

# Prasanth Reddy Nyalapatla

## List of Publications by Year in descending order

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Version: 2024-02-01

18  
papers

520  
citations

840585

11  
h-index

839398

18  
g-index

20  
all docs

20  
docs citations

20  
times ranked

851  
citing authors

#	ARTICLE	IF	CITATIONS
1	Fluorescent Probes for Monitoring Serine Ubiquitination. <i>Biochemistry</i> , 2020, 59, 1309-1313.	1.2	6
2	Single atom changes in newly synthesized HIV protease inhibitors reveal structural basis for extreme affinity, high genetic barrier, and adaptation to the HIV protease plasticity. <i>Scientific Reports</i> , 2020, 10, 10664.	1.6	13
3	Co(II)-salen catalyzed stereoselective cyclopropanation of fluorinated styrenes. <i>Chirality</i> , 2019, 31, 1014-1027.	1.3	4
4	Halogen Bond Interactions of Novel HIV-1 Protease Inhibitors (PI) (GRL-001-15 and GRL-003-15) with the Flap of Protease Are Critical for Their Potent Activity against Wild-Type HIV-1 and Multi-PI-Resistant Variants. <i>Antimicrobial Agents and Chemotherapy</i> , 2019, 63, .	1.4	12
5	A New Synthesis of Gefitinib. <i>Synlett</i> , 2019, 30, 471-476.	1.0	6
6	Design of Highly Potent, Dual-Acting and Central-Nervous-System-Penetrating HIV-1 Protease Inhibitors with Excellent Potency against Multidrug-Resistant HIV-1 Variants. <i>ChemMedChem</i> , 2018, 13, 803-815.	1.6	36
7	Design and Synthesis of Highly Potent HIV-1 Protease Inhibitors Containing Tricyclic Fused Ring Systems as Novel P2 Ligands: Structure-Activity Studies, Biological and X-ray Structural Analysis. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 4561-4577.	2.9	31
8	Total syntheses of both enantiomers of ampirionin 4: A chemoenzymatic based strategy for functionalized tetrahydrofurans. <i>Tetrahedron</i> , 2017, 73, 1820-1830.	1.0	10
9	Design of novel HIV-1 protease inhibitors incorporating isophthalamide-derived P2-P3 ligands: Synthesis, biological evaluation and X-ray structural studies of inhibitor-HIV-1 protease complex. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 5114-5127.	1.4	16
10	Design and Development of Highly Potent HIV-1 Protease Inhibitors with a Crown-Like Oxotricyclic Core as the P2-Ligand To Combat Multidrug-Resistant HIV Variants. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 4267-4278.	2.9	64
11	A novel central nervous system-penetrating protease inhibitor overcomes human immunodeficiency virus 1 resistance with unprecedented aM to pM potency. <i>ELife</i> , 2017, 6, .	2.8	44
12	Enantioselective Total Synthesis of (+)-Ampirionin-4. <i>Organic Letters</i> , 2016, 18, 2296-2299.	2.4	16
13	A Novel Tricyclic Ligand-Containing Nonpeptidic HIV-1 Protease Inhibitor, GRL-0739, Effectively Inhibits the Replication of Multidrug-Resistant HIV-1 Variants and Has a Desirable Central Nervous System Penetration Property <i>In Vitro</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 2625-2635.	1.4	10
14	Ligand-induced Dimerization of Middle East Respiratory Syndrome (MERS) Coronavirus nsp5 Protease (3CLpro). <i>Journal of Biological Chemistry</i> , 2015, 290, 19403-19422.	1.6	134
15	X-ray structure and inhibition of the feline infectious peritonitis virus 3C-like protease: Structural implications for drug design. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 5072-5077.	1.0	19
16	Design, synthesis, biological evaluation and X-ray structural studies of HIV-1 protease inhibitors containing substituted fused-tetrahydropyranyl tetrahydrofuran as P2-ligands. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 11607-11621.	1.5	10
17	Highly Potent HIV-1 Protease Inhibitors with Novel Tricyclic P2 Ligands: Design, Synthesis, and Protein-Ligand X-ray Studies. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 6792-6802.	2.9	42
18	Zinc Catalyzed and Mediated Asymmetric Propargylation of Trifluoromethyl Ketones with a Propargyl Boronate. <i>Journal of Organic Chemistry</i> , 2013, 78, 3592-3615.	1.7	47