Peter A Crooks

List of Publications by Year in descending order

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439 papers

9,208 citations

45 h-index 70 g-index

443 all docs 443 docs citations

times ranked

443

9055 citing authors

#	Article	IF	CITATIONS
1	An orally bioavailable parthenolide analog selectively eradicates acute myelogenous leukemia stem and progenitor cells. Blood, 2007, 110, 4427-4435.	1.4	357
2	High performance liquid chromatographic analysis of the pharmacologically active quinones and related compounds in the oil of the black seed (Nigella sativa L.). Journal of Pharmaceutical and Biomedical Analysis, 1999, 19, 757-762.	2.8	274
3	A novel mechanism of action and potential use for lobeline as a treatment for psychostimulant abuse. Biochemical Pharmacology, 2002, 63, 89-98.	4.4	199
4	Aminoparthenolides as novel anti-leukemic agents: Discovery of the NF-κB inhibitor, DMAPT (LC-1). Bioorganic and Medicinal Chemistry Letters, 2009, 19, 4346-4349.	2.2	168
5	Targeting Aberrant Glutathione Metabolism to Eradicate Human Acute Myelogenous Leukemia Cells. Journal of Biological Chemistry, 2013, 288, 33542-33558.	3.4	163
6	Molecular pathway for thymoquinone-induced cell-cycle arrest and apoptosis in neoplastic keratinocytes. Anti-Cancer Drugs, 2004, 15, 389-399.	1.4	162
7	The NF-κB subunit Rel A is associated with in vitro survival and clinical disease progression in chronic lymphocytic leukemia and represents a promising therapeutic target. Blood, 2008, 111, 4681-4689.	1.4	145
8	A NADPH Oxidase–Dependent Redox Signaling Pathway Mediates the Selective Radiosensitization Effect of Parthenolide in Prostate Cancer Cells. Cancer Research, 2010, 70, 2880-2890.	0.9	117
9	Contribution of CNS nicotine metabolites to the neuropharmacological effects of nicotine and tobacco smoking. Biochemical Pharmacology, 1997, 54, 743-753.	4.4	110
10	Vesicular monoamine transporter 2: Role as a novel target for drug development. AAPS Journal, 2006, 8, E682-E692.	4.4	104
11	Cellulose sulfuric acid: An efficient biodegradable and recyclable solid acid catalyst for the one-pot synthesis of aryl-14H-dibenzo[a.j]xanthenes under solvent-free conditions. Journal of Molecular Catalysis A, 2009, 304, 85-87.	4.8	99
12	Lobeline Displaces [3H]Dihydrotetrabenazine Binding and Releases [3H]Dopamine from Rat Striatal Synaptic Vesicles: Comparison with d-Amphetamine. Journal of Neurochemistry, 2002, 71, 258-265.	3.9	94
13	Antileukemic activity of aminoparthenolide analogs. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 3870-3873.	2.2	85
14	Persistent Activation of NF-κB in BRCA1-Deficient Mammary Progenitors Drives Aberrant Proliferation and Accumulation of DNA Damage. Cell Stem Cell, 2016, 19, 52-65.	11.1	85
15	Extending the analysis of nicotinic receptor antagonists with the study of $\hat{l}\pm 6$ nicotinic receptor subunit chimeras. Neuropharmacology, 2008, 54, 1189-1200.	4.1	82
16	NADPH oxidase activity is essential for Keap1/Nrf2-mediated induction of GCLC in response to 2-indol-3-yl-methylenequinuclidin-3-ols. Cancer Research, 2003, 63, 5636-45.	0.9	80
17	3-[Benzimidazo- and 3-[benzothiadiazoleimidazo-(1,2-c)quinazolin-5-yl]-2H-chromene-2-ones as potent antimicrobial agents. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 524-527.	2.2	77
18	The radiosensitization effect of parthenolide in prostate cancer cells is mediated by nuclear factor- $\hat{\mathbb{P}}$ B inhibition and enhanced by the presence of PTEN. Molecular Cancer Therapeutics, 2007, 6, 2477-2486.	4.1	74

