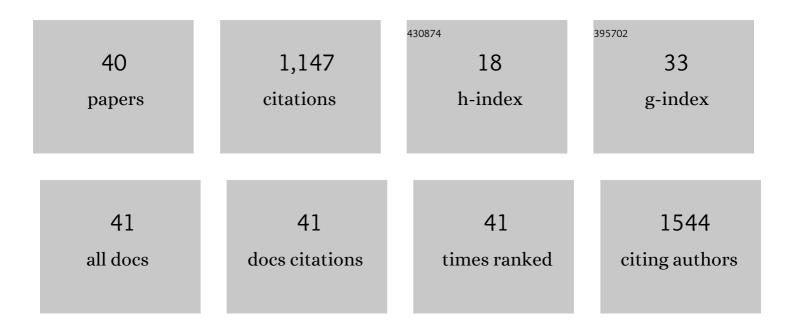
## Carol S Lim

List of Publications by Year in descending order

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CAROL S LIM

#	Article	IF	CITATIONS
1	Differential Localization and Activity of the A- and B-Forms of the Human Progesterone Receptor Using Green Fluorescent Protein Chimeras. Molecular Endocrinology, 1999, 13, 366-375.	3.7	135
2	Trafficking of nuclear receptors in living cells. Journal of Steroid Biochemistry and Molecular Biology, 2000, 74, 249-254.	2.5	118
3	Delivery of drugs and macromolecules to the mitochondria for cancer therapy. Journal of Controlled Release, 2016, 240, 38-51.	9.9	101
4	Resistant mutations in CML and Ph+ALL – role of ponatinib. Biologics: Targets and Therapy, 2014, 8, 243.	3.2	65
5	Controlling Protein Compartmentalization to Overcome Disease. Pharmaceutical Research, 2006, 24, 17-27.	3.5	55
6	Intracellular localization and trafficking of steroid receptors. Cell Biochemistry and Biophysics, 1999, 31, 119-127.	1.8	46
7	Controlling subcellular delivery to optimize therapeutic effect. Therapeutic Delivery, 2010, 1, 169-193.	2.2	45
8	Targeting malignant mitochondria with therapeutic peptides. Therapeutic Delivery, 2012, 3, 961-979.	2.2	39
9	Differential Localization and Activity of the A- and B-Forms of the Human Progesterone Receptor Using Green Fluorescent Protein Chimeras. Molecular Endocrinology, 1999, 13, 366-375.	3.7	38
10	Application of Thiol–yne/Thiol–ene Reactions for Peptide and Protein Macrocyclizations. Chemistry - A European Journal, 2017, 23, 7087-7092.	3.3	36
11	Direct Induction of Apoptosis Using an Optimal Mitochondrially Targeted p53. Molecular Pharmaceutics, 2012, 9, 1449-1458.	4.6	33
12	Disruption of Bcr-Abl Coiled Coil Oligomerization by Design. Journal of Biological Chemistry, 2011, 286, 27751-27760.	3.4	28
13	Model system to study classical nuclear export signals. AAPS PharmSci, 2002, 4, 61-68.	1.3	24
14	Optimizing the protein switch: Altering nuclear import and export signals, and ligand binding domain. Journal of Controlled Release, 2007, 120, 220-232.	9.9	24
15	Improved Coiled-Coil Design Enhances Interaction with Bcr-Abl and Induces Apoptosis. Molecular Pharmaceutics, 2012, 9, 187-195.	4.6	23
16	Controlling subcellular localization to alter function: Sending oncogenic Bcr–Abl to the nucleus causes apoptosis. Journal of Controlled Release, 2009, 140, 245-249.	9.9	21
17	Effect of Initial Subcellular Localization of Progesterone Receptor on Import Kinetics and Transcriptional Activity. Molecular Pharmaceutics, 2005, 2, 509-518.	4.6	20
18	The Androgen Receptor and Its Use in Biological Assays: Looking Toward Effect-Based Testing and Its Applications. Journal of Analytical Toxicology, 2011, 35, 594-607.	2.8	19

