Yonghong Li

List of Publications by Year
in descending order

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48 papers	4,508 citations	172386 29 h-index	233338 45 g-index
48 all docs	48 docs citations	48 times ranked	8880 citing authors

#	Article	IF	Citations
1	Genome-wide association study meta-analysis identifies seven new rheumatoid arthritis risk loci. Nature Genetics, 2010, 42, 508-514.	9.4	1,132
2	A Role for 12-lipoxygenase in Nerve Cell Death Caused by Glutathione Depletion. Neuron, 1997, 19, 453-463.	3.8	460
3	Meta-Analysis of Genome-Wide Association Studies in Celiac Disease and Rheumatoid Arthritis Identifies Fourteen Non-HLA Shared Loci. PLoS Genetics, 2011, 7, e1002004.	1.5	307
4	Genetic variants at CD28, PRDM1 and CD2/CD58 are associated with rheumatoid arthritis risk. Nature Genetics, 2009, 41, 1313-1318.	9.4	306
5	Evidence for novel susceptibility genes for late-onset Alzheimer's disease from a genome-wide association study of putative functional variants. Human Molecular Genetics, 2007, 16, 865-873.	1.4	256
6	A Scan of Chromosome 10 Identifies a Novel Locus Showing Strong Association with Late-Onset Alzheimer Disease. American Journal of Human Genetics, 2006, 78, 78-88.	2.6	157
7	Genome-wide meta-analysis identifies multiple novel associations and ethnic heterogeneity of psoriasis susceptibility. Nature Communications, 2015, 6, 6916.	5.8	154
8	Positive and negative regulation of APP amyloidogenesis by sumoylation. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 259-264.	3.3	140
9	Association of late-onset Alzheimer's disease with genetic variation in multiple members of the GAPD gene family. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 15688-15693.	3.3	134
10	Requirement for cGMP in Nerve Cell Death Caused by Glutathione Depletion. Journal of Cell Biology, 1997, 139, 1317-1324.	2.3	132
11	DAPK1 variants are associated with Alzheimer's disease and allele-specific expression. Human Molecular Genetics, 2006, 15, 2560-2568.	1.4	125
12	SORL1 variants and risk of late-onset Alzheimerâ€™s disease. Neurobiology of Disease, 2008, 29, 293-296.	2.1	78
13	Genetic Improvement ofHelicoverpa zeaNuclear Polyhedrosis Virus as a Biopesticide. Biological Control, 1997, 10, 83-91.	1.4	67
14	Further Genetic Evidence for Three Psoriasis-Risk Genes: ADAM33, CDKAL1, and PTPN22. Journal of Investigative Dermatology, 2009, 129, 629-634.	0.3	67
15	Multiple variants in toll-like receptor 4 gene modulate risk of liver fibrosis in Caucasians with chronic hepatitis C infection. Journal of Hepatology, 2009, 51, 750-757.	1.8	67
16	Carriers of Rare Missense Variants in IFIH1 Are Protected from Psoriasis. Journal of Investigative Dermatology, 2010, 130, 2768-2772.	0.3	65
17	Unraveling the genetics of complex diseases: Susceptibility genes for rheumatoid arthritis and psoriasis. Seminars in Immunology, 2009, 21, 318-327.	2.7	64
18	Isolation and Characterization of Apolipoproteins from Murine Microglia. Journal of Biological Chemistry, 2000, 275, 31770-31777.	1.6	60

