

Carlos Fraga

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

211
papers

5,962
citations

36
h-index

70
g-index

244
ext. papers

6,749
ext. citations

3.8
avg, IF

5.64
L-index

#	Paper	IF	Citations
211	Novel Single Inhibitor of HDAC6/8 and Dual Inhibitor of PI3K/HDAC6 as Potential Alternative Treatments for Prostate Cancer. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	1
210	Multitarget Inhibition of Histone Deacetylase (HDAC) and Phosphatidylinositol-3-kinase (PI3K): Current and Future Prospects. <i>ChemMedChem</i> , 2021 , 16, 448-457	3.7	3
209	Structure–property relationship studies of 3-acyl-substituted furans: the serendipitous identification and characterization of a new non-classical hydrogen bond donor moiety. <i>New Journal of Chemistry</i> , 2020 , 44, 10994-11005	3.6	3
208	Therapeutic Effects of Anti-Inflammatory -Acylhydrazones in the Resolution of Experimental Colitis. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2020 , 374, 420-427	4.7	1
207	Histone deacetylases as targets for the treatment of neurodegenerative disorders: Challenges and future opportunities. <i>Medicinal Research Reviews</i> , 2020 , 40, 2177-2211	14.4	16
206	New 2-amino-pyridinyl-N-acylhydrazones: Synthesis and identification of their mechanism of anti-inflammatory action. <i>Biomedicine and Pharmacotherapy</i> , 2020 , 123, 109739	7.5	3
205	Identification of Novel Functionalized Carbohydrazonamides Designed as Chagas Disease Drug Candidates. <i>Medicinal Chemistry</i> , 2020 , 16, 774-783	1.8	0
204	Investigating the Molecular Basis for the Selective Inhibition of Aldehyde Dehydrogenase 2 by the Isoflavonoid Daidzin. <i>CNS and Neurological Disorders - Drug Targets</i> , 2020 , 19, 437-447	2.6	0
203	Synthesis and trypanocidal activity of novel pyridinyl-1,3,4-thiadiazole derivatives. <i>Biomedicine and Pharmacotherapy</i> , 2020 , 127, 110162	7.5	7
202	Design, Synthesis, and Pharmacological Evaluation of First-in-Class Multitarget N-Acylhydrazone Derivatives as Selective HDAC6/8 and PI3K Inhibitors. <i>ChemMedChem</i> , 2020 , 15, 539-551	3.7	17
201	Gastroprotective effects of N-acylarylhydrazone derivatives on ethanol-induced gastric lesions in mice are dependent on the NO/cGMP/K pathway. <i>Biochemical Pharmacology</i> , 2019 , 169, 113629	6	5
200	Meet Our Section Editor. <i>Current Topics in Medicinal Chemistry</i> , 2019 , 18, 2133-2135	3	
199	Duvelisib: A 2018 Novel FDA-Approved Small Molecule Inhibiting Phosphoinositide 3-Kinases. <i>Pharmaceuticals</i> , 2019 , 12,	5.2	32
198	Evaluation of Functional Selectivity of Haloperidol, Clozapine, and LASSBio-579, an Experimental Compound With Antipsychotic-Like Actions in Rodents, at G Protein and Arrestin Signaling Downstream of the Dopamine D Receptor. <i>Frontiers in Pharmacology</i> , 2019 , 10, 628	5.6	1
197	The Use of Conformational Restriction in Medicinal Chemistry. <i>Current Topics in Medicinal Chemistry</i> , 2019 , 19, 1712-1733	3	9
196	New analogs of LASSBio-1829Cl with anti-inflammatory properties. <i>FASEB Journal</i> , 2019 , 33, lb41	0.9	
195	Design, Synthesis and Pharmacological Evaluation of Novel Antiinflammatory and Analgesic O-Benzoyloxime Compounds Derived From Natural Eugenol. <i>Letters in Drug Design and Discovery</i> , 2019 , 16, 1157-1166	0.8	

