

Francois Cambien

List of Publications by Year in descending order

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145
papers

21,953
citations

14655

66
h-index

9345

143
g-index

147
all docs

147
docs citations

147
times ranked

26518
citing authors

#	ARTICLE	IF	CITATIONS
1	Deletion polymorphism in the gene for angiotensin-converting enzyme is a potent risk factor for myocardial infarction. <i>Nature</i> , 1992, 359, 641-644.	27.8	1,880
2	Genomewide Association Analysis of Coronary Artery Disease. <i>New England Journal of Medicine</i> , 2007, 357, 443-453.	27.0	1,865
3	Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease. <i>Nature Genetics</i> , 2011, 43, 333-338.	21.4	1,685
4	Large-scale association analysis identifies new risk loci for coronary artery disease. <i>Nature Genetics</i> , 2013, 45, 25-33.	21.4	1,439
5	Genome-wide association of early-onset myocardial infarction with single nucleotide polymorphisms and copy number variants. <i>Nature Genetics</i> , 2009, 41, 334-341.	21.4	990
6	Plasma Concentrations and Genetic Variation of Matrix Metalloproteinase 9 and Prognosis of Patients With Cardiovascular Disease. <i>Circulation</i> , 2003, 107, 1579-1585.	1.6	692
7	DNA methylation and body-mass index: a genome-wide analysis. <i>Lancet, The</i> , 2014, 383, 1990-1998.	13.7	686
8	Interleukin-6 receptor pathways in coronary heart disease: a collaborative meta-analysis of 82 studies. <i>Lancet, The</i> , 2012, 379, 1205-1213.	13.7	668
9	Glutathione Peroxidase 1 Activity and Cardiovascular Events in Patients with Coronary Artery Disease. <i>New England Journal of Medicine</i> , 2003, 349, 1605-1613.	27.0	548
10	Genetics and Beyond – The Transcriptome of Human Monocytes and Disease Susceptibility. <i>PLoS ONE</i> , 2010, 5, e10693.	2.5	539
11	Interleukin-18 Is a Strong Predictor of Cardiovascular Death in Stable and Unstable Angina. <i>Circulation</i> , 2002, 106, 24-30.	1.6	534
12	New susceptibility locus for coronary artery disease on chromosome 3q22.3. <i>Nature Genetics</i> , 2009, 41, 280-282.	21.4	440
13	Genome-wide haplotype association study identifies the SLC22A3-LPAL2-LPA gene cluster as a risk locus for coronary artery disease. <i>Nature Genetics</i> , 2009, 41, 283-285.	21.4	427
14	Repeated Replication and a Prospective Meta-Analysis of the Association Between Chromosome 9p21.3 and Coronary Artery Disease. <i>Circulation</i> , 2008, 117, 1675-1684.	1.6	356
15	Asymmetric Dimethylarginine and the Risk of Cardiovascular Events and Death in Patients With Coronary Artery Disease. <i>Circulation Research</i> , 2005, 97, e53-9.	4.5	330
16	Interleukin-18 and the Risk of Coronary Heart Disease in European Men. <i>Circulation</i> , 2003, 108, 2453-2459.	1.6	317
17	A genome-wide association study identifies two loci associated with heart failure due to dilated cardiomyopathy. <i>European Heart Journal</i> , 2011, 32, 1065-1076.	2.2	292
18	Cigarette smoking reduces DNA methylation levels at multiple genomic loci but the effect is partially reversible upon cessation. <i>Epigenetics</i> , 2014, 9, 1382-1396.	2.7	285

