## Jiannis Ragoussis

# List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/6470315/jiannis-ragoussis-publications-by-year.pdf

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

 195
 15,513
 63
 123

 papers
 citations
 h-index
 g-index

 219
 18,179
 9.8
 5.96

 ext. papers
 ext. citations
 avg, IF
 L-index

#	Paper	IF	Citations
195	Single-cell RNA sequencing reveals time- and sex-specific responses of mouse spinal cord microglia to peripheral nerve injury and links ApoE to chronic pain <i>Nature Communications</i> , <b>2022</b> , 13, 843	17.4	5
194	Cohort profile: genomic data for 26 622 individuals from the Canadian Longitudinal Study on Aging (CLSA) <i>BMJ Open</i> , <b>2022</b> , 12, e059021	3	1
193	A small number of early introductions seeded widespread transmission of SARS-CoV-2 in QuBec, Canada. <i>Genome Medicine</i> , <b>2021</b> , 13, 169	14.4	6
192	Lessons learned from understanding chemotherapy resistance in epithelial tubo-ovarian carcinoma from BRCA1and BRCA2mutation carriers. <i>Seminars in Cancer Biology</i> , <b>2021</b> , 77, 110-126	12.7	9
191	Candidate Markers of Olaparib Response from Genomic Data Analyses of Human Cancer Cell Lines. <i>Cancers</i> , <b>2021</b> , 13,	6.6	1
190	Molecular characterization of DICER1-mutated pituitary blastoma. <i>Acta Neuropathologica</i> , <b>2021</b> , 141, 929-944	14.3	6
189	The transcriptional landscape of Shh medulloblastoma. <i>Nature Communications</i> , <b>2021</b> , 12, 1749	17.4	7
188	Nanopore long-read RNA-seq and absolute quantification delineate transcription dynamics in early embryo development of an insect pest. <i>Scientific Reports</i> , <b>2021</b> , 11, 7878	4.9	3
187	Extraction of nuclei from archived postmortem tissues for single-nucleus sequencing applications. <i>Nature Protocols</i> , <b>2021</b> , 16, 2788-2801	18.8	3
186	Locally Adaptive Inversions Modulate Genetic Variation at Different Geographic Scales in a Seaweed Fly. <i>Molecular Biology and Evolution</i> , <b>2021</b> , 38, 3953-3971	8.3	5
185	A coordinated progression of progenitor cell states initiates urinary tract development. <i>Nature Communications</i> , <b>2021</b> , 12, 2627	17.4	5
184	Modeling High-Grade Serous Ovarian Carcinoma Using a Combination of Fallopian Tube Electroporation and CRISPR-Cas9-Mediated Genome Editing. <i>Cancer Research</i> , <b>2021</b> , 81, 5147-5160	10.1	3
183	Rare loss-of-function variants in type I IFN immunity genes are not associated with severe COVID-19. <i>Journal of Clinical Investigation</i> , <b>2021</b> , 131,	15.9	25
182	The genetic landscape of choroid plexus tumors in children and adults. <i>Neuro-Oncology</i> , <b>2021</b> , 23, 650-6	5 <b>6</b> 0	3
181	Whole-exome sequencing of non-BRCA1/BRCA2 mutation carrier cases at high-risk for hereditary breast/ovarian cancer. <i>Human Mutation</i> , <b>2021</b> , 42, 290-299	4.7	9
180	White pupae phenotype of tephritids is caused by parallel mutations of a MFS transporter. <i>Nature Communications</i> , <b>2021</b> , 12, 491	17.4	7
179	Haplotype-resolved de novo assembly of the Vero cell line genome. <i>Npj Vaccines</i> , <b>2021</b> , 6, 106	9.5	5

