Carla Ghelardini

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.

8,997 67 386 45 h-index g-index citations papers 6.07 408 10,442 5.7 L-index ext. citations avg, IF ext. papers

#	Paper	IF	Citations
386	Healthy Properties of a New Formulation of Pomegranate-Peel Extract in Mice Suffering from Experimental Autoimmune Encephalomyelitis <i>Molecules</i> , 2022 , 27,	4.8	3
385	Restorative and pain-relieving effects of fibroin in preclinical models of tendinopathy <i>Biomedicine and Pharmacotherapy</i> , 2022 , 148, 112693	7.5	2
384	Sultam based Carbonic Anhydrase VII inhibitors for the management of neuropathic pain. <i>European Journal of Medicinal Chemistry</i> , 2022 , 227, 113956	6.8	1
383	Effects of Ultramicronized -Palmitoylethanolamine Supplementation on Tramadol and Oxycodone Analgesia and Tolerance Prevention <i>Pharmaceutics</i> , 2022 , 14,	6.4	2
382	Development of Eudragit Nanoparticles for Intranasal Drug Delivery: Preliminary Technological and Toxicological Evaluation. <i>Applied Sciences (Switzerland)</i> , 2022 , 12, 2373	2.6	O
381	Anti-inflammatory Effects of Novel P2X4 Receptor Antagonists, NC-2600 and NP-1815-PX, in a Murine Model of Colitis <i>Inflammation</i> , 2022 , 1	5.1	1
380	Neuronal alarmin IL-1 [®] Levokes astrocyte-mediated protective signals: Effectiveness in chemotherapy-induced neuropathic pain <i>Neurobiology of Disease</i> , 2022 , 168, 105716	7.5	1
379	The Protection of Zinc against Acute Cadmium Exposure: A Morphological and Molecular Study on a BBB In Vitro Model. <i>Cells</i> , 2022 , 11, 1646	7.9	0
378	Beneficial Effect of H2S-Releasing Molecules in an In Vitro Model of Sarcopenia: Relevance of Glucoraphanin. <i>International Journal of Molecular Sciences</i> , 2022 , 23, 5955	6.3	1
377	Inhibitors of Mitochondrial Human Carbonic Anhydrases VA and VB as a Therapeutic Strategy against Paclitaxel-Induced Neuropathic Pain in Mice. <i>International Journal of Molecular Sciences</i> , 2022 , 23, 6229	6.3	2
376	Glyco-Coated CdSe/ZnS Quantum Dots as Nanoprobes for Carbonic Anhydrase IX Imaging in Cancer Cells <i>ACS Applied Nano Materials</i> , 2021 , 4, 14153-14160	5.6	2
375	Precision Medicine in Alzheimer B Disease: Investigating Comorbid Common Biological Substrates in the Rat Model of Amyloid Beta-Induced Toxicity <i>Frontiers in Pharmacology</i> , 2021 , 12, 799561	5.6	1
374	H NMR and HPLC-DAD-MS for the characterization of ellagitannins and triterpenoids of less investigated Anogeissus leiocarpus DC (Combretaceae) stem bark <i>Food Chemistry</i> , 2021 , 375, 131813	8.5	1
373	VEGF-A/VEGFR-1 signalling and chemotherapy-induced neuropathic pain: therapeutic potential of a novel anti-VEGFR-1 monoclonal antibody. <i>Journal of Experimental and Clinical Cancer Research</i> , 2021 , 40, 320	12.8	4
372	Curcumin-in-Cyclodextrins-in-Liposomes: An Alternative for Osteoarthritis Treatment. <i>Proceedings</i> (mdpi), 2021 , 78, 52	0.3	
371	Adenosine A3 agonists reverse neuropathic pain via T cell-mediated production of IL-10. <i>Journal of Clinical Investigation</i> , 2021 , 131,	15.9	11
370	Role of Carbonic Anhydrase in Cerebral Ischemia and Carbonic Anhydrase Inhibitors as Putative Protective Agents. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	3

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369	Oxaliplatin-Induced Neuropathy: Genetic and Epigenetic Profile to Better Understand How to Ameliorate This Side Effect. <i>Frontiers in Molecular Biosciences</i> , 2021 , 8, 643824	5.6	3	
368	Improvement of Butamben Anesthetic Efficacy by the Development of Deformable Liposomes Bearing the Drug as Cyclodextrin Complex. <i>Pharmaceutics</i> , 2021 , 13,	6.4	3	
367	The HS-Donor Erucin Exhibits Protective Effects against Vascular Inflammation in Human Endothelial and Smooth Muscle Cells. <i>Antioxidants</i> , 2021 , 10,	7.1	4	
366	Pain Relieving and Neuroprotective Effects of Non-opioid Compound, DDD-028, in the Rat Model of Paclitaxel-Induced Neuropathy. <i>Neurotherapeutics</i> , 2021 , 18, 2008-2020	6.4	2	
365	Uncovering the Mechanisms of Adenosine Receptor-Mediated Pain Control: Focus on the A Receptor Subtype. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	4	
364	Eruca sativa Mill. seed extract promotes anti-obesity and hypoglycemic effects in mice fed with a high-fat diet. <i>Phytotherapy Research</i> , 2021 , 35, 1983-1990	6.7	5	
363	Therapeutic potential for coxibs-nitric oxide releasing hybrids in cystic fibrosis. <i>European Journal of Medicinal Chemistry</i> , 2021 , 210, 112983	6.8	1	
362	Efficacy of Extract against Inflammatory Pain: In Vivo Studies in Mice. <i>Marine Drugs</i> , 2021 , 19,	6	3	
361	Comparative Assessment of the Activity of Racemic and Dextrorotatory Forms of Thioctic (Alpha-Lipoic) Acid in Low Back Pain: Preclinical Results and Clinical Evidences From an Open Randomized Trial. <i>Frontiers in Pharmacology</i> , 2021 , 12, 607572	5.6	2	
360	Tellurides bearing benzensulfonamide as carbonic anhydrase inhibitors with potent antitumor activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021 , 45, 128147	2.9	2	
359	The Anti-Arthritic Efficacy of Khellin Loaded in Ascorbyl Decanoate Nanovesicles after an Intra-Articular Administration. <i>Pharmaceutics</i> , 2021 , 13,	6.4	2	
358	Lipid Cubic Mesophases Combined with Superparamagnetic Iron Oxide Nanoparticles: A Hybrid Multifunctional Platform with Tunable Magnetic Properties for Nanomedical Applications. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	2	
357	Carbonic Anhydrase IV Selective Inhibitors Counteract the Development of Colitis-Associated Visceral Pain in Rats. <i>Cells</i> , 2021 , 10,	7.9	1	
356	N-Acylethanolamine Acid Amidase Inhibition Potentiates Morphine Analgesia and Delays the Development of Tolerance. <i>Neurotherapeutics</i> , 2021 , 1	6.4	2	
355	Design and synthesis of the first indole-based blockers of Panx-1 channel. <i>European Journal of Medicinal Chemistry</i> , 2021 , 223, 113650	6.8	1	
354	Chalcogenides-incorporating carbonic anhydrase inhibitors concomitantly reverted oxaliplatin-induced neuropathy and enhanced antiproliferative action. <i>European Journal of Medicinal Chemistry</i> , 2021 , 225, 113793	6.8	7	
353	Protective effects of carbonic anhydrase inhibition in brain ischaemia and models. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 964-976	5.6	4	
352	New Perspectives in the Pathophysiology and Treatment of Pain in Patients with Dry Eye Disease Journal of Clinical Medicine, 2021 , 11,	5.1	1	

351	Phenyl(thio)phosphon(amid)ate Benzenesulfonamides as Potent and Selective Inhibitors of Human Carbonic Anhydrases II and VII Counteract Allodynia in a Mouse Model of Oxaliplatin-Induced Neuropathy. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 5185-5200	8.3	6
350	Design, synthesis and biological evaluation of 7-substituted 4-phenyl-6H-imidazo[1,5-a]thieno[3,2-f] [1,4]diazepines as safe anxiolytic agents. <i>European Journal of Medicinal Chemistry</i> , 2020 , 200, 112405	6.8	2
349	Effects of the Combination of EHydroxy-EMethyl Butyrate and R(+) Lipoic Acid in a Cellular Model of Sarcopenia. <i>Molecules</i> , 2020 , 25,	4.8	2
348	The Use of the Selective Imidazoline I Receptor Agonist Carbophenyline as a Strategy for Neuropathic Pain Relief: Preclinical Evaluation in a Mouse Model of Oxaliplatin-Induced Neurotoxicity. <i>Neurotherapeutics</i> , 2020 , 17, 1005-1015	6.4	5
347	Intra-Articular Route for the System of Molecules 14G1862 from: Pain Relieving and Protective Effects in a Rat Model of Osteoarthritis. <i>Nutrients</i> , 2020 , 12,	6.7	3
346	Coronaridine congeners decrease neuropathic pain in mice and inhibit 1010 nicotinic acetylcholine receptors and Ca2.2 channels. <i>Neuropharmacology</i> , 2020 , 175, 108194	5.5	8
345	Functional Selectivity and Antinociceptive Effects of a Novel KOPr Agonist. <i>Frontiers in Pharmacology</i> , 2020 , 11, 188	5.6	18
344	Pomegranate Mesocarp against Colitis-Induced Visceral Pain in Rats: Effects of a Decoction and Its Fractions. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	9
343	The Anti-Inflammatory and Pain-Relieving Effects of AR170, an Adenosine A Receptor Agonist, in a Rat Model of Colitis. <i>Cells</i> , 2020 , 9,	7.9	6
342	ESitosterol Loaded Nanostructured Lipid Carrier: Physical and Oxidative Stability, In Vitro Simulated Digestion and Hypocholesterolemic Activity. <i>Pharmaceutics</i> , 2020 , 12,	6.4	7
341	Novel formyl peptide receptor (FPR) agonists with pyridinone and pyrimidindione scaffolds that are potentially useful for the treatment of rheumatoid arthritis. <i>Bioorganic Chemistry</i> , 2020 , 100, 103880	5.1	9
340	Bioisosteric Development of Multitarget Nonsteroidal Anti-Inflammatory Drug-Carbonic Anhydrases Inhibitor Hybrids for the Management of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 2325-2342	8.3	14
339	The endocannabinoid system dual-target ligand N-cycloheptyl-1,2-dihydro-5-bromo-1-(4-fluorobenzyl)-6-methyl-2-oxo-pyridine-3-carboxamide improves disease severity in a mouse model of multiple sclerosis. <i>European Journal of Medicinal</i>	6.8	8
338	Chemistry, 2020 , 208, 112858 Intranasal Low-Dose Naltrexone Against Opioid Side Effects: A Preclinical Study. <i>Frontiers in Pharmacology</i> , 2020 , 11, 576624	5.6	3
337	()-3-Furan-2-yltolyl-acrylamide and its Derivative DM489 Decrease Neuropathic Pain in Mice Predominantly by Nicotinic Acetylcholine Receptor Potentiation. ACS Chemical Neuroscience, 2020, 11, 3603-3614	5.7	6
336	The active second-generation proteasome inhibitor oprozomib reverts the oxaliplatin-induced neuropathy symptoms. <i>Biochemical Pharmacology</i> , 2020 , 182, 114255	6	3
335	Pain Modulation in WAG/Rij Epileptic Rats (A Genetic Model of Absence Epilepsy): Effects of Biological and Pharmacological Histone Deacetylase Inhibitors. <i>Frontiers in Pharmacology</i> , 2020 , 11, 549	1599	5
334	Development of a stable oral pediatric solution of hydrochlorothiazide by the combined use of cyclodextrins and hydrophilic polymers. <i>International Journal of Pharmaceutics</i> , 2020 , 587, 119692	6.5	2

