	Inherited factor V deficiency associated with a novel heterozygous missense mutation (p.G493R) in a patient with excessive surgical bleeding. Thrombosis and Haemostasis, 2009, 102, 787-789.	1.8	2

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73	Factor XIII Deficiency. Seminars in Thrombosis and Hemostasis, 2009, 35, 426-438.	1.5	197
74	Assessment of thrombotic risk factors predisposing to thromboembolic complications in prosthetic orthopedic surgery. Journal of Orthopaedic Science, 2009, 14, 484-490.	0.5	11
75	Molecular mechanism of the interaction between activated factorÂXIII and its glutamine donor peptide substrate. Journal of Thrombosis and Haemostasis, 2009, 7, 627-633.	1.9	17
76	Binding of plasma factor XIII to thrombin-receptor activated human platelets. Thrombosis and Haemostasis, 2009, 102, 83-89.	1.8	23
77	Severe bleeding complications caused by an autoantibody against the B subunit of plasma factor XIII: a novel form of acquired factor XIII deficiency. Blood, 2009, 113, 723-725.	0.6	69
78	4-Thio-deoxyuridylate-modified thrombin aptamer and its inhibitory effect on fibrin clot formation, platelet aggregation and thrombus growth on subendothelial matrix. Journal of Thrombosis and Haemostasis, 2008, 6, 1764-1771.	1.9	43
79	Decreased factor XIII levels in factor XIII A subunit Leu34 homozygous patients with coronary artery disease. Thrombosis Research, 2008, 121, 469-476.	0.8	14
80	Elevated factor XIII level and the risk of peripheral artery disease. Haematologica, 2008, 93, 1430-1432.	1.7	24
81	Factor XDebrecen: Gly204Arg mutation in factor X causes the synthesis of a non-secretable protein and severe factor X deficiency. Haematologica, 2008, 93, 299-302.	1.7	12
82	The Involvement of Blood Coagulation Factor XIII in Fibrinolysis and Thrombosis. Cardiovascular and Hematological Agents in Medicinal Chemistry, 2008, 6, 190-205.	0.4	110
83	Cleavage of factor XIII by human neutrophil elastase results in a novel active truncated form of factor XIII A subunit. Thrombosis and Haemostasis, 2008, 99, 668-674.	1.8	20
84	Endothelial dysfunction and atherosclerosis in rheumatoid arthritis: a multiparametric analysis using imaging techniques and laboratory markers of inflammation and autoimmunity. Journal of Rheumatology, 2008, 35, 398-406.	1.0	130
85	Elevated factor XIII level and the risk of myocardial infarction in women. Haematologica, 2007, 92, 287-288.	1.7	34
86	Three novel mutations in the glycoprotein IIb gene in a patient with type II Glanzmann thrombasthenia. Haematologica, 2007, 92, 698-701.	1.7	9
87	Modulation of the risk of coronary sclerosis/myocardial infarction by the interaction between factor XIII subunit A Val34Leu polymorphism and fibrinogen concentration in the high risk Hungarian population. Thrombosis Research, 2007, 120, 567-573.	0.8	40
88	International Registry on Factor XIII Deficiency: A basis formed mostly on European data. Thrombosis and Haemostasis, 2007, 97, 914-921.	1.8	129
89	Down-regulation of activated factor XIII by polymorphonuclear granulocyte proteases within fibrin clot. Thrombosis and Haemostasis, 2007, 98, 359-367.	1.8	26
90	Factor XIII: recommended terms and abbreviations1. Journal of Thrombosis and Haemostasis, 2007, 5, 181-183.	1.9	39

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91	A collaborative study to establish the 1st International Standard for factor XIII plasma. Journal of Thrombosis and Haemostasis, 2007, 5, 1923-1929.	1.9	17
92	Factor XIII Val34Leu variant protects against coronary artery disease. Thrombosis and Haemostasis, 2007, 97, 458-463.	1.8	40
93	Establishment of an International Registry of Patients with Inherited FXIII Deficiency Blood, 2007, 110, 2149-2149.	0.6	0
94	Factor XIII Val34Leu variant protects against coronary artery disease. A meta-analysis. Thrombosis and Haemostasis, 2007, 97, 458-63.	1.8	17
95	International registry on factor XIII deficiency: a basis formed mostly on European data. Thrombosis and Haemostasis, 2007, 97, 914-21.	1.8	33
