

Sophie Visvikis-Siest

List of Publications by Year in descending order

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Version: 2024-02-01

290
papers

34,390
citations

22153

59
h-index

4015

176
g-index

313
all docs

313
docs citations

313
times ranked

51587
citing authors

#	ARTICLE	IF	CITATIONS
1	Integrating polymorphism signatures with myocardial perfusion imaging data to improve the prevention of coronary artery disease: Science or science-fiction?. <i>Journal of Nuclear Cardiology</i> , 2022, 29, 2917-2919.	2.1	0
2	The association of vascular endothelial growth factor related SNPs and circulating iron levels might depend on body mass index. <i>Frontiers in Bioscience</i> , 2022, 27, 1.	2.1	1
3	Association of Dietary Patterns with MRI Markers of Hepatic Inflammation and Fibrosis in the MAST4HEALTH Study. <i>International Journal of Environmental Research and Public Health</i> , 2022, 19, 971.	2.6	2
4	VEGF-A-related genetic variants protect against Alzheimer's disease. <i>Aging</i> , 2022, 14, 2524-2536.	3.1	10
5	Sex-dimorphic genetic effects and novel loci for fasting glucose and insulin variability. <i>Nature Communications</i> , 2021, 12, 24.	12.8	87
6	Genome-wide association study of circulating interleukin 6 levels identifies novel loci. <i>Human Molecular Genetics</i> , 2021, 30, 393-409.	2.9	32
7	Heterogeneous contributions of change in population distribution of body mass index to change in obesity and underweight. <i>ELife</i> , 2021, 10, .	6.0	41
8	Effect of Mastiha supplementation on NAFLD: The MAST4HEALTH Randomised, Controlled Trial. <i>Molecular Nutrition and Food Research</i> , 2021, 65, e2001178.	3.3	19
9	Nutrigenetic Interactions Might Modulate the Antioxidant and Anti-Inflammatory Status in Mastiha-Supplemented Patients With NAFLD. <i>Frontiers in Immunology</i> , 2021, 12, 683028.	4.8	12
10	Epigenome-wide association study detects a novel loci associated with central obesity in healthy subjects. <i>BMC Medical Genomics</i> , 2021, 14, 233.	1.5	3
11	Worldwide trends in hypertension prevalence and progress in treatment and control from 1990 to 2019: a pooled analysis of 1201 population-representative studies with 104 million participants. <i>Lancet</i> , The, 2021, 398, 957-980.	13.7	1,289
12	Dietary Patterns, Blood Pressure and the Glycemic and Lipidemic Profile of Two Teenage, European Populations. <i>Nutrients</i> , 2021, 13, 198.	4.1	6
13	A genetic determinant of VEGF-A levels is associated with telomere attrition. <i>Aging</i> , 2021, 13, 23517-23526.	3.1	2
14	National trends in total cholesterol obscure heterogeneous changes in HDL and non-HDL cholesterol and total-to-HDL cholesterol ratio: a pooled analysis of 458 population-based studies in Asian and Western countries. <i>International Journal of Epidemiology</i> , 2020, 49, 173-192.	1.9	44
15	Telomere length determinants in childhood. <i>Clinical Chemistry and Laboratory Medicine</i> , 2020, 58, 162-177.	2.3	33
16	Milestones in Personalized Medicine: From the Ancient Time to Nowadays—the Provocation of COVID-19. <i>Frontiers in Genetics</i> , 2020, 11, 569175.	2.3	17
17	Height and body-mass index trajectories of school-aged children and adolescents from 1985 to 2019 in 200 countries and territories: a pooled analysis of 2181 population-based studies with 65 million participants. <i>Lancet</i> , The, 2020, 396, 1511-1524.	13.7	219
18	Repositioning of the global epicentre of non-optimal cholesterol. <i>Nature</i> , 2020, 582, 73-77.	27.8	138