### (2020-2021)

178	A chromosome-anchored genome assembly for Lake Trout (Salvelinus namaycush). <i>Molecular Ecology Resources</i> , <b>2021</b> ,	8.4	2
177	Oviduct epithelial cells constitute two developmentally distinct lineages that are spatially separated along the distal-proximal axis. <i>Cell Reports</i> , <b>2021</b> , 36, 109677	10.6	5
176	Invasive growth associated with cold-inducible RNA-binding protein expression drives recurrence of surgically resected brain metastases. <i>Neuro-Oncology</i> , <b>2021</b> , 23, 1470-1480	1	10
175	Inferring Copy Number from Triple-Negative Breast Cancer Patient Derived Xenograft scRNAseq Data Using scCNA. <i>Methods in Molecular Biology</i> , <b>2021</b> , 2381, 285-303	1.4	
174	Effects of the Sex Chromosome Complement, XX, XO, or XY, on the Transcriptome and Development of Mouse Oocytes During Follicular Growth <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 792604	4.5	1
173	A functionally impaired missense variant identified in French Canadian families implicates FANCI as a candidate ovarian cancer-predisposing gene. <i>Genome Medicine</i> , <b>2021</b> , 13, 186	14.4	2
172	Single-cell analysis of childhood leukemia reveals a link between developmental states and ribosomal protein expression as a source of intra-individual heterogeneity. <i>Scientific Reports</i> , <b>2020</b> , 10, 8079	4.9	9
171	Model-based analysis of sample index hopping reveals its widespread artifacts in multiplexed single-cell RNA-sequencing. <i>Nature Communications</i> , <b>2020</b> , 11, 2704	17.4	9
170	Single-cell RNA-seq reveals that glioblastoma recapitulates a normal neurodevelopmental hierarchy. <i>Nature Communications</i> , <b>2020</b> , 11, 3406	17.4	88
169	Methodologies for Transcript Profiling Using Long-Read Technologies. <i>Frontiers in Genetics</i> , <b>2020</b> , 11, 606	4.5	24
168	Developmental trajectory of oligodendrocyte progenitor cells in the human brain revealed by single cell RNA sequencing. <i>Glia</i> , <b>2020</b> , 68, 1291-1303	9	22
167	Single-nucleus transcriptomics of the prefrontal cortex in major depressive disorder implicates oligodendrocyte precursor cells and excitatory neurons. <i>Nature Neuroscience</i> , <b>2020</b> , 23, 771-781	25.5	91
166	Single-Cell Transcriptomic Profiling of De Novo and Relapsed Acute Myeloid Leukemia Identifies a Leukemic Stemness Program Shared across Diverse Phenotypes. <i>Blood</i> , <b>2020</b> , 136, 1-1	2.2	
165	DGCR8 microprocessor defect characterizes familial multinodular goiter with schwannomatosis. Journal of Clinical Investigation, <b>2020</b> , 130, 1479-1490	15.9	15
164	Integration of Transcriptome and Metabolome Provides Unique Insights to Pathways Associated With Obese Breast Cancer Patients. <i>Frontiers in Oncology</i> , <b>2020</b> , 10, 804	5.3	13
163	MAFG-driven astrocytes promote CNS inflammation. <i>Nature</i> , <b>2020</b> , 578, 593-599	50.4	125
162	Failure to replicate the association of rare loss-of-function variants in type I IFN immunity genes with severe COVID-19 <b>2020</b> ,		5
161	Metagenomic analysis of planktonic riverine microbial consortia using nanopore sequencing reveals insight into river microbe taxonomy and function. <i>GigaScience</i> , <b>2020</b> , 9,	7.6	11

160	The mevalonate precursor enzyme HMGCS1 is a novel marker and key mediator of cancer stem cell enrichment in luminal and basal models of breast cancer. <i>PLoS ONE</i> , <b>2020</b> , 15, e0236187	3.7	6
159	Regulation of cellular sterol homeostasis by the oxygen responsive noncoding RNA lincNORS. <i>Nature Communications</i> , <b>2020</b> , 11, 4755	17.4	7
158	A Distributed Whole Genome Sequencing Benchmark Study. Frontiers in Genetics, 2020, 11, 612515	4.5	2
157	De novo assembly of the olive fruit fly (Bactrocera oleae) genome with linked-reads and long-read technologies minimizes gaps and provides exceptional Y chromosome assembly. <i>BMC Genomics</i> , <b>2020</b> , 21, 259	4.5	10
156	() orchestrates male sex determination in major agricultural fruit fly pests. <i>Science</i> , <b>2019</b> , 365, 1457-146	6033.3	47
155	Gene Ontology and Expression Studies of Strigolactone Analogues on a Hepatocellular Carcinoma Cell Line. <i>Analytical Cellular Pathology</i> , <b>2019</b> , 2019, 1598182	3.4	2
154	Next-generation HLA typing of 382 International Histocompatibility Working Group reference B-lymphoblastoid cell lines: Report from the 17th International HLA and Immunogenetics Workshop. <i>Human Immunology</i> , <b>2019</b> , 80, 449-460	2.3	13
153	Quality control project of NGS HLA genotyping for the 17th International HLA and Immunogenetics Workshop. <i>Human Immunology</i> , <b>2019</b> , 80, 228-236	2.3	19
152	Analysis of head and neck carcinoma progression reveals novel and relevant stage-specific changes associated with immortalisation and malignancy. <i>Scientific Reports</i> , <b>2019</b> , 9, 11992	4.9	13
151	Exome Sequencing in and -Negative Greek Families Identifies and as Candidate Risk Genes for Hereditary Breast Cancer. <i>Frontiers in Genetics</i> , <b>2019</b> , 10, 1005	4.5	7
150	Genome-wide analysis of androgen receptor binding and transcriptomic analysis in mesenchymal subsets during prostate development. <i>DMM Disease Models and Mechanisms</i> , <b>2019</b> , 12,	4.1	4
149	Stalled developmental programs at the root of pediatric brain tumors. <i>Nature Genetics</i> , <b>2019</b> , 51, 1702-	136.3	58
148	Recurrent noncoding U1´snRNA mutations drive cryptic splicing in SHH medulloblastoma. <i>Nature</i> , <b>2019</b> , 574, 707-711	50.4	78
147	Human candidate gene polymorphisms and risk of severe malaria in children in Kilifi, Kenya: a case-control association study. <i>Lancet Haematology,the</i> , <b>2018</b> , 5, e333-e345	14.6	38
146	Regulators of Asymmetric Cell Division in Breast Cancer. <i>Trends in Cancer</i> , <b>2018</b> , 4, 798-801	12.5	2
145	Global long non-coding RNA expression in the rostral anterior cingulate cortex of depressed suicides. <i>Translational Psychiatry</i> , <b>2018</b> , 8, 224	8.6	30
144	Current and Future Methods for mRNA Analysis: A Drive Toward Single Molecule Sequencing. <i>Methods in Molecular Biology</i> , <b>2018</b> , 1783, 209-241	1.4	22
143	Transcript Profiling Using Long-Read Sequencing Technologies. <i>Methods in Molecular Biology</i> , <b>2018</b> , 1783, 121-147	1.4	24