194	The Chalcone Lonchocarpin Inhibits Wnt/ β -Catenin Signaling and Suppresses Colorectal Cancer Proliferation. <i>Cancers</i> , 2019 , 11,	6.6	13
193	Structural basis for the agonist action at free fatty acid receptor 1 (FFA1R or GPR40). <i>Chemical Biology and Drug Design</i> , 2018 , 91, 668-680	2.9	9
192	Synthesis and pharmacological evaluation of novel isoquinoline N-sulphonylhydrazones designed as ROCK inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1181-1193	5.6	3
191	N-Acylhydrazones as drugs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 2797-2806	2.9	67
190	The novel piperazine-containing compound LQFM018: Necroptosis cell death mechanisms, dopamine D receptor binding and toxicological assessment. <i>Biomedicine and Pharmacotherapy</i> , 2018 , 102, 481-493	7.5	8
189	Eine ungewöhnliche intramolekulare Halogenbindung führt zu konformationeller Selektion. <i>Angewandte Chemie</i> , 2018 , 130, 10120-10126	3.6	
188	An Unusual Intramolecular Halogen Bond Guides Conformational Selection. <i>Angewandte Chemie - International Edition</i> , 2018 , 57, 9970-9975	16.4	9
187	Synergistic interaction between a PDE5 inhibitor (sildenafil) and a new adenosine A2A receptor agonist (LASSBio-1359) improves pulmonary hypertension in rats. <i>PLoS ONE</i> , 2018 , 13, e0195047	3.7	7
186	NF- κ B-IKK β Pathway as a Target for Drug Development: Realities, Challenges and Perspectives. <i>Current Drug Targets</i> , 2018 , 19, 1933-1942	3	26
185	Different molecular conformations in the crystal structures of three 5-nitro-imidazolyl derivatives. <i>Acta Crystallographica Section E: Crystallographic Communications</i> , 2018 , 74, 380-384	0.7	
184	Theoretical and experimental characterization of 1,4-N π S π Hole intramolecular interactions in bioactive N-acylhydrazone derivatives. <i>New Journal of Chemistry</i> , 2018 , 42, 497-505	3.6	13
183	Discovery of naphthyl-N-acylhydrazone p38 β MAPK inhibitors with in vivo anti-inflammatory and anti-TNF α activity. <i>Chemical Biology and Drug Design</i> , 2018 , 91, 391-397	2.9	15
182	Design, Synthesis, Experimental and Theoretical Characterization of a New Multitarget 2-Thienyl-Acylhydrazone Derivative. <i>Pharmaceuticals</i> , 2018 , 11,	5.2	3
181	Modeling zinc-oxygen coordination in histone deacetylase: A comparison of semiempirical methods performance. <i>International Journal of Quantum Chemistry</i> , 2018 , 118, e25720	2.1	3
180	Cardioprotection Induced by Activation of GPER in Ovariectomized Rats With Pulmonary Hypertension. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2018 , 73, 1158-1166	6.4	10
179	LASSBio-1422: a new molecular scaffold with efficacy in animal models of schizophrenia and disorders of attention and cognition. <i>Behavioural Pharmacology</i> , 2017 , 28, 48-62	2.4	7
178	ROCK inhibition with Fasudil induces beta-catenin nuclear translocation and inhibits cell migration of MDA-MB 231 human breast cancer cells. <i>Scientific Reports</i> , 2017 , 7, 13723	4.9	21
177	Adenosine A receptor agonist prevents cardiac remodeling and dysfunction in spontaneously hypertensive male rats after myocardial infarction. <i>Drug Design, Development and Therapy</i> , 2017 , 11, 553-562	4.4	20

