

Geographic Distribution of the 20210 G to A Prothromb

Thrombosis and Haemostasis

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Novel inherited risk factors for venous thrombosis. <i>Translational Research</i> , 1998, 132, 444-445.	2.4	1
2	1 Thrombophilia: disorders predisposing to venous thromboembolism. <i>Best Practice and Research: Clinical Haematology</i> , 1998, 11, 525-540.	1.1	9
3	Prothrombin antigen levels in symptomatic and asymptomatic carriers of the 20210A prothrombin variant. <i>British Journal of Haematology</i> , 1998, 103, 1045-1050.	1.2	54
4	UPDATE ON GENETIC RISK FACTORS FOR THROMBOSIS AND ATHEROSCLEROTIC VASCULAR DISEASE. <i>Hematology/Oncology Clinics of North America</i> , 1998, 12, 1193-1209.	0.9	14
6	Venous Thrombosis – The Interaction of Genes and Environment. <i>New England Journal of Medicine</i> , 1998, 338, 1840-1841.	13.9	55
7	Polymorphisms of Coagulation Factor Genes a Review. <i>Clinical Chemistry and Laboratory Medicine</i> , 1998, 36, 897-906.	1.4	6
8	Prothrombin Gene Mutation (G20210A) in Healthy Centenarians. <i>Thrombosis and Haemostasis</i> , 1999, 81, 990-991.	1.8	6
9	Risk Factors for Venous Thrombotic Disease. <i>Thrombosis and Haemostasis</i> , 1999, 82, 610-619.	1.8	325
10	The 20210 G to A Prothrombin Polymorphism and Late Complications in Type 1 Diabetes Mellitus. <i>Thrombosis and Haemostasis</i> , 1999, 81, 164-164.	1.8	3
11	The Factor II G20210A and Factor V G1691A Gene Transitions and Coronary Heart Disease. <i>Thrombosis and Haemostasis</i> , 1999, 81, 208-213.	1.8	52
12	Compound Heterozygosity for One Novel and One Recurrent Mutation in a Thai Patient with Severe Protein S Deficiency. <i>Thrombosis and Haemostasis</i> , 1999, 81, 189-192.	1.8	22
13	Deep upper limb and jugular venous thrombosis in dilated cardiomyopathy. <i>Journal of the Royal Society of Medicine</i> , 1999, 92, 583-584.	1.1	1
14	The Prothrombin 20210A Allele and Its Association with Myocardial Infarction. <i>Thrombosis and Haemostasis</i> , 1999, 81, 861-864.	1.8	37
15	Absence of the Prothrombin Gene Variant in Koreans. <i>Thrombosis and Haemostasis</i> , 1999, 81, 990-990.	1.8	9
16	Prevalence of the Prothrombin Gene 20210A Mutation in Thrombophilic and Healthy Algerian Subjects. <i>Thrombosis and Haemostasis</i> , 1999, 82, 1554-1555.	1.8	15
17	Cerebral Vein Thrombosis not Related to Use of Oral Contraceptives in a 7-year-old Child Carrier of the Prothrombin 20210A Allele. <i>Thrombosis and Haemostasis</i> , 1999, 81, 991-992.	1.8	2
18	Thrombophilic Polymorphisms Are Common in Women with Fetal Loss without Apparent Cause. <i>Thrombosis and Haemostasis</i> , 1999, 82, 6-9.	1.8	219
19	The VITA Project: Prothrombin G20210A Mutation and Venous Thromboembolism in the General Population. <i>Thrombosis and Haemostasis</i> , 1999, 82, 1395-1398.	1.8	41

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20	Prothrombin G20210A Gene Mutation, Heparin Cofactor II Defects, Primary (Essential) Thrombocythemia, and Thrombohemorrhagic Manifestations. <i>Seminars in Thrombosis and Hemostasis</i> , 1999, 25, 375-386.	1.5	9
21	Genetic Risk Factors in Acute Coronary Disease. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 1999, 29, 212-218.	0.5	9
22	Prothrombin G20210A Gene Mutation and Further Prothrombotic Risk Factors in Childhood Thrombophilia. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 1999, 19, 2568-2572.	1.1	144
23	POSTINFECTION PURPURA FULMINANS IN A PATIENT HETEROZYGOUS FOR PROTHROMBIN G20210A AND ACQUIRED PROTEIN S RESISTANCE. <i>Pediatric Hematology and Oncology</i> , 1999, 16, 561-564.	0.3	18
24	Single and Combined Prothrombotic Factors in Patients With Idiopathic Venous Thromboembolism. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 1999, 19, 511-518.	1.1	244
25	Central Retinal Vein Thrombosis Associated with Prothrombin 20210G/A Gene Variant. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 1999, 5, 190-191.	0.7	0
