

# Kenneth H Pearce

## List of Publications by Year in descending order

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Version: 2024-02-01

22  
papers

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citations

840776

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h-index

677142

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docs citations

23  
times ranked

1327  
citing authors

#	ARTICLE	IF	CITATIONS
1	Discovery of Potent Peptidomimetic Antagonists for Heterochromatin Protein 1 Family Proteins. ACS Omega, 2022, 7, 716-732.	3.5	3
2	Development of Novel IP6K Inhibitors for the Treatment of Obesity and Obesity-Induced Metabolic Dysfunctions. Journal of Medicinal Chemistry, 2022, 65, 6869-6887.	6.4	15
3	Discovery and Structural Basis of the Selectivity of Potent Cyclic Peptide Inhibitors of MAGE-A4. Journal of Medicinal Chemistry, 2022, 65, 7231-7245.	6.4	12
4	Repurposing the Ebola and Marburg Virus Inhibitors Tilorone, Quinacrine, and Pyronaridine: <i>In Vitro</i> Activity against SARS-CoV-2 and Potential Mechanisms. ACS Omega, 2021, 6, 7454-7468.	3.5	56
5	A Peptidomimetic Ligand Targeting the Chromodomain of MPP8 Reveals HRP2's Association with the HUSH Complex. ACS Chemical Biology, 2021, 16, 1721-1736.	3.4	12
6	Discovery and Development of Cyclic Peptide Inhibitors of CIB1. ACS Medicinal Chemistry Letters, 2021, 12, 1832-1839.	2.8	14
7	Small molecule screening identifies inhibitors of the Epstein-Barr virus deubiquitinating enzyme, BPLF1. Antiviral Research, 2020, 173, 104649.	4.1	6
8	Degradation of Polycomb Repressive Complex 2 with an EED-Targeted Bivalent Chemical Degradator. Cell Chemical Biology, 2020, 27, 47-56.e15.	5.2	127
9	A High-Throughput Assay to Identify Allosteric Inhibitors of the PLC- $\beta$ Isozymes Operating at Membranes. Biochemistry, 2020, 59, 4029-4038.	2.5	5
10	Discovery and Characterization of Peptide Inhibitors for Calcium and Integrin Binding Protein 1. ACS Chemical Biology, 2020, 15, 1505-1516.	3.4	11
11	Design and Construction of a Focused DNA-Encoded Library for Multivalent Chromatin Reader Proteins. Molecules, 2020, 25, 979.	3.8	12
12	Application of a MYC degradation screen identifies sensitivity to CDK9 inhibitors in KRAS-mutant pancreatic cancer. Science Signaling, 2019, 12, .	3.6	46
13	Targeting Regorafenib-Induced Toxicity through Inhibition of Gut Microbial $\beta$ -Glucuronidases. ACS Chemical Biology, 2019, 14, 2737-2744.	3.4	41
14	Discovery and Characterization of a Cellular Potent Positive Allosteric Modulator of the Polycomb Repressive Complex 1 Chromodomain, CBX7. Cell Chemical Biology, 2019, 26, 1365-1379.e22.	5.2	38
15	A General TR-FRET Assay Platform for High-Throughput Screening and Characterizing Inhibitors of Methyl-Lysine Reader Proteins. SLAS Discovery, 2019, 24, 693-700.	2.7	25
16	BacMam production and crystal structure of nonglycosylated apo human furin at 1.89 Å resolution. Acta Crystallographica Section F, Structural Biology Communications, 2019, 75, 239-245.	0.8	8
17	Inhibition of Inositol Polyphosphate Kinases by Quercetin and Related Flavonoids: A Structure-Activity Analysis. Journal of Medicinal Chemistry, 2019, 62, 1443-1454.	6.4	38
18	KRAS Suppression-Induced Degradation of MYC Is Antagonized by a MEK5-ERK5 Compensatory Mechanism. Cancer Cell, 2018, 34, 807-822.e7.	16.8	112

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19	Use of Protein Kinase-Focused Compound Libraries for the Discovery of New Inositol Phosphate Kinase Inhibitors. <i>SLAS Discovery</i> , 2018, 23, 982-988.	2.7	15
20	Identification of Cosalane as an Inhibitor of Human and Murine Chemokine Receptor 7 Signaling via a High-Throughput Screen. <i>SLAS Discovery</i> , 2018, 23, 1083-1091.	2.7	10
21	A neomorphic cancer cell-specific role of MAGE-A4 in trans-lesion synthesis. <i>Nature Communications</i> , 2016, 7, 12105.	12.8	52
22	Structure-Activity Relationships and Kinetic Studies of Peptidic Antagonists of CBX Chromodomains. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 8913-8923.	6.4	28