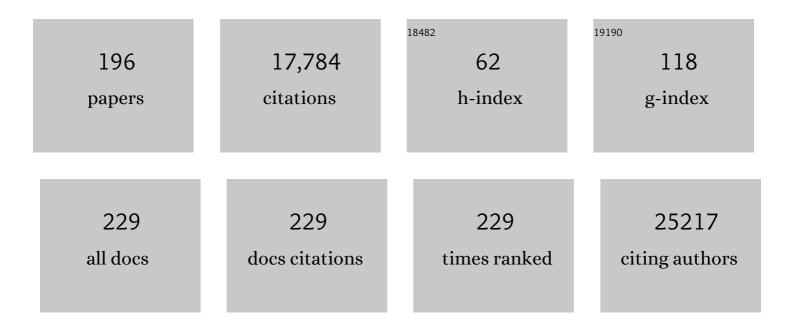
## Koichi K Matsuda

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genome-Wide Association Study of Intracranial Artery Stenosis Followed by Phenome-Wide Association Study. Translational Stroke Research, 2023, 14, 322-333.	4.2	5
2	Population-based Screening for Hereditary Colorectal Cancer Variants in Japan. Clinical Gastroenterology and Hepatology, 2022, 20, 2132-2141.e9.	4.4	20
3	Trans-ethnic Mendelian-randomization study reveals causal relationships between cardiometabolic factors and chronic kidney disease. International Journal of Epidemiology, 2022, 50, 1995-2010.	1.9	39
4	Genome-wide association study of colorectal polyps identified highly overlapping polygenic architecture with colorectal cancer. Journal of Human Genetics, 2022, 67, 149-156.	2.3	5
5	Novel susceptibility loci for steroid-associated osteonecrosis of the femoral head in systemic lupus erythematosus. Human Molecular Genetics, 2022, 31, 1082-1095.	2.9	1
6	Genetic overlap analysis of endometriosis and asthma identifies shared loci implicating sex hormones and thyroid signalling pathways. Human Reproduction, 2022, 37, 366-383.	0.9	19
7	A single-nucleotide-polymorphism in the 5′-flanking region of <i>MSX1</i> gene as a predictive marker candidate for platinum-based therapy of esophageal carcinoma. Therapeutic Advances in Medical Oncology, 2022, 14, 175883592210805.	3.2	0
8	Decoding the diversity of killer immunoglobulin-like receptors by deep sequencing and a high-resolution imputation method. Cell Genomics, 2022, 2, 100101.	6.5	6
9	Large-scale Integrated Analysis of Genetics and Metabolomic Data Reveals Potential Links Between Lipids and Colorectal Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2022, 31, 1216-1226.	2.5	3
10	Leveraging fine-mapping and multipopulation training data to improve cross-population polygenic risk scores. Nature Genetics, 2022, 54, 450-458.	21.4	109
11	Genome-wide risk prediction of common diseases across ancestries in one million people. Cell Genomics, 2022, 2, 100118.	6.5	34
12	Expansion of Cancer Risk Profile for <i>BRCA1</i> and <i>BRCA2</i> Pathogenic Variants. JAMA Oncology, 2022, 8, 871.	7.1	70
13	Genetic analysis of right heart structure and function in 40,000 people. Nature Genetics, 2022, 54, 792-803.	21.4	34
14	Biological insights into systemic lupus erythematosus through an immune cell-specific transcriptome-wide association study. Annals of the Rheumatic Diseases, 2022, 81, 1273-1280.	0.9	9
15	Meta-analysis of 208370 East Asians identifies 113 susceptibility loci for systemic lupus erythematosus. Annals of the Rheumatic Diseases, 2021, 80, 632-640.	0.9	103
16	Genetic analysis of endometriosis and depression identifies shared loci and implicates causal links with gastric mucosa abnormality. Human Genetics, 2021, 140, 529-552.	3.8	36
17	Whole genome sequence analyses of eGFR in 23,732 people representing multiple ancestries in the NHLBI trans-omics for precision medicine (TOPMed) consortium. EBioMedicine, 2021, 63, 103157.	6.1	14
18	Trans-ancestry genome-wide association meta-analysis of prostate cancer identifies new susceptibility loci and informs genetic risk prediction. Nature Genetics, 2021, 53, 65-75.	21.4	264

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19	The simplified Kyoto classification score is consistent with the ABC method of classification as a grading system for endoscopic gastritis. Journal of Clinical Biochemistry and Nutrition, 2021, 68, 101-104.	1.4	10