26	The G20210A Polymorphism in the 3' -Untranslated Region of the Prothrombin Gene in Mexican Mestizo Patients with Primary Antiphospholipid Syndrome. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 1999, 5, 158-160.	0.7	17
27	Complex Association of Protein C Gene Promoter Polymorphism With Circulating Protein C Levels and Thrombotic Risk. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 1999, 19, 1573-1576.	1.1	69
28	Mutations in Promoter Region of Thrombomodulin and Venous Thromboembolic Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 1999, 19, 1098-1104.	1.1	66
29	Interaction Between the G20210A Mutation of the Prothrombin Gene and Oral Contraceptive Use in Deep Vein Thrombosis. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 1999, 19, 700-703.	1.1	170
30	The 20210 Gâ€†â€†A mutation in the 3â€²-untranslated region of the prothrombin gene and the risk for arterial thrombotic disease. <i>British Journal of Haematology</i> , 1999, 104, 50-54.	1.2	99
31	Influence of three potential genetic risk factors for thrombosis in 43 families carrying the factor V Arg 506 to Gln mutation. <i>British Journal of Haematology</i> , 1999, 106, 889-897.	1.2	11
32	The investigation and management of inherited thrombophilia. <i>International Journal of Laboratory Hematology</i> , 1999, 21, 77-92.	0.2	34
33	Development of an internal restriction control in the PCR detection of the prothrombin 20210A mutation. <i>International Journal of Laboratory Hematology</i> , 1999, 21, 281-283.	0.2	4
34	Born to clot: the European burden. <i>British Journal of Haematology</i> , 1999, 105, 564-566.	1.2	100
35	Frequencies of platelet-specific alloantigen systems 1-5 in three distinct ethnic groups in Brazil. <i>International Journal of Immunogenetics</i> , 1999, 26, 355-360.	1.2	37
36	Prevalence of factor V Leiden, prothrombin G20210A, and MTHFR C677T mutations in a Greek population of blood donors. , 1999, 61, 265-267.		59
37	The Risk of Recurrent Deep Venous Thrombosis among Heterozygous Carriers of Both Factor V Leiden and the G20210A Prothrombin Mutation. <i>New England Journal of Medicine</i> , 1999, 341, 801-806.	13.9	467

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38	Venous thrombosis: a multicausal disease. <i>Lancet, The</i> , 1999, 353, 1167-1173.	6.3	1,427
39	Prothrombotic mutations as a risk factor for preterm birth. <i>Lancet, The</i> , 1999, 353, 1411-1412.	6.3	45
40	The prothrombin G20210A mutation: A new high-prevalence congenital risk factor for thrombosis. <i>Gastroenterology</i> , 1999, 116, 213-215.	0.6	17
41	Hypercoagulable state mutation analysis in white patients with early first-trimester recurrent pregnancy loss. <i>Fertility and Sterility</i> , 1999, 71, 1048-1053.	0.5	146
42	Thrombophilia-associated pregnancy wastage. <i>Fertility and Sterility</i> , 1999, 72, 765-774.	0.5	120
43	The molecular genetics of familial venous thrombosis. <i>Best Practice and Research in Clinical Haematology</i> , 1999, 12, 479-503.	0.7	24
44	Venous Thrombosis: Prevalence and Interaction of Risk Factors. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 1999, 29, 1-9.	0.5	26
45	A Novel Clotting Assay for Quantitation of Plasma Prothrombin (Factor II) Using <i>Echis multisquamatus</i> Venom. <i>American Journal of Clinical Pathology</i> , 1999, 112, 705-711.	0.4	7
46	Prothrombin Gene Mutation Uncommon in Pulmonary Embolism. <i>Southern Medical Journal</i> , 2000, 93, 1073-1077.	0.3	3
47	Thrombophilia, polymorphisms, and vascular disease. <i>Journal of Clinical Pathology</i> , 2000, 53, 300-306.	2.1	36
48	Genetic Heterogeneity in Hereditary Thrombophilia. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 2000, 30, 1-10.	0.5	11
49	Inherited Thrombophilia as a Risk Factor for the Development of Ischemic Stroke in Young Adults. <i>Thrombosis and Haemostasis</i> , 2000, 83, 229-233.	1.8	108
