## Jingwen Yan

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1948345/publications.pdf

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63	1,108	17 h-index	31
papers	citations		g-index
69	69	69	1423
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Network approaches to systems biology analysis of complex disease: integrative methods for multi-omics data. Briefings in Bioinformatics, 2018, 19, 1370-1381.	6.5	185
2	Metabolic Network Analysis Reveals Altered Bile Acid Synthesis and Metabolism in Alzheimer's Disease. Cell Reports Medicine, 2020, 1, 100138.	6.5	102
3	Structured sparse canonical correlation analysis for brain imaging genetics: an improved GraphNet method. Bioinformatics, 2016, 32, 1544-1551.	4.1	96
4	Transcriptome-guided amyloid imaging genetic analysis via a novel structured sparse learning algorithm. Bioinformatics, 2014, 30, i564-i571.	4.1	57
5	Progress in Polygenic Composite Scores in Alzheimer's and Other Complex Diseases. Trends in Genetics, 2019, 35, 371-382.	6.7	52
6	Mining Outcome-relevant Brain Imaging Genetic Associations via Three-way Sparse Canonical Correlation Analysis in Alzheimer's Disease. Scientific Reports, 2017, 7, 44272.	3.3	44
7	Cortical surface biomarkers for predicting cognitive outcomes using group l2,1 norm. Neurobiology of Aging, 2015, 36, S185-S193.	3.1	43
8	Identification of associations between genotypes and longitudinal phenotypes via temporally-constrained group sparse canonical correlation analysis. Bioinformatics, 2017, 33, i341-i349.	4.1	42
9	Deep Fusion of Brain Structure-Function in Mild Cognitive Impairment. Medical Image Analysis, 2021, 72, 102082.	11.6	37
10	A Novel Structure-Aware Sparse Learning Algorithm for Brain Imaging Genetics. Lecture Notes in Computer Science, 2014, 17, 329-336.	1.3	36
11	Genome-wide association and interaction studies of CSF T-tau/A $\hat{l}^2$ 42 ratio in ADNI cohort. Neurobiology of Aging, 2017, 57, 247.e1-247.e8.	3.1	34
12	A novel SCCA approach via truncated $\langle i \rangle \langle b \rangle \hat{a}$ ," $\langle b \rangle \langle i \rangle 1$ -norm and truncated group lasso for brain imaging genetics. Bioinformatics, 2018, 34, 278-285.	4.1	31
13	Network-based analysis of genetic variants associated with hippocampal volume in Alzheimer's disease: a study of ADNI cohorts. BioData Mining, 2016, 9, 3.	4.0	28
14	Identifying Multimodal Intermediate Phenotypes Between Genetic Risk Factors and Disease Status in Alzheimer's Disease. Neuroinformatics, 2016, 14, 439-452.	2.8	26
15	Genetic Interactions Explain Variance in Cingulate Amyloid Burden: An AV-45 PET Genome-Wide Association and Interaction Study in the ADNI Cohort. BioMed Research International, 2015, 2015, 1-11.	1.9	24
16	Hippocampal transcriptome-guided genetic analysis of correlated episodic memory phenotypes in Alzheimer's disease. Frontiers in Genetics, 2015, 6, 117.	2.3	23
17	Tissue-specific network-based genome wide study of amygdala imaging phenotypes to identify functional interaction modules. Bioinformatics, 2017, 33, 3250-3257.	4.1	23
18	Regional imaging genetic enrichment analysis. Bioinformatics, 2020, 36, 2554-2560.	4.1	16

