Shuai Wang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2866482/publications.pdf

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18	1,546	933447 10	839539 18
papers	citations	h-index	g-index

22	22	22	5484
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The relationship between interstitial lung abnormalities, mortality, and multimorbidity: a cohort study. Thorax, 2023, 78, 559-565.	5.6	2
2	Impute the missing data using retrieved dropouts. BMC Medical Research Methodology, 2022, 22, 82.	3.1	2
3	Efficacy and Safety of Ertugliflozin in Patients With Diabetes Mellitus Inadequately Controlled by Sulfonylurea Monotherapy: a Substudy of VERTIS CV. Diabetes Therapy, 2021, 12, 1175-1192.	2.5	4
4	Genetic association tests in family samples for multi-category phenotypes. BMC Genomics, 2021, 22, 873.	2.8	1
5	Genetic Studies of Leptin Concentrations Implicate Leptin in the Regulation of Early Adiposity. Diabetes, 2020, 69, 2806-2818.	0.6	26
6	Exome-Derived Adiponectin-Associated Variants Implicate Obesity and Lipid Biology. American Journal of Human Genetics, 2019, 105, 15-28.	6.2	21
7	Exome sequencing of 20,791Âcases of type 2 diabetes and 24,440Âcontrols. Nature, 2019, 570, 71-76.	27.8	248
8	Protein-coding variants implicate novel genes related to lipid homeostasis contributing to body-fat distribution. Nature Genetics, 2019, 51, 452-469.	21.4	89
9	Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. Nature Genetics, 2018, 50, 26-41.	21.4	286
10	Rare and low-frequency coding variants alter human adult height. Nature, 2017, 542, 186-190.	27.8	544
11	Joint association analysis of a binary and a quantitative trait in family samples. European Journal of Human Genetics, 2017, 25, 130-136.	2.8	4
12	A candidate transacting modulator of fetal hemoglobin gene expression in the Arabâ€"Indian haplotype of sickle cell anemia. American Journal of Hematology, 2016, 91, 1118-1122.	4.1	16
13	General Framework for Metaâ€Analysis of Haplotype Association Tests. Genetic Epidemiology, 2016, 40, 244-252.	1.3	0
14	A genomic approach to therapeutic target validation identifies a glucose-lowering GLP1R variant protective for coronary heart disease. Science Translational Medicine, 2016, 8, 341ra76.	12.4	100
15	Comparison of multiple single-nucleotide variant association tests in a meta-analysis of Genetic Analysis Workshop 19 family and unrelated data. BMC Proceedings, 2016, 10, 187-191.	1.6	0
16	Low-frequency and rare exome chip variants associate with fasting glucose and type 2 diabetes susceptibility. Nature Communications, 2015, 6, 5897.	12.8	173
17	Association of Levels of Fasting Glucose and Insulin With Rare Variants at the Chromosome 11p11.2-MADD Locus. Circulation: Cardiovascular Genetics, 2014, 7, 374-382.	5.1	12
18	Comparing baseline and longitudinal measures in association studies. BMC Proceedings, 2014, 8, S84.	1.6	2