

Shing Wan Choi

List of Publications by Year in descending order

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Version: 2024-02-01

23
papers

2,947
citations

516710

16
h-index

713466

21
g-index

36
all docs

36
docs citations

36
times ranked

5901
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | PRSice-2: Polygenic Risk Score software for biobank-scale data. <i>GigaScience</i> , 2019, 8, . | 6.4 | 940 |
| 2 | Tutorial: a guide to performing polygenic risk score analyses. <i>Nature Protocols</i> , 2020, 15, 2759-2772. | 12.0 | 918 |
| 3 | Polygenic scores via penalized regression on summary statistics. <i>Genetic Epidemiology</i> , 2017, 41, 469-480. | 1.3 | 297 |
| 4 | Genome-wide gene-environment analyses of major depressive disorder and reported lifetime traumatic experiences in UK Biobank. <i>Molecular Psychiatry</i> , 2020, 25, 1430-1446. | 7.9 | 116 |
| 5 | Patient-specific induced-pluripotent stem cells-derived cardiomyocytes recapitulate the pathogenic phenotypes of dilated cardiomyopathy due to a novel DES mutation identified by whole exome sequencing. <i>Human Molecular Genetics</i> , 2013, 22, 1395-1403. | 2.9 | 98 |
| 6 | Multi-€“Polygenic Score Approach to Identifying Individual Vulnerabilities Associated With the Risk of Exposure to Bullying. <i>JAMA Psychiatry</i> , 2019, 76, 730. | 11.0 | 65 |
| 7 | Genetic analysis of amyotrophic lateral sclerosis identifies contributing pathways and cell types. <i>Science Advances</i> , 2021, 7, . | 10.3 | 59 |
| 8 | Association of Polygenic Risk for Attention-Deficit/Hyperactivity Disorder With Co-occurring Traits and Disorders. <i>Biological Psychiatry: Cognitive Neuroscience and Neuroimaging</i> , 2018, 3, 635-643. | 1.5 | 57 |
| 9 | Amelioration of X-Linked Related Autophagy Failure in Danon Disease With DNA Methylation Inhibitor. <i>Circulation</i> , 2016, 134, 1373-1389. | 1.6 | 42 |
| 10 | Novel pre-mRNA splicing of intronically integrated HBV generates oncogenic chimera in hepatocellular carcinoma. <i>Journal of Hepatology</i> , 2016, 64, 1256-1264. | 3.7 | 36 |
| 11 | Studying individual risk factors for self-harm in the UK Biobank: A polygenic scoring and Mendelian randomisation study. <i>PLoS Medicine</i> , 2020, 17, e1003137. | 8.4 | 34 |
| 12 | Genetic comorbidity between major depression and cardio-€“metabolic traits, stratified by age at onset of major depression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2020, 183, 309-330. | 1.7 | 33 |
| 13 | Genetic sensitivity analysis: Adjusting for genetic confounding in epidemiological associations. <i>PLoS Genetics</i> , 2021, 17, e1009590. | 3.5 | 30 |
| 14 | Cannabis use, depression and self-€“harm: phenotypic and genetic relationships. <i>Addiction</i> , 2020, 115, 482-492. | 3.3 | 29 |
| 15 | Classical Human Leukocyte Antigen Alleles and C4 Haplotypes Are Not Significantly Associated With Depression. <i>Biological Psychiatry</i> , 2020, 87, 419-430. | 1.3 | 27 |
| 16 | The expression of p53 antigen in primary malignant epithelial tumors of the liver: an immunohistochemical study. <i>Liver</i> , 1993, 13, 172-176. | 0.1 | 23 |
| 17 | Identification of novel locus associated with coronary artery aneurysms and validation of loci for susceptibility to Kawasaki disease. <i>European Journal of Human Genetics</i> , 2021, 29, 1734-1744. | 2.8 | 10 |
| 18 | Familial Influences on Neuroticism and Education in the UK Biobank. <i>Behavior Genetics</i> , 2020, 50, 84-93. | 2.1 | 9 |

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|----|--|-----|-----------|
| 19 | The Value of Rare Genetic Variation in the Prediction of Common Obesity in European Ancestry Populations. <i>Frontiers in Endocrinology</i> , 2022, 13, 863893. | 3.5 | 7 |
| 20 | Disentangling Independent and Mediated Causal Relationships Between Blood Metabolites, Cognitive Factors, and Alzheimer's Disease. <i>Biological Psychiatry Global Open Science</i> , 2022, 2, 167-179. | 2.2 | 6 |
| 21 | Detection of novel tandem duplication with next-generation sequencing. , 2011, , . | | 1 |
| 22 | Patient-specific induced-pluripotent stem cells derived cardiomyocytes recapitulate the pathogenic phenotypes of dilated cardiomyopathy due to a novel DES mutation identified by whole exome sequencing. <i>Human Molecular Genetics</i> , 2014, 23, 2232-2233. | 2.9 | 0 |
| 23 | Summaries of plenary, symposia, and oral sessions at the XXII World Congress of Psychiatric Genetics, Copenhagen, Denmark, 12-16 October 2014. <i>Psychiatric Genetics</i> , 2016, 26, 1-47. | 1.1 | 0 |