Fabrizio Vincenzi

List of Publications by Year
in descending order

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119 papers	4,660 citations	87888 38 h-index	123424 61 g-index
119 all docs	119 docs citations	119 times ranked	4554 citing authors

#	Article	IF	Citations
1	Pulsed Electromagnetic Fields: A Novel Attractive Therapeutic Opportunity for Neuroprotection After Acute Cerebral Ischemia. Neuromodulation, 2022, 25, 1240-1247.	0.8	10
2	A_{2A} Adenosine Receptor Antagonists in Neurodegenerative Diseases. Current Medicinal Chemistry, 2022, 29, 4138-4151.	2.4	18
3	Polypharmacological Approaches for CNS Diseases: Focus on Endocannabinoid Degradation Inhibition. Cells, 2022, 11, 471.	4.1	21
4	Adenosine Receptors in Neuropsychiatric Disorders: Fine Regulators of Neurotransmission and Potential Therapeutic Targets. International Journal of Molecular Sciences, 2022, 23, 1219.	4.1	20
5	Dopamine Transporter, PhosphoSerine129 Î±-Synuclein and Î±-Synuclein Levels in Aged LRRK2 G2019S Knock-In and Knock-Out Mice. Biomedicines, 2022, 10, 881.	3.2	5
6	4-Heteroaryl Substituted Amino-3,5-Dicyanopyridines as New Adenosine Receptor Ligands: Novel Insights on Structure-Activity Relationships and Perspectives. Pharmaceuticals, 2022, 15, 478.	3.8	4
7	Pathophysiological Role and Medicinal Chemistry of A2A Adenosine Receptor Antagonists in Alzheimerâ€™s Disease. Molecules, 2022, 27, 2680.	3.8	17
8	A2A Adenosine Receptor: A Possible Therapeutic Target for Alzheimerâ€™s Disease by Regulating NLRP3 Inflammasome Activity?. International Journal of Molecular Sciences, 2022, 23, 5056.	4.1	9
9	Azetidin-2-one-based small molecules as dual hHDAC6/HDAC8 inhibitors: Investigation of their mechanism of action and impact of dual inhibition profile on cell viability. European Journal of Medicinal Chemistry, 2022, 238, 114409.	5.5	11
10	Behavioral and binding studies on the quinolinyl ester indoles 5F-PB22 (5F-QUPIC) and BB-22 (QUCHIC) in the mouse model. Emerging Trends in Drugs, Addictions, and Health, 2022, 2, 100039.	1.1	4
11	Adenosine A2A receptor inhibition reduces synaptic and cognitive hippocampal alterations in Fmr1 KO mice. Translational Psychiatry, 2021, 11, 112.	4.8	18
12	An Open Question: Is the A2A Adenosine Receptor a Novel Target for Alzheimerâ€™s Disease Treatment?. Frontiers in Pharmacology, 2021, 12, 652455.	3.5	15
13	Upregulation of Cortical A2A Adenosine Receptors Is Reflected in Platelets of Patients with Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2021, 80, 1105-1117.	2.6	21
14	Adenosine and Inflammation: Here, There and Everywhere. International Journal of Molecular Sciences, 2021, 22, 7685.	4.1	63
15	A2A Adenosine Receptor as a Potential Biomarker and a Possible Therapeutic Target in Alzheimerâ€™s Disease. Cells, 2021, 10, 2344.	4.1	15
16	A3 Adenosine and P2X7 Purinergic Receptors as New Targets for an Innovative Pharmacological Therapy of Malignant Pleural Mesothelioma. Frontiers in Oncology, 2021, 11, 679285.	2.8	13
17	l-DOPA promotes striatal dopamine release through D1 receptors and reversal of dopamine transporter. Brain Research, 2021, 1768, 147583.	2.2	9
18	Pulsed Electromagnetic Field Stimulation in Osteogenesis and Chondrogenesis: Signaling Pathways and Therapeutic Implications. International Journal of Molecular Sciences, 2021, 22, 809.	4.1	41

