## Laura I Furlong

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6885259/publications.pdf

Version: 2024-02-01

76 papers 6,630 citations

30 h-index 70 g-index

86 all docs

86 docs citations

86 times ranked 10607 citing authors

#	Article	IF	CITATIONS
1	DisGeNET: a comprehensive platform integrating information on human disease-associated genes and variants. Nucleic Acids Research, 2017, 45, D833-D839.	6.5	1,865
2	The DisGeNET knowledge platform for disease genomics: 2019 update. Nucleic Acids Research, 2020, 48, D845-D855.	6.5	1,083
3	DisGeNET: a discovery platform for the dynamical exploration of human diseases and their genes. Database: the Journal of Biological Databases and Curation, 2015, 2015, bav028-bav028.	1.4	847
4	The DisGeNET cytoscape app: Exploring and visualizing disease genomics data. Computational and Structural Biotechnology Journal, 2021, 19, 2960-2967.	1.9	221
5	Human diseases through the lens of network biology. Trends in Genetics, 2013, 29, 150-159.	2.9	182
6	DisGeNET: a Cytoscape plugin to visualize, integrate, search and analyze gene–disease networks. Bioinformatics, 2010, 26, 2924-2926.	1.8	180
7	Pathway databases and tools for their exploitation: benefits, current limitations and challenges. Molecular Systems Biology, 2009, 5, 290.	3.2	173
8	Extraction of relations between genes and diseases from text and large-scale data analysis: implications for translational research. BMC Bioinformatics, 2015, 16, 55.	1.2	170
9	Gene-Disease Network Analysis Reveals Functional Modules in Mendelian, Complex and Environmental Diseases. PLoS ONE, 2011, 6, e20284.	1.1	153
10	The Semanticscience Integrated Ontology (SIO) for biomedical research and knowledge discovery. Journal of Biomedical Semantics, 2014, 5, 14.	0.9	138
11	The EU-ADR corpus: Annotated drugs, diseases, targets, and their relationships. Journal of Biomedical Informatics, 2012, 45, 879-884.	2.5	99
12	COVID-19 Disease Map, building a computational repository of SARS-CoV-2 virus-host interaction mechanisms. Scientific Data, 2020, 7, 136.	2.4	99
13	PsyGeNET: a knowledge platform on psychiatric disorders and their genes. Bioinformatics, 2015, 31, 3075-3077.	1.8	79
14	Detecting Signs of Depression in Tweets in Spanish: Behavioral and Linguistic Analysis. Journal of Medical Internet Research, 2019, 21, e14199.	2.1	66
15	Personalized Respiratory Medicine: Exploring the Horizon, Addressing the Issues. Summary of a BRN-AJRCCM Workshop Held in Barcelona on June 12, 2014. American Journal of Respiratory and Critical Care Medicine, 2015, 191, 391-401.	2.5	61
16	Identifying temporal patterns in patient disease trajectories using dynamic time warping: A population-based study. Scientific Reports, 2018, 8, 4216.	1.6	61
17	DisGeNET-RDF: harnessing the innovative power of the Semantic Web to explore the genetic basis of diseases. Bioinformatics, 2016, 32, 2236-2238.	1.8	52
18	Network medicine analysis of COPD multimorbidities. Respiratory Research, 2014, 15, 111.	1.4	48

