

Richard J Evans

List of Publications by Year in descending order

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73
papers

4,506
citations

87888

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docs citations

73
times ranked

2625
citing authors

#	ARTICLE	IF	CITATIONS
1	A new class of ligand-gated ion channel defined by P2X receptor for extracellular ATP. <i>Nature</i> , 1994, 371, 516-519.	27.8	1,005
2	A Role of the Fast ATP-gated P2X1 Cation Channel in Thrombosis of Small Arteries In Vivo. <i>Journal of Experimental Medicine</i> , 2003, 198, 661-667.	8.5	191
3	The Role of Positively Charged Amino Acids in ATP Recognition by Human P2X1 Receptors. <i>Journal of Biological Chemistry</i> , 2000, 275, 29361-29367.	3.4	170
4	Molecular properties of P2X receptors. <i>Pflugers Archiv European Journal of Physiology</i> , 2006, 452, 486-500.	2.8	145
5	Physiological role for P2X1 receptors in renal microvascular autoregulatory behavior. <i>Journal of Clinical Investigation</i> , 2003, 112, 1895-1905.	8.2	144
6	Conserved Cysteine Residues in the Extracellular Loop of the Human P2X1 Receptor Form Disulfide Bonds and Are Involved in Receptor Trafficking to the Cell Surface. <i>Molecular Pharmacology</i> , 2002, 61, 303-311.	2.3	137
7	ADP is not an agonist at P2X1 receptors: evidence for separate receptors stimulated by ATP and ADP on human platelets. <i>British Journal of Pharmacology</i> , 2000, 131, 108-114.	5.4	130
8	P2X1 Receptor-Deficient Mice Establish the Native P2X Receptor and a P2Y6-Like Receptor in Arteries. <i>Molecular Pharmacology</i> , 2002, 62, 1438-1445.	2.3	112
9	Molecular properties of ATP-gated P2X receptor ion channels. <i>Trends in Pharmacological Sciences</i> , 2004, 25, 487-493.	8.7	110
10	Synaptic P2X receptors. <i>Current Opinion in Neurobiology</i> , 2001, 11, 378-386.	4.2	109
11	Disruption of Lipid Rafts Inhibits P2X1 Receptor-mediated Currents and Arterial Vasoconstriction. <i>Journal of Biological Chemistry</i> , 2005, 280, 30705-30711.	3.4	91
12	Functional Characterization of a P2X Receptor from <i>Schistosoma mansoni</i> . <i>Journal of Biological Chemistry</i> , 2004, 279, 41650-41657.	3.4	90
13	ATP Binding at Human P2X1 Receptors. <i>Journal of Biological Chemistry</i> , 2004, 279, 9043-9055.	3.4	88
14	P2X1 Ion Channels Promote Neutrophil Chemotaxis through Rho Kinase Activation. <i>Journal of Immunology</i> , 2009, 183, 2801-2809.	0.8	84
15	Inhibition of Platelet Functions and Thrombosis through Selective or Nonselective Inhibition of the Platelet P2 Receptors with Increasing Doses of NF449 [4,4'-bis(4-((2-(Carboxylbis(imino-5,1,3-benzenetriylbis-(carbonylimino)))tetrakis-benzene-1,3-disulfonic Acid Octasodium Salt)). <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2005, 314, 232-243.	2.5	79
16	A study of P2X1 receptor function in murine megakaryocytes and human platelets reveals synergy with P2Y receptors. <i>British Journal of Pharmacology</i> , 2002, 135, 363-372.	5.4	73
17	Emerging roles for P2X1 receptors in platelet activation. <i>Platelets</i> , 2004, 15, 131-144.	2.3	71
18	Functional Regulation of P2X6 Receptors by N-Linked Glycosylation: Identification of a Novel β -Methylene ATP-Sensitive Phenotype. <i>Molecular Pharmacology</i> , 2004, 65, 979-985.	2.3	69