50	Prevalence of the prothrombin G20210A mutation in the Irish populations: use of a novel polymerase chain reaction approach. <i>Blood Coagulation and Fibrinolysis</i> , 2000, 11, 669-672.	0.5	8
51	Prevalence of factor V Leiden and the G20210A prothrombin-gene mutation in inflammatory bowel disease. <i>Blood Coagulation and Fibrinolysis</i> , 2000, 11, 499-503.	0.5	33
52	DNA technology for the detection of common genetic variants that predispose to thrombophilia. <i>Blood Coagulation and Fibrinolysis</i> , 2000, 11, 683-700.	0.5	14
53	Clinical and laboratory expression of associated thrombophilic conditions (homozygous/heterozygous factor V Leiden mutation and heterozygous prothrombin variant 20210A) in an Italian family. <i>Blood Coagulation and Fibrinolysis</i> , 2000, 11, 379-384.	0.5	5
54	Prevalence of the G1691A mutation in the factor V gene (factor V Leiden) and the G20210A prothrombin gene mutation in the Thai population. <i>American Journal of Hematology</i> , 2000, 65, 119-122.	2.0	54
55	The G1691A mutation of factor V, but not the G20210A mutation of factor II or the C677T mutation of methylenetetrahydrofolate reductase genes, is associated with venous thrombosis in patients with lupus anticoagulants. <i>British Journal of Haematology</i> , 2000, 108, 865-870.	1.2	40

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57	Prevalence of methylenetetrahydrofolate reductase C677T and its association with arterial and venous thrombosis in the Chinese population. <i>British Journal of Haematology</i> , 2000, 109, 870-874.	1.2	66
58	Screening for thrombophilic risk factors among 25 German patients with cerebral venous thrombosis. <i>Acta Neurologica Scandinavica</i> , 2000, 102, 31-36.	1.0	31
59	A comparison of polymorphism in the 3' untranslated region of the prothrombin gene between Chinese and Caucasians in Australia. <i>British Journal of Haematology</i> , 2000, 111, 1253-1255.	1.2	0
60	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.	1.2	6
61	The decanucleotide polymorphism in the factor VII promoter predicts factor VII plasma levels but not the risk of acute coronary syndromes. <i>Journal of Thrombosis and Thrombolysis</i> , 2000, 10, 23-28.	1.0	5
63	Prevalence of prothrombin 20210A allele and methylenetetrahydrofolate reductase C677T genetic mutations in the Chinese population. <i>Annals of Hematology</i> , 2000, 79, 239-242.	0.8	17
64	Role of hemostatic gene polymorphisms in venous and arterial thrombotic disease. <i>Blood</i> , 2000, 95, 1517-1532.	0.6	472
65	Linkage analysis demonstrates that the prothrombin G20210A mutation jointly influences plasma prothrombin levels and risk of thrombosis. <i>Blood</i> , 2000, 95, 2780-2785.	0.6	114
66	Lack of Association of the Prothrombin Gene Variant G20210A with Myocardial Infarction in Caucasian Males. <i>Thrombosis and Haemostasis</i> , 2000, 83, 796-797.	1.8	7
67	Venous Thromboembolism and Associated High Plasma Factor VIII Levels: Linked to Cytomegalovirus Infection?. <i>Thrombosis and Haemostasis</i> , 2000, 83, 510-511.	1.8	32
68	Thrombophilic Mutations in Iran. <i>Thrombosis and Haemostasis</i> , 2000, 83, 351-352.	1.8	16
69	Factor II G20210A and Factor V G1691A Gene Mutations and Peripheral Arterial Occlusive Disease. <i>Thrombosis and Haemostasis</i> , 2000, 83, 20-22.	1.8	30
70	Factor V Leiden and Prothrombin G 20210A Variant Are Risk Factors for Venous Thromboembolism in the Argentinean Population. <i>Thrombosis and Haemostasis</i> , 2000, 83, 509-510.	1.8	9
71	Comparison of the Risk of Pulmonary Embolism and Deep Vein Thrombosis in the Presence of Factor V Leiden or Prothrombin G20210A. <i>Thrombosis and Haemostasis</i> , 2000, 83, 352-354.	1.8	15
72	Prothrombin Activation Is Increased among Asymptomatic Carriers of the Prothrombin G20210A and Factor V Arg506Gln Mutations. <i>Thrombosis and Haemostasis</i> , 2000, 84, 396-400.	1.8	26
