

Stephen Safe

List of Publications by Year in descending order

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183
papers

12,633
citations

18482

62
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27406

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224
all docs

224
docs citations

224
times ranked

12049
citing authors

#	ARTICLE	IF	CITATIONS
1	Polychlorinated Biphenyls (PCBs), Dibenzo-p-Dioxins (PCDDs), Dibenzofurans (PCDFs), and Related Compounds: Environmental and Mechanistic Considerations Which Support the Development of Toxic Equivalency Factors (TEFs). <i>Critical Reviews in Toxicology</i> , 1990, 21, 51-88.	3.9	1,702
2	Polychlorinated Biphenyls (PCBs) and Polybrominated Biphenyls (PBBs): Biochemistry, Toxicology, and Mechanism of Action. <i>CRC Critical Reviews in Toxicology</i> , 1984, 13, 319-395.	4.9	714
3	Inhibitory Aryl Hydrocarbon Receptor~Estrogen Receptor ± Cross-Talk and Mechanisms of Action. <i>Chemical Research in Toxicology</i> , 2003, 16, 807-816.	3.3	305
4	Non-classical genomic estrogen receptor (ER)/specificity protein and ER/activating protein-1 signaling pathways. <i>Journal of Molecular Endocrinology</i> , 2008, 41, 263-275.	2.5	278
5	Betulinic Acid Inhibits Prostate Cancer Growth through Inhibition of Specificity Protein Transcription Factors. <i>Cancer Research</i> , 2007, 67, 2816-2823.	0.9	275
6	Microbiome-Derived Tryptophan Metabolites and Their Aryl Hydrocarbon Receptor-Dependent Agonist and Antagonist Activities. <i>Molecular Pharmacology</i> , 2014, 85, 777-788.	2.3	254
7	Cancer chemotherapy with indole-3-carbinol, bis(3- ² -indolyl)methane and synthetic analogs. <i>Cancer Letters</i> , 2008, 269, 326-338.	7.2	237
8	Role of the Aryl Hydrocarbon Receptor in Carcinogenesis and Potential as a Drug Target. <i>Toxicological Sciences</i> , 2013, 135, 1-16.	3.1	230
9	Estrogen-Induced c-fos Protooncogene Expression in MCF-7 Human Breast Cancer Cells: Role of Estrogen Receptor Sp1 Complex Formation*. <i>Endocrinology</i> , 1998, 139, 1981-1990.	2.8	171
10	Inhibition of NF ^κ B and Pancreatic Cancer Cell and Tumor Growth by Curcumin Is Dependent on Specificity Protein Down-regulation. <i>Journal of Biological Chemistry</i> , 2010, 285, 25332-25344.	3.4	165
11	Nur77 Agonists Induce Proapoptotic Genes and Responses in Colon Cancer Cells through Nuclear Receptor~Dependent and Nuclear Receptor~Independent Pathways. <i>Cancer Research</i> , 2007, 67, 674-683.	0.9	160
12	Nuclear receptor 4A (NR4A) family ~orphans no more. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2016, 157, 48-60.	2.5	149
13	Mechanisms of inhibitory aryl hydrocarbon receptor-estrogen receptor crosstalk in human breast cancer cells. <i>Journal of Mammary Gland Biology and Neoplasia</i> , 2000, 5, 295-306.	2.7	147
14	Activation of Nur77 by Selected 1,1-Bis(3- ² -indolyl)-1-(p-substituted phenyl)methanes Induces Apoptosis through Nuclear Pathways. <i>Journal of Biological Chemistry</i> , 2005, 280, 24903-24914.	3.4	145
15	Curcumin and synthetic analogs induce reactive oxygen species and decreases specificity protein (Sp) transcription factors by targeting microRNAs. <i>BMC Cancer</i> , 2012, 12, 564.	2.6	145
16	Inactivation of the Orphan Nuclear Receptor TR3/Nur77 Inhibits Pancreatic Cancer Cell and Tumor Growth. <i>Cancer Research</i> , 2010, 70, 6824-6836.	0.9	139
17	Betulinic acid inhibits colon cancer cell and tumor growth and induces proteasome-dependent and -independent downregulation of specificity proteins (Sp) transcription factors. <i>BMC Cancer</i> , 2011, 11, 371.	2.6	139
18	Endocrine disruptors and human health: is there a problem. <i>Toxicology</i> , 2004, 205, 3-10.	4.2	137