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19	Signaling pathways involved in anti-inflammatory effects of Pulsed Electromagnetic Field in microglial cells. Cytokine, 2020, 125, 154777.	3.2	10
20	A2A adenosine receptors are involved in the reparative response of tendon cells to pulsed electromagnetic fields. PLoS ONE, 2020, 15, e0239807.	2.5	2
21	Targeting Adenosine Receptors: A Potential Pharmacological Avenue for Acute and Chronic Pain. International Journal of Molecular Sciences, 2020, 21, 8710.	4.1	43
22	Piperazine- and Piperidine-Containing Thiazolo[5,4-d]pyrimidine Derivatives as New Potent and Selective Adenosine A2A Receptor Inverse Agonists. Pharmaceuticals, 2020, 13, 161.	3.8	11
23	A1 Adenosine Receptor Partial Agonists and Allosteric Modulators: Advancing Toward the Clinic?. Frontiers in Pharmacology, 2020, 11, 625134.	3.5	8
24	Pulsed Electromagnetic Fields Stimulate HIF-1Î±-Independent VEGF Release in 1321N1 Human Astrocytes Protecting Neuron-like SH-SY5Y Cells from Oxygen-Glucose Deprivation. International Journal of Molecular Sciences, 2020, 21, 8053.	4.1	9
25	The Detrimental Action of Adenosine on Glutamate-Induced Cytotoxicity in PC12 Cells Can Be Shifted towards a Neuroprotective Role through A1AR Positive Allosteric Modulation. Cells, 2020, 9, 1242.	4.1	12
26	Pharmacological data of cannabidiol- and cannabigerol-type phytocannabinoids acting on cannabinoid CB1, CB2 and CB1/CB2 heteromer receptors. Pharmacological Research, 2020, 159, 104940.	7.1	57
27	Structural investigation on thiazolo[5,4-d]pyrimidines to obtain dual-acting blockers of CD73 and adenosine A2A receptor as potential antitumor agents. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 127067.	2.2	12
28	Adenosinergic System Involvement in Ischemic Stroke Patientsâ€™ Lymphocytes. Cells, 2020, 9, 1072.	4.1	7
29	Modifications on the Amino-3,5-dicyanopyridine Core To Obtain Multifaceted Adenosine Receptor Ligands with Antineuropathic Activity. Journal of Medicinal Chemistry, 2019, 62, 6894-6912.	6.4	16
30	Amino-3,5-Dicyanopyridines Targeting the Adenosine Receptors. Ranging from Pan Ligands to Combined A1/A2B Partial Agonists. Pharmaceuticals, 2019, 12, 159.	3.8	9
31	Targeting A3 and A2A adenosine receptors in the fight against cancer. Expert Opinion on Therapeutic Targets, 2019, 23, 669-678.	3.4	32
32	Pulsed electromagnetic field and relief of hypoxiaâ€induced neuronal cell death: The signaling pathway. Journal of Cellular Physiology, 2019, 234, 15089-15097.	4.1	25
33	The aminopyridine-3,5-dicarbonitrile core for the design of new non-nucleoside-like agonists of the human adenosine A2B receptor. European Journal of Medicinal Chemistry, 2018, 150, 127-139.	5.5	30
34	Development of novel pyridazinone-based adenosine receptor ligands. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 1484-1489.	2.2	4
35	Effects of pulsed electromagnetic fields and platelet rich plasma in preventing osteoclastogenesis in an in vitro model of osteolysis. Journal of Cellular Physiology, 2018, 233, 2645-2656.	4.1	14
36	Identification of novel thiazolo[5,4-d]pyrimidine derivatives as human A1 and A2A adenosine receptor antagonists/inverse agonists. Bioorganic and Medicinal Chemistry, 2018, 26, 3688-3695.	3.0	14

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37	Role of Adenosine Receptors in Clinical Biophysics Based on Pulsed Electromagnetic Fields. , 2018, , 557-580.		1
