Dan Yin

List of Publications by Year
in descending order

12	286	1040056 9	1199594 12
papers	citations	h-index	g-index

13	13	13	577
all docs	docs citations	times ranked	citing authors

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