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19	Identification of a Motif within the 5' Regulatory Region of pS2 Which Is Responsible for AP-1 Binding and TCDD-Mediated Suppression. <i>Biochemistry</i> , 1997, 36, 6080-6089.	2.5	135
20	The long non-coding RNA HOTTIP enhances pancreatic cancer cell proliferation, survival and migration. <i>Oncotarget</i> , 2015, 6, 10840-10852.	1.8	134
21	Nuclear Receptor-Mediated Transactivation Through Interaction with Sp Proteins. <i>Progress in Molecular Biology and Translational Science</i> , 2004, 77, 1-36.	1.9	128
22	Minireview: Role Of Orphan Nuclear Receptors in Cancer and Potential as Drug Targets. <i>Molecular Endocrinology</i> , 2014, 28, 157-172.	3.7	128
23	Oncogenic microRNA-27a is a target for anticancer agent methyl 2-cyano-3,11-dioxo-18 β -olean-1,12-dien-30-oate in colon cancer cells. <i>International Journal of Cancer</i> , 2009, 125, 1965-1974.	5.1	125
24	Increased arylhydrocarbon receptor expression offers a potential therapeutic target for pancreatic cancer. <i>Oncogene</i> , 2002, 21, 6059-6070.	5.9	123
25	Role of the aryl hydrocarbon receptor in carcinogenesis and potential as an anti-cancer drug target. <i>Archives of Toxicology</i> , 2017, 91, 2497-2513.	4.2	123
26	Polychlorinated biphenyls: Correlation between in vivo and in vitro quantitative structure-activity relationships (QSARs). <i>Journal of Toxicology and Environmental Health - Part A: Current Issues</i> , 1985, 16, 379-388.	2.3	122
27	The aryl hydrocarbon receptor ligand omeprazole inhibits breast cancer cell invasion and metastasis. <i>BMC Cancer</i> , 2014, 14, 498.	2.6	118
28	Mechanism of Metformin-dependent Inhibition of Mammalian Target of Rapamycin (mTOR) and Ras Activity in Pancreatic Cancer. <i>Journal of Biological Chemistry</i> , 2014, 289, 27692-27701.	3.4	111
29	GT-094, a NO-NSAID, Inhibits Colon Cancer Cell Growth by Activation of a Reactive Oxygen Species-MicroRNA-27a: ZBTB10-Specificity Protein Pathway. <i>Molecular Cancer Research</i> , 2011, 9, 195-202.	3.4	108
30	Short Chain Fatty Acids Enhance Aryl Hydrocarbon (Ah) Responsiveness in Mouse Colonocytes and Caco-2 Human Colon Cancer Cells. <i>Scientific Reports</i> , 2017, 7, 10163.	3.3	103
31	Mechanism of Action of Phenethylisothiocyanate and Other Reactive Oxygen Species-Inducing Anticancer Agents. <i>Molecular and Cellular Biology</i> , 2014, 34, 2382-2395.	2.3	100
32	Crosstalk between estrogen receptor α and the aryl hydrocarbon receptor in breast cancer cells involves unidirectional activation of proteasomes. <i>FEBS Letters</i> , 2000, 478, 109-112.	2.8	98
33	Structure-dependent inhibition of bladder and pancreatic cancer cell growth by 2-substituted glycyrrhetic and ursolic acid derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2633-2639.	2.2	96
34	1,1-Bis(3-indolyl)-1-(p-chlorophenyl)methane activates the orphan nuclear receptor Nurr1 and inhibits bladder cancer growth. <i>Molecular Cancer Therapeutics</i> , 2008, 7, 3825-3833.	4.1	95
35	Drugs that Target Specificity Proteins Downregulate Epidermal Growth Factor Receptor in Bladder Cancer Cells. <i>Molecular Cancer Research</i> , 2010, 8, 739-750.	3.4	95
36	Targeting NR4A1 (TR3) in cancer cells and tumors. <i>Expert Opinion on Therapeutic Targets</i> , 2011, 15, 195-206.	3.4	94