142	Sequencing of DICER1 in sarcomas identifies biallelic somatic DICER1 mutations in an adult-onset embryonal rhabdomyosarcoma. <i>British Journal of Cancer</i> , <b>2017</b> , 116, 1621-1626	8.7	24
141	A Targetable EGFR-Dependent Tumor-Initiating Program in Breast Cancer. <i>Cell Reports</i> , <b>2017</b> , 21, 1140-	114.9	45
140	Update on hypoxia-inducible factors and hydroxylases in oxygen regulatory pathways: from physiology to therapeutics. <i>Hypoxia (Auckland, NZ)</i> , <b>2017</b> , 5, 11-20	2.1	17
139	RNA Sequencing Reveals that Kaposi Sarcoma-Associated Herpesvirus Infection Mimics Hypoxia Gene Expression Signature. <i>PLoS Pathogens</i> , <b>2017</b> , 13, e1006143	7.6	21
138	The empress of subterfuge: cancer of the fallopian tube presenting with malapropism. <i>Lancet, The</i> , <b>2017</b> , 390, 1003-1004	40	
137	H3.1 K36M mutation in a congenital-onset soft tissue neoplasm. <i>Pediatric Blood and Cancer</i> , <b>2017</b> , 64, e26633	3	3
136	Identification of genes expressed in a mesenchymal subset regulating prostate organogenesis using tissue and single cell transcriptomics. <i>Scientific Reports</i> , <b>2017</b> , 7, 16385	4.9	7
135	A multidimensional integration analysis reveals potential bridging targets in the process of colorectal cancer liver metastasis. <i>PLoS ONE</i> , <b>2017</b> , 12, e0178760	3.7	6
134	Survival of metastatic melanoma patients after dendritic cell vaccination correlates with expression of leukocyte phosphatidylethanolamine-binding protein 1/Raf kinase inhibitory protein. <i>Oncotarget</i> , <b>2017</b> , 8, 67439-67456	3.3	12
133	High-sensitivity sequencing reveals multi-organ somatic mosaicism causing DICER1 syndrome. <i>Journal of Medical Genetics</i> , <b>2016</b> , 53, 43-52	5.8	51
132	Weighted gene co-expression network analysis of colorectal cancer liver metastasis genome sequencing data and screening of anti-metastasis drugs. <i>International Journal of Oncology</i> , <b>2016</b> , 49, 1108-18	4.4	19
131	Deep Sequencing Reveals Spatially Distributed Distinct Hot Spot Mutations in DICER1-Related Multinodular Goiter. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2016</b> , 101, 3637-3645	5.6	22
130	Benchmarking of the Oxford Nanopore MinION sequencing for quantitative and qualitative assessment of cDNA populations. <i>Scientific Reports</i> , <b>2016</b> , 6, 31602	4.9	98
129	Live single-cell laser tag. <i>Nature Communications</i> , <b>2016</b> , 7, 11636	17.4	14
128	Pattern recognition receptor mediated downregulation of microRNA-650 fine-tunes MxA expression in dendritic cells infected with influenza A virus. <i>European Journal of Immunology</i> , <b>2016</b> , 46, 167-77	6.1	10
127	Tumor hypoxia induces nuclear paraspeckle formation through HIF-2Idependent transcriptional activation of NEAT1 leading to cancer cell survival. <i>Oncogene</i> , <b>2015</b> , 34, 4482-90	9.2	164
126	De novo point mutations in patients diagnosed with ataxic cerebral palsy. <i>Brain</i> , <b>2015</b> , 138, 1817-32	11.2	101
125	Next-Generation Sequencing Analysis Reveals Differential Expression Profiles of MiRNA-mRNA Target Pairs in KSHV-Infected Cells. <i>PLoS ONE</i> , <b>2015</b> , 10, e0126439	3.7	16

