Juha Kere

List of Publications by Year
in descending order

		2322	4117
568	41,664	98	175
papers	citations	h-index	g-index

595	595	595	50541
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Transcriptomeâ€based identification of novel endotypes in adult atopic dermatitis. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 1486-1498.	5.7	8
2	Characteristics of preeclampsia in donor cell gestations. Pregnancy Hypertension, 2022, 27, 59-61.	1.4	3
3	CRISPR activation enables high-fidelity reprogramming into human pluripotent stem cells. Stem Cell Reports, 2022, 17, 413-426.	4.8	13
4	SkewC: Identifying cells with skewed gene body coverage in single-cell RNA sequencing data. IScience, 2022, 25, 103777.	4.1	4
5	DUX4 is a multifunctional factor priming human embryonic genome activation. IScience, 2022, 25, 104137.	4.1	20
6	Viral infectionâ€related gene upregulation in monocytes in children with signs of Î²â€cell autoimmunity. Pediatric Diabetes, 2022, 23, 703-713.	2.9	3
7	INFLUENCE OF FLG LOSS-OF-FUNCTION MUTATIONS IN HOSTâ€“MICROBE INTERACTIONS DURING ATOPIC SKIN INFLAMMATION. Journal of Dermatological Science, 2022, , .	1.9	0
8	Idiopathic scoliosis: a systematic review and meta-analysis of heritability. EFORT Open Reviews, 2022, 7, 414-421.	4.1	1
9	Transient DUX4 expression in human embryonic stem cells induces blastomere-like expression program that is marked by SLC34A2. Stem Cell Reports, 2022, 17, 1743-1756.	4.8	11
10	Biomarkers of nanomaterials hazard from multi-layer data. Nature Communications, 2022, 13, .	12.8	16
11	Searching for a paternal phenotype for preeclampsia. Acta Obstetricia Et Gynecologica Scandinavica, 2022, 101, 862-870.	2.8	1
12	Genome-wide association study reveals new insights into the heritability and genetic correlates of developmental dyslexia. Molecular Psychiatry, 2021, 26, 3004-3017.	7.9	56
13	Distinct whole-blood transcriptome profile of children with metabolic healthy overweight/obesity compared to metabolic unhealthy overweight/obesity. Pediatric Research, 2021, 89, 1687-1694.	2.3	10
14	Otitis media susceptibility and shifts in the head and neck microbiome due to SPINK5 variants. Journal of Medical Genetics, 2021, 58, 442-452.	3.2	14
15	Association of Maternal DNA Methylation and Offspring Birthweight. Reproductive Sciences, 2021, 28, 218-227.	2.5	2
16	Microbial and transcriptional differences elucidate atopic dermatitis heterogeneity across skin sites. Allergy: European Journal of Allergy and Clinical Immunology, 2021, 76, 1173-1187.	5.7	16
17	DNA Methylation Levels in Mononuclear Leukocytes from the Mother and Her Child Are Associated with IgE Sensitization to Allergens in Early Life. International Journal of Molecular Sciences, 2021, 22, 801.	4.1	18
18	Nasal upregulation of CST1 in dog-sensitised children with severe allergic airway disease. ERJ Open Research, 2021, 7, 00917-2020.	2.6	8

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19	A missense variant in IFT122 associated with a canine model of retinitis pigmentosa. Human Genetics, 2021, 140, 1569-1579.	3.8	4
20	Toxicogenomic Profiling of 28 Nanomaterials in Mouse Airways. Advanced Science, 2021, 8, 2004588.	11.2	15
21	Dysfunction of complement receptors CR3 (CD11b/18) and CR4 (CD11c/18) in preâ€eclampsia: a genetic and functional study. BJOG: an International Journal of Obstetrics and Gynaecology, 2021, 128, 1282-1291.	2.3	9
22	High-resolution targeted bisulfite sequencing reveals blood cell type-specific DNA methylation patterns in IL13 and ORMDL3. Clinical Epigenetics, 2021, 13, 106.	4.1	0
23	HLA â€G expression correlates with histological grade but not with prognosis in colorectal carcinoma. Hla, 2021, 98, 213-217.	0.6	4
24	Discovery of widespread transcription initiation at microsatellites predictable by sequence-based deep neural network. Nature Communications, 2021, 12, 3297.	12.8	11
25	The role of CDHR3 in susceptibility to otitis media. Journal of Molecular Medicine, 2021, 99, 1571-1583.	3.9	4
26	Small RNA expression and miRNA modification dynamics in human oocytes and early embryos. Genome Research, 2021, 31, 1474-1485.	5.5	10
27	Dog colour patterns explained by modular promoters of ancient canid origin. Nature Ecology and Evolution, 2021, 5, 1415-1423.	7.8	24
28	Cystatin B-deficiency triggers ectopic histone H3 tail cleavage during neurogenesis. Neurobiology of Disease, 2021, 156, 105418.	4.4	13
29	Motor Function Deficits in the Estrogen Receptor Beta Knockout Mouse: Role on Excitatory Neurotransmission and Myelination in the Motor Cortex. Neuroendocrinology, 2021, 111, 27-44.	2.5	10
30	Embryonic LTR retrotransposons supply promoter modules to somatic tissues. Genome Research, 2021, 31, 1983-1993.	5.5	7
31	Generation of RNA sequencing libraries for transcriptome analysis of globin-rich tissues of the domestic dog. STAR Protocols, 2021, 2, 100995.	1.2	6
32	Acute wheeze-specific gene module shows correlation with vitamin D and asthma medication. European Respiratory Journal, 2020, 55, 1901330.	6.7	9
33	Nagashima-type palmoplantar keratosis in Finland caused by a SERPINB7 founder mutation. Journal of the American Academy of Dermatology, 2020, 83, 643-645.	1.2	14
34	Single-Cell Analysis of Human Ovarian Cortex Identifies Distinct Cell Populations But No Oogonial Stem Cells. Obstetrical and Gynecological Survey, 2020, 75, 354-355.	0.4	1
35	Multiparametric Profiling of Engineered Nanomaterials: Unmasking the Surface Coating Effect. Advanced Science, 2020, 7, 2002221.	11.2	24
36	Fetal HLA-G mediated immune tolerance and interferon response in preeclampsia. EBioMedicine, 2020, 59, 102872.	6.1	25

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37	Functional annotation of human long noncoding RNAs via molecular phenotyping. Genome Research, 2020, 30, 1060-1072.	5.5	109
38	PCSK2 expression in neuroendocrine tumors points to a midgut, pulmonary, or pheochromocytomaâ€“paraganglioma origin. Apmis, 2020, 128, 563-572.	2.0	5
39	Epigenetic alterations in skin homing CD4+CLA+ T cells of atopic dermatitis patients. Scientific Reports, 2020, 10, 18020.	3.3	23
40	Multi-omic studies on missense PLG variants in families with otitis media. Scientific Reports, 2020, 10, 15035.	3.3	4
41	Differentiation of ciliated human midbrain-derived LUHMES neurons. Journal of Cell Science, 2020, 133, .	2.0	6
42	Rare variants in dynein heavy chain genes in two individuals with situs inversus and developmental dyslexia: a case report. BMC Medical Genetics, 2020, 21, 87.	2.1	5
43	A putative silencer variant in a spontaneous canine model of retinitis pigmentosa. PLoS Genetics, 2020, 16, e1008659.	3.5	9
44	Epigenome-wide meta-analysis of blood DNA methylation in newborns and children identifies numerous loci related to gestational age. Genome Medicine, 2020, 12, 25.	8.2	81
45	Single-cell analysis of human ovarian cortex identifies distinct cell populations but no oogonial stem cells. Nature Communications, 2020, 11, 1147.	12.8	188
46	Novel Hemizygous IL2RG p.(Pro58Ser) Mutation Impairs IL-2 Receptor Complex Expression on Lymphocytes Causing X-Linked Combined Immunodeficiency. Journal of Clinical Immunology, 2020, 40, 503-514.	3.8	11
47	Congenital chloride diarrhea and Pendred syndrome: case report of siblings with two rare recessive disorders of SLC26 family genes. BMC Medical Genetics, 2020, 21, 79.	2.1	1
48	Dyslexia Candidate Gene and Ciliary Gene Expression Dynamics During Human Neuronal Differentiation. Molecular Neurobiology, 2020, 57, 2944-2958.	4.0	11
49	Phenotypic Variability with SLURP1 Mutations and Diffuse Palmoplantar Keratoderma. Acta Dermato-Venereologica, 2020, 100, adv00060.	1.3	0
50	DNA Methylation Trajectories During Pregnancy. Epigenetics Insights, 2019, 12, 251686571986709.	2.0	26
51	Guide for library design and bias correction for large-scale transcriptome studies using highly multiplexed RNAseq methods. BMC Bioinformatics, 2019, 20, 418.	2.6	9
52	Gain-of-function CEBPE mutation causes noncanonical autoinflammatory inflammasomopathy. Journal of Allergy and Clinical Immunology, 2019, 144, 1364-1376.	2.9	37
53	Dominant TOM1 mutation associated with combined immunodeficiency and autoimmune disease. Npj Genomic Medicine, 2019, 4, 14.	3.8	11
54	Microbe-host interplay in atopic dermatitis and psoriasis. Nature Communications, 2019, 10, 4703.	12.8	217

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55	ABOGenotype and Blood Type Are Associated with Otitis Media. Genetic Testing and Molecular Biomarkers, 2019, 23, 823-827.	0.7	4
56	Delineating the Healthy Human Skin UV ResponseÂand Early Induction of Interferon PathwayÂin Cutaneous Lupus Erythematosus. Journal of Investigative Dermatology, 2019, 139, 2058-2061.e4.	0.7	16
57	NET-CAGE characterizes the dynamics and topology of human transcribed cis-regulatory elements. Nature Genetics, 2019, 51, 1369-1379.	21.4	72
58	034 Characterization of novel TMEM173 mutation causing a lupus- and SAVI-like phenotype, modified by polymorphisms in TMEM173 and IFIH1. Journal of Investigative Dermatology, 2019, 139, S220.	0.7	0
59	Complement in Human Pre-implantation Embryos: Attack and Defense. Frontiers in Immunology, 2019, 10, 2234.	4.8	11
60	The human long non-coding RNA gene RMRP has pleiotropic effects and regulates cell-cycle progression at G2. Scientific Reports, 2019, 9, 13758.	3.3	22
61	Epigenome-wide meta-analysis of DNA methylation and childhood asthma. Journal of Allergy and Clinical Immunology, 2019, 143, 2062-2074.	2.9	147
62	Pleomorphic Adenoma Gene 1 Is Needed For Timely Zygotic Genome Activation and Early Embryo Development. Scientific Reports, 2019, 9, 8411.	3.3	16
63	Prenatal Particulate Air Pollution and DNA Methylation in Newborns: An Epigenome-Wide Meta-Analysis. Environmental Health Perspectives, 2019, 127, 57012.	6.0	111
64	Pool-seq driven proteogenomic database for Group G Streptococcus. Journal of Proteomics, 2019, 201, 84-92.	2.4	2
65	A preliminary transcriptome analysis suggests a transitory effect of vitamin D on mitochondrial function in obese young Finnish subjects. Endocrine Connections, 2019, 8, 559-570.	1.9	6
66	A2ML1and otitis media: novel variants, differential expression, and relevant pathways. Human Mutation, 2019, 40, 1156-1171.	2.5	10
67	Meta-analysis of epigenome-wide association studies in neonates reveals widespread differential DNA methylation associated with birthweight. Nature Communications, 2019, 10, 1893.	12.8	140
68	Cationic gold nanoparticles elicit mitochondrial dysfunction: a multi-omics study. Scientific Reports, 2019, 9, 4366.	3.3	54
69	Myoglobinopathy is an adult-onset autosomal dominant myopathy with characteristic sarcoplasmic inclusions. Nature Communications, 2019, 10, 1396.	12.8	11
70	Discovering heritable modes of MEG spectral power. Human Brain Mapping, 2019, 40, 1391-1402.	3.6	17
71	Nocturnal asthma is affected by genetic interactions between RORA and NPSR1. Pediatric Pulmonology, 2019, 54, 847-857.	2.0	9
72	Genome-wide association scan identifies new variants associated with a cognitive predictor of dyslexia. Translational Psychiatry, 2019, 9, 77.	4.8	82

