## Jong-Il Kim

## List of Publications by Year in descending order

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187	9,398	42	89
papers	citations	h-index	g-index
195	195	195	17865
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Digenome-seq: genome-wide profiling of CRISPR-Cas9 off-target effects in human cells. Nature Methods, 2015, 12, 237-243.	19.0	850
2	The transcriptional landscape and mutational profile of lung adenocarcinoma. Genome Research, 2012, 22, 2109-2119.	5 <b>.</b> 5	524
3	A transforming <i>KIF5B</i> and <i>RET</i> gene fusion in lung adenocarcinoma revealed from whole-genome and transcriptome sequencing. Genome Research, 2012, 22, 436-445.	5.5	433
4	Heat Shock Protein 70 Inhibits Apoptosis Downstream of Cytochrome c Release and Upstream of Caspase-3 Activation. Journal of Biological Chemistry, 2000, 275, 25665-25671.	3.4	410
5	A Breakdown in Metabolic Reprogramming Causes Microglia Dysfunction in Alzheimer's Disease. Cell Metabolism, 2019, 30, 493-507.e6.	16.2	374
6	The Fine-Scale and Complex Architecture of Human Copy-Number Variation. American Journal of Human Genetics, 2008, 82, 685-695.	6.2	315
7	A highly annotated whole-genome sequence of a Korean individual. Nature, 2009, 460, 1011-1015.	27.8	295
8	Comprehensive Analysis of the Transcriptional and Mutational Landscape of Follicular and Papillary Thyroid Cancers. PLoS Genetics, 2016, 12, e1006239.	3.5	265
9	A public resource facilitating clinical use of genomes. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 11920-11927.	7.1	194
10	The opsonin MFG-E8 is a ligand for the $\hat{l}\pm v\hat{l}^25$ integrin and triggers DOCK180-dependent Rac1 activation for the phagocytosis of apoptotic cells. Experimental Cell Research, 2004, 292, 403-416.	2.6	193
11	p53-mediated induction of Cox-2 counteracts p53- or genotoxic stress-induced apoptosis. EMBO Journal, 2002, 21, 5635-5644.	7.8	192
12	Genome-wide characterization of the routes to pluripotency. Nature, 2014, 516, 198-206.	27.8	187
13	Viral Load Kinetics of MERS Coronavirus Infection. New England Journal of Medicine, 2016, 375, 1303-1305.	27.0	186
14	Discovery of common Asian copy number variants using integrated high-resolution array CGH and massively parallel DNA sequencing. Nature Genetics, 2010, 42, 400-405.	21.4	179
15	Integrative analysis of genomic and transcriptomic characteristics associated with progression of aggressive thyroid cancer. Nature Communications, 2019, 10, 2764.	12.8	166
16	Gene expression profiling by mRNA sequencing reveals increased expression of immune/inflammation-related genes in the hippocampus of individuals with schizophrenia. Translational Psychiatry, 2013, 3, e321-e321.	4.8	162
17	p53 induction and activation of DDR1 kinase counteract p53-mediated apoptosis and influence p53 regulation through a positive feedback loop. EMBO Journal, 2003, 22, 1289-1301.	7.8	156
18	Genetic alterations of JAK/STAT cascade and histone modification in extranodal NK/T-cell lymphoma nasal type. Oncotarget, 2015, 6, 17764-17776.	1.8	136

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19	Deregulation of Immune Response Genes in Patients With Epstein-Barr Virus-Associated Gastric Cancer and Outcomes. Gastroenterology, 2015, 148, 137-147.e9.	1.3	127
20	Tumor Suppressor miRNA-204-5p Regulates Growth, Metastasis, and Immune Microenvironment Remodeling in Breast Cancer. Cancer Research, 2019, 79, 1520-1534.	0.9	126
