## Zhiyong Lu

## List of Publications by Year in descending order

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172457 265206 4,792 44 29 42 citations h-index g-index papers 45 45 45 3998 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	NCBI disease corpus: A resource for disease name recognition and concept normalization. Journal of Biomedical Informatics, 2014, 47, 1-10.	4.3	525
2	PubTator: a web-based text mining tool for assisting biocuration. Nucleic Acids Research, 2013, 41, W518-W522.	14.5	523
3	DNorm: disease name normalization with pairwise learning to rank. Bioinformatics, 2013, 29, 2909-2917.	4.1	436
4	BioCreative V CDR task corpus: a resource for chemical disease relation extraction. Database: the Journal of Biological Databases and Curation, 2016, 2016, baw068.	3.0	350
5	PubTator central: automated concept annotation for biomedical full text articles. Nucleic Acids Research, 2019, 47, W587-W593.	14.5	248
6	Overview of BioCreative II gene normalization. Genome Biology, 2008, 9, S3.	9.6	237
7	TaggerOne: joint named entity recognition and normalization with semi-Markov Models. Bioinformatics, 2016, 32, 2839-2846.	4.1	221
8	tmVar: a text mining approach for extracting sequence variants in biomedical literature. Bioinformatics, 2013, 29, 1433-1439.	4.1	197
9	LitCovid: an open database of COVID-19 literature. Nucleic Acids Research, 2021, 49, D1534-D1540.	14.5	189
10	The CHEMDNER corpus of chemicals and drugs and its annotation principles. Journal of Cheminformatics, 2015, 7, S2.	6.1	166
11	GNormPlus: An Integrative Approach for Tagging Genes, Gene Families, and Protein Domains. BioMed Research International, 2015, 2015, 1-7.	1.9	155
12	BioC: a minimalist approach to interoperability for biomedical text processing. Database: the Journal of Biological Databases and Curation, 2013, 2013, bat064-bat064.	3.0	123
13	Assessing the state of the art in biomedical relation extraction: overview of the BioCreative V chemical-disease relation (CDR) task. Database: the Journal of Biological Databases and Curation, 2016, 2016, .	3.0	123
14	The gene normalization task in BioCreative III. BMC Bioinformatics, 2011, 12, S2.	2.6	101
15	LitVar: a semantic search engine for linking genomic variant data in PubMed and PMC. Nucleic Acids Research, 2018, 46, W530-W536.	14.5	96
16	On expert curation and scalability: UniProtKB/Swiss-Prot as a case study. Bioinformatics, 2017, 33, 3454-3460.	4.1	91
17	Best Match: New relevance search for PubMed. PLoS Biology, 2018, 16, e2005343.	5.6	90
18	Towards PubMed 2.0. ELife, 2017, 6, .	6.0	86

#	Article	IF	CITATIONS
19	Text Mining Genotype-Phenotype Relationships from Biomedical Literature for Database Curation and Precision Medicine. PLoS Computational Biology, 2016, 12, e1005017.	3.2	81
20	tmVar 2.0: integrating genomic variant information from literature with dbSNP and ClinVar for precision medicine. Bioinformatics, 2018, 34, 80-87.	4.1	79
21	SR4GN: A Species Recognition Software Tool for Gene Normalization. PLoS ONE, 2012, 7, e38460.	2.5	71
22	Biocuration workflows and text mining: overview of the BioCreative 2012 Workshop Track II. Database: the Journal of Biological Databases and Curation, 2012, 2012, bas043-bas043.	3.0	67
23	PMC text mining subset in BioC: about three million full-text articles and growing. Bioinformatics, 2019, 35, 3533-3535.	4.1	51
24	How user intelligence is improving PubMed. Nature Biotechnology, 2018, 36, 937-945.	17.5	46
25	BioConceptVec: Creating and evaluating literature-based biomedical concept embeddings on a large scale. PLoS Computational Biology, 2020, 16, e1007617.	3.2	43
26	Artificial Intelligence in Action: Addressing the COVID-19 Pandemic with Natural Language Processing. Annual Review of Biomedical Data Science, 2021, 4, 313-339.	6.5	38
27	LitSense: making sense of biomedical literature at sentence level. Nucleic Acids Research, 2019, 47, W594-W599.	14.5	37
28	Hybrid curation of gene–mutation relations combining automated extraction and crowdsourcing. Database: the Journal of Biological Databases and Curation, 2014, 2014, .	3.0	35
29	TeamTat: a collaborative text annotation tool. Nucleic Acids Research, 2020, 48, W5-W11.	14.5	34
30	Scaling up data curation using deep learning: An application to literature triage in genomic variation resources. PLoS Computational Biology, 2018, 14, e1006390.	3.2	33
31	Accessing Biomedical Literature in the Current Information Landscape. Methods in Molecular Biology, 2014, 1159, 11-31.	0.9	32
32	Pressing needs of biomedical text mining in biocuration and beyond: opportunities and challenges. Database: the Journal of Biological Databases and Curation, 2016, 2016, baw161.	3.0	30
33	LitSuggest: a web-based system for literature recommendation and curation using machine learning. Nucleic Acids Research, 2021, 49, W352-W358.	14.5	28
34	ezTag: tagging biomedical concepts via interactive learning. Nucleic Acids Research, 2018, 46, W523-W529.	14.5	27
35	NLM-Chem, a new resource for chemical entity recognition in PubMed full text literature. Scientific Data, 2021, 8, 91.	5.3	26
36	SimConcept: A Hybrid Approach for Simplifying Composite Named Entities in Biomedical Text. IEEE Journal of Biomedical and Health Informatics, 2015, 19, 1385-1391.	6.3	18

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37	Deep learning with sentence embeddings pre-trained on biomedical corpora improves the performance of finding similar sentences in electronic medical records. BMC Medical Informatics and Decision Making, 2020, 20, 73.	3.0	15
38	Recent advances of automated methods for searching and extracting genomic variant information from biomedical literature. Briefings in Bioinformatics, $2021, 22, \ldots$	6.5	13
39	Ten tips for a text-mining-ready article: How to improve automated discoverability and interpretability. PLoS Biology, 2020, 18, e3000716.	5.6	10
40	NLM-Gene, a richly annotated gold standard dataset for gene entities that addresses ambiguity and multi-species gene recognition. Journal of Biomedical Informatics, 2021, 118, 103779.	4.3	9
41	MeSH-based dataset for measuring the relevance of text retrieval. , 2018, , .		5
42	Benchmarking Effectiveness and Efficiency of Deep Learning Models for Semantic Textual Similarity in the Clinical Domain: Validation Study. JMIR Medical Informatics, 2021, 9, e27386.	2.6	5
43	Tracking human genes along the translational continuum. Npj Genomic Medicine, 2019, 4, 25.	3.8	2
44	Editor's introduction to the special section on the 7th Biomedical Linked Annotation Hackathon (BLAH7). Genomics and Informatics, 2021, 19, e20.	0.8	0