

Donatella Tondi

List of Publications by Year in descending order

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42
papers

1,210
citations

331670

21
h-index

377865

34
g-index

48
all docs

48
docs citations

48
times ranked

1407
citing authors

#	ARTICLE	IF	CITATIONS
1	Ten Years with New Delhi Metallo- β -lactamase-1 (NDM-1): From Structural Insights to Inhibitor Design. <i>ACS Infectious Diseases</i> , 2019, 5, 9-34.	3.8	123
2	Structure-based discovery and in-parallel optimization of novel competitive inhibitors of thymidylate synthase. <i>Chemistry and Biology</i> , 1999, 6, 319-331.	6.0	103
3	Thymidylate Synthase Structure, Function and Implication in Drug Discovery. <i>Current Medicinal Chemistry</i> , 2005, 12, 2241-2258.	2.4	91
4	Protein-protein interface-binding peptides inhibit the cancer therapy target human thymidylate synthase. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, E542-9.	7.1	77
5	Structure-Based Optimization of a Non- β -lactam Lead Results in Inhibitors That Do Not Up-Regulate β -Lactamase Expression in Cell Culture. <i>Journal of the American Chemical Society</i> , 2005, 127, 4632-4639.	13.7	58
6	Mono- and Disubstituted-3,8-diazabicyclo[3.2.1]octane Derivatives as Analgesics Structurally Related to Epibatidine: Synthesis, Activity, and Modeling. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 674-681.	6.4	56
7	Structure-Based Design of Inhibitors Specific for Bacterial Thymidylate Synthase. <i>Biochemistry</i> , 1999, 38, 1607-1617.	2.5	49
8	Structure-based design and in-parallel synthesis of inhibitors of AmpC β -lactamase. <i>Chemistry and Biology</i> , 2001, 8, 593-610.	6.0	45
9	Targeting Class A and C Serine β -Lactamases with a Broad-Spectrum Boronic Acid Derivative. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 5449-5458.	6.4	45
10	Optimizing Cell Permeation of an Antibiotic Resistance Inhibitor for Improved Efficacy. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 5644-5654.	6.4	41
11	Structural study of phenyl boronic acid derivatives as AmpC β -lactamase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 3416-3419.	2.2	38
12	Structure-Based Virtual Screening for the Discovery of Novel Inhibitors of New Delhi Metallo- β -lactamase-1. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 45-50.	2.8	38
13	Computational and biological profile of boronic acids for the detection of bacterial serine- and metallo- β -lactamases. <i>Scientific Reports</i> , 2017, 7, 17716.	3.3	35
14	Structure-based studies on species-specific inhibition of thymidylate synthase. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2002, 1587, 206-214.	3.8	34
15	SOS response in bacteria: Inhibitory activity of lichen secondary metabolites against <i>Escherichia coli</i> RecA protein. <i>Phytomedicine</i> , 2017, 29, 11-18.	5.3	34
16	X-ray Crystallography Deciphers the Activity of Broad-Spectrum Boronic Acid β -Lactamase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 650-655.	2.8	30
17	Predicting and harnessing protein flexibility in the design of species-specific inhibitors of thymidylate synthase. <i>Escherichia coli</i> thymidylate synthase numbering is used unless otherwise noted. PDB coordinates have been deposited with the RCSB with accession ID: 1JG0.. <i>Chemistry and Biology</i> , 2001, 8, 981-995.	6.0	28
18	Decoding the Structural Basis For Carbapenem Hydrolysis By Class A β -lactamases: Fishing For A Pharmacophore. <i>Current Drug Targets</i> , 2016, 17, 983-1005.	2.1	27

