Nicola Martinelli

List of Publications by Year
in descending order

147 papers	13,943 citations	46984 47 h-index	20943 115 g-index
150 all docs	150 docs citations	150 times ranked	21247 citing authors

#	Article	IF	Citations
1	Plasma HDL cholesterol and risk of myocardial infarction: a mendelian randomisation study. Lancet, The, 2012, 380, 572-580.	6.3	1,937
2	Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease. Nature Genetics, 2011, 43, 333-338.	9.4	1,685
3	Genome-wide association of early-onset myocardial infarction with single nucleotide polymorphisms and copy number variants. Nature Genetics, 2009, 41, 334-341.	9.4	990
4	Loss-of-Function Mutations inAPOC3,Triglycerides, and Coronary Disease. New England Journal of Medicine, 2014, 371, 22-31.	13.9	936
5	Sequence variants affecting eosinophil numbers associate with asthma and myocardial infarction. Nature Genetics, 2009, 41, 342-347.	9.4	709
6	Exome sequencing identifies rare LDLR and APOA5 alleles conferring risk for myocardial infarction. Nature, 2015, 518, 102-106.	13.7	581
7	Identification of ADAMTS7 as a novel locus for coronary atherosclerosis and association of ABO with myocardial infarction in the presence of coronary atherosclerosis: two genome-wide association studies. Lancet, The, 2011, 377, 383-392.	6.3	466
8	Coding Variation inANGPTL4,LPL,andSVEP1and the Risk of Coronary Disease. New England Journal of Medicine, 2016, 374, 1134-1144.	13.9	427
9	Association of Low-Frequency and Rare Coding-Sequence Variants with Blood Lipids and Coronary Heart Disease in 56,000 Whites and Blacks. American Journal of Human Genetics, 2014, 94, 223-232.	2.6	287
10	FADS genotypes and desaturase activity estimated by the ratio of arachidonic acid to linoleic acid are associated with inflammation and coronary artery disease. American Journal of Clinical Nutrition, 2008, 88, 941-949.	2.2	286
11	Air particulate matter and cardiovascular disease: A narrative review. European Journal of Internal Medicine, 2013, 24, 295-302.	1.0	235
12	SNPs of the FADS Gene Cluster are Associated with Polyunsaturated Fatty Acids in a Cohort of Patients with Cardiovascular Disease. Lipids, 2008, 43, 289-299.	0.7	218
13	Systematic Evaluation of Pleiotropy Identifies 6 Further Loci Associated WithÂCoronary ArteryÂDisease. Journal of the American College of Cardiology, 2017, 69, 823-836.	1.2	214
14	Large-Scale Gene-Centric Analysis Identifies Novel Variants for Coronary Artery Disease. PLoS Genetics, 2011, 7, e1002260.	1.5	203
15	Abdominal Aortic Aneurysm Is Associated with a Variant in Low-Density Lipoprotein Receptor-Related Protein 1. American Journal of Human Genetics, 2011, 89, 619-627.	2.6	185
16	Prevalence of Body Iron Excess in the Metabolic Syndrome. Diabetes Care, 2005, 28, 2061-2063.	4.3	181
17	Genome-wide association study identifies a sequence variant within the DAB2IP gene conferring susceptibility to abdominal aortic aneurysm. Nature Genetics, 2010, 42, 692-697.	9.4	181
18	Randomized placebo-controlled trial comparing fluticasone aqueous nasal spray in mono-therapy, fluticasone plus cetirizine, fluticasone plus montelukast and cetirizine plus montelukast for seasonal allergic rhinitis. Clinical and Experimental Allergy, 2004, 34, 259-267.	1.4	162

#	Article	IF	Citations
19	Evaluation of serum s-IgE/total IgE ratio in predicting clinical response to allergen-specific immunotherapy. Journal of Allergy and Clinical Immunology, 2009, 123, 1103-1110.e4.	1.5	161
20	Reduced serum hepcidin levels in patients with chronic hepatitis C. Journal of Hepatology, 2009, 51, 845-852.	1.8	148
21	Randomized placebo-controlled trial comparing desloratadine and montelukast in monotherapy and desloratadine plus montelukast in combined therapy for chronic idiopathic urticaria. Journal of Allergy and Clinical Immunology, 2004, 114, 619-625.	1.5	130