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19	A trans-acting locus regulates an anti-viral expression network and type 1 diabetes risk. <i>Nature</i> , 2010, 467, 460-464.	27.8	271
20	Genome-wide association study of intracranial aneurysm identifies three new risk loci. <i>Nature Genetics</i> , 2010, 42, 420-425.	21.4	262
21	Comparative Impact of Multiple Biomarkers and N-Terminal Pro-Brain Natriuretic Peptide in the Context of Conventional Risk Factors for the Prediction of Recurrent Cardiovascular Events in the Heart Outcomes Prevention Evaluation (HOPE) Study. <i>Circulation</i> , 2006, 114, 201-208.	1.6	236
22	Meta-analysis of 65,734 Individuals Identifies TSPAN15 and SLC44A2 as Two Susceptibility Loci for Venous Thromboembolism. <i>American Journal of Human Genetics</i> , 2015, 96, 532-542.	6.2	222
23	Extensive association analysis between the CETP gene and coronary heart disease phenotypes reveals several putative functional polymorphisms and gene-environment interaction. <i>Genetic Epidemiology</i> , 2000, 19, 64-80.	1.3	205
24	Genetic Analysis of the Interleukin-18 System Highlights the Role of the Interleukin-18 Gene in Cardiovascular Disease. <i>Circulation</i> , 2005, 112, 643-650.	1.6	205
25	Inheritance of coronary artery disease in men: an analysis of the role of the Y chromosome. <i>Lancet, The</i> , 2012, 379, 915-922.	13.7	179
26	The Gln/Arg polymorphism of human paraoxonase (PON 192) is not related to myocardial infarction in the ECTIM Study. <i>Atherosclerosis</i> , 1996, 126, 299-303.	0.8	163
27	Specific haplotypes of the P-selectin gene are associated with myocardial infarction. <i>Human Molecular Genetics</i> , 2002, 11, 2015-2023.	2.9	161
28	Nitric oxide synthase gene polymorphisms, blood pressure and aortic stiffness in normotensive and hypertensive subjects. <i>Journal of Hypertension</i> , 1998, 16, 31-35.	0.5	155
29	Interleukin-6 gene polymorphisms and susceptibility to myocardial infarction: the ECTIM study. <i>Journal of Molecular Medicine</i> , 2001, 79, 300-305.	3.9	155
30	Sequence Diversity in 36 Candidate Genes for Cardiovascular Disorders. <i>American Journal of Human Genetics</i> , 1999, 65, 183-191.	6.2	152
31	Genetic Regulation of Serum Phytosterol Levels and Risk of Coronary Artery Disease. <i>Circulation: Cardiovascular Genetics</i> , 2010, 3, 331-339.	5.1	141
32	A Genome-Wide Association Study Identifies <i>LIPA</i> as a Susceptibility Gene for Coronary Artery Disease. <i>Circulation: Cardiovascular Genetics</i> , 2011, 4, 403-412.	5.1	130
33	The ACE Gene <i>I/D</i> Polymorphism Is Not Associated With the Blood Pressure and Cardiovascular Benefits of ACE Inhibition. <i>Hypertension</i> , 2003, 42, 297-303.	2.7	129
34	Genetics of Venous Thrombosis: Insights from a New Genome Wide Association Study. <i>PLoS ONE</i> , 2011, 6, e25581.	2.5	127
35	Integrating Genome-Wide Genetic Variations and Monocyte Expression Data Reveals Trans-Regulated Gene Modules in Humans. <i>PLoS Genetics</i> , 2011, 7, e1002367.	3.5	126
36	Human CalDAG-GEF1 gene (<i>RASGRP2</i>) mutation affects platelet function and causes severe bleeding. <i>Journal of Experimental Medicine</i> , 2014, 211, 1349-1362.	8.5	117

