

Score Tests for Association between Traits and Haplotypes Ambiguous

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Citation Report

#	ARTICLE	IF	CITATIONS
1	TDT statistics for mapping quantitative trait loci. <i>Annals of Human Genetics</i> , 1998, 62, 431-452.	0.3	68
2	Specific haplotypes of the P-selectin gene are associated with myocardial infarction. <i>Human Molecular Genetics</i> , 2002, 11, 2015-2023.	1.4	161
3	Caution on Pedigree Haplotype Inference with Software That Assumes Linkage Equilibrium. <i>American Journal of Human Genetics</i> , 2002, 71, 992-995.	2.6	88
4	Estrogen Receptor \hat{I}^2 Polymorphisms Are Associated With Bone Mass in Women and Men: The Framingham Study. <i>Journal of Bone and Mineral Research</i> , 2003, 19, 773-781.	3.1	67
5	CTLA4 is associated with susceptibility to multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2003, 134, 133-141.	1.1	73
6	A population-based study of IL4 polymorphisms in multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2003, 137, 134-139.	1.1	45
7	On selecting markers for association studies: Patterns of linkage disequilibrium between two and three diallelic loci. <i>Genetic Epidemiology</i> , 2003, 24, 57-67.	0.6	38
8	Evolutionary-based association analysis using haplotype data. <i>Genetic Epidemiology</i> , 2003, 25, 48-58.	0.6	106
9	Pedigree disequilibrium tests for multilocus haplotypes. <i>Genetic Epidemiology</i> , 2003, 25, 115-121.	0.6	1,094
10	Polygenic inheritance of breast cancer: Implications for design of association studies. <i>Genetic Epidemiology</i> , 2003, 25, 190-202.	0.6	213
11	Polymorphisms in the CYP1A1 gene are associated with prostate cancer risk. <i>International Journal of Cancer</i> , 2003, 106, 375-378.	2.3	56
12	Case-control study of the α -synuclein interacting protein gene and Parkinson's disease. <i>Movement Disorders</i> , 2003, 18, 1233-1239.	2.2	13
13	Nuclear DNA analyses in genetic studies of populations: practice, problems and prospects. <i>Molecular Ecology</i> , 2003, 12, 563-584.	2.0	575
14	Polymorphisms in the CYP1B1 gene are associated with increased risk of prostate cancer. <i>British Journal of Cancer</i> , 2003, 89, 1524-1529.	2.9	59
15	Association of a disintegrin and metalloprotease 33 (ADAM33) gene with asthma in ethnically diverse populations. <i>Journal of Allergy and Clinical Immunology</i> , 2003, 112, 717-722.	1.5	190
16	Detecting Disease Associations due to Linkage Disequilibrium Using Haplotype Tags: A Class of Tests and the Determinants of Statistical Power. <i>Human Heredity</i> , 2003, 56, 18-31.	0.4	392
17	Genetic variation and hematology: single-nucleotide polymorphisms, haplotypes, and complex disease. <i>Seminars in Hematology</i> , 2003, 40, 321-328.	1.8	12
18	Common Sequence Variants of the Macrophage Scavenger Receptor 1 Gene Are Associated with Prostate Cancer Risk. <i>American Journal of Human Genetics</i> , 2003, 72, 208-212.	2.6	94

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19	A Method for the Assessment of Disease Associations with Single-Nucleotide Polymorphism Haplotypes and Environmental Variables in Case-Control Studies. <i>American Journal of Human Genetics</i> , 2003, 72, 1231-1250.	2.6	158
20	Control of Confounding of Genetic Associations in Stratified Populations. <i>American Journal of Human Genetics</i> , 2003, 72, 1492-1504.	2.6	456
21	Inference on Haplotype Effects in Case-Control Studies Using Unphased Genotype Data. <i>American Journal of Human Genetics</i> , 2003, 73, 1316-1329.	2.6	235
22	Haplotype analysis in population genetics and association studies. <i>Pharmacogenomics</i> , 2003, 4, 171-178.	0.6	131
23	Association between Parkinson's disease and polymorphisms in the nNOS and iNOS genes in a community-based case-control study. <i>Human Molecular Genetics</i> , 2003, 12, 79-86.	1.4	108
24	Modeling and E-M Estimation of Haplotype-Specific Relative Risks from Genotype Data for a Case-Control Study of Unrelated Individuals. <i>Human Heredity</i> , 2003, 55, 179-190.	0.4	249