38	Binding Thermodynamic Characteristics of Adenosine Receptor Ligands. , 2018, , 199-215.		1
39	Cannabigerol Action at Cannabinoid CB1 and CB2 Receptors and at CB1â€“CB2 Heteroreceptor Complexes. Frontiers in Pharmacology, 2018, 9, 632.	3.5	88
40	Pharmacology of Adenosine Receptors: The State of the Art. Physiological Reviews, 2018, 98, 1591-1625.	28.8	495
41	Structure-activity relationship studies and pharmacological characterization of N5-heteroarylalkyl-substituted-2-(2-furanyl)thiazolo[5,4-d]pyrimidine-5,7-diamine-based derivatives as inverse agonists at human A2A adenosine receptor. European Journal of Medicinal Chemistry, 2018, 155, 552-561.	5.5	12
42	Pathological overproduction: the bad side of adenosine. British Journal of Pharmacology, 2017, 174, 1945-1960.	5.4	94
43	The role of 5-arylalkylamino- and 5-piperazino- moieties on the 7-aminopyrazolo[4,3-d]pyrimidine core in affecting adenosine A₁ and A_{2A} receptor affinity and selectivity profiles. Journal of Enzyme Inhibition and Medicinal Chemistry, 2017, 32, 248-263.	5.2	14
44	Pharmacoâ€toxicological effects of the novel thirdâ€generation fluorinate synthetic cannabinoids, <scp>5Fâ€ADBINACA</scp>, <scp>ABâ€FUBINACA</scp>, and <scp>STSâ€135</scp> in mice. In vitro and in vivo studies. Human Psychopharmacology, 2017, 32, e2601.	1.5	40
45	Age-dependent dopamine transporter dysfunction and Serine129 phospho-Î±-synuclein overload in G2019S LRRK2 mice. Acta Neuropathologica Communications, 2017, 5, 22.	5.2	73
46	A 2B adenosine receptors stimulate IL-6 production in primary murine microglia through p38 MAPK kinase pathway. Pharmacological Research, 2017, 117, 9-19.	7.1	57
47	Biochemical and Pharmacological Role of A1 Adenosine Receptors and Their Modulation as Novel Therapeutic Strategy. Advances in Experimental Medicine and Biology, 2017, 1051, 193-232.	1.6	40
48	RAW 264.7 coâ€cultured with ultraâ€high molecular weight polyethylene particles spontaneously differentiate into osteoclasts: an in vitro model of periprosthetic osteolysis. Journal of Biomedical Materials Research - Part A, 2017, 105, 510-520.	4.0	16
49	Imidazo[1,2-a]pyrazin-8-amine core for the design of new adenosine receptor antagonists: Structural exploration to target the A3 and A2A subtypes. European Journal of Medicinal Chemistry, 2017, 125, 611-628.	5.5	17
50	Pulsed Electromagnetic Field Exposure Reduces Hypoxia and Inflammation Damage in Neuronâ€Like and Microglial Cells. Journal of Cellular Physiology, 2017, 232, 1200-1208.	4.1	55
51	Binding and Signaling Studies Disclose a Potential Allosteric Site for Cannabidiol in Cannabinoid CB2 Receptors. Frontiers in Pharmacology, 2017, 8, 744.	3.5	134
52	Inhibition of A2A Adenosine Receptor Signaling in Cancer Cells Proliferation by the Novel Antagonist TP455. Frontiers in Pharmacology, 2017, 8, 888.	3.5	48
53	Psychostimulant Effect of the Synthetic Cannabinoid JWH-018 and AKB48: Behavioral, Neurochemical, and Dopamine Transporter Scan Imaging Studies in Mice. Frontiers in Psychiatry, 2017, 8, 130.	2.6	36
54	Role and Function of A2A and A3 Adenosine Receptors in Patients with Ankylosing Spondylitis, Psoriatic Arthritis and Rheumatoid Arthritis. International Journal of Molecular Sciences, 2017, 18, 697.	4.1	46