333	Deepening the Mechanisms of Visceral Pain Persistence: An Evaluation of the Gut-Spinal Cord Relationship. <i>Cells</i> , 2020 , 9,	7.9	11
332	Acute visceral pain relief mediated by A3AR agonists in rats: involvement of N-type voltage-gated calcium channels. <i>Pain</i> , 2020 , 161, 2179-2190	8	11
331	Toxicological Profile of the Pain-Relieving Antioxidant Compound Thioctic Acid in Its Racemic and Enantiomeric Forms. <i>Antioxidants</i> , 2020 , 9,	7.1	3
330	Erucin exhibits vasorelaxing effects and antihypertensive activity by H S-releasing properties. <i>British Journal of Pharmacology</i> , 2020 , 177, 824-835	8.6	29
329	Bacopa monnieri as augmentation therapy in the treatment of anhedonia, preclinical and clinical evaluation. <i>Phytotherapy Research</i> , 2020 , 34, 2331-2340	6.7	7
328	Antioxidant-Conjugated 1,2,4-Triazolo[4,3-]pyrazin-3-one Derivatives: Highly Potent and Selective Human A Adenosine Receptor Antagonists Possessing Protective Efficacy in Neuropathic Pain. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 8511-8531	8.3	9
327	Protective Effects Induced by Two Polyphenolic Liquid Complexes from Olive (, mainly) Pressing Juice in Rat Isolated Tissues Challenged with LPS. <i>Molecules</i> , 2019 , 24,	4.8	13
326	Meal against Diabetic Neuropathic Pain: An HS-Mediated Effect of Glucoerucin. <i>Molecules</i> , 2019 , 24,	4.8	13
325	Synthesis, biological evaluation and molecular modeling of novel selective COX-2 inhibitors: sulfide, sulfoxide, and sulfone derivatives of 1,5-diarylpyrrol-3-substituted scaffold. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 115045	3.4	14
324	Intestinal inflammation increases convulsant activity and reduces antiepileptic drug efficacy in a mouse model of epilepsy. <i>Scientific Reports</i> , 2019 , 9, 13983	4.9	22
323	New Rigid Nicotine Analogues, Carrying a Norbornane Moiety, Are Potent Agonists of II and II* Nicotinic Receptors. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 1887-1901	8.3	4
322	N-aryl-NPureido-O-sulfamates: Potent and selective inhibitors of the human Carbonic Anhydrase VII isoform with neuropathic pain relieving properties. <i>Bioorganic Chemistry</i> , 2019 , 89, 103033	5.1	10
321	Benzensulfonamides bearing spyrohydantoin moieties act as potent inhibitors of human carbonic anhydrases II and VII and show neuropathic pain attenuating effects. <i>European Journal of Medicinal Chemistry</i> , 2019 , 177, 188-197	6.8	19
320	Pain Relieving Effect of-NSAIDs-CAIs Hybrid Molecules: Systemic and Intra-Articular Treatments against Rheumatoid Arthritis. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	17
319	Blueberry juice protects osteocytes and bone precursor cells against oxidative stress partly through SIRT1. <i>FEBS Open Bio</i> , 2019 , 9, 1082-1096	2.7	10
318	Novel 8-amino-1,2,4-triazolo[4,3-a]pyrazin-3-one derivatives as potent human adenosine A and A receptor antagonists. Evaluation of their protective effect against Emyloid-induced neurotoxicity in SH-SY5Y cells. <i>Bioorganic Chemistry</i> , 2019 , 87, 380-394	5.1	11
317	Identification of the First Synthetic Allosteric Modulator of the CB Receptors and Evidence of Its Efficacy for Neuropathic Pain Relief. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 276-287	8.3	28
316	Synthesis of novel tellurides bearing benzensulfonamide moiety as carbonic anhydrase inhibitors with antitumor activity. <i>European Journal of Medicinal Chemistry</i> , 2019 , 181, 111586	6.8	18

315	L. Extract: Alpha-Amylase Inhibition against Metabolic Syndrome in Mice. <i>Nutrients</i> , 2019 , 11,	6.7	10
314	Modifications on the Amino-3,5-dicyanopyridine Core To Obtain Multifaceted Adenosine Receptor Ligands with Antineuropathic Activity. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 6894-6912	8.3	9
313	Synthesis and Evaluation of Carbonic Anhydrase Inhibitors with Carbon Monoxide Releasing Properties for the Management of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 7233-	- <mark>82</mark> 49	26
312	Effect of NIR laser therapy by MLS-MiS source against neuropathic pain in rats: in vivo and ex vivo analysis. <i>Scientific Reports</i> , 2019 , 9, 9297	4.9	10
311	Spirocyclic sulfonamides with carbonic anhydrase inhibitory and anti-neuropathic pain activity. <i>Bioorganic Chemistry</i> , 2019 , 92, 103210	5.1	8
310	Nanostructured lipid carriers for oral delivery of silymarin: Improving its absorption and in vivo efficacy in type 2 diabetes and metabolic syndrome model. <i>International Journal of Pharmaceutics</i> , 2019 , 572, 118838	6.5	19
309	Mesenchymal stem cells, implications for pain therapy. Neural Regeneration Research, 2019, 14, 1915-19	1465	2
308	Researching New Therapeutic Approaches for Abdominal Visceral Pain Treatment: Preclinical Effects of an Assembled System of Molecules of Vegetal Origin. <i>Nutrients</i> , 2019 , 12,	6.7	11
307	Intra-articular mucilages: behavioural and histological evaluations for a new model of articular pain. Journal of Pharmacy and Pharmacology, 2019 , 71, 971-981	4.8	7
306	Cannabidiol Protects Dopaminergic Neuronal Cells from Cadmium. <i>International Journal of Environmental Research and Public Health</i> , 2019 , 16,	4.6	16
305	Adenosine A3 receptor activation inhibits pronociceptive N-type Ca2+ currents and cell excitability in dorsal root ganglion neurons. <i>Pain</i> , 2019 , 160, 1103-1118	8	24
304	Effects of Cadmium on ZO-1 Tight Junction Integrity of the Blood Brain Barrier. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	31
303	Anticancer properties of erucin, an H S-releasing isothiocyanate, on human pancreatic adenocarcinoma cells (AsPC-1). <i>Phytotherapy Research</i> , 2019 , 33, 845-855	6.7	42
302	Selenium and zinc: Two key players against cadmium-induced neuronal toxicity. <i>Toxicology in Vitro</i> , 2018 , 48, 159-169	3.6	43
301	Design, Synthesis, and X-ray of Selenides as New Class of Agents for Prevention of Diabetic Cerebrovascular Pathology. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 462-467	4.3	17
300	Involvement of the N/OFQ-NOP system in rat morphine antinociceptive tolerance: Are astrocytes the crossroad?. <i>European Journal of Pharmacology</i> , 2018 , 823, 79-86	5.3	5
299	Structural investigations on coumarins leading to chromeno[4,3-c]pyrazol-4-ones and pyrano[4,3-c]pyrazol-4-ones: New scaffolds for the design of the tumor-associated carbonic anhydrase isoforms IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2018 , 146, 47-59	6.8	36
298	Discovery of 1,5-Diphenylpyrazole-3-Carboxamide Derivatives as Potent, Reversible, and Selective Monoacylglycerol Lipase (MAGL) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 1340-1354	8.3	28

