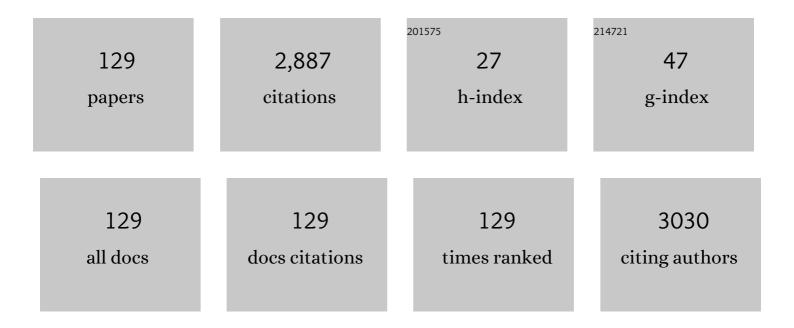
List of Publications by Year in descending order

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ΔΝΝΑ ΙΔΝΕCΚΑ

#	Article	IF	CITATIONS
1	Attachment of Chiral Functional Groups to Modify the Activity of New GPx Mimetics. Materials, 2022, 15, 2068.	1.3	7
2	Synthesis, Pharmacological Evaluation, and Computational Studies of Cyclic Opioid Peptidomimetics Containing β3-Lysine. Molecules, 2022, 27, 151.	1.7	1
3	Synthesis and structure–activity relationship study of novel 3-diethoxyphosphorylfuroquinoline-4,9-diones with potent antitumor efficacy. European Journal of Medicinal Chemistry, 2021, 219, 113429.	2.6	11
4	Harnessing the Anti-Nociceptive Potential of NK2 and NK3 Ligands in the Design of New Multifunctional μ/Π-Opioid Agonist–Neurokinin Antagonist Peptidomimetics. Molecules, 2021, 26, 5406.	1.7	2
5	Pharmacological Characterization of µ-Opioid Receptor Agonists with Biased G Protein or β-Arrestin Signaling, and Computational Study of Conformational Changes during Receptor Activation. Molecules, 2021, 26, 13.	1.7	12
6	Potential of Nociceptin/Orphanin FQ Peptide Analogs for Drug Development. Chemistry and Biodiversity, 2021, 18, e2000871.	1.0	5
7	Synthesis and Cytotoxic Activity of Lepidilines A–D: Comparison with Some 4,5-Diphenyl Analogues and Related Imidazole-2-thiones. Journal of Natural Products, 2021, 84, 3071-3079.	1.5	6
8	Design, Synthesis and Functional Analysis of Cyclic Opioid Peptides with Dmt-Tic Pharmacophore. Molecules, 2020, 25, 4260.	1.7	2
9	Crystal Growth, Single Crystal Structure, and Biological Activity of Thiazolo-Pyridine Dicarboxylic Acid Derivatives. ACS Omega, 2020, 5, 27756-27765.	1.6	5
10	Phenylselanyl Group Incorporation for "Glutathione Peroxidase-Like―Activity Modulation. Molecules, 2020, 25, 3354.	1.7	11
11	Synthesis, Selected Transformations, and Biological Activity of Alkoxy Analogues of Lepidilines A and C. Materials, 2020, 13, 4190.	1.3	6
12	The search for opioid analgesics with limited tolerance liability. Peptides, 2020, 130, 170331.	1.2	11
13	Pharmacological Profile and Molecular Modeling of Cyclic Opioid Analogues Incorporating Various Phenylalanine Derivatives. ChemMedChem, 2020, 15, 1322-1329.	1.6	6
14	New uracil analog U-332 is an inhibitor of NF-κB in 5-fluorouracil-resistant human leukemia HL-60 cell line. BMC Pharmacology & Toxicology, 2020, 21, 18.	1.0	4
15	Seleninic Acid Potassium Salts as Water-Soluble Biocatalysts with Enhanced Bioavailability. Materials, 2020, 13, 661.	1.3	10
16	Molecular mechanisms of apoptosis induced by a novel synthetic quinolinone derivative in HL-60 human leukemia cells. Chemico-Biological Interactions, 2020, 320, 109005.	1.7	9
17	A New Hybrid δ-Lactone Induces Apoptosis and Potentiates Anticancer Activity of Taxol in HL-60 Human Leukemia Cells. Molecules, 2020, 25, 1479.	1.7	4
18	Biased Agonism as an Emerging Strategy in the Search for Better Opioid Analgesics. Current Medicinal Chemistry, 2020, 27, 1562-1575.	1.2	10

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19	New Uracil Analogs with Exocyclic Methylidene Group as Potential Anticancer Agents. Anti-Cancer Agents in Medicinal Chemistry, 2020, 20, 359-368.	0.9	3
20	Synthesis of 2,2,6-Trisubstituted 5-Methylidene-tetrahydropyran-4-ones with Anticancer Activity. Molecules, 2020, 25, 611.	1.7	2
21	Endomorphin-2 analogs containing modified tyrosines: Biological and theoretical investigation of the influence on conformation and pharmacological profile. European Journal of Medicinal Chemistry, 2019, 179, 527-536.	2.6	11
