Xiaoping Miao

List of Publications by Year in descending order

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286 20,413 64 papers citations h-index

g-index
35393

293
all docs doc

293 docs citations 293 times ranked

35393 citing authors

129

#	Article	IF	Citations
1	Neurologic Manifestations of Hospitalized Patients With Coronavirus Disease 2019 in Wuhan, China. JAMA Neurology, 2020, 77, 683.	4.5	5,308
2	Hirschsprung disease, associated syndromes and genetics: a review. Journal of Medical Genetics, 2007, 45, 1-14.	1.5	848
3	Clinical characteristics and risk factors associated with COVID-19 disease severity in patients with cancer in Wuhan, China: a multicentre, retrospective, cohort study. Lancet Oncology, The, 2020, 21, 893-903.	5.1	421
4	Skin damage among health care workers managing coronavirus disease-2019. Journal of the American Academy of Dermatology, 2020, 82, 1215-1216.	0.6	399
5	Genome-wide association study identifies three new susceptibility loci for esophageal squamous-cell carcinoma in Chinese populations. Nature Genetics, 2011, 43, 679-684.	9.4	260
6	A genome-wide association study identifies new susceptibility loci for non-cardia gastric cancer at 3q13.31 and 5p13.1. Nature Genetics, 2011, 43, 1215-1218.	9.4	250
7	Genome-wide association analyses of esophageal squamous cell carcinoma in Chinese identify multiple susceptibility loci and gene-environment interactions. Nature Genetics, 2012, 44, 1090-1097.	9.4	238
8	Pancreatic cancer risk variant in LINC00673 creates a miR-1231 binding site and interferes with PTPN11 degradation. Nature Genetics, 2016, 48, 747-757.	9.4	237
9	Differential Contributions of Rare and Common, Coding and Noncoding Ret Mutations to Multifactorial Hirschsprung Disease Liability. American Journal of Human Genetics, 2010, 87, 60-74.	2.6	230
10	Cohort Profile: The Dongfeng–Tongji cohort study of retired workers. International Journal of Epidemiology, 2013, 42, 731-740.	0.9	219
11	Metaâ€analysis on night shift work and risk of metabolic syndrome. Obesity Reviews, 2014, 15, 709-720.	3.1	218
12	Genetic polymorphisms of interleukin (IL)-1B, IL-1RN, IL-8, IL-10 and tumor necrosis factor \hat{l}_{\pm} and risk of gastric cancer in a Chinese population. Carcinogenesis, 2005, 26, 631-636.	1.3	212
13	Identification of Genetic Variants in Base Excision Repair Pathway and Their Associations with Risk of Esophageal Squamous Cell Carcinoma. Cancer Research, 2004, 64, 4378-4384.	0.4	208
14	Clinical Characteristics of and Medical Interventions for COVID-19 in Hemodialysis Patients in Wuhan, China. Journal of the American Society of Nephrology: JASN, 2020, 31, 1387-1397.	3.0	206
15	IncRNASNP: a database of SNPs in IncRNAs and their potential functions in human and mouse. Nucleic Acids Research, 2015, 43, D181-D186.	6.5	204
16	Identification of Functional Genetic Variants in Cyclooxygenase-2 and Their Association With Risk of Esophageal Cancer. Gastroenterology, 2005, 129, 565-576.	0.6	200
17	PancanQTL: systematic identification of cis-eQTLs and trans-eQTLs in 33 cancer types. Nucleic Acids Research, 2018, 46, D971-D976.	6.5	191
18	Polymorphisms of Death Pathway Genes FAS and FASL in Esophageal Squamous-Cell Carcinoma. Journal of the National Cancer Institute, 2004, 96, 1030-1036.	3.0	185

