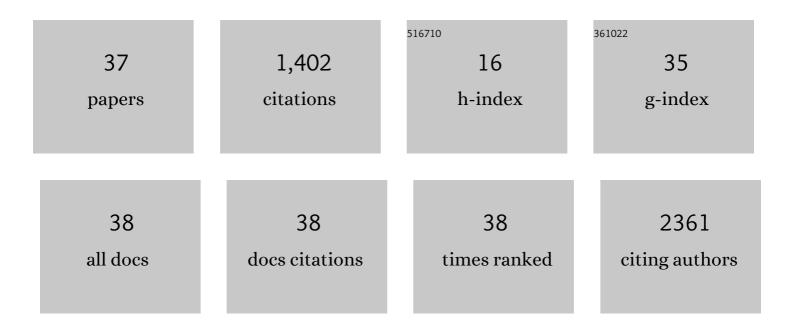
Xin Chen

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Visualization of arrestin recruitment by a G-protein-coupled receptor. Nature, 2014, 512, 218-222.	27.8	433
2	Mechanism of intracellular allosteric β2AR antagonist revealed by X-ray crystal structure. Nature, 2017, 548, 480-484.	27.8	148
3	Allosteric "beta-blocker―isolated from a DNA-encoded small molecule library. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 1708-1713.	7.1	118
4	The Plant Phenolic Compound <i>p</i> -Coumaric Acid Represses Gene Expression in the <i>Dickeya dadantii</i> Type III Secretion System. Applied and Environmental Microbiology, 2009, 75, 1223-1228.	3.1	96
5	Derivatives of Plant Phenolic Compound Affect the Type III Secretion System of Pseudomonas aeruginosa via a GacS-GacA Two-Component Signal Transduction System. Antimicrobial Agents and Chemotherapy, 2012, 56, 36-43.	3.2	75
6	Discovery of Plant Phenolic Compounds That Act as Type III Secretion System Inhibitors or Inducers of the Fire Blight Pathogen, Erwinia amylovora. Applied and Environmental Microbiology, 2013, 79, 5424-5436.	3.1	71
7	Small-Molecule Positive Allosteric Modulators of the <i>β</i> ₂ -Adrenoceptor Isolated from DNA-Encoded Libraries. Molecular Pharmacology, 2018, 94, 850-861.	2.3	66
8	Small-Molecule Inhibitors of the Type III Secretion System. Molecules, 2015, 20, 17659-17674.	3.8	48
9	Synthesis and biological evaluation of novel 4î²-(1,3,4-oxadiazole-2-amino)-podophyllotoxin derivatives. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 4778-4782.	2.2	41
10	Chemoprevention of 7,12-dimethylbenz[<i>a</i>]anthracene (DMBA)-induced Hamster Cheek Pouch Carcinogenesis by a 5-Lipoxygenase Inhibitor, Garcinol. Nutrition and Cancer, 2012, 64, 1211-1218.	2.0	40
11	Derivative of plant phenolic compound inhibits the type <scp>III</scp> secretion system of <i><scp>D</scp>ickeya dadantii</i> via <scp>HrpX</scp> / <scp>HrpY</scp> twoâ€component signal transduction and <scp>R</scp> sm systems. Molecular Plant Pathology, 2015, 16, 150-163.	4.2	33
12	Synthesis and biological evaluation of sulforaphane derivatives as potential antitumor agents. European Journal of Medicinal Chemistry, 2013, 64, 529-539.	5.5	29
13	New Method for the Synthesis of Sulforaphane and Related Isothiocyanates. Synthesis, 2011, 2011, 3991-3996.	2.3	25
14	DeSiphering receptor core-induced and ligand-dependent conformational changes in arrestin via genetic encoded trimethylsilyl 1H-NMR probe. Nature Communications, 2020, 11, 4857.	12.8	25
15	13,14-Dihydroxy groups are critical for the anti-cancer effects of garcinol. Bioorganic Chemistry, 2015, 60, 123-129.	4.1	21
16	The C8 side chain is one of the key functional group of Garcinol for its anti-cancer effects. Bioorganic Chemistry, 2017, 71, 74-80.	4.1	20
17	Visible-Light-Mediated Cyclopropanation Reactions of 3-Diazooxindoles with Arenes. Journal of Organic Chemistry, 2021, 86, 7131-7140.	3.2	16
18	Synthesis and antitumor activity of formononetin nitrogen mustard derivatives. European Journal of Medicinal Chemistry, 2012, 54, 175-187.	5.5	14