22	Bioselectivity Induced by Chirality of New Terpenyl Organoselenium Compounds. Materials, 2019, 12, 3579.	1.3	17
23	Synthesis and Cytotoxic Evaluation of 3-Methylidenechroman-4-ones. Molecules, 2019, 24, 1868.	1.7	4
24	N-Terpenyl Benzisoselenazolones—Evaluation of the Particular Structure-Bioactivity Relationship. Proceedings (mdpi), 2019, 41, 22.	0.2	0
25	New uracil analogs as downregulators of ABC transporters in 5-fluorouracil-resistant human leukemia HL-60 cell line. Molecular Biology Reports, 2019, 46, 5831-5839.	1.0	7
26	Synthesis and Pharmacological Evaluation of Hybrids Targeting Opioid and Neurokinin Receptors. Molecules, 2019, 24, 4460.	1.7	7
27	Helenalin - A Sesquiterpene Lactone with Multidirectional Activity. Current Drug Targets, 2019, 20, 444-452.	1.0	29
28	Antinociceptive potency of a fluorinated cyclopeptide Dmt-c[D-Lys-Phe- <i>p</i> -CF <sub>3</sub> -Phe-Asp]NH <sub>2</sub> . Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 560-566.	2.5	8
29	Involvement of α-methylene-γ- and δ-lactones in the suppression of multidrug resistance in MCF-7 cells. Pharmacological Reports, 2018, 70, 631-638.	1.5	1
30	In vitro and in vivo activity of cyclopeptide Dmt-c[ d -Lys-Phe-Asp]NH 2 , a mu opioid receptor agonist biased toward β-arrestin. Peptides, 2018, 105, 51-57.	1.2	10
31	Cyclopeptide Dmt-[D-Lys-p-CF3-Phe-Phe-Asp]NH2, a novel G protein-biased agonist of the mu opioid receptor. Peptides, 2018, 101, 227-233.	1.2	12
32	Synthesis of 3â€Methylideneâ€1â€tosylâ€2,3â€dihydroquinolinâ€4(1 <i>H</i> )â€ones as Potent Cytotoxic Agen Chemistry and Biodiversity, 2018, 15, e1800242.	ts <sub>1.0</sub>	9
33	Involvement of a coumarin analog AD-013 in the DNA damage response pathways in MCF-7 cells. Molecular Biology Reports, 2018, 45, 1187-1195.	1.0	2
34	Rubiscolins - Highly Potent Peptides Derived from Plant Proteins. Mini-Reviews in Medicinal Chemistry, 2018, 18, 104-112.	1.1	12
35	Anticancer Properties of a New Hybrid Analog AD-013 Combining a Coumarin Scaffold with an α-methylene-δ-lactone Motif. Anti-Cancer Agents in Medicinal Chemistry, 2018, 18, 450-457.	0.9	4
36	Drug resistance in topoisomerase-targeting therapy. Postepy Higieny I Medycyny Doswiadczalnej, 2018, 72, 1073-1083.	0.1	3

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37	Cyclic mu-opioid receptor ligands containing multiple N-methylated amino acid residues. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 1644-1648.	1.0	6
38	Synthesis of 4,4â€Disubstituted 3â€Methylidenechromanâ€2â€ones as Potent Anticancer Agents. ChemMedChem, 2017, 12, 599-605.	1.6	6
39	Design and characterization of opioid ligands based on cycle-in-macrocycle scaffold. Bioorganic and Medicinal Chemistry, 2017, 25, 2399-2405.	1.4	4
40	Evaluation of anticancer properties of a new α-methylene-δ-lactone DL-249 on two cancer cell lines. Open Life Sciences, 2017, 12, 178-189.	0.6	4
41	Synthesis and activity of opioid peptidomimetics with $\hat{I}^2$ 2 - and $\hat{I}^2$ 3 -amino acids. Peptides, 2017, 95, 116-123.	1.2	5
42	Effects of centrally administered endocannabinoids and opioids on orofacial pain perception in rats. British Journal of Pharmacology, 2017, 174, 3780-3789.	2.7	15
43	Synthesis, receptor binding studies, optical spectroscopic and <i>in silico</i> structural characterization of morphiceptin analogs with <i>cis</i> â€4â€amino‣â€proline residues. Journal of Peptide Science, 2017, 23, 864-870.	0.8	2
44	New glutathione peroxidase mimetics—Insights into antioxidant and cytotoxic activity. Bioorganic and Medicinal Chemistry, 2017, 25, 126-131.	1.4	41