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19	Once weekly administration of nicotine produces long-lasting locomotor sensitization in rats via a nicotinic receptor-mediated mechanism. Psychopharmacology, 2001, 156, 469-476.	3.1	69
20	Chemical genomic screening reveals synergism between parthenolide and inhibitors of the PI-3 kinase and mTOR pathways. Blood, 2010, 116, 5983-5990.	1.4	69
21	A pilot study of plasma caffeine concentrations in a US sample of smoker and nonsmoker volunteers. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2003, 27, 165-171.	4.8	68
22	The pharmacological activity of nicotine and nornicotine on nAChRs subtypes: relevance to nicotine dependence and drug discovery. Journal of Neurochemistry, 2007, 101, 160-167.	3.9	66
23	Synthesis and evaluation of chromenyl barbiturates and thiobarbiturates as potential antitubercular agents. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 4329-4331.	2.2	64
24	Lobeline Analogs with Enhanced Affinity and Selectivity for Plasmalemma and Vesicular Monoamine Transporters. Journal of Pharmacology and Experimental Therapeutics, 2004, 310, 1035-1045.	2.5	63
25	Active Transport of High-Affinity Choline and Nicotine Analogs into the Central Nervous System by the Blood-Brain Barrier Choline Transporter. Journal of Pharmacology and Experimental Therapeutics, 2003, 304, 1268-1274.	2.5	61
26	Lobeline inhibits nicotine-evoked [3H]dopamine overflow from rat striatal slices and nicotine-evoked 86Rb+ efflux from thalamic synaptosomes. Neuropharmacology, 2000, 39, 2654-2662.	4.1	60
27	Defunctionalized Lobeline Analogues:Â Structureâ-'Activity of Novel Ligands for the Vesicular Monoamine Transporter. Journal of Medicinal Chemistry, 2005, 48, 5551-5560.	6.4	59
28	Acute and chronic effects of nornicotine on locomotor activity in rats: altered response to nicotine. Psychopharmacology, 1999, 145, 442-451.	3.1	58
29	KEAP1 Is a Redox Sensitive Target That Arbitrates the Opposing Radiosensitive Effects of Parthenolide in Normal and Cancer Cells. Cancer Research, 2013, 73, 4406-4417.	0.9	57
30	Synthesis of thymidine dimers containing internucleoside sulfonate and sulfonamide linkages. Journal of Organic Chemistry, 1992, 57, 2983-2985.	3.2	56
31	Effects of norketamine enantiomers in rodent models of persistent pain. Pharmacology Biochemistry and Behavior, 2008, 90, 676-685.	2.9	56
32	Synthesis and biological evaluation of novel 4,5-disubstituted 2H-1,2,3-triazoles as cis-constrained analogues of combretastatin A-4. European Journal of Medicinal Chemistry, 2015, 103, 123-132.	5. 5	56
33	An in vivo evaluation of the antiseizure activity and acute neurotoxicity of agmatine. Pharmacology Biochemistry and Behavior, 2003, 74, 771-775.	2.9	54
34	Lobelane decreases methamphetamine self-administration in rats. European Journal of Pharmacology, 2007, 571, 33-38.	3.5	54
35	Transdermal Delivery of Naltrexol and Skin Permeability Lifetime after Microneedle Treatment in Hairless Guinea Pigs. Journal of Pharmaceutical Sciences, 2010, 99, 3072-3080.	3.3	54
36	QSAR modeling of mono- and bis-quaternary ammonium salts that act as antagonists at neuronal nicotinic acetylcholine receptors mediating dopamine release. Bioorganic and Medicinal Chemistry, 2006, 14, 3017-3037.	3.0	53

