

Paulo H C Godoi

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

502
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759233

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all docs

21
docs citations

21
times ranked

915
citing authors

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