Jong-Il Kim

List of Publications by Year in Descending Order

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

181
papers7,013
citations38
h-index80
g-index195
ext. papers8,318
ext. citations8
avg, IF5.4
L-index

#	Paper	IF	Citations
181	Experimental development of the epigenomic library construction method to elucidate the epigenetic diversity and causal relationship between epigenome and transcriptome at a single-cell level <i>Genomics and Informatics</i> , 2022 , 20, e2	1.9	
180	Cross-Talk between Wnt Signaling and Src Tyrosine Kinase. <i>Biomedicines</i> , 2022 , 10, 1112	4.8	2
179	Expression-based species deconvolution and realignment removes misalignment error in multispecies single-cell data <i>BMC Bioinformatics</i> , 2022 , 23, 157	3.6	
178	JAK2 regulates paclitaxel resistance in triple negative breast cancers. <i>Journal of Molecular Medicine</i> , 2021 , 99, 1783-1795	5.5	2
177	High prevalence of TP53 loss and whole-genome doubling in early-onset colorectal cancer. <i>Experimental and Molecular Medicine</i> , 2021 , 53, 446-456	12.8	5
176	Sequencing cell-free fetal DNA in pregnant women with GCK-MODY: a proof-of-concept study. Journal of Clinical Endocrinology and Metabolism, 2021,	5.6	2
175	Engineered prime editors with PAM flexibility. <i>Molecular Therapy</i> , 2021 , 29, 2001-2007	11.7	20
174	Thyroid nodules in childhood-onset Hashimoto@thyroiditis: Frequency, risk factors, follow-up course and genetic alterations of thyroid cancer. <i>Clinical Endocrinology</i> , 2021 , 95, 638-648	3.4	2
173	S100A8/A9 mediate the reprograming of normal mammary epithelial cells induced by dynamic cell-cell interactions with adjacent breast cancer cells. <i>Scientific Reports</i> , 2021 , 11, 1337	4.9	4
172	Ablation of STAT3 in Purkinje cells reorganizes cerebellar synaptic plasticity in long-term fear memory network. <i>ELife</i> , 2021 , 10,	8.9	8
171	Conservation of copy number profiles during engraftment and passaging of patient-derived cancer xenografts. <i>Nature Genetics</i> , 2021 , 53, 86-99	36.3	44
170	Genomic profile of metastatic breast cancer patient-derived xenografts established using percutaneous biopsy. <i>Journal of Translational Medicine</i> , 2021 , 19, 7	8.5	
169	Discovery of acquired molecular signature on immune checkpoint inhibitors in paired tumor tissues. <i>Cancer Immunology, Immunotherapy</i> , 2021 , 70, 1755-1769	7.4	2
168	Predictive biomarkers for 5-fluorouracil and oxaliplatin-based chemotherapy in gastric cancers via profiling of patient-derived xenografts. <i>Nature Communications</i> , 2021 , 12, 4840	17.4	3
167	Glucose metabolic profiles evaluated by PET associated with molecular characteristic landscape of gastric cancer. <i>Gastric Cancer</i> , 2021 , 1	7.6	1
166	NTRK and RET fusion-directed therapy in pediatric thyroid cancer yields a tumor response and radioiodine uptake. <i>Journal of Clinical Investigation</i> , 2021 , 131,	15.9	13
165	Vertical sleeve gastrectomy induces distinctive transcriptomic responses in liver, fat and muscle. <i>Scientific Reports</i> , 2021 , 11, 2310	4.9	1