297	Histamine-deficient mice do not respond to the antidepressant-like effects of oleoylethanolamide. <i>Neuropharmacology</i> , 2018 , 135, 234-241	5.5	10
296	Resolution of co-eluting isomers of anti-inflammatory drugs conjugated to carbonic anhydrase inhibitors from plasma in liquid chromatography by energy-resolved tandem mass spectrometry. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 671-679	5.6	14
295	Development of Potent Inhibitors of Fatty Acid Amide Hydrolase Useful for the Treatment of Neuropathic Pain. <i>ChemMedChem</i> , 2018 , 13, 2090-2103	3.7	11
294	Ultramicronized -Palmitoylethanolamine Supplementation for Long-Lasting, Low-Dosed Morphine Antinociception. <i>Frontiers in Pharmacology</i> , 2018 , 9, 473	5.6	12
293	Effect of glucoraphanin and sulforaphane against chemotherapy-induced neuropathic pain: Kv7 potassium channels modulation by H S release in vivo. <i>Phytotherapy Research</i> , 2018 , 32, 2226-2234	6.7	44
292	Tanshinones from Salvia miltiorrhiza Bunge revert chemotherapy-induced neuropathic pain and reduce glioblastoma cells malignancy. <i>Biomedicine and Pharmacotherapy</i> , 2018 , 105, 1042-1049	7.5	26
291	Discovery of Novel Nonsteroidal Anti-Inflammatory Drugs and Carbonic Anhydrase Inhibitors Hybrids (NSAIDs-CAIs) for the Management of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 4961-4977	8.3	37
290	Treatment with acetyl-L-carnitine exerts a neuroprotective effect in the sciatic nerve following loose ligation: a functional and microanatomical study. <i>Neural Regeneration Research</i> , 2018 , 13, 692-698	4.5	7
289	Adipose-derived stem cells decrease pain in a rat model of oxaliplatin-induced neuropathy: Role of VEGF-A modulation. <i>Neuropharmacology</i> , 2018 , 131, 166-175	5.5	21
288	Oxaliplatin-induced blood brain barrier loosening: a new point of view on chemotherapy-induced neurotoxicity. <i>Oncotarget</i> , 2018 , 9, 23426-23438	3.3	33
287	Selective Blockade of HCN1/HCN2 Channels as a Potential Pharmacological Strategy Against Pain. <i>Frontiers in Pharmacology</i> , 2018 , 9, 1252	5.6	26
286	Design, characterization and in vivo evaluation of nanostructured lipid carriers (NLC) as a new drug delivery system for hydrochlorothiazide oral administration in pediatric therapy. <i>Drug Delivery</i> , 2018 , 25, 1910-1921	7	51
285	4-Hydroxy-3-nitro-5-ureido-benzenesulfonamides Selectively Target the Tumor-Associated Carbonic Anhydrase Isoforms IX and XII Showing Hypoxia-Enhanced Antiproliferative Profiles. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 10860-10874	8.3	41
284	Combined Approach of Cyclodextrin Complexationand Nanostructured Lipid Carriers for the Development of a Pediatric Liquid Oral Dosage Form of Hydrochlorothiazide. <i>Pharmaceutics</i> , 2018 , 10,	6.4	13
283	Efficacy of isothiocyanate-based compounds on different forms of persistent pain. <i>Journal of Pain Research</i> , 2018 , 11, 2905-2913	2.9	9
282	Interaction of Half Oxa-/Half -Platin Complex with Human Superoxide Dismutase and Induced Reduction of Neurotoxicity. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 1094-1098	4.3	2
281	Effect of Vitis vinifera hydroalcoholic extract against oxaliplatin neurotoxicity: in vitro and in vivo evidence. <i>Scientific Reports</i> , 2018 , 8, 14364	4.9	10
2 80	Discovery of new 2, 5-disubstituted 1,3-selenazoles as selective human carbonic anhydrase IX inhibitors with potent anti-tumor activity. <i>European Journal of Medicinal Chemistry</i> , 2018 , 157, 1214-122	2 6.8	22

279	Heterocoumarins Are Selective Carbonic Anhydrase IX and XII Inhibitors with Cytotoxic Effects against Cancer Cells Lines. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 947-951	4.3	26
278	Improving the therapeutic efficacy of prilocaine by PLGA microparticles: Preparation, characterization and in vivo evaluation. <i>International Journal of Pharmaceutics</i> , 2018 , 547, 24-30	6.5	14
277	Design, synthesis and X-ray crystallography of selenides bearing benzenesulfonamide moiety with neuropathic pain modulating effects. <i>European Journal of Medicinal Chemistry</i> , 2018 , 154, 210-219	6.8	31
276	Liposomal Formulation to Increase Stability and Prolong Antineuropathic Activity of Verbascoside. <i>Planta Medica</i> , 2017 , 83, 412-419	3.1	25
275	Apoptotic Process Induced by Oxaliplatin in Rat Hippocampus Causes Memory Impairment. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2017 , 120, 14-21	3.1	5
274	Design and Synthesis of Novel Nonsteroidal Anti-Inflammatory Drugs and Carbonic Anhydrase Inhibitors Hybrids (NSAIDs-CAIs) for the Treatment of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 1159-1170	8.3	94
273	HuD-mediated distinct BDNF regulatory pathways promote regeneration after nerve injury. <i>Brain Research</i> , 2017 , 1659, 55-63	3.7	15
272	Inhibition of 91 0 nicotinic acetylcholine receptors prevents chemotherapy-induced neuropathic pain. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, E1825-I	E1832	100
271	Spinal astrocytic c-Jun N-terminal kinase (JNK) activation as counteracting mechanism to the amitriptyline analgesic efficacy in painful peripheral neuropathies. <i>European Journal of Pharmacology</i> , 2017 , 798, 85-93	5.3	8
270	Piperazines as nootropic agents: New derivatives of the potent cognition-enhancer DM235 carrying hydrophilic substituents. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1795-1803	3.4	5
269	Synthesis and pharmacological evaluation of pyrazolo[1,5-a]pyrimidin-7(4H)-one derivatives as potential GABA-R ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1901-1906	3.4	14
268	Astragali radix: could it be an adjuvant for oxaliplatin-induced neuropathy?. <i>Scientific Reports</i> , 2017 , 7, 42021	4.9	41
267	Effects of Hypericum perforatum extract on oxaliplatin-induced neurotoxicity: in vitro evaluations. Zeitschrift Fur Naturforschung - Section C Journal of Biosciences, 2017, 72, 219-226	1.7	10
266	Development and in vivo evaluation of an innovative "Hydrochlorothiazide-in Cyclodextrins-in Solid Lipid Nanoparticles" formulation with sustained release and enhanced oral bioavailability for potential hypertension treatment in pediatrics. <i>International Journal of Pharmaceutics</i> , 2017 , 521, 73-83	6.5	30
265	Synthesis and Pharmacological Evaluation of Novel GABAA Subtype Receptor Ligands with Potential Anxiolytic-like and Anti-hyperalgesic Effect. <i>Journal of Heterocyclic Chemistry</i> , 2017 , 54, 2788-2	2 7 99	7
264	Effects of natural and synthetic isothiocyanate-based HS-releasers against chemotherapy-induced neuropathic pain: Role of Kv7 potassium channels. <i>Neuropharmacology</i> , 2017 , 121, 49-59	5.5	65
263	Behavioural phenotype of histamine H receptor knockout mice: Focus on central neuronal functions. <i>Neuropharmacology</i> , 2017 , 114, 48-57	5.5	32
262	Development and characterization of fast dissolving tablets of oxaprozin based on hybrid systems of the drug with cyclodextrins and nanoclays. <i>International Journal of Pharmaceutics</i> , 2017 , 531, 640-649	96.5	8

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261	Synergic stimulation of serotonin 5-HT receptor and Eddrenoceptors for neuropathic pain relief: Preclinical effects of 2-substituted imidazoline derivatives. <i>European Journal of Pharmacology</i> , 2017 , 810, 128-133	5.3	14
260	Synthesis and Biological Evaluation of Novel Neuroprotective Pyridazine Derivatives as Excitatory Amino Acid Transporter 2 (EAAT2) Activators. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 5216-5221	8.3	12
259	Pain relieving and protective effects of Astragalus hydroalcoholic extract in rat arthritis models. Journal of Pharmacy and Pharmacology, 2017 , 69, 1858-1870	4.8	14
258	Lipoyl-Homotaurine Derivative (ADM_12) Reverts Oxaliplatin-Induced Neuropathy and Reduces Cancer Cells Malignancy by Inhibiting Carbonic Anhydrase IX (CAIX). <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 9003-9011	8.3	6
257	Prophylactic versus Therapeutic Fingolimod: Restoration of Presynaptic Defects in Mice Suffering from Experimental Autoimmune Encephalomyelitis. <i>PLoS ONE</i> , 2017 , 12, e0170825	3.7	22
256	St. Johnß Wort Potentiates anti-Nociceptive Effects of Morphine in Mice Models of Neuropathic Pain. <i>Pain Medicine</i> , 2017 , 18, 1334-1343	2.8	7
255	Effects of a water extract of Lepidium meyenii root in different models of persistent pain in rats. Zeitschrift Fur Naturforschung - Section C Journal of Biosciences, 2017, 72, 449-457	1.7	8
254	Photobiomodulation therapy by NIR laser in persistent pain: an analytical study in the rat. <i>Lasers in Medical Science</i> , 2017 , 32, 1835-1846	3.1	10
253	Calcium alginate microspheres containing metformin hydrochloride niosomes and chitosomes aimed for oral therapy of type 2 diabetes mellitus. <i>International Journal of Pharmaceutics</i> , 2017 , 530, 430-439	6.5	29
252	Discovery of New Selenoureido Analogues of 4-(4-Fluorophenylureido)benzenesulfonamide as Carbonic Anhydrase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 963-968	4.3	51
251	Discovery of New Sulfonamide Carbonic Anhydrase IX Inhibitors Incorporating Nitrogenous Bases. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 1314-1319	4.3	46
250	Identification of a New Pyrazolo[1,5-a]quinazoline Ligand Highly Affine to EAminobutyric Type A (GABA) Receptor Subtype with Anxiolytic-Like and Antihyperalgesic Activity. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 9691-9702	8.3	15
249	A rat model of FOLFOX-induced neuropathy: effects of oral dimiracetam in comparison with duloxetine and pregabalin. <i>Cancer Chemotherapy and Pharmacology</i> , 2017 , 80, 1091-1103	3.5	13
248	3-Hydroxy-1H-quinazoline-2,4-dione as a New Scaffold To Develop Potent and Selective Inhibitors of the Tumor-Associated Carbonic Anhydrases IX and XII. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 6428-	ê439	22
247	Histamine H receptor agonist-induced relief from painful peripheral neuropathy is mediated by inhibition of spinal neuroinflammation and oxidative stress. <i>British Journal of Pharmacology</i> , 2017 , 174, 28-40	8.6	19
246	Effect of amitriptyline treatment on neurofilament-H protein in an experimental model of depression. <i>Brain Research Bulletin</i> , 2017 , 128, 1-6	3.9	10
245	Adrenoceptor: a Target for Neuropathic Pain Treatment. <i>Mini-Reviews in Medicinal Chemistry</i> , 2017 , 17, 95-107	3.2	20
244	Neurorestoration from medicinal plants: an opportunity to treat painful neuropathies. <i>Neural Regeneration Research</i> , 2017 , 12, 403-404	4.5	