20	A deep learning method for HLA imputation and trans-ethnic MHC fine-mapping of type 1 diabetes. Nature Communications, 2021, 12, 1639.	12.8	44
21	Comprehensive molecular and clinicopathological profiling of desmoid tumours. European Journal of Cancer, 2021, 145, 109-120.	2.8	10
22	Leveraging supervised learning for functionallyÂinformed fine-mapping of cis-eQTLs identifies an additional 20,913 putative causal eQTLs. Nature Communications, 2021, 12, 3394.	12.8	44
23	Hematopoietic mosaic chromosomal alterations increase the risk for diverse types of infection. Nature Medicine, 2021, 27, 1012-1024.	30.7	109
24	Combined landscape of single-nucleotide variants and copy number alterations in clonal hematopoiesis. Nature Medicine, 2021, 27, 1239-1249.	30.7	78
25	Collagen XVII inhibits breast cancer cell proliferation and growth through deactivation of the AKT/mTOR signaling pathway. PLoS ONE, 2021, 16, e0255179.	2.5	16
26	Genetic analyses of gynecological disease identify genetic relationships between uterine fibroids and endometrial cancer, and a novel endometrial cancer genetic risk region at the WNT4 1p36.12 locus. Human Genetics, 2021, 140, 1353-1365.	3.8	18
27	Intracellular Accumulation of IFN-λ4 Induces ER Stress and Results in Anti-Cirrhotic but Pro-HCV Effects. Frontiers in Immunology, 2021, 12, 692263.	4.8	6
28	A cross-population atlas of genetic associations for 220 human phenotypes. Nature Genetics, 2021, 53, 1415-1424.	21.4	560
29	A genome-wide association study identifies a novel candidate locus at the DLGAP1 gene with susceptibility to resistant hypertension in the Japanese population. Scientific Reports, 2021, 11, 19497.	3.3	12
30	A single nucleotide polymorphism in <i>Prostate Stem Cell Antigen</i> is associated with endoscopic grading in Kyoto classification of gastritis. Journal of Clinical Biochemistry and Nutrition, 2021, 68, 73-77.	1.4	3
31	Genetic susceptibility to hepatocellular carcinoma in chromosome 22q13.31, findings of a genomeâ€wide association study. JGH Open, 2021, 5, 1363-1372.	1.6	9
32	Germline Pathogenic Variants in 7636 Japanese Patients With Prostate Cancer and 12Â366 Controls. Journal of the National Cancer Institute, 2020, 112, 369-376.	6.3	69
33	Functional variants in ADH1B and ALDH2 are non-additively associated with all-cause mortality in Japanese population. European Journal of Human Genetics, 2020, 28, 378-382.	2.8	14
34	GWAS of five gynecologic diseases and cross-trait analysis in Japanese. European Journal of Human Genetics, 2020, 28, 95-107.	2.8	32
35	Identification of Novel Loci and New Risk Variant in Known Loci for Colorectal Cancer Risk in East Asians. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 477-486.	2.5	25
36	Genetic characterization of pancreatic cancer patients and prediction of carrier status of germline pathogenic variants in cancer-predisposing genes. EBioMedicine, 2020, 60, 103033.	6.1	39

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37	The Polygenic and Monogenic Basis of Blood Traits and Diseases. Cell, 2020, 182, 1214-1231.e11.	28.9	388
38	Population-specific and trans-ancestry genome-wide analyses identify distinct and shared genetic risk loci for coronary artery disease. Nature Genetics, 2020, 52, 1169-1177.	21.4	206
39	Genome-wide association study of intracranial aneurysms identifies 17 risk loci and genetic overlap with clinical risk factors. Nature Genetics, 2020, 52, 1303-1313.	21.4	163
40	Improving the trans-ancestry portability of polygenic risk scores by prioritizing variants in predicted cell-type-specific regulatory elements. Nature Genetics, 2020, 52, 1346-1354.	21.4	126