22	Delta-5 and Delta-6 Desaturases: Crucial Enzymes in Polyunsaturated Fatty Acid-Related Pathways with Pleiotropic Influences in Health and Disease. Advances in Experimental Medicine and Biology, 2014, 824, 61-81.	0.8	128
23	Low plasma vitamin B-6 concentrations and modulation of coronary artery disease risk. American Journal of Clinical Nutrition, 2004, 79, 992-998.	2.2	117
24	Scleroderma patients nailfold videocapillaroscopic patterns are associated with disease subset and disease severity. Rheumatology, 2007, 46, 1566-1569.	0.9	116
25	Association between four SNPs on chromosome 9p21 and myocardial infarction is replicated in an Italian population. Journal of Human Genetics, 2008, 53, 144-150.	1.1	112
26	Apolipoprotein C-III, metabolic syndrome, and risk of coronary artery disease. Journal of Lipid Research, 2003, 44, 2374-2381.	2.0	111
27	Iron deficiency in the elderly population, revisited in the hepcidin era. Frontiers in Pharmacology, 2014, 5, 83.	1.6	97
28	Hepcidin Levels and Their Determinants in Different Types of Myelodysplastic Syndromes. PLoS ONE, 2011, 6, e23109.	1.1	95
29	Interaction of antibodies against cytomegalovirus with heat-shock protein 60 in pathogenesis of atherosclerosis. Lancet, The, 2003, 362, 1971-1977.	6.3	93
30	Exome Sequencing and Directed Clinical Phenotyping Diagnose Cholesterol Ester Storage Disease Presenting as Autosomal Recessive Hypercholesterolemia. Arteriosclerosis, Thrombosis, and Vascular Biology, 2013, 33, 2909-2914.	1.1	87
31	Polymorphisms at LDLR locus may be associated with coronary artery disease through modulation of coagulation factor VIII activity and independently from lipid profile. Blood, 2010, 116, 5688-5697.	0.6	86
32	Lack of Association Between the Trp719Arg Polymorphism in Kinesin-Like Protein-6 and Coronary Artery Disease in 19 Case-Control Studies. Journal of the American College of Cardiology, 2010, 56, 1552-1563.	1.2	84
33	Relations between lipoprotein(a) concentrations, LPA genetic variants, and the risk of mortality in patients with established coronary heart disease: a molecular and genetic association study. Lancet Diabetes and Endocrinology,the, 2017, 5, 534-543.	5.5	84
34	Comparative effect of tacrolimus 0.1% ointment and clobetasol 0.05% ointment in patients with oral lichen planus. Journal of Clinical Periodontology, 2008, 35, 244-249.	2.3	74
35	Increased Serum Hepcidin Levels in Subjects with the Metabolic Syndrome: A Population Study. PLoS ONE, 2012, 7, e48250.	1.1	68
36	Food-Additive-Induced Urticaria: A Survey of 838 Patients with Recurrent Chronic Idiopathic Urticaria. International Archives of Allergy and Immunology, 2005, 138, 235-242.	0.9	67

#	Article	IF	Citations
37	The â˜1131 T>C and S19W APOA5 gene polymorphisms are associated with high levels of triglycerides and apolipoprotein C-III, but not with coronary artery disease: an angiographic study. Atherosclerosis, 2007, 191, 409-417.	0.4	67
38	Differences and Similarities between Allergic and Nonallergic Rhinitis in a Large Sample of Adult Patients with Rhinitis Symptoms. International Archives of Allergy and Immunology, 2011, 155, 263-270.	0.9	66
39	Comparing tacrolimus ointment and oral cyclosporine in adult patients affected by atopic dermatitis: a randomized study. Clinical and Experimental Allergy, 2004, 34, 639-645.	1.4	64
40	Anti-TNFÎ± therapy in rheumatoid arthritis and autoimmunity. Rheumatology International, 2006, 26, 209-214.	1.5	64
41	Global DNA Hypomethylation in Peripheral Blood Mononuclear Cells as a Biomarker of Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 348-355.	1.1	59
