

Jun-Feng Xia

List of Publications by Year in descending order

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Version: 2024-02-01

86
papers

2,823
citations

201385

27
h-index

189595

50
g-index

90
all docs

90
docs citations

90
times ranked

3749
citing authors

#	ARTICLE	IF	CITATIONS
1	Prediction of protein-protein interactions from amino acid sequences with ensemble extreme learning machines and principal component analysis. <i>BMC Bioinformatics</i> , 2013, 14, S10.	1.2	232
2	<i>BRAF</i> L597 Mutations in Melanoma Are Associated with Sensitivity to MEK Inhibitors. <i>Cancer Discovery</i> , 2012, 2, 791-797.	7.7	194
3	APIS: accurate prediction of hot spots in protein interfaces by combining protrusion index with solvent accessibility. <i>BMC Bioinformatics</i> , 2010, 11, 174.	1.2	178
4	Sequence-Based Prediction of Protein-Protein Interactions by Means of Rotation Forest and Autocorrelation Descriptor. <i>Protein and Peptide Letters</i> , 2010, 17, 137-145.	0.4	144
5	Prediction of Protein-Protein Interactions from Protein Sequence Using Local Descriptors. <i>Protein and Peptide Letters</i> , 2010, 17, 1085-1090.	0.4	144
6	Automatic classification for field crop insects via multiple-task sparse representation and multiple-kernel learning. <i>Computers and Electronics in Agriculture</i> , 2015, 119, 123-132.	3.7	120
7	Application of next generation sequencing to human gene fusion detection: computational tools, features and perspectives. <i>Briefings in Bioinformatics</i> , 2013, 14, 506-519.	3.2	102
8	<i>MET</i> Exon 14 Skipping in Non-Small Cell Lung Cancer. <i>Oncologist</i> , 2016, 21, 481-486.	1.9	94
9	Investigating the relationship of DNA methylation with mutation rate and allele frequency in the human genome. <i>BMC Genomics</i> , 2012, 13, S7.	1.2	92
10	A Meta-analysis of Somatic Mutations from Next Generation Sequencing of 241 Melanomas: A Road Map for the Study of Genes with Potential Clinical Relevance. <i>Molecular Cancer Therapeutics</i> , 2014, 13, 1918-1928.	1.9	84
11	Predicting protein-protein interactions from sequence using correlation coefficient and high-quality interaction dataset. <i>Amino Acids</i> , 2010, 38, 891-899.	1.2	82
12	Predicting protein-protein interactions from protein sequences using meta predictor. <i>Amino Acids</i> , 2010, 39, 1595-1599.	1.2	78
13	Comparison and integration of computational methods for deleterious synonymous mutation prediction. <i>Briefings in Bioinformatics</i> , 2020, 21, 970-981.	3.2	56
14	Functional complementation between transcriptional methylation regulation and post-transcriptional microRNA regulation in the human genome. <i>BMC Genomics</i> , 2011, 12, S15.	1.2	52
15	GAERF: predicting lncRNA-disease associations by graph auto-encoder and random forest. <i>Briefings in Bioinformatics</i> , 2021, 22, .	3.2	51
16	BBPpred: Sequence-Based Prediction of Blood-Brain Barrier Peptides with Feature Representation Learning and Logistic Regression. <i>Journal of Chemical Information and Modeling</i> , 2021, 61, 525-534.	2.5	49
17	Next-generation sequencing of paired tyrosine kinase inhibitor-sensitive and -resistant EGFR mutant lung cancer cell lines identifies spectrum of DNA changes associated with drug resistance. <i>Genome Research</i> , 2013, 23, 1434-1445.	2.4	48
18	Computational identification of deleterious synonymous variants in human genomes using a feature-based approach. <i>BMC Medical Genomics</i> , 2019, 12, 12.	0.7	47