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19	The DNA Binding Domain of p53 Is Sufficient To Trigger a Potent Apoptotic Response at the Mitochondria. Molecular Pharmaceutics, 2013, 10, 3592-3602.	4.6	18
20	A Chimeric p53 Evades Mutant p53 Transdominant Inhibition in Cancer Cells. Molecular Pharmaceutics, 2013, 10, 3922-3933.	4.6	18
21	Advances in delivery vectors for gene therapy in liver cancer. Therapeutic Delivery, 2020, 11, 833-850.	2.2	18
22	Roadmap to affinity-tuned antibodies for enhanced chimeric antigen receptor T cell function and selectivity. Trends in Biotechnology, 2022, 40, 875-890.	9.3	17
23	Changing the Subcellular Location of the Oncoprotein Bcr-Abl Using Rationally Designed Capture Motifs. Pharmaceutical Research, 2012, 29, 1098-1109.	3.5	16
24	Delivery of a Monomeric p53 Subdomain with Mitochondrial Targeting Signals from Pro-Apoptotic Bak or Bax. Pharmaceutical Research, 2014, 31, 2503-2515.	3.5	15
25	p53-Bad: A Novel Tumor Suppressor/Proapoptotic Factor Hybrid Directed to the Mitochondria for Ovarian Cancer Gene Therapy. Molecular Pharmaceutics, 2019, 16, 3386-3398.	4.6	15
26	Geldanamycin, an inhibitor of Hsp90, Blocks cytoplasmic retention of progesterone receptors and glucocorticoid receptors via their respective ligand binding domains. AAPS Journal, 2006, 8, E718-E728.	4.4	14
27	Controlled Access of p53 to the Nucleus Regulates Its Proteasomal Degradation by MDM2. Molecular Pharmaceutics, 2013, 10, 1340-1349.	4.6	14
28	The nuclear translocation assay for intracellular protein-protein interactions and its application to the Bcr coiled-coil domain. BioTechniques, 2010, 49, 519-524.	1.8	12
29	Multidomain Targeting of Bcr-Abl by Disruption of Oligomerization and Tyrosine Kinase Inhibition: Toward Eradication of CML. Molecular Pharmaceutics, 2013, 10, 3475-3483.	4.6	12
30	Enhanced and Selective Killing of Chronic Myelogenous Leukemia Cells with an Engineered BCR-ABL Binding Protein and Imatinib. Molecular Pharmaceutics, 2012, 9, 3318-3329.	4.6	11
31	Computational Modeling of Stapled Peptides toward a Treatment Strategy for CML and Broader Implications in the Design of Lengthy Peptide Therapeutics. Journal of Physical Chemistry B, 2018, 122, 3864-3875.	2.6	11
32	Selective Targeting of c-Abl via a Cryptic Mitochondrial Targeting Signal Activated by Cellular Redox Status in Leukemic and Breast Cancer Cells. Pharmaceutical Research, 2012, 29, 2317-2328.	3.5	10
33	A Single Mutant, A276S of p53, Turns the Switch to Apoptosis. Molecular Pharmaceutics, 2013, 10, 1350-1359.	4.6	10
34	Signal Sequences for Targeting of Gene Therapy Products to Subcellular Compartments: The Role Of CRM1 in Nucleocytoplasmic Shuttling of the Protein Switch. Pharmaceutical Research, 2007, 24, 2146-2155.	3.5	9
35	Narrowing the field: cancer-specific promoters for mitochondrially-targeted p53-BH3 fusion gene therapy in ovarian cancer. Journal of Ovarian Research, 2019, 12, 38.	3.0	9
36	Inhibition of Bcr-Abl in Human Leukemic Cells with a Coiled-Coil Protein Delivered by a Leukemia-Specific Cell-Penetrating Peptide. Molecular Pharmaceutics, 2015, 12, 1412-1421.	4.6	8

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37	Correlation among agonist dose, rate of import, and transcriptional activity of liganded progesterone receptor B isoform in living cells. Pharmaceutical Research, 2003, 20, 1574-1580.	3.5	7
38	Re-Engineered p53 Chimera with Enhanced Homo-Oligomerization That Maintains Tumor Suppressor Activity. Molecular Pharmaceutics, 2014, 11, 2442-2452.	4.6	7
39	Mitochondrially targeted p53 or DBD subdomain is superior to wild type p53 in ovarian cancer cells even with strong dominant negative mutant p53. Journal of Ovarian Research, 2019, 12, 45.	3.0	7
40	Utilizing the Estrogen Receptor Ligand-Binding Domain for Controlled Protein Translocation to the Insoluble Fraction. Pharmaceutical Research, 2012, 29, 3455-3463.	3.5	3