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73	A multiethnic meta-analysis defined the association of rs12946942 with severe adolescent idiopathic scoliosis. Journal of Human Genetics, 2019, 64, 493-498.	2.3	11
74	Discovery of increased epidermal DNAH10 expression after regeneration of dermis in a randomized with-in person trial â€" reflections on psoriatic inflammation. Scientific Reports, 2019, 9, 19136.	3.3	3
75	Novel TMEM173 Mutation and the Role of Disease Modifying Alleles. Frontiers in Immunology, 2019, 10, 2770.	4.8	45
76	Impact of obesity on angiogenic and inflammatory markers in the Finnish Genetics of Pre-eclampsia Consortium (FINNPEC) cohort. International Journal of Obesity, 2019, 43, 1070-1081.	3.4	17
77	European families reveal MHC class I and II associations with autoimmune-mediated congenital heart block. Annals of the Rheumatic Diseases, 2018, 77, 1381-1382.	0.9	6
78	DNA methylation in childhood asthma: an epigenome-wide meta-analysis. Lancet Respiratory Medicine,the, 2018, 6, 379-388.	10.7	170
79	Reduced <scp>CDHR</scp>3 expression in children wheezing with rhinovirus. Pediatric Allergy and Immunology, 2018, 29, 200-206.	2.6	20
80	Characterization of the human RFX transcription factor family by regulatory and target gene analysis. BMC Genomics, 2018, 19, 181.	2.8	73
81	Metabolic and functional changes in transgender individuals following cross-sex hormone treatment: Design and methods of the GEnder Dysphoria Treatment in Sweden (GETS) study. Contemporary Clinical Trials Communications, 2018, 10, 148-153.	1.1	27
82	Mlh1 deficiency in normal mouse colon mucosa associates with chromosomally unstable colon cancer. Carcinogenesis, 2018, 39, 788-797.	2.8	18
83	An international meta-analysis confirms the association of BNC2 with adolescent idiopathic scoliosis. Scientific Reports, 2018, 8, 4730.	3.3	20
84	Cohort Profile: Pregnancy And Childhood Epigenetics (PACE) Consortium. International Journal of Epidemiology, 2018, 47, 22-23u.	1.9	105
85	Fetal Microsatellite in the Heme Oxygenase 1 Promoter Is Associated With Severe and Early-Onset Preeclampsia. Hypertension, 2018, 71, 95-102.	2.7	16
86	TAC-seq: targeted DNA and RNA sequencing for precise biomarker molecule counting. Npj Genomic Medicine, 2018, 3, 34.	3.8	26
87	Phylogenetic and mutational analyses of human LEUTX, a homeobox gene implicated in embryogenesis. Scientific Reports, 2018, 8, 17421.	3.3	17
88	A Non-Targeted LC-MS Profiling Reveals Elevated Levels of Carnitine Precursors and Trimethylated Compounds in the Cord Plasma of Pre-Eclamptic Infants. Scientific Reports, 2018, 8, 14616.	3.3	21
89	Singleâ€cell RNAâ€seq analysis reveals the platinum resistance gene COX7B and the surrogate marker CD63. Cancer Medicine, 2018, 7, 6193-6204.	2.8	29
90	FUT2 Variants Confer Susceptibility to Familial Otitis Media. American Journal of Human Genetics, 2018, 103, 679-690.	6.2	40

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91	Genome-wide meta-analysis and replication studies in multiple ethnicities identify novel adolescent idiopathic scoliosis susceptibility loci. Human Molecular Genetics, 2018, 27, 3986-3998.	2.9	34
92	Enrichment of rare copy number variation in children with developmental language disorder. Clinical Genetics, 2018, 94, 313-320.	2.0	19
93	Intracellular signalling pathways and cytoskeletal functions converge on the psoriasis candidate gene CCHCR1 expressed at P-bodies and centrosomes. BMC Genomics, 2018, 19, 432.	2.8	17
94	MANF protects human pancreatic beta cells against stress-induced cell death. Diabetologia, 2018, 61, 2202-2214.	6.3	66
95	The Psoriasis Risk Allele HLA-C*06:02 Shows Evidence of Association with Chronic or Recurrent Streptococcal Tonsillitis. Infection and Immunity, 2018, 86, .	2.2	17
96	A multi-ethnic meta-analysis confirms the association of rs6570507 with adolescent idiopathic scoliosis. Scientific Reports, 2018, 8, 11575.	3.3	33
97	Human pluripotent reprogramming with CRISPR activators. Nature Communications, 2018, 9, 2643.	12.8	128
98	Human ROBO1 regulates white matter structure in corpus callosum. Brain Structure and Function, 2017, 222, 707-716.	2.3	5
99	Damaging heterozygous mutations in NFKB1 lead to diverse immunologic phenotypes. Journal of Allergy and Clinical Immunology, 2017, 140, 782-796.	2.9	113
100	Ketogenic diet attenuates hepatopathy in mouse model of respiratory chain complex III deficiency caused by a Bcs1l mutation. Scientific Reports, 2017, 7, 957.	3.3	27
101	Estrogen receptor Î², a regulator of androgen receptor signaling in the mouse ventral prostate. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E3816-E3822.	7.1	53
102	Optimizing bone morphogenic protein 4-mediated human embryonic stem cell differentiation into trophoblast-like cells using fibroblast growth factor 2 and transforming growth factor-Î²/activin/nodal signalling inhibition. Reproductive BioMedicine Online, 2017, 35, 253-263.	2.4	11
103	Variants in the fetal genome near FLT1 are associated with risk of preeclampsia. Nature Genetics, 2017, 49, 1255-1260.	21.4	205
104	Sequence analysis of pooled bacterial samples enables identification of strain variation in group A streptococcus. Scientific Reports, 2017, 7, 45771.	3.3	3
105	Combined immunodeficiency and hypoglycemia associated with mutations in hypoxia upregulated 1. Journal of Allergy and Clinical Immunology, 2017, 139, 1391-1393.e11.	2.9	14
106	Genome-Wide Interaction Analysis of Air Pollution Exposure and Childhood Asthma with Functional Follow-up. American Journal of Respiratory and Critical Care Medicine, 2017, 195, 1373-1383.	5.6	107
107	A missense mutation in SLC26A3 is associated with human male subfertility and impaired activation of CFTR. Scientific Reports, 2017, 7, 14208.	3.3	20
108	The use of genotyping as a first step in molecular diagnosis of familial hypercholesterolemia. Atherosclerosis, 2017, 263, e62.	0.8	0

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109	FANTOM5 CAGE profiles of human and mouse samples. Scientific Data, 2017, 4, 170112.	5.3	195
110	Acute doses of caffeine shift nervous system cell expression profiles toward promotion of neuronal projection growth. Scientific Reports, 2017, 7, 11458.	3.3	14
111	Identification of NCAN as a candidate gene for developmental dyslexia. Scientific Reports, 2017, 7, 9294.	3.3	15
112	Hypomethylation of HOXA4 promoter is common in Silver-Russell syndrome and growth restriction and associates with stature in healthy children. Scientific Reports, 2017, 7, 15693.	3.3	12
113	Protective Low-Frequency Variants for Preeclampsia in the Fms Related Tyrosine Kinase 1 Gene in the Finnish Population. Hypertension, 2017, 70, 365-371.	2.7	37
114	Comprehensive mapping of the effects of azacitidine on DNA methylation, repressive/permissive histone marks and gene expression in primary cells from patients with MDS and MDS-related disease. Oncotarget, 2017, 8, 28812-28825.	1.8	42
115	The emerging landscape of dynamic DNA methylation in early childhood. BMC Genomics, 2017, 18, 25.	2.8	49
116	Analysis of Complement C3 Gene Reveals Susceptibility to Severe Preeclampsia. Frontiers in Immunology, 2017, 8, 589.	4.8	50
117	Unexpectedly High Prevalence of Common Variable Immunodeficiency in Finland. Frontiers in Immunology, 2017, 8, 1190.	4.8	49
118	Neuropeptide S (NPS) variants modify the signaling and risk effects of NPS Receptor 1 (NPSR1) variants in asthma. PLoS ONE, 2017, 12, e0176568.	2.5	12
119	Exposure to Traffic-Related Air Pollution and Serum Inflammatory Cytokines in Children. Environmental Health Perspectives, 2017, 125, 067007.	6.0	71
120	Epigenome-Wide Meta-Analysis of Methylation in Children Related to Prenatal NO ₂ Air Pollution Exposure. Environmental Health Perspectives, 2017, 125, 104-110.	6.0	176
121	CELSR2 is a candidate susceptibility gene in idiopathic scoliosis. PLoS ONE, 2017, 12, e0189591.	2.5	17
122	Molecular Characterization of Three Canine Models of Human Rare Bone Diseases: Caffey, van den Ende-Gupta, and Raine Syndromes. PLoS Genetics, 2016, 12, e1006037.	3.5	32
123	Evidence for genetic regulation of the human parietoâ€occipital 10â€Hz rhythmic activity. European Journal of Neuroscience, 2016, 44, 1963-1971.	2.6	20
124	NOD-like receptor signaling and inflammasome-related pathways are highlighted in psoriatic epidermis. Scientific Reports, 2016, 6, 22745.	3.3	63
125	The human PRD-like homeobox gene LEUTX has a central role in embryo genome activation. Development (Cambridge), 2016, 143, 3459-3469.	2.5	42
126	High-specificity bioinformatics framework for epigenomic profiling of discordant twins reveals specific and shared markers for ACPA and ACPA-positive rheumatoid arthritis. Genome Medicine, 2016, 8, 124.	8.2	27

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127	Cohort profile: the Finnish Genetics of Pre-eclampsia Consortium (FINNPEC). BMJ Open, 2016, 6, e013148.	1.9	32
128	DNA Methylation in Newborns and Maternal Smoking in Pregnancy: Genome-wide Consortium Meta-analysis. American Journal of Human Genetics, 2016, 98, 680-696.	6.2	717
129	The pruritus- and TH2-associated cytokine IL-31 promotes growth of sensory nerves. Journal of Allergy and Clinical Immunology, 2016, 138, 500-508.e24.	2.9	201
130	Preeclampsia does not share common risk alleles in 9p21 with coronary artery disease and type 2 diabetes. Annals of Medicine, 2016, 48, 330-336.	3.8	2
131	The Hydroxysteroid (17Î²) Dehydrogenase Family Gene HSD17B12 Is Involved in the Prostaglandin Synthesis Pathway, the Ovarian Function, and Regulation of Fertility. Endocrinology, 2016, 157, 3719-3730.	2.8	43
132	An RGS2 3â€²UTR polymorphism is associated with preeclampsia in overweight women. BMC Genetics, 2016, 17, 121.	2.7	13
133	The diagnosis of pre-eclampsia using two revised classifications in the Finnish Pre-eclampsia Consortium (FINNPEC) cohort. BMC Pregnancy and Childbirth, 2016, 16, 221.	2.4	32
134	Ciliary dyslexia candidate genes DYX1C1 and DCDC2 are regulated by Regulatory Factor X (RFX) transcription factors through Xâ€box promoter motifs. FASEB Journal, 2016, 30, 3578-3587.	0.5	28
135	Whole-Exome Sequencing Suggests LAMB3 as a Susceptibility Gene for Morbid Obesity. Diabetes, 2016, 65, 2980-2989.	0.6	16
136	Characterization and target genes of nine human PRD-like homeobox domain genes expressed exclusively in early embryos. Scientific Reports, 2016, 6, 28995.	3.3	33
137	Globin mRNA reduction for whole-blood transcriptome sequencing. Scientific Reports, 2016, 6, 31584.	3.3	42
138	Exome sequencing in pooled DNA samples to identify maternal pre-eclampsia risk variants. Scientific Reports, 2016, 6, 29085.	3.3	19
139	Investigation of rare and low-frequency variants using high-throughput sequencing with pooled DNA samples. Scientific Reports, 2016, 6, 33256.	3.3	13
140	Genome-wide association analysis reveals variants on chromosome 19 that contribute to childhood risk of chronic otitis media with effusion. Scientific Reports, 2016, 6, 33240.	3.3	21
141	Targeted high-throughput sequencing of candidate genes for chronic obstructive pulmonary disease. BMC Pulmonary Medicine, 2016, 16, 146.	2.0	12
142	Genomic sequencing of a dyslexia susceptibility haplotype encompassing ROBO1. Journal of Neurodevelopmental Disorders, 2016, 8, 4.	3.1	8
143	Single-cell transcriptome analysis of endometrial tissue. Human Reproduction, 2016, 31, 844-853.	0.9	95
144	Increased YKL-40 and Chitotriosidase in Asthma and Chronic Obstructive Pulmonary Disease. American Journal of Respiratory and Critical Care Medicine, 2016, 193, 131-142.	5.6	107