45	New Chiral Ebselen Analogues with Antioxidant and Cytotoxic Potential. Molecules, 2017, 22, 492.	1.7	37
46	Anticancer activity of new molecular hybrids combining 1,4-naphthalenedione motif with phosphonic acid moiety in hepatocellular carcinoma HepG2 cells Acta Biochimica Polonica, 2017, 64, 41-48.	0.3	4
47	Novobiocin Analogs as Potential Anticancer Agents. Mini-Reviews in Medicinal Chemistry, 2017, 17, 728-733.	1.1	21
48	Redoubling the ring size of an endomorphinâ€2 analog transforms a centrally acting muâ€opioid receptor agonist into a pure peripheral analgesic. Biopolymers, 2016, 106, 309-317.	1.2	6
49	Anticancer properties of new synthetic hybrid molecules combining naphtho[2,3-b]furan-4,9-dione or benzo[f]indole-4,9-dione motif with phosphonate subunit. European Journal of Medicinal Chemistry, 2016, 120, 51-63.	2.6	28
50	Synthesis of mixed MOR/KOR efficacy cyclic opioid peptide analogs with antinociceptive activity after systemic administration. European Journal of Medicinal Chemistry, 2016, 109, 276-286.	2.6	29
51	Anticancer activity and radiosensitization effect of methyleneisoxazolidin-5-ones in hepatocellular carcinoma HepG2 cells. Chemico-Biological Interactions, 2016, 248, 68-73.	1.7	3
52	Synthesis, biological evaluation and structural analysis of novel peripherally active morphiceptin analogs. Bioorganic and Medicinal Chemistry, 2016, 24, 1582-1588.	1.4	7
53	Opioid and Cannabinoid System in Food Intake. Current Pharmaceutical Design, 2016, 22, 1361-1370.	0.9	12
54	ABC Transporters in the Development of Multidrug Resistance in Cancer Therapy. Current Pharmaceutical Design, 2016, 22, 4705-4716.	0.9	44

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55	Anticancer Properties of Novel 4â€methyleneâ€1,2â€diphenylpyrazolidinâ€3â€ones. Chemical Biology and Drug Design, 2015, 86, 961-968.	1.5	1
56	Synthesis of linear and cyclic opioidâ€based peptide analogs containing multiple <i>N</i> â€methylated amino acid residues. Journal of Peptide Science, 2015, 21, 807-810.	0.8	8
57	The role of oxidative stress in anticancer activity of sesquiterpene lactones. Naunyn-Schmiedeberg's Archives of Pharmacology, 2015, 388, 477-486.	1.4	84
58	Design, synthesis and cytotoxic evaluation of 4-methylidenepyrazolidin-3-ones. European Journal of Medicinal Chemistry, 2015, 92, 565-574.	2.6	13
59	Combined effects of anticancer drugs and new synthetic α-methylene-Î-lactones on MCF-7 cells. Tumor Biology, 2015, 36, 5971-5977.	0.8	5
60	Synthesis of Mixed Opioid Affinity Cyclic Endomorphin-2 Analogues with Fluorinated Phenylalanines. ACS Medicinal Chemistry Letters, 2015, 6, 579-583.	1.3	15
61	Ring size in cyclic endomorphin-2 analogs modulates receptor binding affinity and selectivity. Organic and Biomolecular Chemistry, 2015, 13, 6039-6046.	1.5	16
62	Novel synthesis and cytotoxic activity of 1,4-disubstituted 3-methylidene-3,4-dihydroquinolin-2(1H)-ones. RSC Advances, 2015, 5, 78324-78335.	1.7	7
63	Immunomodulatory Effects of Endogenous and Synthetic Peptides Activating Opioid Receptors. Mini-Reviews in Medicinal Chemistry, 2015, 14, 1148-1155.	1.1	22
64	Trigemino-hypoglossal somatic reflex in the pharmacological studies of nociception in orofacial area. Acta Neurobiologiae Experimentalis, 2015, 75, 253-63.	0.4	2
65	Cyclic endomorphin analogs in targeting opioid receptors to achieve pain relief. Future Medicinal Chemistry, 2014, 6, 2093-2101.	1.1	17
66	Proteomic Analysis of Proteins Engaged in <i>α</i> â€Methyleneâ€ <i>δ</i> â€Lactone Cytotoxic Effects in Hormoneâ€Independent Breast Cancer <scp>MDA</scp> â€ <scp>MB</scp> â€231 Cells. Chemical Biology and Drug Design, 2014, 84, 300-306.	1.5	7