124	Biomarker discovery: quantification of microRNAs and other small non-coding RNAs using next generation sequencing. <i>BMC Medical Genomics</i> , <b>2015</b> , 8, 35	3.7	56
123	Estrogen receptor-Idirectly regulates the hypoxia-inducible factor 1 pathway associated with antiestrogen response in breast cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2015</b> , 112, 15172-7	11.5	74
122	Integrated analysis of microRNA and mRNA expression and association with HIF binding reveals the complexity of microRNA expression regulation under hypoxia. <i>Molecular Cancer</i> , <b>2014</b> , 13, 28	42.1	104
121	Methods of quantifying microRNAs for hypoxia research: classic and next generation. <i>Antioxidants and Redox Signaling</i> , <b>2014</b> , 21, 1239-48	8.4	4
120	Microarray-based ultra-high resolution discovery of genomic deletion mutations. <i>BMC Genomics</i> , <b>2014</b> , 15, 224	4.5	4
119	Microarray-based optimization to detect genomic deletion mutations. <i>Genomics Data</i> , <b>2014</b> , 2, 53-54		4
118	Extensive regulation of the non-coding transcriptome by hypoxia: role of HIF in releasing paused RNApol2. <i>EMBO Reports</i> , <b>2014</b> , 15, 70-6	6.5	125
117	The molecular biology of the olive fly comes of age. <i>BMC Genetics</i> , <b>2014</b> , 15 Suppl 2, S8	2.6	13
116	Intestinal myofibroblast-specific Tpl2-Cox-2-PGE2 pathway links innate sensing to epithelial homeostasis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2014</b> , 111, E4658-67	11.5	52
115	IRF5:RelA interaction targets inflammatory genes in macrophages. <i>Cell Reports</i> , <b>2014</b> , 8, 1308-17	10.6	70
114	Olive fly transcriptomics analysis implicates energy metabolism genes in spinosad resistance. <i>BMC Genomics</i> , <b>2014</b> , 15, 714	4.5	20
113	ROS-mediated vascular homeostatic control of root-to-shoot soil Na delivery in Arabidopsis. <i>EMBO Journal</i> , <b>2013</b> , 32, 914-914	13	5
112	Next-generation sequencing (NGS) as a diagnostic tool for retinal degeneration reveals a much higher detection rate in early-onset disease. <i>European Journal of Human Genetics</i> , <b>2013</b> , 21, 274-80	5.3	107
111	miR-210 is a target of hypoxia-inducible factors 1 and 2 in renal cancer, regulates ISCU and correlates with good prognosis. <i>British Journal of Cancer</i> , <b>2013</b> , 108, 1133-42	8.7	112
110	MicroRNA expression profile in head and neck cancer: HOX-cluster embedded microRNA-196a and microRNA-10b dysregulation implicated in cell proliferation. <i>BMC Cancer</i> , <b>2013</b> , 13, 533	4.8	58
109	Next generation sequencing for molecular diagnosis of neurological disorders using ataxias as a model. <i>Brain</i> , <b>2013</b> , 136, 3106-18	11.2	128
108	High-resolution analysis of cis-acting regulatory networks at the Eglobin locus. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , <b>2013</b> , 368, 20120361	5.8	11
107	GATA-1 genome-wide occupancy associates with distinct epigenetic profiles in mouse fetal liver erythropoiesis. <i>Nucleic Acids Research</i> , <b>2013</b> , 41, 4938-48	20.1	19