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19	G-protein-coupled receptor regulation of P2X1 receptors does not involve direct channel phosphorylation. <i>Biochemical Journal</i> , 2004, 382, 101-110.	3.7	66
20	Agonist-stimulated internalisation of the ligand-gated ion channel P2X1 in rat vas deferens. <i>FEBS Letters</i> , 2001, 489, 154-158.	2.8	62
21	Interplay between P2Y1, P2Y12, and P2X1 receptors in the activation of megakaryocyte cation influx currents by ADP: evidence that the primary megakaryocyte represents a fully functional model of platelet P2 receptor signaling. <i>Blood</i> , 2005, 106, 1644-1651.	1.4	62
22	P2X1 expressed on polymorphonuclear neutrophils and platelets is required for thrombosis in mice. <i>Blood</i> , 2014, 124, 2575-2585.	1.4	58
23	Unique residues in the ATP gated human P2X7 receptor define a novel allosteric binding pocket for the selective antagonist AZ10606120. <i>Scientific Reports</i> , 2017, 7, 725.	3.3	58
24	Cysteine Substitution Mutants Give Structural Insight and Identify ATP Binding and Activation Sites at P2X Receptors. <i>Journal of Neuroscience</i> , 2007, 27, 4072-4082.	3.6	55
25	Direct Voltage Control of Signaling via P2Y1 and Other G \pm q-coupled Receptors. <i>Journal of Biological Chemistry</i> , 2005, 280, 1490-1498.	3.4	52
26	Structural interpretation of P2X receptor mutagenesis studies on drug action. <i>British Journal of Pharmacology</i> , 2010, 161, 961-971.	5.4	52
27	Cysteine Substitution Mutagenesis and the Effects of Methanethiosulfonate Reagents at P2X2 and P2X4 Receptors Support a Core Common Mode of ATP Action at P2X Receptors. <i>Journal of Biological Chemistry</i> , 2008, 283, 20126-20136.	3.4	51
28	The P2X1 receptor and platelet function. <i>Purinergic Signalling</i> , 2011, 7, 341-356.	2.2	51
29	Agonist binding evokes extensive conformational changes in the extracellular domain of the ATP-gated human P2X1 receptor ion channel. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 4663-4667.	7.1	51
30	The P2X1 Receptor Is Required for Neutrophil Extravasation during Lipopolysaccharide-Induced Lethal Endotoxemia in Mice. <i>Journal of Immunology</i> , 2015, 194, 739-749.	0.8	49
31	Heterogeneity of P2X Receptors in Sympathetic Neurons: Contribution of Neuronal P2X1 Receptors Revealed Using Knockout Mice. <i>Molecular Pharmacology</i> , 2004, 65, 139-148.	2.3	46
32	Regulation of Human Recombinant P2X3 Receptors by Ecto-Protein Kinase C. <i>Journal of Neuroscience</i> , 2005, 25, 7734-7742.	3.6	46
33	Lipid Raft Association and Cholesterol Sensitivity of P2X1-4 Receptors for ATP. <i>Journal of Biological Chemistry</i> , 2010, 285, 32770-32777.	3.4	46
34	Contribution of conserved polar glutamine, asparagine and threonine residues and glycosylation to agonist action at human P2X1 receptors for ATP. <i>Journal of Neurochemistry</i> , 2006, 96, 843-852.	3.9	44
35	ATP-Gated P2X Receptor Channels: Molecular Insights into Functional Roles. <i>Annual Review of Physiology</i> , 2019, 81, 43-62.	13.1	44
36	Evidence for P2Y1, P2Y2, P2Y6 and atypical UTP-sensitive receptors coupled to rises in intracellular calcium in mouse cultured superior cervical ganglion neurons and glia. <i>British Journal of Pharmacology</i> , 2004, 143, 525-532.	5.4	40