#	Article	IF	Citations
19	Association of ABCA1 with late-onset Alzheimerâ€™s disease is not observed in a case-control study. Neuroscience Letters, 2004, 366, 268-271.	1.0	58
20	Genetic evidence for ubiquitin-specific proteasesUSP24andUSP40as candidate genes for late-onset Parkinson disease. Human Mutation, 2006, 27, 1017-1023.	1.1	53
21	Genetic Variants in the Apolipoprotein(a) Gene and Coronary Heart Disease. Circulation: Cardiovascular Genetics, 2011, 4, 565-573.	5.1	53
22	KIF6 Polymorphism as a Predictor of Risk of Coronary Events and of Clinical Event Reduction by Statin Therapy. American Journal of Cardiology, 2010, 106, 994-998.	0.7	48
23	A Multigene Test Could Cost-Effectively Help Extend Life Expectancy for Women at Risk of Hereditary Breast Cancer. Value in Health, 2017, 20, 547-555.	0.1	40
24	Expression and functional analysis of a baculovirus gene encodinga truncated protein kinase homolog. Virology, 1995, 206, 314-323.	1.1	39
25	Evidence that common variation in NEDD9 is associated with susceptibility to late-onset Alzheimer's and Parkinson's disease. Human Molecular Genetics, 2008, 17, 759-767.	1.4	39
26	A Case-Control Association Study of the 12 Single-Nucleotide Polymorphisms Implicated in Parkinson Disease by a Recent Genome Scan. American Journal of Human Genetics, 2006, 78, 1090-1092.	2.6	38
27	Ubiquilin 1 polymorphisms are not associated with late-onset Alzheimer's disease. Annals of Neurology, 2006, 59, 21-26.	2.8	37
28	Genetic association of the APP binding protein 2 gene (APBB2) with late onset Alzheimer disease. Human Mutation, 2005, 25, 270-277.	1.1	36
29	Statins Personalized. Medical Clinics of North America, 2012, 96, 123-139.	1.1	36
30	Cost Effectiveness of Sequencing 34 Cancer-Associated Genes as an Aid for Treatment Selection in Patients with Metastatic Melanoma. Molecular Diagnosis and Therapy, 2015, 19, 169-177.	1.6	28
31	The 5q31 variants associated with psoriasis and Crohn's disease are distinct. Human Molecular Genetics, 2008, 17, 2978-2985.	1.4	27
32	Deficiency of TDAG51 Protects Against Atherosclerosis by Modulating Apoptosis, Cholesterol Efflux, and Peroxiredoxinâ€1 Expression. Journal of the American Heart Association, 2013, 2, e000134.	1.6	27
33	The Up-Regulation of Endosomal-Lysosomal Components in Amyloid Î²-Resistant Cells. Journal of Neurochemistry, 2002, 73, 1477-1482.	2.1	25
34	Cost Effectiveness of Karyotyping, Chromosomal Microarray Analysis, and Targeted Next-Generation Sequencing of Patients with Unexplained Global Developmental Delay or Intellectual Disability. Molecular Diagnosis and Therapy, 2018, 22, 129-138.	1.6	23
35	Assessment of the Association of Vitamin D Level With SARS-CoV-2 Seropositivity Among Working-Age Adults. JAMA Network Open, 2021, 4, e2111634.	2.8	23
36	Genetic Risk Factors for Thrombosis in Systemic Lupus Erythematosus. Journal of Rheumatology, 2012, 39, 1603-1610.	1.0	22

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37	Genetic variants in the KIF6 region and coronary event reduction from statin therapy. Human Genetics, 2011, 129, 17-23.	1.8	18
38	Neither Replication nor Simulation Supports a Role for the Axon Guidance Pathway in the Genetics of Parkinson's Disease. PLoS ONE, 2008, 3, e2707.	1.1	17
39	Association of changes in lipid levels with changes in vitamin D levels in a real-world setting. Scientific Reports, 2021, 11, 21536.	1.6	17
40	Genetics of late-onset Alzheimerâ€™s disease: progress and prospect. Pharmacogenomics, 2007, 8, 1747-1755.	0.6	14
41	Brief Report: Singleâ€nucleotide polymorphisms in VKORC1 are risk factors for systemic lupus erythematosus in Asians. Arthritis and Rheumatism, 2013, 65, 211-215.	6.7	10
42	Reply to Bertram et al.. American Journal of Human Genetics, 2006, 79, 183-184.	2.6	4
43	Cost-effectiveness of nucleic acid amplification testing to guide treatment for vaginitis: a decision-modeling analysis. Diagnostic Microbiology and Infectious Disease, 2020, 98, 115119.	0.8	3
44	Analysis of Single Nucleotide Polymorphisms in Caseâ€“Control Studies. Methods in Molecular Biology, 2011, 719, 219-234.	0.4	1
45	A multigene test could cost-effectively help extend life expectancy for women at risk of hereditary breast cancerâ€”Reply to letter to the editor by Petelin et al.. Value in Health, 2018, 21, 893-894.	0.1	1
46	Changes in China call for new health solutions. Nature, 2005, 434, 821-821.	13.7	0
47	PhD: still necessary for independent research leaders. Nature, 2010, 464, 831-831.	13.7	0
48	Outreach and Connection to Care for Chronic Kidney Disease in a Workplace Wellness Setting: A Cost-Effectiveness Analysis. Population Health Management, 2020, 23, 487-494.	0.8	0