

# Jaroslav Bendl

## List of Publications by Year in descending order

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Version: 2024-02-01

30  
papers

4,615  
citations

331259

21  
h-index

476904

29  
g-index

45  
all docs

45  
docs citations

45  
times ranked

8617  
citing authors

#	ARTICLE	IF	CITATIONS
1	Sex Differences in the Human Brain Transcriptome of Cases With Schizophrenia. <i>Biological Psychiatry</i> , 2022, 91, 92-101.	0.7	38
2	Multi-ancestry eQTL meta-analysis of human brain identifies candidate causal variants for brain-related traits. <i>Nature Genetics</i> , 2022, 54, 161-169.	9.4	49
3	Chromatin domain alterations linked to 3D genome organization in a large cohort of schizophrenia and bipolar disorder brains. <i>Nature Neuroscience</i> , 2022, 25, 474-483.	7.1	25
4	Common variants contribute to intrinsic human brain functional networks. <i>Nature Genetics</i> , 2022, 54, 508-517.	9.4	37
5	Impact of schizophrenia GWAS loci converge onto distinct pathways in cortical interneurons vs glutamatergic neurons during development. <i>Molecular Psychiatry</i> , 2022, 27, 4218-4233.	4.1	6
6	ATAC-seq and psychiatric disorders. , 2021, , 143-162.		0
7	Integration of Alzheimer's disease genetics and myeloid genomics identifies disease risk regulatory elements and genes. <i>Nature Communications</i> , 2021, 12, 1610.	5.8	118
8	Common genetic variation influencing human white matter microstructure. <i>Science</i> , 2021, 372, .	6.0	106
9	Neuronal and glial 3D chromatin architecture informs the cellular etiology of brain disorders. <i>Nature Communications</i> , 2021, 12, 3968.	5.8	48
10	Unbiased identification of novel transcription factors in striatal compartmentation and striosome maturation. <i>ELife</i> , 2021, 10, .	2.8	9
11	Common schizophrenia risk variants are enriched in open chromatin regions of human glutamatergic neurons. <i>Nature Communications</i> , 2020, 11, 5581.	5.8	53
12	decorate: differential epigenetic correlation test. <i>Bioinformatics</i> , 2020, 36, 2856-2861.	1.8	11
13	A chromosomal connectome for psychiatric and metabolic risk variants in adult dopaminergic neurons. <i>Genome Medicine</i> , 2020, 12, 19.	3.6	31
14	CommonMind Consortium provides transcriptomic and epigenomic data for Schizophrenia and Bipolar Disorder. <i>Scientific Data</i> , 2019, 6, 180.	2.4	149
15	Functional interpretation of genetic variants using deep learning predicts impact on chromatin accessibility and histone modification. <i>Nucleic Acids Research</i> , 2019, 47, 10597-10611.	6.5	39
16	Transcriptome and epigenome landscape of human cortical development modeled in organoids. <i>Science</i> , 2018, 362, .	6.0	220
17	Integrative functional genomic analysis of human brain development and neuropsychiatric risks. <i>Science</i> , 2018, 362, .	6.0	516
18	Transcriptome-wide isoform-level dysregulation in ASD, schizophrenia, and bipolar disorder. <i>Science</i> , 2018, 362, .	6.0	805

#	ARTICLE	IF	CITATIONS
19	Comprehensive functional genomic resource and integrative model for the human brain. <i>Science</i> , 2018, 362, .	6.0	618
20	An atlas of chromatin accessibility in the adult human brain. <i>Genome Research</i> , 2018, 28, 1243-1252.	2.4	170
21	The Mount Sinai cohort of large-scale genomic, transcriptomic and proteomic data in Alzheimer's disease. <i>Scientific Data</i> , 2018, 5, 180185.	2.4	320
22	NewProt – a protein engineering portal. <i>Protein Engineering, Design and Selection</i> , 2017, 30, 441-447.	1.0	11
23	FireProt: web server for automated design of thermostable proteins. <i>Nucleic Acids Research</i> , 2017, 45, W393-W399.	6.5	104
24	HotSpot Wizard 2.0: automated design of site-specific mutations and smart libraries in protein engineering. <i>Nucleic Acids Research</i> , 2016, 44, W479-W487.	6.5	76
25	PredictSNP2: A Unified Platform for Accurately Evaluating SNP Effects by Exploiting the Different Characteristics of Variants in Distinct Genomic Regions. <i>PLoS Computational Biology</i> , 2016, 12, e1004962.	1.5	149
26	FireProt: Energy- and Evolution-Based Computational Design of Thermostable Multiple-Point Mutants. <i>PLoS Computational Biology</i> , 2015, 11, e1004556.	1.5	144
27	PredictSNP: Robust and Accurate Consensus Classifier for Prediction of Disease-Related Mutations. <i>PLoS Computational Biology</i> , 2014, 10, e1003440.	1.5	593
28	Maximizing the Efficiency of Multienzyme Process by Stoichiometry Optimization. <i>ChemBioChem</i> , 2014, 15, 1891-1895.	1.3	31
29	Computer-Assisted Engineering of the Synthetic Pathway for Biodegradation of a Toxic Persistent Pollutant. <i>ACS Synthetic Biology</i> , 2014, 3, 172-181.	1.9	39
30	Computational Tools for Designing Smart Libraries. <i>Methods in Molecular Biology</i> , 2014, 1179, 291-314.	0.4	21