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37	P2X receptor subtype-specific modulation of excitatory and inhibitory synaptic inputs in the rat brainstem. <i>Journal of Physiology</i> , 2004, 558, 745-757.	2.9	40
38	P2X1 Receptor Subunit Contribution to Gating Revealed by a Dominant Negative PKC Mutant. <i>Biochemical and Biophysical Research Communications</i> , 2002, 291, 611-616.	2.1	38
39	Differential sensitivity of human platelet P2X1 and P2Y1 receptors to disruption of lipid rafts. <i>Biochemical and Biophysical Research Communications</i> , 2006, 343, 415-419.	2.1	38
40	Cysteine Scanning Mutagenesis (Residues Glu52â€“Gly96) of the Human P2X1 Receptor for ATP. <i>Journal of Biological Chemistry</i> , 2011, 286, 29207-29217.	3.4	37
41	Lack of evidence for functional ADP-activated human P2X1 receptors supports a role for ATP during hemostasis and thrombosis. <i>Blood</i> , 2003, 102, 3646-3651.	1.4	34
42	The Intracellular Amino Terminus Plays a Dominant Role in Desensitization of ATP-gated P2X Receptor Ion Channels. <i>Journal of Biological Chemistry</i> , 2011, 286, 44691-44701.	3.4	31
43	Contribution of the region Glu181 to Val200 of the extracellular loop of the human P2X1 receptor to agonist binding and gating revealed using cysteine scanning mutagenesis¹. <i>Journal of Neurochemistry</i> , 2009, 109, 1042-1052.	3.9	30
44	P2X1 receptor mobility and trafficking; regulation by receptor insertion and activation. <i>Journal of Neurochemistry</i> , 2010, 113, 1177-1187.	3.9	30
45	Heat Shock Protein 90 Inhibitors Reduce Trafficking of ATP-gated P2X1 Receptors and Human Platelet Responsiveness*. <i>Journal of Biological Chemistry</i> , 2012, 287, 32747-32754.	3.4	27
46	Contribution of the Juxtatransmembrane Intracellular Regions to the Time Course and Permeation of ATP-gated P2X7 Receptor Ion Channels. <i>Journal of Biological Chemistry</i> , 2015, 290, 14556-14566.	3.4	26
47	Identification of Human P2X1 Receptor-interacting Proteins Reveals a Role of the Cytoskeleton in Receptor Regulation. <i>Journal of Biological Chemistry</i> , 2011, 286, 30591-30599.	3.4	25
48	Mapping the Allosteric Action of Antagonists A740003 and A438079 Reveals a Role for the Left Flipper in Ligand Sensitivity at P2X7 Receptors. <i>Molecular Pharmacology</i> , 2018, 93, 553-562.	2.3	25
49	Mapping the Site of Action of Human P2X7 Receptor Antagonists AZ11645373, Brilliant Blue G, KN-62, Calmidazolium, and ZINC58368839 to the Intersubunit Allosteric Pocket. <i>Molecular Pharmacology</i> , 2019, 96, 355-363.	2.3	25
50	Conserved Negatively Charged Residues Are Not Required for ATP Action at P2X1 Receptors. <i>Biochemical and Biophysical Research Communications</i> , 2001, 289, 700-704.	2.1	21
51	Contribution of conserved glycine residues to ATP action at human P2X1 receptors: mutagenesis indicates that the glycine at position 250 is important for channel function. <i>Journal of Neurochemistry</i> , 2005, 95, 1746-1754.	3.9	21
52	Ca²⁺ Influx through P2X1 Receptors Amplifies P2Y1 Receptor-Evoked Ca²⁺ Signaling and ADP-Evoked Platelet Aggregation. <i>Molecular Pharmacology</i> , 2014, 86, 243-251.	2.3	20
53	Mapping the binding site of the P2X receptor antagonist PPADS reveals the importance of orthosteric site charge and the cysteine-rich head region. <i>Journal of Biological Chemistry</i> , 2018, 293, 12820-12831.	3.4	19
54	Regions of the amino terminus of the P2X₁ receptor required for modification by phorbol ester and mGluR1± receptors. <i>Journal of Neurochemistry</i> , 2009, 108, 331-340.	3.9	18