21	Transforming growth factor-β1 regulates macrophage migration via RhoA. Blood, 2006, 108, 1821-1829.	1.4	124
22	Extensive genomic and transcriptional diversity identified through massively parallel DNA and RNA sequencing of eighteen Korean individuals. Nature Genetics, 2011, 43, 745-752.	21.4	121
23	Conservation of copy number profiles during engraftment and passaging of patient-derived cancer xenografts. Nature Genetics, 2021, 53, 86-99.	21.4	118
24	T Cell Lymphoma in Transgenic Mice Expressing the HumanHsp70Gene. Biochemical and Biophysical Research Communications, 1996, 218, 582-587.	2.1	103
25	RNA editing in <i>RHOQ</i> promotes invasion potential in colorectal cancer. Journal of Experimental Medicine, 2014, 211, 613-621.	8.5	97
26	Targeted disruption of hsp70.1 sensitizes to osmotic stress. EMBO Reports, 2002, 3, 857-861.	4.5	85
27	Genetic diagnosis of Duchenne and Becker muscular dystrophy using next-generation sequencing technology: comprehensive mutational search in a single platform. Journal of Medical Genetics, 2011, 48, 731-736.	3.2	80
28	Novel fusion transcripts in human gastric cancer revealed by transcriptome analysis. Oncogene, 2014, 33, 5434-5441.	5.9	74
29	Targeted Sequencing of Cancer-Related Genes in Colorectal Cancer Using Next-Generation Sequencing. PLoS ONE, 2013, 8, e64271.	2.5	71
30	Cyclooxygenase-2 promotes cell proliferation, migration and invasion in U2OS human osteosarcoma cells. Experimental and Molecular Medicine, 2007, 39, 469-476.	7.7	66
31	FX: an RNA-Seq analysis tool on the cloud. Bioinformatics, 2012, 28, 721-723.	4.1	66
32	MicroRNA Expression Profiles in Gastric Carcinogenesis. Scientific Reports, 2018, 8, 14393.	3.3	65
33	Genome-wide association and expression quantitative trait loci studies identify multiple susceptibility loci for thyroid cancer. Nature Communications, 2017, 8, 15966.	12.8	64
34	Deep resequencing of 131 Crohn's disease associated genes in pooled DNA confirmed three reported variants and identified eight novel variants. Gut, 2016, 65, 788-796.	12.1	62
35	NTRK and RET fusion–directed therapy in pediatric thyroid cancer yields a tumor response and radioiodine uptake. Journal of Clinical Investigation, 2021, 131, .	8.2	62
36	Engineered prime editors with PAM flexibility. Molecular Therapy, 2021, 29, 2001-2007.	8.2	56

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37	SIRT1 regulates tyrosine hydroxylase expression and differentiation of neuroblastoma cells via FOXO3a. FEBS Letters, 2009, 583, 1183-1188.	2.8	55
38	Epigenome signatures landscaped by histone H3K9me3 are associated with the synaptic dysfunction in Alzheimer's disease. Aging Cell, 2020, 19, e13153.	6.7	53
39	COX-2 regulates p53 activity and inhibits DNA damage-induced apoptosis. Biochemical and Biophysical Research Communications, 2005, 328, 1107-1112.	2.1	51
40	Phosphorylation of c-Crk II on the Negative Regulatory Tyr222 Mediates Nerve Growth Factor-induced Cell Spreading and Morphogenesis. Journal of Biological Chemistry, 2000, 275, 24787-24797.	3.4	49
41	Comprehensive genomic analyses associate <i>UGT8</i> variants with musical ability in a Mongolian population. Journal of Medical Genetics, 2012, 49, 747-752.	3.2	48
42	Epigenetic regulation of cholinergic receptor M1 (CHRM1) by histone H3K9me3 impairs Ca2+ signaling in Huntington's disease. Acta Neuropathologica, 2013, 125, 727-739.	7.7	48
43	Whole-Exome Sequencing Identifies Mutations of KIF22 in Spondyloepimetaphyseal Dysplasia with Joint Laxity, Leptodactylic Type. American Journal of Human Genetics, 2011, 89, 760-766.	6.2	46
