

# Jannik N Andersen

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9349493/publications.pdf>

Version: 2024-02-01

34  
papers

4,618  
citations

257450

24  
h-index

434195

31  
g-index

36  
all docs

36  
docs citations

36  
times ranked

6515  
citing authors

#	ARTICLE	IF	CITATIONS
1	Design of BET Inhibitor Bottlebrush Prodrugs with Superior Efficacy and Devoid of Systemic Toxicities. <i>Journal of the American Chemical Society</i> , 2021, 143, 4714-4724.	13.7	18
2	Cross-talk between chromatin acetylation and SUMOylation of tripartite motif-containing protein 24 (TRIM24) impacts cell adhesion. <i>Journal of Biological Chemistry</i> , 2018, 293, 7476-7485.	3.4	27
3	Reduction of liver fibrosis by rationally designed macromolecular telmisartan prodrugs. <i>Nature Biomedical Engineering</i> , 2018, 2, 822-830.	22.5	26
4	TRIM28 multi-domain protein regulates cancer stem cell population in breast tumor development. <i>Oncotarget</i> , 2017, 8, 863-882.	1.8	49
5	Development of a High-Throughput Gene Expression Screen for Modulators of RAS-MAPK Signaling in a Mutant RAS Cellular Context. <i>Journal of Biomolecular Screening</i> , 2016, 21, 989-997.	2.6	5
6	Structure-Guided Design of IACS-9571, a Selective High-Affinity Dual TRIM24-BRPF1 Bromodomain Inhibitor. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1440-1454.	6.4	124
7	Observed bromodomain flexibility reveals histone peptide- and small molecule ligand-compatible forms of ATAD2. <i>Biochemical Journal</i> , 2015, 466, 337-346.	3.7	34
8	Development of novel cellular histone-binding and chromatin-displacement assays for bromodomain drug discovery. <i>Epigenetics and Chromatin</i> , 2015, 8, 37.	3.9	32
9	The SMARCA2/4 ATPase Domain Surpasses the Bromodomain as a Drug Target in SWI/SNF-Mutant Cancers: Insights from cDNA Rescue and PFI-3 Inhibitor Studies. <i>Cancer Research</i> , 2015, 75, 3865-3878.	0.9	202
10	Abstract 4272: Applying TCGA data for breast cancer diagnostics and pathway analysis. , 2014, , .		1
11	Abstract 3714: Characterization of a selective focal adhesion kinase (FAK) inhibitor in a panel of glioblastoma cell lines identify rational drug-drug combination strategies. , 2012, , .		0
12	Conformation-Sensing Antibodies Stabilize the Oxidized Form of PTP1B and Inhibit Its Phosphatase Activity. <i>Cell</i> , 2011, 147, 185-198.	28.9	139
13	Cancer genomics: from discovery science to personalized medicine. <i>Nature Medicine</i> , 2011, 17, 297-303.	30.7	534
14	PDK1 Attenuation Fails to Prevent Tumor Formation in PTEN-Deficient Transgenic Mouse Models. <i>Cancer Research</i> , 2011, 71, 3052-3065.	0.9	30
15	Genetic and Pharmacological Inhibition of PDK1 in Cancer Cells. <i>Journal of Biological Chemistry</i> , 2011, 286, 6433-6448.	3.4	56
16	Identification of Direct Target Engagement Biomarkers for Kinase-Targeted Therapeutics. <i>PLoS ONE</i> , 2011, 6, e26459.	2.5	25
17	A gene expression signature of RAS pathway dependence predicts response to PI3K and RAS pathway inhibitors and expands the population of RAS pathway activated tumors. <i>BMC Medical Genomics</i> , 2010, 3, 26.	1.5	124
18	Pathway-Based Identification of Biomarkers for Targeted Therapeutics: Personalized Oncology with PI3K Pathway Inhibitors. <i>Science Translational Medicine</i> , 2010, 2, 43ra55.	12.4	141

#	ARTICLE	IF	CITATIONS
19	Discovery of PDK1 Kinase Inhibitors with a Novel Mechanism of Action by Ultrahigh Throughput Screening. <i>Journal of Biological Chemistry</i> , 2010, 285, 18838-18846.	3.4	45
20	Abstract 4953: Identification of direct target engagement biomarkers for kinase drug discovery using quantitative mass spectrometry: PDK1 case study. , 2010, , .		0
21	Abstract 5560: Biomarker Discovery and Pathway Mapping using Differential Phosphoproteomics of PI3K-Pathway Inhibitors: PRAS40 Correlates with AKT Activation, but not PTEN Expression in Lung and Breast Cancer. , 2010, , .		0
22	Abstract 4561: Quantitative phosphoproteomics of an AKT inhibitor in a PTEN-LOF breast model by label free phospho-dMS demonstrates modulation of protein transcription, protein translation, and motility. , 2010, , .		0
23	Development of High-Throughput TR-FRET and AlphaScreen <sup>®</sup> Assays for Identification of Potent Inhibitors of PDK1. <i>Journal of Biomolecular Screening</i> , 2009, 14, 1257-1262.	2.6	20
24	Sensitive multiplexed analysis of kinase activities and activity-based kinase identification. <i>Nature Biotechnology</i> , 2009, 27, 933-940.	17.5	99
25	Computational analysis of protein tyrosine phosphatases: practical guide to bioinformatics and data resources. <i>Methods</i> , 2005, 35, 90-114.	3.8	40
26	A genomic perspective on protein tyrosine phosphatases: gene structure, pseudogenes, and genetic disease linkage. <i>FASEB Journal</i> , 2004, 18, 8-30.	0.5	277
27	Redox regulation of protein tyrosine phosphatase 1B involves a sulphenyl-amide intermediate. <i>Nature</i> , 2003, 423, 769-773.	27.8	866
28	Cellular Effects of Small Molecule PTP1B Inhibitors on Insulin Signaling. <i>Biochemistry</i> , 2003, 42, 12792-12804.	2.5	107
29	Enzyme kinetic characterization of protein tyrosine phosphatases. <i>Biochimie</i> , 2003, 85, 527-534.	2.6	24
30	Structural and Evolutionary Relationships among Protein Tyrosine Phosphatase Domains. <i>Molecular and Cellular Biology</i> , 2001, 21, 7117-7136.	2.3	660
31	TYK2 and JAK2 Are Substrates of Protein-tyrosine Phosphatase 1B. <i>Journal of Biological Chemistry</i> , 2001, 276, 47771-47774.	3.4	379
32	Molecular Dynamics Simulations of Protein-Tyrosine Phosphatase 1B. II. Substrate-Enzyme Interactions and Dynamics. <i>Biophysical Journal</i> , 2000, 78, 2191-2200.	0.5	48
33	Molecular Basis for the Dephosphorylation of the Activation Segment of the Insulin Receptor by Protein Tyrosine Phosphatase 1B. <i>Molecular Cell</i> , 2000, 6, 1401-1412.	9.7	432
34	Molecular Dynamics Simulations of Protein-Tyrosine Phosphatase 1B. I. Ligand-Induced Changes in the Protein Motions. <i>Biophysical Journal</i> , 1999, 77, 505-515.	0.5	42