41	Endogenization and excision of human herpesvirus 6 in human genomes. PLoS Genetics, 2020, 16, e1008915.	3.5	22
42	Trans-ethnic and Ancestry-Specific Blood-Cell Genetics in 746,667 Individuals from 5 Global Populations. Cell, 2020, 182, 1198-1213.e14.	28.9	353
43	Association of circulating 25-Hydroxyvitamin D and its related genetic variations with hepatocellular carcinoma incidence and survival. Annals of Translational Medicine, 2020, 8, 1080-1080.	1.7	8
44	Fine Mapping of the Major Histocompatibility Complex Region and Association of the HLA-B*52:01 Allele With Cervical Cancer in Japanese Women. JAMA Network Open, 2020, 3, e2023248.	5.9	7
45	Transethnic Meta-Analysis of Genome-Wide Association Studies Identifies Three New Loci and Characterizes Population-Specific Differences for Coronary Artery Disease. Circulation Genomic and Precision Medicine, 2020, 13, e002670.	3.6	44
46	Chromosomal alterations among age-related haematopoietic clones in Japan. Nature, 2020, 584, 130-135.	27.8	102
47	Large-scale genome-wide association study in a Japanese population identifies novel susceptibility loci across different diseases. Nature Genetics, 2020, 52, 669-679.	21.4	304
48	Identification of novel breast cancer susceptibility loci in meta-analyses conducted among Asian and European descendants. Nature Communications, 2020, 11, 1217.	12.8	46
49	Trans-biobank analysis with 676,000 individuals elucidates the association of polygenic risk scores of complex traits with human lifespan. Nature Medicine, 2020, 26, 542-548.	30.7	74
50	Genetic and phenotypic landscape of the mitochondrial genome in the Japanese population. Communications Biology, 2020, 3, 104.	4.4	32
51	Genome-wide association meta-analysis identifies GP2 gene risk variants for pancreatic cancer. Nature Communications, 2020, 11, 3175.	12.8	34
52	GWAS of 165,084 Japanese individuals identified nine loci associated with dietary habits. Nature Human Behaviour, 2020, 4, 308-316.	12.0	80
53	Identification of a novel uterine leiomyoma GWAS locus in a Japanese population. Scientific Reports, 2020, 10, 1197.	3.3	14
54	Dimensionality reduction reveals fine-scale structure in the Japanese population with consequences for polygenic risk prediction. Nature Communications, 2020, 11, 1569.	12.8	58

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55	Prevalence and Spectrum of Pathogenic Germline Variants in Japanese Patients With Early-Onset Colorectal, Breast, and Prostate Cancer. JCO Precision Oncology, 2020, 4, 183-191.	3.0	6
56	Genome-Wide Natural Selection Signatures Are Linked to Genetic Risk of Modern Phenotypes in the Japanese Population. Molecular Biology and Evolution, 2020, 37, 1306-1316.	8.9	22
57	Claudin-2 deficiency associates with hypercalciuria in mice and human kidney stone disease. Journal of Clinical Investigation, 2020, 130, 1948-1960.	8.2	61
58	Endoscopy-based Kyoto classification score of gastritis related to pathological topography of neutrophil activity. World Journal of Gastroenterology, 2020, 26, 5146-5155.	3.3	11
59	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. Journal of the National Cancer Institute, 2019, 111, 146-157.	6.3	129
60	GWAS of mosaic loss of chromosome Y highlights genetic effects on blood cell differentiation. Nature Communications, 2019, 10, 4719.	12.8	50
61	Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.	12.8	84
62	Antitumor immunity augments the therapeutic effects of p53 activation on acute myeloid leukemia. Nature Communications, 2019, 10, 4869.	12.8	36