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145	The Salivary Scavenger and Agglutinin (SALSA) in Healthy and Complicated Pregnancy. PLoS ONE, 2016, 11, e0147867.	2.5	14
146	Gene-Expression Profiling Suggests Impaired Signaling via the Interferon Pathway in Cstb-/- Microglia. PLoS ONE, 2016, 11, e0158195.	2.5	9
147	Identification of Novel Transcribed Regions in Zebrafish (Danio rerio) Using RNA-Sequencing. PLoS ONE, 2016, 11, e0160197.	2.5	3
148	meQTL analysis of asthma GWAS loci and DNA methylation. , 2016, , .		0
149	Differentially methylated genes related to gestational age are also expressed during fetal lung development. , 2016, , .		0
150	Aberrant splicing of genes involved in haemoglobin synthesis and impaired terminal erythroid maturation in <scp>SF</scp>3B1 mutated refractory anaemia with ring sideroblasts. British Journal of Haematology, 2015, 171, 478-490.	2.5	37
151	Gene expression profiling of pre-eclamptic placentae by RNA sequencing. Scientific Reports, 2015, 5, 14107.	3.3	89
152	Autoimmunity, hypogammaglobulinemia, lymphoproliferation, and mycobacterial disease in patients with activating mutations in STAT3. Blood, 2015, 125, 639-648.	1.4	229
153	High-throughput mutational screening adds clinically important information in myelodysplastic syndromes and secondary or therapy-related acute myeloid leukemia. Haematologica, 2015, 100, e223-e225.	3.5	12
154	Genomeâ€wide association study identifies new susceptibility loci for cutaneous lupus erythematosus. Experimental Dermatology, 2015, 24, 510-515.	2.9	66
155	Discovery of Molecular Markers to Discriminate Corneal Endothelial Cells in the Human Body. PLoS ONE, 2015, 10, e0117581.	2.5	28
156	The Constrained Maximal Expression Level Owing to Haploidy Shapes Gene Content on the Mammalian X Chromosome. PLoS Biology, 2015, 13, e1002315.	5.6	32
157	Application of Gene Expression Trajectories Initiated from ErbB Receptor Activation Highlights the Dynamics of Divergent Promoter Usage. PLoS ONE, 2015, 10, e0144176.	2.5	1
158	Polymorphisms in DCDC2 and S100B associate with developmental dyslexia. Journal of Human Genetics, 2015, 60, 399-401.	2.3	23
159	Associations between the 17q21 region and allergic rhinitis in 5 birth cohorts. Journal of Allergy and Clinical Immunology, 2015, 135, 573-576.e5.	2.9	15
160	Predictors of recurrent cellulitis in five years. Clinical risk factors and the role of PTX3 and CRP. Journal of Infection, 2015, 70, 467-473.	3.3	15
161	The statistical geometry of transcriptome divergence in cell-type evolution and cancer. Nature Communications, 2015, 6, 6066.	12.8	49
162	DCDC2 Mutations Cause a Renal-Hepatic Ciliopathy by Disrupting Wnt Signaling. American Journal of Human Genetics, 2015, 96, 81-92.	6.2	98

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163	A Missense Change in the ATG4D Gene Links Aberrant Autophagy to a Neurodegenerative Vacuolar Storage Disease. PLoS Genetics, 2015, 11, e1005169.	3.5	48
164	Transcriptome analysis of controlled and therapy-resistant childhood asthma reveals distinct gene expression profiles. Journal of Allergy and Clinical Immunology, 2015, 136, 638-648.	2.9	59
165	Helsinki alert of biodiversity and health. Annals of Medicine, 2015, 47, 218-225.	3.8	95
166	135 IN VITRO AZACITIDINE CULTURE INDUCES DNA DEMETHYLATION AND INCREASED MRNA-LEVELS IN PRIMARY MDS PROGENITOR CELLS. Leukemia Research, 2015, 39, S69.	0.8	1
167	Evidence of streptococcal origin of acute non-necrotising cellulitis: a serological study. European Journal of Clinical Microbiology and Infectious Diseases, 2015, 34, 669-672.	2.9	23
168	Age-associated DNA methylation changes in immune genes, histone modifiers and chromatin remodeling factors within 5Âyears after birth in human blood leukocytes. Clinical Epigenetics, 2015, 7, 34.	4.1	65
169	Mutation in CEP63 co-segregating with developmental dyslexia in a Swedish family. Human Genetics, 2015, 134, 1239-1248.	3.8	23
170	Gene expression analysis of skin grafts and cultured keratinocytes using synthetic RNA normalization reveals insights into differentiation and growth control. BMC Genomics, 2015, 16, 476.	2.8	21
171	Candidate gene analysis and exome sequencing confirm LBX1 as a susceptibility gene for idiopathic scoliosis. Spine Journal, 2015, 15, 2239-2246.	1.3	53
172	Novel PRD-like homeodomain transcription factors and retrotransposon elements in early human development. Nature Communications, 2015, 6, 8207.	12.8	100
173	GIMAP GTPase Family Genes: Potential Modifiers in Autoimmune Diabetes, Asthma, and Allergy. Journal of Immunology, 2015, 194, 5885-5894.	0.8	30
174	Variant Profiling of Candidate Genes in Pancreatic Ductal Adenocarcinoma. Clinical Chemistry, 2015, 61, 1408-1416.	3.2	21
175	CTNND2â€"a candidate gene for reading problems and mild intellectual disability. Journal of Medical Genetics, 2015, 52, 111-122.	3.2	35
176	Exome sequencing followed by genotyping suggests SYPL2 as a susceptibility gene for morbid obesity. European Journal of Human Genetics, 2015, 23, 1216-1222.	2.8	21
177	Risk of childhood asthma is associated with CpG-site polymorphisms, regional DNA methylation and mRNA levels at the GSDMB/ORMDL3 locus. Human Molecular Genetics, 2015, 24, 875-890.	2.9	66
178	Dominant NFKB1 Mutations Cause Antibody Deficiency and Autoinflammatory Episodes. Blood, 2015, 126, 206-206.	1.4	1
179	An integrative genomics approach identifies new asthma pathways related to air pollution exposure. , 2015, , .		1
180	Genetic Analysis of Membrane Cofactor Protein (CD46) of the Complement System in Women with and without Preeclamptic Pregnancies. PLoS ONE, 2015, 10, e0117840.	2.5	13

#	Article	IF	Citations
181	Predisposition to Childhood Otitis Media and Genetic Polymorphisms within the Toll-Like Receptor 4 (TLR4) Locus. PLoS ONE, 2015, 10, e0132551.	2.5	35
182	Ceruloplasmin Is a Novel Adipokine Which Is Overexpressed in Adipose Tissue of Obese Subjects and in Obesity-Associated Cancer Cells. PLoS ONE, 2014, 9, e80274.	2.5	50
183	Differences in Gene Expression between Mouse and Human for Dynamically Regulated Genes in Early Embryo. PLoS ONE, 2014, 9, e102949.	2.5	25
184	Differential roles of epigenetic changes and Foxp3 expression in regulatory T cell-specific transcriptional regulation. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 5289-5294.	7.1	111
185	The Roots of Autism and ADHD Twin Study in Sweden (RATSS). Twin Research and Human Genetics, 2014, 17, 164-176.	0.6	62
186	Microsatellite Polymorphism in the Heme Oxygenase-1 Promoter Is Associated With Nonsevere and Late-Onset Preeclampsia. Hypertension, 2014, 64, 172-177.	2.7	37
187	The molecular genetics and neurobiology of developmental dyslexia as model of a complex phenotype. Biochemical and Biophysical Research Communications, 2014, 452, 236-243.	2.1	72
188	<scp>NPSR</scp>1 polymorphisms influence recurrent abdominal pain in children: a populationâ€based study. Neurogastroenterology and Motility, 2014, 26, 1417-1425.	3.0	16
189	Polymorphisms in the Dopamine Receptor 2 Gene Region Influence Improvements during Working Memory Training in Children and Adolescents. Journal of Cognitive Neuroscience, 2014, 26, 54-62.	2.3	58
190	Differentially methylated regions in maternal and paternal uniparental disomy for chromosome 7. Epigenetics, 2014, 9, 351-365.	2.7	41
191	DCDC2Polymorphism Is Associated with Left Temporoparietal Gray and White Matter Structures during Development. Journal of Neuroscience, 2014, 34, 14455-14462.	3.6	35
192	Genetic analysis of dyslexia candidate genes in the European cross-linguistic NeuroDys cohort. European Journal of Human Genetics, 2014, 22, 675-680.	2.8	59
193	A promoter-level mammalian expression atlas. Nature, 2014, 507, 462-470.	27.8	1,838
194	An atlas of active enhancers across human cell types and tissues. Nature, 2014, 507, 455-461.	27.8	2,269
195	Azacitidine induces profound genome-wide hypomethylation in primary myelodysplastic bone marrow cultures but may also reduce histone acetylation. Leukemia, 2014, 28, 411-413.	7.2	14
196	Clonal culturing of human embryonic stem cells on laminin-521/E-cadherin matrix in defined and xeno-free environment. Nature Communications, 2014, 5, 3195.	12.8	248
197	CCL2 enhances pluripotency of human induced pluripotent stem cells by activating hypoxia related genes. Scientific Reports, 2014, 4, 5228.	3.3	21
198	Activating germline mutations in STAT3 cause early-onset multi-organ autoimmune disease. Nature Genetics, 2014, 46, 812-814.	21.4	411

#	Article	IF	Citations
199	Neuropeptide S receptor 1 (NPSR1) activates cancer-related pathways and is widely expressed in neuroendocrine tumors. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2014, 465, 173-183.	2.8	19
200	Enhanced expression of neuropeptide S (NPS) receptor in eosinophils from severe asthmatics and subjects with total IgE above 100IU/ml. Peptides, 2014, 51, 100-109.	2.4	17
201	Dominant Mutations in GRHL3 Cause Van der Woude Syndrome and Disrupt Oral Periderm Development. American Journal of Human Genetics, 2014, 94, 23-32.	6.2	195
202	Sequencing of breast cancer stem cell populations indicates a dynamic conversion between differentiation states in vivo. Breast Cancer Research, 2014, 16, R72.	5.0	54
203	Association and Mutation Analyses of the IRF6 Gene in Families with Nonsyndromic and Syndromic Cleft Lip and/or Cleft Palate. Cleft Palate-Craniofacial Journal, 2014, 51, 49-55.	0.9	20
204	Welander distal myopathy is caused by a mutation in the RNAâ€binding protein TIA1. Annals of Neurology, 2013, 73, 500-509.	5.3	118
205	Drug repositioning: a machine-learning approach through data integration. Journal of Cheminformatics, 2013, 5, 30.	6.1	263
206	Introduction of complementary foods in infancy and atopic sensitization at the age of 5Âyears: timing and food diversity in a Finnish birth cohort. Allergy: European Journal of Allergy and Clinical Immunology, 2013, 68, 507-516.	5.7	77
207	Molecular Networks of DYX1C1 Gene Show Connection to Neuronal Migration Genes and Cytoskeletal Proteins. Biological Psychiatry, 2013, 73, 583-590.	1.3	38
208	Epigenome-wide association data implicate DNA methylation as an intermediary of genetic risk in rheumatoid arthritis. Nature Biotechnology, 2013, 31, 142-147.	17.5	874
209	P-005 Aberrant splicing during erythroid differentiation in SF3B1 mutated sideroblastic anemia. Leukemia Research, 2013, 37, S24-S25.	0.8	0
210	Transcriptome analysis reveals upregulation of bitter taste receptors in severe asthmatics. European Respiratory Journal, 2013, 42, 65-78.	6.7	130
211	Timing of infant feeding in relation to childhood asthma and allergic diseases. Journal of Allergy and Clinical Immunology, 2013, 131, 78-86.	2.9	116
212	Recurrent cellulitis with benzathine penicillin prophylaxis is associated with diabetes and psoriasis. European Journal of Clinical Microbiology and Infectious Diseases, 2013, 32, 369-372.	2.9	12
213	A follow-up linkage study of Finnish pre-eclampsia families identifies a new fetal susceptibility locus on chromosome 18. European Journal of Human Genetics, 2013, 21, 1024-1026.	2.8	26
214	SAMstrt: statistical test for differential expression in single-cell transcriptome with spike-in normalization. Bioinformatics, 2013, 29, 2943-2945.	4.1	114
215	A genome-wide association study of atopic dermatitis identifies loci with overlapping effects on asthma and psoriasis. Human Molecular Genetics, 2013, 22, 4841-4856.	2.9	202
216	Genetic dissection of the pre-eclampsia susceptibility locus on chromosome 2q22 reveals shared novel risk factors for cardiovascular disease. Molecular Human Reproduction, 2013, 19, 423-437.	2.8	54

