

# Tingwei Guo

## List of Publications by Year in descending order

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Version: 2024-02-01

13  
papers

526  
citations

840776

11  
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1199594

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13  
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times ranked

1319  
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#	ARTICLE	IF	CITATIONS
1	Genetic contributors to risk of schizophrenia in the presence of a 22q11.2 deletion. <i>Molecular Psychiatry</i> , 2021, 26, 4496-4510.	7.9	87
2	Genomewide meta-analysis identifies loci associated with IGF and IGFBP levels with impact on age-related traits. <i>Aging Cell</i> , 2016, 15, 811-824.	6.7	83
3	Copy-Number Variation of the Glucose Transporter Gene SLC2A3 and Congenital Heart Defects in the 22q11.2 Deletion Syndrome. <i>American Journal of Human Genetics</i> , 2015, 96, 753-764.	6.2	62
4	Genotype and cardiovascular phenotype correlations with TBX1 in 1,022 velo-cardio-facial/digeorge/22q11.2 deletion syndrome patients. <i>Human Mutation</i> , 2011, 32, 1278-1289.	2.5	57
5	Histone Modifier Genes Alter Conotruncal Heart Phenotypes in 22q11.2 Deletion Syndrome. <i>American Journal of Human Genetics</i> , 2015, 97, 869-877.	6.2	49
6	Rare copy number variants and congenital heart defects in the 22q11.2 deletion syndrome. <i>Human Genetics</i> , 2016, 135, 273-285.	3.8	43
7	Variance of IQ is partially dependent on deletion type among 1,427 22q11.2 deletion syndrome subjects. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 2172-2181.	1.2	33
8	Tbx1 is required autonomously for cell survival and fate in the pharyngeal core mesoderm to form the muscles of mastication. <i>Human Molecular Genetics</i> , 2014, 23, 4215-4231.	2.9	31
9	Reduced dosage of $\beta$ -catenin provides significant rescue of cardiac outflow tract anomalies in a Tbx1 conditional null mouse model of 22q11.2 deletion syndrome. <i>PLoS Genetics</i> , 2017, 13, e1006687.	3.5	27
10	Genome-Wide Association Study to Find Modifiers for Tetralogy of Fallot in the 22q11.2 Deletion Syndrome Identifies Variants in the <i>GPR98</i> Locus on 5q14.3. <i>Circulation: Cardiovascular Genetics</i> , 2017, 10, .	5.1	22
11	Deletion size analysis of 1680 22q11.2DS subjects identifies a new recombination hotspot on chromosome 22q11.2. <i>Human Molecular Genetics</i> , 2018, 27, 1150-1163.	2.9	22
12	HLA-B*07, HLA-DRB1*07, HLA-DRB1*12, and HLA-C*03:02 Strongly Associate With BMI: Data From 1.3 Million Healthy Chinese Adults. <i>Diabetes</i> , 2018, 67, 861-871.	0.6	9
13	Genome-Wide Association Studies of Conotruncal Heart Defects with Normally Related Great Vessels in the United States. <i>Genes</i> , 2021, 12, 1030.	2.4	1