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37	Nicotinic receptor-based therapeutics and candidates for smoking cessation. Biochemical Pharmacology, 2009, 78, 732-743.	4.4	53
38	Flux Across of Microneedle-treated Skin is Increased by Increasing Charge of Naltrexone and Naltrexol In Vitro. Pharmaceutical Research, 2008, 25, 1677-1685.	3.5	52
39	Identification and synthesis of novel alkaloids from the root system of Nicotiana tabacum: Affinity for neuronal nicotinic acetylcholine receptors. Life Sciences, 2005, 78, 495-505.	4.3	50
40	Melampomagnolide B: A new antileukemic sesquiterpene. Bioorganic and Medicinal Chemistry, 2011, 19, 1515-1519.	3.0	50
41	Pharmacologically Distinct Nicotinic Acetylcholine Receptors Drive Efferent-Mediated Excitation in Calyx-Bearing Vestibular Afferents. Journal of Neuroscience, 2015, 35, 3625-3643.	3.6	50
42	Synthesis and anti-cancer screening of novel heterocyclic-(2H)-1,2,3-triazoles as potential anti-cancer agents. MedChemComm, 2015, 6, 1535-1543.	3.4	49
43	In vitro permeation of a pegylated naltrexone prodrug across microneedle-treated skin. Journal of Controlled Release, 2010, 146, 37-44.	9.9	48
44	Synthesis and evaluation of a series of benzothiophene acrylonitrile analogs as anticancer agents. MedChemComm, 2013, 4, 1073.	3.4	48
45	Sodium fluoride as an efficient catalyst for the synthesis of 2,4-disubstituted-1,3-thiazoles and selenazoles at ambient temperature. Chinese Chemical Letters, 2014, 25, 172-175.	9.0	48
46	Inhibition of nicotine-evoked [3H] dopamine release by pyridino N-substituted nicotine analogues: A new class of nicotinic antagonist. Drug Development Research, 1995, 36, 91-102.	2.9	47
47	A simple high performance liquid chromatographic method for the quantification of total cotinine, total 38^{-2} -hydroxycotinine and caffeine in the plasma of smokers. Journal of Pharmaceutical and Biomedical Analysis, 2000, 23, 543-549.	2.8	46
48	Modeling Multiple Species of Nicotine and Deschloroepibatidine Interacting with $\hat{l}\pm4\hat{l}^22$ Nicotinic Acetylcholine Receptor:Â From Microscopic Binding to Phenomenological Binding Affinity. Journal of the American Chemical Society, 2005, 127, 14401-14414.	13.7	46
49	Modeling Subtype-Selective Agonists Binding with $\hat{l}\pm4\hat{l}^22$ and $\hat{l}\pm7$ Nicotinic Acetylcholine Receptors: \hat{A} Effects of Local Binding and Long-Range Electrostatic Interactions. Journal of Medicinal Chemistry, 2006, 49, 7661-7674.	6.4	46
50	Lobelane Inhibits Methamphetamine-Evoked Dopamine Release via Inhibition of the Vesicular Monoamine Transporter-2. Journal of Pharmacology and Experimental Therapeutics, 2010, 332, 612-621.	2.5	45
51	The Vesicular Monoamine Transporter-2. Advances in Pharmacology, 2014, 69, 71-106.	2.0	45
52	Nicotinic Receptor Antagonists as Treatments for Nicotine Abuse. Advances in Pharmacology, 2014, 69, 513-551.	2.0	44
53	bis-Azaaromatic quaternary ammonium analogues: ligands for $\hat{l}\pm4\hat{l}^22^*$ and $\hat{l}\pm7^*$ subtypes of neuronal nicotinic receptors. Bioorganic and Medicinal Chemistry Letters, 2002, 12, 3067-3071.	2.2	43
54	N-n-Alkylnicotinium Analogs, a Novel Class of Nicotinic Receptor Antagonists: Interaction with $\hat{l}\pm4\hat{l}^22^*$ and $\hat{l}\pm7^*$ Neuronal Nicotinic Receptors. Journal of Pharmacology and Experimental Therapeutics, 2003, 304, 400-410.	2.5	43