#	Article	IF	Citations
217	Genes identified in Asian SLE GWASs are also associated with SLE in Caucasian populations. European Journal of Human Genetics, 2013, 21, 994-999.	2.8	90
218	DNA Methylation in the Neuropeptide S Receptor 1 (NPSR1) Promoter in Relation to Asthma and Environmental Factors. PLoS ONE, 2013, 8, e53877.	2.5	35
219	Genetic Susceptibility to Non-Necrotizing Erysipelas/Cellulitis. PLoS ONE, 2013, 8, e56225.	2.5	15
220	Interaction between Retinoid Acid Receptor-Related Orphan Receptor Alpha (RORA) and Neuropeptide S Receptor 1 (NPSR1) in Asthma. PLoS ONE, 2013, 8, e60111.	2.5	28
221	Ultra-Rare Mutation in Long-Range Enhancer Predisposes to Thyroid Carcinoma with High Penetrance. PLoS ONE, 2013, 8, e61920.	2.5	36
222	Altered Methylation of IGF2 Locus 20 Years after Preterm Birth at Very Low Birth Weight. PLoS ONE, 2013, 8, e67379.	2.5	21
223	The Zebrafish Orthologue of the Dyslexia Candidate Gene DYX1C1 Is Essential for Cilia Growth and Function. PLoS ONE, 2013, 8, e63123.	2.5	60
224	Rule-Based Models of the Interplay between Genetic and Environmental Factors in Childhood Allergy. PLoS ONE, 2013, 8, e80080.	2.5	18
225	Erythropoiesis In SF3B1 Mutated RARS Is Disrupted During Terminal Erythroid Maturation. Blood, 2013, 122, 2408-2408.	1.4	0
226	Replication of GWAS-identified systemic lupus erythematosus susceptibility genes affirms B-cell receptor pathway signalling and strengthens the role of IRF5 in disease susceptibility in a Northern European population. Rheumatology, 2012, 51, 87-92.	1.9	68
227	The rs3743205 SNP Is Important for the Regulation of the Dyslexia Candidate GeneDYX1C1by Estrogen Receptor Î² and DNA Methylation. Molecular Endocrinology, 2012, 26, 619-629.	3.7	25
228	Research Resource: Interactome of Human Embryo Implantation: Identification of Gene Expression Pathways, Regulation, and Integrated Regulatory Networks. Molecular Endocrinology, 2012, 26, 203-217.	3.7	107
229	In Vivo Differentiated Human Embryonic Stem Cells Can Acquire Chromosomal Aberrations More Frequently Than In Vitro During the Same Period. Stem Cells and Development, 2012, 21, 3363-3371.	2.1	8
230	Working memory brain activity and capacity link MAOA polymorphism to aggressive behavior during development. Translational Psychiatry, 2012, 2, e85-e85.	4.8	36
231	Conditional analysis identifies three novel major histocompatibility complex loci associated with psoriasis. Human Molecular Genetics, 2012, 21, 5185-5192.	2.9	58
232	Human ROBO1 Regulates Interaural Interaction in Auditory Pathways. Journal of Neuroscience, 2012, 32, 966-971.	3.6	54
233	Genome-Wide Linkage Analysis of Human Auditory Cortical Activation Suggests Distinct Loci on Chromosomes 2, 3, and 8. Journal of Neuroscience, 2012, 32, 14511-14518.	3.6	21
234	Three Dyslexia Susceptibility Genes, DYX1C1, DCDC2, and KIAA0319, Affect Temporo-Parietal White Matter Structure. Biological Psychiatry, 2012, 72, 671-676.	1.3	133

#	Article	IF	Citations
235	Genetic background and the risk of otitis media. International Journal of Pediatric Otorhinolaryngology, 2012, 76, 41-44.	1.0	35
236	Identification of 15 new psoriasis susceptibility loci highlights the role of innate immunity. Nature Genetics, 2012, 44, 1341-1348.	21.4	848
237	Current Knowledge of the Genetics of Otitis Media. Current Allergy and Asthma Reports, 2012, 12, 582-589.	5.3	8
238	A novel screening method detects herpesviral DNA in the idiopathic pulmonary fibrosis lung. Annals of Medicine, 2012, 44, 178-186.	3.8	31
239	Dopamine, working memory, and training induced plasticity: Implications for developmental research.. Developmental Psychology, 2012, 48, 836-843.	1.6	58
240	Genome-Wide Association Scan Identifies a Risk Locus for Preeclampsia on 2q14, Near the Inhibin, Beta B Gene. PLoS ONE, 2012, 7, e33666.	2.5	110
241	Centrosomal Localization of the Psoriasis Candidate Gene Product, CCHCR1, Supports a Role in Cytoskeletal Organization. PLoS ONE, 2012, 7, e49920.	2.5	26
242	Acquisition of Complement Factor H Is Important for Pathogenesis ofStreptococcus pyogenesInfections: Evidence from Bacterial In Vitro Survival and Human Genetic Association. Journal of Immunology, 2012, 188, 426-435.	0.8	36
243	The Aromatase Gene CYP19A1: Several Genetic and Functional Lines of Evidence Supporting a Role in Reading, Speech and Language. Behavior Genetics, 2012, 42, 509-527.	2.1	60
244	<scp>DNA</scp> methylation levels within the <scp>CD</scp>14 promoter region are lower in placentas of mothers living on a farm. Allergy: European Journal of Allergy and Clinical Immunology, 2012, 67, 895-903.	5.7	39
245	aThe dyslexia candidate gene DYX1C1 is a potential marker of poor survival in breast cancer. BMC Cancer, 2012, 12, 79.	2.6	5
246	Differential DNA Methylation in Purified Human Blood Cells: Implications for Cell Lineage and Studies on Disease Susceptibility. PLoS ONE, 2012, 7, e41361.	2.5	860
247	The Dyslexia Candidate Locus on 2p12 Is Associated with General Cognitive Ability and White Matter Structure. PLoS ONE, 2012, 7, e50321.	2.5	41
248	Influence of the COMT Genotype on Working Memory and Brain Activity Changes During Development. Biological Psychiatry, 2011, 70, 222-229.	1.3	139
249	A comprehensive analysis of the COL29A1 gene does not support a role in eczema. Journal of Allergy and Clinical Immunology, 2011, 127, 1187-1194.e7.	2.9	15
250	Methylation of H19 and its imprinted control region (H19 ICR1) in MÃ¼llerian aplasia. Fertility and Sterility, 2011, 95, 2703-2706.	1.0	12
251	P.44 FV Leiden as risk factor for preterm birth â€“ a population-based nested case-control study. Thrombosis Research, 2011, 127, S140.	1.7	0
252	Genome Wide Transcriptome Analysis Suggests Novel Mechanisms In Severe Childhood Asthma. , 2011, , .		0

#	Article	IF	Citations
253	Swedish Population Substructure Revealed by Genome-Wide Single Nucleotide Polymorphism Data. PLoS ONE, 2011, 6, e16747.	2.5	39
254	Increased Expression of the Dyslexia Candidate Gene DCDC2 Affects Length and Signaling of Primary Cilia in Neurons. PLoS ONE, 2011, 6, e20580.	2.5	113
255	Multiple Polymorphisms Affect Expression and Function of the Neuropeptide S Receptor (NPSR1). PLoS ONE, 2011, 6, e29523.	2.5	30
256	Dna Methylation In The Promoter Of Neuropeptide S Receptor 1 (NPSR1) In Relation To Respiratory Syndromes. , 2011, , .		0
257	INVITED SESSION, SESSION 37: COMPREHENSIVE MOLECULAR ANALYSIS OF OOCYTES AND EMBRYOS, Tuesday 5 July 2011 11:45 - 12:45. Human Reproduction, 2011, 26, i56-i56.	0.9	3
258	Genomeâ€wide analysis of extended pedigrees confirms IL2â€“IL21 linkage and shows additional regions of interest potentially influencing coeliac disease risk. Tissue Antigens, 2011, 78, 428-437.	1.0	11
259	FactorÂV Leiden as a risk factor for preterm birth - a population-based nested case-control study. Journal of Thrombosis and Haemostasis, 2011, 9, 71-78.	3.8	20
260	Multiple independent variants in 6q21-22 associated with susceptibility to celiac disease in the Dutch, Finnish and Hungarian populations. European Journal of Human Genetics, 2011, 19, 682-686.	2.8	7
261	Glucocorticoid receptor gene haplotype predicts increased risk of hospital admission for depressive disorders in the Helsinki birth cohort study. Journal of Psychiatric Research, 2011, 45, 1160-1164.	3.1	16
262	LifeGeneâ€”a large prospective population-based study of global relevance. European Journal of Epidemiology, 2011, 26, 67-77.	5.7	91
263	SNP Variations in the 7q33 Region Containing DGKI are Associated with Dyslexia in the Finnish and German Populations. Behavior Genetics, 2011, 41, 134-140.	2.1	25
264	The asthma candidate gene NPSR1 mediates isoform specific downstream signalling. BMC Pulmonary Medicine, 2011, 11, 39.	2.0	20
265	Genome wide association study identifies KCNMA1contributing to human obesity. BMC Medical Genomics, 2011, 4, 51.	1.5	87
266	The zebrafish transcriptome during early development. BMC Developmental Biology, 2011, 11, 30.	2.1	108
267	Molecular genetics and molecular biology of dyslexia. Wiley Interdisciplinary Reviews: Cognitive Science, 2011, 2, 441-448.	2.8	14
268	Genomic strategy identifies a missense mutation in WDâ€repeat domain 65 (WDR65) in an individual with Van der Woude syndrome. American Journal of Medical Genetics, Part A, 2011, 155, 1314-1321.	1.2	12
269	Update on SLC26A3 mutations in congenital chloride diarrhea. Human Mutation, 2011, 32, 715-722.	2.5	103
270	Meta-Analysis Confirms the LCE3C_LCE3B Deletion as a Risk Factor for Psoriasis in Several Ethnic Groups and Finds Interaction with HLA-Cw6. Journal of Investigative Dermatology, 2011, 131, 1105-1109.	0.7	89

#	Article	IF	Citations
271	Study of Estrogen Receptor-Î± and Receptor-Î² Gene Polymorphisms on Alzheimer's Disease. Journal of Alzheimer's Disease, 2011, 26, 431-439.	2.6	18
272	Genetic Association and Gene Expression Analysis IdentifyFGFR1as a New Susceptibility Gene for Human Obesity. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E962-E966.	3.6	25
273	Co-localization of neural cell adhesion molecule and fibroblast growth factor receptor 2 in early embryo development. International Journal of Developmental Biology, 2011, 55, 313-319.	0.6	9
274	Neuropeptide S receptor 1 expression in the intestine and skin â€“ putative role in peptide hormone secretion. Neurogastroenterology and Motility, 2010, 22, 79.	3.0	22
275	Airway Inflammation In COPD And Asthma Is Associated With Elevated Serum Chitotriosidase Activity In A Genotype Dependent Manner. , 2010, , .		0
276	Genetic evidence of multiple loci in dystocia - difficult labour. BMC Medical Genetics, 2010, 11, 105.	2.1	15
277	Factors predisposing to acute and recurrent bacterial non-necrotizing cellulitis in hospitalized patients: a prospective caseâ€“control study. Clinical Microbiology and Infection, 2010, 16, 729-734.	6.0	91
278	Tyrosine kinase 2 and interferon regulatory factor 5 polymorphisms are associated with discoid and subacute cutaneous lupus erythematosus. Experimental Dermatology, 2010, 19, 123-131.	2.9	77
279	Molecular genetic and epigenetic analysis ofNCX2/SLC8A2at 19q13.3 in human gliomas. Neuropathology and Applied Neurobiology, 2010, 36, 198-210.	3.2	13
280	Interaction between early maternal smoking and variants in TNF and GSTP1 in childhood wheezing. Clinical and Experimental Allergy, 2010, 40, 458-467.	2.9	31
281	Genomic landscape of positive natural selection in Northern European populations. European Journal of Human Genetics, 2010, 18, 471-478.	2.8	31
282	Genome scan for loci regulating HDL cholesterol levels in Finnish extended pedigrees with early coronary heart disease. European Journal of Human Genetics, 2010, 18, 604-613.	2.8	7
283	Meta-analysis of 20 genome-wide linkage studies evidenced new regions linked to asthma and atopy. European Journal of Human Genetics, 2010, 18, 700-706.	2.8	54
284	A genome-wide association study identifies new psoriasis susceptibility loci and an interaction between HLA-C and ERAP1. Nature Genetics, 2010, 42, 985-990.	21.4	918
285	Analysis of Neuropeptide S Receptor Gene (NPSR1) Polymorphism in Rheumatoid Arthritis. PLoS ONE, 2010, 5, e9315.	2.5	30
286	ELMOD2, a candidate gene for idiopathic pulmonary fibrosis, regulates antiviral responses. FASEB Journal, 2010, 24, 1167-1177.	0.5	65
287	Dr. Hellquist, et al reply. Journal of Rheumatology, 2010, 37, 678.1-678.	2.0	0
288	Variation in STAT4 is associated with systemic lupus erythematosus in a Finnish family cohort. Annals of the Rheumatic Diseases, 2010, 69, 883-886.	0.9	17

