## Jae Hoon Sul

List of Publications by Year in descending order

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713013 686830 1,607 24 13 21 h-index citations g-index papers 25 25 25 4922 docs citations times ranked citing authors all docs

| #  | Article  | IF           | Citations |
|----|--|--------------|-----------|
| 1  | Accurate diagnosis of atopic dermatitis by combining transcriptome and microbiota data with supervised machine learning. Scientific Reports, 2022, 12, 290.  | 1.6          | 12        |
| 2  | xGAP: a python based efficient, modular, extensible and fault tolerant genomic analysis pipeline for variant discovery. Bioinformatics, 2021, 37, 9-16.  | 1.8          | 0         |
| 3  | Purifying selection on noncoding deletions of human regulatory loci detected using their cellular pleiotropy. Genome Research, 2021, 31, 935-946.  | 2.4          | 5         |
| 4  | Rare variants regulate expression of nearby individual genes in multiple tissues. PLoS Genetics, 2021, 17, e1009596.   | 1.5          | 6         |
| 5  | Rare variants in the endocytic pathway are associated with Alzheimer's disease, its related phenotypes, and functional consequences. PLoS Genetics, 2021, 17, e1009772.  | 1.5          | 1         |
| 6  | Contribution of common and rare variants to bipolar disorder susceptibility in extended pedigrees from population isolates. Translational Psychiatry, 2020, 10, 74.  | 2.4          | 25        |
| 7  | Variant calling and quality control of large-scale human genome sequencing data. Emerging Topics in Life Sciences, 2019, 3, 399-409.   | 1.1          | 1         |
| 8  | Interrogating the Genetic Determinants of Tourette's Syndrome and Other Tic Disorders Through Genome-Wide Association Studies. American Journal of Psychiatry, 2019, 176, 217-227.                             | 4.0          | 242       |
| 9  | ForestQC: Quality control on genetic variants from next-generation sequencing data using random forest. PLoS Computational Biology, 2019, 15, e1007556.  | 1.5          | 17        |
| 10 | Leveraging allelic imbalance to refine fine-mapping for eQTL studies. PLoS Genetics, 2019, 15, e1008481.   | 1.5          | 20        |
| 11 | Multiethnic Meta-Analysis Identifies <i>RAI1</i> as a Possible Obstructive Sleep Apnea–related Quantitative Trait Locus in Men. American Journal of Respiratory Cell and Molecular Biology, 2018, 58, 391-401. | 1.4          | 65        |
| 12 | Population structure in genetic studies: Confounding factors and mixed models. PLoS Genetics, 2018, 14, e1007309.  | 1.5          | 164       |
| 13 | Understanding the Hidden Complexity of Latin American Population Isolates. American Journal of Human Genetics, 2018, 103, 707-726.   | 2.6          | 48        |
| 14 | Negative selection in humans and fruit flies involves synergistic epistasis. Science, 2017, 356, 539-542.  | 6.0          | 103       |
| 15 | Rare Copy Number Variants in NRXN1 and CNTN6 Increase Risk for Tourette Syndrome. Neuron, 2017, 94, 1101-1111.e7.  | 3 <b>.</b> 8 | 137       |
| 16 | Applying meta-analysis to genotype-tissue expression data from multiple tissues to identify eQTLs and increase the number of eGenes. Bioinformatics, 2017, 33, i67-i74.  | 1.8          | 21        |
| 17 | Accounting for Population Structure in Gene-by-Environment Interactions in Genome-Wide Association Studies Using Mixed Models. PLoS Genetics, 2016, 12, e1005849.  | 1.5          | 61        |
| 18 | A genome-wide association analysis of chromosomal aberrations and Hirschsprung disease. Translational Research, 2016, 177, 31-40.e6.   | 2.2          | 10        |

| #  | Article   | IF  | CITATION |
|----|---|-----|----------|
| 19 | A general framework for meta-analyzing dependent studies with overlapping subjects in association mapping. Human Molecular Genetics, 2016, 25, 1857-1866.                         | 1.4 | 42       |
| 20 | Increasing Generality and Power of Rare-Variant Tests by Utilizing Extended Pedigrees. American Journal of Human Genetics, 2016, 99, 846-859.                                     | 2.6 | 26       |
| 21 | Colocalization of GWAS and eQTL Signals Detects Target Genes. American Journal of Human Genetics, 2016, 99, 1245-1260.  | 2.6 | 569      |
| 22 | An efficient linear mixed model framework for meta-analytic association studies across multiple contexts. Leibniz International Proceedings in Informatics, LIPIcs, 2016, 2016, . | 0.0 | 0        |
| 23 | Accurate and Fast Multiple-Testing Correction in eQTL Studies. American Journal of Human Genetics, 2015, 96, 857-868.   | 2.6 | 25       |
| 24 | Gene–Gene Interactions Detection Using a Two-stage Model. Journal of Computational Biology, 2015, 22, 563-576.  | 0.8 | 5        |