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55	S(-)-Nornicotine Increases Dopamine Release in a Calcium-Dependent Manner from Superfused Rat Striatal Slices. Journal of Neurochemistry, 1993, 60, 2167-2174.	3.9	42
56	Total Cotinine in Plasma: A Stable Biomarker for Exposure to Tobacco Smoke. Journal of Clinical Psychopharmacology, 2002, 22, 496-501.	1.4	42
57	Subtype-selective nicotinic receptor antagonists: potential as tobacco use cessation agents. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 1863-1867.	2.2	42
58	Enhancement of transdermal delivery of $6-\hat{l}^2$ -naltrexol via a codrug linked to hydroxybupropion. Journal of Controlled Release, 2006, 113, 137-145.	9.9	42
59	The effects of a novel nicotinic receptor antagonist N,N-dodecane-1,12-diyl-bis-3-picolinium dibromide (bPiDDB) on acute and repeated nicotine-induced increases in extracellular dopamine in rat nucleus accumbens. Neuropharmacology, 2007, 52, 755-763.	4.1	42
60	Combination therapy with 5-fluorouracil and L-canavanine. Anti-Cancer Drugs, 1995, 6, 586-593.	1.4	41
61	Toxicity of Dipyridyl Compounds and Related Compounds. Critical Reviews in Toxicology, 2004, 34, 447-460.	3.9	41
62	One-pot multicomponent synthesis of indole incorporated thiazolylcoumarins and their antibacterial, anticancer and DNA cleavage studies. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 106-112.	2.2	41
63	Recent developments in neuronal nicotinic acetylcholine receptor antagonists. Expert Opinion on Therapeutic Patents, 2000, 10, 1561-1581.	5.0	40
64	Suppression of pancreatic tumor growth by combination chemotherapy with sulindac and LC-1 is associated with cyclin D1 inhibition in vivo. Molecular Cancer Therapeutics, 2007, 6, 1736-1744.	4.1	39
65	Novel Small Molecule $\hat{i}\pm 9\hat{i}\pm 10$ Nicotinic Receptor Antagonist Prevents and Reverses Chemotherapy-Evoked Neuropathic Pain in Rats. Anesthesia and Analgesia, 2012, 115, 713-720.	2.2	39
66	N-n-Alkylnicotinium Analogs, A Novel Class of Nicotinic Receptor Antagonist: Inhibition ofS(â^')-Nicotine-Evoked [3H]Dopamine Overflow from Superfused Rat Striatal Slices. Journal of Pharmacology and Experimental Therapeutics, 2002, 301, 1088-1096.	2.5	38
67	Synthesis and analgesic properties of two leucine-enkephalin analogs containing a conformationally restrained N-terminal tyrosine residue. Journal of Medicinal Chemistry, 1983, 26, 762-765.	6.4	37
68	Novel Chemical Enhancers of Heat Shock Increase Thermal Radiosensitization through a Mitotic Catastrophe Pathway. Cancer Research, 2007, 67, 695-701.	0.9	37
69	<i>N,N</i> ′-Alkane-diyl- <i>bis</i> -3-picoliniums as Nicotinic Receptor Antagonists: Inhibition of Nicotine-Evoked Dopamine Release and Hyperactivity. Journal of Pharmacology and Experimental Therapeutics, 2008, 326, 563-576.	2.5	37
70	4-trimethylammonium antipyrine: A quaternary ammonium nonradionuclide marker for blood-brain barrier integrity during in vivo microdialysis. Journal of Pharmacological and Toxicological Methods, 1992, 28, 129-135.	0.7	36
71	Synthesis and hydrolytic behavior of two novel tripartate codrugs of naltrexone and $6\hat{l}^2$ -naltrexol with hydroxybupropion as potential alcohol abuse and smoking cessation agents. Bioorganic and Medicinal Chemistry, 2006, 14, 7051-7061.	3.0	36
72	Phase I and Phase II Ocular Metabolic Activities and the Role of Metabolism in Ophthalmic Prodrug and Codrug Design and Delivery. Molecules, 2007, 12, 373-388.	3.8	36