67	Cyclic pentapeptide analogs based on endomorphin-2 structure: Cyclization studies using liquid chromatography combined with on-line mass spectrometry and tandem mass spectrometry. Peptides, 2014, 55, 32-40.	1.2	6
68	Pharmacological characterization of endomorphin-2-based cyclic pentapeptides with methylated phenylalanine residues. Peptides, 2014, 55, 145-150.	1.2	22
69	Antinociceptive and antidepressant-like action of endomorphin-2 analogs with proline surrogates in position 2. Bioorganic and Medicinal Chemistry, 2014, 22, 4803-4809.	1.4	13
70	Cyclic side-chain-linked opioid analogs utilizing cis - and trans -4-aminocyclohexyl- d -alanine. Bioorganic and Medicinal Chemistry, 2014, 22, 6545-6551.	1.4	13
71	Endless Peptides - Circular Forms in Nature. Current Medicinal Chemistry, 2014, 22, 352-359.	1.2	4
72	Combined Effect of Parthenolide and Various Anti-cancer Drugs or Anticancer Candidate Substances on Malignant Cells in vitro and in vivo. Mini-Reviews in Medicinal Chemistry, 2014, 14, 222-228.	1.1	16

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73	Pharmacological Properties of Novel Cyclic Pentapeptides with µ-opioid Receptor Agonist Activity. Medicinal Chemistry, 2014, 10, 154-161.	0.7	9
74	α-Methylene-γ-lactones as a Novel Class of Anti-leukemic Agents. Anti-Cancer Agents in Medicinal Chemistry, 2014, 14, 688-694.	0.9	23
75	Novel glycosylated endomorphin-2 analog produces potent centrally-mediated antinociception in mice after peripheral administration. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 6673-6676.	1.0	13
76	Anticancer Activity of New Synthetic αâ€Methyleneâ€Ì ã€Łactones on Two Breast Cancer Cell Lines. Basic and Clinical Pharmacology and Toxicology, 2013, 113, 391-400.	1.2	17
77	First one-pot organocatalytic synthesis of α-methylene-γ-lactones. Chemical Communications, 2013, 49, 1184.	2.2	45
78	Apoptosis-mediated cytotoxic effects of parthenolide and the new synthetic analog MZ-6 on two breast cancer cell lines. Molecular Biology Reports, 2013, 40, 1655-1663.	1.0	26
79	Bioavailability of Endomorphins and the Blood-brain Barrier- A Review. Medicinal Chemistry, 2013, 10, 2-17.	0.7	15
80	Cyclization in Opioid Peptides. Current Drug Targets, 2013, 14, 798-816.	1.0	26
81	Synthesis and biological evaluation of α-methylidene-δ-lactones with 3,4-dihydrocoumarin skeleton. Bioorganic and Medicinal Chemistry, 2012, 20, 5017-5026.	1.4	43
82	Structural comparison of endomorphin-2 and its conformationally restricted analog. Open Chemistry, 2012, 10, 172-179.	1.0	0
83	Kinetic studies of novel inhibitors of endomorphin degrading enzymes. Medicinal Chemistry Research, 2012, 21, 1445-1450.	1.1	8
84	Comparison of Antiâ€Invasive Activity of Parthenolide and 3â€Isopropylâ€2â€Methylâ€4â€Methyleneisoxazolidinâ€5â€One (MZâ€6) – A New Compound with αâ€Metł Motif – on Two Breast Cancer Cell Lines. Chemical Biology and Drug Design, 2012, 79, 112-120.	ıyl <b>a</b> næâ€Ĵ	â€ <b>L9</b> ctone
85	Natural and synthetic α-methylenelactones and α-methylenelactams with anticancer potential. Drug Discovery Today, 2012, 17, 561-572.	3.2	136
86	Opioid-regulated pro- and anti-apoptotic gene expression in cancer cells. Open Life Sciences, 2012, 7, 411-418.	0.6	1
87	Effect of potent endomorphin degradation blockers on analgesic and antidepressant-like responses in mice. Neuropharmacology, 2011, 61, 1229-1238.	2.0	10