243	The Versatile 2-Substituted Imidazoline Nucleus as a Structural Motif of Ligands Directed to the Serotonin 5-HT Receptor. <i>ChemMedChem</i> , 2016 , 11, 2287-2298	3.7	6
242	Anti-neuropathic effects of Rosmarinus officinalis L. terpenoid fraction: relevance of nicotinic receptors. <i>Scientific Reports</i> , 2016 , 6, 34832	4.9	16
241	Synthesis, antiarrhythmic activity, and toxicological evaluation of mexiletine analogues. <i>European Journal of Medicinal Chemistry</i> , 2016 , 121, 300-307	6.8	15
240	Design, Synthesis, and Biological Evaluation of Imidazo[1,5-a]quinoline as Highly Potent Ligands of Central Benzodiazepine Receptors. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 3353-72	8.3	10
239	Altered Expression of Cytoskeletal and Axonal Proteins in Oxaliplatin-Induced Neuropathy. <i>Pharmacology</i> , 2016 , 97, 146-50	2.3	10
238	Effect of the SOD mimetic MnL4 on in vitro and in vivo oxaliplatin toxicity: Possible aid in chemotherapy induced neuropathy. <i>Free Radical Biology and Medicine</i> , 2016 , 93, 67-76	7.8	26
237	Development and Pharmacological Characterization of Selective Blockers of 2-Arachidonoyl Glycerol Degradation with Efficacy in Rodent Models of Multiple Sclerosis and Pain. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 2612-32	8.3	49
236	Synthesis and biological evaluation of fluorinated 1,5-diarylpyrrole-3-alkoxyethyl ether derivatives as selective COX-2 inhibitors endowed with anti-inflammatory activity. <i>European Journal of Medicinal Chemistry</i> , 2016 , 109, 99-106	6.8	24
235	Carbonic anhydrase inhibition for the management of cerebral ischemia: in vivo evaluation of sulfonamide and coumarin inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 894-9	5.6	78
234	Characterisation of the Novel Mixed Mu-NOP Peptide Ligand Dermorphin-N/OFQ (DeNo). <i>PLoS ONE</i> , 2016 , 11, e0156897	3.7	21
233	A Series of COX-2 Inhibitors Endowed with NO-Releasing Properties: Synthesis, Biological Evaluation, and Docking Analysis. <i>ChemMedChem</i> , 2016 , 11, 1804-11	3.7	5
232	Blockade of the spinal BDNF-activated JNK pathway prevents the development of antiretroviral-induced neuropathic pain. <i>Neuropharmacology</i> , 2016 , 105, 543-552	5.5	19
231	Effects of the neutrophil elastase inhibitor EL-17 in rat adjuvant-induced arthritis. <i>Rheumatology</i> , 2016 , 55, 1285-94	3.9	12
230	The B110 nicotinic receptor antagonist Econotoxin RgIA prevents neuropathic pain induced by oxaliplatin treatment. <i>Experimental Neurology</i> , 2016 , 282, 37-48	5.7	43
229	The novel HS-donor 4-carboxyphenyl isothiocyanate promotes cardioprotective effects against ischemia/reperfusion injury through activation of mitoK channels and reduction of oxidative stress. <i>Pharmacological Research</i> , 2016 , 113, 290-299	10.2	50
228	Atomoxetine for hoarding disorder: A pre-clinical and clinical investigation. <i>Journal of Psychiatric Research</i> , 2016 , 83, 240-248	5.2	19
227	Acute effect of Capparis spinosa root extracts on rat articular pain. <i>Journal of Ethnopharmacology</i> , 2016 , 193, 456-465	5	19
226	A class of sulfonamide carbonic anhydrase inhibitors with neuropathic pain modulating effects. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 1828-40	3.4	103

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225	Acute and subchronic antinociceptive effects of nociceptin/orphanin FQ receptor agonists infused by intrathecal route in rats. <i>European Journal of Pharmacology</i> , 2015 , 754, 73-81	5.3	17
224	Oxaliplatin evokes P2X7-dependent glutamate release in the cerebral cortex: A pain mechanism mediated by Pannexin 1. <i>Neuropharmacology</i> , 2015 , 97, 133-41	5.5	40
223	Different apoptotic pathways activated by oxaliplatin in primary astrocytes vs. colo-rectal cancer cells. <i>International Journal of Molecular Sciences</i> , 2015 , 16, 5386-99	6.3	12
222	6-Substituted sulfocoumarins are selective carbonic anhdydrase IX and XII inhibitors with significant cytotoxicity against colorectal cancer cells. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 3975-83	8.3	75
221	Substituted piperazines as nootropic agents: 2- or 3-phenyl derivatives structurally related to the cognition-enhancer DM235. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 1700-1704	2.9	13
220	In Vitro Evidence for the Use of Astragali Radix Extracts as Adjuvant against Oxaliplatin-Induced Neurotoxicity. <i>Planta Medica</i> , 2015 , 81, 1045-55	3.1	25
219	A model of neuropathic pain induced by sorafenib in the rat: Effect of dimiracetam. <i>NeuroToxicology</i> , 2015 , 50, 101-7	4.4	8
218	Inhibition of spinal ERK1/2-c-JUN signaling pathway counteracts the development of low doses morphine-induced hyperalgesia. <i>European Journal of Pharmacology</i> , 2015 , 764, 271-277	5.3	15
217	Intrathecal administration of nociceptin/orphanin FQ receptor agonists in rats: A strategy to relieve chemotherapy-induced neuropathic hypersensitivity. <i>European Journal of Pharmacology</i> , 2015 , 766, 155	5-53	15
216	Nociceptin/orphanin FQ receptor and pain: Feasibility of the fourth opioid family member. <i>European Journal of Pharmacology</i> , 2015 , 766, 151-4	5.3	9
	Luropeuri Journal of Friarmacology, 2013, 100, 131-4	<i>)</i> • <i>)</i>	
215	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. <i>Future Science OA</i> , 2015 , 1, FSO2	2.7	8
215	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management.		
	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. Future Science OA, 2015, 1, FSO2 Synthesis of five and six-membered heterocycles bearing an arylpiperazinylalkyl side chain as orally	2.7	
214	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. Future Science OA, 2015, 1, FSO2 Synthesis of five and six-membered heterocycles bearing an arylpiperazinylalkyl side chain as orally active antinociceptive agents. Bioorganic and Medicinal Chemistry, 2015, 23, 6237-45 Further studies on pyrazolo[1P,5P1,6]pyrimido[4,5-d]pyridazin-4(3H)-ones as potent and selective	2.7 3.4	8
214	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. Future Science OA, 2015, 1, FSO2 Synthesis of five and six-membered heterocycles bearing an arylpiperazinylalkyl side chain as orally active antinociceptive agents. Bioorganic and Medicinal Chemistry, 2015, 23, 6237-45 Further studies on pyrazolo[1P,5P1,6]pyrimido[4,5-d]pyridazin-4(3H)-ones as potent and selective human A1 adenosine receptor antagonists. European Journal of Medicinal Chemistry, 2015, 89, 32-41 Histamine H4 receptor activation alleviates neuropathic pain through differential regulation of	2.7 3.4 6.8	8 7 11
214 213 212	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. <i>Future Science OA</i> , 2015 , 1, FSO2 Synthesis of five and six-membered heterocycles bearing an arylpiperazinylalkyl side chain as orally active antinociceptive agents. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 6237-45 Further studies on pyrazolo[1P,5P1,6]pyrimido[4,5-d]pyridazin-4(3H)-ones as potent and selective human A1 adenosine receptor antagonists. <i>European Journal of Medicinal Chemistry</i> , 2015 , 89, 32-41 Histamine H4 receptor activation alleviates neuropathic pain through differential regulation of ERK, JNK, and P38 MAPK phosphorylation. <i>Pain</i> , 2015 , 156, 2492-2504 Oxidative, metabolic, and apoptotic responses of Schwann cells to high glucose levels. <i>Journal of</i>	2.7 3.4 6.8	8 7 11 39
214 213 212 211	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. Future Science OA, 2015, 1, FSO2 Synthesis of five and six-membered heterocycles bearing an arylpiperazinylalkyl side chain as orally active antinociceptive agents. Bioorganic and Medicinal Chemistry, 2015, 23, 6237-45 Further studies on pyrazolo[1P,5P,1,6]pyrimido[4,5-d]pyridazin-4(3H)-ones as potent and selective human A1 adenosine receptor antagonists. European Journal of Medicinal Chemistry, 2015, 89, 32-41 Histamine H4 receptor activation alleviates neuropathic pain through differential regulation of ERK, JNK, and P38 MAPK phosphorylation. Pain, 2015, 156, 2492-2504 Oxidative, metabolic, and apoptotic responses of Schwann cells to high glucose levels. Journal of Biochemical and Molecular Toxicology, 2015, 29, 274-9 Activation of JNK pathway in spinal astrocytes contributes to acute ultra-low-dose morphine	2.7 3.4 6.8 8	8 7 11 39 21

