Dawei Li

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Meta-analysis shows significant association between dopamine system genes and attention deficit hyperactivity disorder (ADHD). Human Molecular Genetics, 2006, 15, 2276-2284. | 2.9 | 519 |
| 2 | Meta-analysis shows strong positive association of the neuregulin 1 (NRG1) gene with schizophrenia. Human Molecular Genetics, 2006, 15, 1995-2002. | 2.9 | 260 |
| 3 | Strong Association of the Alcohol Dehydrogenase 1B Gene (ADH1B) with Alcohol Dependence and Alcohol-Induced Medical Diseases. Biological Psychiatry, 2011, 70, 504-512. | 1.3 | 150 |
| 4 | Strong protective effect of the aldehyde dehydrogenase gene (ALDH2) 504lys (*2) allele against alcoholism and alcohol-induced medical diseases in Asians. Human Genetics, 2012, 131, 725-737. | 3.8 | 132 |
| 5 | Association of Gamma-Aminobutyric Acid A Receptor α2 Gene (GABRA2) with Alcohol Use Disorder. Neuropsychopharmacology, 2014, 39, 907-918. | 5.4 | 93 |
| 6 | Association study of serotonin 2A receptor (5-HT2A) gene with schizophrenia and suicidal behavior using systematic meta-analysis. Biochemical and Biophysical Research Communications, 2006, 340, 1006-1015. | 2.1 | 86 |
| 7 | Association of the HTR2A gene with alcohol and heroin abuse. Human Genetics, 2014, 133, 357-365. | 3.8 | 56 |
| 8 | Associations of the 5â€hydroxytryptamine (serotonin) Receptor 1B gene (<i>HTR1B</i>) with alcohol, cocaine, and heroin abuse. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 169-176. | 1.7 | 48 |
| 9 | A virome-wide clonal integration analysis platform for discovering cancer viral etiology. Genome Research, 2019, 29, 819-830. | 5.5 | 47 |
| 10 | Multi-Cultural Association of the Serotonin Transporter Gene (SLC6A4) with Substance Use Disorder. Neuropsychopharmacology, 2013, 38, 1737-1747. | 5.4 | 42 |
| 11 | Genome-Wide Association Study of Copy Number Variations (CNVs) with Opioid Dependence. Neuropsychopharmacology, 2015, 40, 1016-1026. | 5.4 | 39 |
| 12 | Comprehensive comparative analysis of methods and software for identifying viral integrations. Briefings in Bioinformatics, 2019, 20, 2088-2097. | 6.5 | 32 |
| 13 | Searching for human oncoviruses: Histories, challenges, and opportunities. Journal of Cellular Biochemistry, 2018, 119, 4897-4906. | 2.6 | 26 |
| 14 | ERVcaller: identifying polymorphic endogenous retrovirus and other transposable element insertions using whole-genome sequencing data. Bioinformatics, 2019, 35, 3913-3922. | 4.1 | 25 |
| 15 | Profile of circulating microRNAs in myalgic encephalomyelitis and their relation to symptom severity, and disease pathophysiology. Scientific Reports, 2020, 10, 19620. | 3.3 | 24 |
| 16 | Atlas of human diseases influenced by genetic variants with extreme allele frequency differences. Human Genetics, 2017, 136, 39-54. | 3.8 | 15 |
| 17 | GACT: a Genome build and Allele definition Conversion Tool for SNP imputation and meta-analysis in genetic association studies. BMC Genomics, 2014, 15, 610. | 2.8 | 7 |
| 18 | Eye color: A potential indicator of alcohol dependence risk in European Americans. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2015, 168, 347-353. | 1.7 | 7 |

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|----|---|-----|-----------|
| 19 | Characterization of Hepatitis B Virus Integrations Identified in Hepatocellular Carcinoma Genomes. Viruses, 2021, 13, 245. | 3.3 | 6 |
| 20 | Vlpower: Simulation-based tool for estimating power of viral integration detection via high-throughput sequencing. Genomics, 2020, 112, 207-211. | 2.9 | 4 |
| 21 | Association of Circulating YKL-40 Levels and CHI3L1 Variants with the Risk of Spinal Deformity Progression in Adolescent Idiopathic Scoliosis. Scientific Reports, 2019, 9, 5712. | 3.3 | 3 |
| 22 | Sequencing facility and DNA source associated patterns of virus-mappable reads in whole-genome sequencing data. Genomics, 2021, 113, 1189-1198. | 2.9 | 3 |
| 23 | Further analyses support the association between light eye color and alcohol dependence. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2015, 168, 757-760. | 1.7 | 1 |