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55	Adenosine Receptors as a Biological Pathway for the Anti-Inflammatory and Beneficial Effects of Low Frequency Low Energy Pulsed Electromagnetic Fields. Mediators of Inflammation, 2017, 2017, 1-11.	3.0	63
56	Anxiolytic properties of A1 adenosine receptor PAMs. Oncotarget, 2017, 8, 7216-7217.	1.8	14
57	Exploring the 2- and 5-positions of the pyrazolo[4,3-d]pyrimidin-7-amino scaffold to target human A1 and A2A adenosine receptors. Bioorganic and Medicinal Chemistry, 2016, 24, 2794-2808.	3.0	14
58	A2A adenosine receptor upregulation correlates with disease activity in patients with systemic lupus erythematosus. Arthritis Research and Therapy, 2016, 18, 192.	3.5	30
59	Effect of the novel synthetic cannabinoids AKB48 and 5F-AKB48 on â€œtetradâ€• sensorimotor, neurological and neurochemical responses in mice. In vitro and in vivo pharmacological studies. Psychopharmacology, 2016, 233, 3685-3709.	3.1	63
60	Positive allosteric modulation of A1 adenosine receptors as a novel and promising therapeutic strategy for anxiety. Neuropharmacology, 2016, 111, 283-292.	4.1	33
61	Design, Synthesis, and Pharmacological Characterization of 2-(2-Furanyl)thiazolo[5,4-d]pyrimidine-5,7-diamine Derivatives: New Highly Potent A_{2A} Adenosine Receptor Inverse Agonists with Antinociceptive Activity. Journal of Medicinal Chemistry, 2016, 59, 10564-10576.	6.4	49
62	Repeated Dosing with NCX1404, a Nitric Oxide-Donating Pregabalin, Re-establishes Normal Nociceptive Responses in Mice with Streptozotocin-Induced Painful Diabetic Neuropathy. Journal of Pharmacology and Experimental Therapeutics, 2016, 357, 240-247.	2.5	4
63	Effect of JWH-250, JWH-073 and their interaction on â€œtetradâ€• sensorimotor, neurological and neurochemical responses in mice. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2016, 67, 31-50.	4.8	62
64	Synthesis and structure activity relationship investigation of triazolo[1,5-a]pyrimidines as CB2 cannabinoid receptor inverse agonists. European Journal of Medicinal Chemistry, 2016, 113, 11-27.	5.5	36
65	Structural refinement of pyrazolo[4,3- d]pyrimidine derivatives to obtain highly potent and selective antagonists for the human A 3 adenosine receptor. European Journal of Medicinal Chemistry, 2016, 108, 117-133.	5.5	18
66	One-Pot Reaction To Obtain N,Nâ€²-Disubstituted Guanidines of Pyrazolo[4,3-e][1,2,4]triazolo[1,5-c]pyrimidine Scaffold as Human A3 Adenosine Receptor Antagonists. Journal of Medicinal Chemistry, 2015, 58, 5355-5360.	6.4	9
67	Synthesis and biological evaluation of a new series of 2-amino-3-aroyl thiophene derivatives as agonist allosteric modulators of the A 1 adenosine receptor. A position-dependent effect study. European Journal of Medicinal Chemistry, 2015, 101, 185-204.	5.5	13
68	Exploring the 7-oxo-thiazolo[5,4-d]pyrimidine core for the design of new human adenosine A3 receptor antagonists. Synthesis, molecular modeling studies and pharmacological evaluation. European Journal of Medicinal Chemistry, 2015, 96, 105-121.	5.5	23
69	Novel halogenated derivates of JWH-018: Behavioral and binding studies in mice. Neuropharmacology, 2015, 95, 68-82.	4.1	81
70	Design, Synthesis, and Biological Evaluation of Novel 2-((2-(4-(Substituted)phenylpiperazin-1-yl)ethyl)amino)-5â€²-N-ethylcarboxamidoadenosines as Potent and Selective Agonists of the A_{2A} Adenosine Receptor. Journal of Medicinal Chemistry, 2015, 58, 3253-3267.	6.4	15