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19	Functional Haplotypes in the Promoter of Matrix Metalloproteinase-2 Predict Risk of the Occurrence and Metastasis of Esophageal Cancer. Cancer Research, 2004, 64, 7622-7628.	0.4	182
20	Genome-wide association study identifies <i>NRG1</i> as a susceptibility locus for Hirschsprung's disease. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 2694-2699.	3.3	171
21	The Role of P53 and MDM2 Polymorphisms in the Risk of Esophageal Squamous Cell Carcinoma. Cancer Research, 2005, 65, 9582-9587.	0.4	169
22	Genome-wide association study identifies five loci associated with susceptibility to pancreatic cancer in Chinese populations. Nature Genetics, 2012, 44, 62-66.	9.4	164
23	Clinical characteristics of 54 medical staff with COVID†19: A retrospective study in a single center in Wuhan, China. Journal of Medical Virology, 2020, 92, 807-813.	2.5	153
24	Joint analysis of three genome-wide association studies of esophageal squamous cell carcinoma in Chinese populations. Nature Genetics, 2014, 46, 1001-1006.	9.4	148
25	Genetic polymorphisms in cell cycle regulatory genesMDM2 andTP53 are associated with susceptibility to lung cancer. Human Mutation, 2006, 27, 110-117.	1.1	142
26	Aurora- A/STK15 T \pm 91A is a general low penetrance cancer susceptibility gene: a meta-analysis of multiple cancer types. Carcinogenesis, 2005, 26, 1368-1373.	1.3	132
27	Polymorphisms in DNA base excision repair genes ADPRT and XRCC1 and risk of lung cancer. Cancer Research, 2005, 65, 722-6.	0.4	127
28	Polymorphisms of DNA repair genesXRCC1 and XPD and their associations with risk of esophageal squamous cell carcinoma in a Chinese population. International Journal of Cancer, 2002, 100, 600-605.	2.3	120
29	Functional polymorphisms in cell death pathway genes FAS and FASL contribute to risk of lung cancer. Journal of Medical Genetics, 2005, 42, 479-484.	1.5	120
30	Genome-wide association study identifies a susceptibility locus for biliary atresia on 10q24.2. Human Molecular Genetics, 2010, 19, 2917-2925.	1.4	117
31	Urinary levels of bisphenol A, F and S and markers of oxidative stress among healthy adult men: Variability and association analysis. Environment International, 2019, 123, 301-309.	4.8	117
32	A novel T-77C polymorphism in DNA repair gene XRCC1 contributes to diminished promoter activity and increased risk of non-small cell lung cancer. Oncogene, 2006, 25, 3613-3620.	2.6	115
33	A genome-wide association study identifies two new cervical cancer susceptibility loci at 4q12 and 17q12. Nature Genetics, 2013, 45, 918-922.	9.4	108
34	Substantial reduction in risk of breast cancer associated with genetic polymorphisms in the promoters of the matrix metalloproteinase-2 and tissue inhibitor of metalloproteinase-2 genes. Carcinogenesis, 2003, 25, 399-404.	1.3	107
35	Meta-analysis of genome-wide association studies of adult height in East Asians identifies 17 novel loci. Human Molecular Genetics, 2015, 24, 1791-1800.	1.4	105
36	FASL –844C polymorphism is associated with increased activation-induced T cell death and risk of cervical cancer. Journal of Experimental Medicine, 2005, 202, 967-974.	4.2	104