88	Effect of 2′,6′-dimethyl-l-tyrosine (Dmt) on pharmacological activity of cyclic endomorphin-2 and morphiceptin analogs. Bioorganic and Medicinal Chemistry, 2011, 19, 6977-6981.	1.4	26
89	Effect of tooth pulp and periaqueductal central gray stimulation on the expression of genes encoding the selected neuropeptides and opioid receptors in the mesencephalon, hypothalamus and thalamus in rats. Brain Research, 2011, 1382, 19-28.	1.1	14
90	Effect of tooth pulp and periaqueductal central gray electrical stimulation on β-endorphin release into the fluid perfusing the cerebral ventricles in rats. Brain Research, 2011, 1405, 15-22.	1.1	9

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91	Opioid-receptor gene expression and localization in cancer cells. Open Life Sciences, 2011, 6, 10-15.	0.6	7
92	The influence of opioids on matrix metalloproteinase-2 and -9 secretion and mRNA levels in MCF-7 breast cancer cell line. Molecular Biology Reports, 2011, 38, 1231-1236.	1.0	48
93	The role of morphine in regulation of cancer cell growth. Naunyn-Schmiedeberg's Archives of Pharmacology, 2011, 384, 221-230.	1.4	114
94	Synthesis and cytotoxic evaluation of β-alkyl or β-aryl-Ĩ´-methyl-α-methylene-Ĩ´-lactones. Comparison with the corresponding γ-lactones. European Journal of Medicinal Chemistry, 2010, 45, 710-718.	2.6	27
95	The Novel Endomorphin Degradation Blockers Tyrâ€Proâ€ <scp>D</scp> ClPheâ€Pheâ€NH <sub>2</sub> (EMDBâ€ and Tyrâ€Proâ€Alaâ€NH <sub>2</sub> (EMDBâ€2) Prolong Endomorphinâ€2 Action in Rat lleum <i>In Vitro</i> . Chemical Biology and Drug Design, 2010, 76, 77-81.	€1) 1.5	6
96	Design, Synthesis and Pharmacological Characterization of Endomorphin Analogues with Non yclic Amino Acid Residues in Position 2. Basic and Clinical Pharmacology and Toxicology, 2010, 106, 106-113.	1.2	17
97	Synthesis and biological evaluation of cyclic endomorphin-2 analogs. Peptides, 2010, 31, 339-345.	1.2	38
98	Synthesis and biological evaluation of novel peripherally active morphiceptin analogs. Peptides, 2010, 31, 1617-1624.	1.2	20
99	A convenient synthesis and cytotoxic evaluation of β-aryl-α-methylidene-γ-lactones and β-aryl-α-methylidene-γ-lactams. New Journal of Chemistry, 2010, 34, 750.	1.4	24
100	The Influence of Opioids on Urokinase Plasminogen Activator on Protein and mRNA Level in MCFâ€7 Breast Cancer Cell Line. Chemical Biology and Drug Design, 2009, 74, 390-396.	1.5	35
101	Biological activity of endomorphin and [Dmt1]endomorphin analogs with six-membered proline surrogates in position 2. Bioorganic and Medicinal Chemistry, 2009, 17, 3789-3794.	1.4	29
102	Synthesis and cytotoxic activity of γ-aryl substituted α-alkylidene-γ-lactones and α-alkylidene-γ-lactams. Bioorganic and Medicinal Chemistry, 2008, 16, 4872-4882.	1.4	51
103	Novel highly potent μ-opioid receptor antagonist based on endomorphin-2 structure. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 1350-1353.	1.0	11
104	Synthesis and Biological Activity of Endomorphinâ€2 Analogs Incorporating Piperidineâ€2â€, 3―or 4â€Carboxylic Acids Instead of Proline in Position 2. Chemical Biology and Drug Design, 2008, 72, 91-94.	1.5	23
105	[Dmt1, d-1-Nal3]morphiceptin, a novel opioid peptide analog with high analgesic activity. Peptides, 2008, 29, 633-638.	1.2	11
106	Enzymatic degradation of endomorphins. Peptides, 2008, 29, 2066-2073.	1.2	84
107	Opioid-induced regulation of µ-opioid receptor gene expression in the MCF-7 breast cancer cell line. Biochemistry and Cell Biology, 2008, 86, 217-226.	0.9	27
108	Endomorphin Analogs. Current Medicinal Chemistry, 2007, 14, 3201-3208.	1.2	49