71	The A₃Adenosine Receptor: History and Perspectives. Pharmacological Reviews, 2015, 67, 74-102.	16.0	204
72	Synthesis and biological evaluation of novel 2-amino-3-aroyl-4-neopentyl-5-substituted thiophene derivatives as allosteric enhancers of the A1 adenosine receptor. Bioorganic and Medicinal Chemistry, 2014, 22, 148-166.	3.0	12

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73	A2Aadenosine receptors and Parkinson's disease severity. Acta Neurologica Scandinavica, 2014, 129, 276-281.	2.1	23
74	7-Amino-2-phenylpyrazolo[4,3-d]pyrimidine derivatives: Structural investigations at the 5-position to target human A1 and A2A adenosine receptors. Molecular modeling and pharmacological studies. European Journal of Medicinal Chemistry, 2014, 84, 614-627.	5.5	22
75	Expression, pharmacology and functional activity of adenosine A1 receptors in genetic models of Huntington's disease. Neurobiology of Disease, 2014, 71, 193-204.	4.4	22
76	Synthesis and Biological Evaluation of Novel Allosteric Enhancers of the A₁ Adenosine Receptor Based on 2-Amino-3-(4â€²-Chlorobenzoyl)-4-Substituted-5-Arylethynyl Thiophene. Journal of Medicinal Chemistry, 2014, 57, 7673-7686.	6.4	26
77	TRR469, a potent A1 adenosine receptor allosteric modulator, exhibits anti-nociceptive properties in acute and neuropathic pain models in mice. Neuropharmacology, 2014, 81, 6-14.	4.1	59
78	Synthesis and biological effects of novel 2-amino-3-(4-chlorobenzoyl)-4-substituted thiophenes as allosteric enhancers ofÂthe A1 adenosine receptor. European Journal of Medicinal Chemistry, 2013, 67, 409-427.	5.5	17
79	The stimulation of A3 adenosine receptors reduces bone-residing breast cancer in a rat preclinical model. European Journal of Cancer, 2013, 49, 482-491.	2.8	40
80	Design, Synthesis, and Pharmacological Properties of New Heteroarylpyridine/Heteroarylpyrimidine Derivatives as CB₂ Cannabinoid Receptor Partial Agonists. Journal of Medicinal Chemistry, 2013, 56, 1098-1112.	6.4	16
81	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. Journal of Medicinal Chemistry, 2013, 56, 2256-2269.	6.4	24
82	Antinociceptive effects of the selective CB2 agonist MT178 in inflammatory and chronic rodent pain models. Pain, 2013, 154, 864-873.	4.2	56
83	Adenosine and adenosine receptors in rheumatoid arthritis. International Journal of Clinical Rheumatology, 2013, 8, 13-25.	0.3	5
84	Functional Tissue Engineering in Articular Cartilage Repair: Is There a Role for Electromagnetic Biophysical Stimulation?. Tissue Engineering - Part B: Reviews, 2013, 19, 353-367.	4.8	51
85	Discovery of 7-Oxopyrazolo[1,5-a]pyrimidine-6-carboxamides as Potent and Selective CB₂ Cannabinoid Receptor Inverse Agonists. Journal of Medicinal Chemistry, 2013, 56, 4482-4496.	6.4	24
86	Multiple sclerosis lymphocytes upregulate <scp>A</scp>_{2A} adenosine receptors that are antiinflammatory when stimulated. European Journal of Immunology, 2013, 43, 2206-2216.	2.9	45
87	Pyrazolo[1,5-c]quinazoline derivatives and their simplified analogues as adenosine receptor antagonists: Synthesis, structureâ€“affinity relationships and molecular modeling studies. Bioorganic and Medicinal Chemistry, 2013, 21, 283-294.	3.0	43