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73	Synthesis and in vitro evaluation of N-alkyl-3-hydroxy-3-(2-imino-3-methyl-5-oxoimidazolidin-4-yl)indolin-2-one analogs as potential anticancer agents. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 4468-4471.	2.2	36
74	A duplex "Gemini―prodrug of naltrexone for transdermal delivery. Journal of Controlled Release, 2004, 97, 283-290.	9.9	35
75	Lobelane analogues as novel ligands for the vesicular monoamine transporter-2. Bioorganic and Medicinal Chemistry, 2005, 13, 3899-3909.	3.0	35
76	In Vitro/in Vivo Correlation of Transdermal Naltrexone Prodrugs in Hairless Guinea Pigs. Pharmaceutical Research, 2005, 22, 981-989.	3.5	35
77	Dimethylaminoparthenolide and gemcitabine: a survival study using a genetically engineered mouse model of pancreatic cancer. BMC Cancer, 2013, 13, 194.	2.6	35
78	1-Benzyl-2-methyl-3-indolylmethylene barbituric acid derivatives: Anti-cancer agents that target nucleophosmin 1 (NPM1). Bioorganic and Medicinal Chemistry, 2015, 23, 7226-7233.	3.0	35
79	Synthesis and anti-proliferative activity of aromatic substituted 5-((1-benzyl-1H-indol-3-yl)methylene)-1,3-dimethylpyrimidine-2,4,6(1H,3H,5H)-trione analogs against human tumor cell lines. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 601-603.	2.2	34
80	MMB triazole analogs are potent NF-κB inhibitors and anti-cancer agents against both hematological and solid tumor cells. European Journal of Medicinal Chemistry, 2018, 157, 562-581.	5.5	34
81	Physicochemical Evaluation, in Vitro Human Skin Diffusion, and Concurrent Biotransformation of 3-O-Alkyl Carbonate Prodrugs of Naltrexone. Pharmaceutical Research, 2004, 21, 1146-1152.	3.5	33
82	Development of subtype-selective ligands as antagonists at nicotinic receptors mediating nicotine-evoked dopamine release. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 1869-1874.	2.2	32
83	Characterization of structurally novel G protein biased CB 1 agonists: Implications for drug development. Pharmacological Research, 2017, 125, 161-177.	7.1	32
84	Canvass: A Crowd-Sourced, Natural-Product Screening Library for Exploring Biological Space. ACS Central Science, 2018, 4, 1727-1741.	11.3	32
85	3D-QSAR study of bis-azaaromatic quaternary ammonium analogs at the blood–brain barrier choline transporter. Bioorganic and Medicinal Chemistry, 2005, 13, 4253-4261.	3.0	31
86	Novel antiglaucoma prodrugs and codrugs of ethacrynic acid. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 3524-3527.	2.2	31
87	In vivo evaluation of 3-0-alkyl ester transdermal prodrugs of naltrexone in hairless guinea pigs. Journal of Controlled Release, 2005, 102, 509-520.	9.9	31
88	Synthesis and evaluation of a series of resveratrol analogues as potent anti-cancer agents that target tubulin. MedChemComm, 2015, 6, 788-794.	3.4	31
89	A novel tetrazole analogue of resveratrol is a potent anticancer agent. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 172-178.	2,2	31
90	Stereospecific in vitro N-methylation of nicotine in guinea pig tissues by an S-adenosylmethionine-dependent N-methyltransferase. Biochemical Pharmacology, 1985, 34, 281-284.	4.4	30

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91	Transdermal Delivery of Naltrexone and its Active Metabolite $6 \cdot \hat{l}^2$ -Naltrexol in Human Skin in Vitro and Guinea Pigs in Vivo. Journal of Pharmaceutical Sciences, 2005, 94, 1965-1975.	3.3	30
92	Novel substituted (Z)-5-((N-benzyl-1H-indol-3-yl)methylene)imidazolidine-2,4-diones and 5-((N-benzyl-1H-indol-3-yl)methylene)pyrimidine-2,4,6(1H,3H,5H)-triones as potent radio-sensitizing agents. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 600-602.	2.2	30
93	Rational Design of a Parthenolide-based Drug Regimen That Selectively Eradicates Acute Myelogenous Leukemia Stem Cells. Journal of Biological Chemistry, 2016, 291, 21984-22000.	3.4	30
94	Formation of quaternary amines by N- methylation of azaheterocycles with homogeneous amine n-methyltransferases. Biochemical Pharmacology, 1988, 37, 1673-1677.	4.4	29
95	A novel technique for visualizing the intracellular localization and distribution of transported polyamines in cultured pulmonary artery smooth muscle cells. Journal of Pharmaceutical and Biomedical Analysis, 1998, 17, 307-320.	2.8	29
96	Human Skin Permeation of Branched-Chain 3-O-Alkyl Ester and Carbonate Prodrugs of Naltrexone. Pharmaceutical Research, 2005, 22, 758-765.	3.5	29
97	Discovery of non-peptide, small molecule antagonists of $\hat{l}\pm 9\hat{l}\pm 10$ nicotinic acetylcholine receptors as novel analgesics for the treatment of neuropathic and tonic inflammatory pain. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 2476-2479.	2.2	29
98	Dimers of Melampomagnolide B Exhibit Potent Anticancer Activity against Hematological and Solid Tumor Cells. Journal of Medicinal Chemistry, 2015, 58, 8896-8906.	6.4	29
99	Pharmacological similarities between native brain and heterologously expressed $\hat{l}\pm4\hat{l}^2$ 2 nicotinic receptors. British Journal of Pharmacology, 1999, 128, 1291-1299.	5.4	28
100	Contributory role for nornicotine in nicotine neuropharmacology: nornicotine-evoked [3H]dopamine overflow from rat nucleus accumbens slices11Abbreviations: DA, dopamine; and DHβE, dihydro-β-erythroidine Biochemical Pharmacology, 2001, 62, 1597-1603.	4.4	28
101	The antiproliferative and immunotoxic effects of L-canavanine and L-canaline. Anti-Cancer Drugs, 2002, 13, 313-320.	1.4	28
102	Indirect Trapping of the Retroconjugate Addition Reaction Intermediate Involved in the Epimerization of Lobeline: Application to the Synthesis of (â^')-Sedamine. Journal of Organic Chemistry, 2004, 69, 8514-8517.	3.2	28
103	Opiate receptor binding properties of morphine-, dihydromorphine-, and codeine 6-O-sulfate ester congeners. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 4291-4295.	2.2	28
104	Development of a GC-MS Assay for the Determination of Fentanyl Pharmacokinetics in Rabbit Plasma after Sublingual Spray Delivery. AAPS Journal, 2008, 10, 261-267.	4.4	28
105	Targeting Nucleophosmin 1 Represents a Rational Strategy for Radiation Sensitization. International Journal of Radiation Oncology Biology Physics, 2014, 89, 1106-1114.	0.8	28
106	DMAPT inhibits NF-κB activity and increases sensitivity of prostate cancer cells to X-rays in vitro and in tumor xenografts in vivo. Free Radical Biology and Medicine, 2017, 112, 318-326.	2.9	28
107	Remarkable substrate-inhibitor properties of nicotine enantiomers towards a guinea pig lung aromatic azaheterocycle N-methyltransferase. Biochemical and Biophysical Research Communications, 1985, 128, 312-316.	2.1	27
108	Computational neural network analysis of the affinity of lobeline and tetrabenazine analogs for the vesicular monoamine transporter-2. Bioorganic and Medicinal Chemistry, 2007, 15, 2975-2992.	3.0	27