164	Epigenome signatures landscaped by histone H3K9me3 are associated with the synaptic dysfunction in Alzheimer@ disease. <i>Aging Cell</i> , 2020 , 19, e13153	9.9	20
163	Predominant Pathogenic Variants in Pediatric Follicular Thyroid Carcinomas. <i>Thyroid</i> , 2020 , 30, 1120-11	36.2	10
162	Outbreak investigation of Serratia marcescens neurosurgical site infections associated with a contaminated shaving razors. <i>Antimicrobial Resistance and Infection Control</i> , 2020 , 9, 64	6.2	5
161	Amplification of transglutaminase 2 enhances tumor-promoting inflammation in gastric cancers. <i>Experimental and Molecular Medicine</i> , 2020 , 52, 854-864	12.8	8
160	A newly developed capture-based sequencing panel for genomic assay of lung cancer. <i>Genes and Genomics</i> , 2020 , 42, 751-759	2.1	1
159	A glioneuronal tumor with fusion. <i>Npj Genomic Medicine</i> , 2020 , 5, 24	6.2	1
158	Downregulated miR-18b-5p triggers apoptosis by inhibition of calcium signaling and neuronal cell differentiation in transgenic SOD1 (G93A) mice and SOD1 (G17S and G86S) ALS patients. Translational Neurodegeneration, 2020, 9, 23	10.3	5
157	Whole genome sequencing of Nontuberculous Mycobacterium (NTM) isolates from sputum specimens of co-habiting patients with NTM pulmonary disease and NTM isolates from their environment. <i>BMC Genomics</i> , 2020 , 21, 322	4.5	11
156	Treatment strategy for papillary renal cell carcinoma type 2: a case series of seven patients treated based on next generation sequencing data. <i>Annals of Translational Medicine</i> , 2020 , 8, 1389-1389	3.2	О
155	DNAJB9 Inhibits p53-Dependent Oncogene-Induced Senescence and Induces Cell Transformation. <i>Molecules and Cells</i> , 2020 , 43, 397-407	3.5	1
154	STAT3 is a key molecule in the oncogenic behavior of diffuse intrinsic pontine glioma. <i>Oncology Letters</i> , 2020 , 20, 1989-1998	2.6	2
153	Protein Phosphatase 1H, Cyclin-Dependent Kinase Inhibitor p27, and Cyclin-Dependent Kinase 2 in Paclitaxel Resistance for Triple Negative Breast Cancers. <i>Journal of Breast Cancer</i> , 2020 , 23, 162-170	3	2
152	Comprehensive Molecular Characterization of Adenocarcinoma of the Gastroesophageal Junction Between Esophageal and Gastric Adenocarcinomas. <i>Annals of Surgery</i> , 2020 ,	7.8	9
151	A genome-wide by PM interaction study identifies novel loci for lung function near BICD1 and IL1RN-IL1F10 genes in Korean adults. <i>Chemosphere</i> , 2020 , 245, 125581	8.4	O
150	Phase II study of durvalumab and tremelimumab in pulmonary sarcomatoid carcinoma: KCSG-LU16-07. <i>Thoracic Cancer</i> , 2020 , 11, 3482-3489	3.2	5
149	Clinical Application of Next-Generation Sequencing-Based Panel to Wild-Type Advanced Melanoma Identifies Key Oncogenic Alterations and Therapeutic Strategies. <i>Molecular Cancer Therapeutics</i> , 2020 , 19, 937-944	6.1	7
148	A population-specific low-frequency variant of SLC22A12 (p.W258*) explains nearby genome-wide association signals for serum uric acid concentrations among Koreans. <i>PLoS ONE</i> , 2020 , 15, e0231336	3.7	О
147	NARD: whole-genome reference panel of 1779 Northeast Asians improves imputation accuracy of rare and low-frequency variants. <i>Genome Medicine</i> , 2019 , 11, 64	14.4	11