88	A_{2A}adenosine receptors are up-regulated in lymphocytes from amyotrophic lateral sclerosis patients. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 406-413.	1.7	34
89	A2A Adenosine Receptors Are Differentially Modulated by Pharmacological Treatments in Rheumatoid Arthritis Patients and Their Stimulation Ameliorates Adjuvant-Induced Arthritis in Rats. PLoS ONE, 2013, 8, e54195.	2.5	43
90	Pulsed Electromagnetic Fields Increased the Anti-Inflammatory Effect of A2A and A3 Adenosine Receptors in Human T/C-28a2 Chondrocytes and hFOB 1.19 Osteoblasts. PLoS ONE, 2013, 8, e65561.	2.5	106

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91	7-Oxo-[1,4]oxazino[2,3,4-ij]quinoline-6-carboxamides as Selective CB₂ Cannabinoid Receptor Ligands: Structural Investigations around a Novel Class of Full Agonists. Journal of Medicinal Chemistry, 2012, 55, 6608-6623.	6.4	36
92	Synthesis and Biological Evaluation of 2-Amino-3-(4-chlorobenzoyl)-4-[(4-arylpiperazin-1-yl)methyl]-5-substituted-thiophenes. Effect of the 5-Modification on Allosteric Enhancer Activity at the A1 Adenosine Receptor. Journal of Medicinal Chemistry, 2012, 55, 7719-7735.	6.4	27
93	A Novel Conjugated Agent between Dopamine and an A_{2A} Adenosine Receptor Antagonist as a Potential Anti-Parkinson Multitarget Approach. Molecular Pharmaceutics, 2012, 9, 591-604.	4.6	29
94	The Anti-Tumor Effect of A3 Adenosine Receptors Is Potentiated by Pulsed Electromagnetic Fields in Cultured Neural Cancer Cells. PLoS ONE, 2012, 7, e39317.	2.5	39
95	Effect of pulsed electromagnetic field exposure on adenosine receptors in rat brain. Bioelectromagnetics, 2012, 33, 279-287.	1.6	31
96	Structureâ€“activity relationships of 2-amino-3-aroyl-4-[(4-arylpiperazin-1-yl)methyl]thiophenes. Part 2: Probing the influence of diverse substituents at the phenyl of the arylpiperazine moiety on allosteric enhancer activity at the A1 adenosine receptor. Bioorganic and Medicinal Chemistry, 2012, 20, 996-1007.	3.0	14
97	Electromagnetic fields (EMFs) and adenosine receptors modulate prostaglandin E₂ and cytokine release in human osteoarthritic synovial fibroblasts. Journal of Cellular Physiology, 2012, 227, 2461-2469.	4.1	85
98	A2A and A3 adenosine receptor expression in rheumatoid arthritis: upregulation, inverse correlation with disease activity score and suppression of inflammatory cytokine and metalloproteinase release. Arthritis Research and Therapy, 2011, 13, R197.	3.5	113
99	Synthesis, structureâ€“affinity relationships, and molecular modeling studies of novel pyrazolo[3,4-c]quinoline derivatives as adenosine receptor antagonists. Bioorganic and Medicinal Chemistry, 2011, 19, 3757-3768.	3.0	21
100	A₃ Receptors Are Overexpressed in Pleura from Patients with Mesothelioma and Reduce Cell Growth via Akt/Nuclear Factor-ÎºB Pathway. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 522-530.	5.6	44
101	The role of adenosine receptors in rheumatoid arthritis. Autoimmunity Reviews, 2010, 10, 61-64.	5.8	40
102	Expression and functional role of adenosine receptors in regulating inflammatory responses in human synoviocytes. British Journal of Pharmacology, 2010, 160, 101-115.	5.4	63
103	P2X₁ and P2X₃ Purinergic Receptors Differentially Modulate the Inflammatory Response in Human Osteoarthritic Synovial Fibroblasts. Cellular Physiology and Biochemistry, 2010, 25, 325-336.	1.6	15