63	12 new susceptibility loci for prostate cancer identified by genome-wide association study in Japanese population. Nature Communications, 2019, 10, 4422.	12.8	49
64	Characterizing rare and low-frequency height-associated variants in the Japanese population. Nature Communications, 2019, 10, 4393.	12.8	123
65	Target genes, variants, tissues and transcriptional pathways influencing human serum urate levels. Nature Genetics, 2019, 51, 1459-1474.	21.4	251
66	Genetic and phenotypic landscape of the major histocompatibilty complex region in the Japanese population. Nature Genetics, 2019, 51, 470-480.	21.4	75
67	Identification of rare coding variants in <i>TYK2</i> protective for rheumatoid arthritis in the Japanese population and their effects on cytokine signalling. Annals of the Rheumatic Diseases, 2019, 78, 1062-1069.	0.9	16
68	A catalog of genetic loci associated with kidney function from analyses of a million individuals. Nature Genetics, 2019, 51, 957-972.	21.4	549
69	Massively parallel sequencing of tenosynovial giant cell tumors reveals novel CSF1 fusion transcripts and novel somatic CBL mutations. International Journal of Cancer, 2019, 145, 3276-3284.	5.1	28
70	INKA2, a novel p53 target that interacts with the serine/threonine kinase PAK4. International Journal of Oncology, 2019, 54, 1907-1920.	3.3	8
71	Novel Risk Loci Identified in a Genome-Wide Association Study of Urolithiasis in a Japanese Population. Journal of the American Society of Nephrology: JASN, 2019, 30, 855-864.	6.1	25
72	GWAS of smoking behaviour in 165,436 Japanese people reveals seven new loci and shared genetic architecture. Nature Human Behaviour, 2019, 3, 471-477.	12.0	54

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73	A Polymorphic Variant in p19Arf Confers Resistance to Chemically Induced Skin Tumors by Activating the p53 Pathway. Journal of Investigative Dermatology, 2019, 139, 1459-1469.	0.7	4
74	Genome-wide meta-analysis identifies multiple novel loci associated with serum uric acid levels in Japanese individuals. Communications Biology, 2019, 2, 115.	4.4	66
75	A multi-ancestry genome-wide study incorporating gene–smoking interactions identifies multiple new loci for pulse pressure and mean arterial pressure. Human Molecular Genetics, 2019, 28, 2615-2633.	2.9	31
76	Large-Scale Genome-Wide Association Study of East Asians Identifies Loci Associated With Risk for Colorectal Cancer. Gastroenterology, 2019, 156, 1455-1466.	1.3	111
77	Genetic variants of calcium and vitamin D metabolism in kidney stone disease. Nature Communications, 2019, 10, 5175.	12.8	69
78	Integrated exome and RNA sequencing of dedifferentiated liposarcoma. Nature Communications, 2019, 10, 5683.	12.8	41
79	Identification of two novel breast cancer loci through large-scale genome-wide association study in the Japanese population. Scientific Reports, 2019, 9, 17332.	3.3	9
80	Clinical and molecular characteristics of <i>MEF2D</i> fusion-positive B-cell precursor acute lymphoblastic leukemia in childhood, including a novel translocation resulting in <i>MEF2D-HNRNPH1</i> gene fusion. Haematologica, 2019, 104, 128-137.	3.5	65
81	Trans-ethnic kidney function association study reveals putative causal genes and effects on kidney-specific disease aetiologies. Nature Communications, 2019, 10, 29.	12.8	113
82	Single Nucleotide Polymorphisms of <i>HAAO</i> and <i>IRX6</i> Genes as Risk Factors for Hypospadias. Journal of Urology, 2019, 201, 386-392.	0.4	9
83	Identification of 28 new susceptibility loci for type 2 diabetes in the Japanese population. Nature Genetics, 2019, 51, 379-386.	21.4	164