96	Coincidence of mutations in different connexin genes in Hungarian patients. International Journal of Molecular Medicine, 2007, 20, 315-21.	1.8	25
97	Down-regulation of activated factor XIII by polymorphonuclear granulocyte proteases within fibrin clot. Thrombosis and Haemostasis, 2007, 98, 359-67.	1.8	8
98	A new mutation in the human pres gene and its effect on prestin function. International Journal of Molecular Medicine, 2007, 20, 545-50.	1.8	17
99	Leukemic lymphoblasts, a novel expression site of coagulation factor XIII subunit A. Thrombosis and Haemostasis, 2006, 96, 176-182.	1.8	24
100	The combined effect of fibrin formation and factor XIII A subunit Val34Leu polymorphism on the activation of factor XIII in whole plasma. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2006, 1764, 1420-1423.	1.1	25
101	Von Willebrand Factor Antigen Latex Immunoassays are Affected to a Different Extent by Rheumatoid Factor. Clinical and Applied Thrombosis/Hemostasis, 2006, 12, 242-243.	0.7	4
102	Leukemic lymphoblasts, a novel expression site of coagulation factor XIII subunit A. Thrombosis and Haemostasis, 2006, 96, 176-82.	1.8	13
103	High-throughput scintillation proximity assay for transglutaminase activity measurement. Analytical Biochemistry, 2005, 343, 256-262.	1.1	15
104	Factor XIII in bronchoalveolar lavage fluid from children with chronic bronchoalveolar inflammation. Journal of Thrombosis and Haemostasis, 2005, 3, 1407-1413.	1.9	18
105	Coagulation factor XIII-A. A flow cytometric intracellular marker in the classification of acute myeloid leukemias. Thrombosis and Haemostasis, 2005, 94, 454-9.	1.8	22
106	Impaired wound healing in factor XIII deficient mice. Thrombosis and Haemostasis, 2005, 94, 432-7.	1.8	102
107	A novel homozygous mutation (1619delC) in GPIIb gene associated with Glanzmann thrombasthenia, the decay of GPIIb-mRNA and the synthesis of a truncated GPIIb unable to form complex with GPIIIa. Thrombosis and Haemostasis, 2005, 93, 904-909.	1.8	7
108	Monitoring platelet function by PFA-100 closure time measurements during thrombolytic therapy of patients with myocardial infarction. Thrombosis Research, 2005, 116, 139-144.	0.8	0

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109	Rapid detection of the factor XIII Val34Leu (163 G→T) polymorphism by real-time PCR using fluorescence resonance energy transfer detection and melting curve analysis. Clinical Chemistry and Laboratory Medicine, 2004, 42, 877-9.	1.4	25
110	Prophylactic and perioperative replacement therapy for acquired factor XIII deficiency: a rebuttal. Journal of Thrombosis and Haemostasis, 2004, 2, 2075-2077.	1.9	43
111	Possible role of factor XIII subunit A in FcÎ ³ and complement receptor-mediated phagocytosis. Cellular Immunology, 2004, 228, 81-90.	1.4	56
112	Phenotype-genotype characterization of 10 families with severe a subunit factor XIII deficiency. Human Mutation, 2004, 23, 98-98.	1.1	45
113	GJB2mutations in patients with non-syndromic hearing loss from Northeastern Hungary. Human Mutation, 2004, 23, 631-632.	1.1	29
114	Possible role of factor XIII subunit A in Fc\$gamma; and complement receptor-mediated phagocytosis. Cellular Immunology, 2004, 228, 81-81.	1.4	1
115	Platelets but not monocytes contribute to the plasma levels of factor XIII subunit A in patients undergoing autologous peripheral blood stem cell transplantation. Blood Coagulation and Fibrinolysis, 2004, 15, 249-253.	0.5	25
116	Anti-factor V auto-antibody in the plasma and platelets of a patient with repeated gastrointestinal bleeding. Journal of Thrombosis and Haemostasis, 2003, 1, 943-949.	1.9	34
117	Evaluation of clinical and laboratory features of antiphospholipid syndrome: a retrospective study of 637 patients. Lupus, 2003, 12, 302-307.	0.8	61
118	Coagulation Factor Deficiencies and Pregnancy Loss. Seminars in Thrombosis and Hemostasis, 2003, 29, 171-174.	1.5	92
119	The mystery of FXIII-A Val34Leu polymorphism. Blood, 2003, 101, 2904-2904.	0.6	0
