

Patricia Busca

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

521
citations

687363

13
h-index

642732

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all docs

30
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30
times ranked

717
citing authors

#	ARTICLE	IF	CITATIONS
1	Tyrosine kinase inhibitor NVP-BGJ398 functionally improves FGFR3-related dwarfism in mouse model. <i>Journal of Clinical Investigation</i> , 2016, 126, 1871-1884.	8.2	84
2	Enantioselective synthesis of non-natural amino acids using phenylalanine dehydrogenases modified by site-directed mutagenesis. <i>Organic and Biomolecular Chemistry</i> , 2004, 2, 2684.	2.8	54
3	Synthesis and biological evaluation of a triazole-based library of pyrido[2,3-d]pyrimidines as FGFR3 tyrosine kinase inhibitors. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 2164.	2.8	53
4	Idiosyncratic features in tRNAs participating in bacterial cell wall synthesis. <i>Nucleic Acids Research</i> , 2007, 35, 6870-6883.	14.5	42
5	A novel tyrosine kinase inhibitor restores chondrocyte differentiation and promotes bone growth in a gain-of-function <i>Fgfr3</i> mouse model. <i>Human Molecular Genetics</i> , 2012, 21, 841-851.	2.9	40
6	Decoding the Logic of the tRNA Regiospecificity of Nonribosomal FemX _{WV} Aminoacyl Transferase. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 5115-5119.	13.8	26
7	Microwave-assisted preparation of 4-amino-3-cyano-5-methoxycarbonyl-N-arylpyrazoles as building blocks for the diversity-oriented synthesis of pyrazole-based polycyclic scaffolds. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 409-423.	2.8	24
8	Synthesis and biological evaluation of new UDP-GalNAc analogues for the study of polypeptide- β -GalNAc-transferases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 1853-1856.	2.2	21
9	A convenient synthesis of β - and β -d-glucosamine-1-phosphate and derivatives. <i>Tetrahedron Letters</i> , 1998, 39, 8101-8104.	1.4	20
10	Synthesis of 2 β -O,3 β -O bicyclic adenosine analogues using ring closing metathesis. <i>Tetrahedron Letters</i> , 2003, 44, 9131-9134.	1.4	20
11	Copper(I)/Copper(II)-Assisted Tandem Catalysis: The Case Study of Ullmann/Chan-Evans-Lam N ¹ ,N ³ -Diarylation of 3-Aminopyrazole. <i>ChemCatChem</i> , 2015, 7, 2433-2436.	3.7	19
12	Regioselective Functionalization of Quinolines through C-H Activation: A Comprehensive Review. <i>Molecules</i> , 2021, 26, 5467.	3.8	15
13	Synthesis of UDP-GalNAc analogues as probes for the study of polypeptide- β -GalNAc-transferases. Part 2. <i>Tetrahedron Letters</i> , 2004, 45, 4433-4436.	1.4	13
14	Synthesis of C-Nucleosidic ATP Mimics as Potential FGFR3 Inhibitors. <i>European Journal of Organic Chemistry</i> , 2006, 2006, 2403-2409.	2.4	13
15	Synthesis of purin-2-yl and purin-6-yl-aminoglucitols as C-nucleosidic ATP mimics and biological evaluation as FGFR3 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 1254-1262.	5.5	11
16	Synthesis and biological evaluation of chemical tools for the study of Dolichol Linked Oligosaccharide Diphosphatase (DLODP). <i>European Journal of Medicinal Chemistry</i> , 2017, 125, 952-964.	5.5	11
17	Demonstration of an oligosaccharide-diphosphodolichol diphosphatase activity whose subcellular localization is different than those of dolichyl-phosphate-dependent enzymes of the dolichol cycle. <i>Journal of Lipid Research</i> , 2016, 57, 1029-1042.	4.2	10
18	Reductive Cleavage of Aromatic and Heteroaromatic Ester Functions via Copper-Catalyzed Proto-Decarbomethoxylation. <i>Organic Letters</i> , 2018, 20, 2724-2727.	4.6	9

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19	Discovery, SAR study and ADME properties of methyl 4-amino-3-cyano-1-(2-benzyloxyphenyl)-1 <i>H</i> -pyrazole-5-carboxylate as an HIV-1 replication inhibitor. RSC Medicinal Chemistry, 2020, 11, 577-582.	3.9	8
20	Destabilization of the human RED ^{SMU1} splicing complex as a basis for host-directed antiinfluenza strategy. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 10968-10977.	7.1	7
21	Brefeldin A promotes the appearance of oligosaccharyl phosphates derived from Glc3Man9GlcNAc2-PP-dolichol within the endomembrane system of HepG2 cells. Journal of Lipid Research, 2016, 57, 1477-1491.	4.2	5
22	Synthesis of Multifunctionalized 2-Iminothiazolidin-4-ones and Their 2-Arylimino Derivatives. Synthesis, 2016, 48, 4569-4579.	2.3	4
23	Molecular modeling study of the induced-fit effect on kinase inhibition: the case of fibroblast growth factor receptor 3 (FGFR3). Journal of Computer-Aided Molecular Design, 2015, 29, 619-641.	2.9	3
24	Bacterial Lipid II Analogs: Novel In Vitro Substrates for Mammalian Oligosaccharyl Diphosphodolichol Diphosphatase (DLODP) Activities. Molecules, 2019, 24, 2135.	3.8	1
25	Synthetic Route to Glycosyl ¹² C-(phosphino)-phosphonates as Unprecedented Stable Glycosyl Diphosphate Analogs and Their Preliminary Biological Evaluation. Molecules, 2020, 25, 4969.	3.8	1
26	Enantioselective Synthesis of Non-Natural Amino Acids Using Phenylalanine Dehydrogenases Modified by Site-Directed Mutagenesis.. ChemInform, 2005, 36, no.	0.0	0