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37	Clinical correlates of environmental endocrine disruptors. Trends in Endocrinology and Metabolism, 2005, 16, 139-144.	7.1	93
38	Inhibition of Bladder Tumor Growth by 1,1-Bis(3-Indolyl)-1-(p-Substitutedphenyl)Methanes: A New Class of Peroxisome Proliferator-Activated Receptor β Agonists. Cancer Research, 2006, 66, 412-418.	0.9	93
39	Methyl 2-Cyano-3,12-dioxooleana-1,9-dien-28-oate Decreases Specificity Protein Transcription Factors and Inhibits Pancreatic Tumor Growth: Role of MicroRNA-27a. Molecular Pharmacology, 2010, 78, 226-236.	2.3	92
40	Transcription factor Sp1, also known as specificity protein 1 as a therapeutic target. Expert Opinion on Therapeutic Targets, 2014, 18, 759-769.	3.4	89
41	Limitations of the toxic equivalency factor approach for risk assessment of TCDD and related compounds. Teratogenesis, Carcinogenesis, and Mutagenesis, 1997, 17, 285-304.	0.8	87
42	The Orphan Nuclear Receptor NR4A1 (Nur77) Regulates Oxidative and Endoplasmic Reticulum Stress in Pancreatic Cancer Cells. Molecular Cancer Research, 2014, 12, 527-538.	3.4	87
43	Aryl Hydrocarbon Receptor Agonists Induce MicroRNA-335 Expression and Inhibit Lung Metastasis of Estrogen Receptor Negative Breast Cancer Cells. Molecular Cancer Therapeutics, 2012, 11, 108-118.	4.1	85
44	Specificity protein (Sp) transcription factors Sp1, Sp3 and Sp4 are non-oncogene addiction genes in cancer cells. Oncotarget, 2016, 7, 22245-22256.	1.8	85
45	Specificity Protein Transcription Factors and Cancer: Opportunities for Drug Development. Cancer Prevention Research, 2018, 11, 371-382.	1.5	84
46	Structure-Dependent Modulation of Aryl Hydrocarbon Receptor-Mediated Activities by Flavonoids. Toxicological Sciences, 2018, 164, 205-217.	3.1	82
47	Estrogen and aryl hydrocarbon receptor expression and crosstalk in human Ishikawa endometrial cancer cells. Journal of Steroid Biochemistry and Molecular Biology, 2000, 72, 197-207.	2.5	80
48	Betulinic Acid Targets YY1 and ErbB2 through Cannabinoid Receptor-Dependent Disruption of MicroRNA-27a:ZBTB10 in Breast Cancer. Molecular Cancer Therapeutics, 2012, 11, 1421-1431.	4.1	79
49	Diindolylmethane Analogs Bind NR4A1 and Are NR4A1 Antagonists in Colon Cancer Cells. Molecular Endocrinology, 2014, 28, 1729-1739.	3.7	79
50	The aryl hydrocarbon receptor as a target for estrogen receptor-negative breast cancer chemotherapy. Endocrine-Related Cancer, 2009, 16, 835-844.	3.1	77
51	Arsenic trioxide downregulates specificity protein (Sp) transcription factors and inhibits bladder cancer cell and tumor growth. Experimental Cell Research, 2010, 316, 2174-2188.	2.6	77
52	Aryl Hydrocarbon Receptor (AHR)-Active Pharmaceuticals Are Selective AHR Modulators in MDA-MB-468 and BT474 Breast Cancer Cells. Journal of Pharmacology and Experimental Therapeutics, 2012, 343, 333-341.	2.5	77
53	Characterization and Biological Potency of Mono- to Tetra-Halogenated Carbazoles. Environmental Science & Technology, 2015, 49, 10658-10666.	10.0	77
54	Aryl Hydrocarbon Receptor Activity of Tryptophan Metabolites in Young Adult Mouse Colonocytes. Drug Metabolism and Disposition, 2015, 43, 1536-1543.	3.3	76