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19	Epigenome-wide association study in healthy individuals identifies significant associations with DNA methylation and PBMC extract VEGF-A concentration. <i>Clinical Epigenetics</i> , 2020, 12, 79.	4.1	4
20	Increased risk of hypercholesterolemia in a French and Lebanese population due to an interaction between rs2569190 in CD14 and gender. <i>Clinica Chimica Acta</i> , 2020, 509, 172-176.	1.1	1
21	TERC Variants Associated with Short Leukocyte Telomeres: Implication of Higher Early Life Leukocyte Telomere Attrition as Assessed by the Blood-and-Muscle Model. <i>Cells</i> , 2020, 9, 1360.	4.1	3
22	Obesity status modifies the association between rs7556897T>C in the intergenic region SLC19A3-CCL20 and blood pressure in French children. <i>Clinical Chemistry and Laboratory Medicine</i> , 2020, 58, 1819-1827.	2.3	3
23	Peripheral blood mononuclear cells extracts VEGF protein levels and VEGF mRNA: Associations with inflammatory molecules in a healthy population. <i>PLoS ONE</i> , 2019, 14, e0220902.	2.5	7
24	Rising rural body-mass index is the main driver of the global obesity epidemic in adults. <i>Nature</i> , 2019, 569, 260-264.	27.8	469
25	Genetic determinants of circulating haptoglobin concentration. <i>Clinica Chimica Acta</i> , 2019, 494, 138-142.	1.1	14
26	Epigenome-Wide Association Study (EWAS) of Blood Lipids in Healthy Population from STANISLAS Family Study (SFS). <i>International Journal of Molecular Sciences</i> , 2019, 20, 1014.	4.1	17
27	Do we need diagnostic strategies enhanced with genetic information for ischemic heart disease?. <i>Journal of Nuclear Cardiology</i> , 2019, 26, 1309-1312.	2.1	1
28	Pleiotropy of ABO gene: correlation of rs644234 with E-selectin and lipid levels. <i>Clinical Chemistry and Laboratory Medicine</i> , 2018, 56, 748-754.	2.3	8
29	Contributions of mean and shape of blood pressure distribution to worldwide trends and variations in raised blood pressure: a pooled analysis of 1018 population-based measurement studies with 88.6 million participants. <i>International Journal of Epidemiology</i> , 2018, 47, 872-883i.	1.9	65
30	The Relationship Between Vascular Endothelial Growth Factor Cis- and Trans-Acting Genetic Variants and Metabolic Syndrome. <i>American Journal of the Medical Sciences</i> , 2018, 355, 559-565.	1.1	8
31	Effect of SLCO1B1 gene polymorphisms and vitamin D on statin-induced myopathy. <i>Drug Metabolism and Personalized Therapy</i> , 2018, 33, 41-47.	0.6	5
32	Effect of <i>LSR</i> polymorphism on blood lipid levels and age-specific epistatic interaction with the <i>APOE</i> common polymorphism. <i>Clinical Genetics</i> , 2018, 93, 846-852.	2.0	5
33	A transnational collaborative network dedicated to the study and applications of the vascular endothelial growth factor-A in medical practice: the VEGF Consortium. <i>Clinical Chemistry and Laboratory Medicine</i> , 2018, 56, 83-86.	2.3	1
34	VEGF-A is associated with early degenerative changes in cartilage and subchondral bone. <i>Growth Factors</i> , 2018, 36, 263-273.	1.7	14
35	Personalised Medicine: The Odyssey from Hope to Practice. <i>Journal of Personalized Medicine</i> , 2018, 8, 31.	2.5	9
36	Genome Analyses of >200,000 Individuals Identify 58 Loci for Chronic Inflammation and Highlight Pathways that Link Inflammation and Complex Disorders. <i>American Journal of Human Genetics</i> , 2018, 103, 691-706.	6.2	326