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19	Improving data and knowledge management to better integrate health care and research. Journal of Internal Medicine, 2013, 274, 321-328.	2.7	44
20	A Knowledge-Driven Approach to Extract Disease-Related Biomarkers from the Literature. BioMed Research International, 2014, 2014, 1-11.	0.9	42
21	Drug-Induced Acute Myocardial Infarction: Identifying â€~Prime Suspects' from Electronic Healthcare Records-Based Surveillance System. PLoS ONE, 2013, 8, e72148.	1.1	41
22	Assessment of NER solutions against the first and second CALBC Silver Standard Corpus. Journal of Biomedical Semantics, 2011, 2, S11.	0.9	39
23	The EUâ€ADR Web Platform: delivering advanced pharmacovigilance tools. Pharmacoepidemiology and Drug Safety, 2013, 22, 459-467.	0.9	36
24	Automatic Filtering and Substantiation of Drug Safety Signals. PLoS Computational Biology, 2012, 8, e1002457.	1.5	34
25	Challenges in the association of human single nucleotide polymorphism mentions with unique database identifiers. BMC Bioinformatics, 2011, 12, S4.	1.2	33
26	Molecular and clinical diseasome of comorbidities in exacerbated COPD patients. European Respiratory Journal, 2015, 46, 1001-1010.	3.1	32
27	Proximal Pathway Enrichment Analysis for Targeting Comorbid Diseases via Network Endopharmacology. Pharmaceuticals, 2018, 11, 61.	1.7	32
28	GUILDify v2.0: A Tool to Identify Molecular Networks Underlying Human Diseases, Their Comorbidities and Their Druggable Targets. Journal of Molecular Biology, 2019, 431, 2477-2484.	2.0	32
29	Evaluation of the proacrosin/acrosin system and its mechanism of activation in human sperm extracts. Journal of Reproductive Immunology, 2002, 54, 43-63.	0.8	31
30	OSIRISv1.2: A named entity recognition system for sequence variants of genes in biomedical literature. BMC Bioinformatics, 2008, 9, 84.	1.2	31
31	Expression of epithelial cadherin in the human male reproductive tract and gametes and evidence of its participation in fertilization. Molecular Human Reproduction, 2008, 14, 561-571.	1.3	31
32	comoRbidity: an R package for the systematic analysis of disease comorbidities. Bioinformatics, 2018, 34, 3228-3230.	1.8	31
33	In silico models in drug development: where we are. Current Opinion in Pharmacology, 2018, 42, 111-121.	1.7	30
34	Uncovering disease mechanisms through network biology in the era of Next Generation Sequencing. Scientific Reports, 2016, 6, 24570.	1.6	29
35	Genetic and functional characterization of disease associations explains comorbidity. Scientific Reports, 2017, 7, 6207.	1.6	28
36	Genetic and Real-World Clinical Data, Combined with Empirical Validation, Nominate Jak-Stat Signaling as a Target for Alzheimer's Disease Therapeutic Development. Cells, 2019, 8, 425.	1.8	27

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37	Binding of recombinant human proacrosin/acrosin to zona pellucida (ZP) glycoproteins. I. Studies with recombinant human ZPA, ZPB, and ZPC. Fertility and Sterility, 2005, 83, 1780-1790.	0.5	25
38	From SNPs to pathways: integration of functional effect of sequence variations on models of cell signalling pathways. BMC Bioinformatics, 2009, 10, S6.	1.2	24
39	OSIRIS: a tool for retrieving literature about sequence variants. Bioinformatics, 2006, 22, 2567-2569.	1.8	22
40	Expression of Human Proacrosin in Escherichia coli and Binding to Zona Pellucida1. Biology of Reproduction, 2000, 62, 606-615.	1.2	21
41	Acrosin antibodies and infertility. I. Detection of antibodies towards proacrosin/acrosin in women consulting for infertility and evaluation of their effects upon the sperm protease activities. Fertility and Sterility, 2009, 91, 1245-1255.	0.5	21
42	Nanopublications: A Growing Resource of Provenance-Centric Scientific Linked Data. , 2018, , .		21
43	Knowledge environments representing molecular entities for the virtual physiological human. Philosophical Transactions Series A, Mathematical, Physical, and Engineering Sciences, 2008, 366, 3091-3110.	1.6	19
44	Publishing DisGeNET as nanopublications. Semantic Web, 2016, 7, 519-528.	1.1	18
45	Network, Transcriptomic and Genomic Features Differentiate Genes Relevant for Drug Response. Frontiers in Genetics, 2018, 9, 412.	1.1	18
46	Proyecto de biomarcadores y perfiles clÃnicos personalizados en la enfermedad pulmonar obstructiva crónica (proyecto BIOMEPOC). Archivos De Bronconeumologia, 2019, 55, 93-99.	0.4	18
47	Comorbidity between Alzheimer's disease and major depression: a behavioural and transcriptomic characterization study in mice. Alzheimer's Research and Therapy, 2021, 13, 73.	3.0	18
48	The eTRANSAFE Project on Translational Safety Assessment through Integrative Knowledge Management: Achievements and Perspectives. Pharmaceuticals, 2021, 14, 237.	1.7	17
49	IDENTIFYING GENE-SPECIFIC VARIATIONS IN BIOMEDICAL TEXT. Journal of Bioinformatics and Computational Biology, 2007, 05, 1277-1296.	0.3	15
50	Gathering and Exploring Scientific Knowledge in Pharmacovigilance. PLoS ONE, 2013, 8, e83016.	1.1	15
51	Combining machine learning, crowdsourcing and expert knowledge to detect chemical-induced diseases in text. Database: the Journal of Biological Databases and Curation, 2016, 2016, baw094.	1.4	14
52	Quantitative Systems Toxicology Modeling To Address Key Safety Questions in Drug Development: A Focus of the TransQST Consortium. Chemical Research in Toxicology, 2020, 33, 7-9.	1.7	14
53	Anti-human proacrosin antibody inhibits the zona pellucida (ZP)–induced acrosome reaction of ZP-bound spermatozoa. Fertility and Sterility, 2010, 93, 2456-2459.	0.5	13
54	Text mining and expert curation to develop a database on psychiatric diseases and their genes. Database: the Journal of Biological Databases and Curation, 2017, 2017, .	1.4	11