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19	Syntheses, structures and luminescence properties of two series of 3D lanthanide coordination polymers based on benzimidazole-5,6-dicarboxylic acid and oxalate. Inorganic Chemistry Communication, 2012, 23, 74-77.	3.9	10
20	Synthesis and antitumor activity of novel 4-aminoquinoline derivatives. Medicinal Chemistry Research, 2013, 22, 2855-2861.	2.4	8
21	Two unusual 12-connected metal–organic coordination polymers with fcu net. Journal of Solid State Chemistry, 2013, 205, 110-115.	2.9	7
22	Alternate Synthesis of HSP90 Inhibitor AT13387. Synthetic Communications, 2014, 44, 2416-2425.	2.1	6
23	Design, synthesis, and functional assessment of Cmpd-15 derivatives as negative allosteric modulators for the β2-adrenergic receptor. Bioorganic and Medicinal Chemistry, 2018, 26, 2320-2330.	3.0	6
24	Synthesis of Fmoc-protected (<i>S</i>)-3,5-Dibromophenylalanine in the Presence of a Phase Transfer Catalyst or a Chiral Catalyst. Chinese Journal of Organic Chemistry, 2016, 36, 2242.	1.3	6
25	A rare 4-connected <i>neb</i> topological framework based on Zn(II). Journal of Coordination Chemistry, 2013, 66, 2843-2851.	2.2	5
26	An ionic lock and a hydrophobic zipper mediate the coupling between an insect pheromone receptor BmOR3 and downstream effectors. Journal of Biological Chemistry, 2021, 297, 101160.	3.4	5
27	Discovery of novel cholic acid derivatives as highly potent agonists for G protein-coupled bile acid receptor. Bioorganic Chemistry, 2022, 120, 105588.	4.1	5
28	Synthesis of Both Enantiomers of Chiral Phenylalanine Derivatives Catalyzed by Cinchona Alkaloid Quaternary Ammonium Salts as Asymmetric Phase Transfer Catalysts. Molecules, 2018, 23, 1421.	3.8	4
29	Asymmetric synthesis of multifunctional aryl allyl ethers by nucleophilic catalysis. RSC Advances, 2019, 9, 11585-11588.	3.6	4
30	Synthesis and Bioactivity of Novel Inhibitors for Type III Secretion System of <i>Pseudomonas aeruginosa</i> PAO1. Chinese Journal of Organic Chemistry, 2013, 33, 1309.	1.3	4
31	Total synthesis of (3 <i>Z</i> ,9 <i>Z</i> ,6 <i>S</i> ,7 <i>R</i>) and (3 <i>Z</i> ,9 <i>Z</i> ,6 <i>R</i> ,7 <i>S</i>)-6,7-epoxy-3,9-octadecadienes. Synthetic Communications, 2017, 47, 1848-1853.	2.1	3
32	Cell active and functionally-relevant small-molecule agonists of calcitonin receptor. Bioorganic Chemistry, 2020, 96, 103596.	4.1	3
33	Two zinc(II) coordination polymers based on a benzenedicarboxylic acid derivative: Synthesis, crystal structures and luminescent properties. Inorganic Chemistry Communication, 2012, 18, 29-33.	3.9	2
34	Synthesis, Biological Evaluation, and Docking Study of Ringâ€Opening Amide Analogs of Matrine as Antitumor Agents. Chemistry and Biodiversity, 2021, 18, e2000979.	2.1	2
35	Synthesis of Bitopic Ligands as Potent Dopamine D ₂ Receptor Agonists. ChemMedChem, 2022, 17, .	3.2	2
36	New method for synthesis of EZH2 methyltransferase inhibitor GSK126. Synthetic Communications, 2016, 46, 1215-1222.	2.1	1

#	Article	IF	CITATIONS
37	Enantioselective benzylation of methyl 4-oxo-3-piperidinecarboxylate with cinchona alkaloids phase-transfer catalysts. Synthetic Communications, 2018, 48, 2260-2271.	2.1	0