120	Severe coagulation factor V deficiency caused by 2 novel frameshift mutations: 2952delT in exon 13 and 5493insG in exon 16 of factor 5 gene. Blood, 2002, 99, 702-705.	0.6	34
121	Organization of the glycoprotein (GP) IIb/IIIa heterodimer on resting human platelets studied by flow cytometric energy transfer. Journal of Photochemistry and Photobiology B: Biology, 2001, 65, 47-58.	1.7	6
122	Enzyme-linked immunosorbent assay for the determination of blood coagulation factor XIII A-subunit in plasma and in cell lysates. Journal of Immunological Methods, 2001, 258, 127-135.	0.6	67
123	Calcium Binding of Transglutaminases: A ⁴³ Ca NMR Study Combined with Surface Polarity Analysis. Journal of Biomolecular Structure and Dynamics, 2001, 19, 59-74.	2.0	32
124	A Simple, Quick One-step ELISA Assay for the Determination of Complex Plasma Factor XIII (A2B2). Thrombosis and Haemostasis, 2000, 83, 268-273.	1.8	88
125	Effect of Val34Leu Polymorphism on the Activation of the Coagulation Factor XIII-A. Thrombosis and Haemostasis, 2000, 84, 595-600.	1.8	92
126	Deficiency Causing Mutations and Common Polymorphisms in the Factor XIII-A Gene. Thrombosis and Haemostasis, 2000, 84, 524-527.	1.8	36

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127	α2-Plasmin Inhibitor is a Substrate for Tissue Transglutaminase. Thrombosis Research, 2000, 99, 399-406.	0.8	16
128	A simple, quick one-step ELISA assay for the determination of complex plasma factor XIII (A2B2). Thrombosis and Haemostasis, 2000, 83, 268-73.	1.8	18
129	Comparison of the O'Brien filter test and the PFA-100 platelet analyzer in the laboratory diagnosis of von Willebrand's disease. Thrombosis and Haemostasis, 2000, 84, 88-92.	1.8	3
130	Val34Leu polymorphism of plasma factor XIII: biochemistry and epidemiology in familial thrombophilia. Blood, 2000, 96, 2479-86.	0.6	23
131	Deficiency causing mutations and common polymorphisms in the factor XIII-A gene. Thrombosis and Haemostasis, 2000, 84, 524-7.	1.8	9
132	Effect of Val34Leu polymorphism on the activation of the coagulation factor XIII-A. Thrombosis and Haemostasis, 2000, 84, 595-600.	1.8	30
133	A modified, optimized kinetic photometric assay for the determination of blood coagulation factor XIII activity in plasma. Clinical Chemistry, 2000, 46, 1946-55.	1.5	20
134	The frequency of the haemochromatosis C282Y mutation in the ethnic Hungarian and Romany populations of eastern Hungary. British Journal of Haematology, 1999, 107, 464-465.	1.2	12
135	Blood Coagulation Factor XIII. Thrombosis Research, 1999, 94, 271-305.	0.8	288
136	Comparison of PFA-100 Closure Time and Template Bleeding Time of Patients with Inherited Disorders Causing Defective Platelet Function. Thrombosis Research, 1999, 96, 487-492.	0.8	52
137	High Prevalence of Factor V Leiden Mutation and 20210A Prothrombin Variant in Hungary. Thrombosis and Haemostasis, 1999, 81, 660-661.	1.8	20
138	High prevalence of factor V Leiden mutation and 20210A prothrombin variant in Hungary. Thrombosis and Haemostasis, 1999, 81, 660-1.	1.8	1
139	High frequency of factor V Leiden mutation and prothrombin 20210A variant in Romanies of Eastern Hungary. Thrombosis and Haemostasis, 1999, 82, 1555-6.	1.8	6
140	A microassay to assess the oxidative resistance of low-density lipoproteins. Clinical Chemistry, 1998, 44, 1762-4.	1.5	4
141	Incompatibility of a recombinant thromboplastin and an APTT reagent on a fully automated coagulation analyzer. Blood Coagulation and Fibrinolysis, 1997, 8, 251-252.	0.5	0
142	Molecular Mechanism of a Mild Phenotype in Coagulation Factor XIII (FXIII) Deficiency: A Splicing Mutation Permitting Partial Correct Splicing of FXIII A-Subunit mRNA. Blood, 1997, 89, 1279-1287.	0.6	44
143	Molecular Mechanisms of Mutations in Factor XIII A-subunit Deficiency: In vitro Expression in COS-cells Demonstrates Intracellular Degradation of the Mutant Proteins. Thrombosis and Haemostasis, 1997, 77, 1068-1072.	1.8	24
144	Molecular mechanism of a mild phenotype in coagulation factor XIII (FXIII) deficiency: a splicing mutation permitting partial correct splicing of FXIII A-subunit mRNA. Blood, 1997, 89, 1279-87.	0.6	7

