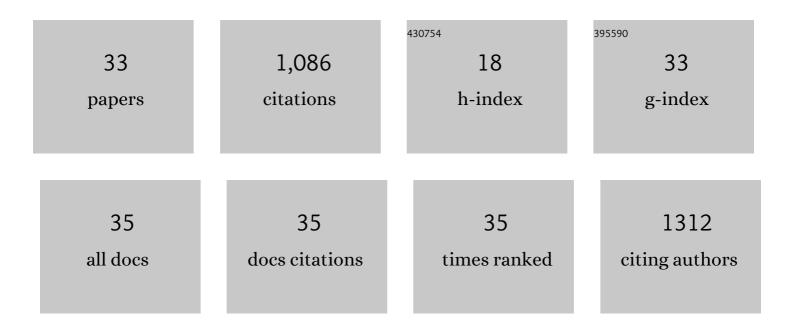
## Niamh M O'boyle

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Colchicine-Binding Site Inhibitors from Chemistry to Clinic: A Review. Pharmaceuticals, 2020, 13, 8.	1.7	187
2	Synthesis and Evaluation of Azetidinone Analogues of Combretastatin A-4 as Tubulin Targeting Agents. Journal of Medicinal Chemistry, 2010, 53, 8569-8584.	2.9	111
3	Skin lipids in health and disease: A review. Chemistry and Physics of Lipids, 2021, 236, 105055.	1.5	72
4	Synthesis, evaluation and structural studies of antiproliferative tubulin-targeting azetidin-2-ones. Bioorganic and Medicinal Chemistry, 2011, 19, 2306-2325.	1.4	62
5	Synthesis and Biochemical Evaluation of 3-Phenoxy-1,4-diarylazetidin-2-ones as Tubulin-Targeting Antitumor Agents. Journal of Medicinal Chemistry, 2016, 59, 90-113.	2.9	57
6	β-Lactam Estrogen Receptor Antagonists and a Dual-Targeting Estrogen Receptor/Tubulin Ligand. Journal of Medicinal Chemistry, 2014, 57, 9370-9382.	2.9	45
7	The vascular targeting agent Combretastatin-A4 directly induces autophagy in adenocarcinoma-derived colon cancer cells. Biochemical Pharmacology, 2012, 84, 612-624.	2.0	44
8	Synthesis, biochemical and molecular modelling studies of antiproliferative azetidinones causing microtubule disruption and mitotic catastrophe. European Journal of Medicinal Chemistry, 2011, 46, 4595-4607.	2.6	41
9	Piperlongumine (piplartine) and analogues: Antiproliferative microtubule-destabilising agents. European Journal of Medicinal Chemistry, 2017, 125, 453-463.	2.6	36
10	β-Lactam analogues of combretastatin A-4 prevent metabolic inactivation by glucuronidation in chemoresistant HT-29 colon cancer cells. European Journal of Medicinal Chemistry, 2017, 130, 261-285.	2.6	35
11	Lead Optimization of Benzoxepin-Type Selective Estrogen Receptor (ER) Modulators and Downregulators with Subtype-Specific ERα and ERβ Activity. Journal of Medicinal Chemistry, 2018, 61, 514-534.	2.9	35
12	Special Issue "Anticancer Drugs― Pharmaceuticals, 2019, 12, 134.	1.7	33
13	The Vascular Targeting Agent Combretastatin-A4 and a Novel <i>cis</i> -Restricted Î <sup>2</sup> -Lactam Analogue, CA-432, Induce Apoptosis in Human Chronic Myeloid Leukemia Cells and Ex Vivo Patient Samples Including Those Displaying Multidrug Resistance. Journal of Pharmacology and Experimental Therapeutics, 2010, 335, 302-313.	1.3	26
14	Analogues of the Epoxy Resin Monomer Diglycidyl Ether of Bisphenol F: Effects on Contact Allergenic Potency and Cytotoxicity. Chemical Research in Toxicology, 2012, 25, 2469-2478.	1.7	25
15	β-Lactams with antiproliferative and antiapoptotic activity in breast and chemoresistant colon cancer cells. European Journal of Medicinal Chemistry, 2020, 189, 112050.	2.6	25
16	Synthesis and biochemical activities of antiproliferative amino acid and phosphate derivatives of microtubule-disrupting β-lactam combretastatins. European Journal of Medicinal Chemistry, 2013, 62, 705-721.	2.6	23