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37	Sequence variations in the DNA repair geneXPD and risk of lung cancer in a Chinese population. International Journal of Cancer, 2003, 105, 669-673.	2.3	102
38	ELABELA and an ELABELA Fragment Protect against AKI. Journal of the American Society of Nephrology: JASN, 2017, 28, 2694-2707.	3.0	101
39	Identification of Functional Genetic Variants in and Their Association With Risk of Esophageal Cancer. Gastroenterology, 2005, 129, 565-576.	0.6	100
40	The Expression Levels of Plasma micoRNAs in Atrial Fibrillation Patients. PLoS ONE, 2012, 7, e44906.	1.1	97
41	Potential Influence of Menstrual Status and Sex Hormones on Female Severe Acute Respiratory Syndrome Coronavirus 2 Infection: A Cross-sectional Multicenter Study in Wuhan, China. Clinical Infectious Diseases, 2021, 72, e240-e248.	2.9	96
42	Imputation and subset-based association analysis across different cancer types identifies multiple independent risk loci in the TERT-CLPTM1L region on chromosome 5p15.33. Human Molecular Genetics, 2014, 23, 6616-6633.	1.4	90
43	DNA repair gene XPC genotypes/haplotypes and risk of lung cancer in a Chinese population. International Journal of Cancer, 2005, 115, 478-483.	2.3	87
44	Analysis of Ovarian Injury Associated With COVID-19 Disease in Reproductive-Aged Women in Wuhan, China: An Observational Study. Frontiers in Medicine, 2021, 8, 635255.	1,2	86
45	Reelin gene variants and risk of autism spectrum disorders: An integrated metaâ€analysis. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2014, 165, 192-200.	1.1	85
46	Common genetic variants regulating ADD3 gene expression alter biliary atresia risk. Journal of Hepatology, 2013, 59, 1285-1291.	1.8	84
47	Identification of a <i>HOXD13</i> mutation in a VACTERL patient. American Journal of Medical Genetics, Part A, 2008, 146A, 3181-3185.	0.7	82
48	Functional Variants in Cell Death Pathway Genes and Risk of Pancreatic Cancer. Clinical Cancer Research, 2008, 14, 3230-3236.	3.2	82
49	Functional STK15 Phe31Ile Polymorphism Is Associated with the Occurrence and Advanced Disease Status of Esophageal Squamous Cell Carcinoma. Cancer Research, 2004, 64, 2680-2683.	0.4	80
50	Dietary legume consumption reduces risk of colorectal cancer: evidence from a meta-analysis of cohort studies. Scientific Reports, 2015, 5, 8797.	1.6	79
51	Functional haplotypes in the promoter of matrix metalloproteinase-2 and lung cancer susceptibility. Carcinogenesis, 2005, 26, 1117-1121.	1.3	78
52	Genetic Variants at 6p21.1 and 7p15.3 Are Associated with Risk of Multiple Cancers in Han Chinese. American Journal of Human Genetics, 2012, 91, 928-934.	2.6	76
53	Exome-wide analyses identify low-frequency variant in CYP26B1 and additional coding variants associated with esophageal squamous cell carcinoma. Nature Genetics, 2018, 50, 338-343.	9.4	75
54	Genetic Variants in the Folate Pathway and the Risk of Neural Tube Defects: A Meta-Analysis of the Published Literature. PLoS ONE, 2013, 8, e59570.	1.1	74