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55	Characterisation of ATP analogues to cross-link and label P2X receptors. <i>Neuropharmacology</i> , 2009, 56, 230-236.	4.1	18
56	Extracellular Ca ²⁺ modulates ADP-evoked aggregation through altered agonist degradation: implications for conditions used to study P2Y receptor activation. <i>British Journal of Haematology</i> , 2011, 153, 83-91.	2.5	18
57	An evaluation of antibody detection of the P2X1 receptor subunit in the CNS of wild type and P2X1-knockout mice. <i>Neuroscience Letters</i> , 2006, 397, 120-125.	2.1	14
58	P2X Receptor Chimeras Highlight Roles of the Amino Terminus to Partial Agonist Efficacy, the Carboxyl Terminus to Recovery from Desensitization, and Independent Regulation of Channel Transitions. <i>Journal of Biological Chemistry</i> , 2013, 288, 21412-21421.	3.4	14
59	Kinetics of Conformational Changes Revealed by Voltage-Clamp Fluorometry Give Insight to Desensitization at ATP-Gated Human P2X1 Receptors. <i>Molecular Pharmacology</i> , 2014, 86, 707-715.	2.3	14
60	Mechanistic insights from resolving ligand-dependent kinetics of conformational changes at ATP-gated P2X1R ion channels. <i>Scientific Reports</i> , 2016, 6, 32918.	3.3	14
61	Mass spectrometry analysis of human P2X1 receptors; insight into phosphorylation, modelling and conformational changes. <i>Journal of Neurochemistry</i> , 2012, 123, 725-735.	3.9	12
62	Calcium Signalling through Ligand-Gated Ion Channels such as P2X1 Receptors in the Platelet and other Non-Excitable Cells. <i>Advances in Experimental Medicine and Biology</i> , 2016, 898, 305-329.	1.6	10
63	P2X1 ion channel deficiency causes massive bleeding in inflamed intestine and increases thrombosis. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 44-56.	3.8	9
64	Mutagenesis studies of conserved proline residues of human P2X1 receptors for ATP indicate that proline 272 contributes to channel function. <i>Journal of Neurochemistry</i> , 2005, 92, 1256-1264.	3.9	8
65	Contribution of the intracellular C terminal domain to regulation of human P2X1 receptors for ATP by phorbol ester and Gq coupled mGlu1± receptors. <i>European Journal of Pharmacology</i> , 2011, 654, 155-159.	3.5	7
66	Use of Chimeras, Point Mutants, and Molecular Modeling to Map the Antagonist-binding Site of 4,4'-((Carboxylbis-(imino-5,1,3-benzenetriylbis(carboxylimino)))tetrakisbenzene-1,3-disulfonic Acid (NF449) at P2X1 Receptors for ATP. <i>Journal of Biological Chemistry</i> , 2015, 290, 1559-1569.	3.4	7
67	Organization of ATP-gated P2X1 receptor intracellular termini in apo and desensitized states. <i>Journal of General Physiology</i> , 2019, 151, 146-155.	1.9	6
68	Cooperation between NMDA-Type Glutamate and P2 Receptors for Neuroprotection during Stroke: Combining Astrocyte and Neuronal Protection. <i>Neuroglia (Basel, Switzerland)</i> , 2018, 1, 30-47.	0.9	5
69	Development of a P2X1-eYFP receptor knock-in mouse to track receptors in real time. <i>Purinergic Signalling</i> , 2019, 15, 397-402.	2.2	5
70	Contribution of P2X1 receptor intracellular basic residues to channel properties. <i>Biochemical and Biophysical Research Communications</i> , 2006, 350, 244-248.	2.1	3
71	Permeability and single-channel properties of mesenteric, basilar, and septal (coronary) artery smooth-muscle P2X receptors. <i>Drug Development Research</i> , 2001, 52, 164-169.	2.9	2
72	Contribution of P2Y1 receptors to ADP signalling in mouse spinal cord cultures. <i>Neuroscience Letters</i> , 2008, 435, 190-193.	2.1	2

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73	Identification of a distinct desensitisation gate in the ATP-gated P2X2 receptor. Biochemical and Biophysical Research Communications, 2020, 523, 190-195.	2.1	2