146	Priming mobilization of hair follicle stem cells triggers permanent loss of regeneration after alkylating chemotherapy. <i>Nature Communications</i> , 2019 , 10, 3694	17.4	13
145	Diagnostic Yield of Epilepsy Panel Testing in Patients With Seizure Onset Within the First Year of Life. <i>Frontiers in Neurology</i> , 2019 , 10, 988	4.1	14
144	Targeted next-generation DNA sequencing identifies Notch signaling pathway mutation as a predictor of radiation response. <i>International Journal of Radiation Biology</i> , 2019 , 95, 1640-1647	2.9	О
143	Genetic variations associated with response to dutasteride in the treatment of male subjects with androgenetic alopecia. <i>PLoS ONE</i> , 2019 , 14, e0222533	3.7	3
142	Unstable Genome and Transcriptome Dynamics during Tumor Metastasis Contribute to Therapeutic Heterogeneity in Colorectal Cancers. <i>Clinical Cancer Research</i> , 2019 , 25, 2821-2834	12.9	8
141	Integrative analysis of genomic and transcriptomic characteristics associated with progression of aggressive thyroid cancer. <i>Nature Communications</i> , 2019 , 10, 2764	17.4	80
140	Tumor Suppressor miRNA-204-5p Regulates Growth, Metastasis, and Immune Microenvironment Remodeling in Breast Cancer. <i>Cancer Research</i> , 2019 , 79, 1520-1534	10.1	83
139	Identifying Pathogenic Variants of Monogenic Diabetes Using Targeted Panel Sequencing in an East Asian Population. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 ,	5.6	16
138	Dissecting the phenotypic and genetic spectrum of early childhood-onset generalized epilepsies. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019 , 71, 222-228	3.2	1
137	A Breakdown in Metabolic Reprogramming Causes Microglia Dysfunction in Alzheimer@ Disease. <i>Cell Metabolism</i> , 2019 , 30, 493-507.e6	24.6	170
137		24.6	170
	Cell Metabolism, 2019, 30, 493-507.e6 The novel high-frequency variant of TRPV3 p.A628T in East Asians showing faster sensitization in		
136	Cell Metabolism, 2019, 30, 493-507.e6 The novel high-frequency variant of TRPV3 p.A628T in East Asians showing faster sensitization in response to chemical agonists. Pflugers Archiv European Journal of Physiology, 2019, 471, 1273-1289 A familial case of limb-girdle muscular dystrophy with CAV3 mutation. Journal of Genetic Medicine,	4.6	2
136 135	Cell Metabolism, 2019, 30, 493-507.e6 The novel high-frequency variant of TRPV3 p.A628T in East Asians showing faster sensitization in response to chemical agonists. Pflugers Archiv European Journal of Physiology, 2019, 471, 1273-1289 A familial case of limb-girdle muscular dystrophy with CAV3 mutation. Journal of Genetic Medicine, 2019, 16, 67-70 Comparative genomics of Mycoplasma pneumoniae isolated from children with pneumonia: South	4.6	2
136 135 134	Cell Metabolism, 2019, 30, 493-507.e6 The novel high-frequency variant of TRPV3 p.A628T in East Asians showing faster sensitization in response to chemical agonists. Pflugers Archiv European Journal of Physiology, 2019, 471, 1273-1289 A familial case of limb-girdle muscular dystrophy with CAV3 mutation. Journal of Genetic Medicine, 2019, 16, 67-70 Comparative genomics of Mycoplasma pneumoniae isolated from children with pneumonia: South Korea, 2010-2016. BMC Genomics, 2019, 20, 910 Identification of African-Specific Admixture between Modern and Archaic Humans. American	4.6 0.2 4.5	2 0
136 135 134	Cell Metabolism, 2019, 30, 493-507.e6 The novel high-frequency variant of TRPV3 p.A628T in East Asians showing faster sensitization in response to chemical agonists. Pflugers Archiv European Journal of Physiology, 2019, 471, 1273-1289 A familial case of limb-girdle muscular dystrophy with CAV3 mutation. Journal of Genetic Medicine, 2019, 16, 67-70 Comparative genomics of Mycoplasma pneumoniae isolated from children with pneumonia: South Korea, 2010-2016. BMC Genomics, 2019, 20, 910 Identification of African-Specific Admixture between Modern and Archaic Humans. American Journal of Human Genetics, 2019, 105, 1254-1261 Interactions of CDH13 gene polymorphisms and ambient PM air pollution exposure with blood	4.6 0.2 4.5	2 0 4
136 135 134 133	Cell Metabolism, 2019, 30, 493-507.e6 The novel high-frequency variant of TRPV3 p.A628T in East Asians showing faster sensitization in response to chemical agonists. Pflugers Archiv European Journal of Physiology, 2019, 471, 1273-1289 A familial case of limb-girdle muscular dystrophy with CAV3 mutation. Journal of Genetic Medicine, 2019, 16, 67-70 Comparative genomics of Mycoplasma pneumoniae isolated from children with pneumonia: South Korea, 2010-2016. BMC Genomics, 2019, 20, 910 Identification of African-Specific Admixture between Modern and Archaic Humans. American Journal of Human Genetics, 2019, 105, 1254-1261 Interactions of CDH13 gene polymorphisms and ambient PM air pollution exposure with blood pressure and hypertension in Korean men. Chemosphere, 2019, 218, 292-298 Alterations in the Rho pathway contribute to Epstein-Barr virus-induced lymphomagenesis in	4.6 0.2 4.5 11 8.4	2 0 4