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55	Estrogen Induces Adenosine Deaminase Gene Expression in MCF-7 Human Breast Cancer Cells: Role of Estrogen Receptor-Sp1 Interactions**This work was supported by the National Institutes of Health (Grant CA-76636), the Welch Foundation, and the Texas Agricultural Experiment Station.. Endocrinology, 1999, 140, 219-227.	2.8	75
56	Aryl hydrocarbon receptor-mediated inhibition of LNCaP prostate cancer cell growth and hormone-induced transactivation. Journal of Steroid Biochemistry and Molecular Biology, 2004, 88, 27-36.	2.5	75
57	Natural Products as Mechanism-based Anticancer Agents: Sp Transcription Factors as Targets. Phytotherapy Research, 2016, 30, 1723-1732.	5.8	75
58	A bioassay to measure energy metabolism in mouse colonic crypts, organoids, and sorted stem cells. American Journal of Physiology - Renal Physiology, 2015, 309, G1-G9.	3.4	72
59	The aryl hydrocarbon receptor (AhR) as a drug target for cancer chemotherapy. Current Opinion in Toxicology, 2017, 2, 24-29.	5.0	72
60	Aryl Hydrocarbon Receptor (AHR) Ligands as Selective AHR Modulators (SAHRMs). International Journal of Molecular Sciences, 2020, 21, 6654.	4.1	69
61	Structure-dependent activation of NR4A2 (Nurr1) by 1,1-bis(3-indolyl)-1-(aromatic)methane analogs in pancreatic cancer cells. Biochemical Pharmacology, 2012, 83, 1445-1455.	4.4	66
62	3',4'-Dimethoxyflavone as an Aryl Hydrocarbon Receptor Antagonist in Human Breast Cancer Cells. Toxicological Sciences, 2000, 58, 235-242.	3.1	65
63	Differential Interaction of the Methoxychlor Metabolite 2,2-Bis-(p-Hydroxyphenyl)-1,1,1-Trichloroethane with Estrogen Receptors \hat{A} and \hat{A} . Endocrinology, 1999, 140, 5746-5753.	2.8	63
64	Aryl hydrocarbon receptor agonists directly activate estrogen receptor $\hat{1}$ in MCF-7 breast cancer cells. Biological Chemistry, 2006, 387, 1209-13.	2.5	62
65	The Nurr1 Activator 1,1-Bis(3-Indolyl)-1-(p-Chlorophenyl)Methane Blocks Inflammatory Gene Expression in BV-2 Microglial Cells by Inhibiting Nuclear Factor $\hat{1}$. Molecular Pharmacology, 2015, 87, 1021-1034.	2.3	62
66	miR-150 regulates obesity-associated insulin resistance by controlling B cell functions. Scientific Reports, 2016, 6, 20176.	3.3	61
67	Development of selective aryl hydrocarbon receptor modulators for treatment of breast cancer. Expert Opinion on Investigational Drugs, 1999, 8, 1385-1396.	4.1	59
68	Omeprazole Inhibits Pancreatic Cancer Cell Invasion through a Nongenomic Aryl Hydrocarbon Receptor Pathway. Chemical Research in Toxicology, 2015, 28, 907-918.	3.3	59
69	Piperlongumine Induces Reactive Oxygen Species (ROS)-Dependent Downregulation of Specificity Protein Transcription Factors. Cancer Prevention Research, 2017, 10, 467-477.	1.5	59
70	Celastrol decreases specificity proteins (Sp) and fibroblast growth factor receptor-3 (FGFR3) in bladder cancer cells. Carcinogenesis, 2012, 33, 886-894.	2.8	57
71	Protective Effects of Ghrelin on Fasting-Induced Muscle Atrophy in Aging Mice. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2020, 75, 621-630.	3.6	56
72	A new class of peroxisome proliferator-activated receptor gamma (PPAR $\hat{\gamma}$) agonists that inhibit growth of breast cancer cells: 1,1-Bis(3'-indolyl)-1-(p-substituted phenyl)methanes. Molecular Cancer Therapeutics, 2004, 3, 247-60.	4.1	55

