Daniel S Himmelstein

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

97
papers

2,057
citations

16
h-index

2,831
ext. papers

2,831
avg, IF

45
g-index

4.74
L-index

#	Paper	IF	Citations
97	PMLB v1.0: An open-source dataset collection for benchmarking machine learning methods. <i>Bioinformatics</i> , 2021 ,	7.2	3
96	Is authorship sufficient for today's collaborative research? A call for contributor roles. <i>Accountability in Research</i> , 2021 , 28, 23-43	1.9	17
95	Analysis of scientific society honors reveals disparities. <i>Cell Systems</i> , 2021 , 12, 900-906.e5	10.6	2
94	An Open-Publishing Response to the COVID-19 Infodemic <i>CEUR Workshop Proceedings</i> , 2021 , 2976, 29-38	0.2	
93	Compressing gene expression data using multiple latent space dimensionalities learns complementary biological representations. <i>Genome Biology</i> , 2020 , 21, 109	18.3	20
92	Open collaborative writing with Manubot. PLoS Computational Biology, 2019, 15, e1007128	5	24
91	Opportunities and obstacles for deep learning in biology and medicine. <i>Journal of the Royal Society Interface</i> , 2018 , 15,	4.1	780
90	Sci-Hub provides access to nearly all scholarly literature. <i>ELife</i> , 2018 , 7,	8.9	59
89	Precision annotation of digital samples in NCBI's gene expression omnibus. <i>Scientific Data</i> , 2017 , 4, 170	18.5	19
88	Systematic integration of biomedical knowledge prioritizes drugs for repurposing. <i>ELife</i> , 2017 , 6,	8.9	151
87	Visualizing the top epilepsy predictions in Cytoscape 2017 ,		2
86	Why we predicted ictogenic tricyclic compounds treat epilepsy? 2017,		2
85	Association of HLA Genetic Risk Burden With Disease Phenotypes in Multiple Sclerosis. <i>JAMA Neurology</i> , 2016 , 73, 795-802	17.2	43
84	Data nomenclature: naming and abbreviating our network types 2016,		2
83	Positive correlations between knockdown and overexpression profiles from LINCS L1000 2016 ,		2
82	Assessing the effectiveness of our hetnet permutations 2016,		2
81	Workshop to analyze LINCS data for the Systems Pharmacology course at UCSF 2016,		2

(2016-2016)

80	Announcing PharmacotherapyDB: the Open Catalog of Drug Therapies for Disease 2016,	2
79	Assessing the imputation quality of gene expression in LINCS L1000 2016 ,	3
78	Incorporating DrugCentral data in our network 2016 ,	2
77	Estimating the complexity of hetnet traversal 2016 ,	2
76	Transforming DWPCs for hetnet edge prediction 2016 ,	2
75	Network Edge Prediction: how to deal with self-testing 2016 ,	2
74	Measuring user contribution and content creation 2016,	2
73	Network Edge Prediction: Estimating the prior 2016 ,	2
72	Predictions of whether a compound treats a disease 2016 ,	2
71	Computing standardized logistic regression coefficients 2016 ,	3
70	Our hetnet edge prediction methodology: the modeling framework for Project Rephetio 2016,	2
7º 69	Our hetnet edge prediction methodology: the modeling framework for Project Rephetio 2016 , Cataloging drugdisease therapies in the ClinicalTrials.gov database 2016 ,	2
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69	Cataloging drugdisease therapies in the ClinicalTrials.gov database 2016 ,	2
69	Cataloging drugdisease therapies in the ClinicalTrials.gov database 2016, Sounding the alarm on DrugBankd new license and terms of use 2016,	2
69 68 67	Cataloging drugdisease therapies in the ClinicalTrials.gov database 2016, Sounding the alarm on DrugBankd new license and terms of use 2016, Edge dropout contamination in hetnet edge prediction 2016,	2 2
69 68 67 66	Cataloging drugflisease therapies in the ClinicalTrials.gov database 2016, Sounding the alarm on DrugBanklinew license and terms of use 2016, Edge dropout contamination in hetnet edge prediction 2016, Hosting Hetionet in the cloud: creating a public Neo4j instance 2016,	2 2 2