73	Management of Patients with Hereditary Hypercoagulable Disorders. <i>Annual Review of Medicine</i> , 2000, 51, 169-185.	5.0	115
74	Factor V Leiden and prothrombin G20210A mutations, but not methylenetetrahydrofolate reductase C677T, are associated with recurrent miscarriages. <i>Human Reproduction</i> , 2000, 15, 458-462.	0.4	162

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75	Familial thrombophilia and retinal vein occlusion. <i>Eye</i> , 2000, 14, 13-17.	1.1	27
76	Association of the Prothrombin G20210A Mutation With Factor V Leiden in a Midwestern American Population. <i>American Journal of Clinical Pathology</i> , 2000, 114, 272-275.	0.4	12
77	A Randomized Trial of Oral Contraceptive and Hormone Replacement Therapy on Bone Mineral Density and Coronary Heart Disease Risk Factors in Postmenopausal Women. <i>Obstetrics and Gynecology</i> , 2000, 95, 87-94.	1.2	0
78	Mutations in Coagulation Factors in Women with Unexplained Late Fetal Loss. <i>New England Journal of Medicine</i> , 2000, 343, 1015-1018.	13.9	296
79	Venous Thromboembolism: Risk Factors and Prophylaxis. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2000, 21, 477-486.	0.8	5
80	Smoking: back to the futureâ€”again?. <i>European Heart Journal</i> , 2000, 21, 1570-1571.	1.0	1
81	Acute Myocardial Infarction Associated With the Prothrombin G20210A Mutation. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2000, 6, 111-112.	0.7	0
82	Prevalence of Genetic Mutations That Predispose to Thrombophilia in a Greek Cypriot Population. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2000, 6, 104-107.	0.7	32
83	Genetic Risk Factors for Venous Thrombosis. <i>Molecular Genetics and Metabolism</i> , 2000, 71, 51-61.	0.5	16
85	A randomized trial of oral contraceptive and hormone replacement therapy on bone mineral density and coronary heart disease risk factors in postmenopausal women. <i>Obstetrics and Gynecology</i> , 2000, 95, 87-94.	1.2	24
86	Prothrombin G20210A, Factor V Leiden, and Factor XIII Val34Leu. <i>Thrombosis Research</i> , 2000, 99, 35-39.	0.8	58
87	The Allele Frequency of Mutations in Four Genes that Confer Enhanced Susceptibility to Venous Thromboembolism in an Unselected Group of New York State Newborns. <i>Thrombosis Research</i> , 2000, 99, 317-324.	0.8	39
88	Prothrombin Activity and Concentration in Healthy Subjects with and without the Prothrombin G20210A Mutation. <i>Thrombosis Research</i> , 2000, 99, 549-556.	0.8	18
90	Prothrombin G20210A Polymorphism and Thrombophilia. <i>Mayo Clinic Proceedings</i> , 2000, 75, 595-604.	1.4	49
92	Prevalence of factor v leiden and prothrombin variant g20210a in patients age <50 years with no significant stenoses at angiography three to four weeks after myocardial infarction. <i>Journal of the American College of Cardiology</i> , 2000, 36, 717-722.	1.2	71
93	Genetic Susceptibility to Venous Thrombosis. <i>New England Journal of Medicine</i> , 2001, 344, 1222-1231.	13.9	763
94	Prothrombotic inherited abnormalities other than factor v leiden mutation do not play a role in venous thrombosis in inflammatory bowel disease. <i>American Journal of Gastroenterology</i> , 2001, 96, 1448-1454.	0.2	36
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97	When to suspect hypercoagulability and how to investigate it. <i>Annals of Diagnostic Pathology</i> , 2001, 5, 177-183.	0.6	4
98	Major and Potential Prothrombotic Genotypes in a Cohort of Patients With Venous Thromboembolism. <i>Thrombosis Research</i> , 2001, 104, 317-324.	0.8	38
99	Simultaneous allele-specific amplification: A strategy using modified primer-template mismatches for SNP detection[mdash]Application to prothrombin 20210A (factor II) and factor V Leiden (1691A) gene mutations. <i>Molecular Diagnosis and Therapy</i> , 2001, 6, 201-209.	1.3	7