84	The Association of Single Nucleotide Polymorphisms with Cancer Risk. , 2019, , 87-144.		0
85	GWAS identifies two novel colorectal cancer loci at 16q24.1 and 20q13.12. Carcinogenesis, 2018, 39, 652-660.	2.8	52
86	Deep whole-genome sequencing reveals recent selection signatures linked to evolution and disease risk of Japanese. Nature Communications, 2018, 9, 1631.	12.8	132
87	Genetic analysis of quantitative traits in the Japanese population links cell types to complex human diseases. Nature Genetics, 2018, 50, 390-400.	21.4	613
88	Citrullination of RGG Motifs in FET Proteins by PAD4 Regulates Protein Aggregation and ALS Susceptibility. Cell Reports, 2018, 22, 1473-1483.	6.4	85
89	Decrease in <i>PSCA</i> expression caused by <i>Helicobacter pylori</i> infection may promote progression to severe gastritis. Oncotarget, 2018, 9, 3936-3945.	1.8	21
90	Genome-wide association study (GWAS) of ovarian cancer in Japanese predicted regulatory variants in 22q13.1. PLoS ONE, 2018, 13, e0209096.	2.5	8

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91	Genomeâ€wide association study identifies gastric cancer susceptibility loci at 12q24.11â€12 and 20q11.21. Cancer Science, 2018, 109, 4015-4024.	3.9	39
92	Interethnic analyses of blood pressure loci in populations of East Asian and European descent. Nature Communications, 2018, 9, 5052.	12.8	75
93	Germline pathogenic variants of 11 breast cancer genes in 7,051 Japanese patients and 11,241 controls. Nature Communications, 2018, 9, 4083.	12.8	179
94	Elucidating the genetic architecture of reproductive ageing in the Japanese population. Nature Communications, 2018, 9, 1977.	12.8	44
95	The p53 activator overcomes resistance to ALK inhibitors by regulating p53-target selectivity in ALK-driven neuroblastomas. Cell Death Discovery, 2018, 4, 56.	4.7	23
96	Risk prediction models for mortality in patients with cardiovascular disease: The BioBank Japan project. Journal of Epidemiology, 2017, 27, S71-S76.	2.4	11
97	EPSIN 3, A Novel p53 Target, Regulates the Apoptotic Pathway and Gastric Carcinogenesis. Neoplasia, 2017, 19, 185-195.	5.3	14
98	GALNT6 Stabilizes GRP78 Protein by O-glycosylation and Enhances its Activity to Suppress Apoptosis Under Stress Condition. Neoplasia, 2017, 19, 43-53.	5.3	23
99	Characteristics and prognosis of Japanese colorectal cancer patients: The BioBank Japan Project. Journal of Epidemiology, 2017, 27, S36-S42.	2.4	38
100	Characteristics of patients with liver cancer in the BioBank Japan project. Journal of Epidemiology, 2017, 27, S43-S48.	2.4	17
101	Survival of macrovascular disease, chronic kidney disease, chronic respiratory disease, cancer and smoking in patients with type 2 diabetes: BioBank Japan cohort. Journal of Epidemiology, 2017, 27, S98-S106.	2.4	20
102	Statin use and all-cause and cancer mortality: BioBank Japan cohort. Journal of Epidemiology, 2017, 27, S84-S91.	2.4	25
103	Characteristics and prognosis of Japanese female breast cancer patients: The BioBank Japan project. Journal of Epidemiology, 2017, 27, S58-S64.	2.4	27
104	Demographic and lifestyle factors and survival among patients with esophageal and gastric cancer: The Biobank Japan Project. Journal of Epidemiology, 2017, 27, S29-S35.	2.4	32
105	Cross-sectional analysis of BioBank Japan clinical data: A large cohort of 200,000 patients with 47 common diseases. Journal of Epidemiology, 2017, 27, S9-S21.	2.4	133
106	Argininosuccinate synthase 1 is an intrinsic Akt repressor transactivated by p53. Science Advances, 2017, 3, e1603204.	10.3	40
