Masaaki Sato

List of Publications by Year in descending order

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MASAAKI SATO

#	Article	IF	CITATIONS
1	Ligandâ€directed signalling at βâ€adrenoceptors. British Journal of Pharmacology, 2010, 159, 1022-1038.	5.4	141
2	Glucose uptake in brown fat cells is dependent on mTOR complex 2–promoted GLUT1 translocation. Journal of Cell Biology, 2014, 207, 365-374.	5.2	138
3	Improving Type 2 Diabetes Through a Distinct Adrenergic Signaling Pathway Involving mTORC2 That Mediates Clucose Uptake in Skeletal Muscle. Diabetes, 2014, 63, 4115-4129.	0.6	101
4	Ligand-Directed Signaling at the β ₃ -Adrenoceptor Produced by 3-(2-Ethylphenoxy)-1-[(1, <i>S</i>)-1,2,3,4-tetrahydronapth-1-ylamino]-2 <i>S</i> -2-propanol oxalate (SR59230A) Relative to Receptor Agonists. Molecular Pharmacology, 2007, 72, 1359-1368.	2.3	80
5	4-[[(Hexylamino)carbonyl]amino]-N-[4-[2-[[(2S)-2-hydroxy-3-(4-hydroxyphenoxy)propyl]amino]ethyl]-phenyl]-ben (L755507) and Antagonist (S)-N-[4-[2-[[3-[3-(Acetamidomethyl)phenoxy]-2-hydroxypropyl]amino]-ethyl]phenyl]benzenesulfonamide (L748337) Activate Different Signaling Pathways in Chinese Hamster Ovary-K1 Cells Stably Expressing	zenesulfo 2.3	namide 47
6	the Human 123-Adrenoceptor. Molecular Pharmacology, 2008, 74, 1417-1428. Could burning fat start with a brite spark? Pharmacological and nutritional ways to promote thermogenesis. Molecular Nutrition and Food Research, 2016, 60, 18-42.	3.3	39
7	The PPARÎ ³ agonist rosiglitazone promotes the induction of brite adipocytes, increasing β-adrenoceptor-mediated mitochondrial function and glucose uptake. Cellular Signalling, 2018, 42, 54-66.	3.6	38
8	Adrenoceptors promote glucose uptake into adipocytes and muscle by an insulin-independent signaling pathway involving mechanistic target of rapamycin complex 2. Pharmacological Research, 2017, 116, 87-92.	7.1	30
9	Functional Domains of the Mouse β3-Adrenoceptor Associated with Differential G Protein Coupling. Journal of Pharmacology and Experimental Therapeutics, 2005, 315, 1354-1361.	2.5	25
10	β ₂ â€Adrenoceptors increase translocation of GLUT4 via GPCR kinase sites in the receptor Câ€ŧerminal tail. British Journal of Pharmacology, 2012, 165, 1442-1456.	5.4	25
11	Rosiglitazone and a β3-Adrenoceptor Agonist Are Both Required for Functional Browning of White Adipocytes in Culture. Frontiers in Endocrinology, 2018, 9, 249.	3.5	25
12	Factors influencing biased agonism in recombinant cells expressing the human α _{1A} â€adrenoceptor. British Journal of Pharmacology, 2017, 174, 2318-2333.	5.4	24
13	Interaction with Caveolin-1 Modulates G Protein Coupling of Mouse β3-Adrenoceptor. Journal of Biological Chemistry, 2012, 287, 20674-20688.	3.4	23
14	α 1A -Adrenoceptors activate mTOR signalling and glucose uptake in cardiomyocytes. Biochemical Pharmacology, 2018, 148, 27-40.	4.4	20
15	BRL37344 stimulates GLUT4 translocation and glucose uptake in skeletal muscle via β ₂ -adrenoceptors without causing classical receptor desensitization. American Journal of Physiology - Regulatory Integrative and Comparative Physiology, 2019, 316, R666-R677.	1.8	16
16	Adrenoceptor regulation of the mechanistic target of rapamycin in muscle and adipose tissue. British Journal of Pharmacology, 2019, 176, 2433-2448.	5.4	9
17	The metabolic effects of mirabegron are mediated primarily by β 3 â€adrenoceptors. Pharmacology Research and Perspectives, 2020, 8, e00643.	2.4	9
18	Response to Comment on Sato et al. Improving Type 2 Diabetes Through a Distinct Adrenergic Signaling Pathway Involving mTORC2 That Mediates Glucose Uptake in Skeletal Muscle. Diabetes 2014;63:4115–4129. Diabetes, 2014, 63, e22-e23.	0.6	7

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19	GPR55 regulates the responsiveness to, but does not dimerise with, α1A-adrenoceptors. Biochemical Pharmacology, 2021, 188, 114560.	4.4	0
20	α _{1A} -adrenoceptor stimulation promotes glucose uptake and cell survival in cardiomyocytes - role of mTOR. Proceedings for Annual Meeting of the Japanese Pharmacological Society, 2018, WCP2018, PO1-2-28.	0.0	0
21	Metabolic effects of mirabegron in mice: implications for use in diabetes. Proceedings for Annual Meeting of the Japanese Pharmacological Society, 2018, WCP2018, PO1-5-25.	0.0	Ο