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19	Virtual screening identifies broad-spectrum \hat{I}^2 -lactamase inhibitors with activity on clinically relevant serine- and metallo-carbapenemases. <i>Scientific Reports</i> , 2020, 10, 12763.	3.3	25
20	Phenylboronic Acid Derivatives as Validated Leads Active in Clinical Strains Overexpressing KPC \hat{I}^2 : A Step against Bacterial Resistance. <i>ChemMedChem</i> , 2018, 13, 713-724.	3.2	24
21	Phthalein Derivatives as a New Tool for Selectivity in Thymidylate Synthase Inhibition. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 2112-2124.	6.4	23
22	The Inhibition of Extended Spectrum \hat{I}^2 -Lactamases: Hits and Leads. <i>Current Medicinal Chemistry</i> , 2014, 21, 1405-1434.	2.4	23
23	Improving Specificity vs Bacterial Thymidylate Synthases through N-Dansyl Modulation of Didansyltyrosine. <i>Journal of Medicinal Chemistry</i> , 2005, 48, 913-916.	6.4	16
24	Inhibition of the transcriptional repressor LexA: Withstanding drug resistance by inhibiting the bacterial mechanisms of adaptation to antimicrobials. <i>Life Sciences</i> , 2020, 241, 117116.	4.3	16
25	Conformational analysis of phthalein derivatives acting as thymidylate synthase inhibitors by means of 1H NMR and quantum chemical calculations. <i>Bioorganic and Medicinal Chemistry</i> , 1996, 4, 1783-1794.	3.0	14
26	An Improved Synthesis of CENTA, a Chromogenic Substrate for \hat{I}^2 -Lactamases. <i>Synlett</i> , 2016, 27, 2447-2450.	1.8	13
27	4-Amino-1,2,4-triazole-3-thione as a Promising Scaffold for the Inhibition of Serine and Metallo- \hat{I}^2 -Lactamases. <i>Pharmaceuticals</i> , 2020, 13, 52.	3.8	13
28	Design, synthesis and biological evaluation of non-covalent AmpC \hat{I}^2 -lactamases inhibitors. <i>Medicinal Chemistry Research</i> , 2017, 26, 975-986.	2.4	11
29	Separation, structural determination and biological evaluation of the thymidylate synthase inhibitor 3,3-di-(4-hydroxyphenyl)-6-(7-chloro-1-oxo-1,3-dihydro-1H-naphtho[1,8-cd]pyrazolo[4,5-b]pyridin-2-yl)pyrazolo[4,5-b]pyridine. <i>Journal of Heterocyclic Chemistry</i> , 1999, 36, 1043-1048.		
30	In silico identification and experimental validation of hits active against KPC-2 \hat{I}^2 -lactamase. <i>PLoS ONE</i> , 2018, 13, e0203241.	2.5	9
31	Phenylboronic Acids Probing Molecular Recognition against Class A and Class C \hat{I}^2 -lactamases. <i>Antibiotics</i> , 2019, 8, 171.	3.7	9
32	First virtual screening and experimental validation of inhibitors targeting GES-5 carbapenemase. <i>Journal of Computer-Aided Molecular Design</i> , 2019, 33, 295-305.	2.9	9
33	Naphthalimido derivatives as antifolate thymidylate synthase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 1996, 31, 1011-1016.	5.5	8
34	ortho-Halogen naphthaleins as specific inhibitors of <i>Lactobacillus casei</i> thymidylate synthase. Conformational properties and biological activity. <i>Bioorganic and Medicinal Chemistry</i> , 2003, 11, 951-963.	3.0	8
35	Conformational Propensity and Biological Studies of Proline Mutated LR Peptides Inhibiting Human Thymidylate Synthase and Ovarian Cancer Cell Growth. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 7374-7380.	6.4	6
36	Can We Exploit \hat{I}^2 -Lactamases Intrinsic Dynamics for Designing More Effective Inhibitors?. <i>Antibiotics</i> , 2020, 9, 833.	3.7	6

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37	Novel Targets and Mechanisms in Antimicrobial Drug Discovery. <i>Antibiotics</i> , 2021, 10, 141.	3.7	5
38	Constrained Dansyl Derivatives Reveal Bacterial Specificity of Highly Conserved Thymidylate Synthases. <i>ChemBioChem</i> , 2008, 9, 779-790.	2.6	4
39	A step further in the discovery of phthalein derivatives as Thymidylate Synthase inhibitors. <i>Arkivoc</i> , 2004, 2004, 382-396.	0.5	4
40	Protocetraric and Salazinic Acids as Potential Inhibitors of SARS-CoV-2 3CL Protease: Biochemical, Cytotoxic, and Computational Characterization of Depsidones as Slow-Binding Inactivators. <i>Pharmaceuticals</i> , 2022, 15, 714.	3.8	2
41	Targeting the Class A Carbapenemase GES-5 via Virtual Screening. <i>Biomolecules</i> , 2020, 10, 304.	4.0	1
42	Crystallographic studies of novel inhibitors of Î²-lactamases. <i>Acta Crystallographica Section A: Foundations and Advances</i> , 2005, 61, c248-c248.	0.3	0