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55	Pancreatic cancer and autoimmune diseases: An association sustained by computational and epidemiological case–control approaches. International Journal of Cancer, 2019, 144, 1540-1549.	2.3	11
56	An ensemble learning approach for modeling the systems biology of drug-induced injury. Biology Direct, 2021, 16, 5.	1.9	11
57	ResMarkerDB: a database of biomarkers of response to antibody therapy in breast and colorectal cancer. Database: the Journal of Biological Databases and Curation, 2019, 2019, .	1.4	10
58	Reliable Granular References to Changing Linked Data. Lecture Notes in Computer Science, 2017, , 436-451.	1.0	10
59	A crowdsourcing workflow for extracting chemical-induced disease relations from free text. Database: the Journal of Biological Databases and Curation, 2016, 2016, baw051.	1.4	9
60	Comorbidity4j: a tool for interactive analysis of disease comorbidities over large patient datasets. Bioinformatics, 2019, 35, 3530-3532.	1.8	9
61	Binding of recombinant human proacrosin/acrosin to zona pellucida glycoproteins. II. Participation of mannose residues in the interaction. Fertility and Sterility, 2005, 83, 1791-1796.	0.5	8
62	A system-level analysis of patient disease trajectories based on clinical, phenotypic and molecular similarities. Bioinformatics, 2021, 37, 1435-1443.	1.8	8
63	Antiacrosin antibodies and infertility. II. Gene immunization with human proacrosin to assess the effect of immunity toward proacrosin/acrosin upon protein activities and animal fertility. Fertility and Sterility, 2009, 91, 1256-1268.	0.5	7
64	Mining the Modular Structure of Protein Interaction Networks. PLoS ONE, 2015, 10, e0122477.	1.1	7
65	CDH1/E-cadherin and solid tumors. An updated gene-disease association analysis using bioinformatics tools. Computational Biology and Chemistry, 2016, 60, 9-20.	1.1	7
66	The BIOMEPOC Project: Personalized Biomarkers and Clinical Profiles in Chronic Obstructive Pulmonary Disease. Archivos De Bronconeumologia, 2019, 55, 93-99.	0.4	5
67	The ELIXIR Human Copy Number Variations Community: building bioinformatics infrastructure for research. F1000Research, 2020, 9, 1229.	0.8	5
68	Rcupcake: an R package for querying and analyzing biomedical data through the BD2K PIC-SURE RESTful API. Bioinformatics, 2018, 34, 1431-1432.	1.8	4
69	Digging for knowledge with information extraction. , 2010, , .		3
70	psygenet2r: a R/Bioconductor package for the analysis of psychiatric disease genes. Bioinformatics, 2017, 33, 4004-4006.	1.8	3
71	Exploring the Association of Cancer and Depression in Electronic Health Records: Combining Encoded Diagnosis and Mining Free-Text Clinical Notes. JMIR Cancer, 2022, 8, e39003.	0.9	1
72	Functional Genomics Analysis to Disentangle the Role of Genetic Variants in Major Depression. Genes, 2022, 13, 1259.	1.0	1

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73	Embracing the Dark Side: Computational Approaches to Unveil the Functionality of Genes Lacking Biological Annotation in Drug-Induced Liver Injury. Frontiers in Genetics, 2018, 9, 527.	1.1	О
74	Identification of Sequence Variants of Genes from Biomedical Literature., 2009,, 289-300.		0
75	Abstract 1085: A bioinformatics approach to evaluate the involvement of CDH1/E-cadherin in solid tumors and to identify breast cancer biomarkers. , 2015, , .		O
76	Evaluating Behavioral and Linguistic Changes During Drug Treatment for Depression Using Tweets in Spanish: Pairwise Comparison Study. Journal of Medical Internet Research, 2020, 22, e20920.	2.1	0