44	Genomic alterations in <i>BCL2L1</i> and <i>DLC1</i> contribute to drug sensitivity in gastric cancer. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 12492-12497.	7.1	46
45	Association between salivary amylase ( $<$ i $><$ scp $>$ AMY $<$ /scp $>$ 1 $<$ /i $>)$ gene copyÂnumbers and insulin resistance in asymptomatic Korean men. Diabetic Medicine, 2015, 32, 1588-1595.	2.3	44
46	Dysregulated Wnt signalling and recurrent mutations of the tumour suppressor <i><scp>RNF43</scp></i> in early gastric carcinogenesis. Journal of Pathology, 2016, 240, 304-314.	<b>4.</b> 5	44
47	RNA-Seq Analysis of Frontal Cortex and Cerebellum from 5XFAD Mice at Early Stage of Disease Pathology. Journal of Alzheimer's Disease, 2012, 29, 793-808.	2.6	43
48	Whole-exome and transcriptome sequencing of refractory diffuse large B-cell lymphoma. Oncotarget, 2016, 7, 86433-86445.	1.8	43
49	Noninvasive Prenatal Diagnosis of Duchenne Muscular Dystrophy: Comprehensive Genetic Diagnosis in Carrier, Proband, and Fetus. Clinical Chemistry, 2015, 61, 829-837.	3.2	42
50	Transcriptome analyses of chronic traumatic encephalopathy show alterations in protein phosphatase expression associated with tauopathy. Experimental and Molecular Medicine, 2017, 49, e333-e333.	7.7	41
51	Recurrent fusion transcripts detected by wholeâ€transcriptome sequencing of 120 primary breast cancer samples. Genes Chromosomes and Cancer, 2015, 54, 681-691.	2.8	38
52	Epidemiologic Characteristics of Intraocular Pressure in the Korean and Mongolian Populations: The Healthy Twin and the GENDISCAN Study. Ophthalmology, 2012, 119, 450-457.	5.2	37
53	High prevalence of TP53 mutations is associated with poor survival and an EMT signature in gliosarcoma patients. Experimental and Molecular Medicine, 2017, 49, e317-e317.	7.7	37
54	Nonsynonymous Variants in <i>PAX4</i> and <i>GLP1R</i> Are Associated With Type 2 Diabetes in an East Asian Population. Diabetes, 2018, 67, 1892-1902.	0.6	36

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55	Rho Is Involved in Superoxide Formation during Phagocytosis of Opsonized Zymosans. Journal of Biological Chemistry, 2004, 279, 21589-21597.	3.4	35
56	A multifunctional protein EWS regulates the expression of Drosha and microRNAs. Cell Death and Differentiation, 2014, 21, 136-145.	11.2	34
57	Findings of a 1303 Korean whole-exome sequencing study. Experimental and Molecular Medicine, 2017, 49, e356-e356.	7.7	34
58	Neuronal loss in primary long-term cortical culture involves neurodegeneration-like cell death via calpain and p35 processing, but not developmental apoptosis or aging. Experimental and Molecular Medicine, 2007, 39, 14-26.	7.7	33
59	Higher mitochondrial DNA copy number is associated with lower prevalence of microalbuminuria. Experimental and Molecular Medicine, 2009, 41, 253.	7.7	33
60	Downstream components of RhoA required for signal pathway of superoxide formation during phagocytosis of serum opsonized zymosans in macrophages. Experimental and Molecular Medicine, 2005, 37, 575-587.	7.7	32
61	HSP90 protects apoptotic cleavage of vimentin in geldanamycin-induced apoptosis. Molecular and Cellular Biochemistry, 2006, 281, 111-121.	3.1	31
62	Clinical whole exome sequencing in early onset diabetes patients. Diabetes Research and Clinical Practice, 2016, 122, 71-77.	2.8	31
63	Clinical application of genomic profiling to find druggable targets for adolescent and young adult (AYA) cancer patients with metastasis. BMC Cancer, 2016, 16, 170.	2.6	30
