

Sai P Pydi

List of Publications by Year in descending order

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papers

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361413

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951
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#	ARTICLE	IF	CITATIONS
1	Amino Acid Derivatives as Bitter Taste Receptor (T2R) Blockers. <i>Journal of Biological Chemistry</i> , 2014, 289, 25054-25066.	3.4	78
2	Structural Basis of Activation of Bitter Taste Receptor T2R1 and Comparison with Class A G-protein-coupled Receptors (GPCRs). <i>Journal of Biological Chemistry</i> , 2011, 286, 36032-36041.	3.4	74
3	Bitter taste receptor T2R1 is activated by dipeptides and tripeptides. <i>Biochemical and Biophysical Research Communications</i> , 2010, 398, 331-335.	2.1	67
4	Abscisic Acid Acts as a Blocker of the Bitter Taste G Protein-Coupled Receptor T2R4. <i>Biochemistry</i> , 2015, 54, 2622-2631.	2.5	49
5	Adipocyte β -arrestin-2 is essential for maintaining whole body glucose and energy homeostasis. <i>Nature Communications</i> , 2019, 10, 2936.	12.8	43
6	Constitutively active mutant gives novel insights into the mechanism of bitter taste receptor activation. <i>Journal of Neurochemistry</i> , 2012, 122, 537-544.	3.9	36
7	Characterization of the Direct Interaction between Hybrid Sensor Kinases PA1611 and RetS That Controls Biofilm Formation and the Type III Secretion System in <i>Pseudomonas aeruginosa</i> . <i>ACS Infectious Diseases</i> , 2017, 3, 162-175.	3.8	35
8	The third intracellular loop plays a critical role in bitter taste receptor activation. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2014, 1838, 231-236.	2.6	34
9	Lack of adipocyte purinergic P2Y ₆ receptor greatly improves whole body glucose homeostasis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 30763-30774.	7.1	34
10	Hepatic Gi signaling regulates whole-body glucose homeostasis. <i>Journal of Clinical Investigation</i> , 2018, 128, 746-759.	8.2	34
11	Dextromethorphan Mediated Bitter Taste Receptor Activation in the Pulmonary Circuit Causes Vasoconstriction. <i>PLoS ONE</i> , 2014, 9, e110373.	2.5	33
12	Adipocyte Gi signaling is essential for maintaining whole-body glucose homeostasis and insulin sensitivity. <i>Nature Communications</i> , 2020, 11, 2995.	12.8	27
13	Recent Advances in Structure and Function Studies on Human Bitter Taste Receptors. <i>Current Protein and Peptide Science</i> , 2012, 13, 501-508.	1.4	26
14	Use of DREADD Technology to Identify Novel Targets for Antidiabetic Drugs. <i>Annual Review of Pharmacology and Toxicology</i> , 2021, 61, 421-440.	9.4	26
15	Selective activation of Gs signaling in adipocytes causes striking metabolic improvements in mice. <i>Molecular Metabolism</i> , 2019, 27, 83-91.	6.5	25
16	Serum thrombospondin 2 is a novel predictor for the severity in the patients with NAFLD. <i>Liver International</i> , 2021, 41, 505-514.	3.9	25
17	Metabolic roles of G protein-coupled receptor signaling in obesity and type 2 diabetes. <i>FEBS Journal</i> , 2021, 288, 2622-2644.	4.7	25
18	New Insights into Structural Determinants for Prostanoid Thromboxane A2 Receptor- and Prostacyclin Receptor-G Protein Coupling. <i>Molecular and Cellular Biology</i> , 2013, 33, 184-193.	2.3	23