207	Antineuropathic profile of N-palmitoylethanolamine in a rat model of oxaliplatin-induced neurotoxicity. <i>PLoS ONE</i> , 2015 , 10, e0128080	3.7	63
206	Delay of morphine tolerance by palmitoylethanolamide. <i>BioMed Research International</i> , 2015 , 2015, 89	47332	22
205	☑ Nicotinic Receptor Promotes the Neuroprotective Functions of Astrocytes against Oxaliplatin Neurotoxicity. <i>Neural Plasticity</i> , 2015 , 2015, 396908	3.3	16
204	Widespread pain reliever profile of a flower extract of Tanacetum parthenium. <i>Phytomedicine</i> , 2015 , 22, 752-8	6.5	12
203	Differential contribution of GIJo subunits in the response to food deprivation. <i>European Journal of Pharmacology</i> , 2015 , 750, 27-31	5.3	
202	Lipoic-based TRPA1/TRPV1 antagonist to treat orofacial pain. ACS Chemical Neuroscience, 2015, 6, 380-	· 5 5.7	13
201	Synthesis, biological evaluation and docking analysis of a new series of methylsulfonyl and sulfamoyl acetamides and ethyl acetates as potent COX-2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 810-20	3.4	18
2 00	Peroxisome determination in optical microscopy: a useful tool derived by a simplification of an old ultrastructural technique. <i>Acta Histochemica</i> , 2014 , 116, 863-70	2	2
199	Broad spectrum and prolonged efficacy of dimiracetam in models of neuropathic pain. <i>Neuropharmacology</i> , 2014 , 81, 85-94	5.5	7
198	Development of a chitosan-derivative micellar formulation to improve celecoxib solubility and bioavailability. <i>Drug Development and Industrial Pharmacy</i> , 2014 , 40, 1494-502	3.6	15
197	Econotoxin RgIA protects against the development of nerve injury-induced chronic pain and prevents both neuronal and glial derangement. <i>Pain</i> , 2014 , 155, 1986-95	8	73
196	Glial role in oxaliplatin-induced neuropathic pain. Experimental Neurology, 2014, 261, 22-33	5.7	105
195	Regionally selective activation of ERK and JNK in morphine paradoxical hyperalgesia: a step toward improving opioid pain therapy. <i>Neuropharmacology</i> , 2014 , 86, 67-77	5.5	29
194	PKC-mediated HuD-GAP43 pathway activation in a mouse model of antiretroviral painful neuropathy. <i>Pharmacological Research</i> , 2014 , 81, 44-53	10.2	24
193	Enhancing the pharmacodynamic profile of a class of selective COX-2 inhibiting nitric oxide donors. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 772-86	3.4	20
192	Involvement of II nAChR subtype in rat oxaliplatin-induced neuropathy: effects of selective activation. <i>Neuropharmacology</i> , 2014 , 79, 37-48	5.5	64
191	A smart platform for hyperthermia application in cancer treatment: cobalt-doped ferrite nanoparticles mineralized in human ferritin cages. <i>ACS Nano</i> , 2014 , 8, 4705-19	16.7	154
190	Serotonergic modulation in neuropathy induced by oxaliplatin: effect on the 5HT2C receptor. <i>European Journal of Pharmacology</i> , 2014 , 735, 141-9	5.3	29

189	PKC-mediated potentiation of morphine analgesia by St. Johnß Wort in rodents and humans. <i>Journal of Pharmacological Sciences</i> , 2014 , 124, 409-17	3.7	13
188	PPAR- Impairment alters peroxisome functionality in primary astrocyte cell cultures. <i>BioMed Research International</i> , 2014 , 2014, 546453	3	12
187	BDNF and Artemin are increased in drug-naWe non-depressed GAD patients: preliminary data. <i>International Journal of Psychiatry in Clinical Practice</i> , 2014 , 18, 255-60	2.4	10
186	Spinal administration of mGluR5 antagonist prevents the onset of bortezomib induced neuropathic pain in rat. <i>Neuropharmacology</i> , 2014 , 86, 294-300	5.5	17
185	St. Johnß Wort seed and feverfew flower extracts relieve painful diabetic neuropathy in a rat model of diabetes. <i>Floterap</i> [12014, 92, 23-33	3.2	14
184	Prophylactic role of acetyl-l-carnitine on knee lesions and associated pain in a rat model of osteoarthritis. <i>Life Sciences</i> , 2014 , 106, 32-9	6.8	12
183	Oxaliplatin neurotoxicity involves peroxisome alterations. PPARlagonism as preventive pharmacological approach. <i>PLoS ONE</i> , 2014 , 9, e102758	3.7	54
182	Reversal of NO-induced nociceptive hypersensitivity by St. JohnB wort and hypericin: NF-B, CREB and STAT1 as molecular targets. <i>Psychopharmacology</i> , 2013 , 227, 149-63	4.7	15
181	Nicotine is a pain reliever in trauma- and chemotherapy-induced neuropathy models. <i>European Journal of Pharmacology</i> , 2013 , 711, 87-94	5.3	15
180	Morphologic features and glial activation in rat oxaliplatin-dependent neuropathic pain. <i>Journal of Pain</i> , 2013 , 14, 1585-600	5.2	122
179	Differentiation state affects morphine induced cell regulation in neuroblastoma cultured cells. <i>Neuroscience Letters</i> , 2013 , 555, 51-6	3.3	6
178	Low dose native type II collagen prevents pain in a rat osteoarthritis model. <i>BMC Musculoskeletal Disorders</i> , 2013 , 14, 228	2.8	27
177	Pleiotropic effect of histamine H4 receptor modulation in the central nervous system. <i>Neuropharmacology</i> , 2013 , 71, 141-7	5.5	39
176	A class of pyrrole derivatives endowed with analgesic/anti-inflammatory activity. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 3695-701	3.4	54
175	2-Arylpyrazolo[4,3-d]pyrimidin-7-amino derivatives as new potent and selective human A3 adenosine receptor antagonists. Molecular modeling studies and pharmacological evaluation. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 2256-69	8.3	24
174	Synthesis of novel cognition enhancers with pyrazolo[5,1-c][1,2,4]benzotriazine core acting at Eminobutyric acid type A (GABA(A)) receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 2186-2198	3.4	23
173	Inhibition of the PKCIIpathway relieves from meningeal nociception in an animal model: an innovative perspective for migraine therapy?. <i>Neurotherapeutics</i> , 2013 , 10, 329-39	6.4	11
172	Novel analgesic/anti-inflammatory agents: 1,5-diarylpyrrole nitrooxyalkyl ethers and related compounds as cyclooxygenase-2 inhibiting nitric oxide donors. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 3191-206	8.3	37

171	St. JohnB wort reversal of meningeal nociception: a natural therapeutic perspective for migraine pain. <i>Phytomedicine</i> , 2013 , 20, 930-8	6.5	14
170	Flow Synthesis and Biological Studies of an Analgesic Adamantane Derivative That Inhibits P2X7-Evoked Glutamate Release. <i>ACS Medicinal Chemistry Letters</i> , 2013 , 4, 704-9	4.3	12
169	Neuroprotective activity of thioctic acid in central nervous system lesions consequent to peripheral nerve injury. <i>BioMed Research International</i> , 2013 , 2013, 985093	3	16
168	Therapeutic effects of the superoxide dismutase mimetic compound MnIIMe2DO2A on experimental articular pain in rats. <i>Mediators of Inflammation</i> , 2013 , 2013, 905360	4.3	42
167	St. JohnB wort relieves pain in an animal model of migraine. European Journal of Pain, 2013, 17, 369-81	3.7	12
166	Dual effect of morphine in long-term social memory in rat. <i>British Journal of Pharmacology</i> , 2013 , 168, 1786-93	8.6	8
165	A TRPA1 antagonist reverts oxaliplatin-induced neuropathic pain. Scientific Reports, 2013, 3, 2005	4.9	47
164	PKC-Dependent Signaling Pathways within PAG and Thalamus Contribute to the Nitric Oxide-Induced Nociceptive Behavior. <i>ISRN Pain</i> , 2013 , 2013, 471378		2
163	Aminopyrrolic synthetic receptors for monosaccharides: a class of carbohydrate-binding agents endowed with antibiotic activity versus pathogenic yeasts. <i>Chemistry - A European Journal</i> , 2012 , 18, 500	6 4: 82	26
162	Selective modulation of the PKCe/p38MAP kinase signalling pathway for the antidepressant-like activity of amitriptyline. <i>Neuropharmacology</i> , 2012 , 62, 289-96	5.5	19
161	Oxaliplatin-induced neuropathy: oxidative stress as pathological mechanism. Protective effect of silibinin. <i>Journal of Pain</i> , 2012 , 13, 276-84	5.2	127
160	3-Hydroxy-1H-quinazoline-2,4-dione derivatives as new antagonists at ionotropic glutamate receptors: molecular modeling and pharmacological studies. <i>European Journal of Medicinal Chemistry</i> , 2012 , 54, 470-82	6.8	28
159	Improving the solubility of a new class of antiinflammatory pharmacodynamic hybrids, that release nitric oxide and inhibit cycloxygenase-2 isoenzyme. <i>European Journal of Medicinal Chemistry</i> , 2012 , 58, 287-98	6.8	14
158	Synthesis and toxicopharmacological evaluation of m-hydroxymexiletine, the first metabolite of mexiletine more potent than the parent compound on voltage-gated sodium channels. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 1418-22	8.3	21
157	Regionally selective activation and differential regulation of ERK, JNK and p38 MAP kinase signalling pathway by protein kinase C in mood modulation. <i>International Journal of Neuropsychopharmacology</i> , 2012 , 15, 781-93	5.8	38
156	New insight into the central benzodiazepine receptor-ligand interactions: design, synthesis, biological evaluation, and molecular modeling of 3-substituted 6-phenyl-4H-imidazo[1,5-a][1,4]benzodiazepines and related compounds. <i>Journal of Medicinal</i>	8.3	37
155	Synthesis and biological evaluation of 3,7-diazabicyclo[4.3.0]nonan-8-ones as potential nootropic and analgesic drugs. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 2512-6	8.3	6
154	Synthesis and structure-activity relationship studies in translocator protein ligands based on a pyrazolo[3,4-b]quinoline scaffold. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 7165-75	8.3	21