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19	Global Brain Functional Network Connectivity in Infants With Prenatal Opioid Exposure. Frontiers in Pediatrics, 2022, 10, 847037.	1.9	15
20	IDENTIFICATION OF DISCRIMINATIVE IMAGING PROTEOMICS ASSOCIATIONS IN ALZHEIMER'S DISEASE VIA A NOVEL SPARSE CORRELATION MODEL. , 2017, 22, 94-104.		14
21	GN-SCCA: GraphNet Based Sparse Canonical Correlation Analysis for Brain Imaging Genetics. Lecture Notes in Computer Science, 2015, 9250, 275-284.	1.3	14
22	Genome-wide network-based pathway analysis of CSF t-tau/A $\hat{l}^2$ 1-42 ratio in the ADNI cohort. BMC Genomics, 2017, 18, 421.	2.8	13
23	Two-dimensional enrichment analysis for mining high-level imaging genetic associations. Brain Informatics, 2017, 4, 27-37.	3.0	13
24	Identifying Associations Between Brain Imaging Phenotypes and Genetic Factors via a Novel Structured SCCA Approach. Lecture Notes in Computer Science, 2017, 10265, 543-555.	1.3	12
25	Quantitative trait loci identification for brain endophenotypes via new additive model with random networks. Bioinformatics, 2018, 34, i866-i874.	4.1	11
26	Genome-wide Network-assisted Association and Enrichment Study of Amyloid Imaging Phenotype in Alzheimer's Disease. Current Alzheimer Research, 2020, 16, 1163-1174.	1.4	11
27	Multivariate genome wide association and network analysis of subcortical imaging phenotypes in Alzheimer's disease. BMC Genomics, 2020, 21, 896.	2.8	11
28	Structured sparse CCA for brain imaging genetics via graph OSCAR. BMC Systems Biology, 2016, 10, 68.	3.0	9
29	Pattern Discovery in Brain Imaging Genetics via SCCA Modeling with a Generic Non-convex Penalty. Scientific Reports, 2017, 7, 14052.	3.3	9
30	Integrative-omics for discovery of network-level disease biomarkers: a case study in Alzheimer's disease. Briefings in Bioinformatics, 2021, 22, .	6.5	8
31	Heritability Estimation of Reliable Connectomic Features. Lecture Notes in Computer Science, 2018, 11083, 58-66.	1.3	8
32	Longitudinal Genotype-Phenotype Association Study via Temporal Structure Auto-learning Predictive Model. Lecture Notes in Computer Science, 2017, 10229, 287-302.	1.3	8
33	Data synthesis and method evaluation for brain imaging genetics. , 2014, 2014, 1202-1205.		6
34	Sparse Canonical Correlation Analysis via truncated â,, " <inf>1</inf> -norm with application to brain imaging genetics., 2016, 2016, 707-711.		6
35	Longitudinal Genotype–Phenotype Association Study through Temporal Structure Auto-Learning Predictive Model. Journal of Computational Biology, 2018, 25, 809-824.	1.6	6
36	Identification of functionally connected multi-omic biomarkers for Alzheimer's disease using modularity-constrained Lasso. PLoS ONE, 2020, 15, e0234748.	2.5	6

