

Beatriz Baragaña

List of Publications by Year in descending order

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26
papers

1,411
citations

471509

17
h-index

526287

27
g-index

30
all docs

30
docs citations

30
times ranked

2089
citing authors

#	ARTICLE	IF	CITATIONS
1	Chemogenomics identifies acetyl-coenzyme A synthetase as a target for malaria treatment and prevention. <i>Cell Chemical Biology</i> , 2022, 29, 191-201.e8.	5.2	39
2	High-Throughput Screening Platform To Identify Inhibitors of Protein Synthesis with Potential for the Treatment of Malaria. <i>Antimicrobial Agents and Chemotherapy</i> , 2022, 66, .	3.2	10
3	<i>Mycobacterium tuberculosis</i> Phe-tRNA synthetase: structural insights into tRNA recognition and aminoacylation. <i>Nucleic Acids Research</i> , 2021, 49, 5351-5368.	14.5	1
4	MalDA, Accelerating Malaria Drug Discovery. <i>Trends in Parasitology</i> , 2021, 37, 493-507.	3.3	51
5	Prioritization of Molecular Targets for Antimalarial Drug Discovery. <i>ACS Infectious Diseases</i> , 2021, 7, 2764-2776.	3.8	35
6	Substituted Aminoacetamides as Novel Leads for Malaria Treatment. <i>ChemMedChem</i> , 2019, 14, 1329-1335.	3.2	5
7	Lysyl-tRNA synthetase as a drug target in malaria and cryptosporidiosis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 7015-7020.	7.1	94
8	Validation of Plasmodium falciparum dUTPase as the target of 5 ² -tritylated deoxyuridine analogues with anti-malarial activity. <i>Malaria Journal</i> , 2019, 18, 392.	2.3	7
9	Challenges and recent progress in drug discovery for tropical diseases. <i>Nature</i> , 2018, 559, 498-506.	27.8	164
10	Biochemical and Structural Characterization of Selective Allosteric Inhibitors of the Plasmodium falciparum Drug Target, Prolyl-tRNA-synthetase. <i>ACS Infectious Diseases</i> , 2017, 3, 34-44.	3.8	45
11	Screening a protein kinase inhibitor library against Plasmodium falciparum. <i>Malaria Journal</i> , 2017, 16, 446.	2.3	12
12	Discovery of a Quinoline-4-carboxamide Derivative with a Novel Mechanism of Action, Multistage Antimalarial Activity, and Potent in Vivo Efficacy. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9672-9685.	6.4	66
13	Trisubstituted Pyrimidines as Efficacious and Fast-Acting Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 6101-6120.	6.4	13
14	A novel multiple-stage antimalarial agent that inhibits protein synthesis. <i>Nature</i> , 2015, 522, 315-320.	27.8	353
15	2-Branched acyclic nucleoside analogues as inhibitors of Plasmodium falciparum dUTPase. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 2378-2391.	3.0	24
16	Design, Synthesis, and Evaluation of 5 ² -Diphenyl Nucleoside Analogues as Inhibitors of the Plasmodium falciparum dUTPase. <i>ChemMedChem</i> , 2011, 6, 1816-1831.	3.2	30
17	Selective delivery of 2-hydroxy APA to Trypanosoma brucei using the melamine motif. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 4364-4366.	2.2	14
18	Differentially-protected steroidal triamines; scaffolds with potential for medicinal, supramolecular, and combinatorial chemistry. <i>Organic and Biomolecular Chemistry</i> , 2004, 2, 3320-3328.	2.8	54

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19	Protease-Catalyzed Peptide Synthesis on Solid Support. <i>Journal of the American Chemical Society</i> , 2002, 124, 10988-10989.	13.7	107
20	Enantioselective Transport by a Steroidal Guanidinium Receptor. <i>Chemistry - A European Journal</i> , 2002, 8, 2931.	3.3	64
21	Synthesis of allylamines in enantiomerically pure form. <i>Tetrahedron Letters</i> , 2000, 41, 4361-4362.	1.4	7
22	Synthesis of Enantiopure α -Amino β -Epoxy Ketones from α -Amino Bromomethyl Ketones. <i>Journal of Organic Chemistry</i> , 1999, 64, 5048-5052.	3.2	27
23	Preparation and Synthetic Applications of Enantiopure (2S,3S)- or (2R,3S)-2-Halomethyl-1,2-epoxyalkan-3-amines. <i>Journal of Organic Chemistry</i> , 1999, 64, 2843-2846.	3.2	16
24	Synthetic Applications of 1-Aminoalkyl Chloromethyl Ketones. Synthesis of Enantiopure 3-Azetidinols and Aminoalkyl Epoxides. <i>Journal of Organic Chemistry</i> , 1997, 62, 5974-5977.	3.2	23
25	High Diastereoselective Synthesis of Threo or Erythro Aminoalkyl Epoxides from α -Amino Acids. <i>Journal of Organic Chemistry</i> , 1995, 60, 6696-6699.	3.2	98
26	The first direct preparation of chiral functionalised ketones and their synthetic uses. <i>Journal of the Chemical Society Chemical Communications</i> , 1994, , 969-970.	2.0	48