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109	Antidepressant-Like Effect of Endomorphin-1 and Endomorphin-2 in Mice. Neuropsychopharmacology, 2007, 32, 813-821.	2.8	50
110	The Endomorphin System and Its Evolving Neurophysiological Role. Pharmacological Reviews, 2007, 59, 88-123.	7.1	217
111	Synthesis and Characterization of Potent and Selective μ-Opioid Receptor Antagonists, [Dmt, d-2-Nal4]endomorphin-1 (Antanal-1) and [Dmt1, d-2-Nal4]endomorphin-2 (Antanal-2). Journal of Medicinal Chemistry, 2007, 50, 512-520.	2.9	40
112	Enzymatic degradation studies of endomorphin-2 and its analogs containing N-methylated amino acids. Peptides, 2006, 27, 131-135.	1.2	41
113	In vitro Characterization of Novel Peptide Inhibitors of Endomorphin-degrading Enzymes in the Rat Brain. Chemical Biology and Drug Design, 2006, 68, 173-175.	1.5	16
114	4-Methylideneisoxazolidin-5-ones—A new class of α-methylidene-γ-lactones with high cytostatic activity. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 1430-1433.	1.0	26
115	Synthesis and biological activity of N-methylated analogs of endomorphin-2. Bioorganic and Medicinal Chemistry, 2005, 13, 6713-6717.	1.4	27
116	Synthesis and antinociceptive activity of cyclic endomorphin-2 and morphiceptin analogs. Biochemical Pharmacology, 2005, 71, 188-195.	2.0	25
117	Enantioselective Synthesis and Cytotoxic Evaluation of 4,5-Dihydro-5-[aryl(hydroxy)methyl]-3-methylidenefuran-2(3H)-ones. Chemistry and Biodiversity, 2005, 2, 1256-1265.	1.0	10
118	Conformationally Restricted Peptides as Tools in Opioid Receptor Studies. Current Medicinal Chemistry, 2005, 12, 471-481.	1.2	49
119	Novel Synthesis, Cytotoxic Evaluation, and Structureâ <sup>~,</sup> Activity Relationship Studies of a Series of α-Alkylidene-Î <sup>3</sup> -lactones and Lactams. Journal of Medicinal Chemistry, 2005, 48, 3516-3521.	2.9	104
120	Comparison of antagonist activity of spantide family at human neurokinin receptors measured by aequorin luminescence-based functional calcium assay. Regulatory Peptides, 2005, 131, 23-28.	1.9	15
121	Opioid Receptors and their Ligands. Current Topics in Medicinal Chemistry, 2004, 4, 1-17.	1.0	240
122	Opioid peptides in cancer. Cancer and Metastasis Reviews, 2004, 23, 351-366.	2.7	43
123	Opioid receptor binding and in vivo antinociceptive activity of position 3-substituted morphiceptin analogs. Biochemical and Biophysical Research Communications, 2004, 320, 531-536.	1.0	35
124	Binding of the new morphiceptin analogs to human MCF-7 breast cancer cells and their effect on growth. Regulatory Peptides, 2004, 120, 237-241.	1.9	9
125	Structure-activity Relationship, Conformation and Pharmacology Studies of Morphiceptin Analogues - Selective μ-Opioid Receptor Ligands. Mini-Reviews in Medicinal Chemistry, 2002, 2, 565-572.	1.1	13
126	New Stereocontrolled Synthesis and Biological Evaluation of 5-(1â€~-Hydroxyalkyl)-3-methylidenetetrahydro-2-furanones as Potential Cytotoxic Agents. Journal of Medicinal Chemistry, 2002, 45, 1142-1145.	2.9	30

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127	Substance P content in the cerebrospinal fluid and fluid perfusing cerebral ventricles during elicitation and inhibition of trigemino-hypoglossal reflex in rats. Brain Research, 2002, 941, 29-33.	1.1	8
128	Effect of cerebral ventricles perfusion with naloxone on trigemino-hypoglossal reflex in rats. Regulatory Peptides, 2001, 97, 7-13.	1.9	7
129	Inhibition of tongue reflex in rats by tooth pulp stimulation during cerebral ventricle perfusion with (6–11) substance P analogs. Brain Research, 1997, 753, 128-132.	1.1	13