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19	Prediction of circRNA-disease associations based on inductive matrix completion. BMC Medical Genomics, 2020, 13, 42.	0.7	42
20	Do cancer proteins really interact strongly in the human protein-protein interaction network?. Computational Biology and Chemistry, 2011, 35, 121-125.	1.1	38
21	Inconsistency and features of single nucleotide variants detected in whole exome sequencing versus transcriptome sequencing: A case study in lung cancer. Methods, 2015, 83, 118-127.	1.9	33
22	dbDSM: a manually curated database for deleterious synonymous mutations. Bioinformatics, 2016, 32, 1914-1916.	1.8	33
23	NGS catalog: A database of next generation sequencing studies in humans. Human Mutation, 2012, 33, E2341-E2355.	1.1	32
24	A Sequence-Based Dynamic Ensemble Learning System for Protein Ligand-Binding Site Prediction. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2016, 13, 901-912.	1.9	32
25	Cancer Subtype Discovery Based on Integrative Model of Multigenomic Data. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2017, 14, 1115-1121.	1.9	32
26	A feature-based approach to predict hot spots in protein-DNA binding interfaces. Briefings in Bioinformatics, 2020, 21, 1038-1046.	3.2	31
27	Prediction of Neuropeptides from Sequence Information Using Ensemble Classifier and Hybrid Features. Journal of Proteome Research, 2020, 19, 3732-3740.	1.8	31
28	A Deep Learning-Based Method for Identification of Bacteriophage-Host Interaction. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2021, 18, 1801-1810.	1.9	31
29	Exploiting a Reduced Set of Weighted Average Features to Improve Prediction of DNA-Binding Residues from 3D Structures. PLoS ONE, 2011, 6, e28440.	1.1	30
30	Identifying multi-functional bioactive peptide functions using multi-label deep learning. Briefings in Bioinformatics, 2022, 23, .	3.2	30
31	PredCID: prediction of driver frameshift indels in human cancer. Briefings in Bioinformatics, 2021, 22, .	3.2	29
32	LNDriver: identifying driver genes by integrating mutation and expression data based on gene-gene interaction network. BMC Bioinformatics, 2016, 17, 467.	1.2	28
33	CINOEDV: a co-information based method for detecting and visualizing n-order epistatic interactions. BMC Bioinformatics, 2016, 17, 214.	1.2	27
34	Computational Methods for the Prediction of Protein-Protein Interactions. Protein and Peptide Letters, 2010, 17, 1069-1078.	0.4	23
35	Prioritizing Cancer Genes Based on an Improved Random Walk Method. Frontiers in Genetics, 2020, 11, 377.	1.1	22
36	Predicting hot spots in protein interfaces based on protrusion index, pseudo hydrophobicity and electron-ion interaction pseudopotential features. Oncotarget, 2016, 7, 18065-18075.	0.8	21

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37	Somatic synonymous mutations in regulatory elements contribute to the genetic aetiology of melanoma. <i>BMC Medical Genomics</i> , 2020, 13, 43.	0.7	20
38	Identification of ovarian cancer subtype-specific network modules and candidate drivers through an integrative genomics approach. <i>Oncotarget</i> , 2016, 7, 4298-4309.	0.8	20
39	Virus interactions with human signal transduction pathways. <i>International Journal of Computational Biology and Drug Design</i> , 2011, 4, 83.	0.3	19
40	Extra Trees Method for Predicting LncRNA-Disease Association Based on Multi-layer Graph Embedding Aggregation. <i>IEEE/ACM Transactions on Computational Biology and Bioinformatics</i> , 2021, PP, 1-1.	1.9	19
41	Identification of breast cancer candidate genes using gene co-expression and protein-protein interaction information. <i>Oncotarget</i> , 2016, 7, 36092-36100.	0.8	19
42	Identification of mutated driver pathways in cancer using a multi-objective optimization model. <i>Computers in Biology and Medicine</i> , 2016, 72, 22-29.	3.9	18
43	Prediction of hot spots in protein-DNA binding interfaces based on supervised isometric feature mapping and extreme gradient boosting. <i>BMC Bioinformatics</i> , 2020, 21, 381.	1.2	18
44	An analysis of mutational signatures of synonymous mutations across 15 cancer types. <i>BMC Medical Genetics</i> , 2019, 20, 190.	2.1	15
45	dbCPG: A web resource for cancer predisposition genes. <i>Oncotarget</i> , 2016, 7, 37803-37811.	0.8	15
46	OCCGene: a database of experimentally verified ovarian cancer-related genes with precomputed regulation information. <i>Cell Death and Disease</i> , 2015, 6, e2036-e2036.	2.7	13
47	Concordance between somatic copy number loss and down-regulated expression: A pan-cancer study of cancer predisposition genes. <i>Scientific Reports</i> , 2016, 6, 37358.	1.6	13
48	Module Based Differential Coexpression Analysis Method for Type 2 Diabetes. <i>BioMed Research International</i> , 2015, 2015, 1-8.	0.9	12
49	DriverFinder: A Gene Length-Based Network Method to Identify Cancer Driver Genes. <i>Complexity</i> , 2017, 2017, 1-10.	0.9	12
50	usDSM: a novel method for deleterious synonymous mutation prediction using undersampling scheme. <i>Briefings in Bioinformatics</i> , 2021, 22, .	3.2	12
51	Prediction of α -Hairpins in Proteins Using Physicochemical Properties and Structure Information. <i>Protein and Peptide Letters</i> , 2010, 17, 1123-1128.	0.4	11
52	Improved Method for Predicting β -Turns in Proteins Using a Two-Stage Classifier. <i>Protein and Peptide Letters</i> , 2010, 17, 1117-1122.	0.4	11
53	MGDB: a comprehensive database of genes involved in melanoma. <i>Database: the Journal of Biological Databases and Curation</i> , 2015, 2015, bav097.	1.4	11
54	Complete Genome Sequence of <i>Clostridium kluveri</i> JZZ Applied in Chinese Strong-Flavor Liquor Production. <i>Current Microbiology</i> , 2018, 75, 1429-1433.	1.0	11