64	Comprehensive Molecular Characterization of Adenocarcinoma of the Gastroesophageal Junction Between Esophageal and Gastric Adenocarcinomas. Annals of Surgery, 2022, 275, 706-717.	4.2	30
65	Upregulation of neuronal nitric oxide synthase in the periphery promotes pain hypersensitivity after peripheral nerve injury. Neuroscience, 2011, 190, 367-378.	2.3	29
66	A role of placental growth factor in hair growth. Journal of Dermatological Science, 2014, 74, 125-134.	1.9	29
67	Targeted linked-read sequencing for direct haplotype phasing of maternal DMD alleles: a practical and reliable method for noninvasive prenatal diagnosis. Scientific Reports, 2018, 8, 8678.	3.3	29
68	Predominant <i>DICER1</i> Pathogenic Variants in Pediatric Follicular Thyroid Carcinomas. Thyroid, 2020, 30, 1120-1131.	4.5	29
69	NARD: whole-genome reference panel of $1779$ Northeast Asians improves imputation accuracy of rare and low-frequency variants. Genome Medicine, $2019$ , $11$ , $64$ .	8.2	28
70	COX-2 inhibits anoikis by activation of the PI-3K/Akt pathway in human bladder cancer cells. Experimental and Molecular Medicine, 2005, 37, 199-203.	7.7	27
71	I148 <scp>M</scp> variant in <i><scp>PNPLA</scp>3</i> reduces central adiposity and metabolic disease risks while increasing nonalcoholic fatty liver disease. Liver International, 2015, 35, 2537-2546.	3.9	27
72	Identifying Pathogenic Variants of Monogenic Diabetes Using Targeted Panel Sequencing in an East Asian Population. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4188-4198.	3.6	27

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73	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. BMC Genomics, 2020, 21, 322.	2.8	27
74	Predictive biomarkers for 5-fluorouracil and oxaliplatin-based chemotherapy in gastric cancers via profiling of patient-derived xenografts. Nature Communications, 2021, 12, 4840.	12.8	27
75	Diagnostic Yield of Epilepsy Panel Testing in Patients With Seizure Onset Within the First Year of Life. Frontiers in Neurology, 2019, 10, 988.	2.4	26
76	Phagocytosis of serum-and IgG-opsonized zymosan particles induces apoptosis through superoxide but not nitric oxide in macrophage J774A.1. Experimental and Molecular Medicine, 2003, 35, 211-221.	7.7	25
77	A genome-wide Asian genetic map and ethnic comparison: The GENDISCAN study. BMC Genomics, 2008, 9, 554.	2.8	25
78	Somatic deletions implicated in functional diversity of brain cells of individuals with schizophrenia and unaffected controls. Scientific Reports, 2015, 4, 3807.	3.3	25
79	Genetic association of APOA5 and APOE with metabolic syndrome and their interaction with health-related behavior in Korean men. Lipids in Health and Disease, 2015, 14, 105.	3.0	25
80	Priming mobilization of hair follicle stem cells triggers permanent loss of regeneration after alkylating chemotherapy. Nature Communications, 2019, 10, 3694.	12.8	25
81	Altered nucleocytoplasmic proteome and transcriptome distributions in an in vitro model of amyotrophic lateral sclerosis. PLoS ONE, 2017, 12, e0176462.	2.5	24
82	$\mbox{\ensuremath{\mbox{\scriptsize ci}}}\mbox{NTRK1}\mbox{\ensuremath{\mbox{\scriptsize los}}}\mbox{\ensuremath{\mbox{\scriptsize colon}}}\mbox{\ensuremath{\mbox{\scriptsize cancer.}}}\mbox{\ensuremath{\mbox{\scriptsize Oncotarget}}}\mbox{\ensuremath{\mbox{\scriptsize colon}}}\mbox{\ensuremath{\mbox{\scriptsize cancer.}}}\mbox{\ensuremath{\mbox{\scriptsize colon}}}\mbox{\ensuremath{\mbox{\scriptsize cancer.}}}\mbox{\ensuremath{\mbox{\scriptsize cancer.}}}\ensuremath{\mbox{\scriptsize ca$	1.8	24