42	Promoter methylation in coagulationF7gene influences plasma FVII concentrations and relates to coronary artery disease. Journal of Medical Genetics, 2012, 49, 192-199.	1.5	57
43	A relative ADAMTS13 deficiency supports the presence of a secondary microangiopathy in COVID 19. Thrombosis Research, 2020, 193, 170-172.	0.8	57
44	Gene sequence variations of the platelet P2Y12 receptor are associated with coronary artery disease. BMC Medical Genetics, 2007, 8, 59.	2.1	53
45	Apolipoprotein Câ€III predicts cardiovascular mortality in severe coronary artery disease and is associated with an enhanced plasma thrombin generation. Journal of Thrombosis and Haemostasis, 2010, 8, 463-471.	1.9	53
46	Antibodies against cyclic citrullinated peptides in patients affected by rheumatoid arthritis before and after infliximab treatment. Rheumatology International, 2005, 26, 58-62.	1.5	52
47	A1298C methylenetetrahydrofolate reductase mutation and coronary artery disease: relationships with C677T polymorphism and homocysteine/folate metabolism. Clinical and Experimental Medicine, 2002, 2, 7-12.	1.9	50
48	Interaction between smoking and PON2 Ser311Cys polymorphism as a determinant of the risk of myocardial infarction. European Journal of Clinical Investigation, 2004, 34, 14-20.	1.7	49
49	Monosodium benzoate hypersensitivity in subjects with persistent rhinitis. Allergy: European Journal of Allergy and Clinical Immunology, 2004, 59, 192-197.	2.7	47
50	Apolipoprotein C-III, n-3 Polyunsaturated Fatty Acids, and â€œInsulin-Resistantâ€•Tâ˜455C APOC3 Gene Polymorphism in Heart Disease Patients: Example of Gene-Diet Interaction. Clinical Chemistry, 2005, 51, 360-367.	1.5	47
51	On the association of the oxidised LDL receptor 1 (OLR1) gene in patients with acute myocardial infarction or coronary artery disease. European Journal of Human Genetics, 2006, 14, 127-130.	1.4	45
52	Folic Acid Effects on S-Adenosylmethionine, S-Adenosylhomocysteine, and DNA Methylation in Patients with Intermediate Hyperhomocysteinemia. Journal of the American College of Nutrition, 2011, 30, 11-18.	1.1	45
53	Access Rate to the Emergency Department for Venous Thromboembolism in Relationship with Coarse and Fine Particulate Matter Air Pollution. PLoS ONE, 2012, 7, e34831.	1.1	44
54	Influence of polymorphisms in the factor VII gene promoter on activated factor VII levels and on the risk of myocardial infarction in advanced coronary atherosclerosis. Thrombosis and Haemostasis, 2004, 92, 541-549.	1.8	43

#	Article	IF	Citations
55	ApoE Î¼2/Î¼3/Î¼4 polymorphism, ApoC-III/ApoE ratio and metabolic syndrome. Clinical and Experimental Medicine, 2007, 7, 164-172.	1.9	39
56	A score of risk factors associated with ischemic digital ulcers in patients affected by systemic sclerosis treated with iloprost. Clinical Rheumatology, 2009, 28, 807-813.	1.0	39
57	ALOX5AP gene variants and risk of coronary artery disease: an angiography-based study. European Journal of Human Genetics, 2007, 15, 959-966.	1.4	37
58	Whole-genome sequencing analysis of semi-supercentenarians. ELife, 2021, 10, .	2.8	37
59	Genetic Architecture of Coronary Artery Disease in the Genome-Wide Era: Implications for the Emerging â€œGolden Dozenâ€•Loci. Seminars in Thrombosis and Hemostasis, 2009, 35, 671-682.	1.5	36
60	Combined Effect of Hemostatic Gene Polymorphisms and the Risk of Myocardial Infarction in Patients with Advanced Coronary Atherosclerosis. PLoS ONE, 2008, 3, e1523.	1.1	35
61	Serum DNase I, soluble Fas/FasL levels and cell surface Fas expression in patients with SLE: a possible explanation for the lack of efficacy of hrDNase I treatment. International Immunology, 2009, 21, 237-243.	1.8	35
62	Factor H interferes with the adhesion of sickle red cells to vascular endothelium: a novel disease-modulating molecule. Haematologica, 2019, 104, 919-928.	1.7	34