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37	The future of telomere length in personalized medicine. <i>Frontiers in Bioscience - Landmark</i> , 2018, 23, 1628-1654.	3.0	33
38	The polymorphism rs6918289 located in the downstream region of the TREM2 gene is associated with TNF- α levels and IMT-F. <i>Scientific Reports</i> , 2018, 8, 7160.	3.3	3
39	Epistatic interaction of apolipoprotein E and lipolysis-stimulated lipoprotein receptor genetic variants is associated with Alzheimer's disease. <i>Neurobiology of Aging</i> , 2018, 69, 292.e1-292.e5.	3.1	4
40	VEGF-related polymorphisms identified by GWAS and risk for major depression. <i>Translational Psychiatry</i> , 2017, 7, e1055-e1055.	4.8	34
41	VEGF, the underlying factor for metabolic syndrome; fact or fiction?. <i>Diabetes and Metabolic Syndrome: Clinical Research and Reviews</i> , 2017, 11, S61-S64.	3.6	30
42	Worldwide trends in body-mass index, underweight, overweight, and obesity from 1975 to 2016: a pooled analysis of 2416 population-based measurement studies in 128.9 million children, adolescents, and adults. <i>Lancet, The</i> , 2017, 390, 2627-2642.	13.7	5,010
43	Next generation sequencing and immuno-histochemistry profiling identify numerous biomarkers for personalized therapy of endometrioid endometrial carcinoma. <i>Clinical Chemistry and Laboratory Medicine</i> , 2017, 56, e19-e22.	2.3	1
44	IL6R haplotype rs4845625*T/rs4537545*C is a risk factor for simultaneously high CRP, LDL and ApoB levels. <i>Genes and Immunity</i> , 2017, 18, 163-169.	4.1	7
45	Worldwide trends in blood pressure from 1975 to 2015: a pooled analysis of 1479 population-based measurement studies with 19.1 million participants. <i>Lancet, The</i> , 2017, 389, 37-55.	13.7	1,667
46	Pharmacogenomic Challenges in Cardiovascular Diseases: Examples of Drugs and Considerations for Future Integration in Clinical Practice. <i>Current Pharmaceutical Biotechnology</i> , 2017, 18, 231-241.	1.6	8
47	TREM-1 SNP rs2234246 regulates TREM-1 protein and mRNA levels and is associated with plasma levels of L-selectin. <i>PLoS ONE</i> , 2017, 12, e0182226.	2.5	31
48	Cardiovascular diseases-related GNB3 C825T polymorphism has a significant sex-specific effect on serum soluble E-selectin levels. <i>Journal of Inflammation</i> , 2016, 13, 39.	3.4	5
49	Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4.4 million participants. <i>Lancet, The</i> , 2016, 387, 1513-1530.	13.7	2,842
50	Trends in adult body-mass index in 200 countries from 1975 to 2014: a pooled analysis of 1698 population-based measurement studies with 19.2 million participants. <i>Lancet, The</i> , 2016, 387, 1377-1396.	13.7	3,941
51	Pro- and anti-angiogenic VEGF mRNAs in autoimmune thyroid diseases. <i>Autoimmunity</i> , 2016, 49, 366-372.	2.6	6
52	Plasma VEGF-related polymorphisms are implicated in autoimmune thyroid diseases. <i>Autoimmunity</i> , 2016, 49, 229-235.	2.6	6
53	Six Novel Loci Associated with Circulating VEGF Levels Identified by a Meta-analysis of Genome-Wide Association Studies. <i>PLoS Genetics</i> , 2016, 12, e1005874.	3.5	56
54	De lâ€™ α -cog α n α tique α la pharmacog α nomique par le stress oxydant. <i>HEGEL - HEpato-GastroEntαrologie, Libαrale</i> , 2016, N α o 2, 217a-218.	0,0	0