### (2011-2013)

106	Smooth muscle cells differentiated from reprogrammed embryonic lung fibroblasts through DKK3 signaling are potent for tissue engineering of vascular grafts. <i>Circulation Research</i> , <b>2013</b> , 112, 1433-43	15.7	63
105	Cellular interference in craniofrontonasal syndrome: males mosaic for mutations in the X-linked EFNB1 gene are more severely affected than true hemizygotes. <i>Human Molecular Genetics</i> , <b>2013</b> , 22, 1654-62	5.6	50
104	Cross-species analysis reveals evolving and conserved features of the nuclear factor <b>B</b> (NF- <b>B</b> ) proteins. <i>Journal of Biological Chemistry</i> , <b>2013</b> , 288, 11546-54	5.4	13
103	Regulation of Hypoxia Responses by MicroRNA Expression <b>2013</b> , 267-285		
102	TGF-¶Smad2/3 signaling directly regulates several miRNAs in mouse ES cells and early embryos. <i>PLoS ONE</i> , <b>2013</b> , 8, e55186	3.7	12
101	ROS-mediated vascular homeostatic control of root-to-shoot soil Na delivery in Arabidopsis. <i>EMBO Journal</i> , <b>2012</b> , 31, 4359-70	13	120
100	A novel approach of homozygous haplotype sharing identifies candidate genes in autism spectrum disorder. <i>Human Genetics</i> , <b>2012</b> , 131, 565-79	6.3	150
99	Direct reprogramming of fibroblasts into endothelial cells capable of angiogenesis and reendothelialization in tissue-engineered vessels. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2012</b> , 109, 13793-8	11.5	194
98	Association of microRNA-221/222 and -323-3p with rheumatoid arthritis via predictions using the human TNF transgenic mouse model. <i>Arthritis Research and Therapy</i> , <b>2012</b> , 14,	5.7	1
97	Polycomb associates genome-wide with a specific RNA polymerase II variant, and regulates metabolic genes in ESCs. <i>Cell Stem Cell</i> , <b>2012</b> , 10, 157-70	18	221
96	Algorithm implementation for CNV discovery using Affymetrix and Illumina SNP array data. <i>Methods in Molecular Biology</i> , <b>2012</b> , 838, 291-310	1.4	1
95	Identification of microRNA-221/222 and microRNA-323-3p association with rheumatoid arthritis via predictions using the human tumour necrosis factor transgenic mouse model. <i>Annals of the Rheumatic Diseases</i> , <b>2012</b> , 71, 1716-23	2.4	83
94	Three-dimensional hydrogel structures as optical sensor arrays, for the detection of specific DNA sequences. <i>Analytical Biochemistry</i> , <b>2012</b> , 421, 1-8	3.1	13
93	Recessive mutations in SPTBN2 implicate III spectrin in both cognitive and motor development. <i>PLoS Genetics</i> , <b>2012</b> , 8, e1003074	6	74
92	Preparation of high-quality next-generation sequencing libraries from picogram quantities of target DNA. <i>Genome Research</i> , <b>2012</b> , 22, 125-33	9.7	45
91	Individual common variants exert weak effects on the risk for autism spectrum disorders. <i>Human Molecular Genetics</i> , <b>2012</b> , 21, 4781-92	5.6	279
90	Genome-wide analysis of mutations in mutant lineages selected following fast-neutron irradiation mutagenesis of Arabidopsis thaliana. <i>Genome Research</i> , <b>2012</b> , 22, 1306-15	9.7	87
89	Principles of dimer-specific gene regulation revealed by a comprehensive characterization of NF- <b>B</b> family DNA binding. <i>Nature Immunology</i> , <b>2011</b> , 13, 95-102	19.1	143

88	Direct targeting of Sec23a by miR-200s influences cancer cell secretome and promotes metastatic colonization. <i>Nature Medicine</i> , <b>2011</b> , 17, 1101-8	50.5	486
87	Animal models for arthritis: innovative tools for prevention and treatment. <i>Annals of the Rheumatic Diseases</i> , <b>2011</b> , 70, 1357-62	2.4	78
86	Extensive characterization of NF- <b>B</b> binding uncovers non-canonical motifs and advances the interpretation of genetic functional traits. <i>Genome Biology</i> , <b>2011</b> , 12, R70	18.3	110
85	High-resolution genome-wide mapping of HIF-binding sites by ChIP-seq. <i>Blood</i> , <b>2011</b> , 117, e207-17	2.2	484
84	The small-nucleolar RNAs commonly used for microRNA normalisation correlate with tumour pathology and prognosis. <i>British Journal of Cancer</i> , <b>2011</b> , 104, 1168-77	8.7	217
83	Regenerant Arabidopsis lineages display a distinct genome-wide spectrum of mutations conferring variant phenotypes. <i>Current Biology</i> , <b>2011</b> , 21, 1385-90	6.3	65
82	microRNA-associated progression pathways and potential therapeutic targets identified by integrated mRNA and microRNA expression profiling in breast cancer. <i>Cancer Research</i> , <b>2011</b> , 71, 5635-	4 <sup>1</sup> 5 <sup>0.1</sup>	253
81	Foxp2 regulates gene networks implicated in neurite outgrowth in the developing brain. <i>PLoS Genetics</i> , <b>2011</b> , 7, e1002145	6	198
80	Substantial histone reduction modulates genomewide nucleosomal occupancy and global transcriptional output. <i>PLoS Biology</i> , <b>2011</b> , 9, e1001086	9.7	140
79	The use of genome-wide eQTL associations in lymphoblastoid cell lines to identify novel genetic pathways involved in complex traits. <i>PLoS ONE</i> , <b>2011</b> , 6, e22070	3.7	35
78	An oncogenic role of eIF3e/INT6 in human breast cancer. <i>Oncogene</i> , <b>2010</b> , 29, 4080-9	9.2	41
77	Functional impact of global rare copy number variation in autism spectrum disorders. <i>Nature</i> , <b>2010</b> , 466, 368-72	50.4	1499
76	A large fraction of extragenic RNA pol II transcription sites overlap enhancers. <i>PLoS Biology</i> , <b>2010</b> , 8, e1000384	9.7	617
75	MicroRNA-210 regulates mitochondrial free radical response to hypoxia and krebs cycle in cancer cells by targeting iron sulfur cluster protein ISCU. <i>PLoS ONE</i> , <b>2010</b> , 5, e10345	3.7	243
74	A genome-wide scan for common alleles affecting risk for autism. <i>Human Molecular Genetics</i> , <b>2010</b> , 19, 4072-82	5.6	443
73	The histone demethylase JMJD2B is regulated by estrogen receptor alpha and hypoxia, and is a key mediator of estrogen induced growth. <i>Cancer Research</i> , <b>2010</b> , 70, 6456-66	10.1	141
72	Characterization of a family with rare deletions in CNTNAP5 and DOCK4 suggests novel risk loci for autism and dyslexia. <i>Biological Psychiatry</i> , <b>2010</b> , 68, 320-8	7.9	103
71	Association of the aromatase gene with Alzheimer's disease in women. <i>Neuroscience Letters</i> , <b>2010</b> , 468, 202-6	3.3	31