83	Heritabilities of Facial Measurements and Their Latent Factors in Korean Families. Genomics and Informatics, $2013,11,83.$	0.8	24
84	Transcriptional regulation of Zic3 by heterodimeric AP-1(c-Jun/c-Fos) during Xenopus development. Experimental and Molecular Medicine, 2004, 36, 468-475.	7.7	23
85	Glutaminase 2 expression is associated with regional heterogeneity of 5-aminolevulinic acid fluorescence in glioblastoma. Scientific Reports, 2017, 7, 12221.	3.3	23
86	Exoenzyme Tat-C3 inhibits association of zymosan particles, phagocytosis, adhesion, and complement binding in macrophage cells. Molecules and Cells, 2003, 16, 216-23.	2.6	23
87	Reference-unbiased copy number variant analysis using CGH microarrays. Nucleic Acids Research, 2010, 38, e190-e190.	14.5	22
88	Amplification of transglutaminase 2 enhances tumor-promoting inflammation in gastric cancers. Experimental and Molecular Medicine, 2020, 52, 854-864.	7.7	22
89	Ablation of STAT3 in Purkinje cells reorganizes cerebellar synaptic plasticity in long-term fear memory network. ELife, 2021, 10, .	6.0	22
90	Analysis of Gene Expression in Cyclooxygenase-2-Overexpressed Human Osteosarcoma Cell Lines. Genomics and Informatics, 2014, 12, 247.	0.8	22

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91	Combined linkage and association analyses identify a novel locus for obesity near $\langle i \rangle PROX1 \langle  i \rangle$ in Asians. Obesity, 2013, 21, 2405-2412.	3.0	21
92	RANBP2-ALK fusion combined with monosomy 7 in acute myelomonocytic leukemia. Cancer Genetics, 2014, 207, 40-45.	0.4	21
93	FARS2 mutation and epilepsy: Possible link with early-onset epileptic encephalopathy. Epilepsy Research, 2017, 129, 118-124.	1.6	21
94	Replication of a Glaucoma Candidate Gene on 5q22.1 for Intraocular Pressure in Mongolian Populations: The GENDISCAN Project., 2010, 51, 1335.		20
95	Markers of disease and steroid responsiveness in paediatric idiopathic nephrotic syndrome: Whole-transcriptome sequencing of peripheral blood mononuclear cells. Journal of International Medical Research, 2017, 45, 948-963.	1.0	20
96	Targeted resequencing of candidate genes reveals novel variants associated with severe Behçet's uveitis. Experimental and Molecular Medicine, 2013, 45, e49-e49.	7.7	18
97	Isolation of Middle East Respiratory Syndrome Coronavirus from a Patient of the 2015 Korean Outbreak. Journal of Korean Medical Science, 2016, 31, 315.	2.5	18
98	Collagen Vlâ€related myopathy: Expanding the clinical and genetic spectrum. Muscle and Nerve, 2018, 58, 381-388.	2.2	18
99	Exomic Sequencing of Immune-Related Genes Reveals Novel Candidate Variants Associated with Alopecia Universalis. PLoS ONE, 2013, 8, e53613.	2.5	18
100	A Novel Combination Treatment Targeting BCL-XL and MCL1 for <i>KRAS/BRAF</i> -mutated and <i>BCL2L1</i> -amplified Colorectal Cancers. Molecular Cancer Therapeutics, 2017, 16, 2178-2190.	4.1	17
101	Interactions of CDH13 gene polymorphisms and ambient PM10 air pollution exposure with blood pressure and hypertension in Korean men. Chemosphere, 2019, 218, 292-298.	8.2	17
102	JAK2 regulates paclitaxel resistance in triple negative breast cancers. Journal of Molecular Medicine, 2021, 99, 1783-1795.	3.9	17
103	p21WAF/CIP1/SDI1 is upregulated due to increased mRNA stability during hydroxyurea-induced senescence of human fibroblasts. Mechanisms of Ageing and Development, 2005, 126, 1255-1261.	4.6	16