(2017-2018)

128	Development of a common platform for the noninvasive prenatal diagnosis of X-linked diseases. <i>Prenatal Diagnosis</i> , 2018 , 38, 835-840	3.2	О	
127	Genome-Wide Association Study Reveals Distinct Genetic Susceptibility of Thyroid Nodules From Thyroid Cancer. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018 , 103, 4384-4394	5.6	5	
126	Targeted linked-read sequencing for direct haplotype phasing of maternal DMD alleles: a practical and reliable method for noninvasive prenatal diagnosis. <i>Scientific Reports</i> , 2018 , 8, 8678	4.9	15	
125	3-Oxoacid CoA transferase 1 as a therapeutic target gene for cisplatin-resistant ovarian cancer. <i>Oncology Letters</i> , 2018 , 15, 2611-2618	2.6	9	
124	Editor@Introduction to This Issue (G&I 16:1, 2018). Genomics and Informatics, 2018, 16, 1	1.9	1	
123	Genomic characterization of clonal evolution during oropharyngeal carcinogenesis driven by human papillomavirus 16. <i>BMB Reports</i> , 2018 , 51, 584-589	5.5	4	
122	Editor@Introduction to This Issue (G&I 16:2, 2018). Genomics and Informatics, 2018, 16, 21	1.9		
121	MicroRNA Expression Profiles in Gastric Carcinogenesis. <i>Scientific Reports</i> , 2018 , 8, 14393	4.9	46	
120	Nonsynonymous Variants in and Are Associated With Type 2 Diabetes in an East Asian Population. <i>Diabetes</i> , 2018 , 67, 1892-1902	0.9	23	
119	FARS2 mutation and epilepsy: Possible link with early-onset epileptic encephalopathy. <i>Epilepsy Research</i> , 2017 , 129, 118-124	3	15	
118	CDH13 gene-by-PM interaction effect on lung function decline in Korean men. <i>Chemosphere</i> , 2017 , 168, 583-589	8.4	9	
117	High prevalence of TP53 mutations is associated with poor survival and an EMT signature in gliosarcoma patients. <i>Experimental and Molecular Medicine</i> , 2017 , 49, e317	12.8	24	
116	Abdominal adiposity intensifies the negative effects of ambient air pollution on lung function in Korean men. <i>International Journal of Obesity</i> , 2017 , 41, 1218-1223	5.5	13	
115	A Novel Combination Treatment Targeting BCL-X and MCL1 for -mutated and -amplified Colorectal Cancers. <i>Molecular Cancer Therapeutics</i> , 2017 , 16, 2178-2190	6.1	12	
114	Transcriptome analyses of chronic traumatic encephalopathy show alterations in protein phosphatase expression associated with tauopathy. <i>Experimental and Molecular Medicine</i> , 2017 , 49, e33	3 ^{12.8}	21	
113	Glutaminase 2 expression is associated with regional heterogeneity of 5-aminolevulinic acid fluorescence in glioblastoma. <i>Scientific Reports</i> , 2017 , 7, 12221	4.9	17	
112	Altered nucleocytoplasmic proteome and transcriptome distributions in an in vitro model of amyotrophic lateral sclerosis. <i>PLoS ONE</i> , 2017 , 12, e0176462	3.7	13	
111	Findings of a 1303 Korean whole-exome sequencing study. <i>Experimental and Molecular Medicine</i> , 2017 , 49, e356	12.8	23	