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55	Common genetic variation in ETV6 is associated with colorectal cancer susceptibility. Nature Communications, 2016, 7, 11478.	5.8	73
56	A functional polymorphism in the matrix metalloproteinase-2 gene promoter (-1306C/T) is associated with risk of development but not metastasis of gastric cardia adenocarcinoma. Cancer Research, 2003, 63, 3987-90.	0.4	73
57	Association between serum uric acid and the metabolic syndrome among a middle- and old-age Chinese population. European Journal of Epidemiology, 2013, 28, 669-676.	2.5	72
58	Exposure to the Chinese Famine in Childhood Increases Type 2 Diabetes Risk in Adults. Journal of Nutrition, 2016, 146, 2289-2295.	1.3	70
59	Glyceraldehydeâ€3â€phosphate dehydrogenase promotes liver tumorigenesis by modulating phosphoglycerate dehydrogenase. Hepatology, 2017, 66, 631-645.	3.6	70
60	Environmental exposure to metals and the risk of hypertension: A cross-sectional study in China. Environmental Pollution, 2018, 233, 670-678.	3.7	70
61	Genetic variations in the TGFÂ signaling pathway, smoking and risk of colorectal cancer in a Chinese population. Carcinogenesis, 2013, 34, 936-942.	1.3	69
62	Evaluation of Genetic Susceptibility Loci for Chronic Hepatitis B in Chinese: Two Independent Case-Control Studies. PLoS ONE, 2011, 6, e17608.	1.1	69
63	A common polymorphism in the 3′UTR of cyclooxygenase 2/prostaglandin synthase 2 gene and risk of lung cancer in a Chinese population. Lung Cancer, 2005, 48, 11-17.	0.9	68
64	Interaction of P53 Arg72Pro and MDM2 T309G polymorphisms and their associations with risk of gastric cardia cancer. Carcinogenesis, 2007, 28, 1996-2001.	1.3	68
65	A functional polymorphism in <i>lnc-LAMC2-1:1</i> confers risk of colorectal cancer by affecting miRNA binding. Carcinogenesis, 2016, 37, 443-451.	1.3	68
66	Identification of new susceptibility loci for gastric non-cardia adenocarcinoma: pooled results from two Chinese genome-wide association studies. Gut, 2017, 66, 581-587.	6.1	68
67	A genome wide association study of genetic loci that influence tumour biomarkers cancer antigen 19-9, carcinoembryonic antigen and \hat{l}_{\pm} fetoprotein and their associations with cancer risk. Gut, 2014, 63, 143-151.	6.1	67
68	Longer Sleep Duration and Midday Napping Are Associated with a Higher Risk of CHD Incidence in Middle-Aged and Older Chinese: the Dongfeng-Tongji Cohort Study. Sleep, 2016, 39, 645-652.	0.6	64
69	Non-linear dose–response relationship between cigarette smoking and pancreatic cancer risk: Evidence from a meta-analysis of 42 observational studies. European Journal of Cancer, 2014, 50, 193-203.	1.3	63
70	The effect of laparoscopic excision vs open excision in children with choledochal cyst: a midterm follow-up study. Journal of Pediatric Surgery, 2011, 46, 662-665.	0.8	61
71	Identification of Genes Associated With Hirschsprung Disease, Based on Whole-Genome Sequence Analysis, and Potential Effects on Enteric Nervous System Development. Gastroenterology, 2018, 155, 1908-1922.e5.	0.6	61
72	CancerSplicingQTL: a database for genome-wide identification of splicing QTLs in human cancer. Nucleic Acids Research, 2019, 47, D909-D916.	6.5	61

