

# Yufeng Wei

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/769885/publications.pdf>

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16  
papers

521  
citations

1305906

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h-index

1336881

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all docs

17  
docs citations

17  
times ranked

755  
citing authors

#	ARTICLE	IF	CITATIONS
1	PEA-15 engages in allosteric interactions using a common scaffold in a phosphorylation-dependent manner. <i>Scientific Reports</i> , 2022, 12, 116.	1.6	1
2	PEA-15 Uses a Common Scaffold to Interact with Different Binding Partners in a Phosphorylation-Dependent Manner. <i>FASEB Journal</i> , 2021, 35, .	0.2	0
3	Substance Use Disorder in the COVID-19 Pandemic: A Systematic Review of Vulnerabilities and Complications. <i>Pharmaceuticals</i> , 2020, 13, 155.	1.7	88
4	PEA-15 C-Terminal Tail Allosterically Modulates Death-Effector Domain Conformation and Facilitates Protein-Protein Interactions. <i>International Journal of Molecular Sciences</i> , 2019, 20, 3335.	1.8	3
5	Phosphorylation States of PEA-15 Control Binding Specificity and Regulate Cell Proliferation and Apoptosis. <i>FASEB Journal</i> , 2019, 33, 631.12.	0.2	0
6	Crossroad control of cell proliferation and apoptosis by PEA-15 phosphorylation homeostasis and allosteric regulation of protein conformations and interactions. <i>FASEB Journal</i> , 2018, 32, 792.30.	0.2	0
7	Phosphorylation of PEA-15 allosterically induces conformational change suited for FADD binding and negatively regulates apoptosis. <i>FASEB Journal</i> , 2018, 32, 652.32.	0.2	0
8	Involvement of the Hippocampus in Binge Ethanol-Induced Spleen Atrophy in Adolescent Rats. <i>Alcoholism: Clinical and Experimental Research</i> , 2016, 40, 1489-1500.	1.4	12
9	On the Quest of Cellular Functions of PEA-15 and the Therapeutic Opportunities. <i>Pharmaceuticals</i> , 2015, 8, 455-473.	1.7	9
10	NeuroHIV and Use of Addictive Substances. <i>International Review of Neurobiology</i> , 2014, 118, 403-440.	0.9	38
11	Substantial Conformational Change Mediated by Charge-Triad Residues of the Death Effector Domain in Protein-Protein Interactions. <i>PLoS ONE</i> , 2013, 8, e83421.	1.1	12
12	High-definition NMR structure of PED/PEA-15 death effector domain reveals details of key polar side chain interactions. <i>Biochemical and Biophysical Research Communications</i> , 2012, 424, 141-146.	1.0	11
13	Profound conformational changes of PED/PEA-15 in ERK2 complex revealed by NMR backbone dynamics. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2012, 1824, 1382-1393.	1.1	9
14	The Structure of FADD and Its Mode of Interaction with Procaspase-8. <i>Molecular Cell</i> , 2006, 22, 599-610.	4.5	154
15	PISEMA Solid-State NMR Spectroscopy. <i>Annual Reports on NMR Spectroscopy</i> , 2004, 52, 1-52.	0.7	165
16	One-dimensional <sup>1</sup> H-detected solid-state NMR experiment to determine amide- <sup>1</sup> H chemical shifts in peptides. <i>Chemical Physics Letters</i> , 2002, 351, 42-46.	1.2	16