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109	Synthesis and evaluation of a series of homologues of lobelane at the vesicular monoamine transporter-2. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 6509-6512.	2.2	27
110	The effect of a novel VMAT2 inhibitor, GZ-793A, on methamphetamine reward in rats. Psychopharmacology, 2012, 220, 395-403.	3.1	27
111	5-((1-Aroyl-1H-indol-3-yl)methylene)-2-thioxodihydropyrimidine-4,6(1H,5H)-diones as potential anticancer agents with anti-inflammatory properties. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 1442-1446.	2.2	27
112	Anti-cancer activity of carbamate derivatives of melampomagnolide B. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 3499-3502.	2.2	27
113	A Small-Molecule Inhibitor of Human DNA Polymerase η Potentiates the Effects of Cisplatin in Tumor Cells. Biochemistry, 2018, 57, 1262-1273.	2.5	27
114	Novel substituted (Z)-2-(N-benzylindol-3-ylmethylene)quinuclidin-3-one and (Z)-(±)-2-(N-benzylindol-3-ylmethylene)quinuclidin-3-ol derivatives as potent thermal sensitizing agents. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 6821-6824.	2.2	26
115	Effect of Celecoxib and the Novel Anti-Cancer Agent, Dimethylamino-Parthenolide, in a Developmental Model of Pancreatic Cancer. Pancreas, 2008, 37, e45-e53.	1.1	26
116	Mecamylamine, dihydro- \hat{l}^2 -erythroidine, and dextromethorphan block conditioned responding evoked by the conditional stimulus effects of nicotine. Pharmacology Biochemistry and Behavior, 2009, 94, 319-328.	2.9	26
117	Microwave assisted synthesis and in vitro cytotoxicities of substituted (Z)-2-amino-5-(1-benzyl-1H-indol-3-yl)methylene-1-methyl-1H-imidazol-4(5H)-ones against human tumor cell lines. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 591-593.	2.2	26
118	Dimethylamino Parthenolide Enhances the Inhibitory Effects of Gemcitabine in Human Pancreatic Cancer Cells. Journal of Gastrointestinal Surgery, 2012, 16, 1333-1340.	1.7	26
119	Multiple Modes of <i>α</i> 7 nAChR Noncompetitive Antagonism of Control Agonist-Evoked and Allosterically Enhanced Currents. Molecular Pharmacology, 2013, 84, 459-475.	2.3	26
120	Identification of resveratrol analogs as potent antiâ€dengue agents using a cellâ€based assay. Journal of Medical Virology, 2017, 89, 397-407.	5.0	26
121	Design of novel prodrugs for the enhancement of the transdermal penetration of indomethacin. International Journal of Pharmaceutics, 1995, 123, 127-136.	5.2	25
122	Neuronal nicotinic acetylcholine receptor binding affinities of boron-containing nicotine analogues. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 1245-1248.	2.2	25
123	Transdermal Delivery of Bupropion and its Active Metabolite, Hydroxybupropion: A Prodrug Strategy as an Alternative Approach. Journal of Pharmaceutical Sciences, 2009, 98, 583-594.	3.3	25
124	Cellulose Sulfuric Acid: An Efficient Biodegradable and Recyclable Solid Acid Catalyst for the One-Pot Synthesis of 3,4-Dihydropyrimidine-2(1 <i>H</i>)-ones. Synthetic Communications, 2009, 39, 1257-1263.	2.1	25
125	Aplysinopsin analogs: Synthesis and anti-proliferative activity of substituted (Z)-5-(N-benzylindol-3-ylmethylene)imidazolidine-2,4-diones. Bioorganic and Medicinal Chemistry, 2010, 18, 3570-3574.	3.0	25
126	bPiDI: a novel selective $\hat{l}\pm6\hat{l}^22^*$ nicotinic receptor antagonist and preclinical candidate treatment for nicotine abuse. British Journal of Pharmacology, 2011, 163, 346-357.	5.4	25