176	Design, Synthesis, and Trypanocidal Activity of Novel 5-Nitroimidazolyl O-Benzylloxime Ethers. <i>Journal of Heterocyclic Chemistry</i> , 2017 , 54, 3626-3631	1.9	3
175	LASSBio-897 Reduces Lung Injury Induced by Silica Particles in Mice: Potential Interaction with the A Receptor. <i>Frontiers in Pharmacology</i> , 2017 , 8, 778	5.6	3
174	LASSBio-579, a prototype antipsychotic drug, and clozapine are effective in novel object recognition task, a recognition memory model. <i>Behavioural Pharmacology</i> , 2016 , 27, 339-49	2.4	6
173	Filtering promiscuous compounds in early drug discovery: is it a good idea?. <i>Drug Discovery Today</i> , 2016 , 21, 868-72	8.8	54
172	Design, Synthesis, and Pharmacological Evaluation of Novel N-Acylhydrazone Derivatives as Potent Histone Deacetylase 6/8 Dual Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 655-70	8.3	61
171	LASSBio-1425, an analog of thalidomide, decreases triglyceride and increases HDL cholesterol levels by inhibition of TNF- α production. <i>International Journal of Cardiology</i> , 2016 , 202, 497-9	3.2	7
170	Discovery of Novel Orally Active Tetrahydro-Naphthyl-N-Acylhydrazones with In Vivo Anti-TNF- α Effect and Remarkable Anti-Inflammatory Properties. <i>PLoS ONE</i> , 2016 , 11, e0156271	3.7	14
169	Beyond the Selective Inhibition of Histone Deacetylase 6. <i>Mini-Reviews in Medicinal Chemistry</i> , 2016 , 16, 1175-84	3.2	12
168	Understanding the Structural Basis of ALDH-2 Inhibition by Molecular Docking. <i>Medicinal Chemistry</i> , 2016 , 12, 506-12	1.8	0
167	LASSBio-1829 Hydrochloride: Development of a New Orally Active N-Acylhydrazone IKK2 Inhibitor with Anti-inflammatory Properties. <i>ChemMedChem</i> , 2016 , 11, 234-44	3.7	7
166	Treatment with Adenosine Receptor Agonist Ameliorates Pain Induced by Acute and Chronic Inflammation. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2016 , 358, 315-23	4.7	16
165	Novel agonist of adenosine receptor induces relaxation of corpus cavernosum in guinea pigs: an in vitro and in vivo study. <i>Urology</i> , 2015 , 85, 1214.e17-1214.e21	1.6	4
164	Partial agonism and fast dissociation of LASSBio-579 at dopamine D2 receptor. <i>Progress in Neuro-Psychopharmacology and Biological Psychiatry</i> , 2015 , 62, 1-6	5.5	3
163	In vivo effect of LASSBio-785, a lipid-lowering and anti-inflammatory agent, on cardiac Ca(2+)-ATPases from hypercholesterolemic rats. <i>International Journal of Cardiology</i> , 2015 , 201, 282-4	3.2	2
162	Structural characterization of LASSBio-1289: a new vasoactive N-methyl-N-acylhydrazone derivative. <i>CrystEngComm</i> , 2015 , 17, 165-173	3.3	10
161	Synthesis and Biological Evaluation of Pyrazolo[3,4-b]pyridin-4-ones as a New Class of Topoisomerase II Inhibitors. <i>Medicinal Chemistry</i> , 2015 , 11, 342-53	1.8	6
160	BUBONIC PLAGUE: HISTORICAL ASPECTS AND THERAPY. <i>Military Medical Science Letters (Vojenske Zdravotnicke Listy)</i> , 2015 , 84, 67-75	0.2	1
159	Acylhydrazone derivatives: a patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2014 , 24, 1161-70	6.8	36