25	A haplotype at the <i>PARK3</i> locus influences onset age for Parkinson's disease. <i>Neurology</i> , 2003, 61, 1557-1561.	1.5	49
26	Hepatic CYP2B6 Expression: Gender and Ethnic Differences and Relationship to CYP2B6 Genotype and CAR (Constitutive Androstane Receptor) Expression. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2003, 307, 906-922.	1.3	367
27	Functional haplotypes of the RET proto-oncogene promoter are associated with Hirschsprung disease (HSCR). <i>Human Molecular Genetics</i> , 2003, 12, 3207-3214.	1.4	67
28	Association Between Genetic Polymorphisms in the Prostate-Specific Antigen Gene Promoter and Serum Prostate-Specific Antigen Levels. <i>Journal of the National Cancer Institute</i> , 2003, 95, 1044-1053.	3.0	83
29	Choosing Haplotype-Tagging SNPS Based on Unphased Genotype Data Using a Preliminary Sample of Unrelated Subjects with an Example from the Multiethnic Cohort Study. <i>Human Heredity</i> , 2003, 55, 27-36.	0.4	386
30	Estimation and Tests of Haplotype-Environment Interaction when Linkage Phase Is Ambiguous. <i>Human Heredity</i> , 2003, 55, 56-65.	0.4	423
31	Polymorphisms in the Insulin-Degrading Enzyme Gene Are Associated With Type 2 Diabetes in Men From the NHLBI Framingham Heart Study. <i>Diabetes</i> , 2003, 52, 1562-1567.	0.3	100
32	Genetic Variation at the Scavenger Receptor Class B Type I Gene Locus Determines Plasma Lipoprotein Concentrations and Particle Size and Interacts with Type 2 Diabetes: The Framingham Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 2869-2879.	1.8	108
33	Association of tumor necrosis factor gene polymorphisms and prolonged mechanical ventilation after coronary artery bypass surgery. <i>Critical Care Medicine</i> , 2003, 31, 133-140.	0.4	28
35	The role of genetic polymorphisms in environmental health.. <i>Environmental Health Perspectives</i> , 2003, 111, 1055-1064.	2.8	127
36	Bayesian Spatial Modeling of Haplotype Associations. <i>Human Heredity</i> , 2003, 56, 32-40.	0.4	44
37	Disease Associations and Family-Based Tests. <i>Current Protocols in Human Genetics</i> , 2003, 38, Unit 1.12.	3.5	0

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38	Detailed Analysis of Allelic Variation in the ABCA4 Gene in Age-Related Maculopathy. , 2003, 44, 2868.		33
39	Haplotype analysis of the matrix metalloproteinase 3 gene and myocardial infarction in a Chinese Han population. Thrombosis and Haemostasis, 2004, 92, 867-873.	1.8	41
40	Sequence Variants of Toll-Like Receptor 4 Are Associated with Prostate Cancer Risk. Cancer Research, 2004, 64, 2918-2922.	0.4	214
41	2LD, GENECOUNTING and HAP: computer programs for linkage disequilibrium analysis. Bioinformatics, 2004, 20, 1325-1326.	1.8	157
42	Complex Haplotypes of the PGC-1 α Gene Are Associated With Carbohydrate Metabolism and Type 2 Diabetes. Diabetes, 2004, 53, 1385-1393.	0.3	99
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45	Genetic Analysis of the RNASEL Gene in Hereditary, Familial, and Sporadic Prostate Cancer. Clinical Cancer Research, 2004, 10, 7150-7156.	3.2	87
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47	Polymorphism in the Calsequestrin 1 (CASQ1) Gene on Chromosome 1q21 Is Associated With Type 2 Diabetes in the Old Order Amish. Diabetes, 2004, 53, 3292-3299.	0.3	44
48	Association of Protein Tyrosine Phosphatase 1B Gene Polymorphisms With Type 2 Diabetes. Diabetes, 2004, 53, 3007-3012.	0.3	113
49	Haplotypic analyses of the IGF2-INS-TH gene cluster in relation to cardiovascular risk traits. Human Molecular Genetics, 2004, 13, 715-725.	1.4	57
50	Corticosteroid pharmacogenetics: association of sequence variants in CRHR1 with improved lung function in asthmatics treated with inhaled corticosteroids. Human Molecular Genetics, 2004, 13, 1353-1359.	1.4	315
51	Association of late-onset Alzheimer's disease with genetic variation in multiple members of the GAPD gene family. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 15688-15693.	3.3	134
52	A Note on Inference of Trait Associations with SNP Haplotypes and Other Attributes in Generalized Linear Models. Human Heredity, 2004, 57, 200-206.	0.4	38
