

Geoffrey Greene

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/906759/publications.pdf>

Version: 2024-02-01

55
papers

11,249
citations

126708

33
h-index

168136

53
g-index

61
all docs

61
docs citations

61
times ranked

9581
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular basis of agonism and antagonism in the oestrogen receptor. <i>Nature</i> , 1997, 389, 753-758.	13.7	3,139
2	The Structural Basis of Estrogen Receptor/Coactivator Recognition and the Antagonism of This Interaction by Tamoxifen. <i>Cell</i> , 1998, 95, 927-937.	13.5	2,441
3	Sequence and expression of human estrogen receptor complementary DNA. <i>Science</i> , 1986, 231, 1150-1154.	6.0	1,223
4	ESR1 ligand-binding domain mutations in hormone-resistant breast cancer. <i>Nature Genetics</i> , 2013, 45, 1439-1445.	9.4	960
5	Overcoming mutation-based resistance to antiandrogens with rational drug design. <i>ELife</i> , 2013, 2, e00499.	2.8	334
6	Activating ESR1 Mutations Differentially Affect the Efficacy of ER Antagonists. <i>Cancer Discovery</i> , 2017, 7, 277-287.	7.7	286
7	Structural Basis for an Unexpected Mode of SERM-Mediated ER Antagonism. <i>Molecular Cell</i> , 2005, 18, 413-424.	4.5	225
8	Estrogen receptor alpha somatic mutations Y537S and D538G confer breast cancer endocrine resistance by stabilizing the activating function-2 binding conformation. <i>ELife</i> , 2016, 5, .	2.8	212
9	MicroRNA-30c inhibits human breast tumour chemotherapy resistance by regulating TWF1 and IL-11. <i>Nature Communications</i> , 2013, 4, 1393.	5.8	209
10	Structural characterization of a subtype-selective ligand reveals a novel mode of estrogen receptor antagonism. <i>Nature Structural Biology</i> , 2002, 9, 359-64.	9.7	188
11	NF κ B selectivity of estrogen receptor ligands revealed by comparative crystallographic analyses. <i>Nature Chemical Biology</i> , 2008, 4, 241-247.	3.9	149
12	Structural underpinnings of oestrogen receptor mutations in endocrine therapy resistance. <i>Nature Reviews Cancer</i> , 2018, 18, 377-388.	12.8	148
13	Neutrophil elastase selectively kills cancer cells and attenuates tumorigenesis. <i>Cell</i> , 2021, 184, 3163-3177.e21.	13.5	119
14	14q32-encoded microRNAs mediate an oligometastatic phenotype. <i>Oncotarget</i> , 2015, 6, 3540-3552.	0.8	103
15	Allosteric Control of Ligand Selectivity between Estrogen Receptors ER α and ER β . <i>Molecular Cell</i> , 2004, 13, 317-327.	4.5	100
16	Genomic agonism and phenotypic antagonism between estrogen and progesterone receptors in breast cancer. <i>Science Advances</i> , 2016, 2, e1501924.	4.7	100
17	MicroRNA-30c targets cytoskeleton genes involved in breast cancer cell invasion. <i>Breast Cancer Research and Treatment</i> , 2013, 137, 373-382.	1.1	90
18	Stapled Peptides with Methylated Hydrocarbon Chains for the Estrogen Receptor/Coactivator Interaction. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 4252-4255.	7.2	73

#	ARTICLE	IF	CITATIONS
19	Analysis of the Structural Core of the Human Estrogen Receptor Ligand Binding Domain by Selective Proteolysis/Mass Spectrometric Analysis. <i>Biochemistry</i> , 1995, 34, 12605-12615.	1.2	72
20	The SERM/SERD bazedoxifene disrupts ESR1 helix 12 to overcome acquired hormone resistance in breast cancer cells. <i>ELife</i> , 2018, 7, .	2.8	72
21	Identification of cysteine 530 as the covalent attachment site of an affinity-labeling estrogen (ketononestrol aziridine) and antiestrogen (tamoxifen aziridine) in the human estrogen receptor. <i>Journal of Biological Chemistry</i> , 1989, 264, 17476-85.	1.6	65
22	Estrogen and Progesterin Receptors and Aromatase Activity in Rhesus Monkey Prostate*. <i>Endocrinology</i> , 1988, 123, 2312-2322.	1.4	62
23	Characterization of the Subunit Nature of Nuclear Estrogen Receptors by Chemical Cross-Linking and Dense Amino Acid Labeling*. <i>Endocrinology</i> , 1985, 117, 515-522.	1.4	57
24	Green tea catechins inhibit angiogenesis through suppression of STAT3 activation. <i>Breast Cancer Research and Treatment</i> , 2009, 117, 505-515.	1.1	56
25	Estrogen Inhibits Vascular Smooth Muscle Cell-Dependent Adventitial Fibroblast Migration In Vitro. <i>Circulation</i> , 1999, 100, 1639-1645.	1.6	55
26	Identification of Ligands with Bicyclic Scaffolds Provides Insights into Mechanisms of Estrogen Receptor Subtype Selectivity*. <i>Journal of Biological Chemistry</i> , 2006, 281, 17909-17919.	1.6	51
27	Discovery of a Glucocorticoid Receptor (GR) Activity Signature Using Selective GR Antagonism in ER-Negative Breast Cancer. <i>Clinical Cancer Research</i> , 2018, 24, 3433-3446.	3.2	49
28	Progesterone receptor isoforms, agonists and antagonists differentially reprogram estrogen signaling. <i>Oncotarget</i> , 2018, 9, 4282-4300.	0.8	49
29	Inhibition of mammary tumorigenesis in the C3(1)/SV40 mouse model by green tea. <i>Breast Cancer Research and Treatment</i> , 2008, 107, 359-369.	1.1	42
30	Specific stereochemistry of OP-1074 disrupts estrogen receptor alpha helix 12 and confers pure antiestrogenic activity. <i>Nature Communications</i> , 2018, 9, 2368.	5.8	42
31	Next-Generation ER \pm Inhibitors for Endocrine-Resistant ER+ Breast Cancer. <i>Endocrinology</i> , 2019, 160, 759-769.	1.4	42
32	Lasofoxifene as a potential treatment for therapy-resistant ER-positive metastatic breast cancer. <i>Breast Cancer Research</i> , 2021, 23, 54.	2.2	38
33	Recruitment of Histone Deacetylase 4 to the N-Terminal Region of Estrogen Receptor $\hat{\pm}$. <i>Molecular Endocrinology</i> , 2005, 19, 2930-2942.	3.7	37
34	Molecular characterization of a B-ring unsaturated estrogen: Implications for conjugated equine estrogen components of Premarin. <i>Steroids</i> , 2008, 73, 59-68.	0.8	32
35	Versatile Peptide Macrocyclization with Diels-Alder Cycloadditions. <i>Journal of the American Chemical Society</i> , 2019, 141, 16374-16381.	6.6	32
36	The NF- $\hat{\pm}$ B Pathway Promotes Tamoxifen Tolerance and Disease Recurrence in Estrogen Receptor-Positive Breast Cancers. <i>Molecular Cancer Research</i> , 2020, 18, 1018-1027.	1.5	31

