

# Biao Jiang

## List of Publications by Year in descending order

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19  
papers

527  
citations

687363  
13  
h-index

794594  
19  
g-index

19  
all docs

19  
docs citations

19  
times ranked

655  
citing authors

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Construction of an IMiD-based azide library as a kit for PROTAC research. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 166-170.  | 2.8 | 21        |
| 2  | Combretastatin A4-derived payloads for antibody-drug conjugates. <i>European Journal of Medicinal Chemistry</i> , 2021, 216, 113355.  | 5.5 | 7         |
| 3  | Structure-based discovery of SIAIS001 as an oral bioavailability ALK degrader constructed from Alectinib. <i>European Journal of Medicinal Chemistry</i> , 2021, 217, 113335.                                 | 5.5 | 26        |
| 4  | Discovery of a Brigatinib Degradar SIAIS164018 with Destroying Metastasis-Related Oncoproteins and a Reshuffling Kinome Profile. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9152-9165.                 | 6.4 | 23        |
| 5  | Accurate Retention Time Prediction Based on Monolinked Peptide Information to Confidently Identify Cross-Linked Peptides. <i>Journal of the American Society for Mass Spectrometry</i> , 2021, 32, 2410-2416. | 2.8 | 1         |
| 6  | Site-specific construction of triptolide-based antibody-drug conjugates. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 51, 116497.  | 3.0 | 5         |
| 7  | Divinylsulfonamides enable the construction of homogeneous antibody-drug conjugates. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115793.  | 3.0 | 4         |
| 8  | Decision Tree Searching Strategy to Boost the Identification of Cross-Linked Peptides. <i>Analytical Chemistry</i> , 2020, 92, 13702-13710.   | 6.5 | 3         |
| 9  | Development of a Brigatinib degrader (SIAIS117) as a potential treatment for ALK positive cancer resistance. <i>European Journal of Medicinal Chemistry</i> , 2020, 193, 112190.                              | 5.5 | 50        |
| 10 | Bis(vinylsulfonyl)piperazines as efficient linkers for highly homogeneous antibody-drug conjugates. <i>European Journal of Medicinal Chemistry</i> , 2020, 190, 112080.                                       | 5.5 | 15        |
| 11 | A novel mass spectrometry-cleavable, phosphate-based enrichable and multi-targeting protein cross-linker. <i>Chemical Science</i> , 2019, 10, 6443-6447.  | 7.4 | 19        |
| 12 | Chemoselective Synthesis of Lenalidomide-Based PROTAC Library Using Alkylation Reaction. <i>Organic Letters</i> , 2019, 21, 3838-3841.  | 4.6 | 48        |
| 13 | H <sub>2</sub> O-Regulated Chemoselectivity in Oxa- Versus Aza-Michael Reactions. <i>Organic Letters</i> , 2019, 21, 4159-4162.   | 4.6 | 9         |
| 14 | N-Phenyl-N-acetylvinylsulfonamides as Efficient and Chemoselective Handles for N-Terminal Modification of Peptides and Proteins. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 829-836.          | 2.4 | 16        |
| 15 | Proteolysis Targeting Chimeras (PROTACs) of Anaplastic Lymphoma Kinase (ALK). <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 304-314.  | 5.5 | 165       |
| 16 | N-Methyl-N-phenylvinylsulfonamides for Cysteine-Selective Conjugation. <i>Organic Letters</i> , 2018, 20, 6526-6529.  | 4.6 | 20        |
| 17 | Intramolecular Aza-Piancatelli Rearrangement of Alkyl- or Arylamines Promoted by PPh <sub>3</sub> /Diethyl Azodicarboxylate. <i>Organic Letters</i> , 2017, 19, 1028-1031.                                    | 4.6 | 35        |
| 18 | Selective lysine modification of native peptides via aza-Michael addition. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 7339-7345.   | 2.8 | 28        |

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 19 | Divinylsulfonamides as Specific Linkers for Stapling Disulfide Bonds in Peptides. Organic Letters, 2017, 19, 4972-4975. | 4.6 | 32        |