53	A Haplotype-Based Test of Association Using Data from Cohort and Nested Case-Control Epidemiologic Studies. Human Heredity, 2004, 58, 18-29.	0.4	18
54	Association in Multifactorial Traits: How to Deal with Rare Observations?. Human Heredity, 2004, 58, 73-81.	0.4	8
55	Genotypes and haplotypes predisposing to myocardial infarction: a multilocus case-control study. European Heart Journal, 2004, 25, 459-467.	1.0	133

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56	Multigenic control of serum adiponectin levels: evidence for a role of the APM1 gene and a locus on 14q13. <i>Physiological Genomics</i> , 2004, 19, 170-174.	1.0	67
57	Cat Red Blood Cell Thiopurine S-Methyltransferase: Companion Animal Pharmacogenetics. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2004, 308, 617-626.	1.3	17
58	A common haplotype at the CD36 locus is associated with high free fatty acid levels and increased cardiovascular risk in Caucasians. <i>Human Molecular Genetics</i> , 2004, 13, 2197-2205.	1.4	161
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60	H6D Polymorphism in Macrophage-Inhibitory Cytokine-1 Gene Associated With Prostate Cancer. <i>Journal of the National Cancer Institute</i> , 2004, 96, 1248-1254.	3.0	111
61	Large-Scale Association Study Identifies ICAM Gene Region as Breast and Prostate Cancer Susceptibility Locus. <i>Cancer Research</i> , 2004, 64, 8906-8910.	0.4	92
62	The transforming growth factor- β 1 (TGFB1) gene is associated with chronic obstructive pulmonary disease (COPD). <i>Human Molecular Genetics</i> , 2004, 13, 1649-1656.	1.4	203
63	Common DNase I polymorphism associated with autoantibody production among systemic lupus erythematosus patients. <i>Human Molecular Genetics</i> , 2004, 13, 2343-2350.	1.4	62
64	Association of <i>APOE</i> polymorphisms with disease severity in MS is limited to women. <i>Neurology</i> , 2004, 62, 811-814.	1.5	75
65	Haplotypic analyses of the aldosterone synthase gene CYP11B2 associated with stage-2 hypertension in northern Han Chinese. <i>Clinical Genetics</i> , 2004, 66, 409-416.	1.0	24
66	A ₁₆ C haplotype in the <i>FCμRI²</i> gene confers a higher risk for atopic asthma in the Indian population. <i>Clinical Genetics</i> , 2004, 66, 417-425.	1.0	16
67	Cox proportional hazards survival regression in haplotype-based association analysis using the Stochastic-EM algorithm. <i>European Journal of Human Genetics</i> , 2004, 12, 971-974.	1.4	61
68	Interleukin-10 promoter single-nucleotide polymorphisms as markers for disease susceptibility and disease severity in leprosy. <i>Genes and Immunity</i> , 2004, 5, 592-595.	2.2	69
69	Human catechol O-methyltransferase genetic variation: gene resequencing and functional characterization of variant allozymes. <i>Molecular Psychiatry</i> , 2004, 9, 151-160.	4.1	146
70	Association of a corticotropin-releasing hormone receptor 1 haplotype and antidepressant treatment response in Mexican-Americans. <i>Molecular Psychiatry</i> , 2004, 9, 1075-1082.	4.1	159
71	CD95 polymorphisms are associated with susceptibility to MS in women. <i>Journal of Neuroimmunology</i> , 2004, 146, 162-170.	1.1	30
72	Haplotype Association Analysis of Discrete and Continuous Traits Using Mixture of Regression Models. <i>Behavior Genetics</i> , 2004, 34, 207-214.	1.4	43
73	Comprehensive association analysis of APOE regulatory region polymorphisms in Alzheimer disease. <i>Neurogenetics</i> , 2004, 5, 201-208.	0.7	22

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74	Haplotype analysis of VDR gene polymorphisms: a meta-analysis. <i>Osteoporosis International</i> , 2004, 15, 729-734.	1.3	78
75	Human catecholamine sulfotransferase (SULT1A3) pharmacogenetics: functional genetic polymorphism. <i>Journal of Neurochemistry</i> , 2004, 87, 809-819.	2.1	55
76	A new algorithm for haplotype-based association analysis: the Stochastic-EM algorithm. <i>Annals of Human Genetics</i> , 2004, 68, 165-177.	0.3	258
77	Introducing the Multivariate Dale Model in Population-Based Genetic Association Studies. <i>Biometrical Journal</i> , 2004, 46, 187-202.	0.6	2