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55	Le Professeur Gérard Siest nous a quittés (1936-2016). HEGEL - Hépatogastroentérologie Libérale, 2016, N° 2, 96-97.	0,0	0
56	Angiogenesis related genes NOS3, CD14, MMP3 and IL4R are associated to VEGF gene expression and circulating levels in healthy adults. BMC Medical Genetics, 2015, 16, 90.	2.1	27
57	Beyond genome-wide association studies: identifying variants using -omics approaches. Personalized Medicine, 2015, 12, 529-531.	1.5	0
58	Effects of diabetes definition on global surveillance of diabetes prevalence and diagnosis: a pooled analysis of 96 population-based studies with 331 288 participants. Lancet Diabetes and Endocrinology, 2015, 3, 624-637.	11.4	139
59	Genetic Determinants of Leucocyte Telomere Length in Children: a Neglected and Challenging Field. Paediatric and Perinatal Epidemiology, 2015, 29, 146-150.	1.7	10
60	Genetic determined low response to thienopyridines is associated with higher systemic inflammation in smokers. Pharmacogenomics, 2015, 16, 459-469.	1.3	0
61	Public Biobanks: Calculation and Recovery of Costs. Science Translational Medicine, 2014, 6, 261fs45.	12.4	47
62	Conference Scene: Pharmacogenomics: from cell to clinic (Part 2). Pharmacogenomics, 2014, 15, 739-744.	1.3	1
63	Influence of inflammation on cardiovascular protective effects of cytochrome P450 epoxygenase-derived epoxyeicosatrienoic acids. Drug Metabolism Reviews, 2014, 46, 33-56.	3.6	24
64	Low copy number of the salivary amylase gene predisposes to obesity. Nature Genetics, 2014, 46, 492-497.	21.4	214
65	Human cytochrome P450 epoxygenases: Variability in expression and role in inflammation-related disorders. , 2014, 144, 134-161.		74
66	Epistatic study reveals two genetic interactions in blood pressure regulation. BMC Medical Genetics, 2013, 14, 2.	2.1	13
67	Associations of vascular endothelial growth factor (VEGF) with adhesion and inflammation molecules in a healthy population. Cytokine, 2013, 61, 602-607.	3.2	24
68	Dairy product consumption, calcium intakes, and metabolic syndrome-related factors over 5 years in the STANISLAS study. Nutrition, 2013, 29, 519-524.	2.4	60
69	Relationship between catalase haplotype and arterial aging. Atherosclerosis, 2013, 227, 100-105.	0.8	14
70	Newly identified synergy between clopidogrel and calcium-channel blockers for blood pressure regulation possibly involves CYP2C19 rs4244285. International Journal of Cardiology, 2013, 168, 3057-3058.	1.7	2
71	A common variant highly associated with plasma VEGFA levels also contributes to the variation of both LDL-C and HDL-C. Journal of Lipid Research, 2013, 54, 535-541.	4.2	28
72	Causal Relationship between Obesity and Vitamin D Status: Bi-Directional Mendelian Randomization Analysis of Multiple Cohorts. PLoS Medicine, 2013, 10, e1001383.	8.4	753

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73	Mining the Human Phenome Using Allelic Scores That Index Biological Intermediates. <i>PLoS Genetics</i> , 2013, 9, e1003919.	3.5	84
74	Influence of Genetic Variations on Levels of Inflammatory Markers of Healthy Subjects at Baseline and One Week after Clopidogrel Therapy; Results of a Preliminary Study. <i>International Journal of Molecular Sciences</i> , 2013, 14, 16402-16413.	4.1	1
75	What Is the Contribution of Two Genetic Variants Regulating VEGF Levels to Type 2 Diabetes Risk and to Microvascular Complications?. <i>PLoS ONE</i> , 2013, 8, e55921.	2.5	35
76	Association of TERC and OBFC1 Haplotypes with Mean Leukocyte Telomere Length and Risk for Coronary Heart Disease. <i>PLoS ONE</i> , 2013, 8, e83122.	2.5	42
77	Genome-Wide Association for Abdominal Subcutaneous and Visceral Adipose Reveals a Novel Locus for Visceral Fat in Women. <i>PLoS Genetics</i> , 2012, 8, e1002695.	3.5	245
78	Novel Loci for Adiponectin Levels and Their Influence on Type 2 Diabetes and Metabolic Traits: A Multi-Ethnic Meta-Analysis of 45,891 Individuals. <i>PLoS Genetics</i> , 2012, 8, e1002607.	3.5	419
79	Common mutations and polymorphisms predicting adverse cardiovascular events: current view. <i>Pharmacogenomics</i> , 2012, 13, 1875-1878.	1.3	1
80	Influence of Pre-analytical Variables on VEGF Gene Expression and Circulating Protein Concentrations. <i>Biopreservation and Biobanking</i> , 2012, 10, 454-461.	1.0	15
81	Novel association approach for variable number tandem repeats (VNTRs) identifies DOCK5 as a susceptibility gene for severe obesity. <i>Human Molecular Genetics</i> , 2012, 21, 3727-3738.	2.9	37
82	Alcohol Consumption, Beverage Preference, and Diet in Middle-Aged Men from the STANISLAS Study. <i>Journal of Nutrition and Metabolism</i> , 2012, 2012, 1-6.	1.8	21
83	Need for pharmacogenomic information also for generic medications: Recommendation of the European Society of Pharmacogenomics and Theranostics (ESPT). <i>Drug Metabolism and Drug Interactions</i> , 2012, 27, 119.	0.3	0
84	Genome-wide meta-analysis points to CTC1 and ZNF676 as genes regulating telomere homeostasis in humans. <i>Human Molecular Genetics</i> , 2012, 21, 5385-5394.	2.9	210
85	Clinical interest of point-of-care pharmacogenomic testing: clopidogrel behind warfarin. <i>Pharmacogenomics</i> , 2012, 13, 1215-1218.	1.3	5
86	Genetic biomarkers of hypertension and future challenges integrating epigenomics. <i>Clinica Chimica Acta</i> , 2012, 414, 259-265.	1.1	32
87	Human formyl peptide receptor 1 C32T SNP interacts with age and is associated with blood pressure levels. <i>Clinica Chimica Acta</i> , 2012, 413, 34-38.	1.1	15
88	Clinical necessity of partitioning of human plasma haptoglobin reference intervals by recently-discovered rs2000999. <i>Clinica Chimica Acta</i> , 2012, 413, 1618-1624.	1.1	15
89	High Prevalence of Metabolic Syndrome in Iran in Comparison with France: What Are the Components That Explain This?. <i>Metabolic Syndrome and Related Disorders</i> , 2012, 10, 181-188.	1.3	51
90	A Genome-Wide Association Search for Type 2 Diabetes Genes in African Americans. <i>PLoS ONE</i> , 2012, 7, e29202.	2.5	197