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73	Histone Deacetylase Inhibitors Inhibit Rhabdomyosarcoma by Reactive Oxygen Species-Dependent Targeting of Specificity Protein Transcription Factors. <i>Molecular Cancer Therapeutics</i> , 2015, 14, 2143-2153.	4.1	53
74	The Paradoxical Roles of Orphan Nuclear Receptor 4A (NR4A) in Cancer. <i>Molecular Cancer Research</i> , 2021, 19, 180-191.	3.4	52
75	Polychlorinated dibenzo-p-dioxins and related compounds: Sources, environmental distribution and risk assessment. <i>Journal of Environmental Science and Health, Part C: Environmental Carcinogenesis and Ecotoxicology Reviews</i> , 1991, 9, 261-302.	2.9	51
76	Nuclear Receptor 4A1 (NR4A1) as a Drug Target for Renal Cell Adenocarcinoma. <i>PLoS ONE</i> , 2015, 10, e0128308.	2.5	51
77	Nuclear receptor 4A1 as a drug target for breast cancer chemotherapy. <i>Endocrine-Related Cancer</i> , 2015, 22, 831-840.	3.1	51
78	Metformin-induced anticancer activities: recent insights. <i>Biological Chemistry</i> , 2018, 399, 321-335.	2.5	51
79	NR4A1 Antagonists Inhibit β 1-Integrin-Dependent Breast Cancer Cell Migration. <i>Molecular and Cellular Biology</i> , 2016, 36, 1383-1394.	2.3	49
80	Neuroprotective Efficacy and Pharmacokinetic Behavior of Novel Anti-Inflammatory <i>Para</i> -Phenyl Substituted Diindolylmethanes in a Mouse Model of Parkinson's Disease. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2013, 345, 125-138.	2.5	48
81	Estrogen-Induced <i>c-fos</i> Protooncogene Expression in MCF-7 Human Breast Cancer Cells: Role of Estrogen Receptor Sp1 Complex Formation. <i>Endocrinology</i> , 1998, 139, 1981-1990.	2.8	47
82	Transforming Growth Factor β 2/NR4A1-Inducible Breast Cancer Cell Migration and Epithelial-to-Mesenchymal Transition Is p38 (Mitogen-Activated Protein Kinase 14) Dependent. <i>Molecular and Cellular Biology</i> , 2017, 37, .	2.3	45
83	Penfluridol Represses Integrin Expression in Breast Cancer through Induction of Reactive Oxygen Species and Downregulation of Sp Transcription Factors. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 205-216.	4.1	45
84	Pharmacological Activators of the NR4A Nuclear Receptors Enhance LTP in a CREB/CBP-Dependent Manner. <i>Neuropsychopharmacology</i> , 2017, 42, 1243-1253.	5.4	45
85	The selective aryl hydrocarbon receptor modulator 6-methyl-1,3,8-trichlorodibenzofuran inhibits prostate tumor metastasis in TRAMP mice. <i>Biochemical Pharmacology</i> , 2009, 77, 1151-1160.	4.4	44
86	Benzyl Isothiocyanate (BITC) Induces Reactive Oxygen Species-dependent Repression of STAT3 Protein by Down-regulation of Specificity Proteins in Pancreatic Cancer. <i>Journal of Biological Chemistry</i> , 2016, 291, 27122-27133.	3.4	44
87	Flavonoids: structure-function and mechanisms of action and opportunities for drug development. <i>Toxicological Research</i> , 2021, 37, 147-162.	2.1	44
88	Novel <i>Para</i> -Phenyl Substituted Diindolylmethanes Protect Against MPTP Neurotoxicity and Suppress Glial Activation in a Mouse Model of Parkinson's Disease. <i>Toxicological Sciences</i> , 2015, 143, 360-373.	3.1	43
89	Aryl hydrocarbon receptor (AhR) ligands as selective AhR modulators: Genomic studies. <i>Current Opinion in Toxicology</i> , 2018, 11-12, 10-20.	5.0	43
90	Editor's Highlight: Microbial-Derived 1,4-Dihydroxy-2-naphthoic Acid and Related Compounds as Aryl Hydrocarbon Receptor Agonists/Antagonists: Structure-Activity Relationships and Receptor Modeling. <i>Toxicological Sciences</i> , 2017, 155, 458-473.	3.1	40