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145	Molecular mechanisms of mutations in factor XIII A-subunit deficiency: in vitro expression in COS-cells demonstrates intracellular degradation of the mutant proteins. Thrombosis and Haemostasis, 1997, 77, 1068-72.	1.8	4
146	Novel Aspects of Blood Coagulation Factor XIII. I. Structure, Distribution, Activation, and Function. Critical Reviews in Clinical Laboratory Sciences, 1996, 33, 357-421.	2.7	182
147	ETRO Working Party on Factor XIII Questionnaire on Congenital Factor XIII Deficiency in Europe: Status and Perspectives. Seminars in Thrombosis and Hemostasis, 1996, 22, 415-418.	1.5	70
148	Promotion of the Crosslinking of Fibrin and α2-antiplasmin by Platelets. Thrombosis and Haemostasis, 1996, 75, 161-167.	1.8	31
149	Promotion of the crosslinking of fibrin and alpha 2-antiplasmin by platelets. Thrombosis and Haemostasis, 1996, 75, 161-7.	1.8	8
150	Localization of transglutaminase in human lenses Journal of Histochemistry and Cytochemistry, 1995, 43, 1173-1177.	1.3	11
151	Transformation of Cellular Factor XIII into an Active Zymogen Transglutaminase in Thrombin-Stimulated Platelets. Thrombosis and Haemostasis, 1995, 73, 702-705.	1.8	37
152	Transformation of cellular factor XIII into an active zymogen transglutaminase in thrombin-stimulated platelets. Thrombosis and Haemostasis, 1995, 73, 702-5.	1.8	11
153	Covalent binding of arachidonate to G protein alpha subunits of human platelets. Journal of Biological Chemistry, 1994, 269, 4713-6.	1.6	91
154	Platelet Factor XIII Becomes Active without the Release of Activation Peptide during Platelet Activation. Thrombosis and Haemostasis, 1993, 69, 282-285.	1.8	58
155	P-selectin is acylated with palmitic acid and stearic acid at cysteine 766 through a thioester linkage. Journal of Biological Chemistry, 1993, 268, 11394-400.	1.6	45
156	Platelet factor XIII becomes active without the release of activation peptide during platelet activation. Thrombosis and Haemostasis, 1993, 69, 282-5.	1.8	19
157	Covalent modification of proteins by arachidonate and eicosapentaenoate in platelets. Journal of Biological Chemistry, 1993, 268, 18243-8.	1.6	39
158	Myristoylation of proteins in platelets occurs predominantly through thioester linkages. Journal of Biological Chemistry, 1993, 268, 8251-5.	1.6	27
159	European Thrombosis Research Organization Working Party on Factor XIII Abstracts of the 1st Meeting 2???4 September, 1992 Debrecen, Hungary. Blood Coagulation and Fibrinolysis, 1992, 3, 803.	0.5	0
160	The role of factor XIII in intratumoural fibrin formation. Blood Coagulation and Fibrinolysis, 1992, 3, 808.	0.5	0
161	Marker profile, enzyme activity, and function of a human myelomonocytic leukemia cell line. Cellular Immunology, 1992, 139, 531-540.	1.4	17
162	Translocation of pp60c-src to the cytoskeleton during platelet aggregation. EMBO Journal, 1992, 11, 855-61.	3.5	21

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163	Isolation and partial characterization of a cystine-rich basic heparin-binding protein from bovine platelets. Biochemical and Biophysical Research Communications, 1991, 174, 70-76.	1.0	1
164	Non-proteolytic activation of cellular protransglutaminase (placenta macrophage factor XIII). Biochemical Journal, 1990, 267, 557-560.	1.7	76
165	Covalent modification of platelet proteins by palmitate. Blood, 1989, 74, 1339-1347.	0.6	27
166	Immunohistochemical detection of factor XIII subunit a in histiocytes of human uterus. Histochemistry, 1989, 91, 169-174.	1.9	22
167	Characterization of rapidly adhering amniotic fluid cells by combined immunofluorescence and phagocytosis assays. American Journal of Human Genetics, 1989, 45, 786-92.	2.6	34
168	Factors of the extrinsic pathway of blood coagulation in tumour associated macrophages. Thrombosis and Haemostasis, 1989, 62, 850-5.	1.8	3
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