158	N-acylhydrazone derivative ameliorates monocrotaline-induced pulmonary hypertension through the modulation of adenosine AA2R activity. <i>International Journal of Cardiology</i> , 2014 , 173, 154-62	3.2	28
157	N-acylhydrazone improves exercise intolerance in rats submitted to myocardial infarction by the recovery of calcium homeostasis in skeletal muscle. <i>Life Sciences</i> , 2014 , 94, 30-6	6.8	6
156	Antiprotozoal activity of (E)-cinnamic N-acylhydrazone derivatives. <i>Molecules</i> , 2014 , 19, 20374-81	4.8	9
155	Novel potent imidazo[1,2-a]pyridine-N-Glyciny-hydrazone inhibitors of TNF- α production: in vitro and in vivo studies. <i>PLoS ONE</i> , 2014 , 9, e91660	3.7	11
154	Crystal structures of 1-hydroxyimidazole and imidazole 1-oxide derivatives. <i>Zeitschrift Fur Kristallographie - Crystalline Materials</i> , 2014 , 229,	1	3
153	Vasodilator and antihypertensive effects of a novel N-acylhydrazone derivative mediated by the inhibition of L-type Ca ²⁺ channels. <i>Fundamental and Clinical Pharmacology</i> , 2014 , 28, 29-41	3.1	6
152	LASSBio-1135: a dual TRPV1 antagonist and anti-TNF-alpha compound orally effective in models of inflammatory and neuropathic pain. <i>PLoS ONE</i> , 2014 , 9, e99510	3.7	11
151	Biotransformation of LASSBio-579 and pharmacological evaluation of p-hydroxylated metabolite a N-phenylpiperazine antipsychotic lead compound. <i>European Journal of Medicinal Chemistry</i> , 2013 , 62, 214-21	6.8	10
150	A novel Ca ²⁺ channel antagonist reverses cardiac hypertrophy and pulmonary arteriolar remodeling in experimental pulmonary hypertension. <i>European Journal of Pharmacology</i> , 2013 , 702, 316-22	5.3	13
149	P.3.c.001 Pharmacological evaluation of new N-phenylpiperazine derivatives designed as homologues of the antipsychotic lead compound LASSBio-579. <i>European Neuropsychopharmacology</i> , 2013 , 23, S453	1.2	
148	Synthesis and pharmacological evaluation of new N-phenylpiperazine derivatives designed as homologues of the antipsychotic lead compound LASSBio-579. <i>European Journal of Medicinal Chemistry</i> , 2013 , 66, 122-34	6.8	23
147	Beneficial effects of a novel agonist of the adenosine A2A receptor on monocrotaline-induced pulmonary hypertension in rats. <i>British Journal of Pharmacology</i> , 2013 , 169, 953-62	8.6	30
146	New insights into pharmacological profile of LASSBio-579, a multi-target N-phenylpiperazine derivative active on animal models of schizophrenia. <i>Behavioural Brain Research</i> , 2013 , 237, 86-95	3.4	22
145	Antihyperalgesic effects of a novel muscarinic agonist (LASSBio-873) in spinal nerve ligation in rats. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2013 , 40, 404-11	3	3
144	Anti-atherogenic effects of a new thienylacylhydrazone derivative, LASSBio-788, in rats fed a hypercholesterolemic diet. <i>Journal of Pharmacological Sciences</i> , 2013 , 123, 47-57	3.7	11
143	Characterization of amide bond conformers for a novel heterocyclic template of N-acylhydrazone derivatives. <i>Molecules</i> , 2013 , 18, 11683-704	4.8	58
142	Synthesis and trypanocidal activity of novel 2,4,5-triaryl-N-hydroxyimidazole derivatives. <i>Molecules</i> , 2013 , 18, 3445-57	4.8	17
141	Impairment of locomotor activity induced by the novel N-acylhydrazone derivatives LASSBio-785 and LASSBio-786 in mice. <i>Brazilian Journal of Medical and Biological Research</i> , 2013 , 46, 263-9	2.8	2