78	?-Synuclein promoter confers susceptibility to Parkinson's disease. <i>Annals of Neurology</i> , 2004, 56, 591-595.	2.8	200
79	Family-based tests for associating haplotypes with general phenotype data: Application to asthma genetics. <i>Genetic Epidemiology</i> , 2004, 26, 61-69.	0.6	395
80	IL10 gene polymorphisms are associated with asthma phenotypes in children. <i>Genetic Epidemiology</i> , 2004, 26, 155-165.	0.6	86
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82	Haplotype-based association analysis in cohort studies of unrelated individuals. <i>Genetic Epidemiology</i> , 2004, 26, 255-264.	0.6	45
83	Comparison of prospective and retrospective methods for haplotype inference in case-control studies. <i>Genetic Epidemiology</i> , 2004, 27, 192-201.	0.6	82
84	Genetic association mapping under founder heterogeneity via weighted haplotype similarity analysis in candidate genes. <i>Genetic Epidemiology</i> , 2004, 27, 182-191.	0.6	32
85	Tag SNP selection for association studies. <i>Genetic Epidemiology</i> , 2004, 27, 365-374.	0.6	165
86	Use of unphased multilocus genotype data in indirect association studies. <i>Genetic Epidemiology</i> , 2004, 27, 415-428.	0.6	178
87	Evaluating associations of haplotypes with traits. <i>Genetic Epidemiology</i> , 2004, 27, 348-364.	0.6	293
88	SNPs, haplotypes, and model selection in a candidate gene region: The SIMPLe analysis for multilocus data. <i>Genetic Epidemiology</i> , 2004, 27, 429-441.	0.6	40
89	Two-Stage sampling designs for gene association studies. <i>Genetic Epidemiology</i> , 2004, 27, 401-414.	0.6	88
90	Molecular genetics of myocardial infarction: many genes, more questions than answers. <i>European Heart Journal</i> , 2004, 25, 451-453.	1.0	9
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94	Association of a variation in the promoter region of the brain-derived neurotrophic factor gene with familial Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2004, 10, 213-219.	1.1	31
95	Association of ABCA1 with late-onset Alzheimer's disease is not observed in a case-control study. <i>Neuroscience Letters</i> , 2004, 366, 268-271.	1.0	58
96	The Future of Association Studies: Gene-Based Analysis and Replication. <i>American Journal of Human Genetics</i> , 2004, 75, 353-362.	2.6	598
97	Association of the DTNBP1 Locus with Schizophrenia in a U.S. Population. <i>American Journal of Human Genetics</i> , 2004, 75, 891-898.	2.6	155
98	Glutathione-S-transferase M1, M3, P1 and T1 polymorphisms and severity of lung disease in children with cystic fibrosis. <i>Pharmacogenetics and Genomics</i> , 2004, 14, 295-301.	5.7	40
99	Angiotensin II type I receptor gene and myocardial infarction. <i>Pharmacogenetics and Genomics</i> , 2004, 14, 673-681.	5.7	16
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104	Influence of the endothelial nitric oxide synthase gene on conventional and ambulatory blood pressure: sib-pair analysis and haplotype study. <i>Journal of Hypertension</i> , 2005, 23, 759-765.	0.3	16
105	Finding and using haplotype blocks in candidate gene association studies. , 2005, , .		0
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107	Thrombomodulin gene polymorphisms or haplotypes as potential risk factors for venous thromboembolism: a population-based case-control study. <i>Journal of Thrombosis and Haemostasis</i> , 2005, 3, 710-717.	1.9	43
108	Cyclooxygenase-1 haplotype modulates platelet response to aspirin. <i>Journal of Thrombosis and Haemostasis</i> , 2005, 3, 2340-2345.	1.9	187
109	Common genomic sequence variation of the prothrombin gene and risk of non-fatal myocardial infarction in white women. <i>Journal of Thrombosis and Haemostasis</i> , 2005, 3, 2809-2811.	1.9	2
110	C Protein β ₃ Subunit Gene Variants and Essential Hypertension in the Northern Chinese Han Population. <i>Annals of Human Genetics</i> , 2005, 69, 468-473.	0.3	23