104	A _{2A} adenosine receptor overexpression and functionality, as well as TNFâ€Î± levels, correlate with motor symptoms in Parkinson's disease. FASEB Journal, 2010, 24, 587-598.	0.5	107
105	Oxidative/nitrosative stress selectively altered A_{2B}adenosine receptors in chronic obstructive pulmonary disease. FASEB Journal, 2010, 24, 1192-1204.	0.5	15
106	Adenosine analogs and electromagnetic fields inhibit prostaglandin E2Ârelease in bovine synovial fibroblasts. Osteoarthritis and Cartilage, 2009, 17, 252-262.	1.3	79
107	Normalization of A_{2A} and A₃ adenosine receptor upâ€regulation in rheumatoid arthritis patients by treatment with antiâ€“tumor necrosis factor Î± but not methotrexate. Arthritis and Rheumatism, 2009, 60, 2880-2891.	6.7	74
108	Novel potent and highly selective human A3 adenosine receptor antagonists belonging to the 4-amido-2-arylpyrazolo[3,4-c]quinoline series: Molecular docking analysis and pharmacological studies. Bioorganic and Medicinal Chemistry, 2009, 17, 401-410.	3.0	21

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109	Deficiency of polycystic kidney disease-1 gene (PKD1) expression increases A3 adenosine receptors in human renal cells: Implications for cAMP-dependent signalling and proliferation of PKD1-mutated cystic cells. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2009, 1792, 531-540.	3.8	22
110	2-Phenylpyrazolo[4,3-d]pyrimidin-7-one as a New Scaffold To Obtain Potent and Selective Human A₃ Adenosine Receptor Antagonists: New Insights into the Receptorâ"Antagonist Recognition. Journal of Medicinal Chemistry, 2009, 52, 7640-7652.	6.4	51
111	Binding thermodynamic characterization of human P2X1 and P2X3 purinergic receptors. Biochemical Pharmacology, 2008, 75, 1198-1208.	4.4	9
112	Characterization of adenosine receptors in bovine chondrocytes and fibroblast-like synoviocytes exposed to low frequency low energy pulsed electromagnetic fields. Osteoarthritis and Cartilage, 2008, 16, 292-304.	1.3	110
113	Pharmacological characterization of P2X1 and P2X3 purinergic receptors in bovine chondrocytes. Osteoarthritis and Cartilage, 2008, 16, 1421-1429.	1.3	39
114	Synthesis and Biological Evaluation of 2-Amino-3-(4-Chlorobenzoyl)-4-[N-(Substituted) Piperazin-1-yl]Thiophenes as Potent Allosteric Enhancers of the A_₁ Adenosine Receptor. Journal of Medicinal Chemistry, 2008, 51, 5875-5879.	6.4	46
115	Scouting Human A3 Adenosine Receptor Antagonist Binding Mode Using a Molecular Simplification Approach: From Triazoloquinoxaline to a Pyrimidine Skeleton as a Key Study. Journal of Medicinal Chemistry, 2007, 50, 6596-6606.	6.4	30
116	Alteration of Adenosine Receptors in Patients with Chronic Obstructive Pulmonary Disease. American Journal of Respiratory and Critical Care Medicine, 2006, 173, 398-406.	5.6	101
117	Synthesis and biological characterization of [3H] (2-amino-4,5,6,7-tetrahydrobenzo[b]thiophen-3-yl)-(4-chlorophenyl)-methanone, the first radiolabelled adenosine A1 allosteric enhancer. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 1402-1404.	2.2	14
118	ST 1535: a preferential A2A adenosine receptor antagonist. International Journal of Neuropsychopharmacology, 2006, 9, 575.	2.1	37
119	Pharmacological characterization of novel adenosine ligands in recombinant and native human A2B receptors. Biochemical Pharmacology, 2005, 70, 1601-1612.	4.4	53