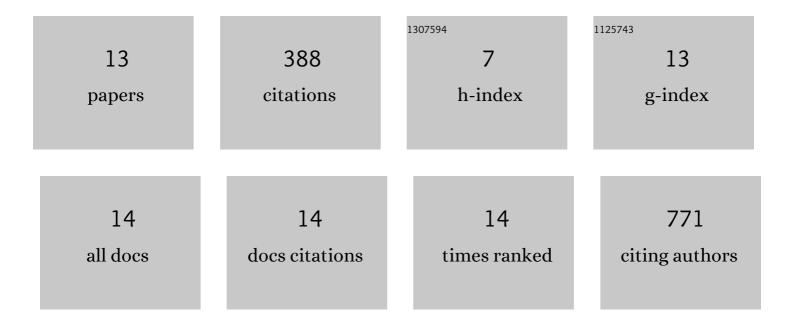
Mary E Law

List of Publications by Year in descending order

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MADYFLAW

#	Article	IF	CITATIONS
1	Inhibitors of ERp44, PDIA1, and AGR2 induce disulfide-mediated oligomerization of Death Receptors 4 and 5 and cancer cell death. Cancer Letters, 2022, 534, 215604.	7.2	4
2	Anticancer Agents Derived from Cyclic Thiosulfonates: Structureâ€Reactivity and Structureâ€Activity Relationships. ChemMedChem, 2022, 17, .	3.2	1
3	Repurposing Tranexamic Acid as an Anticancer Agent. Frontiers in Pharmacology, 2021, 12, 792600.	3.5	4
4	Disulfide bond-disrupting agents activate the tumor necrosis family-related apoptosis-inducing ligand/death receptor 5 pathway. Cell Death Discovery, 2019, 5, 153.	4.7	9
5	A novel proteotoxic combination therapy for EGFR+ and HER2+ cancers. Oncogene, 2019, 38, 4264-4282.	5.9	8
6	Loss of sirtuin 1 and mitofusin 2 contributes to enhanced ischemia/reperfusion injury in aged livers. Aging Cell, 2018, 17, e12761.	6.7	60
7	The unfolded protein response as a target for anticancer therapeutics. Critical Reviews in Oncology/Hematology, 2018, 127, 66-79.	4.4	102
8	Disulfide bond disrupting agents activate the unfolded protein response in EGFR- and HER2-positive breast tumor cells. Oncotarget, 2017, 8, 28971-28989.	1.8	11
9	CUB domain-containing protein 1 and the epidermal growth factor receptor cooperate to induce cell detachment. Breast Cancer Research, 2016, 18, 80.	5.0	25
10	Cyclin-Dependent Kinase Inhibitors as Anticancer Therapeutics. Molecular Pharmacology, 2015, 88, 846-852.	2.3	79
11	Signaling Mechanisms that Suppress the Cytostatic Actions of Rapamycin. PLoS ONE, 2014, 9, e99927.	2.5	3
12	Identification of a small molecule inhibitor of serine 276 phosphorylation of the p65 subunit of NF-κB using in silico molecular docking. Cancer Letters, 2010, 291, 217-224.	7.2	18
13	Rapamycin Disrupts Cyclin/Cyclin-Dependent Kinase/p21/Proliferating Cell Nuclear Antigen Complexes and Cyclin D1 Reverses Rapamycin Action by Stabilizing These Complexes. Cancer Research, 2006, 66, 1070-1080.	0.9	63