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111	Using Tree-Based Recursive Partitioning Methods to Group Haplotypes for Increased Power in Association Studies. <i>Annals of Human Genetics</i> , 2005, 69, 577-589.	0.3	24
112	Association of CommonCRP Gene Variants with CRP Levels and Cardiovascular Events. <i>Annals of Human Genetics</i> , 2005, 69, 623-638.	0.3	199
113	Tests of Association Between Quantitative Traits and Haplotypes In A Reduced-Dimensional Space. <i>Annals of Human Genetics</i> , 2005, 69, 715-732.	0.3	15
114	Interleukin-1 gene cluster polymorphisms predict risk of ESRD. <i>Kidney International</i> , 2005, 68, 278-284.	2.6	34
115	Polymorphisms in the 5-lipoxygenase activating protein (ALOX5AP) gene are not associated with asthma in an Australian population. <i>Clinical and Experimental Allergy</i> , 2005, 35, 332-338.	1.4	12
116	Interleukin-10 promoter polymorphisms and atopic asthma in North Indians. <i>Clinical and Experimental Allergy</i> , 2005, 35, 914-919.	1.4	62
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118	Interleukin-1 receptor antagonist haplotype associated with prostate cancer risk. <i>British Journal of Cancer</i> , 2005, 93, 493-497.	2.9	51
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120	Genome-wide association studies for common diseases and complex traits. <i>Nature Reviews Genetics</i> , 2005, 6, 95-108.	7.7	2,717
121	IFNG polymorphisms are associated with gender differences in susceptibility to multiple sclerosis. <i>Genes and Immunity</i> , 2005, 6, 153-161.	2.2	57
122	Promoter region polymorphism of macrophage migration inhibitory factor is strong risk factor for young onset of extensive alopecia areata. <i>Genes and Immunity</i> , 2005, 6, 285-289.	2.2	36
123	IL-6 gene variation is not associated with increased serum levels of IL-6, muscle, weakness, or frailty in older women. <i>Experimental Gerontology</i> , 2005, 40, 344-352.	1.2	59
124	Decision Forest Analysis of 61 Single Nucleotide Polymorphisms in a Case-Control Study of Esophageal Cancer; a novel method. <i>BMC Bioinformatics</i> , 2005, 6, S4.	1.2	48
125	Common variation in EMSY and risk of breast and ovarian cancer: a case-control study using HapMap tagging SNPs. <i>BMC Cancer</i> , 2005, 5, 81.	1.1	14
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127	Accounting for haplotype uncertainty in matched association studies: A comparison of simple and flexible techniques. <i>Genetic Epidemiology</i> , 2005, 28, 261-272.	0.6	131
128	Direct analysis of unphased SNP genotype data in population-based association studies via Bayesian partition modelling of haplotypes. <i>Genetic Epidemiology</i> , 2005, 29, 91-107.	0.6	41

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129	Asymptotic equivalence between two score tests for haplotype-specific risk in general linear models. <i>Genetic Epidemiology</i> , 2005, 29, 166-170.	0.6	18
130	Maximum likelihood estimation of haplotype effects and haplotype-environment interactions in association studies. <i>Genetic Epidemiology</i> , 2005, 29, 299-312.	0.6	109
131	Global transmission/disequilibrium tests based on haplotype sharing in multiple candidate genes. <i>Genetic Epidemiology</i> , 2005, 29, 323-335.	0.6	7
132	Assessment and implications of linkage disequilibrium in genome-wide single-nucleotide polymorphism and microsatellite panels. <i>Genetic Epidemiology</i> , 2005, 29, S72-S76.	0.6	12
133	Genetic association of the APP binding protein 2 gene (APBB2) with late onset Alzheimer disease. <i>Human Mutation</i> , 2005, 25, 270-277.	1.1	36
134	CYP1A1 variants and smoking-related lung cancer in San Francisco bay area Latinos and African Americans. <i>International Journal of Cancer</i> , 2005, 113, 141-147.	2.3	39
135	Polymorphisms in GLTSCR1 and ERCC2 are associated with the development of oligodendrogliomas. <i>Cancer</i> , 2005, 103, 2363-2372.	2.0	60
136	Mutational screening and association study of glutamate decarboxylase 1 as a candidate susceptibility gene for bipolar affective disorder and schizophrenia. , 2005, 135B, 94-101.		44