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73	A functional polymorphism located at transcription factor binding sites, rs6695837 near LAMC1 gene, confers risk of colorectal cancer in Chinese populations. Carcinogenesis, 2016, 38, bgw204.	1.3	59
74	Exome Array Analysis Identifies Variants in SPOCD1 and BTN3A2 That Affect Risk for Gastric Cancer. Gastroenterology, 2017, 152, 2011-2021.	0.6	58
75	Severe acute respiratory syndrome coronavirus 2 detection in the female lower genital tract. American Journal of Obstetrics and Gynecology, 2020, 223, 131-134.	0.7	58
76	Functional Phe31lle polymorphism in Aurora A and risk of breast carcinoma. Carcinogenesis, 2004, 25, 2225-2230.	1.3	57
77	A Genome Wide Association Study Identifies Common Variants Associated with Lipid Levels in the Chinese Population. PLoS ONE, 2013, 8, e82420.	1.1	57
78	Genetic Polymorphisms in Methylenetetrahydrofolate Reductase and Thymidylate Synthase and Risk of Pancreatic Cancer. Clinical Gastroenterology and Hepatology, 2005, 3, 743-751.	2.4	56
79	Association between polymorphisms of <i>DRD2</i> and <i>DRD4</i> and opioid dependence: Evidence from the current studies. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 661-670.	1.1	55
80	Prevalence and Associated Risk Factors of Dyslexic Children in a Middle-Sized City of China: A Cross-Sectional Study. PLoS ONE, 2013, 8, e56688.	1.1	55
81	Polymorphisms of XRCC1 genes and risk of nasopharyngeal carcinoma in the Cantonese population. BMC Cancer, 2006, 6, 167.	1.1	54
82	A Rare Missense Variant in TCF7L2 Associates with Colorectal Cancer Risk by Interacting with a GWAS-Identified Regulatory Variant in the MYC Enhancer. Cancer Research, 2018, 78, 5164-5172.	0.4	54
83	Inactivation of DNA repair gene O6-methylguanine-DNA methyltransferase by promoter hypermethylation and its relation to p53 mutations in esophageal squamous cell carcinoma. Carcinogenesis, 2003, 24, 1039-1044.	1.3	53
84	Adenosine Diphosphate Ribosyl Transferase and X-Ray Repair Cross-Complementing 1 Polymorphisms in Gastric Cardia Cancer. Gastroenterology, 2006, 131, 420-427.	0.6	53
85	Genome-Wide Interrogation Identifies <i>YAP1</i> Variants Associated with Survival of Small-Cell Lung Cancer Patients. Cancer Research, 2010, 70, 9721-9729.	0.4	53
86	Allium Vegetables and Garlic Supplements Do Not Reduce Risk of Colorectal Cancer, Based on Meta-analysis of Prospective Studies. Clinical Gastroenterology and Hepatology, 2014, 12, 1991-2001.e4.	2.4	53
87	AWESOME: a database of SNPs that affect protein post-translational modifications. Nucleic Acids Research, 2019, 47, D874-D880.	6.5	53
88	N ⁶ -methyladenosine mRNA methylation of <i>PIK3CB</i> regulates AKT signalling to promote PTEN-deficient pancreatic cancer progression. Gut, 2020, 69, 2180-2192.	6.1	52
89	A functional polymorphism in the SULT1A1 gene (G638A) is associated with risk of lung cancer in relation to tobacco smoking. Carcinogenesis, 2003, 25, 773-778.	1.3	51
90	Genome-Wide Copy Number Analysis Uncovers a New HSCR Gene: NRG3. PLoS Genetics, 2012, 8, e1002687.	1.5	51

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91	Mutations in the NRG1 gene are associated with Hirschsprung disease. Human Genetics, 2012, 131, 67-76.	1.8	51
92	Risk SNP-Mediated Enhancer–Promoter Interaction Drives Colorectal Cancer through Both <i>FADS2</i> and <i>AP002754.2</i> Cancer Research, 2020, 80, 1804-1818.	0.4	50
93	Perturbation of Hoxb5 Signaling in Vagal Neural Crests Down-Regulates Ret Leading to Intestinal Hypoganglionosis in Mice. Gastroenterology, 2008, 134, 1104-1115.	0.6	49
94	A polymorphic MYC response element in KBTBD11 influences colorectal cancer risk, especially in interaction with an MYC-regulated SNP rs6983267. Annals of Oncology, 2018, 29, 632-639.	0.6	49
95	A lowâ \in frequency variant in SMAD7 modulates TGFâ \in fi ² signaling and confers risk for colorectal cancer in Chinese population. Molecular Carcinogenesis, 2017, 56, 1798-1807.	1.3	48
96	Reducing protein regulator of cytokinesis 1 as a prospective therapy for hepatocellular carcinoma. Cell Death and Disease, 2018, 9, 534.	2.7	48
97	Bidirectional association between nonalcoholic fatty liver disease and type 2 diabetes in Chinese population: Evidence from the Dongfeng-Tongji cohort study. PLoS ONE, 2017, 12, e0174291.	1.1	48
98	Genetic polymorphism in myeloperoxidase but not GSTM1 is associated with risk of lung squamous cell carcinoma in a Chinese population. International Journal of Cancer, 2002, 102, 275-279.	2.3	47
99	Dinucleotide polymorphism of p73 gene is associated with a reduced risk of lung cancer in a Chinese population. International Journal of Cancer, 2005, 114, 455-460.	2.3	47
100	Implications of Endocrine Gland–Derived Vascular Endothelial Growth Factor/Prokineticin-1 Signaling in Human Neuroblastoma Progression. Clinical Cancer Research, 2007, 13, 868-875.	3.2	47
101	Reduced RET expression in gut tissue of individuals carrying risk alleles of Hirschsprung's disease. Human Molecular Genetics, 2010, 19, 1461-1467.	1.4	47
102	Integrative expression quantitative trait locus–based analysis of colorectal cancer identified a functional polymorphism regulating SLC22A5 expression. European Journal of Cancer, 2018, 93, 1-9.	1.3	47
103	Urinary biomarkers of phthalates exposure and risks of thyroid cancer and benign nodule. Journal of Hazardous Materials, 2020, 383, 121189.	6.5	46
104	Association of candidate genetic variations with gastric cardia adenocarcinoma in Chinese population: a multiple interaction analysis. Carcinogenesis, 2011, 32, 336-342.	1.3	45
105	Pancan-meQTL: a database to systematically evaluate the effects of genetic variants on methylation in human cancer. Nucleic Acids Research, 2019, 47, D1066-D1072.	6.5	45
106	Genetic variations in <i>TERTâ€CLPTM111</i>	1.3	44
107	COVID-19 in Peritoneal Dialysis Patients. Clinical Journal of the American Society of Nephrology: CJASN, 2021, 16, 121-123.	2.2	44
108	<scp><i>Helicobacter pylori</i></scp> infection is associated with type 2 diabetes among a middle―and old―age Chinese population. Diabetes/Metabolism Research and Reviews, 2016, 32, 95-101.	1.7	43

