

Adam B Keeton

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/3857673/adam-b-keeton-publications-by-year.pdf>

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

25
papers

606
citations

12
h-index

24
g-index

25
ext. papers

795
ext. citations

4.7
avg, IF

3.25
L-index

#	Paper	IF	Citations
25	The path to the clinic: a comprehensive review on direct KRAS inhibitors.. <i>Journal of Experimental and Clinical Cancer Research</i> , 2022 , 41, 27	12.8	7
24	Pan-RAS inhibitors: Hitting multiple RAS isozymes with one stone.. <i>Advances in Cancer Research</i> , 2022 , 153, 131-168	5.9	0
23	From Celecoxib to a Novel Class of Phosphodiesterase 5 Inhibitors: Trisubstituted Pyrazolines as Novel Phosphodiesterase 5 Inhibitors with Extremely High Potency and Phosphodiesterase Isozyme Selectivity. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 4462-4477	8.3	1
22	Suppression of Colon Tumorigenesis in Mutant Mice by a Novel PDE10 Inhibitor that Reduces Oncogenic β Catenin. <i>Cancer Prevention Research</i> , 2021 , 14, 995-1008	3.2	2
21	Novel thiazolidine derivatives as potent selective pro-apoptotic agents. <i>Bioorganic Chemistry</i> , 2021 , 114, 105143	5.1	0
20	Discovery of trisubstituted pyrazolines as a novel scaffold for the development of selective phosphodiesterase 5 inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 104, 104322	5.1	3
19	Extending the use of tadalafil scaffold: Development of novel selective phosphodiesterase 5 inhibitors and histone deacetylase inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 98, 103742	5.1	9
18	Targeting cGMP/PKG signaling for the treatment or prevention of colorectal cancer with novel sulindac derivatives lacking cyclooxygenase inhibitory activity 2020 , 3, 1-6		1
17	PDE5 and PDE10 inhibition activates cGMP/PKG signaling to block Wnt/ β catenin transcription, cancer cell growth, and tumor immunity. <i>Drug Discovery Today</i> , 2020 , 25, 1521-1527	8.8	7
16	New Isatin-Indole Conjugates: Synthesis, Characterization, and a Plausible Mechanism of Their in vitro Antiproliferative Activity. <i>Drug Design, Development and Therapy</i> , 2020 , 14, 483-495	4.4	7
15	Enhancing anticancer activity of checkpoint immunotherapy by targeting RAS. <i>MedComm</i> , 2020 , 1, 121-128		6
14	Antiproliferative activity and possible mechanism of action of certain 5-methoxyindole tethered C-5 functionalized isatins. <i>Drug Design, Development and Therapy</i> , 2019 , 13, 3069-3078	4.4	2
13	Abstract 2707: A novel RAS inhibitor, MCI-062, inhibits colon tumor growth in vivo and activates antitumor immunity 2019 ,		3
12	Exploiting RAS Nucleotide Cycling as a Strategy for Drugging RAS-Driven Cancers. <i>International Journal of Molecular Sciences</i> , 2019 , 21,	6.3	12
11	Synthesis and biological evaluation of certain hydrazonoindolin-2-one derivatives as new potent anti-proliferative agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 867-878	5.6	24
10	New hydrazonoindolin-2-ones: Synthesis, exploration of the possible anti-proliferative mechanism of action and encapsulation into PLGA microspheres. <i>PLoS ONE</i> , 2017 , 12, e0181241	3.7	23
9	Phosphodiesterase 10A is overexpressed in lung tumor cells and inhibitors selectively suppress growth by blocking β catenin and MAPK signaling. <i>Oncotarget</i> , 2017 , 8, 69264-69280	3.3	16

8	β-catenin nuclear translocation in colorectal cancer cells is suppressed by PDE10A inhibition, cGMP elevation, and activation of PKG. <i>Oncotarget</i> , 2016 , 7, 5353-65	3.3	28
7	Suppression of β-catenin/TCF transcriptional activity and colon tumor cell growth by dual inhibition of PDE5 and 10. <i>Oncotarget</i> , 2015 , 6, 27403-15	3.3	27
6	Sulindac selectively inhibits colon tumor cell growth by activating the cGMP/PKG pathway to suppress Wnt/β-catenin signaling. <i>Molecular Cancer Therapeutics</i> , 2013 , 12, 1848-59	6.1	83
5	A novel sulindac derivative that potently suppresses colon tumor cell growth by inhibiting cGMP phosphodiesterase and β-catenin transcriptional activity. <i>Cancer Prevention Research</i> , 2012 , 5, 822-33	3.2	56
4	Inhibition of PDE5 by sulindac sulfide selectively induces apoptosis and attenuates oncogenic Wnt/β-catenin-mediated transcription in human breast tumor cells. <i>Cancer Prevention Research</i> , 2011 , 4, 1275-84	3.2	77
3	Colon tumor cell growth-inhibitory activity of sulindac sulfide and other nonsteroidal anti-inflammatory drugs is associated with phosphodiesterase 5 inhibition. <i>Cancer Prevention Research</i> , 2010 , 3, 1303-13	3.2	62
2	A novel sulindac derivative that does not inhibit cyclooxygenases but potently inhibits colon tumor cell growth and induces apoptosis with antitumor activity. <i>Cancer Prevention Research</i> , 2009 , 2, 572-80	3.2	68
1	Sulindac sulfide selectively inhibits growth and induces apoptosis of human breast tumor cells by phosphodiesterase 5 inhibition, elevation of cyclic GMP, and activation of protein kinase G. <i>Molecular Cancer Therapeutics</i> , 2009 , 8, 3331-40	6.1	82