110	Genome-wide association and expression quantitative trait loci studies identify multiple susceptibility loci for thyroid cancer. <i>Nature Communications</i> , 2017 , 8, 15966	17.4	46
109	Markers of disease and steroid responsiveness in paediatric idiopathic nephrotic syndrome: Whole-transcriptome sequencing of peripheral blood mononuclear cells. <i>Journal of International Medical Research</i> , 2017 , 45, 948-963	1.4	9
108	A copy number variation in PKD1L2 is associated with colorectal cancer predisposition in korean population. <i>International Journal of Cancer</i> , 2017 , 140, 86-94	7·5	4
107	Analysis of Gene Expression in Human Dermal Fibroblasts Treated with Senescence-Modulating COX Inhibitors. <i>Genomics and Informatics</i> , 2017 , 15, 56-64	1.9	1
106	Dysregulated Wnt signalling and recurrent mutations of the tumour suppressor RNF43 in early gastric carcinogenesis. <i>Journal of Pathology</i> , 2016 , 240, 304-314	9.4	35
105	Clinical whole exome sequencing in early onset diabetes patients. <i>Diabetes Research and Clinical Practice</i> , 2016 , 122, 71-77	7.4	23
104	Integrated analysis of omics data using microRNA-target mRNA network and PPI network reveals regulation of Gnai1 function in the spinal cord of Ews/Ewsr1 KO mice. <i>BMC Medical Genomics</i> , 2016 , 9 Suppl 1, 33	3.7	1
103	Deep resequencing of 131 Crohn@ disease associated genes in pooled DNA confirmed three reported variants and identified eight novel variants. <i>Gut</i> , 2016 , 65, 788-96	19.2	54
102	Clinical application of genomic profiling to find druggable targets for adolescent and young adult (AYA) cancer patients with metastasis. <i>BMC Cancer</i> , 2016 , 16, 170	4.8	24
101	Comprehensive Analysis of the Transcriptional and Mutational Landscape of Follicular and Papillary Thyroid Cancers. <i>PLoS Genetics</i> , 2016 , 12, e1006239	6	178
100	xCyp26c Induced by Inhibition of BMP Signaling Is Involved in Anterior-Posterior Neural Patterning of Xenopus laevis. <i>Molecules and Cells</i> , 2016 , 39, 352-7	3.5	7
99	Whole-exome and transcriptome sequencing of refractory diffuse large B-cell lymphoma. <i>Oncotarget</i> , 2016 , 7, 86433-86445	3.3	31
98	NTRK1 fusions for the therapeutic intervention of Korean patients with colon cancer. <i>Oncotarget</i> , 2016 , 7, 8399-412	3.3	16
97	Draft Genome of , a Pathogen Responsible for Visceral Larva Migrans. <i>Korean Journal of Parasitology</i> , 2016 , 54, 751-758	1.7	3
96	Isolation of Middle East Respiratory Syndrome Coronavirus from a Patient of the 2015 Korean Outbreak. <i>Journal of Korean Medical Science</i> , 2016 , 31, 315-20	4.7	13
95	Estimation of Prognostic Marker Genes by Public Microarray Data in Patients with Ovarian Serous Cystadenocarcinoma. <i>Yonsei Medical Journal</i> , 2016 , 57, 872-8	3	
94	Viral Load Kinetics of MERS Coronavirus Infection. New England Journal of Medicine, 2016, 375, 1303-5	59.2	155
93	Noninvasive prenatal diagnosis of duchenne muscular dystrophy: comprehensive genetic diagnosis in carrier, proband, and fetus. <i>Clinical Chemistry</i> , 2015 , 61, 829-37	5.5	35