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109	Functional evaluation of missense variations in the human MAD1L1 and MAD2L1 genes and their impact on susceptibility to lung cancer. Journal of Medical Genetics, 2010, 47, 616-622.	1.5	42
110	Associations of environmental exposure to metals with the risk of hypertension in China. Science of the Total Environment, 2018, 622-623, 184-191.	3.9	42
111	Interactions between Genetic Variants in the Adiponectin, Adiponectin Receptor 1 and Environmental Factors on the Risk of Colorectal Cancer. PLoS ONE, 2011, 6, e27301.	1.1	41
112	Exposure to the Chinese famine in early life and hypertension prevalence risk in adults. Journal of Hypertension, 2017, 35, 63-68.	0.3	41
113	Association of co-exposure to heavy metals with renal function in a hypertensive population. Environment International, 2018, 112, 198-206.	4.8	41
114	A Rare Variant P507L in TPP1 Interrupts TPP1–TIN2 Interaction, Influences Telomere Length, and Confers Colorectal Cancer Risk in Chinese Population. Cancer Epidemiology Biomarkers and Prevention, 2018, 27, 1029-1035.	1.1	41
115	Systematic Functional Interrogation of Genes in GWAS Loci Identified ATF1 as a Key Driver in Colorectal Cancer Modulated by a Promoter-Enhancer Interaction. American Journal of Human Genetics, 2019, 105, 29-47.	2.6	41
116	Circulating essential metals and lung cancer: Risk assessment and potential molecular effects. Environment International, 2019, 127, 685-693.	4.8	41
117	Dietary Mushroom Intake May Reduce the Risk of Breast Cancer: Evidence from a Meta-Analysis of Observational Studies. PLoS ONE, 2014, 9, e93437.	1.1	40
118	Comparative Evaluation of Participation and Diagnostic Yield of Colonoscopy vs Fecal Immunochemical Test vs Risk-Adapted Screening in Colorectal Cancer Screening: Interim Analysis of a Multicenter Randomized Controlled Trial (TARGET-C). American Journal of Gastroenterology, 2020, 115, 1264-1274.	0.2	40
119	Loss of myeloid-related proteins 8 and myeloid-related proteins 14 expression in human esophageal squamous cell carcinoma correlates with poor differentiation. World Journal of Gastroenterology, 2004, 10, 1093.	1.4	40
120	The SLC4A7 variant rs4973768 is associated with breast cancer risk: evidence from a case–control study and a meta-analysis. Breast Cancer Research and Treatment, 2012, 136, 847-857.	1.1	39
121	Association between bilirubin and risk of Non-Alcoholic Fatty Liver Disease based on a prospective cohort study. Scientific Reports, 2016, 6, 31006.	1.6	39
122	Potential of Arbidol for Post-exposure Prophylaxis of COVID-19 Transmission: A Preliminary Report of a Retrospective Cohort Study. Current Medical Science, 2020, 40, 480-485.	0.7	39
123	Combined Effect of Genetic Polymorphisms in P53, P73, and MDM2 on Non-small Cell Lung Cancer Survival. Journal of Thoracic Oncology, 2011, 6, 1793-1800.	0.5	38
124	Genetic variant in <i>KIAA0319</i> , but not in <i>DYX1C1</i> , is associated with risk of dyslexia: An integrated metaâ€analysis. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 970-976.	1.1	38
125	Epigenomic landscape of 5-hydroxymethylcytosine reveals its transcriptional regulation of lncRNAs in colorectal cancer. British Journal of Cancer, 2017, 116, 658-668.	2.9	38
126	Using different anthropometric indices to assess prediction ability of type 2 diabetes in elderly population: a 5Âyear prospective study. BMC Geriatrics, 2018, 18, 218.	1.1	38