104	Abdominal adiposity intensifies the negative effects of ambient air pollution on lung function in Korean men. International Journal of Obesity, 2017, 41, 1218-1223.	3.4	16
105	Identification of African-Specific Admixture between Modern and Archaic Humans. American Journal of Human Genetics, 2019, 105, 1254-1261.	6.2	16
106	Phase <scp>II</scp> study of durvalumab and tremelimumab in pulmonary sarcomatoid carcinoma: <scp>KCSG‣U16</scp> â€07. Thoracic Cancer, 2020, 11, 3482-3489.	1.9	16
107	TIARA: a database for accurate analysis of multiple personal genomes based on cross-technology. Nucleic Acids Research, 2011, 39, D883-D888.	14.5	15
108	S100A8/A9 mediate the reprograming of normal mammary epithelial cells induced by dynamic cell–cell interactions with adjacent breast cancer cells. Scientific Reports, 2021, 11, 1337.	3.3	15

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109	Genomic Copy Number Variations Characterize the Prognosis of Both P16-Positive and P16-Negative Oropharyngeal Squamous Cell Carcinoma After Curative Resection. Medicine (United States), 2015, 94, e2187.	1.0	14
110	Clinical Application of Next-Generation Sequencing–Based Panel to ⟨i⟩BRAF⟨/i⟩ Wild-Type Advanced Melanoma Identifies Key Oncogenic Alterations and Therapeutic Strategies. Molecular Cancer Therapeutics, 2020, 19, 937-944.	4.1	14
111	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.	8.0	14
112	Association of HLA Genotype and Fulminant Type 1 Diabetes in Koreans. Genomics and Informatics, 2015, 13, 126.	0.8	14
113	Transcriptional regulation of Xbr-1a/Xvent-2 homeobox gene: analysis of its promoter region. Biochemical and Biophysical Research Communications, 2002, 298, 815-823.	2.1	13
114	Molecular diagnosis of congenital muscular dystrophies with defective glycosylation of alpha-dystroglycan using next-generation sequencing technology. Neuromuscular Disorders, 2013, 23, 337-344.	0.6	13
115	A family-based association study after genome-wide linkage analysis identified two genetic loci for renal function in a Mongolian population. Kidney International, 2013, 83, 285-292.	5.2	13
116	Colorectal cancerâ€susceptibility singleâ€nucleotide polymorphisms in <scp>K</scp> orean population. Journal of Gastroenterology and Hepatology (Australia), 2015, 30, 849-857.	2.8	13
117	High prevalence of TP53 loss and whole-genome doubling in early-onset colorectal cancer. Experimental and Molecular Medicine, 2021, 53, 446-456.	7.7	13
118	Family-Based Association Study of Pulmonary Function in a Population in Northeast Asia. PLoS ONE, 2015, 10, e0139716.	2.5	13
119	Hypoxia-induced cell death of HepG2 cells involves a necrotic cell death mediated by calpain. Apoptosis: an International Journal on Programmed Cell Death, 2007, 12, 707-718.	4.9	12
120	Hoyeraal–Hreidarsson syndrome with a DKC1 mutation identified by whole-exome sequencing. Gene, 2014, 546, 425-429.	2.2	12
121	A Common Variant of NGEF Is Associated with Abdominal Visceral Fat in Korean Men. PLoS ONE, 2015, 10, e0137564.	2.5	12
122	3â€'Oxoacid CoA transferase 1 as a therapeutic target gene for cisplatin-resistant ovarian cancer. Oncology Letters, 2018, 15, 2611-2618.	1.8	12
123	Unstable Genome and Transcriptome Dynamics during Tumor Metastasis Contribute to Therapeutic Heterogeneity in Colorectal Cancers. Clinical Cancer Research, 2019, 25, 2821-2834.	7.0	12
124	A copy number variation in <i>PKD1L2</i> is associated with colorectal cancer predisposition in korean population. International Journal of Cancer, 2017, 140, 86-94.	5.1	11