140	Novel furfurylidene N-acylhydrazones derived from natural safrole: discovery of LASSBio-1215, a new potent antiplatelet prototype. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 101-9	5.6	6
139	Phenylpiperazine derivatives: a patent review (2006 - present). <i>Expert Opinion on Therapeutic Patents</i> , 2012 , 22, 1169-78	6.8	14
138	Design, synthesis, and pharmacological evaluation of N-acylhydrazones and novel conformationally constrained compounds as selective and potent orally active phosphodiesterase-4 inhibitors. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 7525-45	8.3	82
137	Design and synthesis of new (E)-cinnamic N-acylhydrazones as potent antitrypanosomal agents. <i>European Journal of Medicinal Chemistry</i> , 2012 , 54, 512-21	6.8	56
136	Antihypertensive profile of 2-thienyl-3,4-methylenedioxybenzoylhydrazone is mediated by activation of the A _{2A} adenosine receptor. <i>European Journal of Medicinal Chemistry</i> , 2012 , 55, 49-57	6.8	26
135	Discovery of novel orally active anti-inflammatory N-phenylpyrazolyl-N-glycinyldihydrazone derivatives that inhibit TNF- α production. <i>PLoS ONE</i> , 2012 , 7, e46925	3.7	18
134	Molecular docking and molecular dynamic studies of semi-synthetic piperidine alkaloids as acetylcholinesterase inhibitors. <i>Journal of the Brazilian Chemical Society</i> , 2012 , 23, 163-170	1.5	5
133	Synthesis and characterization of the atropisomeric relationships of a substituted N-phenyl-bipyrzole derivative with anti-inflammatory properties. <i>Chirality</i> , 2012 , 24, 463-70	2.1	1
132	Natural Products as Lead Compounds in Medicinal Chemistry 2012 , 81-126		1
131	Combination of docking, molecular dynamics and quantum mechanical calculations for metabolism prediction of 3,4-methylenedioxybenzoyl-2-thienylhydrazone. <i>Journal of Molecular Modeling</i> , 2012 , 18, 2065-78	2	17
130	(2E)-NR[(E)-Benzyl-idene]-3-phenyl-prop-2-enohydrazide from synchrotron radiation. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2012 , 68, o2255-6		2
129	(2E)-NR[(E)-2-Hydroxy-benzyl-idene]-3-phenyl-prop-2-enohydrazide. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2012 , 68, o2253-4		2
128	Molecular modeling studies of <i>Yersinia pestis</i> dihydrofolate reductase. <i>Journal of Biomolecular Structure and Dynamics</i> , 2011 , 29, 351-67	3.6	11
127	The methylation effect in medicinal chemistry. <i>Chemical Reviews</i> , 2011 , 111, 5215-46	68.1	485
126	Discovery of LASSBio-772, a 1,3-benzodioxole N-phenylpiperazine derivative with potent α 1A/D-adrenergic receptor blocking properties. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 3000-12	6.8	26
125	CYP1A2-mediated biotransformation of cardioactive 2-thienylidene-3,4-methylenedioxybenzoylhydrazone (LASSBio-294) by rat liver microsomes and human recombinant CYP enzymes. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 349-55	6.8	7
124	Structure-based design and biological profile of (E)-N-(4-Nitrobenzylidene)-2-naphthohydrazide, a novel small molecule inhibitor of IB kinase- β . <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 1245-53	6.8	16
123	Determination of the cardioactive prototype LASSBio-294 and its metabolites in dog plasma by LC-MS/MS: application for a pharmacokinetic study. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2011 , 55, 1024-30	3.5	5