153	Contribution of G inhibitory protein alpha subunits in paradoxical hyperalgesia elicited by exceedingly low doses of morphine in mice. <i>Life Sciences</i> , 2011 , 89, 918-25	6.8	6	
152	Antidepressant phenotype by inhibiting the phospholipase C[1]protein kinase C[pathway in the forced swim test. <i>Neuropharmacology</i> , 2011 , 60, 937-43	5.5	19	
151	Antihyperalgesic activity of verbascoside in two models of neuropathic pain. <i>Journal of Pharmacy and Pharmacology</i> , 2011 , 63, 594-601	4.8	37	
150	Novel analgesic/anti-inflammatory agents: diarylpyrrole acetic esters endowed with nitric oxide releasing properties. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 7759-71	8.3	36	
149	Analgesic and antineuropathic drugs acting through central cholinergic mechanisms. <i>Recent Patents on CNS Drug Discovery</i> , 2011 , 6, 119-40		44	
148	Salvianolic acid B and its liposomal formulations: anti-hyperalgesic activity in the treatment of neuropathic pain. <i>European Journal of Pharmaceutical Sciences</i> , 2011 , 44, 552-8	5.1	36	
147	Development of ligands at Elaminobutyrric acid type A (GABAA) receptor subtype as new agents for pain relief. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 7441-52	3.4	15	
146	Antidepressant-like effect of artemin in mice: a mechanism for acetyl-L-carnitine activity on depression. <i>Psychopharmacology</i> , 2011 , 218, 347-56	4.7	20	
145	Effects of dietary extra-virgin olive oil on behaviour and brain biochemical parameters in ageing rats. <i>British Journal of Nutrition</i> , 2010 , 103, 1674-83	3.6	51	
144	A prolonged protein kinase C-mediated, opioid-related antinociceptive effect of st Johnß Wort in mice. <i>Journal of Pain</i> , 2010 , 11, 149-59	5.2	34	
143	Liposomal formulations of prilocaine: effect of complexation with hydroxypropyl-Etyclodextrin on drug anesthetic efficacy. <i>Journal of Liposome Research</i> , 2010 , 20, 315-22	6.1	33	
142	Novel ester and acid derivatives of the 1,5-diarylpyrrole scaffold as anti-inflammatory and analgesic agents. Synthesis and in vitro and in vivo biological evaluation. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 723-33	8.3	36	
141	New fluoro derivatives of the pyrazolo[5,1-c][1,2,4]benzotriazine 5-oxide system: evaluation of fluorine binding properties in the benzodiazepine site on Eminobutyrric acid type A (GABA(A)) receptor. Design, synthesis, biological, and molecular modeling investigation. <i>Journal of Medicinal</i>	8.3	22	
140	Chemistry, 2010 , 53, 7532-48 Protective effect of alpha7 nAChR: behavioural and morphological features on neuropathy. <i>Pain</i> , 2010 , 150, 542-549	8	45	
139	St. Johnß Wort reduces neuropathic pain through a hypericin-mediated inhibition of the protein kinase Cgamma and epsilon activity. <i>Biochemical Pharmacology</i> , 2010 , 79, 1327-36	6	43	
138	Chiral aryloxyalkylamines: Selective 5-HT(1B/1D) activation and analgesic activity. <i>ChemMedChem</i> , 2010 , 5, 696-704	3.7	10	
137	Supraspinal role of protein kinase C in oxaliplatin-induced neuropathy in rat. <i>Pain</i> , 2009 , 146, 141-7	8	42	
136	Neuroprotective effects of acetyl-L-carnitine on neuropathic pain and apoptosis: a role for the nicotinic receptor. <i>Journal of Neuroscience Research</i> , 2009 , 87, 200-7	4.4	39	

135	Alpha2-agonists as analgesic agents. Medicinal Research Reviews, 2009, 29, 339-68	14.4	39
134	The central analgesia induced by antimigraine drugs is independent from Gi proteins: superiority of a fixed combination of indomethacin, prochlorperazine and caffeine, compared to sumatriptan, in an in vivo model. <i>Journal of Headache and Pain</i> , 2009 , 10, 435-40	8.8	5
133	Design, synthesis and nootropic activity of new analogues of sunifiram and sapunifiram, two potent cognition-enhancers. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 7606-14	3.4	8
132	Supraspinal Gbetagamma-dependent stimulation of PLCbeta originating from G inhibitory protein-mu opioid receptor-coupling is necessary for morphine induced acute hyperalgesia. <i>Journal of Neurochemistry</i> , 2009 , 111, 171-80	6	28
131	[35S]GTP gamma S binding studies of amphiphilic drugs-activated Gi proteins: a caveat. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 2224-9	2.9	
130	A novel manganese complex effective as superoxide anion scavenger and therapeutic agent against cell and tissue oxidative injury. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 7273-83	8.3	34
129	Synthesis, in vivo evaluation, and molecular modeling studies of new pyrazolo[5,1-c][1,2,4]benzotriazine 5-oxide derivatives. Identification of a bifunctional hydrogen bond area related to the inverse agonism. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 4668-82	8.3	19
128	Effect of preparation technique on the properties and in vivo efficacy of benzocaine-loaded ethosomes. <i>Journal of Liposome Research</i> , 2009 , 19, 253-60	6.1	56
127	Receptor-independent modulation of reconstituted Galpha(i) protein mediated by liposomes. <i>Molecular BioSystems</i> , 2009 , 5, 356-67		2
126	Acetyl-L-carnitine in the management of pain during methadone withdrawal syndrome. <i>Clinical Neuropharmacology</i> , 2009 , 32, 35-40	1.4	6
125	Molecular interaction in the mouse PAG between NMDA and opioid receptors in morphine-induced acute thermal nociception. <i>Journal of Neurochemistry</i> , 2008 , 105, 91-100	6	18
124	Dopamine is necessary to endogenous morphine formation in mammalian brain in vivo. <i>Journal of Neurochemistry</i> , 2008 , 106, 2337-44	6	21
123	A gene-specific cerebral types 1, 2, and 3 RyR protein knockdown induces an antidepressant-like effect in mice. <i>Journal of Neurochemistry</i> , 2008 , 106, 2385-94	6	24
122	Synthesis, biological evaluation, and enzyme docking simulations of 1,5-diarylpyrrole-3-alkoxyethyl ethers as selective cyclooxygenase-2 inhibitors endowed with anti-inflammatory and antinociceptive activity. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 4476-81	8.3	46
121	An antidepressant behaviour in mice carrying a gene-specific InsP3R1, InsP3R2 and InsP3R3 protein knockdown. <i>Neuropharmacology</i> , 2008 , 55, 1156-64	5.5	18
120	Ethyl 8-fluoro-6-(3-nitrophenyl)-4H-imidazo[1,5-a][1,4]benzodiazepine-3-carboxylate as novel, highly potent, and safe antianxiety agent. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 4730-43	8.3	32
119	Different involvement of type 1, 2, and 3 ryanodine receptors in memory processes. <i>Learning and Memory</i> , 2008 , 15, 315-23	2.8	41
118	Design, synthesis and preliminary pharmacological evaluation of new piperidine and piperazine derivatives as cognition-enhancers. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 1431-43	3.4	11

