

# Yang You

## List of Publications by Year in descending order

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Version: 2024-02-01

15  
papers

577  
citations

933447

10  
h-index

996975

15  
g-index

16  
all docs

16  
docs citations

16  
times ranked

893  
citing authors

#	ARTICLE	IF	CITATIONS
1	Activated human astrocyte-derived extracellular vesicles modulate neuronal uptake, differentiation and firing. <i>Journal of Extracellular Vesicles</i> , 2020, 9, 1706801.	12.2	116
2	Proteomic and biological profiling of extracellular vesicles from Alzheimer's disease human brain tissues. <i>Alzheimer's and Dementia</i> , 2020, 16, 896-907.	0.8	105
3	Emerging roles of extracellular vesicles in neurodegenerative disorders. <i>Neurobiology of Disease</i> , 2019, 130, 104512.	4.4	78
4	Schizophrenia Related Variants in CACNA1C also Confer Risk of Autism. <i>PLoS ONE</i> , 2015, 10, e0133247.	2.5	55
5	Human neural cell type-specific extracellular vesicle proteome defines disease-related molecules associated with activated astrocytes in Alzheimer's disease brain. <i>Journal of Extracellular Vesicles</i> , 2022, 11, e12183.	12.2	54
6	Inhibition of colony stimulating factor 1 receptor corrects maternal inflammation-induced microglial and synaptic dysfunction and behavioral abnormalities. <i>Molecular Psychiatry</i> , 2021, 26, 1808-1831.	7.9	44
7	Growth arrest specific gene 7 is associated with schizophrenia and regulates neuronal migration and morphogenesis. <i>Molecular Brain</i> , 2016, 9, 54.	2.6	23
8	The Schizophrenia Susceptibility Gene OPCML Regulates Spine Maturation and Cognitive Behaviors through Eph-Cofilin Signaling. <i>Cell Reports</i> , 2019, 29, 49-61.e7.	6.4	20
9	Genetic Evidence for Possible Involvement of the Calcium Channel Gene CACNA1A in Autism Pathogenesis in Chinese Han Population. <i>PLoS ONE</i> , 2015, 10, e0142887.	2.5	18
10	Protein phosphatase 2A and complement component 4 are linked to the protective effect of <i>APOE</i> $\epsilon$ 2 for Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2022, 18, 2042-2054.	0.8	18
11	<i>Auts2</i> deletion involves in DG hypoplasia and social recognition deficit: The developmental and neural circuit mechanisms. <i>Science Advances</i> , 2022, 8, eabk1238.	10.3	14
12	Chromatin remodeling gene EZH2 involved in the genetic etiology of autism in Chinese Han population. <i>Neuroscience Letters</i> , 2016, 610, 182-186.	2.1	12
13	Genetic variants in the transcription regulatory region of MEGF10 are associated with autism in Chinese Han population. <i>Scientific Reports</i> , 2017, 7, 2292.	3.3	7
14	Alzheimer's disease associated AKAP9 I2558M mutation alters posttranslational modification and interactome of tau and cellular functions in CRISPR-edited human neuronal cells. <i>Aging Cell</i> , 2022, 21, e13617.	6.7	7
15	Cre-inducible Adeno Associated Virus-mediated Expression of P301L Mutant Tau Causes Motor Deficits and Neuronal Degeneration in the Substantia Nigra. <i>Neuroscience</i> , 2019, 422, 65-74.	2.3	6