63	Evaluation of Hepcidin Isoforms in Hemodialysis Patients by a Proteomic Approach Based on SELDI-TOF MS. Journal of Biomedicine and Biotechnology, 2010, 2010, 1-7.	3.0	33
64	Association and Functional Analyses of MEF2A as a Susceptibility Gene for Premature Myocardial Infarction and Coronary Artery Disease. Circulation: Cardiovascular Genetics, 2009, 2, 165-172.	5.1	32
65	Novel serum paraoxonase activity assays are associated with coronary artery disease. Clinical Chemistry and Laboratory Medicine, 2009, 47, 432-40.	1.4	29
66	Serum levels of the hepcidin-20 isoform in a large general population: The Val Borbera study. Journal of Proteomics, 2012, 76, 28-35.	1.2	29
67	Paraoxonases. Advances in Clinical Chemistry, 2013, 59, 65-100.	1.8	29
68	Infections and autoimmunity: role of human cytomegalovirus in autoimmune endothelial cell damage. Lupus, 2015, 24, 419-432.	0.8	29
69	Impact of Selection Bias on Estimation of Subsequent Event Risk. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	28
70	Modulation of Factor V Levels in Plasma by Polymorphisms in the C2 Domain. Arteriosclerosis, Thrombosis, and Vascular Biology, 2004, 24, 200-206.	1.1	27
71	Factor II Activity is Similarly Increased in Patients With Elevated Apolipoprotein CIII and in Carriers of the Factor II 20210A Allele. Journal of the American Heart Association, 2013, 2, e000440.	1.6	27
72	Low Levels of Serum Paraoxonase Activities are Characteristic of Metabolic Syndrome and May Influence the Metabolic-Syndrome-Related Risk of Coronary Artery Disease. Experimental Diabetes Research, 2012, 2012, 1-9.	3.8	26

#	Article	IF	Citations
73	A â€˜Desaturase Hypothesisâ€™ for Atherosclerosis: Janus-Faced Enzymes in Ï‰â€“6 and Ï‰â€“3 Polyunsaturated Fatty Acid Metabolism. Journal of Nutrigenetics and Nutrigenomics, 2009, 2, 129-139.	1.8	24
74	Myristic acid induces proteomic and secretomic changes associated with steatosis, cytoskeleton remodeling, endoplasmic reticulum stress, protein turnover and exosome release in HepG2 cells. Journal of Proteomics, 2018, 181, 118-130.	1.2	24
75	Additive effect of LRP8/APOER2 R952Q variant to APOE Îµ2/Îµ3/Îµ4 genotype in modulating apolipoprotein E concentration and the risk of myocardial infarction: a case-control study. BMC Medical Genetics, 2009, 10, 41.	2.1	23
76	Improvement of maternal and fetal outcomes in women with sickle cell disease treated with early prophylactic erythrocytapheresis. Transfusion, 2018, 58, 2192-2201.	0.8	22
77	Association of Chromosome 9p21 With Subsequent Coronary Heart Disease Events. Circulation Genomic and Precision Medicine, 2019, 12, e002471.	1.6	22
78	Tacrolimus ointment in nickel sulphateâ€“induced steroid-resistant allergic contact dermatitis. Allergy and Asthma Proceedings, 2006, 27, 527-531.	1.0	21
79	A Novel Molecular Diagnostic Marker for Familial and Early-Onset Coronary Artery Disease and Myocardial Infarction in the LRP8 Gene. Circulation: Cardiovascular Genetics, 2014, 7, 514-520.	5.1	21
80	Activated factor VIIâ€“antithrombin complex predicts mortality in patients with stable coronary artery disease: a cohort study. Journal of Thrombosis and Haemostasis, 2016, 14, 655-666.	1.9	21
81	Interaction between metabolic syndrome and PON1 polymorphisms as a determinant of the risk of coronary artery disease. Clinical and Experimental Medicine, 2005, 5, 20-30.	1.9	20
82	Paraoxonase-1 status in patients with hereditary hemochromatosis. Journal of Lipid Research, 2013, 54, 1484-1492.	2.0	20
83	One-carbon genetic variants and the role of MTHFD1 1958G>A in liver and colon cancer risk according to global DNA methylation. PLoS ONE, 2017, 12, e0185792.	1.1	19