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37	Gene polymorphisms of the renin-angiotensin system in relation to hypertension and parental history of myocardial infarction and stroke. <i>Journal of Hypertension</i> , 1998, 16, 37-44.	0.5	114
38	Genetic Association Study Identifies HSPB7 as a Risk Gene for Idiopathic Dilated Cardiomyopathy. <i>PLoS Genetics</i> , 2010, 6, e1001167.	3.5	110
39	Angiotensin II type 1 receptor $\alpha 1$ 153A/G and 1166A/C gene polymorphisms and increase in aortic stiffness with age in hypertensive subjects. <i>Journal of Hypertension</i> , 2001, 19, 407-413.	0.5	108
40	MORGAM (an international pooling of cardiovascular cohorts). <i>International Journal of Epidemiology</i> , 2004, 34, 21-27.	1.9	105
41	Characterization of a Unique Genetic Variant in the $\beta 1$ -adrenoceptor Gene and Evaluation of its Role in Idiopathic Dilated Cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , 1999, 31, 1025-1032.	1.9	103
42	Glutathione Peroxidase-1 and Homocysteine for Cardiovascular Risk Prediction. <i>Journal of the American College of Cardiology</i> , 2005, 45, 1631-1637.	2.8	103
43	Statistical colocalization of monocyte gene expression and genetic risk variants for type 1 diabetes. <i>Human Molecular Genetics</i> , 2012, 21, 2815-2824.	2.9	103
44	Impact of pathogen burden in patients with coronary artery disease in relation to systemic inflammation and variation in genes encoding cytokines. <i>American Journal of Cardiology</i> , 2003, 92, 515-521.	1.6	100
45	Common variant near the endothelin receptor type A (<i>EDNRA</i>) gene is associated with intracranial aneurysm risk. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 19707-19712.	7.1	100
46	Long-range DNA looping and gene expression analyses identify DEXI as an autoimmune disease candidate gene. <i>Human Molecular Genetics</i> , 2012, 21, 322-333.	2.9	100
47	B-Type Natriuretic Peptide and the Risk of Cardiovascular Events and Death in Patients With Stable Angina. <i>Journal of the American College of Cardiology</i> , 2006, 47, 552-558.	2.8	99
48	Multiple marker approach to risk stratification in patients with stable coronary artery disease. <i>European Heart Journal</i> , 2010, 31, 3024-3031.	2.2	97
49	In-Depth Haplotype Analysis of ABCA1 Gene Polymorphisms in Relation to Plasma ApoA1 Levels and Myocardial Infarction. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2004, 24, 775-781.	2.4	96
50	Exome-wide association study reveals novel susceptibility genes to sporadic dilated cardiomyopathy. <i>PLoS ONE</i> , 2017, 12, e0172995.	2.5	92
51	Angiotensin converting enzyme gene polymorphism and ACE inhibition in diabetic nephropathy. <i>Kidney International</i> , 1998, 53, 1002-1006.	5.2	91
52	Platelet-activating factor-acetylhydrolase and PAF-receptor gene haplotypes in relation to future cardiovascular event in patients with coronary artery disease. <i>Human Molecular Genetics</i> , 2004, 13, 1341-1351.	2.9	91
53	Analysis of N-terminal-pro-brain natriuretic peptide and C-reactive protein for risk stratification in stable and unstable coronary artery disease: results from the AtheroGene study. <i>European Heart Journal</i> , 2005, 26, 241-249.	2.2	90
54	Genome-wide association study for circulating levels of PAI-1 provides novel insights into its regulation. <i>Blood</i> , 2012, 120, 4873-4881.	1.4	90