#	Article	IF	Citations
289	Submicroscopic genomic alterations in Silver-Russell syndrome and Silver-Russell-like patients. Journal of Medical Genetics, 2010, 47, 816-822.	3.2	76
290	Assessment of the Neuropeptide S System in Anxiety Disorders. Biological Psychiatry, 2010, 68, 474-483.	1.3	79
291	The SNAP25 Gene Is Linked to Working Memory Capacity and Maturation of the Posterior Cingulate Cortex During Childhood. Biological Psychiatry, 2010, 68, 1120-1125.	1.3	56
292	A genome-wide association scan on estrogen receptor-negative breast cancer. Breast Cancer Research, 2010, 12, R93.	5.0	35
293	Genetics of complex disorders. Biochemical and Biophysical Research Communications, 2010, 396, 143-146.	2.1	10
294	Factor V Leiden as risk factor for unexplained stillbirth â€“ a population-based nested case-control study. Thrombosis Research, 2010, 125, 505-510.	1.7	11
295	A Large-Scale, Consortium-Based Genomewide Association Study of Asthma. New England Journal of Medicine, 2010, 363, 1211-1221.	27.0	1,762
296	A Teratocarcinoma-Like Human Embryonic Stem Cell (hESC) Line and Four hESC Lines Reveal Potentially Oncogenic Genomic Changes. PLoS ONE, 2010, 5, e10263.	2.5	45
297	Polymorphisms of the ITGAM Gene Confer Higher Risk of Discoid Cutaneous Than of Systemic Lupus Erythematosus. PLoS ONE, 2010, 5, e14212.	2.5	39
298	Neuropeptide S Receptor 1: an Asthma Susceptibility Gene. , 2010, , 191-205.		1
299	Transcriptome Analysis of Differentiating Erythroid Progenitors in Refractory Anemia with Ringed Sideroblasts. Blood, 2010, 116, 1864-1864.	1.4	0
300	CCHCR1 Is Up-Regulated in Skin Cancer and Associated with EGFR Expression. PLoS ONE, 2009, 4, e6030.	2.5	33
301	Transcriptome Profiling of Human Pre-Implantation Development. PLoS ONE, 2009, 4, e7844.	2.5	103
302	Evidence for Genetic Association and Interaction Between the TYK2 and IRF5 Genes in Systemic Lupus Erythematosus. Journal of Rheumatology, 2009, 36, 1631-1638.	2.0	71
303	Functional interaction of DYX1C1 with estrogen receptors suggests involvement of hormonal pathways in dyslexia. Human Molecular Genetics, 2009, 18, 2802-2812.	2.9	56
304	MMP12,Lung Function, and COPD in High-Risk Populations. New England Journal of Medicine, 2009, 361, 2599-2608.	27.0	315
305	Clinically Distinct Epigenetic Subgroups in Silver-Russell Syndrome: The Degree ofH19Hypomethylation Associates with Phenotype Severity and Genital and Skeletal Anomalies. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 579-587.	3.6	78
306	ROCK2 allelic variants are not associated with pre-eclampsia susceptibility in the Finnish population. Molecular Human Reproduction, 2009, 15, 443-449.	2.8	7

#	Article	IF	Citations
307	Association study of the IL18RAP locus in three European populations with coeliac disease. Human Molecular Genetics, 2009, 18, 1148-1155.	2.9	29
308	A Susceptibility Locus for Papillary Thyroid Carcinoma on Chromosome 8q24. Cancer Research, 2009, 69, 625-631.	0.9	133
309	No Association Between the Eczema Genes COL29A1 and IL31 and Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2009, 15, 961-962.	1.9	3
310	PepT1 oligopeptide transporter (SLC15A1) gene polymorphism in inflammatory bowel disease. Inflammatory Bowel Diseases, 2009, 15, 1562-1569.	1.9	51
311	IL23R in the Swedish, Finnish, Hungarian and Italian populations: association with IBD and psoriasis, and linkage to celiac disease. BMC Medical Genetics, 2009, 10, 8.	2.1	61
312	Association of psoriasis to PGLYRP and SPRR genes at PSORS4 locus on 1q shows heterogeneity between Finnish, Swedish and Irish families. Experimental Dermatology, 2009, 18, 109-115.	2.9	37
313	Fine mapping of the CELIAC2 locus on chromosome 5q31â€q33 in the Finnish and Hungarian populations. Tissue Antigens, 2009, 74, 408-416.	1.0	16
314	Population Structure in Contemporary Swedenâ€"A Yâ€€chromosomal and Mitochondrial DNA Analysis. Annals of Human Genetics, 2009, 73, 61-73.	0.8	36
315	Blood group AB and factor V Leiden as risk factors for pre-eclampsia: A population-based nested case-control study. Thrombosis Research, 2009, 124, 167-173.	1.7	55
316	Further evidence for DYX1C1 as a susceptibility factor for dyslexia. Psychiatric Genetics, 2009, 19, 59-63.	1.1	62
317	Identification of MAMDC1 as a Candidate Susceptibility Gene for Systemic Lupus Erythematosus (SLE). PLoS ONE, 2009, 4, e8037.	2.5	14
318	Population substructure in Finland and Sweden revealed by the use of spatial coordinates and a small number of unlinked autosomal SNPs. BMC Genetics, 2008, 9, 54.	2.7	31
319	Association analysis of positional obesity candidate genes based on integrated data from transcriptomics and linkage analysis. International Journal of Obesity, 2008, 32, 816-825.	3.4	17
320	Novel and recurrent STAT3 mutations in hyper-IgE syndrome patients from different ethnic groups. Molecular Immunology, 2008, 46, 202-206.	2.2	82
321	D.P.3.07 Welander distal myopathy: The evasive gene. Neuromuscular Disorders, 2008, 18, 767.	0.6	0
322	Restriction Siteâ€"Specific Methylation Studies of Imprinted Genes with Quantitative Real-Time PCR. Clinical Chemistry, 2008, 54, 491-499.	3.2	31
323	The complex of TFIIâ€‰, PARP1, and SFPQ proteins regulates the DYX1C1 gene implicated in neuronal migration and dyslexia. FASEB Journal, 2008, 22, 3001-3009.	0.5	70
324	The protective effect of farm animal exposure on childhood allergy is modified by NPSR1 polymorphisms. Journal of Medical Genetics, 2008, 46, 159-167.	3.2	30

#	Article	IF	Citations
325	Biological and genetic interaction between Tenascin C and Neuropeptide S receptor 1 in allergic diseases. Human Molecular Genetics, 2008, 17, 1673-1682.	2.9	28
326	Acute Bacterial, Nonnecrotizing Cellulitis in Finland: Microbiological Findings. Clinical Infectious Diseases, 2008, 46, 855-861.	5.8	56
327	The impact of sodium chloride and volume depletion in the chronic kidney disease of congenital chloride diarrhea. Kidney International, 2008, 74, 1085-1093.	5.2	33
328	Familial non-syndromic cleft lip and palate--analysis of the IRF6 gene and clinical phenotypes. European Journal of Orthodontics, 2008, 30, 169-175.	2.4	33
329	Overview of the SLC26 Family and Associated Diseases. Novartis Foundation Symposium, 2008, , 2-18.	1.1	20
330	Investigation of the DCDC2 intron 2 deletion/compound short tandem repeat polymorphism in a large German dyslexia sample. Psychiatric Genetics, 2008, 18, 310-312.	1.1	46
331	Mapping Genes for Asthma and Psoriasis. Novartis Foundation Symposium, 2008, , 46-56.	1.1	1
332	Expression Analysis of the NLRP Gene Family Suggests a Role in Human Preimplantation Development. PLoS ONE, 2008, 3, e2755.	2.5	109
333	Genome-Wide Analysis of Single Nucleotide Polymorphisms Uncovers Population Structure in Northern Europe. PLoS ONE, 2008, 3, e3519.	2.5	112
334	Interactions between Glutathione S- Transferase P1, Tumor Necrosis Factor, and Traffic-Related Air Pollution for Development of Childhood Allergic Disease. Environmental Health Perspectives, 2008, 116, 1077-1084.	6.0	115
335	Lack of Association between Neuropeptide S Receptor 1 Gene (NPSR1) and Eczema in Five European Populations. Acta Dermato-Venereologica, 2008, 89, 115-121.	1.3	4
336	Myosin IXB gene region and gluten intolerance: linkage to coeliac disease and a putative dermatitis herpetiformis association. Journal of Medical Genetics, 2007, 45, 222-227.	3.2	35
337	The PSORS1 locus gene CCHCR1 affects keratinocyte proliferation in transgenic mice. Human Molecular Genetics, 2007, 17, 1043-1051.	2.9	38
338	Expression of ion transport-associated proteins in human efferent and epididymal ducts. Reproduction, 2007, 133, 775-784.	2.6	60
339	Clinical Associations of the Risk Alleles of HLA-Cw6 and CCHCR1*WWCC in Psoriasis. Acta Dermato-Venereologica, 2007, 87, 127-134.	1.3	21
340	Regulation of the basolateral chloride/base exchangers AE1 and SLC26A7 in the kidney collecting duct in potassium depletion. Nephrology Dialysis Transplantation, 2007, 22, 3462-3470.	0.7	13
341	The human GIMAP5 gene has a common polyadenylation polymorphism increasing risk to systemic lupus erythematosus. Journal of Medical Genetics, 2007, 44, 314-321.	3.2	70
342	A locus on 2p12 containing the co-regulated MRPL19 and C2ORF3 genes is associated to dyslexia. Human Molecular Genetics, 2007, 16, 667-677.	2.9	102

#	Article	IF	Citations
343	Large-Scale Zygosity Testing Using Single Nucleotide Polymorphisms. Twin Research and Human Genetics, 2007, 10, 604-625.	0.6	110
344	An unfavorable combination of factor V Leiden with age, weight, and blood group causes high risk of pregnancy-associated venous thrombosisâ€"a population-based nested case-control study. Thrombosis Research, 2007, 119, 423-432.	1.7	22
345	Exclusion of coding-region mutations in luteinizing hormone and follicle-stimulating hormone receptor genes as the cause of ovarian hyperstimulation syndrome. Fertility and Sterility, 2007, 87, 603-606.	1.0	34
346	Distinct sets of developmentally regulated genes that are expressed by human oocytes and human embryonic stem cells. Fertility and Sterility, 2007, 87, 677-690.	1.0	39
347	A Common Haplotype in the G-Proteinâ€"Coupled Receptor Gene GPR74 Is Associated with Leanness and Increased Lipolysis. American Journal of Human Genetics, 2007, 80, 1115-1124.	6.2	20
348	Physical Exerciseâ€"Induced Hypoglycemia Caused by Failed Silencing of Monocarboxylate Transporter 1 in Pancreatic Î² Cells. American Journal of Human Genetics, 2007, 81, 467-474.	6.2	213
349	Neuropeptide S Receptor 1 Gene Polymorphism Is Associated With Susceptibility to Inflammatory Bowel Disease. Gastroenterology, 2007, 133, 808-817.	1.3	87
350	Mechanisms of inactivation of MLH1 in hereditary nonpolyposis colorectal carcinoma: a novel approach. Oncogene, 2007, 26, 4541-4549.	5.9	56
351	Mutations in the gene encoding the 3â€²-5â€² DNA exonuclease TREX1 are associated with systemic lupus erythematosus. Nature Genetics, 2007, 39, 1065-1067.	21.4	590
352	Evaluation of STOX1 as a preeclampsia candidate gene in a population-wide sample. European Journal of Human Genetics, 2007, 15, 494-497.	2.8	45
353	A recurrent mutation in PALB2 in Finnish cancer families. Nature, 2007, 446, 316-319.	27.8	402
354	Chromosome 7p linkage and GPR154 gene association in Italian families with allergic asthma. Clinical and Experimental Allergy, 2007, 37, 83-89.	2.9	43
355	The CCHCR1 (HCR) gene is relevant for skin steroidogenesis and downregulated in cultured psoriatic keratinocytes. Journal of Molecular Medicine, 2007, 85, 589-601.	3.9	49
356	Expression of Allograft Inflammatory Factor-1 in Inflammatory Skin Disorders. Acta Dermato-Venereologica, 2007, 87, 223-227.	1.3	20
357	Two Chinese families with pulverulent congenital cataracts and deltaG91 CRYBA1 mutations. Molecular Vision, 2007, 13, 1154-60.	1.1	35
358	G proteinâ€coupled receptor for asthma susceptibility associates with respiratory distress syndrome. Annals of Medicine, 2006, 38, 357-366.	3.8	31
359	Evaluation of the role of Finnish ataxia-telangiectasia mutations in hereditary predisposition to breast cancer. Carcinogenesis, 2006, 28, 1040-1045.	2.8	21
360	Strong Genetic Evidence of DCDC2 as a Susceptibility Gene for Dyslexia. American Journal of Human Genetics, 2006, 78, 52-62.	6.2	211

