JoÃ£o Neres

List of Publications by Year in descending order

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Version: 2024-02-01

32 papers	2,082 citations	257357 24 h-index	414303 32 g-index
32 all docs	32 docs citations	32 times ranked	2273 citing authors

#	Article	IF	Citations
1	Mechanism of a Standalone Î²â€Lactone Synthetase: Newâ€‰Continuous Assay for a Widespread ANL Superfamily Enzyme. ChemBioChem, 2019, 20, 1701-1711.	1.3	5
2	Fluorescent Benzothiazinone Analogues Efficiently and Selectively Label Dpre1 in Mycobacteria and Actinobacteria. ACS Chemical Biology, 2018, 13, 3184-3192.	1.6	16
3	Nitroarenes as Antitubercular Agents: Stereoelectronic Modulation to Mitigate Mutagenicity. ChemMedChem, 2016, 11, 331-339.	1.6	19
4	Characterization of DprE1-Mediated Benzothiazinone Resistance in Mycobacterium tuberculosis. Antimicrobial Agents and Chemotherapy, 2016, 60, 6451-6459.	1.4	36
5	The 8-Pyrrole-Benzothiazinones Are Noncovalent Inhibitors of DprE1 from Mycobacterium tuberculosis. Antimicrobial Agents and Chemotherapy, 2015, 59, 4446-4452.	1.4	85
6	DprE1 Is a Vulnerable Tuberculosis Drug Target Due to Its Cell Wall Localization. ACS Chemical Biology, 2015, 10, 1631-1636.	1.6	123
7	Discovery of benzothiazoles as antimycobacterial agents: Synthesis, structureâ€“activity relationships and binding studies with Mycobacterium tuberculosis decaprenylphosphoryl-Î²-d-ribose 2â€²-oxidase. Bioorganic and Medicinal Chemistry, 2015, 23, 7694-7710.	1.4	44
8	2-Carboxyquinoxalines Kill Mycobacterium tuberculosis through Noncovalent Inhibition of DprE1. ACS Chemical Biology, 2015, 10, 705-714.	1.6	116
9	Selective detection of epimeric pentose saccharides at physiological pH using a fluorescent receptor. Carbohydrate Research, 2014, 391, 61-65.	1.1	1
10	Pyridomycin bridges the NADH- and substrate-binding pockets of the enoyl reductase InhA. Nature Chemical Biology, 2014, 10, 96-98.	3.9	63
11	Towards a new combination therapy for tuberculosis with next generation benzothiazinones. EMBO Molecular Medicine, 2014, 6, 372-383.	3.3	311
12	4-Aminoquinolone Piperidine Amides: Noncovalent Inhibitors of DprE1 with Long Residence Time and Potent Antimycobacterial Activity. Journal of Medicinal Chemistry, 2014, 57, 5419-5434.	2.9	97
13	Insights into the Activity and Specificity of Trypanosoma cruzi trans-Sialidase from Molecular Dynamics Simulations. Biochemistry, 2013, 52, 3740-3751.	1.2	14
14	Non-Nucleoside Inhibitors of BasE, an Adenylating Enzyme in the Siderophore Biosynthetic Pathway of the Opportunistic Pathogen Acinetobacter baumannii. Journal of Medicinal Chemistry, 2013, 56, 2385-2405.	2.9	48
15	DprE1 - from the Discovery to the Promising Tuberculosis Drug Target. Current Pharmaceutical Design, 2013, 20, 4379-4403.	0.9	47
16	Towards a new tuberculosis drug: pyridomycin â€“ nature's isoniazid. EMBO Molecular Medicine, 2012, 4, 1032-1042.	3.3	175
17	Structural Basis for Benzothiazinone-Mediated Killing of Mycobacterium tuberculosis. Science Translational Medicine, 2012, 4, 150ra121.	5.8	159
18	Inhibitors of the Salicylate Synthase (MbtI) from Mycobacterium tuberculosis Discovered by Highâ€Throughput Screening. ChemMedChem, 2010, 5, 2079-2087.	1.6	41

#	Article	IF	Citations
19	Biological and structural characterization of the Mycobacterium smegmatis nitroreductase NfnB, and its role in benzothiazinone resistance. Molecular Microbiology, 2010, 77, 1172-1185.	1.2	63
20	Tryptophan as a Molecular Shovel in the Glycosyl Transfer Activity ofÂTrypanosoma cruzi Trans-sialidase. Biophysical Journal, 2010, 98, L38-L40.	0.2	26
21	Biophysical and X-ray Crystallographic Analysis of Mps1 Kinase Inhibitor Complexes[,]. Biochemistry, 2010, 49, 1689-1701.	1.2	35
22	Biochemical and Structural Characterization of Bisubstrate Inhibitors of BasE, the Self-Standing Nonribosomal Peptide Synthetase Adenylate-Forming Enzyme of Acinetobactin Synthesis,. Biochemistry, 2010, 49, 9292-9305.	1.2	52
23	The Global Virulence Regulators VsrAD and PhcA Control Secondary Metabolism in the Plant Pathogen Ralstonia solanacearum. ChemBioChem, 2009, 10, 2730-2732.	1.3	38
24	Discovery of novel inhibitors of Trypanosoma cruzi trans-sialidase from in silico screening. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 589-596.	1.0	68
25	Rational drug design in parasitology: trans-sialidase as a case study for Chagas disease. Drug Discovery Today, 2008, 13, 110-117.	3.2	46
26	Inhibition of Siderophore Biosynthesis by 2-Triazole Substituted Analogues of 5â€²-O-[N-(Salicyl)sulfamoyl]adenosine: Antibacterial Nucleosides Effective against Mycobacterium tuberculosis. Journal of Medicinal Chemistry, 2008, 51, 7495-7507.	2.9	83
27	Aryl Acid Adenylating Enzymes Involved in Siderophore Biosynthesis: Fluorescence Polarization Assay, Ligand Specificity, and Discovery of Non-nucleoside Inhibitors via High-Throughput Screening. Biochemistry, 2008, 47, 11735-11749.	1.2	43
28	Inhibition of Siderophore Biosynthesis in Mycobacterium tuberculosis with Nucleoside Bisubstrate Analogues: Structureâ”Activity Relationships of the Nucleobase Domain of 5â€²-O-[N-(Salicyl)sulfamoyl]adenosine. Journal of Medicinal Chemistry, 2008, 51, 5349-5370.	2.9	118
29	Benzoic acid and pyridine derivatives as inhibitors of Trypanosoma cruzi trans-sialidase. Bioorganic and Medicinal Chemistry, 2007, 15, 2106-2119.	1.4	41
30	Continuous fluorimetric assay for high-throughput screening of inhibitors of trans-sialidase from Trypanosoma cruzi. Analytical Biochemistry, 2006, 357, 302-304.	1.1	19
31	Design, Synthesis, and Enzymatic Evaluation ofN1-Acyloxyalkyl-andN1-Oxazolidin-2,4-dion-5-yl-Substituted Î²-lactams as Novel Inhibitors of Human Leukocyte Elastase. Journal of Medicinal Chemistry, 2005, 48, 4861-4870.	2.9	33
32	Synthesis, Stability and In Vitro Dermal Evaluation of Aminocarbonyloxymethyl Esters as Prodrugs of Carboxylic Acid Agents. Bioorganic and Medicinal Chemistry, 2002, 10, 809-816.	1.4	17