(2014-2015)

92	Genomic alterations in BCL2L1 and DLC1 contribute to drug sensitivity in gastric cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015 , 112, 12492-7	11.5	36
91	The full-length DNA sequence of Epstein Barr virus from a human gastric carcinoma cell line, SNU-719. <i>Virus Genes</i> , 2015 , 51, 329-37	2.3	9
90	Deregulation of immune response genes in patients with Epstein-Barr virus-associated gastric cancer and outcomes. <i>Gastroenterology</i> , 2015 , 148, 137-147.e9	13.3	85
89	Genetic association of APOA5 and APOE with metabolic syndrome and their interaction with health-related behavior in Korean men. <i>Lipids in Health and Disease</i> , 2015 , 14, 105	4.4	15
88	Association between salivary amylase (AMY1) gene copy numbers and insulin resistance in asymptomatic Korean men. <i>Diabetic Medicine</i> , 2015 , 32, 1588-95	3.5	34
87	I148M variant in PNPLA3 reduces central adiposity and metabolic disease risks while increasing nonalcoholic fatty liver disease. <i>Liver International</i> , 2015 , 35, 2537-46	7.9	24
86	Recurrent fusion transcripts detected by whole-transcriptome sequencing of 120 primary breast cancer samples. <i>Genes Chromosomes and Cancer</i> , 2015 , 54, 681-91	5	30
85	A Common Variant of NGEF Is Associated with Abdominal Visceral Fat in Korean Men. <i>PLoS ONE</i> , 2015 , 10, e0137564	3.7	11
84	Genetic alterations of JAK/STAT cascade and histone modification in extranodal NK/T-cell lymphoma nasal type. <i>Oncotarget</i> , 2015 , 6, 17764-76	3.3	104
83	Genomic Copy Number Variations Characterize the Prognosis of Both P16-Positive and P16-Negative Oropharyngeal Squamous Cell Carcinoma After Curative Resection. <i>Medicine (United States)</i> , 2015 , 94, e2187	1.8	9
82	Digenome-seq: genome-wide profiling of CRISPR-Cas9 off-target effects in human cells. <i>Nature Methods</i> , 2015 , 12, 237-43, 1 p following 243	21.6	652
81	Colorectal cancer-susceptibility single-nucleotide polymorphisms in Korean population. <i>Journal of Gastroenterology and Hepatology (Australia)</i> , 2015 , 30, 849-57	4	12
80	Family-Based Association Study of Pulmonary Function in a Population in Northeast Asia. <i>PLoS ONE</i> , 2015 , 10, e0139716	3.7	6
79	Association of HLA Genotype and Fulminant Type 1 Diabetes in Koreans. <i>Genomics and Informatics</i> , 2015 , 13, 126-31	1.9	9
78	A multifunctional protein EWS regulates the expression of Drosha and microRNAs. <i>Cell Death and Differentiation</i> , 2014 , 21, 136-45	12.7	28
77	Novel fusion transcripts in human gastric cancer revealed by transcriptome analysis. <i>Oncogene</i> , 2014 , 33, 5434-41	9.2	57
76	Hoyeraal-Hreidarsson syndrome with a DKC1 mutation identified by whole-exome sequencing. <i>Gene</i> , 2014 , 546, 425-9	3.8	9
75	A role of placental growth factor in hair growth. <i>Journal of Dermatological Science</i> , 2014 , 74, 125-34	4.3	23