100	Factor V Leiden, Prothrombin Gene G20210A Variant, and Methylenetetrahydrofolate Reductase C677T Genotype in Young Adults With Ischemic Stroke. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2001, 7, 346-350.	0.7	55
101	Prevalence of Molecular Risk Factors FV Leiden, FV HR2, FII 20210G>A and MTHFR 677C>T in Different Populations and Ethnic Groups of Germany, Costa Rica and India. <i>International Journal of Human Genetics</i> , 2001, 1, 33-39.	0.1	16
102	Polymorphisms in the Prothrombin Gene and their Association with Plasma Prothrombin Levels. <i>Thrombosis and Haemostasis</i> , 2001, 85, 1066-1070.	1.8	51
103	Combined Effect of Factor V Leiden and Prothrombin 20210A on the Risk of Venous Thromboembolism. <i>Thrombosis and Haemostasis</i> , 2001, 86, 809-816.	1.8	301
104	Prevalence of Factor V Leiden and Prothrombin 20210A Mutations in Indigenous Australians. <i>Thrombosis and Haemostasis</i> , 2001, 86, 1592-1593.	1.8	16
105	Coexistence of Factor V G1691A and Factor II G20210A Gene Mutations in a Thrombotic Family Is Associated with Recurrence and Early Onset of Venous Thrombosis. <i>Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research</i> , 2001, 31, 99-105.	0.5	8
106	Laboratory Investigation of Thrombophilia. <i>Clinical Chemistry</i> , 2001, 47, 1597-1606.	1.5	116
107	Heterozygous carrier of G20210A prothrombin mutation used oral contraceptive treatment for 23 years without thrombotic events, and developed cerebral venous thrombosis 1 month after resumption of the medication at the age of 50. <i>Blood Coagulation and Fibrinolysis</i> , 2001, 12, 161-162.	0.5	1
108	Thrombogenic alleles, Escherichia coli O157:H7 infections, and hemolytic uremic syndrome. <i>Blood Coagulation and Fibrinolysis</i> , 2001, 12, 283-288.	0.5	5
109	An Update on Hypercoagulable Disorders. <i>Archives of Internal Medicine</i> , 2001, 161, 1051.	4.3	61
110	Genetic hypercoagulability: screening should be an informed choice. <i>Blood</i> , 2001, 98, 20-20.	0.6	29
111	Primary Prevention of Ischemic Stroke. <i>Stroke</i> , 2001, 32, 280-299.	1.0	512
112	Genetic Approach to Thrombophilia. <i>Thrombosis and Haemostasis</i> , 2001, 86, 92-103.	1.8	78
113	Risk Factors in Venous Thromboembolism. <i>Thrombosis and Haemostasis</i> , 2001, 86, 395-403.	1.8	176

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115	Die Bedeutung von Mutationen in den Genen fÄ¼r Faktor V, Faktor II und der Methylenetetrahydrofolatreduktase bei habituellen Aborten. <i>Reproduktionsmedizin</i> , 2001, 17, 42-47.	0.1	2
116	The heterozygous 20210 G/A genotype prevalence in patients affected by central and branch retinal vein occlusion: a pilot study. <i>Graefe's Archive for Clinical and Experimental Ophthalmology</i> , 2001, 239, 251-256.	1.0	15
117	Genetic risk factors of venous thrombosis. <i>Human Genetics</i> , 2001, 109, 369-384.	1.8	252
118	Genetics University of Toronto Thrombophilia Study in Women (GUTTSI): genetic and other risk factors for venous thromboembolism in women. <i>Current Controlled Trials in Cardiovascular Medicine</i> , 2001, 2, 141.	1.5	31
119	Gene polymorphisms of the haemostatic system and the risk of arterial thrombotic disease. <i>British Journal of Haematology</i> , 2001, 115, 491-506.	1.2	46
120	Primary thrombophilia in Mexico. II. Factor V G1691A (Leiden), prothrombin G20210A, and methylenetetrahydrofolate reductase C677T polymorphism in thrombophilic Mexican mestizos. <i>American Journal of Hematology</i> , 2001, 66, 28-31.	2.0	34
121	Update on selected inherited venous thrombotic disorders. <i>American Journal of Hematology</i> , 2001, 68, 256-268.	2.0	51
122	Two common genetic thrombotic risk factors: Factor V Leiden and prothrombin G20210A in adult Turkish patients with thrombosis. <i>American Journal of Hematology</i> , 2001, 67, 107-111.	2.0	37
123	Genotyping of thrombotic risk factors by maldi-tof mass spectrometry. <i>Clinical Biochemistry</i> , 2001, 34, 531-536.	0.8	29