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91	Compensatory Expression of Nur77 and Nurr1 Regulates NF- κ B-Dependent Inflammatory Signaling in Astrocytes. <i>Molecular Pharmacology</i> , 2018, 94, 1174-1186.	2.3	40
92	Transcriptional Activation of Insulin-Like Growth Factor-Binding Protein-4 by 17 β -Estradiol in MCF-7 Cells: Role of Estrogen Receptor-Sp1 Complexes. <i>Endocrinology</i> , 1999, 140, 2501-2508.	2.8	38
93	Estrogen and aryl hydrocarbon responsiveness of ECC-1 endometrial cancer cells. <i>Molecular and Cellular Endocrinology</i> , 1999, 150, 11-21.	3.2	36
94	The Transcriptional Repressor ZBTB4 Regulates EZH2 Through a MicroRNA-ZBTB4-Specificity Protein Signaling Axis. <i>Neoplasia</i> , 2014, 16, 1059-1069.	5.3	36
95	A novel synthetic activator of Nurr1 induces dopaminergic gene expression and protects against 6-hydroxydopamine neurotoxicity in vitro. <i>Neuroscience Letters</i> , 2015, 607, 83-89.	2.1	36
96	Role of metastasis-associated lung adenocarcinoma transcript-1 (MALAT-1) in pancreatic cancer. <i>PLoS ONE</i> , 2018, 13, e0192264.	2.5	36
97	Inhibition of pituitary tumor-transforming gene-1 in thyroid cancer cells by drugs that decrease specificity proteins. <i>Molecular Carcinogenesis</i> , 2011, 50, 655-667.	2.7	35
98	Inhibition of rhabdomyosarcoma cell and tumor growth by targeting specificity protein (Sp) transcription factors. <i>International Journal of Cancer</i> , 2013, 132, 795-806.	5.1	35
99	Estrogenic Activity of a Dieldrin/Toxaphene Mixture in the Mouse Uterus, MCF-7 Human Breast Cancer Cells, and Yeast-Based Estrogen Receptor Assays: No Apparent Synergism. <i>Endocrinology</i> , 1997, 138, 1520-1527.	2.8	35
100	The Nurr1 Ligand, 1,1-bis(3-Indolyl)-1-(4-Chlorophenyl)Methane, Modulates Glial Reactivity and Is Neuroprotective in MPTP-Induced Parkinsonism. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2018, 365, 636-651.	2.5	34
101	The aryl hydrocarbon receptor is a tumor suppressor-like gene in glioblastoma. <i>Journal of Biological Chemistry</i> , 2019, 294, 11342-11353.	3.4	33
102	Mechanism of action and development of selective aryl hydrocarbon receptor modulators for treatment of hormone-dependent cancers (Review). <i>International Journal of Oncology</i> , 2002, 20, 1123-8.	3.3	32
103	Dietary β -Tocopherol-Rich Mixture Inhibits Estrogen-Induced Mammary Tumorigenesis by Modulating Estrogen Metabolism, Antioxidant Response, and PPAR γ . <i>Cancer Prevention Research</i> , 2015, 8, 807-816.	1.5	30
104	Strong adsorption of Polychlorinated Biphenyls by processed montmorillonite clays: Potential applications as toxin enterosorbents during disasters and floods. <i>Environmental Pollution</i> , 2019, 255, 113210.	7.5	30
105	Ultra-flexible nanocarriers for enhanced topical delivery of a highly lipophilic antioxidative molecule for skin cancer chemoprevention. <i>Colloids and Surfaces B: Biointerfaces</i> , 2016, 143, 156-167.	5.0	29
106	Sulindac sulfide inhibits colon cancer cell growth and downregulates specificity protein transcription factors. <i>BMC Cancer</i> , 2015, 15, 974.	2.6	27
107	TGF β -Induced Lung Cancer Cell Migration Is NR4A1-Dependent. <i>Molecular Cancer Research</i> , 2018, 16, 1991-2002.	3.4	27
108	Specificity protein (Sp) transcription factors and metformin regulate expression of the long non-coding RNA HULC. <i>Oncotarget</i> , 2015, 6, 26359-26372.	1.8	27