#### (2009-2010)

70	ATR-X syndrome protein targets tandem repeats and influences allele-specific expression in a size-dependent manner. <i>Cell</i> , <b>2010</b> , 143, 367-78	56.2	297
69	The role of hypoxia regulated microRNAs in cancer. <i>Current Topics in Microbiology and Immunology</i> , <b>2010</b> , 345, 47-70	3.3	33
68	A statistical approach for detecting genomic aberrations in heterogeneous tumor samples from single nucleotide polymorphism genotyping data. <i>Genome Biology</i> , <b>2010</b> , 11, R92	18.3	111
67	hsa-mir-210 is a marker of tumor hypoxia and a prognostic factor in head and neck cancer. <i>Cancer</i> , <b>2010</b> , 116, 2148-58	6.4	193
66	Identification and characterization of enhancers controlling the inflammatory gene expression program in macrophages. <i>Immunity</i> , <b>2010</b> , 32, 317-28	32.3	497
65	Chemokine gene expression in lung CD8 T cells correlates with protective immunity in mice immunized intra-nasally with Adenovirus-85A. <i>BMC Medical Genomics</i> , <b>2010</b> , 3, 46	3.7	10
64	Altered intra-nuclear organisation of heterochromatin and genes in ICF syndrome. <i>PLoS ONE</i> , <b>2010</b> , 5, e11364	3.7	22
63	Genome-wide association of hypoxia-inducible factor (HIF)-1alpha and HIF-2alpha DNA binding with expression profiling of hypoxia-inducible transcripts. <i>Journal of Biological Chemistry</i> , <b>2009</b> , 284, 16767-	1 <i>§7</i> 75	406
62	Phase I/II trial of bevacizumab and radiotherapy for locally advanced inoperable colorectal cancer: vasculature-independent radiosensitizing effect of bevacizumab. <i>Clinical Cancer Research</i> , <b>2009</b> , 15, 706	5 <del>9-7</del> 8	46
61	Comparative analysis of methods for gene transcription profiling data derived from different microarray technologies in rat and mouse models of diabetes. <i>BMC Genomics</i> , <b>2009</b> , 10, 63	4.5	15
60	Transcriptional signature of human adipose tissue-derived stem cells (hASCs) preconditioned for chondrogenesis in hypoxic conditions. <i>Experimental Cell Research</i> , <b>2009</b> , 315, 1937-52	4.2	37
59	Seeing clearly: the dominant and recessive nature of FOXE3 in eye developmental anomalies. <i>Human Mutation</i> , <b>2009</b> , 30, 1378-86	4.7	78
58	MicroRNA-125a is over-expressed in insulin target tissues in a spontaneous rat model of Type 2 Diabetes. <i>BMC Medical Genomics</i> , <b>2009</b> , 2, 54	3.7	96
57	A 15q13.3 microdeletion segregating with autism. European Journal of Human Genetics, 2009, 17, 687-9	25.3	114
56	Failed gene conversion leads to extensive end processing and chromosomal rearrangements in fission yeast. <i>EMBO Journal</i> , <b>2009</b> , 28, 3400-12	13	34
55	Genome-wide and fine-resolution association analysis of malaria in West Africa. <i>Nature Genetics</i> , <b>2009</b> , 41, 657-65	36.3	297
54	CMIP and ATP2C2 modulate phonological short-term memory in language impairment. <i>American Journal of Human Genetics</i> , <b>2009</b> , 85, 264-72	11	142
53	CpG methylation profiling in VHL related and VHL unrelated renal cell carcinoma. <i>Molecular Cancer</i> , <b>2009</b> , 8, 31	42.1	56