117	Synthesis and biological evaluation of novel dimiracetam derivatives useful for the treatment of neuropathic pain. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 3224-32	3.4	12
116	Novel 3-aroylpyrazolo[5,1-c][1,2,4]benzotriazine 5-oxides 8-substituted, ligands at GABAA/benzodiazepine receptor complex: synthesis, pharmacological and molecular modeling studies. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 4471-89	3.4	23
115	Synthesis, in vitro, and in vivo biological evaluation and molecular docking simulations of chiral alcohol and ether derivatives of the 1,5-diarylpyrrole scaffold as novel anti-inflammatory and analgesic agents. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 8072-81	3.4	16
114	Design, synthesis and preliminary pharmacological evaluation of new analogues of DM232 (unifiram) and DM235 (sunifiram) as cognition modulators. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 10034-42	3.4	9
113	Cyclooxygenase-2 inhibitors. 1,5-diarylpyrrol-3-acetic esters with enhanced inhibitory activity toward cyclooxygenase-2 and improved cyclooxygenase-2/cyclooxygenase-1 selectivity. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 5403-11	8.3	52
112	4-amino-5-substituted-3(2H)-pyridazinones as orally active antinociceptive agents: synthesis and studies on the mechanism of action. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3945-53	8.3	19
111	Design, synthesis, and preliminary pharmacological evaluation of new quinoline derivatives as nicotinic ligands. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 4993-5002	8.3	11
110	Synthesis and biological evaluation of chiral alpha-aminoanilides with central antinociceptive activity. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 1907-15	8.3	10
109	Structure-activity relationships of methoctramine-related polyamines as muscarinic antagonist: effect of replacing the inner polymethylene chain with cyclic moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 2312-21	3.4	8
108	Novel 3-iodo-8-ethoxypyrazolo[5,1-c][1,2,4]benzotriazine 5-oxide as promising lead for design of alpha5-inverse agonist useful tools for therapy of mnemonic damage. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 2573-86	3.4	18
107	New 1,8-naphthyridine and quinoline derivatives as CB2 selective agonists. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 6505-10	2.9	58
106	Development, characterization and in vivo evaluation of benzocaine-loaded liposomes. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2007 , 67, 86-95	5.7	116
105	Activity and expression of semicarbazide-sensitive benzylamine oxidase in a rodent model of diabetes: interactive effects with methylamine and alpha-aminoguanidine. <i>European Journal of Pharmacology</i> , 2006 , 529, 179-87	5.3	11
104	Arylpiperazinylalkylpyridazinones and analogues as potent and orally active antinociceptive agents: synthesis and studies on mechanism of action. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 7826-35	8.3	34
103	Structural investigation of the 7-chloro-3-hydroxy-1H-quinazoline-2,4-dione scaffold to obtain AMPA and kainate receptor selective antagonists. Synthesis, pharmacological, and molecular modeling studies. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 6015-26	8.3	41
102	A recombinant transductor-effector system: in vitro study of G inhibitory protein (G-alpha-i1) direct activators. <i>Archives of Biochemistry and Biophysics</i> , 2006 , 453, 151-60	4.1	2
101	Signaling pathway of morphine induced acute thermal hyperalgesia in mice. <i>Pain</i> , 2006 , 123, 294-305	8	60
100	Gi/o proteins: expression for direct activation enquiry. <i>Protein Expression and Purification</i> , 2006 , 47, 303	3-10	6

99	Blockade of intracellular calcium release induces an antidepressant-like effect in the mouse forced swimming test. <i>Neuropharmacology</i> , 2006 , 50, 309-16	5.5	38
98	Pharmacological characterization of DM232 (unifiram) and DM235 (sunifiram), new potent cognition enhancers. <i>CNS Neuroscience & Therapeutics</i> , 2006 , 12, 39-52		12
97	4-methyl benzylamine stimulates food consumption and counteracts the hypophagic effects of amphetamine acting on brain Shaker-like Kv1.1 channels. <i>British Journal of Pharmacology</i> , 2006 , 147, 218-24	8.6	3
96	Ryanodine receptors are involved in muscarinic antinociception in mice. <i>Behavioural Brain Research</i> , 2005 , 164, 165-71	3.4	6
95	Design, synthesis, and preliminary pharmacological evaluation of a set of small molecules that directly activate gi proteins. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 6491-503	8.3	7
94	Enantioselective synthesis and preliminary pharmacological evaluation of the enantiomers of unifiram (DM232), a potent cognition-enhancing agent. <i>Medicinal Chemistry</i> , 2005 , 1, 473-80	1.8	7
93	Effects of endogenous morphine deprivation on memory retention of passive avoidance learning in mice. <i>International Journal of Neuropsychopharmacology</i> , 2004 , 7, 311-9	5.8	17
92	Methylamine, but not ammonia, is hypophagic in mouse by interaction with brain Kv1.6 channel subtype. <i>British Journal of Pharmacology</i> , 2004 , 142, 381-9	8.6	11
91	Structure-activity relationship studies on unifiram (DM232) and sunifiram (DM235), two novel and potent cognition enhancing drugs. <i>Bioorganic and Medicinal Chemistry</i> , 2004 , 12, 71-85	3.4	17
90	Design, synthesis, and preliminary pharmacological evaluation of 4-aminopiperidine derivatives as N-type calcium channel blockers active on pain and neuropathic pain. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 6070-81	8.3	24
89	Prochlorperazine induces central antinociception mediated by the muscarinic system. <i>Pharmacological Research</i> , 2004 , 50, 351-8	10.2	15
88	Indomethacin, alone and combined with prochlorperazine and caffeine, but not sumatriptan, abolishes peripheral and central sensitization in in vivo models of migraine. <i>Journal of Pain</i> , 2004 , 5, 413	3- 5 ²	15
87	Alpha-2 agonist-induced memory impairment is mediated by the alpha-2A-adrenoceptor subtype. <i>Behavioural Brain Research</i> , 2004 , 153, 409-17	3.4	35
86	H1-receptor stimulation induces hyperalgesia through activation of the phospholipase C-PKC pathway. <i>Neuropharmacology</i> , 2004 , 47, 295-303	5.5	27
85	Acetyl-L-carnitine requires phospholipase C-IP3 pathway activation to induce antinociception. <i>Neuropharmacology</i> , 2004 , 47, 286-94	5.5	22
84	Role of intracellular calcium in acute thermal pain perception. <i>Neuropharmacology</i> , 2004 , 47, 935-44	5.5	14
83	Synthesis, biological evaluation, and receptor docking simulations of 2-[(acylamino)ethyl]-1,4-benzodiazepines as kappa-opioid receptor agonists endowed with antinociceptive and antiamnesic activity. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 3853-64	8.3	27
82	Antisense knockdown of the Shaker-like Kv1.1 gene abolishes the central stimulatory effects of amphetamines in mice and rats. <i>Neuropsychopharmacology</i> , 2003 , 28, 1096-105	8.7	13

(2002-2003)

81	AMPA-receptor activation is involved in the antiamnesic effect of DM 232 (unifiram) and DM 235 (sunifiram). <i>Naunyn-SchmiedebergmArchives of Pharmacology</i> , 2003 , 368, 538-45	3.4	12	
80	2-pyrrolidinone moiety is not critical for the cognition-enhancing activity of piracetam-like drugs. <i>Il Farmaco</i> , 2003 , 58, 715-22		6	
79	4-Aminopiperidine derivatives as a new class of potent cognition enhancing drugs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003 , 13, 2303-6	2.9	16	
78	[(3-Chlorophenyl)piperazinylpropyl]pyridazinones and analogues as potent antinociceptive agents. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 1055-9	8.3	98	
77	The phospholipase C-IP3 pathway is involved in muscarinic antinociception. Neuropsychopharmacology, 2003 , 28, 888-97	8.7	44	
76	Role of Gi proteins in the antidepressant-like effect of amitriptyline and clomipramine. <i>Neuropsychopharmacology</i> , 2002 , 27, 554-64	8.7	12	
75	The novel nootropic compound DM232 (UNIFIRAM) ameliorates memory impairment in mice and rats. <i>Drug Development Research</i> , 2002 , 56, 23-32	5.1	7	
74	DM235 (sunifiram): a novel nootropic with potential as a cognitive enhancer. <i>Naunyn-Schmiedebergm Archives of Pharmacology</i> , 2002 , 365, 419-26	3.4	15	
73	Endogenous morphine modulates acute thermonociception in mice. <i>Journal of Neurochemistry</i> , 2002 , 80, 271-7	6	18	
72	Novel potent 5-HT(3) receptor ligands based on the pyrrolidone structure: synthesis, biological evaluation, and computational rationalization of the ligand-receptor interaction modalities. <i>Bioorganic and Medicinal Chemistry</i> , 2002 , 10, 779-801	3.4	32	
71	Central cholinergic challenging of migraine by testing second-generation anticholinesterase drugs. Headache, 2002 , 42, 596-602	4.2	32	
70	Indomethacin, caffeine and prochlorperazine alone and combined revert hyperalgesia in in vivo models of migraine. <i>Pharmacological Research</i> , 2002 , 46, 245-50	10.2	21	
69	Design and study of piracetam-like nootropics, controversial members of the problematic class of cognition-enhancing drugs. <i>Current Pharmaceutical Design</i> , 2002 , 8, 125-38	3.3	94	
68	Inactivation of Gi proteins induces an antidepressant-like effect in the mouse forced-swimming test. <i>Neuropharmacology</i> , 2002 , 43, 457-65	5.5		
67	Acetyl-l-carnitine induces muscarinic antinocieption in mice and rats. <i>Neuropharmacology</i> , 2002 , 43, 11	80 5 75	43	
66	Amitriptyline and clomipramine activate Gi-protein signaling pathway in the induction of analgesia. <i>Naunyn-Schmiedebergm Archives of Pharmacology</i> , 2002 , 365, 1-7	3.4	15	
65	Antihistamine antinociception is mediated by Gi-protein activation. <i>Neuroscience</i> , 2002 , 109, 811-8	3.9	9	
64	Menthol: a natural analgesic compound. <i>Neuroscience Letters</i> , 2002 , 322, 145-8	3.3	257	

