

Roberta Tesch

List of Publications by Citations

Source: <https://exaly.com/author-pdf/8782967/roberta-tesch-publications-by-citations.pdf>

Version: 2024-04-27

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.

16
papers

182
citations

9
h-index

13
g-index

17
ext. papers

253
ext. citations

6
avg, IF

2.57
L-index

#	Paper	IF	Citations
16	Acylhydrazone derivatives: a patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2014 , 24, 1161-70	6.8	36
15	N-acylhydrazone derivative ameliorates monocrotaline-induced pulmonary hypertension through the modulation of adenosine AA2R activity. <i>International Journal of Cardiology</i> , 2014 , 173, 154-62	3.2	28
14	Antihypertensive profile of 2-thienyl-3,4-methylenedioxybenzoylhydrazone is mediated by activation of the A2A adenosine receptor. <i>European Journal of Medicinal Chemistry</i> , 2012 , 55, 49-57	6.8	26
13	Pyridinylimidazoles as dual glycogen synthase kinase 3 β /p38 β mitogen-activated protein kinase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019 , 175, 309-329	6.8	15
12	Identification of molecular targets for the targeted treatment of gastric cancer using dasatinib. <i>Oncotarget</i> , 2020 , 11, 535-549	3.3	15
11	Phenylpiperazine derivatives: a patent review (2006 - present). <i>Expert Opinion on Therapeutic Patents</i> , 2012 , 22, 1169-78	6.8	14
10	Novel potent imidazo[1,2-a]pyridine-N-Glyciny-hydrazone inhibitors of TNF- α production: in vitro and in vivo studies. <i>PLoS ONE</i> , 2014 , 9, e91660	3.7	11
9	Fast Iterative Synthetic Approach toward Identification of Novel Highly Selective p38 MAP Kinase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 10757-10782	8.3	10
8	An Unusual Intramolecular Halogen Bond Guides Conformational Selection. <i>Angewandte Chemie - International Edition</i> , 2018 , 57, 9970-9975	16.4	9
7	A Highly Selective Chemical Probe for Activin Receptor-like Kinases ALK4 and ALK5. <i>ACS Chemical Biology</i> , 2020 , 15, 862-870	4.9	7
6	Structure-Based Design of Selective Salt-Inducible Kinase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 8142-8160	8.3	4
5	LASSBio-897 Reduces Lung Injury Induced by Silica Particles in Mice: Potential Interaction with the A Receptor. <i>Frontiers in Pharmacology</i> , 2017 , 8, 778	5.6	3
4	Addressing a Trapped High-Energy Water: Design and Synthesis of Highly Potent Pyrimidoindole-Based Glycogen Synthase Kinase-3 Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 8142-8160	8.3	3
3	The Small-Molecule Inhibitor MR1A9 Reveals Novel Insights into the Cell Cycle Roles of SIK2 in Ovarian Cancer Cells. <i>Cancers</i> , 2021 , 13,	6.6	1
2	Eine ungewöhnliche intramolekulare Halogenbindung führt zu konformationeller Selektion. <i>Angewandte Chemie</i> , 2018 , 130, 10120-10126	3.6	
1	Design, Synthesis and Pharmacological Evaluation of Novel Antiinflammatory and Analgesic O-Benzoyloxime Compounds Derived From Natural Eugenol. <i>Letters in Drug Design and Discovery</i> , 2019 , 16, 1157-1166	0.8	