52	Analysis of FGGY as a risk factor for sporadic amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , <b>2009</b> , 10, 441-7		13
51	Genotyping technologies for genetic research. <i>Annual Review of Genomics and Human Genetics</i> , <b>2009</b> , 10, 117-33	9.7	164
50	Comparing CNV detection methods for SNP arrays. <i>Briefings in Functional Genomics &amp; Proteomics</i> , <b>2009</b> , 8, 353-66		150
49	Variations within oxygen-regulated gene expression in humans. <i>Journal of Applied Physiology</i> , <b>2009</b> , 106, 212-20	3.7	31
48	MicroRNA-10b and breast cancer metastasis. <i>Nature</i> , <b>2008</b> , 455, E8-9; author reply E9	50.4	113
47	A commonly occurring polymorphism upstream of the estrogen receptor alpha alters transcription and is associated with increased HDL. <i>Atherosclerosis</i> , <b>2008</b> , 199, 354-61	3.1	5
46	GenoSNP: a variational Bayes within-sample SNP genotyping algorithm that does not require a reference population. <i>Bioinformatics</i> , <b>2008</b> , 24, 2209-14	7.2	56
45	hsa-miR-210 Is induced by hypoxia and is an independent prognostic factor in breast cancer. <i>Clinical Cancer Research</i> , <b>2008</b> , 14, 1340-8	12.9	555
44	VACTERL/caudal regression/Currarino syndrome-like malformations in mice with mutation in the proprotein convertase Pcsk5. <i>Genes and Development</i> , <b>2008</b> , 22, 1465-77	12.6	99
43	Association of the KIAA0319 dyslexia susceptibility gene with reading skills in the general population. <i>American Journal of Psychiatry</i> , <b>2008</b> , 165, 1576-84	11.9	104
42	BeadArray-based genotyping. <i>Methods in Molecular Biology</i> , <b>2008</b> , 439, 53-74	1.4	12
41	Evaluating the effects of imputation on the power, coverage, and cost efficiency of genome-wide SNP platforms. <i>American Journal of Human Genetics</i> , <b>2008</b> , 83, 112-9	11	80
40	Analysis of DNA Methylation at the Human Alpha Globin Cluster during Hematopoiesis <i>Blood</i> , <b>2008</b> , 112, 1861-1861	2.2	
39	Functional conservation of Rel binding sites in drosophilid genomes. <i>Genome Research</i> , <b>2007</b> , 17, 1327-3	35 <sub>).7</sub>	13
38	QuantiSNP: an Objective Bayes Hidden-Markov Model to detect and accurately map copy number variation using SNP genotyping data. <i>Nucleic Acids Research</i> , <b>2007</b> , 35, 2013-25	20.1	462
37	Accuracy and reproducibility of protein-DNA microarray technology. <i>Advances in Biochemical Engineering/Biotechnology</i> , <b>2007</b> , 104, 87-110	1.7	9
36	Quantitative profiling of protein-DNA binding on microarrays. <i>Methods in Molecular Biology</i> , <b>2006</b> , 338, 261-80	1.4	4
35	Genomic reconstruction by serial mitotic recombination of yeast artificial chromosomes. <i>Methods in Molecular Biology</i> , <b>2006</b> , 349, 117-26	1.4	

