

# Dan Yin

## List of Publications by Year in descending order

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12  
papers

286  
citations

1040056

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1199594

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#	ARTICLE	IF	CITATIONS
1	Molecular Basis of Gene-Gene Interaction: Cyclic Cross-Regulation of Gene Expression and Post-GWAS Gene-Gene Interaction Involved in Atrial Fibrillation. <i>PLoS Genetics</i> , 2015, 11, e1005393.	3.5	47
2	Genomic variant in CAV1 increases susceptibility to coronary artery disease and myocardial infarction. <i>Atherosclerosis</i> , 2016, 246, 148-156.	0.8	44
3	Minor Allele C of Chromosome 1p32 Single Nucleotide Polymorphism rs11206510 Confers Risk of Ischemic Stroke in the Chinese Han Population. <i>Stroke</i> , 2010, 41, 1587-1592.	2.0	35
4	Genomic Variant in IL-37 Confers A Significant Risk of Coronary Artery Disease. <i>Scientific Reports</i> , 2017, 7, 42175.	3.3	31
5	Candidate Pathway-Based Genome-Wide Association Studies Identify Novel Associations of Genomic Variants in the Complement System Associated With Coronary Artery Disease. <i>Circulation: Cardiovascular Genetics</i> , 2014, 7, 887-894.	5.1	30
6	BRG1 variant rs1122608 on chromosome 19p13.2 confers protection against stroke and regulates expression of pre-mRNA-splicing factor SFRS3. <i>Human Genetics</i> , 2014, 133, 499-508.	3.8	24
7	Meta-analysis identifies robust association between SNP rs17465637 in MIA3 on chromosome 1q41 and coronary artery disease. <i>Atherosclerosis</i> , 2013, 231, 136-140.	0.8	22
8	Sigma-2 Receptor A Potential Target for Cancer/Alzheimer's Disease Treatment via Its Regulation of Cholesterol Homeostasis. <i>Molecules</i> , 2020, 25, 5439.	3.8	21
9	Raised serum ferritin concentration in hereditary hyperferritinemia cataract syndrome is not a marker for iron overload. <i>Hepatology</i> , 2014, 59, 1204-1206.	7.3	15
10	Significant Association between OPG/TNFRSF11B Variant and Common Complex Ischemic Stroke. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2018, 27, 1683-1691.	1.6	7
11	Demonstration of the Presence of the "Deleted" MIR122 Gene in HepG2 Cells. <i>PLoS ONE</i> , 2015, 10, e0122471.	2.5	6
12	Targeted next-generation sequencing identifies nine novel filaggrin gene variants in Chinese Han patients with ichthyosis vulgaris. <i>British Journal of Dermatology</i> , 2017, 177, e202-e203.	1.5	4