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127	Common genetic polymorphisms in the 5′-flanking Region of the SULT1A1 gene: haplotypes and their association with platelet enzymatic activity. Pharmacogenetics and Genomics, 2005, 15, 465-473.	0.7	37
128	Significant increase in risk of gastroesophageal cancer is associated with interaction between promoter polymorphisms in thymidylate synthase and serum folate status. Carcinogenesis, 2005, 26, 1430-1435.	1.3	37
129	Risk prediction of esophageal squamous-cell carcinoma with common genetic variants and lifestyle factors in Chinese population. Carcinogenesis, 2013, 34, 1782-1786.	1.3	37
130	CYP2C19 polymorphism and clinical outcomes among patients of different races treated with clopidogrel: A systematic review and meta-analysis. Journal of Huazhong University of Science and Technology [Medical Sciences], 2015, 35, 147-156.	1.0	37
131	Fine Mapping of the NRG1 Hirschsprung's Disease Locus. PLoS ONE, 2011, 6, e16181.	1.1	37
132	Different Physical Activity Subtypes and Risk of Metabolic Syndrome in Middle-Aged and Older Chinese People. PLoS ONE, 2013, 8, e53258.	1.1	36
133	Platelet 12-lipoxygenase Arg261Gln polymorphism: functional characterization and association with risk of esophageal squamous cell carcinoma in combination with COX-2 polymorphisms. Pharmacogenetics and Genomics, 2007, 17, 197-205.	0.7	35
134	A functional polymorphism (â^1607 1Gâ†2G) in the matrix metalloproteinase†promoter is associated with development and progression of lung cancer. Cancer, 2011, 117, 5172-5181.	2.0	35
135	Genetic variants in the SWI/SNF complex and smoking collaborate to modify the risk of pancreatic cancer in a Chinese population. Molecular Carcinogenesis, 2015, 54, 761-768.	1.3	35
136	Sleep Duration and Midday Napping with 5-Year Incidence and Reversion of Metabolic Syndrome in Middle-Aged and Older Chinese. Sleep, 2016, 39, 1911-1918.	0.6	35
137	<i>ANKLE1</i> N ⁶ â€Methyladenosineâ€related variant is associated with colorectal cancer risk by maintaining the genomic stability. International Journal of Cancer, 2020, 146, 3281-3293.	2.3	35
138	Genetic variants in m6A modification genes are associated with esophageal squamous-cell carcinoma in the Chinese population. Carcinogenesis, 2020, 41, 761-768.	1.3	35
139	Liver Intestine-Cadherin (CDH17) Haplotype Is Associated with Increased Risk of Hepatocellular Carcinoma. Clinical Cancer Research, 2006, 12, 5248-5252.	3.2	34
140	A Solute Carrier Family 22 Member 3 Variant rs3088442 Gâ†'A Associated with Coronary Heart Disease Inhibits Lipopolysaccharide-induced Inflammatory Response. Journal of Biological Chemistry, 2015, 290, 5328-5340.	1.6	34
141	Green tea consumption is associated with reduced incident CHD and improved CHD-related biomarkers in the Dongfeng-Tongji cohort. Scientific Reports, 2016, 6, 24353.	1.6	34
142	Long sleep duration and afternoon napping are associated with higher risk of incident diabetes in middle-aged and older Chinese: the Dongfeng-Tongji cohort study. Annals of Medicine, 2016, 48, 216-223.	1.5	34
143	Susceptibility to gastric cardia adenocarcinoma and genetic polymorphisms in methylenetetrahydrofolate reductase in an at-risk Chinese population. Cancer Epidemiology Biomarkers and Prevention, 2002, 11, 1454-8.	1.1	34
144	MNX1 (HLXB9) mutations in Currarino patients. Journal of Pediatric Surgery, 2009, 44, 1892-1898.	0.8	33

