

# Rajendra Prasad Yejella

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3089379/publications.pdf>

Version: 2024-02-01

9  
papers

118  
citations

1478505

6  
h-index

1474206

9  
g-index

9  
all docs

9  
docs citations

9  
times ranked

168  
citing authors

#	ARTICLE	IF	CITATIONS
1	Design, Synthesis, and Antibacterial and Antifungal Activities of Novel Trifluoromethyl and Trifluoromethoxy Substituted Chalcone Derivatives. <i>Pharmaceuticals</i> , 2020, 13, 375.	3.8	29
2	Molecular docking based screening of novel designed chalcone series of compounds for their anti-cancer activity targeting EGFR kinase domain. <i>Bioinformation</i> , 2015, 11, 322-329.	0.5	22
3	Novel indolizine derivatives lowers blood glucose levels in streptozotocin-induced diabetic rats: A histopathological approach. <i>Pharmacological Reports</i> , 2019, 71, 233-242.	3.3	21
4	A Study of Anti-inflammatory and Analgesic Activity of New 2,4,6-Trisubstituted Pyrimidines. <i>Chemical and Pharmaceutical Bulletin</i> , 2011, 59, 1079-1082.	1.3	20
5	Development and Validation of a Stability Indicating RP-HPLC Method for Simultaneous Estimation of Teneligliptin and Metformin. <i>Turkish Journal of Pharmaceutical Sciences</i> , 2020, 17, 141-147.	1.4	11
6	Synthesis, Antimicrobial, and Computational Evaluation of Novel Isobutylchalcones as Antimicrobial Agents. <i>International Journal of Medicinal Chemistry</i> , 2017, 2017, 1-14.	2.2	7
7	Antitubercular activity assessment of fluorinated chalcones, 2-aminopyridine-3-carbonitrile and 2-amino-4H-pyran-3-carbonitrile derivatives: In vitro, molecular docking and in-silico drug likeliness studies. <i>PLoS ONE</i> , 2022, 17, e0265068.	2.5	4
8	De novo Based Ligand generation and Docking studies of PPAR $\gamma$ Agonists: Correlations between Predicted Biological activity vs. Biopharmaceutical Descriptors. <i>Chem-Bio Informatics Journal</i> , 2010, 10, 74-86.	0.3	3
9	Molecular docking based screening of G6PS with 1, 5 Benzothiazepine derivates for a potential inhibitor. <i>Bioinformation</i> , 2015, 11, 525-528.	0.5	1