#	ARTICLE	IF	CITATIONS
37	Interferon-Stimulated Genes Are Transcriptionally Repressed by PR in Breast Cancer. <i>Molecular Cancer Research</i> , 2017, 15, 1331-1340.	1.5	29
38	RAC3 is a pro-migratory co-activator of ER α . <i>Oncogene</i> , 2011, 30, 1984-1994.	2.6	28
39	Molecular characterization by mass spectrometry of the human estrogen receptor ligand-binding domain expressed in <i>Escherichia coli</i> . <i>Molecular Endocrinology</i> , 1995, 9, 647-658.	3.7	26
40	A α -cross-stitched peptide with improved helicity and proteolytic stability. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 3702-3706.	1.5	26
41	Endoxifen, 4-Hydroxytamoxifen and an Estrogenic Derivative Modulate Estrogen Receptor Complex Mediated Apoptosis in Breast Cancer. <i>Molecular Pharmacology</i> , 2018, 94, 812-822.	1.0	24
42	Removal of lactate dehydrogenase-elevating virus from human-in-mouse breast tumor xenografts by cell-sorting. <i>Journal of Virological Methods</i> , 2011, 173, 266-270.	1.0	22
43	A small-molecule activator of the unfolded protein response eradicates human breast tumors in mice. <i>Science Translational Medicine</i> , 2021, 13, .	5.8	20
44	The Structure-Function Relationship of Angular Estrogens and Estrogen Receptor Alpha to Initiate Estrogen-Induced Apoptosis in Breast Cancer Cells. <i>Molecular Pharmacology</i> , 2020, 98, 24-37.	1.0	19
45	Rapid Induction of the Unfolded Protein Response and Apoptosis by Estrogen Mimic TTC-352 for the Treatment of Endocrine-Resistant Breast Cancer. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 11-25.	1.9	11
46	Mapping ER α Genomic Binding Sites Reveals Unique Genomic Features and Identifies EBF1 as an ER α Interactor. <i>PLoS ONE</i> , 2013, 8, e71355.	1.1	11
47	Stereospecific lasofoxifene derivatives reveal the interplay between estrogen receptor alpha stability and antagonistic activity in ESR1 mutant breast cancer cells. <i>ELife</i> , 2022, 11, .	2.8	11
48	Antagonists for Constitutively Active Mutant Estrogen Receptors: Insights into the Roles of Antiestrogen-Core and Side-Chain. <i>ACS Chemical Biology</i> , 2018, 13, 3374-3384.	1.6	8
49	Facilitating Drug Discovery in Breast Cancer by Virtually Screening Patients Using In Vitro Drug Response Modeling. <i>Cancers</i> , 2021, 13, 885.	1.7	6
50	Defining the Energetic Basis for a Conformational Switch Mediating Ligand-Independent Activation of Mutant Estrogen Receptors in Breast Cancer. <i>Molecular Cancer Research</i> , 2021, 19, 1559-1570.	1.5	6
51	Selective pressure of endocrine therapy activates the integrated stress response through NF κ B signaling in a subpopulation of ER positive breast cancer cells. <i>Breast Cancer Research</i> , 2022, 24, 19.	2.2	6
52	Endocrine Therapy-Resistant Breast Cancer Cells Are More Sensitive to Ceramide Kinase Inhibition and Elevated Ceramide Levels Than Therapy-Sensitive Breast Cancer Cells. <i>Cancers</i> , 2022, 14, 2380.	1.7	4
53	A Structural Explanation for ER α /ER β SERM Discrimination. , 2004, , 33-45.		2
54	Labeling of a Mutant Estrogen Receptor with an Affimer in a Breast Cancer Cell Line. <i>Biophysical Journal</i> , 2022, , .	0.2	1

#	ARTICLE	IF	CITATIONS
55	Reinvestigating the acyl cyclization to the precursor of diptoindonesin G. Tetrahedron Letters, 2021, 69, 152980.	0.7	0