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55	Common genetic variation of the cholesteryl ester transfer protein gene strongly predicts future cardiovascular death in patients with coronary artery disease. <i>Journal of the American College of Cardiology</i> , 2003, 41, 1983-1989.	2.8	89
56	Genetics of Cardiovascular Diseases. <i>Circulation</i> , 2007, 116, 1714-1724.	1.6	86
57	Lack of Association Between the Trp719Arg Polymorphism in Kinesin-Like Protein-6 and Coronary Artery Disease in 19 Case-Control Studies. <i>Journal of the American College of Cardiology</i> , 2010, 56, 1552-1563.	2.8	84
58	Physical Activity May Modulate Effects of ApoE Genotype on Lipid Profile. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2002, 22, 133-140.	2.4	82
59	Progression of diabetic nephropathy in normotensive type 1 diabetic patients. <i>Kidney International</i> , 1999, 56, S101-S105.	5.2	80
60	Cardiovascular risk factors and alcohol consumption in France and Northern Ireland. <i>Atherosclerosis</i> , 1995, 115, 225-232.	0.8	78
61	CORONARY HEART DISEASE IN MIDDLE-AGED FRENCHMEN. <i>Lancet, The</i> , 1980, 315, 1346-1350.	13.7	77
62	The impact of newly identified loci on coronary heart disease, stroke and total mortality in the MORGAM prospective cohorts. <i>Genetic Epidemiology</i> , 2009, 33, 237-246.	1.3	77
63	Lack of association between polymorphisms of eight candidate genes and idiopathic dilated cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2000, 35, 29-35.	2.8	74
64	Analysis of 14 Candidate Genes for Diabetic Nephropathy on Chromosome 3q in European Populations. <i>Diabetes</i> , 2006, 55, 3166-3174.	0.6	74
65	New polymorphisms of the angiotensin II type 1 receptor gene and their associations with myocardial infarction and blood pressure. <i>Journal of Hypertension</i> , 1998, 16, 1443-1447.	0.5	72
66	Bayesian Detection of Expression Quantitative Trait Loci Hot Spots. <i>Genetics</i> , 2011, 189, 1449-1459.	2.9	70
67	Natural genetic variation of the cardiac transcriptome in non-diseased donors and patients with dilated cardiomyopathy. <i>Genome Biology</i> , 2017, 18, 170.	8.8	70
68	Prognostic value of tissue inhibitor of metalloproteinase-1 for cardiovascular death among patients with cardiovascular disease: results from the AtheroGene study. <i>European Heart Journal</i> , 2006, 27, 150-156.	2.2	69
69	Polymorphisms of the β_2 -adrenoceptor (ADRB2) gene and essential hypertension: the ECTIM and PEGASE studies. <i>Journal of Hypertension</i> , 2002, 20, 229-235.	0.5	65
70	Long-Term Renoprotective Effects of Losartan in Diabetic Nephropathy. <i>Diabetes Care</i> , 2003, 26, 1501-1506.	8.6	60
71	Polymorphisms of the endothelin-A and -B receptor genes in relation to blood pressure and myocardial infarction The etude cas-tmoins sur l'infarctus du myocarde (ECTIM) study. <i>American Journal of Hypertension</i> , 1999, 12, 304-310.	2.0	59
72	Cytokine Polymorphisms Associated With Carotid Intima-Media Thickness in Stroke Patients. <i>Stroke</i> , 2006, 37, 1691-1696.	2.0	59

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73	GUESS-ing Polygenic Associations with Multiple Phenotypes Using a GPU-Based Evolutionary Stochastic Search Algorithm. <i>PLoS Genetics</i> , 2013, 9, e1003657.	3.5	58
74	A Follow-Up Study of a Genome-wide Association Scan Identifies a Susceptibility Locus for Venous Thrombosis on Chromosome 6p24.1. <i>American Journal of Human Genetics</i> , 2010, 86, 592-595.	6.2	57
75	Sequence polymorphisms in the apolipoprotein(a) gene and their association with lipoprotein(a) levels and myocardial infarction. The ECTIM Study. <i>Atherosclerosis</i> , 1999, 144, 323-333.	0.8	55
76	Heterogeneity of linkage disequilibrium in human genes has implications for association studies of common diseases. <i>Human Molecular Genetics</i> , 2002, 11, 419-429.	2.9	54
77	Genome-Wide Haplotype Analysis of Cis Expression Quantitative Trait Loci in Monocytes. <i>PLoS Genetics</i> , 2013, 9, e1003240.	3.5	53
78	Endothelin gene variants and aortic and cardiac structure in never-treated hypertensives. <i>American Journal of Hypertension</i> , 2001, 14, 755-760.	2.0	50
79	Genome-wide association analysis in dilated cardiomyopathy reveals two new players in systolic heart failure on chromosomes 3p25.1 and 22q11.23. <i>European Heart Journal</i> , 2021, 42, 2000-2011.	2.2	49
80	A pharmacogenetic analysis of determinants of hypertension and blood pressure response to angiotensin-converting enzyme inhibitor therapy in patients with vascular disease and healthy individuals. <i>Journal of Hypertension</i> , 2011, 29, 509-519.	0.5	47
81	The angiotensin-converting enzyme (ACE) genetic polymorphism: its relationship with plasma ACE level and myocardial infarction. <i>Clinical Genetics</i> , 1994, 46, 94-101.	2.0	46
82	Polymorphism R92Q of the tumour necrosis factor receptor 1 gene is associated with myocardial infarction and carotid intima-media thickness – The ECTIM, AXA, EVA and GENIC Studies. <i>European Journal of Human Genetics</i> , 2004, 12, 213-219.	2.8	45
83	Adverse Associations Between CX3CR1 Polymorphisms and Risk of Cardiovascular or Cerebrovascular Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2005, 25, 847-853.	2.4	44
84	Genome-Wide Association Study for Circulating Tissue Plasminogen Activator Levels and Functional Follow-Up Implicates Endothelial <i>STXBP5</i> and <i>STX2</i> . <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2014, 34, 1093-1101.	2.4	43
85	High expressor paraoxonase PON1 gene promoter polymorphisms are associated with reduced risk of vascular disease in younger coronary patients. <i>Atherosclerosis</i> , 2002, 161, 463-467.	0.8	40
86	Renin-angiotensin-aldosterone system in brain infarction and vascular death. <i>Annals of Neurology</i> , 2005, 58, 131-138.	5.3	40
87	Gene Polymorphisms of the Renin-Angiotensin System and Early Development of Hypertension. <i>American Journal of Hypertension</i> , 2006, 19, 837-842.	2.0	40
88	G/T Substitution in Intron 1 of the UNC13B Gene Is Associated With Increased Risk of Nephropathy in Patients With Type 1 Diabetes. <i>Diabetes</i> , 2008, 57, 2843-2850.	0.6	39
89	Automated Detection of Informative Combined Effects in Genetic Association Studies of Complex Traits. <i>Genome Research</i> , 2003, 13, 1952-1960.	5.5	38
90	Family history, longevity, and risk of coronary heart disease: the PRIME Study. <i>International Journal of Epidemiology</i> , 2003, 32, 71-77.	1.9	37