#	Article	IF	Citations
361	Elevated Expression and Genetic Association Links the SOCS3 Gene to Atopic Dermatitis. American Journal of Human Genetics, 2006, 78, 1060-1065.	6.2	47
362	ELMOD2 Is a Candidate Gene for Familial Idiopathic Pulmonary Fibrosis. American Journal of Human Genetics, 2006, 79, 149-154.	6.2	115
363	Absence of a Paternally Inherited FOXP2 Gene in Developmental Verbal Dyspraxia. American Journal of Human Genetics, 2006, 79, 965-972.	6.2	170
364	DYX1C1 functions in neuronal migration in developing neocortex. Neuroscience, 2006, 143, 515-522.	2.3	131
365	Liver X receptor gene polymorphisms and adipose tissue expression levels in obesity. Pharmacogenetics and Genomics, 2006, 16, 881-889.	1.5	53
366	Interaction between variants in the interleukin-4 receptor ? and interleukin-9 receptor genes in childhood wheezing: evidence from a birth cohort study. Clinical and Experimental Allergy, 2006, 36, 1391-1398.	2.9	16
367	The role of inflammatory bowel disease susceptibility loci in multiple sclerosis and systemic lupus erythematosus. Genes and Immunity, 2006, 7, 327-334.	4.1	29
368	Genome scan on Swedish Alzheimer's disease families. Molecular Psychiatry, 2006, 11, 182-186.	7.9	40
369	Heterogeneity-based genome search meta-analysis for preeclampsia. Human Genetics, 2006, 120, 360-370.	3.8	32
370	A quality assessment survey of SNP genotyping laboratories. Human Mutation, 2006, 27, 711-714.	2.5	18
371	Associations of Body Size at Birth with Late-Life Cortisol Concentrations and Glucose Tolerance Are Modified by Haplotypes of the Glucocorticoid Receptor Gene. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 4544-4551.	3.6	44
372	Expression of SLC26A3, CFTR and NHE3 in the human male reproductive tract: role in male subfertility caused by congenital chloride diarrhoea. Molecular Human Reproduction, 2006, 12, 107-111.	2.8	98
373	Neuropeptide S and G protein-coupled receptor 154 modulate macrophage immune responses. Human Molecular Genetics, 2006, 15, 1667-1679.	2.9	64
374	Variation in DNA Repair Genes ERCC2, XRCC1, and XRCC3 and Risk of Follicular Lymphoma. Cancer Epidemiology Biomarkers and Prevention, 2006, 15, 258-265.	2.5	61
375	Downstream target genes of the neuropeptide Sâ€“NPSR1 pathway. Human Molecular Genetics, 2006, 15, 2923-2935.	2.9	32
376	Nordic collaborative study of the BARD1 Cys557Ser allele in 3956 patients with cancer: enrichment in familial BRCA1/BRCA2 mutation-negative breast cancer but not in other malignancies. Journal of Medical Genetics, 2006, 43, 856-862.	3.2	38
377	Overview of the SLC26 family and associated diseases. Novartis Foundation Symposium, 2006, 273, 2-11; discussion 11-8, 261-4.	1.1	10
378	The CIDEA Gene V115F Polymorphism Is Associated With Obesity in Swedish Subjects. Diabetes, 2005, 54, 3032-3034.	0.6	51

#	Article	IF	Citations
379	A Common Î²₂â€Adrenoceptor Gene Haplotype Protects against Obesity in Swedish Women. Obesity, 2005, 13, 1645-1650.	4.0	6
380	MHC2TA is associated with differential MHC molecule expression and susceptibility to rheumatoid arthritis, multiple sclerosis and myocardial infarction. Nature Genetics, 2005, 37, 486-494.	21.4	276
381	Family-based association study of DYX1C1 variants in autism. European Journal of Human Genetics, 2005, 13, 127-130.	2.8	16
382	Haplotype construction of the FRDA gene and evaluation of its role in type II diabetes. European Journal of Human Genetics, 2005, 13, 849-855.	2.8	10
383	Novel and de novo mutations of the IRF6 gene detected in patients with Van der Woude or popliteal pterygium syndrome. European Journal of Human Genetics, 2005, 13, 1261-1267.	2.8	27
384	Fine Mapping of the Psoriasis Susceptibility Gene PSORS1: A Reassessment of Risk Associated with a Putative Risk Haplotype Lacking HLA-Cw6. Journal of Investigative Dermatology, 2005, 124, 921-930.	0.7	18
385	Absence of Association Between Asthma and High Serum Immunoglobulin E Associated GPRA Haplotypes and Adult Atopic Dermatitis. Journal of Investigative Dermatology, 2005, 125, 399-401.	0.7	23
386	BMPR2 mutations have short lifetime expectancy in primary pulmonary hypertension. Human Mutation, 2005, 26, 119-124.	2.5	30
387	Association analysis of the R620W polymorphism of protein tyrosine phosphatase PTPN22 in systemic lupus erythematosus families: Increased t allele frequency in systemic lupus erythematosus patients with autoimmune thyroid disease. Arthritis and Rheumatism, 2005, 52, 2396-2402.	6.7	80
388	Mutational analysis of the human SLC26A8 gene: exclusion as a candidate for male infertility due to primary spermatogenic failure. Molecular Human Reproduction, 2005, 11, 129-132.	2.8	25
389	Subpopulation difference scanning: a strategy for exclusion mapping of susceptibility genes. Journal of Medical Genetics, 2005, 43, 590-597.	3.2	2
390	The Axon Guidance Receptor Gene ROBO1 Is a Candidate Gene for Developmental Dyslexia. PLoS Genetics, 2005, 1, e50.	3.5	276
391	A genome-wide scan of non-syndromic cleft palate only (CPO) in Finnish multiplex families. Journal of Medical Genetics, 2005, 42, 177-184.	3.2	25
392	Characterization of GPRA, a Novel G Proteinâ€“Coupled Receptor Related to Asthma. American Journal of Respiratory Cell and Molecular Biology, 2005, 33, 262-270.	2.9	96
393	Phenylketonuria screening registry as a resource for population genetic studies. Journal of Medical Genetics, 2005, 42, e60-e60.	3.2	43
394	Mapping and identifying genes for asthma and psoriasis. Philosophical Transactions of the Royal Society B: Biological Sciences, 2005, 360, 1551-1561.	4.0	15
395	Functional Comparison of Mouse slc26a6 Anion Exchanger with Human SLC26A6 Polypeptide Variants. Journal of Biological Chemistry, 2005, 280, 8564-8580.	3.4	137
396	SLC26A6 and SLC26A7 Anion Exchangers Have a Distinct Distribution in Human Kidney. Nephron Experimental Nephrology, 2005, 101, e50-e58.	2.2	24

#	Article	IF	Citations
397	Global analysis of uniparental disomy using high density genotyping arrays. Journal of Medical Genetics, 2005, 42, 847-851.	3.2	37
398	Haplotypes of G Proteinâ€“coupled Receptor 154 Are Associated with Childhood Allergy and Asthma. American Journal of Respiratory and Critical Care Medicine, 2005, 171, 1089-1095.	5.6	111
399	Gene mapping with pooled samples on three genotyping platforms. Molecular and Cellular Probes, 2005, 19, 408-416.	2.1	5
400	Polymorphisms in the Tyrosine Kinase 2 and Interferon Regulatory Factor 5 Genes Are Associated with Systemic Lupus Erythematosus. American Journal of Human Genetics, 2005, 76, 528-537.	6.2	526
401	Association analysis of common variants of STAT6, GATA3, and STAT4 to asthma and high serum IgE phenotypes. Journal of Allergy and Clinical Immunology, 2005, 115, 80-87.	2.9	91
402	Spectrum of mutations in CFTR in Finland: 18 years follow-up study and identification of two novel mutations. Journal of Cystic Fibrosis, 2005, 4, 233-237.	0.7	23
403	Mapping genes for asthma and psoriasis. Novartis Foundation Symposium, 2005, 267, 46-52; discussion 52-6.	1.1	1
404	Sex specific protective effects of interleukin-9 receptor haplotypes on childhood wheezing and sensitisation. Journal of Medical Genetics, 2004, 41, e123-e123.	3.2	19
405	The Effects of the ACE Gene Insertion/Deletion Polymorphism on Glucose Tolerance and Insulin Secretion in Elderly People Are Modified by Birth Weight. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 5738-5741.	3.6	46
406	Differential Regulation of Basolateral Clâ^'/HCO3 â^' Exchangers SLC26A7 and AE1 in Kidney Outer Medullary Collecting Duct. Journal of the American Society of Nephrology: JASN, 2004, 15, 2002-2011.	6.1	37
407	Linkage mapping of systemic lupus erythematosus (SLE) in Finnish families multiply affected by SLE. Journal of Medical Genetics, 2004, 41, 2e-5.	3.2	18
408	Characterization of a Common Susceptibility Locus for Asthma-Related Traits. Science, 2004, 304, 300-304.	12.6	442
409	Transgenic mouse models support HCR as an effector gene in the PSORS1 locus. Human Molecular Genetics, 2004, 13, 1551-1561.	2.9	35
410	Anti-telomere antibodies in systemic lupus erythematosus (SLE): a comparison with five antinuclear antibody assays in 430 patients with SLE and other rheumatic diseases. Annals of the Rheumatic Diseases, 2004, 63, 1250-1254.	0.9	11
411	Two functional variants of the superoxide dismutase genes in Finnish families with asthma. Thorax, 2004, 59, 116-119.	5.6	45
412	The association of antibodies to cardiolipin,Î²2â€glycoprotein I, prothrombin, and oxidized lowâ€density lipoprotein with thrombosis in 292 patients with familial and sporadic systemic lupus erythematosus. Scandinavian Journal of Rheumatology, 2004, 33, 246-252.	1.1	19
413	Influence of male sex and parental allergic disease on childhood wheezing: role of interactions. Clinical and Experimental Allergy, 2004, 34, 839-844.	2.9	62
414	Association study of 15 novel single-nucleotide polymorphisms of the T-bet locus among Finnish asthma families. Clinical and Experimental Allergy, 2004, 34, 1049-1055.	2.9	19

#	Article	IF	Citations
415	Haplotype associations define target regions for susceptibility loci in systemic lupus erythematosus. European Journal of Human Genetics, 2004, 12, 489-494.	2.8	9
416	Fine mapping of the 2p11 dyslexia locus and exclusion of TACR1 as a candidate gene. Human Genetics, 2004, 114, 510-516.	3.8	27
417	?2-Heremans?Schmid glycoprotein gene polymorphisms are associated with adipocyte insulin action. Diabetologia, 2004, 47, 1974-1979.	6.3	62
418	Positionally cloned susceptibility genes in allergy and asthma. Current Opinion in Immunology, 2004, 16, 689-694.	5.5	35
419	SLC26A7: a basolateral Cl⁻/HCO₃⁻exchanger specific to intercalated cells of the outer medullary collecting duct. American Journal of Physiology - Renal Physiology, 2004, 286, F161-F169.	2.7	79
420	Genetic Analysis of PSORS1 Distinguishes Guttate Psoriasis and Palmoplantar Pustulosis. Journal of Investigative Dermatology, 2003, 120, 627-632.	0.7	190
421	Psoriasis Susceptibility Locus on 18p Revealed by Genome Scan in Finnish Families Not Associated with PSORS1. Journal of Investigative Dermatology, 2003, 121, 735-740.	0.7	44
422	HCR, a Candidate Gene for Psoriasis, Is Expressed Differently in Psoriasis and Other Hyperproliferative Skin Disorders and Is Downregulated by Interferon-Î³ in Keratinocytes. Journal of Investigative Dermatology, 2003, 121, 1360-1364.	0.7	26
423	Collagen XI sequence variations in nonsyndromic cleft palate, Robin sequence and micrognathia. European Journal of Human Genetics, 2003, 11, 265-270.	2.8	57
424	Identification of a basolateral Cl-/HCO3- exchanger specific to gastric parietal cells. Gastroenterology, 2003, 124, A3.	1.3	0
425	Susceptibility Loci for Preeclampsia on Chromosomes 2p25 and 9p13 in Finnish Families. American Journal of Human Genetics, 2003, 72, 168-177.	6.2	151
426	Human Chromosome 7: DNA Sequence and Biology. Science, 2003, 300, 767-772.	12.6	185
427	Physical Exercise-Induced Hyperinsulinemic Hypoglycemia Is an Autosomal-Dominant Trait Characterized by Abnormal Pyruvate-Induced Insulin Release. Diabetes, 2003, 52, 199-204.	0.6	135
428	A genome scan for developmental dyslexia confirms linkage to chromosome 2p11 and suggests a new locus on 7q32. Journal of Medical Genetics, 2003, 40, 340-345.	3.2	101
429	A candidate gene for developmental dyslexia encodes a nuclear tetratricopeptide repeat domain protein dynamically regulated in brain. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 11553-11558.	7.1	319
430	Linkage and linkage disequilibrium searched for between non-syndromic cleft palate and four candidate loci. Journal of Medical Genetics, 2003, 40, 464-468.	3.2	4
431	Genomics and Pediatric Research. Pediatric Research, 2003, 53, 4-9.	2.3	19
432	Isoforms of SLC26A6 mediate anion transport and have functional PDZ interaction domains. American Journal of Physiology - Cell Physiology, 2003, 284, C769-C779.	4.6	125