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145	Serologic Detection of SARS-CoV-2 Infections in Hemodialysis Centers: A Multicenter Retrospective Study in Wuhan, China. American Journal of Kidney Diseases, 2020, 76, 490-499.e1.	2.1	33
146	A functional â^'77T>C polymorphism in XRCC1 is associated with risk of breast cancer. Breast Cancer Research and Treatment, 2011, 125, 479-487.	1.1	32
147	A Genetic Variant rs1801274 in FCGR2A as a Potential Risk Marker for Kawasaki Disease: A Case-Control Study and Meta-Analysis. PLoS ONE, 2014, 9, e103329.	1.1	32
148	Exome-wide analysis identifies three low-frequency missense variants associated with pancreatic cancer risk in Chinese populations. Nature Communications, 2018, 9, 3688.	5.8	32
149	Assessment of XPD Lys751Gln and XRCC1 T–77C polymorphisms in advanced non-small-cell lung cancer patients treated with platinum-based chemotherapy. Lung Cancer, 2011, 73, 110-115.	0.9	31
150	Systematic Confirmation Study of GWAS-Identified Genetic Variants for Kawasaki Disease in A Chinese Population. Scientific Reports, 2015, 5, 8194.	1.6	31
151	Variations in $\langle i \rangle$ HSPA1B $\langle i \rangle$ at 6p21.3 Are Associated with Lung Cancer Risk and Prognosis in Chinese Populations. Cancer Research, 2011, 71, 7576-7586.	0.4	30
152	Exome Sequencing Identified NRG3 as a Novel Susceptible Gene of Hirschsprung's Disease in a Chinese Population. Molecular Neurobiology, 2013, 47, 957-966.	1.9	30
153	Functional Polymorphisms in FAS/FASL System Increase the Risk of Neuroblastoma in Chinese Population. PLoS ONE, 2013, 8, e71656.	1.1	30
154	Prenatal phthalate exposure, birth outcomes and DNA methylation of Alu and LINE-1 repetitive elements: A pilot study in China. Chemosphere, 2018, 206, 759-765.	4.2	30
155	GSK-3Î ² Polymorphism Discriminates Bipolar Disorder and Schizophrenia: A Systematic Meta-Analysis. Molecular Neurobiology, 2013, 48, 404-411.	1.9	29
156	Independent and joint effects of moderate alcohol consumption and smoking on the risks of non-alcoholic fatty liver disease in elderly Chinese men. PLoS ONE, 2017, 12, e0181497.	1.1	28
157	Correlation Between Genetic Variations in Hox Clusters and Hirschsprung's Disease. Annals of Human Genetics, 2007, 71, 526-536.	0.3	27
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159	Polymorphisms of death pathway genes FAS and FASL and risk of nasopharyngeal carcinoma. Molecular Carcinogenesis, 2010, 49, 944-950.	1.3	27
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