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91	Runs of Homozygosity: Association with Coronary Artery Disease and Gene Expression in Monocytes and Macrophages. <i>American Journal of Human Genetics</i> , 2015, 97, 228-237.	6.2	37
92	Graphical Modeling of Gene Expression in Monocytes Suggests Molecular Mechanisms Explaining Increased Atherosclerosis in Smokers. <i>PLoS ONE</i> , 2013, 8, e50888.	2.5	36
93	Powerful Identification of Cis-regulatory SNPs in Human Primary Monocytes Using Allele-Specific Gene Expression. <i>PLoS ONE</i> , 2012, 7, e52260.	2.5	36
94	The renin-angiotensin-aldosterone system in cerebral small vessel disease. <i>Journal of Neurology</i> , 2008, 255, 993-1000.	3.6	31
95	Renoprotective effects of losartan in diabetic nephropathy: Interaction with ACE insertion/deletion genotype?. <i>Kidney International</i> , 2002, 62, 192-198.	5.2	30
96	European rational approach for the genetics of diabetic complications EURAGEDIC: patient populations and strategy. <i>Nephrology Dialysis Transplantation</i> , 2007, 23, 161-168.	0.7	30
97	Differential haplotypic expression of the interleukin-18 gene. <i>European Journal of Human Genetics</i> , 2007, 15, 856-863.	2.8	29
98	Four Genetic Loci Influencing Electrocardiographic Indices of Left Ventricular Hypertrophy. Circulation: <i>Cardiovascular Genetics</i> , 2011, 4, 626-635.	5.1	28
99	Polymorphisms of genes of the cardiac calcineurin pathway and cardiac hypertrophy. <i>European Journal of Human Genetics</i> , 2003, 11, 659-664.	2.8	27
100	A meta-analysis of genome-wide association studies identifies ORM1 as a novel gene controlling thrombin generation potential. <i>Blood</i> , 2014, 123, 777-785.	1.4	27
101	Haplotypes of the Caspase-1 Gene, Plasma Caspase-1 Levels, and Cardiovascular Risk. <i>Circulation Research</i> , 2006, 99, 102-108.	4.5	26
102	Identification of two polymorphisms in the promoter of the microsomal triglyceride transfer protein (MTP) gene: lack of association with lipoprotein profiles. <i>Journal of Lipid Research</i> , 1998, 39, 2432-2435.	4.2	26
103	Role of lipid phosphate phosphatase 3 in human aortic endothelial cell function. <i>Cardiovascular Research</i> , 2016, 112, 702-713.	3.8	25
104	SASH1, a new potential link between smoking and atherosclerosis. <i>Atherosclerosis</i> , 2015, 242, 571-579.	0.8	24
105	A polymorphism in the endothelin-A receptor gene is linked to baroreflex sensitivity. <i>Journal of Hypertension</i> , 2005, 23, 2019-2026.	0.5	23
106	Influence of sex and genetic variability on expression of X-linked genes in human monocytes. <i>Genomics</i> , 2011, 98, 320-326.	2.9	23
107	The Choice of the Filtering Method in Microarrays Affects the Inference Regarding Dosage Compensation of the Active X-Chromosome. <i>PLoS ONE</i> , 2011, 6, e23956.	2.5	23
108	Allelic expression mapping across cellular lineages to establish impact of non-coding <sc>SNP</sc>s. <i>Molecular Systems Biology</i> , 2014, 10, 754.	7.2	21