122	MAOS and medicinal chemistry: some important examples from the last years. <i>Molecules</i> , 2011 , 16, 9274-9747	4.7	17
121	The role of natural products in the discovery of new drug candidates for the treatment of neurodegenerative disorders II: Alzheimer's disease. <i>CNS and Neurological Disorders - Drug Targets</i> , 2011 , 10, 251-70	2.6	67
120	The role of natural products in the discovery of new drug candidates for the treatment of neurodegenerative disorders I: Parkinson's disease. <i>CNS and Neurological Disorders - Drug Targets</i> , 2011 , 10, 239-50	2.6	29
119	LASSBio-881: an N-acylhydrazone transient receptor potential vanilloid subfamily type 1 antagonist orally effective against the hypernociception induced by capsaicin or partial sciatic ligation. <i>British Journal of Pharmacology</i> , 2010 , 159, 1716-23	8.6	10
118	(2E)-NR[(E)-4-Chloro-benzyl-idene]-3-phenyl-prop-2-enohydrazide monohydrate. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2010 , 66, o2410-1		4
117	Pharmacological characterization of (3-thienylidene)-3,4-methylenedioxybenzoylhydrazide: a novel muscarinic agonist with antihypertensive profile. <i>American Journal of Hypertension</i> , 2010 , 23, 135-41	2.3	16
116	LASSBio-294, A compound with inotropic and lusitropic activity, decreases cardiac remodeling and improves Ca ²⁺ influx into sarcoplasmic reticulum after myocardial infarction. <i>American Journal of Hypertension</i> , 2010 , 23, 1220-7	2.3	15
115	Cardiovascular effects induced by N-(4Rdihydro)-piperoylthiomorpholine in normotensive rats. <i>Journal of Pharmacy and Pharmacology</i> , 2010 , 62, 1794-800	4.8	
114	In vitro and in vivo activities of 1,3,4-thiadiazole-2-arylhydrazone derivatives of megazol against <i>Trypanosoma cruzi</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2010 , 54, 2023-31	5.9	30
113	Discovery of Dual Chemotherapy Drug Candidates Designed by Molecular Hybridization. <i>Current Enzyme Inhibition</i> , 2010 , 6, 171-182	0.5	13
112	Searching for multi-target antipsychotics: Discovery of orally active heterocyclic N-phenylpiperazine ligands of D2-like and 5-HT1A receptors. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 1925-35	3.4	50
111	Novel thienylacylhydrazone derivatives inhibit platelet aggregation through cyclic nucleotides modulation and thromboxane A2 synthesis inhibition. <i>European Journal of Pharmacology</i> , 2010 , 638, 5-12	5.3	21
110	Characterization of the conformational ensemble from bioactive N-acylhydrazone derivatives. <i>Journal of Molecular Graphics and Modelling</i> , 2010 , 28, 446-54	2.8	10
109	Design of new dopamine D2 receptor ligands: biosynthesis and pharmacological evaluation of the hydroxylated metabolite of LASSBio-581. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 2888-91	2.9	6
108	Microwave-assisted synthesis and structure-activity relationships of neuroactive pyrazolo[3,4-b]pyrrolo[3,4-d]pyridine derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 74-7	2.9	35
107	Structure-based prediction and biosynthesis of the major mammalian metabolite of the cardioactive prototype LASSBio-294. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 3734-6	2.9	10
106	Antimycobacterial Profile of 5-phenyl-1,3,4-thiadiazole-2-arylhydrazone Derivatives. <i>Letters in Drug Design and Discovery</i> , 2010 , 7, 606-609	0.8	4
105	Extração e purificação de fármacos anti-inflamatórios não esteroidais ciclo-oxigenase-2 seletivos. <i>Química Nova</i> , 2009 , 32, 1324-1328	1.6	4

104	5-Phenyl-2-(benzalhydrazonyl)-1,3,4-thiadiazoles, potential trypanocidal agents: consistent dimer formation via NâHâ⋯N intermolecular hydrogen bonds. <i>Zeitschrift für Kristallographie</i> , 2009 , 224, 598-606		12
103	Sedation and antinociception induced by a new pyrazolo[3,4-b]pyrrolo[3,4-d]pyridine derivative (LASSBio-873) is modulated by activation of muscarinic receptors. <i>Pharmacology Biochemistry and Behavior</i> , 2009 , 94, 70-4	3.9	16
102	Studies towards the identification of putative bioactive conformation of potent vasodilator arylidene N-acylhydrazone derivatives. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 4004-9	6.8	60
101	Synthesis, pharmacological evaluation and docking studies of new sulindac analogues. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 1959-71	6.8	7
100	Development of CoMFA and CoMSIA models of affinity and selectivity for indole ligands of cannabinoid CB1 and CB2 receptors. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 2482-96	6.8	7
99	Discovery of novel analgesic and anti-inflammatory 3-arylamino-imidazo[1,2-a]pyridine symbiotic prototypes. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 74-84	3.4	149
98	Novel 6-methanesulfonamide-3,4-methylenedioxyphenyl-N-acylhydrazones: orally effective anti-inflammatory drug candidates. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 1125-31	3.4	29
97	Synthesis and analgesic profile of conformationally constrained N-acylhydrazone analogues: discovery of novel N-arylideneamino quinazolin-4(3H)-one compounds derived from natural safrole. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 6517-25	3.4	18
96	Design, synthesis and analgesic properties of novel conformationally-restricted N-acylhydrazones (NAH). <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 4963-6	2.9	43
95	Structural insights into IKKbeta inhibition by natural products staurosporine and quercetin. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 6907-10	2.9	10
94	Drug hybridization strategies: before or after lead identification?. <i>Expert Opinion on Drug Discovery</i> , 2009 , 4, 605-9	6.2	47
93	From nature to drug discovery: the indole scaffold as a privileged structure. <i>Mini-Reviews in Medicinal Chemistry</i> , 2009 , 9, 782-93	3.2	406
92	Pharmacokinetic evaluation of LASSBio-579: an N-phenylpiperazine antipsychotic prototype. <i>Journal of Pharmacy and Pharmacology</i> , 2008 , 60, 699-707	4.8	26
91	CNS-selective noncompetitive cholinesterase inhibitors derived from the natural piperidine alkaloid (-)-spectaline. <i>European Journal of Pharmacology</i> , 2008 , 580, 339-49	5.3	30
90	Serotonergic neurotransmission mediates hypothermia induced by the N-phenylpiperazine antipsychotic prototypes LASSBio-579 and LASSBio-581. <i>Pharmacology Biochemistry and Behavior</i> , 2008 , 89, 23-30	3.9	13
89	Microbial reduction of alpha-substituted-alpha-acetyl-gamma-butyrolactones. <i>Catalysis Communications</i> , 2008 , 9, 1782-1786	3.2	6
88	Improved Solvent-Free Dakin Oxidation Protocol. <i>Synthetic Communications</i> , 2008 , 38, 784-788	1.7	20
87	Antinociceptive profile of 2,3,6-trisubstituted piperidine alkaloids: 3-O-acetyl-spectaline and semi-synthetic derivatives of (-)-spectaline. <i>Chemical and Pharmaceutical Bulletin</i> , 2008 , 56, 407-12	1.9	18

