The IUPHAR/BPS Guide to PHARMACOLOGY: an exper and their ligands

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Citation Report

#	Article	IF	Citations
1	<scp> L</scp> â€ 1^2 and <scp> L</scp> â€ 18 : inflammatory markers or mediators of hypertension?. British Journal of Pharmacology, 2014, 171, 5589-5602.	2.7	168
2	Alteration of vascular reactivity in heart failure: role of phosphodiesterases 3 and 4. British Journal of Pharmacology, 2014, 171, 5361-5375.	2.7	19
3	Structural basis for constitutive activity and agonistâ€induced activation of the enteroendocrine fat sensor <scp>GPR</scp> 119. British Journal of Pharmacology, 2014, 171, 5774-5789.	2.7	23
4	The ceramide kinase inhibitor <scp>NVP</scp> â€231 inhibits breast and lung cancer cell proliferation by inducing <scp>M</scp> phase arrest and subsequent cell death. British Journal of Pharmacology, 2014, 171, 5829-5844.	2.7	56
5	Different apoptotic effects of [<scp><i>O< i>< scp><i>O< i><scp><i>O< i><scp><i>o< i><scp><i>i>o< i>< scp><i>i>o< i>< scp><i>i>o< i>o< i>o< i>o< i>o< i>o< i>o< i</i></i></i></scp></i></scp></i></scp></i></i></scp>	ɔ>â€acac) 2.7	(<scp>DMS<!--</td--></scp>
6	Therapeutic use of botulinum toxin in migraine: mechanisms of action. British Journal of Pharmacology, 2014, 171, 4177-4192.	2.7	78
7	Epigenetic pathway targets for the treatment of disease: accelerating progress in the development of pharmacological tools: <scp>IUPHAR</scp> Review 11. British Journal of Pharmacology, 2014, 171, 4981-5010.	2.7	23
8	Lipoxin <scp>A</scp> ₄ suppresses the development of endometriosis in an <scp>ALX</scp> receptorâ€dependent manner via the p38 <scp>MAPK</scp> pathway. British Journal of Pharmacology, 2014, 171, 4927-4940.	2.7	44
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10	<scp>CFTR</scp> potentiators partially restore channel function to <scp>A</scp> 561 <scp>E</scp> â€CFTR, a cystic fibrosis mutant with a similar mechanism of dysfunction as <scp>F</scp> 508delâ€ <scp>CFTR</scp> . British Journal of Pharmacology, 2014, 171, 4490-4503.	2.7	23
11	An <i>in vivo</i> role for <scp>R</scp> ho kinase activation in the tumour vascular disrupting activity of combretastatin <scp>A</scp> â€4 3â€ <scp><i>O</i></scp> â€phosphate. British Journal of Pharmacology, 2014, 171, 4902-4913.	2.7	14
12	Buprenorphine signalling is compromised at the <scp>N</scp> 40 <scp>D</scp> polymorphism of the human ν opioid receptor <i>iin vitro. British Journal of Pharmacology, 2014, 171, 4273-4288.</i>	2.7	24
13	<scp>C</scp> elastrol protects ischaemic myocardium through a heat shock response with upâ€regulation of haeme oxygenaseâ€1. British Journal of Pharmacology, 2014, 171, 5265-5279.	2.7	52
14	Quercetin attenuates doxorubicin cardiotoxicity by modulating <scp>B</scp> miâ€1 expression. British Journal of Pharmacology, 2014, 171, 4440-4454.	2.7	107
15	Purinergic neuromuscular transmission in the gastrointestinal tract; functional basis for future clinical and pharmacological studies. British Journal of Pharmacology, 2014, 171, 4360-4375.	2.7	36
16	Novel coumarin modified <scp>GLP</scp> â€1 derivatives with enhanced plasma stability and prolonged <i>in vivo</i> glucoseâ€lowering ability. British Journal of Pharmacology, 2014, 171, 5252-5264.	2.7	41
17	Identification of an old antibiotic clofoctol as a novel activator of unfolded protein response pathways and an inhibitor of prostate cancer. British Journal of Pharmacology, 2014, 171, 4478-4489.	2.7	27
18	Fluoxetine elevates allopregnanolone in female rat brain but inhibits a steroid microsomal dehydrogenase rather than activating an aldoâ€keto reductase. British Journal of Pharmacology, 2014, 171, 5870-5880.	2.7	29

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20	Multiple roles of the <scp>PGE</scp> ₂ â€ <scp>EP</scp> receptor signal in vascular permeability. British Journal of Pharmacology, 2014, 171, 4879-4889.	2.7	50
21	4â€bromopropofol decreases action potential generation in spinal neurons by inducing a glycine receptorâ€mediated tonic conductance. British Journal of Pharmacology, 2014, 171, 5790-5801.	2.7	5
23	Pharmacological manipulations in animal models of anorexia and binge eating in relation to humans. British Journal of Pharmacology, 2014, 171, 4767-4784.	2.7	20
24	Molecular basis of agonist docking in a human <scp>GPR</scp> 103 homology model by siteâ€directed mutagenesis and structureâ€"activity relationship studies. British Journal of Pharmacology, 2014, 171, 4425-4439.	2.7	13
25	<scp>PAR</scp> 1â€dependent <scp>COX</scp> â€2/ <scp>PGE₂</scp> production contributes to cell proliferation via <scp>EP₂</scp> receptors in primary human cardiomyocytes. British Journal of Pharmacology, 2014, 171, 4504-4519.	2.7	19
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30	Molecular targets of the multifunctional ironâ€chelating drug, <scp>M</scp> 30, in the brains of mouse models of type 2 diabetes mellitus. British Journal of Pharmacology, 2014, 171, 5636-5649.	2.7	9
31	Emodin inhibits tonic tension through suppressing <scp>PKCÎ′</scp> â€mediated inhibition of myosin phosphatase in rat isolated thoracic aorta. British Journal of Pharmacology, 2014, 171, 4300-4310.	2.7	27
32	Ulinastatin activates haem oxygenase 1 antioxidant pathway and attenuates allergic inflammation. British Journal of Pharmacology, 2014, 171, 4399-4412.	2.7	10
33	Pharmacological bronchodilation is partially mediated by reduced airway wall stiffness. British Journal of Pharmacology, 2014, 171, 4376-4384.	2.7	19
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39	Molecular mechanism of allosteric modulation at <scp>GPCRs</scp> : insight from a binding kinetics study at the human <scp>A</scp> ₁ adenosine receptor. British Journal of Pharmacology, 2014, 171, 5295-5312.	2.7	20
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41	Nicotinic acetylcholine receptors control acetylcholine and noradrenaline release in the rodent habenuloâ€interpeduncular complex. British Journal of Pharmacology, 2014, 171, 5209-5224.	2.7	20
42	Identification of novel insulin mimetic drugs by quantitative total internal reflection fluorescence (<scp>TIRF</scp>) microscopy. British Journal of Pharmacology, 2014, 171, 5237-5251.	2.7	28
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50	$$$ < scp>PKC inhibition results in a < scp>K< sub>v1.5 + < scp>K< sub>v ^21.3 pharmacology closer to < scp>K< sub>v1.5 channels. British Journal of Pharmacology, 2014, 171, 4914-4926.$	2.7	3
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