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91	A Genome-Wide Association Study Identifies rs2000999 as a Strong Genetic Determinant of Circulating Haptoglobin Levels. <i>PLoS ONE</i> , 2012, 7, e32327.	2.5	34
92	Functional Epistatic Interaction between rs6046G>A in F7 and rs5355C>T in SELE Modifies Systolic Blood Pressure Levels. <i>PLoS ONE</i> , 2012, 7, e40777.	2.5	8
93	Functional genomics towards personalized healthcare and systems medicine. <i>Personalized Medicine</i> , 2011, 8, 227-242.	1.5	1
94	Childhood Obesity Is Associated with Shorter Leukocyte Telomere Length. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011, 96, 1500-1505.	3.6	127
95	Mirror extreme BMI phenotypes associated with gene dosage at the chromosome 16p11.2 locus. <i>Nature</i> , 2011, 478, 97-102.	27.8	394
96	Genome-wide association study identifies loci influencing concentrations of liver enzymes in plasma. <i>Nature Genetics</i> , 2011, 43, 1131-1138.	21.4	501
97	Klotho KL-VS genotype is involved in blood pressure regulation. <i>Clinica Chimica Acta</i> , 2011, 412, 1773-1777.	1.1	19
98	Cardiovascular diseases and genome-wide association studies. <i>Clinica Chimica Acta</i> , 2011, 412, 1697-1701.	1.1	22
99	The Lin28/let-7 Axis Regulates Glucose Metabolism. <i>Cell</i> , 2011, 147, 81-94.	28.9	812
100	Genome-Wide Association Identifies Nine Common Variants Associated With Fasting Proinsulin Levels and Provides New Insights Into the Pathophysiology of Type 2 Diabetes. <i>Diabetes</i> , 2011, 60, 2624-2634.	0.6	335
101	Genome-Wide Association Analysis Identifies Variants Associated with Nonalcoholic Fatty Liver Disease That Have Distinct Effects on Metabolic Traits. <i>PLoS Genetics</i> , 2011, 7, e1001324.	3.5	796
102	Substantial variation in qPCR measured mean blood telomere lengths in young men from eleven European countries. <i>American Journal of Human Biology</i> , 2011, 23, 228-231.	1.6	37
103	Availability of pharmacogenetic and pharmacogenomic information in anticancer drug monographs in France: personalized cancer therapy. <i>Pharmacogenomics</i> , 2011, 12, 681-691.	1.3	4
104	Identification of <i>cis</i> - and <i>trans</i> -Acting Genetic Variants Explaining Up to Half the Variation in Circulating Vascular Endothelial Growth Factor Levels. <i>Circulation Research</i> , 2011, 109, 554-563.	4.5	72
105	Association of Genetic Loci With Glucose Levels in Childhood and Adolescence. <i>Diabetes</i> , 2011, 60, 1805-1812.	0.6	103
106	Biological and genetic factors associated with ABCB1 and pregnane-X-receptor expressions in peripheral blood mononuclear cells in the STANISLAS cohort. <i>Drug Metabolism and Drug Interactions</i> , 2011, 26, 27-32.	0.3	3
107	Association Between Angiotensin II Type 1 Receptor Gene Polymorphism and Metabolic Syndrome in a Young Female Iranian Population. <i>Archives of Medical Research</i> , 2010, 41, 343-349.	3.3	17
108	A multi-stage multi-design strategy provides strong evidence that the BAI3 locus is associated with early-onset venous thromboembolism. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 2671-2679.	3.8	42

