Iwao Ojima

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36 papers 1,204 18 h-index g-index

37 ext. papers ext. citations 4.9 avg, IF L-index

| # | Paper | IF | Citations |
|----|---|------|-----------|
| 36 | Fatty acid-binding proteins (FABPs) are intracellular carriers for B -tetrahydrocannabinol (THC) and cannabidiol (CBD). <i>Journal of Biological Chemistry</i> , 2015 , 290, 8711-21 | 5.4 | 150 |
| 35 | Exploration of fluorine chemistry at the multidisciplinary interface of chemistry and biology. Journal of Organic Chemistry, 2013 , 78, 6358-83 | 4.2 | 149 |
| 34 | Taxane anticancer agents: a patent perspective. Expert Opinion on Therapeutic Patents, 2016, 26, 1-20 | 6.8 | 119 |
| 33 | Identification of a New Class of Antifungals Targeting the Synthesis of Fungal Sphingolipids. <i>MBio</i> , 2015 , 6, e00647 | 7.8 | 94 |
| 32 | Inhibition of fatty acid binding proteins elevates brain anandamide levels and produces analgesia. <i>PLoS ONE</i> , 2014 , 9, e94200 | 3.7 | 85 |
| 31 | Poly(2-oxazoline) based micelles with high capacity for 3rd generation taxoids: preparation, in vitro and in vivo evaluation. <i>Journal of Controlled Release</i> , 2015 , 208, 67-75 | 11.7 | 68 |
| 30 | Recent advances in the discovery and development of antibacterial agents targeting the cell-division protein FtsZ. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 6354-6369 | 3.4 | 58 |
| 29 | SAR studies on trisubstituted benzimidazoles as inhibitors of Mtb FtsZ for the development of novel antitubercular agents. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 9756-70 | 8.3 | 55 |
| 28 | Drug discovery targeting cell division proteins, microtubules and FtsZ. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 5060-77 | 3.4 | 55 |
| 27 | Recent progress in the strategic incorporation of fluorine into medicinally active compounds. Journal of Fluorine Chemistry, 2019 , 217, 29-40 | 2.1 | 43 |
| 26 | Acylhydrazones as Antifungal Agents Targeting the Synthesis of Fungal Sphingolipids. <i>Antimicrobial Agents and Chemotherapy</i> , 2018 , 62, | 5.9 | 39 |
| 25 | Design, synthesis and evaluation of novel 2,5,6-trisubstituted benzimidazoles targeting FtsZ as antitubercular agents. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 2602-12 | 3.4 | 35 |
| 24 | Benzimidazole-based antibacterial agents against Francisella tularensis. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 3318-26 | 3.4 | 28 |
| 23 | Quest for Efficacious Next-Generation Taxoid Anticancer Agents and Their Tumor-Targeted Delivery. <i>Journal of Natural Products</i> , 2018 , 81, 703-721 | 4.9 | 23 |
| 22 | Fluorine-Containing Taxoid Anticancer Agents and Their Tumor-Targeted Drug Delivery. <i>Journal of Fluorine Chemistry</i> , 2013 , 152, 157-165 | 2.1 | 23 |
| 21 | Design, Synthesis and Application of Fluorine-Labeled Taxoids as F NMR Probes for the Metabolic Stability Assessment of Tumor-Targeted Drug Delivery Systems. <i>Journal of Fluorine Chemistry</i> , 2015 , 171, 148-161 | 2.1 | 23 |
| 20 | Targeting the Hemopexin-like Domain of Latent Matrix Metalloproteinase-9 (proMMP-9) with a Small Molecule Inhibitor Prevents the Formation of Focal Adhesion Junctions. <i>ACS Chemical Biology</i> , 2017 , 12, 2788-2803 | 4.9 | 21 |

