

Shengzheng Wang

List of Publications by Year in descending order

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26
papers

987
citations

471371

17
h-index

526166

27
g-index

28
all docs

28
docs citations

28
times ranked

1469
citing authors

#	ARTICLE	IF	CITATIONS
1	Discovery of novel tubulin inhibitors targeting the colchicine binding site via virtual screening, structural optimization and antitumor evaluation. <i>Bioorganic Chemistry</i> , 2022, 118, 105486.	2.0	11
2	Recent Progress in the Discovery of Antifungal Agents Targeting the Cell Wall. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 12429-12459.	2.9	37
3	Resveratrol: Multi-Targets Mechanism on Neurodegenerative Diseases Based on Network Pharmacology. <i>Frontiers in Pharmacology</i> , 2020, 11, 694.	1.6	25
4	Antifungal Activity of Crude Extract from the Rhizome and Root of <i>Smilacina japonica</i> A. Gray. Evidence-based Complementary and Alternative Medicine, 2019, 2019, 1-9.	0.5	1
5	Structural simplification: an efficient strategy in lead optimization. <i>Acta Pharmaceutica Sinica B</i> , 2019, 9, 880-901.	5.7	49
6	Design, synthesis and biological evaluation of novel antitumor spiro-dihydrothiopyran-oxindole derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 1636-1642.	1.0	15
7	Organocatalytic asymmetric synthesis of highly functionalized spiro-thiazolone-cyclopropane-oxindoles bearing two vicinal spiro quaternary centers. <i>Organic Chemistry Frontiers</i> , 2019, 6, 1442-1447.	2.3	19
8	Structural Simplification of Natural Products. <i>Chemical Reviews</i> , 2019, 119, 4180-4220.	23.0	157
9	Synthesis of spiro-tetrahydrothiopyran-oxindoles by Michael-aldol cascade reactions: discovery of potential P53-MDM2 inhibitors with good antitumor activity. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 625-634.	1.5	25
10	Organocatalytic Asymmetric Synthesis of Spiro-tetrahydrothiophene Oxindoles Bearing Four Contiguous Stereocenters by One-Pot Michael-Henry-Cascade-Rearrangement Reactions. <i>Chemistry - A European Journal</i> , 2018, 24, 62-66.	1.7	24
11	Design, synthesis and biological evaluation of novel antitumor spiro-tetrahydrothiopyran-oxindole derivatives as potent p53-MDM2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 5268-5277.	1.4	17
12	Facile construction of pyrrolo[1,2-b]isoquinolin-10(5H)-ones via a redox-amination-aromatization-Friedel-Crafts acylation cascade reaction and discovery of novel topoisomerase inhibitors. <i>Chemical Communications</i> , 2016, 52, 9593-9596.	2.2	8
13	Meeting Organocatalysis with Drug Discovery: Asymmetric Synthesis of 3,3-Spirooxindoles Fused with Tetrahydrothiopyrans as Novel p53-MDM2 Inhibitors. <i>Organic Letters</i> , 2016, 18, 1028-1031.	2.4	77
14	Divergent Cascade Construction of Skeletally Diverse Privileged-Pyrazole-Derived Molecular Architectures. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 2030-2037.	1.2	67
15	Scaffold Diversity Inspired by the Natural Product Evodiamine: Discovery of Highly Potent and Multitargeting Antitumor Agents. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6678-6696.	2.9	156
16	Facile Assembly of Chiral Tetrahydrothiopyrans Containing Four Consecutive Stereocenters via an Organocatalytic Enantioselective Michael-Michael Cascade. <i>Organic Letters</i> , 2014, 16, 692-695.	2.4	24
17	Novel Carboline Derivatives as Potent Antifungal Lead Compounds: Design, Synthesis, and Biological Evaluation. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 506-511.	1.3	49
18	Discovery of 1-arylpyrrolidone derivatives as potent p53-MDM2 inhibitors based on molecule fusing strategy. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 2648-2650.	1.0	14

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19	Design, synthesis and antifungal activity of novel triazole derivatives containing substituted 1,2,3-triazole-piperidine side chains. <i>European Journal of Medicinal Chemistry</i> , 2014, 82, 490-497.	2.6	68
20	Design, synthesis and biological activity of piperlongumine derivatives as selective anticancer agents. <i>European Journal of Medicinal Chemistry</i> , 2014, 82, 545-551.	2.6	33
21	Asymmetric Synthesis of Chiral Dihydrothiopyrans via an Organocatalytic Enantioselective Formal Thio [3 + 3] Cycloaddition Reaction with Binucleophilic Bisketone Thioethers. <i>Organic Letters</i> , 2013, 15, 5570-5573.	2.4	31
22	Structure-activity relationships of tetrahydrocarbazole derivatives as antifungal lead compounds. <i>MedChemComm</i> , 2013, 4, 353-362.	3.5	13
23	Facile Construction of Structurally Diverse Thiazolidinedione-Derived Compounds via Divergent Stereoselective Cascade Organocatalysis and Their Biological Exploratory Studies. <i>ACS Combinatorial Science</i> , 2013, 15, 298-308.	3.8	41
24	Asymmetric Synthesis, Antifungal Activity and Molecular Modeling of Iodiconazole Isomers. <i>Chinese Journal of Chemistry</i> , 2013, 31, 1139-1143.	2.6	2
25	Design, synthesis and structure-activity relationships of new triazole derivatives containing N-substituted phenoxypropylamino side chains. <i>European Journal of Medicinal Chemistry</i> , 2012, 53, 292-299.	2.6	15
26	Discovery of highly potent antifungal triazoles by structure-based lead fusion. <i>MedChemComm</i> , 2011, 2, 1066.	3.5	8