

# Gao-Feng Zha

## List of Publications by Year in descending order

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29  
papers

1,124  
citations

448610

19  
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536525

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docs citations

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times ranked

1348  
citing authors

#	ARTICLE	IF	CITATIONS
1	Benzimidazole analogues as efficient arsenals in war against methicillin-resistance staphylococcus aureus (MRSA) and its SAR studies. <i>Bioorganic Chemistry</i> , 2021, 115, 105175.	2.0	49
2	Structure-activity relationships (SAR) of triazine derivatives: Promising antimicrobial agents. <i>European Journal of Medicinal Chemistry</i> , 2020, 185, 111804.	2.6	80
3	Stereoselective Construction of Nitrile-Substituted Cyclopropanes from 2-Substituted Ethenesulfonyl Fluorides <i>via</i> Carbon-Sulfur Bond Cleavage. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 4596-4601.	2.1	16
4	A SO <sub>2</sub> F <sub>2</sub> mediated mild, practical, and gram-scale dehydroxylative transforming primary alcohols to quaternary ammonium salts. <i>Tetrahedron</i> , 2019, 75, 4648-4656.	1.0	6
5	Making Carbonyls of Amides Nucleophilic and Hydroxyls of Alcohols Electrophilic Mediated by SO <sub>2</sub> F <sub>2</sub> for Synthesis of Esters from Amides. <i>Organic Letters</i> , 2019, 21, 8657-8661.	2.4	17
6	Regioselective installation of fluorosulfate (OSO <sub>2</sub> F) functionality into aromatic C(sp <sup>2</sup> )-H bonds for the construction of para-amino-arylfluorosulfates. <i>Chemical Communications</i> , 2019, 55, 6273-6276.	2.2	16
7	A Simple, Mild and General Oxidation of Alcohols to Aldehydes or Ketones by SO <sub>2</sub> F <sub>2</sub> /K <sub>2</sub> CO <sub>3</sub> Using DMSO as Solvent and Oxidant. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 2262-2267.	2.1	38
8	Rh-Catalyzed Highly Enantioselective Synthesis of Aliphatic Sulfonyl Fluorides. <i>IScience</i> , 2019, 21, 695-705.	1.9	32
9	Discovery of novel arylenesulfonyl fluorides as potential candidates against methicillin-resistant of <i>Staphylococcus aureus</i> (MRSA) for overcoming multidrug resistance of bacterial infections. <i>European Journal of Medicinal Chemistry</i> , 2019, 162, 364-377.	2.6	72
10	Pharmaceutical significance of azepane based motifs for drug discovery: A critical review. <i>European Journal of Medicinal Chemistry</i> , 2019, 162, 465-494.	2.6	55
11	Construction of Di(hetero)arylmethanes Through Pd-Catalyzed Direct Dehydroxylative Cross-Coupling of Benzylic Alcohols and Aryl Boronic Acids Mediated by Sulfonyl Fluoride (SO <sub>2</sub> F <sub>2</sub> ). <i>European Journal of Organic Chemistry</i> , 2019, 2019, 1801-1807.	1.2	20
12	SO <sub>2</sub> F <sub>2</sub> -Mediated Oxidative Dehydrogenation and Dehydration of Alcohols to Alkynes. <i>Journal of the American Chemical Society</i> , 2018, 140, 17666-17673.	6.6	69
13	Ethenesulfonyl fluoride derivatives as telomerase inhibitors: structure-based design, SAR, and anticancer evaluation <i>in vitro</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018, 33, 1266-1270.	2.5	19
14	Cooperativity of axial and centre chirality in the biaryl disulfoxide/Rh-catalysed asymmetric 1,4-addition of arylboronic acids to 2-cyclohexenone: a DFT study. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 2226-2233.	1.5	4
15	Discovery of potential anticancer multi-targeted ligustrazine based cyclohexanone and oxime analogs overcoming the cancer multidrug resistance. <i>European Journal of Medicinal Chemistry</i> , 2017, 135, 34-48.	2.6	61
16	Synthesis, SAR and molecular docking studies of benzo[d]thiazole-hydrazones as potential antibacterial and antifungal agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3148-3155.	1.0	70
17	Synthesis of benzo[d]thiazole-hydrazone analogues: molecular docking and SAR studies of potential H <sup>+</sup> /K <sup>+</sup> -ATPase inhibitors and anti-inflammatory agents. <i>MedChemComm</i> , 2017, 8, 1173-1189.	3.5	43
18	Palladium-Catalyzed Fluorosulfonylvinylation of Organic Iodides. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 4849-4852.	7.2	95

#	ARTICLE	IF	CITATIONS
19	Palladium-catalyzed Fluorosulfonylvinilation of Organic Iodides. <i>Angewandte Chemie</i> , 2017, 129, 4927-4930.	1.6	31
20	Synthesis of a Class of Fused $\beta$ -Sultone Heterocycles <i>via</i> DBU-catalyzed Direct Annulative SuFEx Click of Ethenesulfonyl Fluorides and Pyrazolones or 1,3-dicarbonyl Compounds. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 3254-3260.	2.1	36
21	Gram-scale Synthesis of $\beta$ -(Hetero)arylethenesulfonyl Fluorides <i>via</i> a Pd(OAc) <sub>2</sub> -catalyzed Oxidative Heck Process with DDQ or AgNO <sub>3</sub> as an Oxidant. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 3237-3242.	2.1	55
22	Biological evaluation of synthetic $\alpha,\beta$ -unsaturated carbonyl based cyclohexanone derivatives as neuroprotective novel inhibitors of acetylcholinesterase, butyrylcholinesterase and amyloid- $\beta$ aggregation. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 2352-2359.	1.4	31
23	Silver-mediated direct trifluoromethoxylation of $\beta$ -diazo esters via the <sup>+</sup> OCF <sub>3</sub> <sup>-</sup> anion. <i>Chemical Communications</i> , 2016, 52, 7458-7461.	2.2	77
24	Visible-light-mediated aerobic oxidative dimerizative annulation of $\beta$ -carbonylketones: a facile strategy to construct highly functionalized furans. <i>Tetrahedron Letters</i> , 2016, 57, 4680-4683.	0.7	7
25	Perfluoroalkanesulfonylation of Alkynyl(phenyl)iodonium Tosylates by the Weakly Nucleophilic Sodium Perfluoroalkanesulfinates. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 4119-4124.	2.1	10
26	Expeditious trifluoromethylthiolation and trifluoromethylselenolation of alkynyl(phenyl)iodoniums by [XCF <sub>3</sub> ] <sup>+</sup> (X = S, Se) anions. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 11502-11509.	1.5	52
27	[Ru(bpy) <sub>3</sub> ]Cl <sub>2</sub> -catalyzed aerobic oxidative cleavage $\beta$ -diketones to carboxylic acids under visible light irradiation. <i>Tetrahedron Letters</i> , 2016, 57, 5628-5631.	0.7	13
28	Ruthenium-catalyzed direct arylations with aryl chlorides. <i>RSC Advances</i> , 2016, 6, 30875-30885.	1.7	49
29	A simple synthetic approach for the transformation of (S)-Ugi's amine. <i>Chinese Chemical Letters</i> , 2014, 25, 1301-1304.	4.8	1