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109	Contribution of Rare and Common Genetic Variants to Plasma Lipid Levels and Carotid Stiffness and Geometry. <i>Circulation: Cardiovascular Genetics</i> , 2015, 8, 628-636.	5.1	21
110	Coronary heart disease and genetics an epidemiologist's view. <i>Trends in Molecular Medicine</i> , 1997, 3, 197-203.	2.6	20
111	Lack of association between complement factor H polymorphisms and coronary artery disease or myocardial infarction. <i>Journal of Molecular Medicine</i> , 2007, 85, 771-775.	3.9	19
112	Molecular genetic analysis of a human insulin-like growth factor 1 promoter P1 variation. <i>FASEB Journal</i> , 2009, 23, 1303-1313.	0.5	19
113	Coronary heart disease and polymorphisms in genes affecting lipid metabolism and inflammation. <i>Current Atherosclerosis Reports</i> , 2005, 7, 188-195.	4.8	18
114	Lack of association between polymorphisms of the IL18R1 and IL18RAP genes and cardiovascular risk: the MORGAM Project. <i>BMC Medical Genetics</i> , 2009, 10, 44.	2.1	18
115	No Physical Activity ?? CETP 1b.-629 Interaction Effects on Lipid Profile. <i>Medicine and Science in Sports and Exercise</i> , 2003, 35, 1124-1129.	0.4	17
116	Caution in Interpreting Results from Imputation Analysis When Linkage Disequilibrium Extends over a Large Distance: A Case Study on Venous Thrombosis. <i>PLoS ONE</i> , 2012, 7, e38538.	2.5	17
117	Distribution of apolipoprotein E between apo B- and non apo B-containing lipoproteins according to apo E phenotype. <i>Atherosclerosis</i> , 1997, 131, 257-262.	0.8	16
118	Molecular and functional characterization of polymorphisms in the secreted phospholipase A2 group X gene: relevance to coronary artery disease. <i>Journal of Molecular Medicine</i> , 2009, 87, 723-733.	3.9	16
119	Preservation Analysis of Macrophage Gene Coexpression Between Human and Mouse Identifies PARK2 as a Genetically Controlled Master Regulator of Oxidative Phosphorylation in Humans. <i>G3: Genes, Genomes, Genetics</i> , 2016, 6, 3361-3371.	1.8	15
120	Identification of two polymorphisms in the early growth response protein-1 gene: possible association with lipid variables. <i>Journal of Molecular Medicine</i> , 2000, 78, 81-86.	3.9	14
121	Cytokine phenotype, genotype, and renal outcomes at cardiac surgery. <i>Cytokine</i> , 2013, 61, 275-284.	3.2	14
122	Automated detection of informative combined effects in genetic association studies of complex traits. <i>Genome Research</i> , 2003, 13, 1952-60.	5.5	14
123	Atherosclerosis: From Genetic Polymorphisms to System Genetics. <i>Cardiovascular Toxicology</i> , 2005, 5, 143-152.	2.7	12
124	Haplotypic analysis of tag SNPs of the interleukin-18 gene in relation to cardiovascular disease events: the MORGAM Project. <i>European Journal of Human Genetics</i> , 2008, 16, 1512-1520.	2.8	12
125	Adrenomedullin and Arterial Stiffness. <i>Circulation: Cardiovascular Genetics</i> , 2014, 7, 634-641.	5.1	12
126	SAH gene variants are associated with obesity-related hypertension in Caucasians: the PEGASE Study. <i>Journal of Hypertension</i> , 2007, 25, 557-564.	0.5	11