107	The Transcriptional Landscape of p53 Signalling Pathway. EBioMedicine, 2017, 20, 109-119.	6.1	47
108	Meta-analysis identifies five novel loci associated with endometriosis highlighting key genes involved in hormone metabolism. Nature Communications, 2017, 8, 15539.	12.8	230

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109	Overview of the BioBank Japan Project: Study design and profile. Journal of Epidemiology, 2017, 27, S2-S8.	2.4	451
110	Overview of BioBank Japan follow-up data in 32 diseases. Journal of Epidemiology, 2017, 27, S22-S28.	2.4	47
111	Cholesterol levels of Japanese dyslipidaemic patients with various comorbidities: BioBank Japan. Journal of Epidemiology, 2017, 27, S77-S83.	2.4	3
112	Clinical and histopathological characteristics of patients with prostate cancer in the BioBank Japan project. Journal of Epidemiology, 2017, 27, S65-S70.	2.4	11
113	Characteristics and prognosis of Japanese male and female lung cancer patients: The BioBank Japan Project. Journal of Epidemiology, 2017, 27, S49-S57.	2.4	17
114	Genome-wide Trans-ethnic Meta-analysis Identifies Seven Genetic Loci Influencing Erythrocyte Traits and a Role for RBPMS in Erythropoiesis. American Journal of Human Genetics, 2017, 100, 51-63.	6.2	45
115	Identification of a p53 target, CD137L, that mediates growth suppression and immune response of osteosarcoma cells. Scientific Reports, 2017, 7, 10739.	3.3	3
116	Genome-wide association study identifies 112 new loci for body mass index in the Japanese population. Nature Genetics, 2017, 49, 1458-1467.	21.4	380
117	Serum glucose, cholesterol and blood pressure levels in Japanese type 1 and 2 diabetic patients: BioBank Japan. Journal of Epidemiology, 2017, 27, S92-S97.	2.4	12
118	Regulation of tubular recycling endosome biogenesis by the p53-MICALL1 pathway. International Journal of Oncology, 2017, 51, 724-736.	3.3	6
119	Identification of a novel p53 target, COL17A1, that inhibits breast cancer cell migration and invasion. Oncotarget, 2017, 8, 55790-55803.	1.8	58
120	Identification of a p53-repressed gene module in breast cancer cells. Oncotarget, 2017, 8, 55821-55836.	1.8	6
121	Adjustment of Cell-Type Composition Minimizes Systematic Bias in Blood DNA Methylation Profiles Derived by DNA Collection Protocols. PLoS ONE, 2016, 11, e0147519.	2.5	21
122	Association Study of a Functional Variant on ABCG2 Gene with Sunitinib-Induced Severe Adverse Drug Reaction. PLoS ONE, 2016, 11, e0148177.	2.5	39
123	Trans-ethnic Fine Mapping Highlights Kidney-Function Genes Linked to Salt Sensitivity. American Journal of Human Genetics, 2016, 99, 636-646.	6.2	67
124	Genetic risk score based on the prevalence of vertebral fracture in Japanese women with osteoporosis. Bone Reports, 2016, 5, 168-172.	0.4	15
125	Contribution of a Non-classical HLA Gene, HLA-DOA, to the Risk of Rheumatoid Arthritis. American Journal of Human Genetics, 2016, 99, 366-374.	6.2	68
126	Cystatin C as a p53â€inducible apoptotic mediator that regulates cathepsin L activity. Cancer Science, 2016, 107, 298-306.	3.9	38

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127	Regulation of myo-inositol biosynthesis by p53-ISYNA1 pathway. International Journal of Oncology, 2016, 48, 2415-2424.	3.3	41
128	Genome-wide association study in East Asians identifies two novel breast cancer susceptibility loci. Human Molecular Genetics, 2016, 25, 3361-3371.	2.9	40
129	Identification of Susceptibility Loci and Genes for Colorectal Cancer Risk. Gastroenterology, 2016, 150, 1633-1645.	1.3	97