74	Somatic deletions implicated in functional diversity of brain cells of individuals with schizophrenia and unaffected controls. <i>Scientific Reports</i> , 2014 , 4, 3807	4.9	20
73	The First Kazakh Whole Genomes: The First Report of NGS Data. <i>Central Asian Journal of Global Health</i> , 2014 , 3, 146	0.8	
72	RNA editing in RHOQ promotes invasion potential in colorectal cancer. <i>Journal of Experimental Medicine</i> , 2014 , 211, 613-21	16.6	67
71	Genome-wide characterization of the routes to pluripotency. <i>Nature</i> , 2014 , 516, 198-206	50.4	153
70	RANBP2-ALK fusion combined with monosomy 7 in acute myelomonocytic leukemia. <i>Cancer Genetics</i> , 2014 , 207, 40-5	2.3	15
69	Targeted next-generation sequencing at copy-number breakpoints for personalized analysis of rearranged ends in solid tumors. <i>PLoS ONE</i> , 2014 , 9, e100089	3.7	3
68	Analysis of gene expression in cyclooxygenase-2-overexpressed human osteosarcoma cell lines. <i>Genomics and Informatics</i> , 2014 , 12, 247-53	1.9	20
67	RNA Editing in RHOQ Promotes Invasion Potential in Colorectal Cancer. <i>Journal of Cell Biology</i> , 2014 , 204, 2047OIA60	7.3	1
66	Epigenetic regulation of cholinergic receptor M1 (CHRM1) by histone H3K9me3 impairs Ca(2+) signaling in Huntington@ disease. <i>Acta Neuropathologica</i> , 2013 , 125, 727-39	14.3	38
65	Combined linkage and association analyses identify a novel locus for obesity near PROX1 in Asians. <i>Obesity</i> , 2013 , 21, 2405-12	8	15
64	Molecular diagnosis of congenital muscular dystrophies with defective glycosylation of alpha-dystroglycan using next-generation sequencing technology. <i>Neuromuscular Disorders</i> , 2013 , 23, 337-44	2.9	11
63	TIARA genome database: update 2013. <i>Database: the Journal of Biological Databases and Curation</i> , 2013 , 2013, bat003	5	5
62	Targeted resequencing of candidate genes reveals novel variants associated with severe BehëtQ uveitis. Experimental and Molecular Medicine, 2013, 45, e49	12.8	13
61	Gene expression profiling by mRNA sequencing reveals increased expression of immune/inflammation-related genes in the hippocampus of individuals with schizophrenia. <i>Translational Psychiatry</i> , 2013 , 3, e321	8.6	116
60	A family-based association study after genome-wide linkage analysis identified two genetic loci for renal function in a Mongolian population. <i>Kidney International</i> , 2013 , 83, 285-92	9.9	12
59	Targeted sequencing of cancer-related genes in colorectal cancer using next-generation sequencing. <i>PLoS ONE</i> , 2013 , 8, e64271	3.7	61
58	Exomic sequencing of immune-related genes reveals novel candidate variants associated with alopecia universalis. <i>PLoS ONE</i> , 2013 , 8, e53613	3.7	12
57	Heritabilities of facial measurements and their latent factors in korean families. <i>Genomics and Informatics</i> , 2013 , 11, 83-92	1.9	13