84	Comparison of topical tacrolimus 0.1 % in pectin ointment with clobetasol 0.5% ointment in adults with moderate to severe desquamative gingivitis: A 4-week, randomized, double-blind clinical trial. Clinical Therapeutics, 2006, 28, 1296-1302.	1.1	18
85	Relationship Between Human Leucocyte Antigen Class I and Class II and Chronic Idiopathic Urticaria Associated With Aspirin and/or NSAIDs Hypersensitivity. Mediators of Inflammation, 2006, 2006, 1-5.	1.4	18
86	Hyperhomocysteinemia and Mortality after Coronary Artery Bypass Grafting. PLoS ONE, 2006, 1, e83.	1.1	17
87	Subsequent Event Risk in Individuals With Established Coronary Heart Disease. Circulation Genomic and Precision Medicine, 2019, 12, e002470.	1.6	17
88	Apolipoprotein C-III Strongly Correlates with Activated Factor VIIâ€“Anti-Thrombin Complex: An Additional Link between Plasma Lipids and Coagulation. Thrombosis and Haemostasis, 2019, 119, 192-202.	1.8	17
89	Basophil Blood Cell Count Is Associated With Enhanced Factor II Plasma Coagulant Activity and Increased Risk of Mortality in Patients With Stable Coronary Artery Disease: Not Only Neutrophils as Prognostic Marker in Ischemic Heart Disease. Journal of the American Heart Association, 2021, 10, e018243.	1.6	17
90	Serum levels of soluble CD30 in adult patients affected by atopic dermatitis and its relation to age, duration of disease and Scoring Atopic Dermatitis index. Mediators of Inflammation, 2003, 12, 123-125.	1.4	15

#	Article	IF	Citations
91	Clinical Importance of Eosinophil Count in Nasal Fluid in Patients with Allergic and Non-Allergic Rhinitis. International Journal of Immunopathology and Pharmacology, 2009, 22, 1077-1087.	1.0	15
92	Functional polymorphisms in the LDLR and pharmacokinetics of Factor VIII concentrates. Journal of Thrombosis and Haemostasis, 2019, 17, 1288-1296.	1.9	15
93	Deep vein thrombosis in SARS-CoV-2 pneumonia-affected patients within standard care units: Exploring a submerged portion of the iceberg. Thrombosis Research, 2020, 194, 216-219.	0.8	15
94	A decade of progress on the genetic basis of coronary artery disease. Practical insights for the internist. European Journal of Internal Medicine, 2017, 41, 10-17.	1.0	14
95	Increased plasma thrombin potential is associated with stable coronary artery disease: An angiographically-controlled study. Thrombosis Research, 2017, 155, 16-22.	0.8	14
96	Multi-allelic haplotype association identifies novel information different from single-SNP analysis: A new protective haplotype in the LRP8 gene is against familial and early-onset CAD and MI. Gene, 2013, 521, 78-81.	1.0	13
97	An integrated genomic-transcriptomic approach supports a role for the proto-oncogene BCL3 in atherosclerosis. Thrombosis and Haemostasis, 2015, 113, 655-663.	1.8	13
98	The Positive Association between Plasma Myristic Acid and ApoCIII Concentrations in Cardiovascular Disease Patients Is Supported by the Effects of Myristic Acid in HepG2 Cells. Journal of Nutrition, 2020, 150, 2707-2715.	1.3	11
99	Reply to Novelli. European Journal of Human Genetics, 2006, 14, 895-895.	1.4	10
100	Sialylated isoforms of apolipoprotein C-III and plasma lipids in subjects with coronary artery disease. Clinical Chemistry and Laboratory Medicine, 2018, 56, 1542-1550.	1.4	10
101	Serum Hepcidin Levels Correlate with Phenotypes of the Metabolic Syndrome At Population Level. Blood, 2011, 118, 348-348.	0.6	10
102	Results of Double-Blind Placebo-Controlled Challenge with Nickel Salts in Patients Affected by Recurrent Aphthous Stomatitis. International Archives of Allergy and Immunology, 2003, 131, 296-300.	0.9	9
103	Reply to J Dierkes et al. American Journal of Clinical Nutrition, 2005, 81, 727-728.	2.2	9
