Paulo H C Godoi

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8071378/publications.pdf

Version: 2024-02-01

759233 940533 17 502 12 16 citations h-index g-index papers 21 21 21 915 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Discovery and Characterization of Selective and Ligand-Efficient DYRK Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 11709-11728.	6.4	11
2	Binding and structural analyses of potent inhibitors of the human Ca2+/calmodulin dependent protein kinase kinase 2 (CAMKK2) identified from a collection of commercially-available kinase inhibitors. Scientific Reports, 2019, 9, 16452.	3.3	16
3	Validation of the protein kinase <i>Pf</i> CLK3 as a multistage cross-species malarial drug target. Science, 2019, 365, .	12.6	51
4	SGC-GAK-1: A Chemical Probe for Cyclin G Associated Kinase (GAK). Journal of Medicinal Chemistry, 2019, 62, 2830-2836.	6.4	56
5	WNT Activates the AAK1 Kinase to Promote Clathrin-Mediated Endocytosis of LRP6 and Establish a Negative Feedback Loop. Cell Reports, 2019, 26, 79-93.e8.	6.4	68
6	Identification and Optimization of 4â€Anilinoquinolines as Inhibitors of Cyclinâ€G Associated Kinase. ChemMedChem, 2018, 13, 48-66.	3.2	51
7	Structural Analysis of Inhibitor Binding to CAMKK1 Identifies Features Necessary for Design of Specific Inhibitors. Scientific Reports, 2018, 8, 14800.	3.3	13
8	1,2,6-Thiadiazinones as Novel Narrow Spectrum Calcium/Calmodulin-Dependent Protein Kinase Kinase 2 (CaMKK2) Inhibitors. Molecules, 2018, 23, 1221.	3.8	23
9	Structural characterization of human Vaccinia-Related Kinases (VRK) bound to small-molecule inhibitors identifies different P-loop conformations. Scientific Reports, 2017, 7, 7501.	3.3	21
10	Orphan Nuclear Receptor NR4A1 Binds a Novel Protein Interaction Site on Anti-apoptotic B Cell Lymphoma Gene 2 Family Proteins. Journal of Biological Chemistry, 2016, 291, 14072-14084.	3.4	17
11	TR-FRET-Based High-Throughput Screening Assay for Identification of UBC13 Inhibitors. Journal of Biomolecular Screening, 2012, 17, 163-176.	2.6	34
12	High-Throughput Fluorescence Polarization Assay for Chemical Library Screening against Anti-Apoptotic Bcl-2 Family Member Bfl-1. Journal of Biomolecular Screening, 2012, 17, 350-360.	2.6	22
13	A TR3/Nur77 Peptide-Based High-Throughput Fluorescence Polarization Screen for Small Molecule Bcl-B Inhibitors. Journal of Biomolecular Screening, 2008, 13, 665-673.	2.6	12
14	Structure of the Thiazole Biosynthetic Enzyme THI1 from Arabidopsis thaliana. Journal of Biological Chemistry, 2006, 281, 30957-30966.	3.4	72
15	Crystal structure of human phosphoglucose isomerase and analysis of the initial catalytic steps. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2003, 1645, 117-122.	2.3	27
16	Human phosphoglucose isomerase: expression, purification, crystallization and preliminary crystallographic analysis. Acta Crystallographica Section D: Biological Crystallography, 2001, 57, 592-595.	2.5	4
17	A Transcription-uncoupled Negative Feedback Loop for the $1\mathrm{WNT}$ Pathway: WNT Activates the AAK1 Kinase to Promote Clathrin-mediated Endocytosis of LRP6. SSRN Electronic Journal, $0, , .$	0.4	O