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109	Expression of inflammatory molecules and associations with BMI in children. <i>European Journal of Clinical Investigation</i> , 2010, 40, 388-392.	3.4	23
110	Sex-dependent Associations of Leptin With Metabolic Syndrome-related Variables: The Stanislas Study. <i>Obesity</i> , 2010, 18, 196-201.	3.0	24
111	Association analyses of 249,796 individuals reveal 18 new loci associated with body mass index. <i>Nature Genetics</i> , 2010, 42, 937-948.	21.4	2,634
112	Prevalence of Loss-of-Function FTO Mutations in Lean and Obese Individuals. <i>Diabetes</i> , 2010, 59, 311-318.	0.6	93
113	Interactions of Dietary Whole-Grain Intake With Fasting Glucose- and Insulin-Related Genetic Loci in Individuals of European Descent: A meta-analysis of 14 cohort studies. <i>Diabetes Care</i> , 2010, 33, 2684-2691.	8.6	127
114	Association Study of Gene Polymorphisms Involved in Vascular Alterations in Elderly Hypertensives with Subjective Memory Complaints. <i>Dementia and Geriatric Cognitive Disorders</i> , 2010, 30, 440-448.	1.5	11
115	Metabolic syndrome-related composite factors over 5 years in the STANISLAS Family Study: Genetic heritability and common environmental influences. <i>Clinica Chimica Acta</i> , 2010, 411, 833-839.	1.1	14
116	MS64 CAPILLARY ISOTACHOPHORESIS AS AN OVERALL TOOL TO REVEAL ATHEROGENIC LIPOPROTEIN PHENOTYPE. <i>Atherosclerosis Supplements</i> , 2010, 11, 123.	1.2	0
117	New genetic loci implicated in fasting glucose homeostasis and their impact on type 2 diabetes risk. <i>Nature Genetics</i> , 2010, 42, 105-116.	21.4	1,982
118	Visfatin: The Link Between Inflammation and Childhood Obesity. <i>Diabetes Care</i> , 2009, 32, e71-e71.	8.6	20
119	Personalized therapy and pharmacogenomics: future perspective. <i>Pharmacogenomics</i> , 2009, 10, 927-930.	1.3	9
120	Capillary isotachopheresis study of lipoprotein network sensitive to apolipoprotein E phenotype. 2. ApoE and apoC-III relations in triglyceride clearance. <i>Molecular and Cellular Biochemistry</i> , 2009, 325, 25-40.	3.1	3
121	Capillary isotachopheresis study of lipoprotein network sensitive to apolipoprotein E phenotype. 1. ApoE distribution between lipoproteins. <i>Molecular and Cellular Biochemistry</i> , 2009, 325, 41-51.	3.1	3
122	Genetic profiling of human cell lines used as in vitro model to study cardiovascular pathophysiology and pharmacotoxicology. <i>Cell Biology and Toxicology</i> , 2009, 25, 561-571.	5.3	6
123	A variant near MTNR1B is associated with increased fasting plasma glucose levels and type 2 diabetes risk. <i>Nature Genetics</i> , 2009, 41, 89-94.	21.4	540
124	Genome-wide association study for early-onset and morbid adult obesity identifies three new risk loci in European populations. <i>Nature Genetics</i> , 2009, 41, 157-159.	21.4	585
125	Association between EGF and lipid concentrations: A benefit role in the atherosclerotic process?. <i>Clinica Chimica Acta</i> , 2009, 402, 196-198.	1.1	8
126	Association of ABCB1 gene polymorphisms with plasma lipid and apolipoprotein concentrations in the STANISLAS cohort. <i>Clinica Chimica Acta</i> , 2009, 403, 198-202.	1.1	23