130	Regulation of iron homeostasis by the p53-ISCU pathway. Scientific Reports, 2015, 5, 16497.	3.3	68
131	Influence of Genetic Variants in EGF and Other Genes on Hematological Traits in Korean Populations by a Genome-Wide Approach. BioMed Research International, 2015, 2015, 1-9.	1.9	6
132	Construction of a population-specific HLA imputation reference panel and its application to Graves' disease risk in Japanese. Nature Genetics, 2015, 47, 798-802.	21.4	119
133	Identification of novel epigenetically inactivated gene PAMR1 in breast carcinoma. Oncology Reports, 2015, 33, 267-273.	2.6	31
134	A genome-wide association study identifies PLCL2 and AP3D1-DOT1L-SF3A2 as new susceptibility loci for myocardial infarction in Japanese. European Journal of Human Genetics, 2015, 23, 374-380.	2.8	48
135	Directional dominance on stature and cognition inÂdiverse human populations. Nature, 2015, 523, 459-462.	27.8	173
136	Large-scale association analysis in Asians identifies new susceptibility loci for prostate cancer. Nature Communications, 2015, 6, 8469.	12.8	51
137	Genome-wide association study identified SNP on 15q24 associated with bladder cancer risk in Japanese population. Human Molecular Genetics, 2015, 24, 1177-1184.	2.9	38
138	Antitumor Activity and Induction of TP53-Dependent Apoptosis toward Ovarian Clear Cell Adenocarcinoma by the Dual PI3K/mTOR Inhibitor DS-7423. PLoS ONE, 2014, 9, e87220.	2.5	40
139	Quantitative T cell repertoire analysis by deep cDNA sequencing of T cell receptor α and β chains using next-generation sequencing (NCS). Oncolmmunology, 2014, 3, e968467.	4.6	68
140	Late Cornified Envelope Group I, a Novel Target of p53, Regulates PRMT5 Activity. Neoplasia, 2014, 16, 656-664.	5.3	16
141	Downregulation of the tumor suppressor HSPB7, involved in the p53 pathway, in renal cell carcinoma by hypermethylation. International Journal of Oncology, 2014, 44, 1490-1498.	3.3	25
142	A Rare Polymorphic Variant of <i>NBS1</i> Reduces DNA Repair Activity and Elevates Chromosomal Instability. Cancer Research, 2014, 74, 3707-3715.	0.9	5
143	Genomeâ€wide association study identifies a new SMAD7 risk variant associated with colorectal cancer risk in East Asians. International Journal of Cancer, 2014, 135, 948-955.	5.1	52
144	Large-scale genetic study in East Asians identifies six new loci associated with colorectal cancer risk. Nature Genetics, 2014, 46, 533-542.	21.4	212

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145	Genome-wide association analysis in East Asians identifies breast cancer susceptibility loci at 1q32.1, 5q14.3 and 15q26.1. Nature Genetics, 2014, 46, 886-890.	21.4	135
146	Trans-ethnic meta-analysis of white blood cell phenotypes. Human Molecular Genetics, 2014, 23, 6944-6960.	2.9	60
147	Identification of a nuclear protein, LRRC42, involved in lung carcinogenesis. International Journal of Oncology, 2014, 45, 147-156.	3.3	7
148	A replication study for three nephrolithiasis loci at 5q35.3, 7p14.3 and 13q14.1 in the Japanese population. Journal of Human Genetics, 2013, 58, 588-593.	2.3	24
149	A genome-wide association study of HCV-induced liver cirrhosis in the Japanese population identifies novel susceptibility loci at the MHC region. Journal of Hepatology, 2013, 58, 875-882.	3.7	58
150	Impact of polymorphisms in drug pathway genes on disease-free survival in adults with acute myeloid leukemia. Journal of Human Genetics, 2013, 58, 353-361.	2.3	38
151	The histone methyltransferase Wolf–Hirschhorn syndrome candidate 1â€like 1 (WHSC1L1) is involved in human carcinogenesis. Genes Chromosomes and Cancer, 2013, 52, 126-139.	2.8	64