#	Article	IF	Citations
433	Identification of a basolateral Cl^{â^'}/HCO 3 â^' exchanger specific to gastric parietal cells. American Journal of Physiology - Renal Physiology, 2003, 284, G1093-G1103.	3.4	81
434	Genomics and Pediatric Research. Pediatric Research, 2003, 53, 4-9.	2.3	0
435	Coding haplotype analysis supports HCR as the putative susceptibility gene for psoriasis at the MHC PSORS1 locus. Human Molecular Genetics, 2002, 11, 589-597.	2.9	131
436	Functional Characterization of Three Novel Tissue-specific Anion Exchangers SLC26A7, -A8, and -A9. Journal of Biological Chemistry, 2002, 277, 14246-14254.	3.4	200
437	Genetic Screening for Maternal Uniparental Disomy of Chromosome 7 in Prenatal and Postnatal Growth Retardation of Unknown Cause. Pediatrics, 2002, 109, 441-448.	2.1	39
438	Long-range control of expression in yeast. Bioinformatics, 2002, 18, 482-483.	4.1	7
439	Down-regulated in adenoma mediates apical Clâ^'/HCO3â^' exchange in rabbit, rat, and human duodenum. Gastroenterology, 2002, 122, 709-724.	1.3	162
440	Healing a natural knockout of epithelial organogenesis. Trends in Molecular Medicine, 2002, 8, 197-200.	6.7	9
441	Matrix metalloproteinase-21, the human orthologue for XMMP, is expressed during fetal development and in cancer. Gene, 2002, 301, 31-41.	2.2	43
442	Genome segmentation using piecewise constant intensity models and reversible jump MCMC. Bioinformatics, 2002, 18, S211-S218.	4.1	10
443	Upregulation of CFTR expression but not SLC26A3 and SLC9A3 in ulcerative colitis. American Journal of Physiology - Renal Physiology, 2002, 283, G567-G575.	3.4	53
444	Familial dyslexia: neurocognitive and genetic correlation in a large Finnish family. Developmental Medicine and Child Neurology, 2002, 44, 580-6.	2.1	13
445	Recurrent DNA sequence copy losses on chromosomal arm 6q in capillary hemangioblastoma. Cancer Genetics and Cytogenetics, 2002, 133, 174-178.	1.0	19
446	SLC26A3 mutations in congenital chloride diarrhea. Human Mutation, 2002, 20, 425-438.	2.5	131
447	A new locus for coeliac disease mapped to chromosome 15 in a population isolate. Human Genetics, 2002, 111, 40-45.	3.8	27
448	Investigatory and analytical approaches to differential gene expression profiling in mantle cell lymphoma. British Journal of Haematology, 2002, 119, 905-915.	2.5	49
449	Physical map of an asthma susceptibility locus in 7p15-p14 and an association study of TCRG. European Journal of Human Genetics, 2002, 10, 658-665.	2.8	5
450	Familial dyslexia: neurocognitive and genetic correlation in a large Finnish family. Developmental Medicine and Child Neurology, 2002, 44, 580-586.	2.1	21

#	Article	IF	Citations
451	Association analysis for quantitative traits by data mining: QHPM. Annals of Human Genetics, 2002, 66, 419-29.	0.8	6
452	Maternal and Paternal Chromosomes 7 Show Differential Methylation of Many Genes in Lymphoblast DNA. Genomics, 2001, 73, 1-9.	2.9	24
453	A Second-Generation Association Study of the 5q31 Cytokine Gene Cluster and the Interleukin-4 Receptor in Asthma. Genomics, 2001, 77, 35-42.	2.9	46
454	A Narrow Segment of Maternal Uniparental Disomy of Chromosome 7q31-qter in Silver-Russell Syndrome Delimits a Candidate Gene Region. American Journal of Human Genetics, 2001, 68, 247-253.	6.2	127
455	PPP2R1BGene in Chronic Lymphocytic Leukemias and Mantle Cell Lymphomas. Leukemia and Lymphoma, 2001, 41, 177-183.	1.3	16
456	Concomitant DNA copy number amplification at 17q and 22q in dermatofibrosarcoma protuberans. Cytogenetic and Genome Research, 2001, 92, 192-195.	1.1	49
457	Mining Associations Between Genetic Markers, Phenotypes, and Covariates. Genetic Epidemiology, 2001, 21, S588-S593.	1.3	2
458	Identification of seven novel mutations including the first two genomic rearrangements in SLC26A3 mutated in congenital chloride diarrhea. Human Mutation, 2001, 18, 233-242.	2.5	29
459	The mutation spectrum of the EDA gene in X-linked anhidrotic ectodermal dysplasia. Human Mutation, 2001, 17, 349-349.	2.5	60
460	Involvement ofBRCA1 andBRCA2 in breast cancer in a western Finnish sub-population. Genetic Epidemiology, 2001, 20, 239-246.	1.3	7
461	A susceptibility locus for asthma-related traits on chromosome 7 revealed by genome-wide scan in a founder population. Nature Genetics, 2001, 28, 87-91.	21.4	168
462	Mapping of the second locus for the Van der Woude syndrome to chromosome 1p34. European Journal of Human Genetics, 2001, 9, 747-752.	2.8	63
463	Haplotype analysis in Icelandic and Finnish BRCA2 999del5 breast cancer families. European Journal of Human Genetics, 2001, 9, 773-779.	2.8	23
464	Distinct outcomes of chloride diarrhoea in two siblings with identical genetic background of the disease: implications for early diagnosis and treatment. Gut, 2001, 48, 724-727.	12.1	46
465	Ultrastructural features resembling those of harlequin ichthyosis in patients with severe congenital ichthyosiform erythroderma.. British Journal of Dermatology, 2001, 145, 480-483.	1.5	16
466	SLC26A2 (Diastrophic Dysplasia Sulfate Transporter) is Expressed in Developing and Mature Cartilage But Also in Other Tissues and Cell Types. Journal of Histochemistry and Cytochemistry, 2001, 49, 973-982.	2.5	80
467	Autosomal dominant midfrequency hearing impairment. Scandinavian Audiology, 2001, 30, 85-87.	0.5	2
468	Ectodysplasin is released by proteolytic shedding and binds to the EDAR protein. Human Molecular Genetics, 2001, 10, 953-962.	2.9	97

#	Article	IF	Citations
469	HUMANPOPULATIONGENETICS: Lessons from Finland. Annual Review of Genomics and Human Genetics, 2001, 2, 103-128.	6.2	77
470	A dominant gene for developmental dyslexia on chromosome 3. Journal of Medical Genetics, 2001, 38, 658-664.	3.2	135
471	Title is missing!. Nature Genetics, 2001, 28, 87-91.	21.4	75
472	Cystic Fibrosis Gene Mutations Î”F508 and 394delTT in Patients with Chronic Sinusitis in Finland. Acta Oto-Laryngologica, 2001, 121, 945-947.	0.9	16
473	Genetic heterogeneity and exclusion of a modifying locus at 17p11.2-p11.1 in Finnish families with van der Woude syndrome. Journal of Medical Genetics, 2001, 38, 198-202.	3.2	7
474	Do patients with maternal uniparental disomy for chromosome 7 have a distinct mild Silver-Russell phenotype?. Journal of Medical Genetics, 2001, 38, 273-278.	3.2	62
475	Clinical Findings in Mosaic Carriers of Hypohidrotic Ectodermal Dysplasia. Archives of Dermatology, 2000, 136, 217-24.	1.4	59
476	Chapter 8 Anion absorption in the intestine: Anion transporters, short-chain fatty acids, and role of the DRA gene product. Current Topics in Membranes, 2000, 50, 301-328.	0.9	7
477	Gene Mapping in Isolated Populations: New Roles for Old Friends?. Human Heredity, 2000, 50, 57-65.	0.8	65
478	Stromelysin-2 is Upregulated During Normal Wound Repair and is Induced by Cytokines. Journal of Investigative Dermatology, 2000, 115, 778-787.	0.7	84
479	Juxtaposed regions of extensive and minimal linkage disequilibrium in human Xq25 and Xq28. Nature Genetics, 2000, 25, 324-328.	21.4	272
480	Multiple founder effects and geographical clustering of BRCA1 and BRCA2 families in Finland. European Journal of Human Genetics, 2000, 8, 757-763.	2.8	75
481	The IL9R region contribution in asthma is supported by genetic association in an isolated population. European Journal of Human Genetics, 2000, 8, 788-792.	2.8	20
482	The congenital chloride diarrhea gene is expressed in seminal vesicle, sweat gland, inflammatory colon epithelium, and in some dysplastic colon cells. Histochemistry and Cell Biology, 2000, 113, 279-286.	1.7	59
483	Two translocations of chromosome 15q associated with dyslexia. Journal of Medical Genetics, 2000, 37, 771-775.	3.2	81
484	Hereditary Hearing Loss - the Role of Environmental Factors. Acta Oto-Laryngologica, 2000, 120, 70-72.	0.9	3
485	Association Study of the Chromosomal Region Containing the FCER2 Gene Suggests It Has a Regulatory Role in Atopic Disorders. American Journal of Respiratory and Critical Care Medicine, 2000, 161, 700-706.	5.6	40
486	A candidate gene for psoriasis near HLA-C, HCR (Pg8), is highly polymorphic with a disease-associated susceptibility allele. Human Molecular Genetics, 2000, 9, 1533-1542.	2.9	120

#	Article	IF	Citations
487	Mapping of Five New Putative Anion Transporter Genes in Human and Characterization of SLC26A6, A Candidate Gene for Pancreatic Anion Exchanger. Genomics, 2000, 70, 102-112.	2.9	187
488	Assignment of a Novel Locus for Autosomal Recessive Congenital Ichthyosis to Chromosome 19p13.1-p13.2. American Journal of Human Genetics, 2000, 66, 1132-1137.	6.2	49
489	Data Mining Applied to Linkage Disequilibrium Mapping. American Journal of Human Genetics, 2000, 67, 133-145.	6.2	114
490	Inherited disorders of ion transport in the intestine. Current Opinion in Genetics and Development, 2000, 10, 306-309.	3.3	25
491	Dominantly inherited hyperinsulinism caused by a mutation in the sulfonylurea receptor type 1. Journal of Clinical Investigation, 2000, 106, 897-906.	8.2	237
492	III. Congenital chloride diarrhea. American Journal of Physiology - Renal Physiology, 1999, 276, G7-G13.	3.4	48
493	Downregulated in adenoma gene encodes a chloride transporter defective in congenital chloride diarrhea. American Journal of Physiology - Renal Physiology, 1999, 276, G185-G192.	3.4	103
494	A point mutation inactivating the sulfonylurea receptor causes the severe form of persistent hyperinsulinemic hypoglycemia of infancy in Finland. Diabetes, 1999, 48, 408-415.	0.6	144
495	Functional Characterization of the Promoter of the X-linked Ectodermal Dysplasia Gene. Journal of Biological Chemistry, 1999, 274, 26477-26484.	3.4	8
496	Ectodysplasin is a collagenous trimeric type II membrane protein with a tumor necrosis factor-like domain and co-localizes with cytoskeletal structures at lateral and apical surfaces of cells. Human Molecular Genetics, 1999, 8, 2079-2086.	2.9	112
497	Kidney kinetics and chloride ion pumps. Nature Genetics, 1999, 21, 67-68.	21.4	11
498	Clinical and morphological correlations for transglutaminase 1 gene mutations in autosomal recessive congenital ichthyosis. European Journal of Human Genetics, 1999, 7, 625-632.	2.8	46
499	Deletions at 14q in malignant mesothelioma detected by microsatellite marker analysis. British Journal of Cancer, 1999, 81, 1111-1115.	6.4	32
500	Long-Term Prognosis of Haemangioblastoma of the CNS: Impact of von Hippel-Lindau Disease. Acta Neurochirurgica, 1999, 141, 1147-1156.	1.7	75
501	Myotilin, a novel sarcomeric protein with two Ig-like domains, is encoded by a candidate gene for limb-girdle muscular dystrophy. Human Molecular Genetics, 1999, 8, 1329-1336.	2.9	181
502	Ectodysplasin, a protein required for epithelial morphogenesis, is a novel TNF homologue and promotes cell-matrix adhesion. Mechanisms of Development, 1999, 88, 133-146.	1.7	115
503	Arrhythmic disorder mapped to chromosome 1q42â€“q43 causes malignant polymorphic ventricular tachycardia in structurally normal hearts. Journal of the American College of Cardiology, 1999, 34, 2035-2042.	2.8	321
504	Cusp Patterning Defect in Tabby Mouse Teeth and Its Partial Rescue by FGF. Developmental Biology, 1999, 216, 521-534.	2.0	174