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127	<i>N</i> -Aroyl Indole Thiobarbituric Acids as Inhibitors of DNA Repair and Replication Stress Response Polymerases. ACS Chemical Biology, 2013, 8, 1722-1729.	3.4	25
128	Chiral purity determination of tobacco alkaloids and nicotine-like compounds by 1H NMR spectroscopy in the presence of 1,1?-binaphthyl-2,2?-diylphosphoric acid. Chirality, 1996, 8, 295-299.	2.6	24
129	Ceric Ammonium Nitrate (CAN): An Efficient Catalyst for the Coumarin Synthesis via Pechmann Condensation using Conventional Heating and Microwave Irradiation. Synthetic Communications, 2008, 38, 2082-2088.	2.1	24
130	Synthesis and in vitro screening of novel N-benzyl aplysinopsin analogs as potential anticancer agents. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 1411-1413.	2.2	24
131	Synthesis and evaluation of conformationally restricted pyridinoN-alkylated nicotine analogs as nicotinic acetylcholine receptor antagonists. Drug Development Research, 2002, 55, 173-186.	2.9	23
132	Bioconversion of Naltrexone and Its 3-O-Alkyl-Ester Prodrugs in a Human Skin Equivalent. Journal of Pharmaceutical Sciences, 2005, 94, 828-836.	3.3	23
133	Cellulose Sulfuric Acid: An Efficient Biodegradable and Recyclable Solid Acid Catalyst for the Synthesis of 1-Oxo-hexahydroxanthene. Synthetic Communications, 2011, 41, 1719-1724.	2.1	23
134	The Novel Chemical Entity YTR107 Inhibits Recruitment of Nucleophosmin to Sites of DNA Damage, Suppressing Repair of DNA Double-Strand Breaks and Enhancing Radiosensitization. Clinical Cancer Research, 2011, 17, 6490-6499.	7.0	23
135	Efficacy of Dimethylaminoparthenolide and Sulindac in Combination With Gemcitabine in a Genetically Engineered Mouse Model of Pancreatic Cancer. Pancreas, 2013, 42, 160-167.	1.1	23
136	Indole carboxylic acid esters of melampomagnolide B are potent anticancer agents against both hematological and solid tumor cells. European Journal of Medicinal Chemistry, 2017, 136, 393-405.	5.5	23
137	Structural modeling of GSK3 \hat{l}^2 implicates the inactive (DFG-out) conformation as the target bound by TDZD analogs. Scientific Reports, 2020, 10, 18326.	3.3	23
138	Carrier-Mediated Transport of the Quaternary Ammonium Neuronal Nicotinic Receptor Antagonist ⟨i⟩N⟨ i⟩′-Dodecylbispicolinium Dibromide at the Blood-Brain Barrier. Journal of Pharmacology and Experimental Therapeutics, 2008, 324, 244-250.	2.5	22
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