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19	Metabolic Functions of G Protein-Coupled Receptors in Hepatocytesâ€”Potential Applications for Diabetes and NAFLD. <i>Biomolecules</i> , 2020, 10, 1445.	4.0	23
20	Constitutive Activity of Bitter Taste Receptors (T2Rs). <i>Advances in Pharmacology</i> , 2014, 70, 303-326.	2.0	22
21	Cholesterol modulates bitter taste receptor function. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2016, 1858, 2081-2087.	2.6	22
22	Beneficial metabolic role of Î²-arrestin-1 expressed by AgRP neurons. <i>Science Advances</i> , 2020, 6, eaaz1341.	10.3	17
23	Site-Directed Mutations and the Polymorphic Variant Ala160Thr in the Human Thromboxane Receptor Uncover a Structural Role for Transmembrane Helix 4. <i>PLoS ONE</i> , 2012, 7, e29996.	2.5	16
24	Î²-arrestin-1 suppresses myogenic reprogramming of brown fat to maintain euglycemia. <i>Science Advances</i> , 2020, 6, eaba1733.	10.3	15
25	Adipocyte P2Y14 receptors play a key role in regulating whole-body glucose and lipid homeostasis. <i>JCI Insight</i> , 2021, 6, .	5.0	15
26	Î²-Arrestins as Important Regulators of Glucose and Energy Homeostasis. <i>Annual Review of Physiology</i> , 2022, 84, 17-40.	13.1	14
27	Î²-Arrestin-1 is required for adaptive Î²-cell mass expansion during obesity. <i>Nature Communications</i> , 2021, 12, 3385.	12.8	13
28	Adipocyte Gq signaling is a regulator of glucose and lipid homeostasis in mice. <i>Nature Communications</i> , 2022, 13, 1652.	12.8	13
29	Role of rhodopsin N-terminus in structure and function of rhodopsin-bitter taste receptor chimeras. <i>Biochemical and Biophysical Research Communications</i> , 2013, 430, 179-182.	2.1	9
30	Chemogenetic approaches to identify metabolically important GPCR signaling pathways: Therapeutic implications. <i>Journal of Neurochemistry</i> , 2021, 158, 603-620.	3.9	8
31	2â€Step PLT16â€AST44 method: Simplified liver fibrosis detection system in patients with nonâ€alcoholic fatty liver disease. <i>Hepatology Research</i> , 2022, 52, 352-363.	3.4	8
32	Key Metabolic Functions of Î²-Arrestins: Studies with Novel Mouse Models. <i>Trends in Endocrinology and Metabolism</i> , 2021, 32, 118-129.	7.1	7
33	Identification of a high affinity selective inhibitor of Polo-like kinase 1 for cancer chemotherapy by computational approach. <i>Journal of Molecular Graphics and Modelling</i> , 2014, 51, 104-112.	2.4	4
34	Expression and Role of Î²3-Adrenergic Receptor during the Differentiation of 3T3-L1 Preadipocytes into Adipocytes. <i>Biology</i> , 2022, 11, 772.	2.8	3
35	Serine Phosphorylation: An Important Post-translational Modification For Functional Regulation Of Smooth Muscle Thromboxane Receptor. , 2010, , .		0
36	32-OR: Receptor-Mediated Gq Signaling in Adipocytes as a Critical Modulator of Systemic Glucose and Lipid Homeostasis. <i>Diabetes</i> , 2021, 70, 32-OR.	0.6	0

#	ARTICLE	IF	CITATIONS
37	STRUCTURE & FUNCTION ANALYSIS OF THE LIGAND BINDING POCKET OF BITTER TASTE RECEPTOR T2R4. FASEB Journal, 2013, 27, 883.6.	0.5	0
38	Adipocyte-Selective Deletion of β -Arrestin-1 in Mice Causes Adiposity, Impaired Glucose Tolerance, and Reduced Insulin Sensitivity. Diabetes, 2018, 67, .	0.6	0
39	1797-P: β -arrestin-1 in AgRP Neurons Plays Crucial Role in Maintaining Whole Body Glucose Homeostasis. Diabetes, 2019, 68, 1797-P.	0.6	0
40	38-OR: Beta-Arrestin-1 Regulates Pancreatic Beta-Cell Function and Plays a Key Role in Maintaining Whole Body Glucose Homeostasis. Diabetes, 2019, 68, 38-OR.	0.6	0
41	1781-P: Adipocyte GI Signaling Regulates Whole-Body Glucose Homeostasis and Insulin Sensitivity. Diabetes, 2019, 68, 1781-P.	0.6	0
42	1971-P: Activation of Adipocyte Gq Signaling Causes Improved Whole-Body Glucose Homeostasis. Diabetes, 2020, 69, 1971-P.	0.6	0
43	2129-P: Glucagon Secretion from Pancreatic Islets Is Regulated by Beta-Arrestin-1. Diabetes, 2020, 69, .	0.6	0
44	1694-P: Beta-Arrestin 1 Suppresses Myogenic Reprogramming of Brown Fat to Maintain Euglycemia. Diabetes, 2020, 69, 1694-P.	0.6	0
45	β -Arrestins as regulators of key metabolic processes. , 2022, , 69-85.		0