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109	Loss of Aryl Hydrocarbon Receptor Promotes Colon Tumorigenesis in <i>ApcS580/+; KrasG12D/+</i> Mice. <i>Molecular Cancer Research</i> , 2021, 19, 771-783.	3.4	26
110	Interferon Tau Alleviates Obesity-Induced Adipose Tissue Inflammation and Insulin Resistance by Regulating Macrophage Polarization. <i>PLoS ONE</i> , 2014, 9, e98835.	2.5	26
111	CHARACTERIZATION OF A BOTTLENOSE DOLPHIN (<i>TURSIOPS TRUNCATUS</i>) KIDNEY EPITHELIAL CELL LINE. <i>Marine Mammal Science</i> , 1994, 10, 52-69.	1.8	25
112	Isoflavones as Ah Receptor Agonists in Colon-Derived Cell Lines: Structure-Activity Relationships. <i>Chemical Research in Toxicology</i> , 2019, 32, 2353-2364.	3.3	25
113	A Bis-Indole-Derived NR4A1 Antagonist Induces PD-L1 Degradation and Enhances Antitumor Immunity. <i>Cancer Research</i> , 2020, 80, 1011-1023.	0.9	25
114	PAX3-FOXO1A Expression in Rhabdomyosarcoma Is Driven by the Targetable Nuclear Receptor NR4A1. <i>Cancer Research</i> , 2017, 77, 732-741.	0.9	24
115	Potent inhibition of breast cancer by bis-indole-derived nuclear receptor 4A1 (NR4A1) antagonists. <i>Breast Cancer Research and Treatment</i> , 2019, 177, 29-40.	2.5	24
116	Pharmacological activation of Nr4a rescues age-associated memory decline. <i>Neurobiology of Aging</i> , 2020, 85, 140-144.	3.1	24
117	Flavonoids kaempferol and quercetin are nuclear receptor 4A1 (NR4A1, Nur77) ligands and inhibit rhabdomyosarcoma cell and tumor growth. <i>Journal of Experimental and Clinical Cancer Research</i> , 2021, 40, 392.	8.6	24
118	Transcriptional activation of rat creatine kinase B by 17 β -estradiol in MCF-7 cells involves an estrogen responsive element and GC-rich sites. <i>Journal of Cellular Biochemistry</i> , 2002, 84, 156-172.	2.6	23
119	Nuclear receptor 4A1 (NR4A1) as a drug target for treating rhabdomyosarcoma (RMS). <i>Oncotarget</i> , 2016, 7, 31257-31269.	1.8	23
120	Diet-Host-Microbiota Interactions Shape Aryl Hydrocarbon Receptor Ligand Production to Modulate Intestinal Homeostasis. <i>Annual Review of Nutrition</i> , 2021, 41, 455-478.	10.1	23
121	Gas chromatographic/mass spectrometric analysis of specific isomers of polychlorodibenzofurans. <i>Biological Mass Spectrometry</i> , 1985, 12, 247-253.	0.5	22
122	Inactivation of the orphan nuclear receptor NR4A1 contributes to apoptosis induction by fangchinoline in pancreatic cancer cells. <i>Toxicology and Applied Pharmacology</i> , 2017, 332, 32-39.	2.8	22
123	Reactive Oxygen Species (ROS)-Inducing Triterpenoid Inhibits Rhabdomyosarcoma Cell and Tumor Growth through Targeting Sp Transcription Factors. <i>Molecular Cancer Research</i> , 2019, 17, 794-805.	3.4	22
124	Dual targeting of Nur77 and AMPK α by isolantolactone inhibits adipogenesis in vitro and decreases body fat mass in vivo. <i>International Journal of Obesity</i> , 2019, 43, 952-962.	3.4	22
125	MicroRNA-Specificity Protein (Sp) Transcription Factor Interactions and Significance in Carcinogenesis. <i>Current Pharmacology Reports</i> , 2015, 1, 73-78.	3.0	21
126	The role of xenoestrogenic compounds in the development of breast cancer. <i>Trends in Pharmacological Sciences</i> , 2006, 27, 447-454.	8.7	20