17	Combretazet-3 a novel synthetic cis-stable combretastatin A-4- azetidinone hybrid with enhanced stability and therapeutic efficacy in colon cancer. Oncology Reports, 2013, 29, 2451-2458.	1.2	22
18	Novel cis-restricted β-lactam combretastatin A-4 analogues display anti-vascular and anti-metastatic properties in vitro. Oncology Reports, 2013, 29, 585-594.	1.2	19

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19	Lead identification of β-lactam and related imine inhibitors of the molecular chaperone heat shock protein 90. Bioorganic and Medicinal Chemistry, 2011, 19, 6055-6068.	1.4	18
20	Synthesis and evaluation of antiproliferative microtubule-destabilising combretastatin A-4 piperazine conjugates. Organic and Biomolecular Chemistry, 2019, 17, 6184-6200.	1.5	15
21	Lead identification of conformationally restricted benzoxepin type combretastatin analogs: synthesis, antiproliferative activity, and tubulin effects. Journal of Enzyme Inhibition and Medicinal Chemistry, 2010, 25, 180-194.	2.5	13
22	Combretastatin (CA)-4 and its novel analogue CA-432 impair T-cell migration through the Rho/ROCK signalling pathway. Biochemical Pharmacology, 2014, 92, 544-557.	2.0	13
23	Assessment of crossâ€reactivity of new less sensitizing epoxy resin monomers in epoxy resinâ€allergic individuals. Contact Dermatitis, 2016, 75, 144-150.	0.8	13
24	Epoxy Resin Monomers with Reduced Skin Sensitizing Potency. Chemical Research in Toxicology, 2014, 27, 1002-1010.	1.7	12
25	Potent Quinoline-Containing Combretastatin A-4 Analogues: Design, Synthesis, Antiproliferative, and Anti-Tubulin Activity. Pharmaceuticals, 2020, 13, 393.	1.7	12
26	3-Vinylazetidin-2-Ones: Synthesis, Antiproliferative and Tubulin Destabilizing Activity in MCF-7 and MDA-MB-231 Breast Cancer Cells. Pharmaceuticals, 2019, 12, 56.	1.7	10
27	â€~ <i>tieredScreen</i> ' – Layered Virtual Screening Tool for the Identification of Novel Estrogen Receptor Alpha Modulators. Molecular Informatics, 2010, 29, 421-430.	1.4	7
28	Synthesis and Antiproliferative Evaluation of 3-Chloroazetidin-2-ones with Antimitotic Activity: Heterocyclic Bridged Analogues of Combretastatin A-4. Pharmaceuticals, 2021, 14, 1119.	1.7	7
29	Nature-derived epoxy resins: Synthesis, allergenicity, and thermosetting properties of pinoresinol diglycidyl ether. Toxicology and Industrial Health, 2022, 38, 259-269.	0.6	7
30	Synthesis and Biological Evaluation of 1-(Diarylmethyl)-1H-1,2,4-triazoles and 1-(Diarylmethyl)-1H-imidazoles as a Novel Class of Anti-Mitotic Agent for Activity in Breast Cancer. Pharmaceuticals, 2021, 14, 169.	1.7	5
31	Involvement of NF-κB in mediating the anti-tumour effects of combretastatins in T cells. Investigational New Drugs, 2018, 36, 523-535.	1.2	4
32	Special Issue "Anticancer Drugs 2021― Pharmaceuticals, 2022, 15, 479.	1.7	2
33	Azetidin-2-ones: structures of antimitotic compounds based on the 1-(3,4,5-trimethoxyphenyl)azetidin-2-one core. Acta Crystallographica Section E: Crystallographic Communications, 2020, 76, 1187-1194	0.2	1