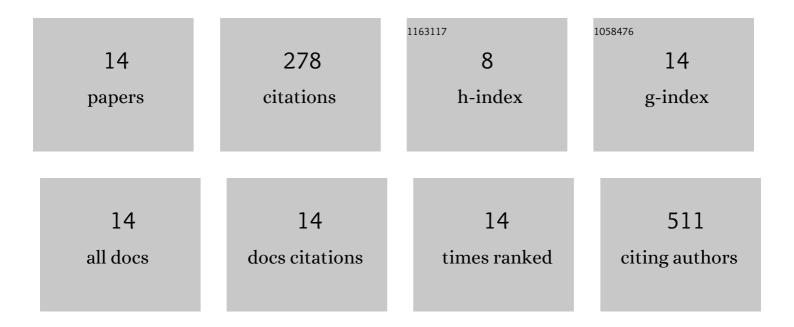
Suzannah J Harnor

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Cyclin-Dependent Kinase (CDK) Inhibitors: Structure–Activity Relationships and Insights into the CDK-2 Selectivity of 6-Substituted 2-Arylaminopurines. Journal of Medicinal Chemistry, 2017, 60, 1746-1767.	6.4	77
2	Targeting DNAâ€Dependent Protein Kinase for Cancer Therapy. ChemMedChem, 2017, 12, 895-900.	3.2	63
3	Structure-Based Design of Potent and Orally Active Isoindolinone Inhibitors of MDM2-p53 Protein–Protein Interaction. Journal of Medicinal Chemistry, 2021, 64, 4071-4088.	6.4	30
4	Human Toxicity Caused by Indole and Indazole Carboxylate Synthetic Cannabinoid Receptor Agonists: From Horizon Scanning to Notification. Clinical Chemistry, 2018, 64, 346-354.	3.2	23
5	Structure-activity analysis of CJ-15,801 analogues that interact with Plasmodium falciparum pantothenate kinase and inhibit parasite proliferation. European Journal of Medicinal Chemistry, 2018, 143, 1139-1147.	5.5	16
6	Identification of a novel orally bioavailable ERK5 inhibitor with selectivity over p38 \hat{I}_{\pm} and BRD4. European Journal of Medicinal Chemistry, 2019, 178, 530-543.	5.5	15
7	The Enhanced In Vivo Activity of the Combination of a MEK and a PI3K Inhibitor Correlates with [18F]-FLT PET in Human Colorectal Cancer Xenograft Tumour-Bearing Mice. PLoS ONE, 2013, 8, e81763.	2.5	13
8	An Alkynylpyrimidine-Based Covalent Inhibitor That Targets a Unique Cysteine in NF-κB-Inducing Kinase. Journal of Medicinal Chemistry, 2021, 64, 10001-10018.	6.4	9
9	Chronic glucokinase activator treatment activates liver Carbohydrate response element binding protein and improves hepatocyte <scp>ATP</scp> homeostasis during substrate challenge. Diabetes, Obesity and Metabolism, 2020, 22, 1985-1994.	4.4	8
10	2-Arylamino-6-ethynylpurines are cysteine-targeting irreversible inhibitors of Nek2 kinase. RSC Medicinal Chemistry, 2020, 11, 707-731.	3.9	8
11	Photochemical synthesis of benz[h]isoquinolines. Tetrahedron, 2008, 64, 5072-5078.	1.9	7
12	Concise syntheses of bridged morpholines. RSC Advances, 2016, 6, 53955-53957.	3.6	5
13	Parallel Optimization of Potency and Pharmacokinetics Leading to the Discovery of a Pyrrole Carboxamide ERK5 Kinase Domain Inhibitor. Journal of Medicinal Chemistry, 2022, 65, 6513-6540.	6.4	3
14	Synthesis of 3′-deoxy-3′-fluorothymidine (FLT) 5′-O-glucuronide: a reference standard for imaging studies with [¹⁸ F]FLT. MedChemComm, 2014, 5, 984-988.	3.4	1