125	STAT3 is a key molecule in the oncogenic behavior of diffuse intrinsic pontine glioma. Oncology Letters, 2020, 20, 1989-1998.	1.8	11
126	Cross-Talk between Wnt Signaling and Src Tyrosine Kinase. Biomedicines, 2022, 10, 1112.	3.2	11

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127	Analysis of genetic and non-genetic factors that affect the QTc interval in a Mongolian population: the GENDISCAN study. Experimental and Molecular Medicine, 2009, 41, 841.	7.7	10
128	Copy Number Variation of Age-Related Macular Degeneration Relevant Genes in the Korean Population. PLoS ONE, 2012, 7, e31243.	2.5	10
129	The full-length DNA sequence of Epstein Barr virus from a human gastric carcinoma cell line, SNU-719. Virus Genes, 2015, 51, 329-337.	1.6	10
130	CDH13 gene-by-PM10 interaction effect on lung function decline in Korean men. Chemosphere, 2017, 168, 583-589.	8.2	10
131	Genomic characterization of clonal evolution during oropharyngeal carcinogenesis driven by human papillomavirus 16. BMB Reports, 2018, 51, 584-589.	2.4	10
132	Calsequestrin 2 overexpression in breast cancer increases tumorigenesis and metastasis by modulating the tumor microenvironment. Molecular Oncology, 2022, 16, 466-484.	4.6	10
133	CYP1A1 gene polymorphisms modify the association between PM10 exposure and lung function. Chemosphere, 2018, 203, 353-359.	8.2	9
134	Genome-Wide Association Study Reveals Distinct Genetic Susceptibility of Thyroid Nodules From Thyroid Cancer. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 4384-4394.	3.6	9
135	Outbreak investigation of Serratia marcescens neurosurgical site infections associated with a contaminated shaving razors. Antimicrobial Resistance and Infection Control, 2020, 9, 64.	4.1	9
136	The Function of Heterodimeric AP-1 Comprised of c-Jun and c-Fos in Activin Mediated Spemann Organizer Gene Expression. PLoS ONE, 2011, 6, e21796.	2.5	8
137	Gene mapping study for constitutive skin color in an isolated Mongolian population. Experimental and Molecular Medicine, 2012, 44, 241.	7.7	8
138	Vertical sleeve gastrectomy induces distinctive transcriptomic responses in liver, fat and muscle. Scientific Reports, 2021, 11, 2310.	3.3	8
139	Linkage and association scan for tanning ability in an isolated Mongolian population. BMB Reports, 2011, 44, 741-746.	2.4	8
140	Genome-wide combination profiling of copy number and methylation offers an approach for deciphering misregulation and development in cancer cells. Gene, 2008, 407, 139-147.	2,2	7
141	Heritability and linkage study on heart rates in a Mongolian population. Experimental and Molecular Medicine, 2008, 40, 558.	7.7	7
142	Alterations in the Rho pathway contribute to Epstein-Barr virus–induced lymphomagenesis in immunosuppressed environments. Blood, 2018, 131, 1931-1941.	1.4	7
143	Comparative genomics of Mycoplasma pneumoniae isolated from children with pneumonia: South Korea, 2010–2016. BMC Genomics, 2019, 20, 910.	2.8	7
144	xCyp26cInduced by Inhibition of BMP Signaling Is Involved in Anterior-Posterior Neural Patterning of Xenopus laevis. Molecules and Cells, 2016, 39, 352-357.	2.6	7

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145	Early onset female pattern hair loss: A case–control study for analyzing clinical features and genetic variants. Journal of Dermatological Science, 2022, 106, 21-28.	1.9	7