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127	Adipokine expression in adipose tissue and in peripheral blood mononuclear cells in children. <i>Clinica Chimica Acta</i> , 2009, 410, 85-89.	1.1	7
128	Association of human cathelicidin (hCAP-18/LL-37) gene expression with cardiovascular disease risk factors. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2009, 19, 720-728.	2.6	28
129	Human formyl peptide receptor 1 (<i>FPR1</i>) c.32C>T SNP is associated with decreased soluble E-selectin levels. <i>Pharmacogenomics</i> , 2009, 10, 951-959.	1.3	8
130	Parental precocious influences on offspring cardiovascular risk markers: an exploratory study in the STANISLAS Cohort. <i>Personalized Medicine</i> , 2009, 6, 343-352.	1.5	0
131	Candidate gene microarray analysis in peripheral blood cells for studying hypertension/obesity. <i>Personalized Medicine</i> , 2009, 6, 269-291.	1.5	5
132	Functional genomics towards personalized healthcare. <i>Personalized Medicine</i> , 2009, 6, 19-32.	1.5	3
133	Drug Metabolizing Enzymes and Transporters mRNA in Peripheral Blood Mononuclear Cells of Healthy Subjects: Biological Variations and Importance of Preanalytical Steps. <i>Current Drug Metabolism</i> , 2009, 10, 410-419.	1.2	5
134	Genomics and the Prospects of Existing and Emerging Therapeutics for Cardiovascular Diseases. <i>Current Pharmaceutical Design</i> , 2009, 15, 3193-3206.	1.9	12
135	The association of telomere length with paternal history of premature myocardial infarction in the European Atherosclerosis Research Study II. <i>Journal of Molecular Medicine</i> , 2008, 86, 815-824.	3.9	54
136	Five-year alterations in BMI are associated with clustering of changes in cardiovascular risk factors in a gender-dependant way: the Stanislas study. <i>International Journal of Obesity</i> , 2008, 32, 1279-1288.	3.4	27
137	Genetic Polymorphism of <i>CYP2C19</i> Gene in the Stanislas Cohort. A link with Inflammation. <i>Annals of Human Genetics</i> , 2008, 72, 178-183.	0.8	35
138	P-selectin polymorphisms'™ influences on P-selectin serum concentrations and on their familial correlation: the STANISLAS family study. <i>Journal of Thrombosis and Haemostasis</i> , 2008, 6, 920-927.	3.8	9
139	Visfatin, low'grade inflammation and body mass index (BMI). <i>Clinical Endocrinology</i> , 2008, 69, 568-574.	2.4	38
140	Statins as effectors of key activities involved in apoE-dependent VLDL metabolism: Review and hypothesis. <i>Vascular Pharmacology</i> , 2008, 48, 70-75.	2.1	10
141	Pharmacogenomics and Cardiovascular Drugs. <i>Methods in Pharmacology and Toxicology</i> , 2008, , 413-446.	0.2	0
142	Leptin expression in Peripheral Blood Mononuclear Cells (PBMCs) is related with blood pressure variability. <i>Clinica Chimica Acta</i> , 2008, 395, 47-50.	1.1	15
143	Transcription Factor and Drug-Metabolizing Enzyme Gene Expression in Lymphocytes from Healthy Human Subjects. <i>Drug Metabolism and Disposition</i> , 2008, 36, 182-189.	3.3	80
144	A new single nucleotide polymorphism genotyping method based on gap ligase chain reaction and a microsphere detection assay. <i>Clinical Chemistry and Laboratory Medicine</i> , 2008, 46, 486-9.	2.3	4

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145	The STANISLAS Cohort: a 10-year follow-up of supposed healthy families. Gene-environment interactions, reference values and evaluation of biomarkers in prevention of cardiovascular diseases. <i>Clinical Chemistry and Laboratory Medicine</i> , 2008, 46, 733-47.	2.3	50
146	Genetic profiling in healthy subjects from the Stanislas cohort based on 24 polymorphisms: effects on biological variables. <i>Clinical Chemistry and Laboratory Medicine</i> , 2008, 46, 64-72.	2.3	1
147	Association of classical and related inflammatory markers with high-sensitivity C-reactive protein in healthy individuals: results from the Stanislas cohort. <i>Clinical Chemistry and Laboratory Medicine</i> , 2007, 45, 1339-46.	2.3	11
148	Effect of HMGCoA Reductase Inhibitors on Cytochrome P450 Expression in Endothelial Cell Line. <i>Journal of Cardiovascular Pharmacology</i> , 2007, 49, 306-315.	1.9	23
149	Enzymes and pharmacogenetics of cardiovascular drugs. <i>Clinica Chimica Acta</i> , 2007, 381, 26-31.	1.1	40
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