63	Hypofunctionality of Gi proteins as aetiopathogenic mechanism for migraine and cluster headache. <i>Cephalalgia</i> , 2001 , 21, 38-45	6.1	30
62	Antiamnesic effect of the two novel Eppioid agonists, VA-100 and VA-101, in the mouse passive avoidance test. <i>Drug Development Research</i> , 2001 , 54, 12-18	5.1	2
61	Antinociceptive profile of the natural cholinesterase inhibitor huperzine A. <i>Drug Development Research</i> , 2001 , 54, 19-26	5.1	1
60	Differential prevention of morphine amnesia by antisense oligodeoxynucleotides directed against various Gi-protein alpha subunits. <i>British Journal of Pharmacology</i> , 2001 , 133, 267-74	8.6	13
59	Methylamine and benzylamine induced hypophagia in mice: modulation by semicarbazide-sensitive benzylamine oxidase inhibitors and aODN towards Kv1.1 channels. <i>British Journal of Pharmacology</i> , 2001 , 134, 880-6	8.6	20
58	Selective inhibition of amine oxidases differently potentiate the hypophagic effect of benzylamine in mice. <i>European Journal of Pharmacology</i> , 2001 , 413, 91-9	5.3	7
57	M1 receptor activation is a requirement for arecoline analgesia. <i>Il Farmaco</i> , 2001 , 56, 383-5		27
56	Local anaesthetic activity of beta-caryophyllene. <i>Il Farmaco</i> , 2001 , 56, 387-9		185
55	Local anaesthetic activity of monoterpenes and phenylpropanes of essential oils. <i>Planta Medica</i> , 2001 , 67, 564-6	3.1	54
54	Local anaesthetic activity of (+)- and (-)-menthol. <i>Planta Medica</i> , 2001 , 67, 174-6	3.1	44
53	Structure-affinity relationships of a unique nicotinic ligand: N(1)-dimethyl-N(4)-phenylpiperazinium iodide (DMPP). <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 3946-55	8.3	35
52	Involvement of potassium channels in amitriptyline and clomipramine analgesia. <i>Neuropharmacology</i> , 2001 , 40, 75-84	5.5	49
51	Design, synthesis, and biological activity of methoctramine-related polyamines as putative G(i) protein activators. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 4035-8	8.3	9
50	AG-4: a nicotinic agonist endowed with antiamnesic properties. <i>Drug Development Research</i> , 2000 , 51, 191-196	5.1	
49	Loss of muscarinic antinociception by antisense inhibition of M(1) receptors. <i>British Journal of Pharmacology</i> , 2000 , 129, 1633-40	8.6	44
48	Pharmacological identification of SM-21, the novel sigma(2) antagonist. <i>Pharmacology Biochemistry and Behavior</i> , 2000 , 67, 659-62	3.9	28
47	Local anaesthetic, antibacterial and antifungal properties of sesquiterpenes from myrrh. <i>Planta Medica</i> , 2000 , 66, 356-8	3.1	108
46	Molecular simplification of 1,4-diazabicyclo[4.3.0]nonan-9-ones gives piperazine derivatives that maintain high nootropic activity. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 4499-507	8.3	28

(1998-2000)

45	Design, synthesis, and preliminary pharmacological evaluation of 1, 4-diazabicyclo[4.3.0]nonan-9-ones as a new class of highly potent nootropic agents. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 1969-74	8.3	25	
44	Antinociception induced by amitriptyline and imipramine is mediated by alpha2A-adrenoceptors. <i>The Japanese Journal of Pharmacology</i> , 2000 , 82, 130-7		50	
43	Role of 5-HT1A receptors in a mouse passive avoidance paradigm. <i>The Japanese Journal of Pharmacology</i> , 2000 , 84, 418-24		14	
42	Local anaesthetic activity of the essential oil of Lavandula angustifolia. <i>Planta Medica</i> , 1999 , 65, 700-3	3.1	134	
41	Hybridized and isosteric analogues of N1-acetyl-N4-dimethyl-piperazinium iodide (ADMP) and N1-phenyl-N4-dimethyl-piperazinium iodide (DMPP) with central nicotinic action. <i>Bioorganic and Medicinal Chemistry</i> , 1999 , 7, 457-65	3.4	21	
40	Role of potassium channels in the antinociception induced by agonists of alpha2-adrenoceptors. <i>British Journal of Pharmacology</i> , 1999 , 126, 1214-20	8.6	33	
39	Effect of potassium channel modulators in mouse forced swimming test. <i>British Journal of Pharmacology</i> , 1999 , 126, 1653-9	8.6	47	
38	Improvement of cognitive functions by the acetylcholine releaser SM 21 1999 , 47, 118-126		2	
37	Antisense RnockdownsPof M1 receptors induces transient anterograde amnesia in mice. <i>Neuropharmacology</i> , 1999 , 38, 339-48	5.5	23	
36	The role of potassium channels in antihistamine analgesia. <i>Neuropharmacology</i> , 1999 , 38, 1893-901	5.5	18	
35	Antinociceptive effect of R-(+)-hyoscyamine on the conjunctival reflex test in rabbits. <i>The Japanese Journal of Pharmacology</i> , 1999 , 81, 34-7		2	
34	Influence of potassium channel modulators on cognitive processes in mice. <i>British Journal of Pharmacology</i> , 1998 , 123, 1079-84	8.6	42	
33	Further structure-activity relationships in the series of tropanyl esters endowed with potent antinociceptive activity. <i>Il Farmaco</i> , 1998 , 53, 764-72		1	
32	Antiamnesic activity of the nicotinic agonist DBO-83 in mice. <i>Drug Development Research</i> , 1998 , 45, 45-	55.1	2	
31	No development of tolerance to analgesia by repeated administration of H1 antagonists. <i>Life Sciences</i> , 1998 , 63, PL 317-22	6.8	5	
30	Mono- and disubstituted-3,8-diazabicyclo[3.2.1]octane derivatives as analgesics structurally related to epibatidine: synthesis, activity, and modeling. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 674-81	8.3	51	
29	Kv1.1 channel antisense attenuates learning and modulation of dentate polysialylated NCAM. <i>NeuroReport</i> , 1998 , 9, 2727-31	1.7	10	
28	Memory facilitation and stimulation of endogenous nerve growth factor synthesis by the acetylcholine releaser PG-9. <i>The Japanese Journal of Pharmacology</i> , 1998 , 78, 245-51		15	

27	Effect of K+ channel modulation on mouse feeding behaviour. <i>European Journal of Pharmacology</i> , 1997 , 329, 1-8	5.3	16
26	Blockade of clomipramine and amitriptyline analgesia by an antisense oligonucleotide to mKv1.1, a mouse Shaker-like K+ channel. <i>European Journal of Pharmacology</i> , 1997 , 330, 15-25	5.3	20
25	In vitro characterization of a novel, potent and selective M3 antagonist. <i>Life Sciences</i> , 1997 , 61, 1217-26	6.8	4
24	Antinociception induced by SM 32 depends on a central cholinergic mechanism. <i>Pharmacological Research</i> , 1997 , 35, 141-7	10.2	4
23	Antiamnesic activity of metoclopramide, cisapride and SR-17 in the mouse passive avoidance test. <i>Pharmacological Research</i> , 1997 , 36, 59-67	10.2	6
22	Reversible antisense inhibition of Shaker-like Kv1.1 potassium channel expression impairs associative memory in mouse and rat. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1997 , 94, 4430-4	11.5	71
21	5-HT1A agonists induce central cholinergic antinociception. <i>Pharmacology Biochemistry and Behavior</i> , 1997 , 57, 835-41	3.9	24
20	Effects of two histamine-N-methyltransferase inhibitors, SKF 91488 and BW 301 U, in rodent antinociception. <i>Naunyn-SchmiedebergmArchives of Pharmacology</i> , 1997 , 355, 354-60	3.4	22
19	Caffeine induces central cholinergic analgesia. <i>Naunyn-Schmiedebergm Archives of Pharmacology</i> , 1997 , 356, 590-5	3.4	48
18	SAR studies on the potent and selective muscarinic antagonist 2-ethylthio-2,2-diphenylacetic acid N,N-diethylaminoethyl ester. <i>Archiv Der Pharmazie</i> , 1997 , 330, 122-8	4.3	4
17	Antinociceptive profile of the new nicotinic agonist DBO-83 1997 , 40, 251-258		14
16	Antinociceptive property of the nicotinic agonist AG-4 in rodents 1997 , 41, 1-9		13
15	Central muscarinic antinociception induced by ET-142 and SS-20 in rodents 1997 , 42, 26-34		5
14	S-(-)-ET 126: a potent and selective M1 antagonist in vitro and in vivo. <i>Life Sciences</i> , 1996 , 58, 991-1000	6.8	14
13	Synthesis, characterization and pharmacological profile of tropicamide enantiomers. <i>Life Sciences</i> , 1996 , 58, 2147-53	6.8	15
12	Central cholinergic antinociception induced by 5HT4 agonists: BIMU 1 and BIMU 8. <i>Life Sciences</i> , 1996 , 58, 2297-309	6.8	38
11	Effect of pertussis toxin on morphine, diphenhydramine, baclofen, clomipramine and physostigmine antinociception. <i>European Journal of Pharmacology</i> , 1996 , 308, 125-33	5.3	22
10	Investigation into the role of histamine receptors in rodent antinociception. <i>Pharmacology Biochemistry and Behavior</i> , 1996 , 53, 567-74	3.9	35

LIST OF PUBLICATIONS

9	Synthesis and enantioselectivity of the enantiomers of PG9 and SM21, new potent analgesic and cognition-enhancing drugs. <i>Chirality</i> , 1996 , 8, 225-33	2.1	10
8	Chiral synthesis and pharmacological evaluation of the enantiomers of SM32, a new analgesic and cognition-enhancing agent. <i>Chirality</i> , 1996 , 8, 579-84	2.1	5
7	Reduced flexibility analogs of analgesic and cognition enhancing alpha-tropanyl esters. <i>Archiv Der Pharmazie</i> , 1996 , 329, 105-11	4.3	4
6	Analgesic effects of myrrh. <i>Nature</i> , 1996 , 379, 29	50.4	87
5	Presynaptic cholinergic modulators as potent cognition enhancers and analgesic drugs. 2. 2-Phenoxy-, 2-(phenylthio)-, and 2-(phenylamino)alkanoic acid esters. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 1712-9	8.3	30
4	Presynaptic cholinergic modulators as potent cognition enhancers and analgesic drugs. 1. Tropic and 2-phenylpropionic acid esters. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 1704-11	8.3	31
3	Role of histamine in rodent antinociception. <i>British Journal of Pharmacology</i> , 1994 , 111, 1269-79	8.6	88
2	Stereoselectivity in central analgesic action of tocainide and its analogs. <i>Chirality</i> , 1993 , 5, 135-42	2.1	12
1	Presynaptic cholinergic mechanism of R-(+) hyoscyamine activity: In vitro studies. <i>Pharmacological Research</i> , 1992 , 26, 322	10.2	1