Niamh M O boyle

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

26 709 17 31 h-index g-index citations papers 889 35 4.72 4.9 avg, IF L-index ext. papers ext. citations

#	Paper	IF	Citations
31	Nature-derived epoxy resins: Synthesis, allergenicity, and thermosetting properties of pinoresinol diglycidyl ether <i>Toxicology and Industrial Health</i> , 2022 , 7482337221089595	1.8	O
30	Synthesis and Antiproliferative Evaluation of 3-Chloroazetidin-2-ones with Antimitotic Activity: Heterocyclic Bridged Analogues of Combretastatin A-4. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	1
29	Skin lipids in health and disease: A review. <i>Chemistry and Physics of Lipids</i> , 2021 , 236, 105055	3.7	17
28	Synthesis and Biological Evaluation of 1-(Diarylmethyl)-1-1,2,4-triazoles and 1-(Diarylmethyl)-1-imidazoles as a Novel Class of Anti-Mitotic Agent for Activity in Breast Cancer. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	2
27	Lactams with antiproliferative and antiapoptotic activity in breast and chemoresistant colon cancer cells. <i>European Journal of Medicinal Chemistry</i> , 2020 , 189, 112050	6.8	16
26	Colchicine-Binding Site Inhibitors from Chemistry to Clinic: A Review. <i>Pharmaceuticals</i> , 2020 , 13,	5.2	79
25	Azetidin-2-ones: structures of anti-mitotic compounds based on the 1-(3,4,5-tri-meth-oxy-phen-yl)azetidin-2-one core. <i>Acta Crystallographica Section E: Crystallographic Communications</i> , 2020 , 76, 1187-1194	0.7	O
24	Potent Quinoline-Containing Combretastatin A-4 Analogues: Design, Synthesis, Antiproliferative, and Anti-Tubulin Activity. <i>Pharmaceuticals</i> , 2020 , 13,	5.2	4
23	Synthesis and evaluation of antiproliferative microtubule-destabilising combretastatin A-4 piperazine conjugates. <i>Organic and Biomolecular Chemistry</i> , 2019 , 17, 6184-6200	3.9	8
22	3-Vinylazetidin-2-Ones: Synthesis, Antiproliferative and Tubulin Destabilizing Activity in MCF-7 and MDA-MB-231 Breast Cancer Cells. <i>Pharmaceuticals</i> , 2019 , 12,	5.2	5
21	Involvement of NF- B in mediating the anti-tumour effects of combretastatins in T cells. Investigational New Drugs, 2018 , 36, 523-535	4.3	3
20	Lead Optimization of Benzoxepin-Type Selective Estrogen Receptor (ER) Modulators and Downregulators with Subtype-Specific ERIand ERIActivity. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 514-534	8.3	20
19	Lactam analogues of combretastatin A-4 prevent metabolic inactivation by glucuronidation in chemoresistant HT-29 colon cancer cells. <i>European Journal of Medicinal Chemistry</i> , 2017 , 130, 261-285	6.8	25
18	Piperlongumine (piplartine) and analogues: Antiproliferative microtubule-destabilising agents. <i>European Journal of Medicinal Chemistry</i> , 2017 , 125, 453-463	6.8	26
17	Synthesis and Biochemical Evaluation of 3-Phenoxy-1,4-diarylazetidin-2-ones as Tubulin-Targeting Antitumor Agents. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 90-113	8.3	45
16	Assessment of cross-reactivity of new less sensitizing epoxy resin monomers in epoxy resin-allergic individuals. <i>Contact Dermatitis</i> , 2016 , 75, 144-50	2.7	11
15	Lactam estrogen receptor antagonists and a dual-targeting estrogen receptor/tubulin ligand. Journal of Medicinal Chemistry, 2014 , 57, 9370-82	8.3	37

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14	Epoxy resin monomers with reduced skin sensitizing potency. <i>Chemical Research in Toxicology</i> , 2014 , 27, 1002-10	4	8
13	Combretastatin (CA)-4 and its novel analogue CA-432 impair T-cell migration through the Rho/ROCK signalling pathway. <i>Biochemical Pharmacology</i> , 2014 , 92, 544-57	6	12
12	Synthesis and biochemical activities of antiproliferative amino acid and phosphate derivatives of microtubule-disrupting Elactam combretastatins. <i>European Journal of Medicinal Chemistry</i> , 2013 , 62, 705-21	6.8	18
11	Novel cis-restricted Elactam combretastatin A-4 analogues display anti-vascular and anti-metastatic properties in vitro. <i>Oncology Reports</i> , 2013 , 29, 585-94	3.5	18
10	Combretazet-3 a novel synthetic cis-stable combretastatin A-4-azetidinone hybrid with enhanced stability and therapeutic efficacy in colon cancer. <i>Oncology Reports</i> , 2013 , 29, 2451-8	3.5	20
9	Analogues of the epoxy resin monomer diglycidyl ether of bisphenol F: effects on contact allergenic potency and cytotoxicity. <i>Chemical Research in Toxicology</i> , 2012 , 25, 2469-78	4	20
8	The vascular targeting agent Combretastatin-A4 directly induces autophagy in adenocarcinoma-derived colon cancer cells. <i>Biochemical Pharmacology</i> , 2012 , 84, 612-24	6	36
7	Synthesis, biochemical and molecular modelling studies of antiproliferative azetidinones causing microtubule disruption and mitotic catastrophe. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 459	5 ⁶ 607	36
6	Lead identification of Elactam and related imine inhibitors of the molecular chaperone heat shock protein 90. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 6055-68	3.4	17
5	Synthesis, evaluation and structural studies of antiproliferative tubulin-targeting azetidin-2-ones. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 2306-25	3.4	53
4	The vascular targeting agent combretastatin-A4 and a novel cis-Restricted {beta}-Lactam Analogue, CA-432, induce apoptosis in human chronic myeloid leukemia cells and ex vivo patient samples including those displaying multidrug resistance. <i>Journal of Pharmacology and Experimental</i>	4.7	21
3	Lead identification of conformationally restricted benzoxepin type combretastatin analogs: synthesis, antiproliferative activity, and tubulin effects. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2010 , 25, 180-94	5.6	12
2	Synthesis and evaluation of azetidinone analogues of combretastatin A-4 as tubulin targeting agents. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 8569-84	8.3	91
1	dieredScreena- Layered Virtual Screening Tool for the Identification of Novel Estrogen Receptor Alpha Modulators. <i>Molecular Informatics</i> , 2010 , 29, 421-30	3.8	6