#	Article	IF	Citations
505	Erythropoietin Receptor Mutations Associated With Familial Erythrocytosis Cause Hypersensitivity to Erythropoietin in the Heterozygous State. Blood, 1999, 94, 2530-2532.	1.4	54
506	Expression of the anhidrotic ectodermal dysplasia gene is reduced in skin cancer coinciding with reduced E-cadherin. Experimental Dermatology, 1998, 7, 168-174.	2.9	8
507	Clustering of private mutations in the congenital chloride diarrhea/down-regulated in adenoma gene. Human Mutation, 1998, 11, 321-327.	2.5	27
508	Genomic structure of the human ezrin gene. Human Genetics, 1998, 103, 662-665.	3.8	12
509	Phenotyping asthma patients for a gene mapping study in Finland. Clinical and Experimental Allergy, 1998, 28, 40-42.	2.9	2
510	22-Mb integrated physical and genetic map based on YAC/STS content spanning the interval DXS1125â€“DXS95 in human Xq12â€“q21.31. Gene, 1998, 208, 147-156.	2.2	8
511	Genomic structure of the human congenital chloride diarrhea (CLD) gene. Gene, 1998, 214, 87-93.	2.2	33
512	Verification of self-reported asthma and allergy in subjects and their family members volunteering for gene mapping studies. Respiratory Medicine, 1998, 92, 1281-1288.	2.9	44
513	Genetic Background of Congenital Chloride Diarrhea in High-Incidence Populations: Finland, Poland, and Saudi Arabia and Kuwait. American Journal of Human Genetics, 1998, 63, 760-768.	6.2	78
514	Distinct expression profiles of stromelysin-2 (MMP-10), collagenase-3 (MMP-13), macrophage metalloelastase (HME, MMP-12) and TIMP-3 in intestinal ulcerations. Gastroenterology, 1998, 114, A1064.	1.3	1
515	Human TIMP-3 Is Expressed During Fetal Development, Hair Growth Cycle, and Cancer Progression. Journal of Histochemistry and Cytochemistry, 1998, 46, 437-447.	2.5	48
516	The Anhidrotic Ectodermal Dysplasia Gene (EDA) Undergoes Alternative Splicing and Encodes Ectodysplasin-A with Deletion Mutations in Collagenous Repeats. Human Molecular Genetics, 1998, 7, 1661-1669.	2.9	193
517	The Gene Defective in Anhidrotic Ectodermal Dysplasia Is Expressed in the Developing Epithelium, Neuroectoderm, Thymus, and Bone. Journal of Histochemistry and Cytochemistry, 1998, 46, 281-289.	2.5	52
518	Clustering of private mutations in the congenital chloride diarrhea/downâ€regulated in adenoma gene. Human Mutation, 1998, 11, 321-327.	2.5	1
519	Anhidrotic ectodermal dysplasia (EDA) protein expressed in MCF-7 cells associates with cell membrane and induces rounding. Human Molecular Genetics, 1997, 6, 1581-1587.	2.9	19
520	Genetic Control of Serum IgE Levels and Asthma: Linkage and Linkage Disequilibrium Studies in an Isolated Population. Human Molecular Genetics, 1997, 6, 2069-2076.	2.9	83
521	X chromosome map at 75-kb STS resolution, revealing extremes of recombination and GC content.. Genome Research, 1997, 7, 210-222.	5.5	109
522	The Tabby phenotype is caused by mutation in a mouse homologue of theEDAgene that reveals novel mouse and human exons and encodes a protein (ectodysplasin-A) withâ€‰collagenousâ€‰domains. Proceedings of the National Academy of Sciences of the United States of America, 1997, 94, 13069-13074.	7.1	282

#	Article	IF	Citations
523	Monoallelic Expression of HumanPEG1/MESTIs Paralleled by Parent-Specific Methylation in Fetuses. Genomics, 1997, 42, 236-244.	2.9	91
524	Transglutaminase 1 Mutations in Autosomal Recessive Congenital Ichthyosis: Private and Recurrent Mutations in an Isolated Population. American Journal of Human Genetics, 1997, 61, 529-538.	6.2	111
525	Microsatellite diversity and the demographic history ofâ€‰modernâ€‰humans. Proceedings of the National Academy of Sciences of the United States of America, 1997, 94, 3100-3103.	7.1	268
526	Evidence for two molecular steps in the pathogenesis of myeloid disorders associated with deletion of chromosome 7 long arm. Leukemia, 1997, 11, 2097-2104.	7.2	18
527	Xâ€“linked anhidrotic (hypohidrotic) ectodermal dysplasia is caused by mutation in a novel transmembrane protein. Nature Genetics, 1996, 13, 409-416.	21.4	691
528	Mutations of the Downâ€“regulated in adenoma (DRA) gene cause congenital chloride diarrhoea. Nature Genetics, 1996, 14, 316-319.	21.4	394
529	Split hand/split foot malformation, deafness, and mental retardation with a complex cytogenetic rearrangement involving 7q21.3.. Journal of Medical Genetics, 1996, 33, 507-510.	3.2	38
530	Cloning and characterization of DXS6673E, a candidate gene for X-linked mental retardation in Xq13.1. Human Molecular Genetics, 1996, 5, 887-897.	2.9	69
531	Positional candidate genes for congenital chloride diarrhea suggested by high-resolution physical mapping in chromosome region 7q31.. Genome Research, 1996, 6, 202-210.	5.5	36
532	Patterns of matrix metalloproteinase and TIMP-1 expression in chronic and normally healing human cutaneous wounds. British Journal of Dermatology, 1996, 135, 52-59.	1.5	121
533	Report of the Second International Workshop on Human Chromosome 7 Mapping 1994. Cytogenetic and Genome Research, 1995, 71, 1-31.	1.1	27
534	A 6-Mb YAC contig in Xp22.1â€“p22.2 spanning the DXS69E, XE59, GLRA2, PIGA, GRPR, CALB3, and PHKA2 genes. Genomics, 1995, 25, 691-700.	2.9	29
535	Physical mapping of the split hand/split foot locus on chromosome 7 and implication in syndromic ectrodactyly. Human Molecular Genetics, 1994, 3, 1345-1354.	2.9	125
536	Characterization of four human YAC libraries for clone size, chimerism and X chromosome sequence representation. Nucleic Acids Research, 1994, 22, 3406-3411.	14.5	29
537	Sequence-tagged sites (STSs) from YAC insert-ends and X-specific flow-sorted chromosomes. Mammalian Genome, 1994, 5, 511-514.	2.2	2
538	Cystic fibrosis in a low-incidence population: two major mutations in Finland. Human Genetics, 1994, 93, 162-166.	3.8	35
539	A rare reciprocal translocation (12;21) segregating for nine generations. Human Genetics, 1993, 92, 509-512.	3.8	4
540	Anhidrotic Ectodermal Dysplasia Gene Region Cloned in Yeast Artificial Chromosomes. Genomics, 1993, 16, 305-310.	2.9	21

#	Article	IF	Citations
541	Ordered Shotgun Sequencing, a Strategy for Integrated Mapping and Sequencing of YAC Clones. Genomics, 1993, 17, 651-656.	2.9	50
542	Haplotype analysis to determine the position of a mutation among closely linked DNA markers. Human Molecular Genetics, 1993, 2, 1007-1014.	2.9	19
543	The gene for congenital chloride diarrhea maps close to but is distinct from the gene for cystic fibrosis transmembrane conductance regulator.. Proceedings of the National Academy of Sciences of the United States of America, 1993, 90, 10686-10689.	7.1	40
544	Conserved sequence-tagged sites: a phylogenetic approach to genome mapping.. Proceedings of the National Academy of Sciences of the United States of America, 1992, 89, 3681-3685.	7.1	24
545	Mapping human chromosomes by walking with sequence-tagged sites from end fragments of yeast artificial chromosome inserts. Genomics, 1992, 14, 241-248.	2.9	110
546	Gelsolinâ€“derived familial amyloidosis caused by asparagine or tyrosine substitution for aspartic acid at residue 187. Nature Genetics, 1992, 2, 157-160.	21.4	163
547	Familial amyloidosis, Finnish type: G654 â†' A mutation of the gelsolin gene in Finnish families and an unrelated American family. Genomics, 1992, 13, 898-901.	2.9	59
548	Homozygosity for the Asn187 gelsolin mutation in Finnish-type familial amyloidosis is associated with severe renal disease. Genomics, 1992, 13, 902-903.	2.9	60
549	Screening for Defined Cystic Fibrosis Mutations by Solid-Phase Minisequencing. Clinical Chemistry, 1992, 38, 39-43.	3.2	39
550	Abnormalities of chromosomes 7 and 22 in human malignant pleural mesothelioma: Correlation between southern blot and cytogenetic analyses. Genes Chromosomes and Cancer, 1992, 4, 176-182.	2.8	16
551	Yeast artificial chromosome-based genome mapping: Some lessons from Xq24â€“q28. Genomics, 1991, 11, 783-793.	2.9	71
552	DNA discontinuities in the domain of amplified humanMYC oncogenes. Genes Chromosomes and Cancer, 1991, 3, 136-141.	2.8	6
553	Direct selection: a method for the isolation of cDNAs encoded by large genomic regions.. Proceedings of the National Academy of Sciences of the United States of America, 1991, 88, 9628-9632.	7.1	302
554	Refinement of human chromosome 7 map around the proalpha2(I)collagen gene by long-range restriction mapping. Nucleic Acids Research, 1991, 19, 2755-2759.	14.5	4
555	Cystic fibrosis mutation Î”F508 in Finland: other mutations predominate. Human Genetics, 1990, 85, 413-415.	3.8	12
556	Finnish hereditary amyloidosis is caused by a single nucleotide substitution in the gelsolin gene. FEBS Letters, 1990, 276, 75-77.	2.8	118
557	Craniofrontonasal dysostosis: variable expression in a threeâ€generation family. Clinical Genetics, 1990, 38, 441-446.	2.0	22
558	Chromosome 7 long arm deletion breakpoints in preleukemia: mapping by pulsed field gel electrophresis. Nucleic Acids Research, 1989, 17, 1511-1520.	14.5	25

#	Article	IF	Citations
559	Cystic fibrosis in Finland: a molecular and genealogical study. Human Genetics, 1989, 83, 20-25.	3.8	25
560	Detection of a rare allele with the pMP6d-9/MspI RFLP near the cystic fibrosis locus. Human Genetics, 1989, 83, 305-306.	3.8	2
561	132 Malignant pleural mesothelioma: Chromosome and DNA analysis. Cancer Genetics and Cytogenetics, 1989, 38, 204.	1.0	0
562	Determination of deletion sizes in the MHC-linked complement C4 and steroid 21-hydroxylase genes by pulsed-field gel electrophoresis. Genomics, 1989, 5, 345-349.	2.9	24
563	Chromosome 7 long-arm deletions in myeloid disorders: terminal DNA sequences are commonly conserved and breakpoints vary. Cytogenetic and Genome Research, 1989, 50, 226-229.	1.1	39
564	Monosomy 7 in Granulocytes and Monocytes in Myelodysplastic Syndrome. New England Journal of Medicine, 1987, 316, 499-503.	27.0	100
565	Prenatal diagnosis of X-linked chronic granulomatous disease using restriction fragment length polymorphism analysis. Genomics, 1987, 1, 87-92.	2.9	12
566	Characterization of chromosome 7 long arm deletions by DNA probes in myelodysplastic syndrome. European Journal of Cancer & Clinical Oncology, 1987, 23, 1767.	0.7	0
567	Synthesis of soluble conducting polymers: polyacetylenes and polyromatics. Journal of the Chemical Society Chemical Communications, 1984, , 255.	2.0	14
568	Gene mapping by haplotype pattern mining. , 0, , .		4