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55	dbCID: a manually curated resource for exploring the driver indels in human cancer. Briefings in Bioinformatics, 2019, 20, 1925-1933.	3.2	11
56	Prediction of cancer cell sensitivity to natural products based on genomic and chemical properties. PeerJ, 2015, 3, e1425.	0.9	11
57	dbCPM: a manually curated database for exploring the cancer passenger mutations. Briefings in Bioinformatics, 2018, , .	3.2	10
58	Pan-Cancer Analysis of Radiotherapy Benefits and Immune Infiltration in Multiple Human Cancers. Cancers, 2020, 12, 957.	1.7	10
59	Double matrix completion for circRNA-disease association prediction. BMC Bioinformatics, 2021, 22, 307.	1.2	10
60	PredAPP: Predicting Anti-Parasitic Peptides with Undersampling and Ensemble Approaches. Interdisciplinary Sciences, Computational Life Sciences, 2022, 14, 258-268.	2.2	10
61	WEIGHTED NEIGHBORHOOD CLASSIFIER FOR THE CLASSIFICATION OF IMBALANCED TUMOR DATASET. Journal of Circuits, Systems and Computers, 2010, 19, 259-273.	1.0	9
62	dbCRSR: a manually curated database for regulation of cancer radiosensitivity. Database: the Journal of Biological Databases and Curation, 2018, 2018, .	1.4	8
63	Prediction of Radiosensitivity in Head and Neck Squamous Cell Carcinoma Based on Multiple Omics Data. Frontiers in Genetics, 2020, 11, 960.	1.1	8
64	Prediction of circRNA-Disease Associations Based on the Combination of Multi-Head Graph Attention Network and Graph Convolutional Network. Biomolecules, 2022, 12, 932.	1.8	8
65	Efficient Ensemble Schemes for Protein Secondary Structure Prediction. Protein and Peptide Letters, 2008, 15, 488-493.	0.4	7
66	Prediction of proteinâ€™protein interaction sites by means of ensemble learning and weighted feature descriptor. Journal of Biological Research, 2016, 23, 10.	2.2	7
67	SPHot: Prediction of Hot Spots in Protein-RNA Complexes by Protein Sequence Information and Ensemble Classifier. IEEE Access, 2019, 7, 104941-104946.	2.6	7
68	Biomedical Data Integration, Modeling, and Simulation in the Era of Big Data and Translational Medicine. BioMed Research International, 2014, 2014, 1-1.	0.9	6
69	An Ensemble Framework for Improving the Prediction of Deleterious Synonymous Mutation. IEEE Transactions on Circuits and Systems for Video Technology, 2022, 32, 2603-2611.	5.6	6
70	schFC: a hybrid fuzzy clustering method for single-cell RNA-seq data optimized by natural computation. Briefings in Bioinformatics, 2022, 23, .	3.2	6
71	Identification of driver pathways in cancer based on combinatorial patterns of somatic gene mutations. Neoplasia, 2016, 63, 57-63.	0.7	5
72	An improved DNA-binding hot spot residues prediction method by exploring interfacial neighbor properties. BMC Bioinformatics, 2021, 22, 253.	1.2	4

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73	MECoRank: cancer driver genes discovery simultaneously evaluating the impact of SNVs and differential expression on transcriptional networks. BMC Medical Genomics, 2019, 12, 140.	0.7	3
74	Identification of driver genes based on gene mutational effects and network centrality. BMC Bioinformatics, 2021, 22, 457.	1.2	3
75	dbBIP: a comprehensive bipolar disorder database for genetic research. Database: the Journal of Biological Databases and Curation, 2022, 2022, .	1.4	3
76	Network analysis of gene fusions in human cancer. BMC Bioinformatics, 2013, 14, A13.	1.2	2
77	Comprehensive Analysis of Alzheimer's Disease Biologically Candidate Causal Genes Revealed by Function Association Study With GWAS. IEEE Access, 2019, 7, 114236-114245.	2.6	2
78	Inferring Strengths of Protein-Protein Interaction Using Artificial Neural Network. Neural Networks (IJCNN), International Joint Conference on, 2007, , .	0.0	1
79	Differential coexpression analysis in gene modules level and its application to type 2 diabetes. , 2013, , .		1
80	Prediction of cytochrome P450 inhibition using ensemble of extreme learning machine. , 2013, , .		0
81	Cancer genes discovery based on integrating transcriptomic data and the impact of gene length. , 2016, , .		0
82	Sequence-Based Prediction of Transmembrane Protein Crystallization Propensity. Interdisciplinary Sciences, Computational Life Sciences, 2021, 13, 693-702.	2.2	0
83	Prediction of Protein-Protein Interaction Sites by Using Autocorrelation Descriptor and Support Vector Machine. Lecture Notes in Computer Science, 2010, , 76-82.	1.0	0
84	Potential Driver Genes Regulated by OncomiRNA Are Associated with Druggability in Pan-Negative Melanoma. Lecture Notes in Computer Science, 2014, , 315-321.	1.0	0
85	Comparative Assessment of Data Sets of Protein Interaction Hot Spots Used in the Computational Method. Lecture Notes in Computer Science, 2014, , 478-486.	1.0	0
86	Computational Prediction of Driver Missense Mutations in Melanoma. Lecture Notes in Computer Science, 2018, , 438-447.	1.0	0