86	1-Methyl-7-(4-nitro-phen-yl)-3-phenyl-pyrazolo[3,4-b]pyrrolo[3,4-d]pyridine-6,8(3H,7H)-dione. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2008 , 64, o2356		
85	New Insights for Multifactorial Disease Therapy: The Challenge of the Symbiotic Drugs. <i>Current Drug Therapy</i> , 2008 , 3, 1-13	0.7	16
84	Synthesis and anti-platelet activity of novel arylsulfonate--acylhydrazone derivatives, designed as antithrombotic candidates. <i>European Journal of Medicinal Chemistry</i> , 2008 , 43, 348-56	6.8	52
83	NSAIDs revisited: putative molecular basis of their interactions with peroxisome proliferator-activated gamma receptor (PPARgamma). <i>European Journal of Medicinal Chemistry</i> , 2008 , 43, 1918-25	6.8	7
82	Studies toward the structural optimization of new brazilzone-related trypanocidal 1,3,4-thiadiazole-2-arylhydrazone derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 413-21	3.4	38
81	Atropoisomerismo: o efeito da quiralidade axial em substâncias bioativas. <i>Quimica Nova</i> , 2007 , 30, 125-135.6		13
80	Synthesis, pharmacological evaluation and electrochemical studies of novel 6-nitro-3,4-methylenedioxyphenyl-N-acylhydrazone derivatives: Discovery of LASSBio-881, a new ligand of cannabinoid receptors. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 2421-33	3.4	53
79	Development and validation of a LC-MS/MS method with electrospray ionization for determination of LASSBio-579 in rat plasma. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2007 , 43, 677-82	3.5	4
78	Molecular hybridization: a useful tool in the design of new drug prototypes. <i>Current Medicinal Chemistry</i> , 2007 , 14, 1829-52	4.3	685
77	Privileged structures: a useful concept for the rational design of new lead drug candidates. <i>Mini-Reviews in Medicinal Chemistry</i> , 2007 , 7, 1108-19	3.2	219
76	The Molecular Basis of COX-2 Versus COX-1 Selectivity of Lumiracoxib by Molecular Docking Studies. <i>Letters in Drug Design and Discovery</i> , 2007 , 4, 422-425	0.8	2
75	Medicinal chemistry of N-acylhydrazones: new lead-compounds of analgesic, antiinflammatory and antithrombotic drugs. <i>Current Medicinal Chemistry</i> , 2006 , 13, 167-98	4.3	78
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