124	Impact of the factor II: G20210A variant on the risk of venous thromboembolism in relatives from families with the factor V: R506Q mutation. <i>European Journal of Haematology</i> , 2001, 67, 165-169.	1.1	6
125	G20210A prothrombin gene mutation identified in patients with venous leg ulcers. <i>Journal of Cellular and Molecular Medicine</i> , 2001, 5, 397-401.	1.6	14
127	High prevalence of thrombophilic genotypes in patients with acute mesenteric vein thrombosis. <i>American Journal of Gastroenterology</i> , 2001, 96, 146-149.	0.2	52
128	Thrombophilia: genetic polymorphisms and their association with retinal vascular occlusive disease. <i>British Journal of Ophthalmology</i> , 2001, 85, 883-886.	2.1	22
129	Venous Thromboembolism in Young Patients From Western India: A Study. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2001, 7, 158-165.	0.7	62
130	Primary Prevention of Ischemic Stroke. <i>Circulation</i> , 2001, 103, 163-182.	1.6	340
131	Hypercoagulability Syndromes. <i>Archives of Internal Medicine</i> , 2001, 161, 2433-2439.	4.3	116
132	G20210A Prothrombin Gene Polymorphism and Prothrombin Activity in Subjects With or Without Angiographically Documented Coronary Artery Disease. <i>Circulation</i> , 2001, 103, 2436-2440.	1.6	44

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134	Venous Thromboembolic Disease: Risk Factors and Laboratory Investigation. <i>Seminars in Vascular Medicine</i> , 2001, 01, 081-088.	2.1	12
135	Factor V and Thrombotic Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2002, 22, 530-538.	1.1	188
136	Venous thromboembolism in young women. Role of thrombophilic mutations and oral contraceptive use. <i>European Heart Journal</i> , 2002, 23, 984-990.	1.0	55
137	Portal Vein Thrombosis after Variceal Endoscopic Sclerotherapy in Cirrhotic Patients: Role of Genetic Thrombophilia. <i>Endoscopy</i> , 2002, 34, 535-538.	1.0	89
138	Hereditary thrombophilia in elite athletes. <i>Medicine and Science in Sports and Exercise</i> , 2002, 34, 218-221.	0.2	20
139	The G20210A mutation does not affect the stability of prothrombin mRNA in vivo. <i>Blood</i> , 2002, 100, 359-362.	0.6	44
140	Increasing Mortality From Pulmonary Embolism in Japan, 1951-2000.. <i>Circulation Journal</i> , 2002, 66, 1144-1149.	0.7	63
141	Hypercoagulability: Too Many Tests, Too Much Conflicting Data. <i>Hematology American Society of Hematology Education Program</i> , 2002, 2002, 353-368.	0.9	41
142	The prothrombin G20210A polymorphism in patients with myocardial infarction. <i>Blood Coagulation and Fibrinolysis</i> , 2002, 13, 603-608.	0.5	8
143	Risk factors for thromboembolism in teens: when should I test?. <i>Current Opinion in Pediatrics</i> , 2002, 14, 370-378.	1.0	26
144	Arterial wall thickness and the risk of recurrent ischemic events in carriers of the prothrombin G20210A mutation with clinical manifestations of atherosclerosis. <i>Atherosclerosis</i> , 2002, 163, 135-140.	0.4	29
145	Female Hormones and Thrombosis. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2002, 22, 201-210.	1.1	239
146	Factor V Leiden and prothrombin G20210A mutations in pregnancies with adverse outcome. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2002, 12, 267-273.	0.7	39
147	Recurrent pulmonary embolism in a 13-year-old male homozygous for the prothrombin G20210A mutation combined with protein S deficiency and increased lipoprotein (a). <i>Thrombosis Research</i> , 2002, 105, 49-53.	0.8	19
148	Markers of activated coagulation in patients with factor V Leiden and/or G20210A prothrombin gene mutation. <i>Thrombosis Research</i> , 2002, 107, 7-11.	0.8	15
149	The PORtromb Project: prothrombin G20210A mutation and venous thromboembolism in young people. <i>Vascular</i> , 2002, 10, 45-48.	0.5	7
150	Update in pulmonary medicine. <i>Clinics in Podiatric Medicine and Surgery</i> , 2002, 19, 1-22.	0.2	2

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