

# Minghao Xu

## List of Publications by Year in descending order

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17  
papers

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citations

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18  
docs citations

18  
times ranked

1008  
citing authors

#	ARTICLE	IF	CITATIONS
1	Bioorthogonal chemistry. <i>Nature Reviews Methods Primers</i> , 2021, 1, .	11.8	201
2	Bioorthogonal Removal of 3-Isocyanopropyl Groups Enables the Controlled Release of Fluorophores and Drugs in Vivo. <i>Journal of the American Chemical Society</i> , 2018, 140, 8410-8414.	6.6	103
3	Dissociative Bioorthogonal Reactions. <i>ChemBioChem</i> , 2019, 20, 1615-1627.	1.3	61
4	Rapid and efficient tetrazine-induced drug release from highly stable benzonorbornadiene derivatives. <i>Chemical Communications</i> , 2017, 53, 6271-6274.	2.2	55
5	Discovery and Rational Design of Natural-Product-Derived 2-Phenyl-3,4-dihydro-2 <i>H</i> -benzo[ <i>f</i> ]chromen-3-amine Analogs as Novel and Potent Dipeptidyl Peptidase 4 (DPP-4) Inhibitors for the Treatment of Type 2 Diabetes. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 6772-6790.	2.9	49
6	Discovery of Diverse Human Dihydroorotate Dehydrogenase Inhibitors as Immunosuppressive Agents by Structure-Based Virtual Screening. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 8341-8349.	2.9	47
7	Novel Selective and Potent Inhibitors of Malaria Parasite Dihydroorotate Dehydrogenase: Discovery and Optimization of Dihydrothiophenone Derivatives. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 7911-7924.	2.9	47
8	Design, Synthesis, X-ray Crystallographic Analysis, and Biological Evaluation of Thiazole Derivatives as Potent and Selective Inhibitors of Human Dihydroorotate Dehydrogenase. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 1123-1139.	2.9	47
9	Rationally Designed Multitarget Anticancer Agents. <i>Current Medicinal Chemistry</i> , 2013, 20, 1694-1714.	1.2	42
10	Isonitrile-responsive and bioorthogonally removable tetrazine protecting groups. <i>Chemical Science</i> , 2020, 11, 169-179.	3.7	41
11	Dissociative reactions of benzonorbornadienes with tetrazines: scope of leaving groups and mechanistic insights. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 9855-9865.	1.5	28
12	Tuning Isonitrile/Tetrazine Chemistry for Accelerated Deprotection and Formation of Stable Conjugates. <i>Journal of Organic Chemistry</i> , 2019, 84, 15520-15529.	1.7	22
13	Rational Design of Benzylidenehydrazinyl-Substituted Thiazole Derivatives as Potent Inhibitors of Human Dihydroorotate Dehydrogenase with in Vivo Anti-arthritic Activity. <i>Scientific Reports</i> , 2015, 5, 14836.	1.6	19
14	Synthesis, activity evaluation, and docking analysis of barbituric acid aryl hydrazone derivatives as RSK2 inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013, 28, 747-752.	2.5	6
15	Cytotoxicity and DNA Binding Ability of Two Novel Gold(III) Complexes. <i>Journal of Applied Spectroscopy</i> , 2019, 86, 618-622.	0.3	6
16	A Stable Precursor for Bioorthogonally Removable 3-Isocyanopropoxyloxycarbonyl (ICPrC) Protecting Groups. <i>Synlett</i> , 2020, 31, 1701-1706.	1.0	1
17	Abstract 4772: Novel drug discovery approach targeting STAT3 for breast cancer therapy using MLSD and drug repositioning. , 2012, , .		1