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| 19 | Design, Synthesis, and Biological Evaluations of Asymmetric Bow-Tie PAMAM Dendrimer-Based Conjugates for Tumor-Targeted Drug Delivery. <i>ACS Omega</i> , 2018 , 3, 3717-3736 | 3.9 | 21 |
|----|---|---------------------|-----|
| 18 | Design, synthesis and biological evaluation of a highly-potent and cancer cell selective folate-taxoid conjugate. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2187-94 | 3.4 | 15 |
| 17 | Taxol Analogues Exhibit Differential Effects on Photoaffinity Labeling of Tubulin and the Multidrug Resistance Associated P-Glycoprotein. <i>Journal of Natural Products</i> , 2018 , 81, 600-606 | 4.9 | 14 |
| 16 | SAR studies on truxillic acid mono esters as a new class of antinociceptive agents targeting fatty acid binding proteins. <i>European Journal of Medicinal Chemistry</i> , 2018 , 154, 233-252 | 6.8 | 14 |
| 15 | Computer-aided identification, synthesis, and biological evaluation of novel inhibitors for botulinum neurotoxin serotype A. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 5489-95 | 3.4 | 11 |
| 14 | Synthesis of Colchicinoids and Allocolchicinoids through Rh(I)-Catalyzed [2+2+2+1] and [2+2+2] Cycloadditions of o-Phenylenetriynes with and without CO. <i>Journal of Organic Chemistry</i> , 2018 , 83, 116. | 2 3 -116 | 549 |
| 13 | SAR Studies on Aromatic Acylhydrazone-Based Inhibitors of Fungal Sphingolipid Synthesis as Next-Generation Antifungal Agents. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 8249-8273 | 8.3 | 9 |
| 12 | Contribution of diacylglycerol lipase Ito pain after surgery. <i>Journal of Pain Research</i> , 2018 , 11, 473-482 | 2.9 | 9 |
| 11 | Substituents at the C3Zand C3ZN positions are critical for taxanes to overcome acquired resistance of cancer cells to paclitaxel. <i>Toxicology and Applied Pharmacology</i> , 2018 , 347, 79-91 | 4.6 | 8 |
| 10 | Synthesis of a Next-Generation Taxoid by Rapid Methylation Amenable for C-Labeling. <i>Journal of Organic Chemistry</i> , 2018 , 83, 2847-2857 | 4.2 | 5 |
| 9 | Design, synthesis and SAR study of 3rd-generation taxoids bearing 3-CH, 3-CFO and 3-CHFO groups at the C2-benzoate position. <i>Bioorganic Chemistry</i> , 2020 , 95, 103523 | 5.1 | 5 |
| 8 | Pd-catalyzed asymmetric allylic amination with BOP ligands and its applications to the synthesis of fused polycyclic alkaloids. <i>Tetrahedron Letters</i> , 2015 , 56, 3288-3292 | 2 | 4 |
| 7 | Incarvillateine produces antinociceptive and motor suppressive effects via adenosine receptor activation. <i>PLoS ONE</i> , 2019 , 14, e0218619 | 3.7 | 3 |
| 6 | A novel taxane, difluorovinyl-ortataxel, effectively overcomes paclitaxel-resistance in breast cancer cells. <i>Cancer Letters</i> , 2020 , 491, 36-49 | 9.9 | 3 |
| 5 | Structure-activity relationship studies on 2,5,6-trisubstituted benzimidazoles targeting -FtsZ as antitubercular agents. <i>RSC Medicinal Chemistry</i> , 2021 , 12, 78-94 | 3.5 | 3 |
| 4 | Computational Design and Synthesis of Novel Fluoro-Analogs of Combretastatins A-4 and A-1. <i>Journal of Fluorine Chemistry</i> , 2017 , 203, 193-199 | 2.1 | 2 |
| 3 | Structure and inhibition of Cryptococcus neoformans sterylglucosidase to develop antifungal agents. <i>Nature Communications</i> , 2021 , 12, 5885 | 17.4 | 1 |
| 2 | Design, synthesis and SAR study of Fluorine-containing 3rd-generation taxoids <i>Bioorganic Chemistry</i> , 2021 , 119, 105578 | 5.1 | О |

Potent antitumor activity of novel taxoids in anaplastic thyroid cancer. *Endocrine*, **2021**, 1

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