104	Detection of a large deletion in the P-selectin (SELP) gene. Molecular and Cellular Probes, 2010, 24, 161-165.	0.9	9
105	Polymorphisms of cyclo-oxygenases and 5-lipo-oxygenase-activating protein are associated with chronic spontaneous urticaria and urinary leukotriene E4. European Journal of Dermatology, 2011, 21, 47-52.	0.3	9
106	Not Just Arterial Damage: Increased Incidence of Venous Thromboembolic Events in Cardiovascular Patients With Elevated Plasma Levels of Apolipoprotein CIII. Journal of the American Heart Association, 2019, 8, e010973.	1.6	9
107	Rare variants lowering the levels of coagulation factor X are protective against ischemic heart disease. Haematologica, 2020, 105, e365-e369.	1.7	9
108	The Asialoglycoprotein Receptor Minor Subunit Gene Contributes to Pharmacokinetics of Factor VIII Concentrates in Hemophilia A. Thrombosis and Haemostasis, 2022, 122, 715-725.	1.8	9

#	Article	IF	Citations
109	Evaluation of finger skin temperature in scleroderma patients cyclically treated with iloprost. Joint Bone Spine, 2006, 73, 57-61.	0.8	7
110	Increased factor VIII coagulant activity levels in male carriers of the factor V R2 polymorphism. Blood Coagulation and Fibrinolysis, 2007, 18, 125-129.	0.5	7
111	rs629301 CELSR2 polymorphism confers a ten-year equivalent risk of critical stenosis assessed by coronary angiography. Nutrition, Metabolism and Cardiovascular Diseases, 2021, 31, 1542-1547.	1.1	7
112	Long-term evaluation of lung function in patients affected by scleroderma treated with cyclic iloprost infusions. Rheumatology International, 2005, 25, 250-254.	1.5	6
113	Homocysteine, traditional risk factors and impaired renal function in coronary artery disease. European Journal of Clinical Investigation, 2006, 36, 698-704.	1.7	6
114	Aptamer-modified FXa generation assays to investigate hypercoagulability in plasma from patients with ischemic heart disease. Thrombosis Research, 2020, 189, 140-146.	0.8	6
115	An Unusual Heart Failure. Circulation, 2011, 123, e583-4.	1.6	5
116	The RFC1 80G>A, among Common One-Carbon Polymorphisms, Relates to Survival Rate According to DNA Global Methylation in Primary Liver Cancers. PLoS ONE, 2016, 11, e0167534.	1.1	5
117	Increased Incidence of Ischemic Cerebrovascular Events in Cardiovascular Patients With Elevated Apolipoprotein CIII. Stroke, 2020, 51, 61-68.	1.0	5
118	Therapeutic oligonucleotides in cardiovascular and metabolic diseases: insights for the internist. Internal and Emergency Medicine, 2018, 13, 313-318.	1.0	4
119	Detection of Urinary Exosomal HSD11B2 mRNA Expression: A Useful Novel Tool for the Diagnostic Approach of Dysfunctional 11Î²-HSD2-Related Hypertension. Frontiers in Endocrinology, 2021, 12, 681974.	1.5	4
120	Randomized placebo-controlled trial comparing fluticasone aqueous nasal spray in mono-therapy, fluticasone plus cetirizine, fluticasone plus montelukast and cetirizine plus montelukast for seasonal allergic rhinitis. Clinical and Experimental Allergy, 2004, 34, 1329-1329.	1.4	3
121	Altered renal folate handling in hypertensive patients with nephroangiosclerotic damage. Journal of Human Hypertension, 2007, 21, 327-329.	1.0	3
122	DISHphagia: An Unusual Cause of Dysphagia. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 2573-2574.	1.8	3
123	Plasma Proteome Profiles of Stable CAD Patients Stratified According to Total Apo Câ€III Levels. Proteomics - Clinical Applications, 2019, 13, e1800023.	0.8	3
124	Serum Uric Acid Levels, but Not rs7442295 Polymorphism of SCL2A9 Gene, Predict Mortality in Clinically Stable Coronary Artery Disease. Current Problems in Cardiology, 2021, 46, 100798.	1.1	3
125	Combination of CLEC4M rs868875 G-Carriership and ABO O Genotypes May Predict Faster Decay of FVIII Infused in Hemophilia A Patients. Journal of Clinical Medicine, 2022, 11, 733.	1.0	3