137	Effects of common atopy-associated amino acid substitutions in the IL-4 receptor alpha chain on IL-4 induced phenotypes. <i>Immunogenetics</i> , 2005, 56, 808-817.	1.2	22
138	Polymorphisms in the human surfactant protein-D (SFTPD) gene: strong evidence that serum levels of surfactant protein-D (SP-D) are genetically influenced. <i>Immunogenetics</i> , 2005, 57, 1-7.	1.2	65
139	717A>G polymorphism of human C-reactive protein gene associated with coronary heart disease in ethnic Han Chinese: the Beijing atherosclerosis study. <i>Journal of Molecular Medicine</i> , 2005, 83, 72-78.	1.7	60
140	Association between polymorphisms in the nuclear respiratory factor 1 gene and type 2 diabetes mellitus in the Korean population. <i>Diabetologia</i> , 2005, 48, 2033-2038.	2.9	18
141	Association of a polymorphism in the gene encoding phosphoenolpyruvate carboxykinase 1 with high-density lipoprotein and triglyceride levels. <i>Diabetologia</i> , 2005, 48, 2025-2032.	2.9	14
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143	An analysis of genetic variation across the MBL2 locus in Dutch Caucasians indicates that 3 haplotypes could modify circulating levels of mannose-binding lectin. <i>Human Genetics</i> , 2005, 118, 404-415.	1.8	39
144	Comprehensive genetic evaluation of common E-cadherin sequence variants and prostate cancer risk: strong confirmation of functional promoter SNP. <i>Human Genetics</i> , 2005, 118, 339-347.	1.8	29
145	The EPAS1 gene influences the aerobic/anaerobic contribution in elite endurance athletes. <i>Human Genetics</i> , 2005, 118, 416-423.	1.8	54
146	Association analysis of interleukin 5 receptor alpha subunit (IL5RA) polymorphisms and asthma. <i>Journal of Human Genetics</i> , 2005, 50, 628-634.	1.1	25

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147	Haplotype Effects on Human Survival: Logistic Regression Models Applied to Unphased Genotype Data. <i>Annals of Human Genetics</i> , 2005, 69, 168-175.	0.3	4
148	Linkage disequilibrium mapping via cladistic analysis of phase-unknown genotypes and inferred haplotypes in the Genetic Analysis Workshop 14 simulated data. <i>BMC Genetics</i> , 2005, 6, S100.	2.7	9
149	Fine Mapping Functional Sites or Regions from Case-Control Data Using Haplotypes of Multiple Linked SNPs. <i>Annals of Human Genetics</i> , 2005, 69, 102-112.	0.3	21
150	Polymorphisms of the Flavin containing monooxygenase 3 (FMO3) gene do not predispose to essential hypertension in Caucasians. <i>BMC Medical Genetics</i> , 2005, 6, 41.	2.1	18
151	Genetic analysis of the GLUT10 glucose transporter (SLC2A10) polymorphisms in Caucasian American type 2 diabetes. <i>BMC Medical Genetics</i> , 2005, 6, 42.	2.1	24
152	Tests of trait-haplotype association when linkage phase is ambiguous, appropriate for matched case-control and cohort studies with competing risks. <i>Statistics in Medicine</i> , 2005, 24, 2299-2316.	0.8	2
153	Retrospective analysis of case-control studies when the population is in Hardy-Weinberg equilibrium. <i>Statistics in Medicine</i> , 2005, 24, 3289-3310.	0.8	9
156	Genetic polymorphisms in DPF3 associated with risk of breast cancer and lymph node metastases. <i>Journal of Carcinogenesis</i> , 2005, 4, 13.	2.5	18
157	Common Polymorphisms in ERCC2 (Xeroderma pigmentosum D) are not Associated with Breast Cancer Risk. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2005, 14, 1828-1831.	1.1	23
158	ERCC1 and ERCC2 polymorphisms and adult glioma. <i>Neuro-Oncology</i> , 2005, 7, 495-507.	0.6	71
159	COMT Polymorphisms and Anxiety-Related Personality Traits. <i>Neuropsychopharmacology</i> , 2005, 30, 2092-2102.	2.8	199
160	Cyclooxygenase-1 gene polymorphisms in patients with different asthma phenotypes and atopy. <i>European Respiratory Journal</i> , 2005, 26, 249-256.	3.1	29
161	Haplotype association analysis of human disease traits using genotype data of unrelated individuals. <i>Genetical Research</i> , 2005, 86, 223-231.	0.3	18
162	CDX2 Polymorphisms, RNA Expression, and Risk of Colorectal Cancer. <i>Cancer Research</i> , 2005, 65, 5488-5492.	0.4	29
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