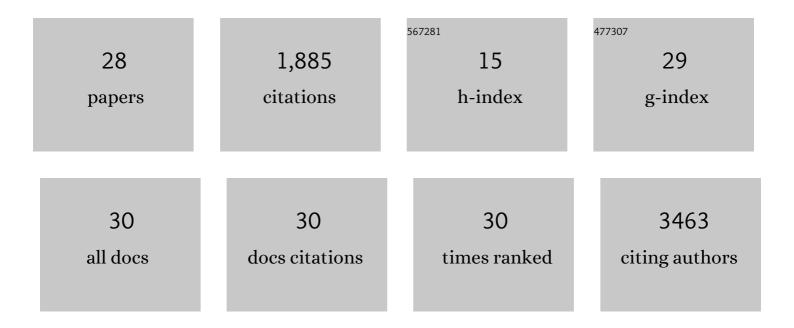
## Maureen Higgins

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/166051/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	The isoquinoline PRL-295 increases the thermostability of Keap1 and disrupts its interaction with Nrf2. IScience, 2022, 25, 103703.	4.1	11
2	Pirin, an Nrf2-Regulated Protein, Is Overexpressed in Human Colorectal Tumors. Antioxidants, 2022, 11, 262.	5.1	8
3	Nrf2 activation reprograms macrophage intermediary metabolism and suppresses the type I interferon response. IScience, 2022, 25, 103827.	4.1	51
4	The synthetic triterpenoids CDDO-TFEA and CDDO-Me, but not CDDO, promote nuclear exclusion of BACH1 impairing its activity. Redox Biology, 2022, 51, 102291.	9.0	12
5	Phenyl Bis-Sulfonamide Keap1-Nrf2 Protein–Protein Interaction Inhibitors with an Alternative Binding Mode. Journal of Medicinal Chemistry, 2022, 65, 7380-7398.	6.4	14
6	Nrf2 is activated by disruption of mitochondrial thiol homeostasis but not by enhanced mitochondrial superoxide production. Journal of Biological Chemistry, 2021, 296, 100169.	3.4	25
7	Clinically relevant aberrant Filip1l DNA methylation detected in a murine model of cutaneous squamous cell carcinoma. EBioMedicine, 2021, 67, 103383.	6.1	4
8	Novel iodinated quinazolinones bearing sulfonamide as new scaffold targeting radiation induced oxidative stress. Bioorganic and Medicinal Chemistry Letters, 2021, 42, 128002.	2.2	6
9	Nrf2 activation does not affect adenoma development in a mouse model of colorectal cancer. Communications Biology, 2021, 4, 1081.	4.4	1
10	Isomeric O-methyl cannabidiolquinones with dual BACH1/NRF2 activity. Redox Biology, 2020, 37, 101689.	9.0	23
11	Downregulation of Keap1 Confers Features of a Fasted Metabolic State. IScience, 2020, 23, 101638.	4.1	21
12	Radiomodulatory effect of a non-electrophilic NQO1 inducer identified in a screen of new 6, 8-diiodoquinazolin-4(3H)-ones carrying a sulfonamide moiety. European Journal of Medicinal Chemistry, 2020, 200, 112467.	5.5	10
13	Itaconate is an anti-inflammatory metabolite that activates Nrf2 via alkylation of KEAP1. Nature, 2018, 556, 113-117.	27.8	1,115
14	A DHODH inhibitor increases p53 synthesis and enhances tumor cell killing by p53 degradation blockage. Nature Communications, 2018, 9, 1107.	12.8	63
15	KEAP1 inhibition is neuroprotective and suppresses the development of epilepsy. Brain, 2018, 141, 1390-1403.	7.6	99
16	KEAP1-modifying small molecule reveals muted NRF2 signaling responses in neural stem cells from Huntington's disease patients. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E4676-E4685.	7.1	119
17	Whole-Exome Sequencing Validates a Preclinical Mouse Model for the Prevention and Treatment of Cutaneous Squamous Cell Carcinoma. Cancer Prevention Research, 2017, 10, 67-75.	1.5	17
18	NAD(P)H:quinone oxidoreductase 1 inducer activity of some novel anilinoquinazoline derivatives. Drug Design, Development and Therapy, 2016, Volume 10, 2515-2524.	4.3	5

MAUREEN HIGGINS

#	Article	IF	CITATIONS
19	Loss of Nrf2 abrogates the protective effect of Keap1 downregulation in a preclinical model of cutaneous squamous cell carcinoma. Scientific Reports, 2016, 6, 25804.	3.3	28
20	Synthesis, molecular modeling and NAD(P)H:quinone oxidoreductase 1 inducer activity of novel 2-phenylquinazolin-4-amine derivatives. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1612-1618.	5.2	4
21	Synthesis and biological evaluation of novel 2-phenylquinazoline-4-amine derivatives: identification of 6-phenyl-8H-benzo[g]quinazolino[4,3-b]quinazolin-8-one as a highly potent inducer of NAD(P)H quinone oxidoreductase 1. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 34-39.	5.2	4
22	Potency of extracts from selected Egyptian plants as inducers of the Nrf2-dependent chemopreventive enzyme NQO1. Journal of Natural Medicines, 2016, 70, 683-688.	2.3	9
23	NAD(P)H: quinone oxidoreductase 1 inducer activity of novel 4-aminoquinazoline derivatives. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1369-1374.	5.2	3
24	Redox effects and cytotoxic profiles of MJ25 and auranofin towards malignant melanoma cells. Oncotarget, 2015, 6, 16488-16506.	1.8	30
25	New Monocyclic, Bicyclic, and Tricyclic Ethynylcyanodienones as Activators of the Keap1/Nrf2/ARE Pathway and Inhibitors of Inducible Nitric Oxide Synthase. Journal of Medicinal Chemistry, 2015, 58, 4738-4748.	6.4	26
26	Nrf2 Activation Protects against Solar-Simulated Ultraviolet Radiation in Mice and Humans. Cancer Prevention Research, 2015, 8, 475-486.	1.5	94
27	N1-Benzyl substituted cambinol analogues as isozyme selective inhibitors of the sirtuin family of protein deacetylases. MedChemComm, 2011, 2, 611.	3.4	16
28	Evaluation of an Actinomycin D/VX-680 aurora kinase inhibitor combination in p53-based cyclotherapy. Oncotarget, 2010, 1, 639-650.	1.8	65