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127	High expression of orphan nuclear receptor NR4A1 in a subset of ovarian tumors with worse outcome. <i>Gynecologic Oncology</i> , 2016, 141, 348-356.	1.4	20
128	Orphan nuclear receptor 4A1 (NR4A1) and novel ligands. <i>Essays in Biochemistry</i> , 2021, 65, 877-886.	4.7	20
129	Endocrine disruptors and falling sperm counts: lessons learned or not!. <i>Asian Journal of Andrology</i> , 2013, 15, 191-194.	1.6	19
130	CDODA-Me decreases specificity protein transcription factors and induces apoptosis in bladder cancer cells through induction of reactive oxygen species. <i>Urologic Oncology: Seminars and Original Investigations</i> , 2016, 34, 337.e11-337.e18.	1.6	18
131	The nuclear orphan receptor NR4A1 regulates β 1 integrin expression in pancreatic and colon cancer cells and can be targeted by NR4A1 antagonists. <i>Molecular Carcinogenesis</i> , 2017, 56, 2066-2075.	2.7	18
132	Nuclear receptor 4A2 (NR4A2) is a druggable target for glioblastomas. <i>Journal of Neuro-Oncology</i> , 2020, 146, 25-39.	2.9	18
133	Ah receptor ligands and their impacts on gut resilience: structure-activity effects. <i>Critical Reviews in Toxicology</i> , 2020, 50, 463-473.	3.9	18
134	CF3DODA-Me induces apoptosis, degrades Sp1, and blocks the transformation phase of the blebbistatin emergency program. <i>Apoptosis: an International Journal on Programmed Cell Death</i> , 2017, 22, 719-729.	4.9	17
135	Suppression of aberrant choroidal neovascularization through activation of the aryl hydrocarbon receptor. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2018, 1864, 1583-1595.	3.8	17
136	Inhibition of pancreatic cancer Panc1 cell migration by omeprazole is dependent on aryl hydrocarbon receptor activation of JNK. <i>Biochemical and Biophysical Research Communications</i> , 2018, 501, 751-757.	2.1	17
137	Bis-Indole-Derived NR4A1 Ligands and Metformin Exhibit NR4A1-Dependent Glucose Metabolism and Uptake in C2C12 Cells. <i>Endocrinology</i> , 2018, 159, 1950-1963.	2.8	17
138	2,3,7,8-Tetrachlorodibenzo-p-dioxin has both pro-carcinogenic and anti-carcinogenic effects on neuroendocrine prostate carcinoma formation in TRAMP mice. <i>Toxicology and Applied Pharmacology</i> , 2016, 305, 242-249.	2.8	16
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