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127	Polymorphisms in the genes encoding platelet-derived growth factor A and α_2 receptor. <i>Journal of Molecular Medicine</i> , 2000, 78, 287-292.	3.9	10
128	Influence of cholesteryl ester transfer protein, peroxisome proliferator-activated receptor α , apolipoprotein E, and apolipoprotein A-I polymorphisms on high-density lipoprotein cholesterol, apolipoprotein A-I, lipoprotein A-I, and lipoprotein A-I:A-II concentrations: the Prospective Epidemiological Study of Myocardial Infarction study. <i>Metabolism: Clinical and Experimental</i> , 2009, 58, 283-289.	3.4	9
129	Osteopontin gene variation and cardio/cerebrovascular disease phenotypes. <i>Atherosclerosis</i> , 2009, 206, 209-215.	0.8	9
130	Single Nucleotide Polymorphisms with Cis-Regulatory Effects on Long Non-Coding Transcripts in Human Primary Monocytes. <i>PLoS ONE</i> , 2014, 9, e102612.	2.5	9
131	Renin-angiotensin system genes as candidate genes in cardiovascular diseases. <i>Trends in Cardiovascular Medicine</i> , 1993, 3, 250-258.	4.9	8
132	P-selectin gene polymorphisms and risk of coronary heart disease among Tunisians. <i>Journal of Thrombosis and Thrombolysis</i> , 2009, 28, 314-319.	2.1	8
133	Comprehensive Exploration of the Effects of miRNA SNPs on Monocyte Gene Expression. <i>PLoS ONE</i> , 2012, 7, e45863.	2.5	8
134	Identification and Functional Analyses of Molecular Haplotypes of the Human Osteoprotegerin Gene Promoter. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2009, 29, 1638-1643.	2.4	7
135	Patterns of Alcohol Consumption and Cardiovascular Risk in Northern Ireland and France. <i>Annals of Epidemiology</i> , 2007, 17, S75-S80.	1.9	6
136	Polymorphisms in 33 inflammatory genes and risk of myocardial infarction—a system genetics approach. <i>Journal of Molecular Medicine</i> , 2007, 85, 1271-1280.	3.9	6
137	Lack of association of genetic variants in the LRP8 gene with familial and sporadic myocardial infarction. <i>Journal of Molecular Medicine</i> , 2008, 86, 1163-1170.	3.9	6
138	Association of three polymorphisms selected from a genome-wide association study with coronary heart disease in the Tunisian population. <i>Journal of Thrombosis and Thrombolysis</i> , 2010, 29, 114-118.	2.1	6
139	Heritability, Weak Effects, and Rare Variants in Genomewide Association Studies. <i>Clinical Chemistry</i> , 2011, 57, 1263-1266.	3.2	5
140	Molecular investigation of the functional relevance of missense variants of ICAM-1. <i>Pharmacogenetics and Genomics</i> , 2008, 18, 1017-1019.	1.5	3
141	Neutrophil elastase gene variation and coronary heart disease. <i>Pharmacogenetics and Genomics</i> , 2007, 17, 629-637.	1.5	2
142	Coronary heart disease aetiology: associations and causality. <i>Comptes Rendus - Biologies</i> , 2007, 330, 299-305.	0.2	1
143	Effects of Alcohol on Lipids and Lipoprotein Metabolism. <i>Medical Science Symposia Series</i> , 2000, , 11-22.	0.0	0
144	The genetic contribution to the onset of acute coronary heart disease. <i>Developments in Cardiovascular Medicine</i> , 1996, , 111-133.	0.1	0

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145	First case of a human <i>RASGRP2</i> mutation affecting Rap1 activation in platelets and causing severe bleeding.. Journal of Cell Biology, 2014, 206, 2061OIA111.	5.2	0