152	Meta-Analysis of Genome-Wide Association Studies Identifies Six New Loci for Serum Calcium Concentrations. PLoS Genetics, 2013, 9, e1003796.	3.5	142
153	Germline Variants and Advanced Colorectal Adenomas: Adenoma Prevention with Celecoxib Trial Genome-wide Association Study. Clinical Cancer Research, 2013, 19, 6430-6437.	7.0	9
154	Impact of four loci on serum tamsulosin hydrochloride concentration. Journal of Human Genetics, 2013, 58, 21-26.	2.3	2
155	Impact of PSCA Variation on Gastric Ulcer Susceptibility. PLoS ONE, 2013, 8, e63698.	2.5	15
156	Identification of a Functional Variant in the MICA Promoter Which Regulates MICA Expression and Increases HCV-Related Hepatocellular Carcinoma Risk. PLoS ONE, 2013, 8, e61279.	2.5	59
157	Genome Wide Association Study of Age at Menarche in the Japanese Population. PLoS ONE, 2013, 8, e63821.	2.5	26
158	A Genome-Wide Association Study Identified AFF1 as a Susceptibility Locus for Systemic Lupus Eyrthematosus in Japanese. PLoS Genetics, 2012, 8, e1002455.	3.5	115
159	A Genome-Wide Association Study of Nephrolithiasis in the Japanese Population Identifies Novel Susceptible Loci at 5q35.3, 7p14.3, and 13q14.1. PLoS Genetics, 2012, 8, e1002541.	3.5	63
160	Critical Function for Nuclear Envelope Protein TMEM209 in Human Pulmonary Carcinogenesis. Cancer Research, 2012, 72, 4110-4118.	0.9	27
161	Common variation near CDKN1A, POLD3 and SHROOM2 influences colorectal cancer risk. Nature Genetics, 2012, 44, 770-776.	21.4	210
162	A genome-wide association study identifies two susceptibility loci for duodenal ulcer in the Japanese population. Nature Genetics, 2012, 44, 430-434.	21.4	114

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163	No association for Chinese HBV-related hepatocellular carcinoma susceptibility SNP in other East Asian populations. BMC Medical Genetics, 2012, 13, 47.	2.1	40
164	Quantitative Structural Characterization of Local N-Glycan Microheterogeneity in Therapeutic Antibodies by Energy-Resolved Oxonium Ion Monitoring. Analytical Chemistry, 2012, 84, 9655-9662.	6.5	49
165	CLCA2 as a p53-Inducible Senescence Mediator. Neoplasia, 2012, 14, 141-IN9.	5.3	32
166	Association of Common Variants in TNFRSF13B, TNFSF13, and ANXA3 with Serum Levels of Non-Albumin Protein and Immunoglobulin Isotypes in Japanese. PLoS ONE, 2012, 7, e32683.	2.5	34
167	Soluble MICA and a MICA Variation as Possible Prognostic Biomarkers for HBV-Induced Hepatocellular Carcinoma. PLoS ONE, 2012, 7, e44743.	2.5	81
168	Regulation of histone modification and chromatin structure by the p53–PADI4 pathway. Nature Communications, 2012, 3, 676.	12.8	128
169	Novel susceptibility loci for hepatocellular carcinoma in chronic HBV carriers. Hepatobiliary Surgery and Nutrition, 2012, 1, 59-60.	1.5	7
170	Germline Genetic Polymorphisms Are Associated with Disease-Free Survival in Adults with Acute Myeloid Leukemia (AML): A Genomewide Association Study From the Pgrn-Riken Global Alliance Blood, 2012, 120, 2548-2548.	1.4	0
171	Genome-wide association study for C-reactive protein levels identified pleiotropic associations in the IL6 locus. Human Molecular Genetics, 2011, 20, 1224-1231.	2.9	82
172	Genome-wide association study identifies loci influencing concentrations of liver enzymes in plasma. Nature Genetics, 2011, 43, 1131-1138.	21.4	501
173	Large-scale genome-wide association studies in east Asians identify new genetic loci influencing metabolic traits. Nature Genetics, 2011, 43, 990-995.	21.4	270
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