146	Phosphorylation of 46-kDa protein of synaptic vesicle membranes is stimulated by GTP and Ca2+/calmodulin. Experimental and Molecular Medicine, 2002, 34, 434-443.	7.7	6
147	Genome-wide linkage analysis for ocular and nasal anthropometric traits in a Mongolian population. Experimental and Molecular Medicine, 2010, 42, 799.	7.7	6
148	TIARA genome database: update 2013. Database: the Journal of Biological Databases and Curation, 2013, 2013, bat003-bat003.	3.0	6
149	Sequencing Cell-free Fetal DNA in Pregnant Women With <i>GCK</i> -MODY. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 2678-2689.	<b>3.</b> 6	6
150	Thyroid nodules in childhoodâ€onset Hashimoto's thyroiditis: Frequency, risk factors, followâ€up course and genetic alterations of thyroid cancer. Clinical Endocrinology, 2021, 95, 638-648.	2.4	6
151	Nerve growth factor induces proliferation of PC12 cells through Cdc42. NeuroReport, 2003, 14, 1277-1281.	1.2	5
152	DNAJB9 Inhibits p53-Dependent Oncogene-Induced Senescence and Induces Cell Transformation. Molecules and Cells, 2020, 43, 397-407.	2.6	5
153	Draft Genome of Toxocara canis, a Pathogen Responsible for Visceral Larva Migrans. Korean Journal of Parasitology, 2016, 54, 751-758.	1.3	5
154	Protein Phosphatase 1H, Cyclin-Dependent Kinase Inhibitor p27, and Cyclin-Dependent Kinase 2 in Paclitaxel Resistance for Triple Negative Breast Cancers. Journal of Breast Cancer, 2020, 23, 162.	1.9	5
155	The first Irish genome and ways of improving sequence accuracy. Genome Biology, 2010, 11, 132.	9.6	4
156	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. BMC Medical Genomics, 2016, 9, 33.	1.5	4
157	Genetic variations associated with response to dutasteride in the treatment of male subjects with androgenetic alopecia. PLoS ONE, 2019, 14, e0222533.	2.5	4
158	Discovery of acquired molecular signature on immune checkpoint inhibitors in paired tumor tissues. Cancer Immunology, Immunotherapy, 2021, 70, 1755-1769.	4.2	4
159	Radiation Response Prediction Model Based on Integrated Clinical and Genomic Data Analysis. Cancer Research and Treatment, 2022, 54, 383-395.	3.0	4
160	Targeted Next-Generation Sequencing at Copy-Number Breakpoints for Personalized Analysis of Rearranged Ends in Solid Tumors. PLoS ONE, 2014, 9, e100089.	2.5	4
161	Endoscopic treatment of interstitial pregnancy. Acta Obstetricia Et Gynecologica Scandinavica, 2003, 82, 189-191.	2.8	3
162	Development of a common platform for the noninvasive prenatal diagnosis of Xâ€linked diseases. Prenatal Diagnosis, 2018, 38, 835-840.	2.3	3

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163	A glioneuronal tumor with CLIP2-MET fusion. Npj Genomic Medicine, 2020, 5, 24.	3.8	3
164	Image Correlation-Based Method to Assess Ciliary Beat Frequency in Human Airway Organoids. IEEE Transactions on Medical Imaging, 2022, 41, 374-382.	8.9	3
165	Genomic Landscape of Young-Onset Bladder Cancer and Its Prognostic Implications on Adult Bladder Cancer. Cancers, 2020, 12, 307.	3.7	3
166	Treatment strategy for papillary renal cell carcinoma type 2: a case series of seven patients treated based on next generation sequencing data. Annals of Translational Medicine, 2020, 8, 1389-1389.	1.7	3
167	xCITED2 Induces Neural Genes in Animal Cap Explants of Xenopus Embryos. Experimental Neurobiology, 2011, 20, 123-129.	1.6	2
168	Dissecting the phenotypic and genetic spectrum of early childhood-onset generalized epilepsies. Seizure: the Journal of the British Epilepsy Association, 2019, 71, 222-228.	2.0	2
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