56	FX: an RNA-Seq analysis tool on the cloud. <i>Bioinformatics</i> , 2012 , 28, 721-3	7.2	57
55	The transcriptional landscape and mutational profile of lung adenocarcinoma. <i>Genome Research</i> , 2012 , 22, 2109-19	9.7	435
54	Epidemiologic characteristics of intraocular pressure in the Korean and Mongolian populations: the Healthy Twin and the GENDISCAN study. <i>Ophthalmology</i> , 2012 , 119, 450-7	7.3	29
53	A transforming KIF5B and RET gene fusion in lung adenocarcinoma revealed from whole-genome and transcriptome sequencing. <i>Genome Research</i> , 2012 , 22, 436-45	9.7	367
52	Copy number variation of age-related macular degeneration relevant genes in the Korean population. <i>PLoS ONE</i> , 2012 , 7, e31243	3.7	10
51	RNA-Seq analysis of frontal cortex and cerebellum from 5XFAD mice at early stage of disease pathology. <i>Journal of Alzheimerjs Disease</i> , 2012 , 29, 793-808	4.3	31
50	A public resource facilitating clinical use of genomes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012 , 109, 11920-7	11.5	154
49	Gene mapping study for constitutive skin color in an isolated Mongolian population. <i>Experimental and Molecular Medicine</i> , 2012 , 44, 241-9	12.8	8
48	Comprehensive genomic analyses associate UGT8 variants with musical ability in a Mongolian population. <i>Journal of Medical Genetics</i> , 2012 , 49, 747-52	5.8	38
47	Extensive genomic and transcriptional diversity identified through massively parallel DNA and RNA sequencing of eighteen Korean individuals. <i>Nature Genetics</i> , 2011 , 43, 745-52	36.3	110
46	Upregulation of neuronal nitric oxide synthase in the periphery promotes pain hypersensitivity after peripheral nerve injury. <i>Neuroscience</i> , 2011 , 190, 367-78	3.9	28
45	xCITED2 Induces Neural Genes in Animal Cap Explants of Xenopus Embryos. <i>Experimental Neurobiology</i> , 2011 , 20, 123-9	4	2
44	The function of heterodimeric AP-1 comprised of c-Jun and c-Fos in activin mediated Spemann organizer gene expression. <i>PLoS ONE</i> , 2011 , 6, e21796	3.7	6
43	Whole-exome sequencing identifies mutations of KIF22 in spondyloepimetaphyseal dysplasia with joint laxity, leptodactylic type. <i>American Journal of Human Genetics</i> , 2011 , 89, 760-6	11	34
42	Genetic diagnosis of Duchenne and Becker muscular dystrophy using next-generation sequencing technology: comprehensive mutational search in a single platform. <i>Journal of Medical Genetics</i> , 2011 , 48, 731-6	5.8	70
41	TIARA: a database for accurate analysis of multiple personal genomes based on cross-technology. Nucleic Acids Research, 2011 , 39, D883-8	20.1	15
40	Linkage and association scan for tanning ability in an isolated Mongolian population. <i>BMB Reports</i> , 2011 , 44, 741-6	5.5	7
39	Discovery of common Asian copy number variants using integrated high-resolution array CGH and massively parallel DNA sequencing. <i>Nature Genetics</i> , 2010 , 42, 400-5	36.3	167

38	Reference-unbiased copy number variant analysis using CGH microarrays. <i>Nucleic Acids Research</i> , 2010 , 38, e190	20.1	20
37	Replication of a glaucoma candidate gene on 5q22.1 for intraocular pressure in mongolian populations: the GENDISCAN Project 2010 , 51, 1335-40		15
36	Genome-wide linkage analysis for ocular and nasal anthropometric traits in a Mongolian population. <i>Experimental and Molecular Medicine</i> , 2010 , 42, 799-804	12.8	5
35	The first Irish genome and ways of improving sequence accuracy. <i>Genome Biology</i> , 2010 , 11, 132	18.3	4
34	Analysis of genetic and non-genetic factors that affect the QTc interval in a Mongolian population: the GENDISCAN study. <i>Experimental and Molecular Medicine</i> , 2009 , 41, 841-8	12.8	10
33	Higher mitochondrial DNA copy number is associated with lower prevalence of microalbuminuria. <i>Experimental and Molecular Medicine</i> , 2009 , 41, 253-8	12.8	27
32	SIRT1 regulates tyrosine hydroxylase expression and differentiation of neuroblastoma cells via FOXO3a. <i>FEBS Letters</i> , 2009 , 583, 1183-8	3.8	47
31	A highly annotated whole-genome sequence of a Korean individual. <i>Nature</i> , 2009 , 460, 1011-5	50.4	265
30	Detection of hydin Gene Duplication in Personal Genome Sequence Data. <i>Genomics and Informatics</i> , 2009 , 7, 159-162	1.9	2
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18	Downstream components of RhoA required for signal pathway of superoxide formation during phagocytosis of serum opsonized zymosans in macrophages. <i>Experimental and Molecular Medicine</i> , 2005 , 37, 575-87	12.8	27
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	Phosphorylation of 46-kDa protein of synaptic vesicle membranes is stimulated by GTP and		
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Whole-genome reference panel of 1,781 Northeast Asians improves imputation accuracy of rare and low-frequency variants

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