#### (1999-2006)

34	Matrix-assisted laser desorption/ionisation, time-of-flight mass spectrometry in genomics research. <i>PLoS Genetics</i> , <b>2006</b> , 2, e100	6	92
33	The leukocyte receptor complex in chicken is characterized by massive expansion and diversification of immunoglobulin-like Loci. <i>PLoS Genetics</i> , <b>2006</b> , 2, e73	6	58
32	Concordant regulation of gene expression by hypoxia and 2-oxoglutarate-dependent dioxygenase inhibition: the role of HIF-1alpha, HIF-2alpha, and other pathways. <i>Journal of Biological Chemistry</i> , <b>2006</b> , 281, 15215-26	5.4	351
31	Affymetrix GeneChip system: moving from research to the clinic. <i>Expert Review of Molecular Diagnostics</i> , <b>2006</b> , 6, 145-52	3.8	31
30	Genotyping technologies for all. <i>Drug Discovery Today: Technologies</i> , <b>2006</b> , 3, 115-22	7.1	20
29	Alpha-haemoglobin stabilising protein is a quantitative trait gene that modifies the phenotype of beta-thalassaemia. <i>British Journal of Haematology</i> , <b>2006</b> , 133, 675-82	4.5	70
28	SW-ARRAY: a dynamic programming solution for the identification of copy-number changes in genomic DNA using array comparative genome hybridization data. <i>Nucleic Acids Research</i> , <b>2005</b> , 33, 345	5 <del>2</del> -6 <del>4</del>	79
27	Single nucleotide polymorphisms in TNFSF15 confer susceptibility to Crohns disease. <i>Human Molecular Genetics</i> , <b>2005</b> , 14, 3499-506	5.6	376
26	Development and evaluation of real competitive PCR for high-throughput quantitative applications. <i>Analytical Biochemistry</i> , <b>2005</b> , 339, 231-41	3.1	28
25	Genetically indistinguishable SNPs and their influence on inferring the location of disease-associated variants. <i>Genome Research</i> , <b>2005</b> , 15, 1503-10	9.7	22
24	Dissecting the genetic complexity of human 6p deletion syndromes by using a region-specific, phenotype-driven mouse screen. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2005</b> , 102, 12477-82	11.5	26
23	An Entirely Novel Form of IThalassemia in Patients from the South Pacific Linked to Chromosome 16 <i>Blood</i> , <b>2005</b> , 106, 2688-2688	2.2	
22	Quantitative high-throughput analysis of transcription factor binding specificities. <i>Nucleic Acids Research</i> , <b>2004</b> , 32, e44	20.1	67
21	Refined genotype-phenotype correlations in cases of chromosome 6p deletion syndromes. <i>European Journal of Human Genetics</i> , <b>2004</b> , 12, 718-28	5.3	49
20	Positional cloning identifies a novel cyclophilin as a candidate amplified oncogene in 1q21. <i>Oncogene</i> , <b>2002</b> , 21, 2261-9	9.2	50
19	Subchromosomal positioning of the epidermal differentiation complex (EDC) in keratinocyte and lymphoblast interphase nuclei. <i>Experimental Cell Research</i> , <b>2002</b> , 272, 163-75	4.2	170
18	Comparison of human chromosome 6p25 with mouse chromosome 13 reveals a greatly expanded ov-serpin gene repertoire in the mouse. <i>Genomics</i> , <b>2002</b> , 79, 349-62	4.3	55
17	Human epidermal differentiation complex in a single 2.5 Mbp long continuum of overlapping DNA cloned in bacteria integrating physical and transcript maps. <i>Journal of Investigative Dermatology</i> , <b>1999</b> , 112, 910-8	4.3	48

16	Isolation and characterization of human and mouse ZIRTL, a member of the IRT1 family of transporters, mapping within the epidermal differentiation complex. <i>Genomics</i> , <b>1999</b> , 62, 272-80	4.3	50
15	Human ovalbumin serpin evolution: phylogenic analysis, gene organization, and identification of new PI8-related genes suggest that two interchromosomal and several intrachromosomal duplications generated the gene clusters at 18q21-q23 and 6p25. <i>Genomics</i> , <b>1999</b> , 62, 490-9	4.3	40
14	Characterization and mapping of the human SOX4 gene. Mammalian Genome, 1993, 4, 577-84	3.2	47
13	New vector for transfer of yeast artificial chromosomes to mammalian cells. <i>Somatic Cell and Molecular Genetics</i> , <b>1993</b> , 19, 161-9		22
12	Mitotic recombination of yeast artificial chromosomes. <i>Nucleic Acids Research</i> , <b>1992</b> , 20, 3135-8	20.1	14
11	Genomic organization of the human folate receptor genes on chromosome 11q13. <i>Genomics</i> , <b>1992</b> , 14, 423-30	4.3	63
10	A testis-expressed Zn finger gene (ZNF76) in human 6p21.3 centromeric to the MHC is closely linked to the human homolog of the t-complex gene tcp-11. <i>Genomics</i> , <b>1992</b> , 14, 673-9	4.3	47
9	Single-cell RNA-seq reveals that glioblastoma recapitulates normal brain development		1
8	Statistical modeling, estimation, and remediation of sample index hopping in multiplexed droplet-based single-cell RNA-seq data		1
7	The genetic analysis of a founder Northern American population of European descent identifies FANCI as a candidate familial ovarian cancer risk gene		1
6	Oviduct epithelial cells constitute two developmentally distinct lineages that are spatially separated along the distal-proximal axis		1
5	Single-cell RNA sequencing reveals time- and sex-specific responses of spinal cord microglia to peripheral nerve injury and links ApoE to neuropathic pain		1
4	Single-nucleus RNA sequencing shows convergent evidence from different cell types for altered synaptic plasticity in major depressive disorder		5
3	Transcriptome landscape of the developing olive fruit fly embryo delineated by Oxford Nanopore long-read RNA-Seq		5
2	De novo genome assembly of the olive fruit fly (Bactrocera oleae) developed through a combination of linked-reads and long-read technologies		2
1	Maleness-on-the-Y (MoY) orchestrates male sex determination in major agricultural fruit fly pests		1