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37	Network-Guided Sparse Learning for Predicting Cognitive Outcomes from MRI Measures. Lecture Notes in Computer Science, 2013, 8159, 202-210.	1.3	5
38	Transcriptome-Guided Imaging Genetic Analysis via a Novel Sparse CCA Algorithm. Lecture Notes in Computer Science, 2017, 10551, 220-229.	1.3	5
39	DIAGNOSIS-GUIDED METHOD FOR IDENTIFYING MULTI-MODALITY NEUROIMAGING BIOMARKERS ASSOCIATED WITH GENETIC RISK FACTORS IN ALZHEIMER'S DISEASE. , 2016, , .		4
40	Differential co-expression analysis reveals early stage transcriptomic decoupling in alzheimer's disease. BMC Medical Genomics, 2020, 13, 53.	1.5	4
41	Predicting Interrelated Alzheimer's Disease Outcomes via New Self-learned Structured Low-Rank Model. Lecture Notes in Computer Science, 2017, 10265, 198-209.	1.3	4
42	MoNET: an R package for multi-omic network analysis. Bioinformatics, 2021, , .	4.1	2
43	Integrative analysis of eQTL and GWAS summary statistics reveals transcriptomic alteration in Alzheimer brains. BMC Medical Genomics, 2022, 15, 93.	1.5	2
44	Joint identification of imaging and proteomics biomarkers of Alzheimer's disease using network-guided sparse learning., 2014, 2014, 665-668.		1
45	Network-based genome wide study of hippocampal imaging phenotype in Alzheimer's Disease to identify functional interaction modules., 2017, 2017, 6170-6174.		1
46	IC-O1-03: Hippocampal transcriptome-guided gene-gene interaction of memory phenotype in MCI and Alzheimer's disease., 2013, 9, P4-P4.		0
47	IC-P-172: GENOME-WIDE PROTEIN INTERACTION GUIDED EPISTATIC ANALYSIS ON MEMORY PERFORMANCE: AN ADNI STUDY. , 2014, 10, P95-P96.		O
48	IC-P-173: EFFECTS OF NEWLY IDENTIFIED TOP AD CANDIDATE GENES ON MEMORY PERFORMANCE: SNP, GENE, AND EPISTASIS ANALYSES IN ADNI. , 2014, 10, P96-P97.		0
49	P1-230: EFFECTS OF NEWLY IDENTIFIED TOP AD CANDIDATE GENES ON MEMORY PERFORMANCE: SNP, GENE, AND EPISTASIS ANALYSES IN ADNI. , 2014, 10, P388-P388.		O
50	P1-213: GENOME-WIDE PROTEIN INTERACTION-GUIDED EPISTATIC ANALYSIS ON MEMORY PERFORMANCE: AN ADNI STUDY. , 2014, 10, P381-P382.		0
51	P4-002: Genome-wide network-based pathway analysis of CSF biomarker t-tau in the ADNI cohort. , 2015, 11, P765-P765.		O
52	P1-009: The nav2 (neuron navigator 2) gene as a common genetic influence across correlated episodic memory performances., 2015, 11, P339-P340.		0
53	[P2–120]: INVESTIGATION OF GENETIC INFLUENCES ON ATROPHY RATE DURING THE MCI DISEASE STAGE USING A BOOTSTRAPâ€ENHANCED SPARSE ASSOCIATION MODEL. Alzheimer's and Dementia, 2017, 13, P653.	0.8	O
54	[P2–220]: GENETIC FINDINGS USING ADNI MULTIMODAL QUANTITATIVE PHENOTYPES: A 2016 UPDATE. Alzheimer's and Dementia, 2017, 13, P694.	0.8	О

#	Article	IF	Citations
55	[F1–02–04]: INTEGRATING MULTIâ€MODALITY IMAGING AND MULTIâ€LAYER â€OMICS TO ADVANCE THE S BIOLOGY OF ALZHEIMER's DISEASE. Alzheimer's and Dementia, 2017, 13, P175.	YSŢĘMS	0
56	P2â€253: <i>EP300</i> IS ASSOCIATED WITH ALTERED BILE ACIDS IN ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2018, 14, P772.	0.8	0
57	ICâ€Pâ€075: GENETIC FINDINGS USING ADNI MULTIMODAL QUANTITATIVE PHENOTYPES: A 2017 UPDATE. Alzheimer's and Dementia, 2018, 14, P66.	0.8	0
58	Endophenotype driven polygenic risk scores for Alzheimer's disease. Alzheimer's and Dementia, 2020, 16, e046766.	0.8	0
59	Genetic Influence Underlying Brain Connectivity Phenotype: A Study on Two Age-Specific Cohorts. Frontiers in Genetics, 2021, 12, 782953.	2.3	0
60	Biomarkerâ€based polygenic risk scores for profiling genetic susceptibility in Alzheimer's disease. Alzheimer's and Dementia, 2021, 17, .	0.8	0
61	Integrative analysis of eQTL and GWAS summary statistics reveals novel genes related to Alzheimer's disease. Alzheimer's and Dementia, 2021, 17, .	0.8	0
62	Integrative â€omics for discovery of networkâ€level disease biomarkers for Alzheimer's disease. Alzheimer's and Dementia, 2021, 17, .	0.8	0
63	Gene co-expression changes underlying the functional connectomic alterations in Alzheimer's disease. BMC Medical Genomics, 2022, 15, 92.	1.5	0