62	Decomposing the DWPC to assess intermediate node or edge contributions 2016,		2
61	Decomposing predictions into their network support 2016 ,		2
60	Genetic Association-Guided Analysis of Gene Networks for the Study of Complex Traits. <i>Circulation: Cardiovascular Genetics</i> , 2016 , 9, 179-84		7
59	Meta-analysis of genome-wide association studies reveals genetic overlap between Hodgkin lymphoma and multiple sclerosis. <i>International Journal of Epidemiology</i> , 2016 , 45, 728-40	7.8	16
58	Understanding multicellular function and disease with human tissue-specific networks. <i>Nature Genetics</i> , 2015 , 47, 569-76	36.3	473
57	iCTNet2: integrating heterogeneous biological interactions to understand complex traits. <i>F1000Research</i> , 2015 , 4, 485	3.6	7
56	iCTNet2: integrating heterogeneous biological interactions to understand complex traits. <i>F1000Research</i> , 2015 , 4, 485	3.6	6
55	Heterogeneous Network Edge Prediction: A Data Integration Approach to Prioritize Disease-Associated Genes. <i>PLoS Computational Biology</i> , 2015 , 11, e1004259	5	69
54	One network to rule them all 2015 ,		2
53	Renaming fleterogeneous networksIto a more concise and catchy term 2015,		2
52	Processing DisGeNET for disease-gene relationships 2015,		2
51	Processing the DISEASES resource for diseasegene relationships 2015 ,		2
50	Integrating resources with disparate licensing into an open network 2015,		4
49	MSigDB licensing 2015 ,		2
48	LINCS L1000 licensing 2015 ,		2
47	Incomplete Interactome licensing 2015,		2
46	Using the neo4j graph database for hetnets 2015 ,		2
45	Assessing the informativeness of features 2015 ,		2

(2015-2015)

44	Processing Bgee for tissue-specific gene presence and over/under-expression 2015,	2
43	Permuting hetnets and implementing randomized edge swaps in cypher 2015,	2
42	How should we construct a catalog of drug indications? 2015,	2
41	Enabling reproducibility and reuse 2015,	2
40	Using Entrez Gene as our gene vocabulary 2015 ,	2
39	Compiling Gene Ontology annotations into an easy-to-use format 2015 ,	2
38	Unifying drug vocabularies 2015 ,	2
37	Tissue Node 2015 ,	2
36	Computing consensus transcriptional profiles for LINCS L1000 perturbations 2015,	2
35	Unifying disease vocabularies 2015 ,	2
34	Processing LabeledIn to extract indications 2015,	2
33	Integrating drug target information from BindingDB 2015 ,	2
32	Selecting informative ERC (evolutionary rate covariation) values between genes 2015,	2
31	Extracting indications from the ehrlink resource 2015 ,	2
30	Protein (target, carrier, transporter, and enzyme) interactions in DrugBank 2015,	2
29	Mining knowledge from MEDLINE articles and their indexed MeSH terms 2015 ,	5
28	Disease Ontology feature requests 2015 ,	2
27	Calculating molecular similarities between DrugBank compounds 2015,	2

26	Calculating genomic windows for GWAS lead SNPs 2015 ,		2
25	Adding pathway resources to your network 2015 ,		2
24	Extracting disease-gene associations from the GWAS Catalog 2015,		5
23	Tissue-specific gene expression resources 2015 ,		2
22	Creating a catalog of protein interactions 2015 ,		2
21	The TISSUES resource for the tissue-specificity of genes 2015 ,		2
20	Disease similarity from MEDLINE topic cooccurrence 2015 ,		2
19	Functional disease annotations for genes using DOAF 2015 ,		2
18	Expert curation of our indication catalog for disease-modifying treatments 2015,		2
17	STARGEO: expression signatures for disease using crowdsourced GEO annotation 2015,		2
16	Extracting side effects from SIDER 4 2015 ,		2
15	Lung cancer incidence decreases with elevation: evidence for oxygen as an inhaled carcinogen. <i>PeerJ</i> , 2015 , 3, e705	3.1	19
14	Evolving hard problems: Generating human genetics datasets with a complex etiology. <i>BioData Mining</i> , 2011 , 4, 21	4.3	12
13	Multifactor dimensionality reduction for graphics processing units enables genome-wide testing of epistasis in sporadic ALS. <i>Bioinformatics</i> , 2010 , 26, 694-5	7.2	60
12	Enabling personal genomics with an explicit test of epistasis. <i>Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing</i> , 2010 , 327-36	1.3	31
11	The Informative Extremes: Using Both Nearest and Farthest Individuals Can Improve Relief Algorithms in the Domain of Human Genetics. <i>Lecture Notes in Computer Science</i> , 2010 , 182-193	0.9	16
10	A Model Free Method to Generate Human Genetics Datasets with Complex Gene-Disease Relationships. <i>Lecture Notes in Computer Science</i> , 2010 , 74-85	0.9	2
9	Rephetio: Repurposing drugs on a hetnet [project]		3

LIST OF PUBLICATIONS

8	Rephetio: Repurposing drugs on a hetnet [proposal]	2
7	Sci-Hub provides access to nearly all scholarly literature	10
6	Sci-Hub provides access to nearly all scholarly literature	5
5	Systematic integration of biomedical knowledge prioritizes drugs for repurposing	1
4	Opportunities and obstacles for deep learning in biology and medicine	45
3	Analysis of ISCB honorees and keynotes reveals disparities	7
2	Sequential compression of gene expression across dimensionalities and methods reveals no single best method or dimensionality	3
1	Expanding a Database-derived Biomedical Knowledge Graph via Multi-relation Extraction from Biomedical Abstracts	1