126	Identification of a Novel Serological Marker in Seronegative Rheumatoid Arthritis Using the Peptide Library Approach. Frontiers in Immunology, 2021, 12, 753400.	2.2	2

#	Article	IF	Citations
127	Acute haemolysis by cold antibody during SARS-CoV-2 infection in a patient with Evans syndrome: a case report and literature review. Blood Transfusion, 2021, , .	0.3	2
128	Nickel sulphate, food additives and seronegative arthritis: is there any relationship?. Nickel sulphates are triggers or aggravating factors for seronegative arthritis. Allergy: European Journal of Allergy and Clinical Immunology, 2003, 58, 958-959.	2.7	1
129	Ã‰valuation de la tempÃ©rature cutanÃ©e des patients atteints de sclÃ©rodermie traitÃ©s pÃ©riodiquement par iloprost. Revue Du Rhumatisme (Edition Francaise), 2006, 73, 53-57.	0.0	1
130	Portal hypertensive biliopathy and bile duct varices presenting as jaundice. Endoscopy, 2021, 53, E442-E443.	1.0	1
131	Trace Elements Status and Metallothioneins DNA Methylation Influence Human Hepatocellular Carcinoma Survival Rate. Frontiers in Oncology, 2020, 10, 596040.	1.3	1
132	Abstract 1229: Clonal hematopoiesis of indeterminate potential (CHIP), centenarians and age-related cardiovascular risk: Is TET2 the culprit. , 2018, , .		1
133	High Plasma Concentration of Apolipoprotein C-III Confers an Increased Risk of Cerebral Ischemic Events on Cardiovascular Patients Anticoagulated With Warfarin. Frontiers in Cardiovascular Medicine, 2021, 8, 781383.	1.1	1
134	Novel protein-truncating variant in the APOB gene may protect from coronary artery disease and adverse cardiovascular events. Atherosclerosis Plus, 2022, 49, 42-46.	0.3	1
135	Infective endocarditis with lung and systemic embolization in an injection drug user. European Heart Journal, 2006, 27, 2938-2938.	1.0	0
136	Two new highly polymorphic markers in the 3' UTR region of the PLA2G7 gene. International Journal of Immunogenetics, 2007, 34, 465-468.	0.8	0
137	Response to Letter Regarding Article, â€œAn Unusual Heart Failure: Cardiac Amyloidosis Due to Light Chain Myelomaâ€• Circulation, 2011, 124, .	1.6	0
138	Apolipoprotein C-III glycoforms correlate heterogeneously with plasma lipid profile in subjects with coronary artery disease. Atherosclerosis, 2016, 252, e100.	0.4	0
139	A dangerous onychodystrophy. American Journal of Hematology, 2021, 96, 891-892.	2.0	0
140	Hyperhomocysteinemia in Relation to Total and Cardiovascular Death after Coronary Artery Bypass Grafting. A Prospective Study.. Blood, 2005, 106, 1640-1640.	0.6	0
141	ALOX5AP Gene Variants and Risk of Coronary Artery Disease in Italy. An Angiography-Based Study.. Blood, 2006, 108, 1459-1459.	0.6	0
142	Hepcidin Levels and Their Determinants In Different Types of Myelodysplastic Syndromes. Blood, 2010, 116, 4250-4250.	0.6	0
143	Global DNA hypomethylation in peripheral blood mononuclear cells as a biomarker of cancer risk. FASEB Journal, 2013, 27, 248.1.	0.2	0
144	26. An 83-year-old Man with Metabolic Syndrome, Coronary Artery Disease, and Dysphagia. , 2013, , 168-171.		0

#	Article	IF	Citations
145	Iron Status Independently Associates With Bone Mineral Density At Population Level. Insights From The Val Borbera Study. Blood, 2013, 122, 4672-4672.	0.6	0
146	Activated Factor VIIâ€“Antithrombin Complex Plasma Concentration Is An Independent Predictor Of Total and Cardiovascular Mortality In Patients With Coronary Artery Disease and Its Prognostic Significance Is Improved By Using Factor VII Genotype-Specific Threshold Levels. Blood, 2013, 122, 2339-2339.	0.6	0
147	The MTHFD1 1958G>A Relates to Survival Rate According to PBMCs DNA Global Methylation in Cancer. FASEB Journal, 2015, 29, 749.3.	0.2	0