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List of Publications by Year
in descending order

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141 papers	13,808 citations	25034 57 h-index	21540 114 g-index
142 all docs	142 docs citations	142 times ranked	12156 citing authors

#	Article	IF	Citations
1	Glucagon-like peptide 1 (GLP-1). Molecular Metabolism, 2019, 30, 72-130.	6.5	850
2	Ghrelin. Molecular Metabolism, 2015, 4, 437-460.	6.5	810
3	Truncated glucagonâ€like peptide I, an insulinâ€releasing hormone from the distal gut. FEBS Letters, 1987, 211, 169-174.	2.8	498
4	Somatostatin cell processes as pathways for paracrine secretion. Science, 1979, 205, 1393-1395.	12.6	497
5	High Constitutive Signaling of the Ghrelin Receptorâ€"Identification of a Potent Inverse Agonist. Molecular Endocrinology, 2003, 17, 2201-2210.	3.7	455
6	GPR41/FFAR3 and GPR43/FFAR2 as Cosensors for Short-Chain Fatty Acids in Enteroendocrine Cells vs FFAR3 in Enteric Neurons and FFAR2 in Enteric Leukocytes. Endocrinology, 2013, 154, 3552-3564.	2.8	436
7	GPCR-Mediated Signaling of Metabolites. Cell Metabolism, 2017, 25, 777-796.	16.2	403
8	International Union of Basic and Clinical Pharmacology. XCIV. Adhesion G Proteinâ€"Coupled Receptors. Pharmacological Reviews, 2015, 67, 338-367.	16.0	392
9	MOLECULAR MECHANISM OF 7TM RECEPTOR ACTIVATIONâ€"A GLOBAL TOGGLE SWITCH MODEL. Annual Review of Pharmacology and Toxicology, 2006, 46, 481-519.	9.4	382
10	GPR39 Signaling Is Stimulated by Zinc Ions But Not by Obestatin. Endocrinology, 2007, 148, 13-20.	2.8	371
11	Ligand binding and micro-switches in 7TM receptor structures. Trends in Pharmacological Sciences, 2009, 30, 249-259.	8.7	310
12	Common Structural Basis for Constitutive Activity of the Ghrelin Receptor Family. Journal of Biological Chemistry, 2004, 279, 53806-53817.	3.4	303
13	Locating ligand-binding sites in 7tm receptors by protein engineering. Current Opinion in Biotechnology, 1994, 5, 434-444.	6.6	290
14	A Major Lineage of Enteroendocrine Cells Coexpress CCK, Secretin, GIP, GLP-1, PYY, and Neurotensin but Not Somatostatin. Endocrinology, 2012, 153, 5782-5795.	2.8	269
15	Seven transmembrane G protein-coupled receptor repertoire of gastric ghrelin cells. Molecular Metabolism, 2013, 2, 376-392.	6.5	261
16	Different binding epitopes on the NK1 receptor for substance P and a non-peptide antagonist. Nature, 1993, 362, 345-348.	27.8	241
17	The processing of peptide precursors. FEBS Letters, 1986, 200, 1-10.	2.8	235
18	snRNA-seq reveals a subpopulation of adipocytes that regulates thermogenesis. Nature, 2020, 587, 98-102.	27.8	221

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19	Expression of the short chain fatty acid receptor GPR41/FFAR3 in autonomic and somatic sensory ganglia. Neuroscience, 2015, 290, 126-137.	2.3	192
20	Y1 and Y2 receptors for neuropeptide Y. FEBS Letters, 1989, 245, 209-214.	2.8	178
21	GPR40 (FFAR1) â€“ Combined Gs and Gq signaling inÂvitro is associated with robust incretin secretagogue action exÂvivo and inÂvivo. Molecular Metabolism, 2015, 4, 3-14.	6.5	175
22	Conversion of antagonist-binding site to metal-ion site in the tachykinin NK-1 receptor. Nature, 1995, 374, 74-77.	27.8	173
23	GPR119 as a fat sensor. Trends in Pharmacological Sciences, 2012, 33, 374-381.	8.7	165
24	Allosteric enhancers, allosteric agonists and ago-allosteric modulators: where do they bind and how do they act?. Trends in Pharmacological Sciences, 2007, 28, 366-373.	8.7	161
25	Enterochromaffin 5-HT cells â€“ A major target for GLP-1 and gut microbial metabolites. Molecular Metabolism, 2018, 11, 70-83.	6.5	160
26	Biosynthesis of peptide precursors and protease inhibitors using new constitutive and inducible eukaryotic expression vectors. FEBS Letters, 1990, 267, 289-294.	2.8	159
27	A Highly Selective Cc Chemokine Receptor (Ccr)8 Antagonist Encoded by the Poxvirus Molluscum Contagiosum. Journal of Experimental Medicine, 2000, 191, 171-180.	8.5	144
28	Regional Distribution of Putative NPY Y1Receptors and Neurons Expressing Y1mRNA in Forebrain Areas of the Rat Central Nervous System. European Journal of Neuroscience, 1993, 5, 1622-1637.	2.6	140
29	The Melanocortin-4 Receptor Is Expressed in Enteroendocrine L Cells and Regulates the Release of Peptide YY and Glucagon-like Peptide 1 InÂVivo. Cell Metabolism, 2014, 20, 1018-1029.	16.2	139
30	A Gut Feeling for Obesity: 7TM Sensors on Enteroendocrine Cells. Cell Metabolism, 2008, 8, 447-449.	16.2	128
31	Is there a â€˜lockâ€™ for all agonist â€˜keysâ€™ in 7TM receptors?. Trends in Pharmacological Sciences, 1996, 17, 213-216.	8.7	127
32	A Conserved Aromatic Lock for the Tryptophan Rotameric Switch in TM-VI of Seven-transmembrane Receptors. Journal of Biological Chemistry, 2010, 285, 3973-3985.	3.4	126
33	Profiling of G protein-coupled receptors in vagal afferents reveals novel gut-to-brain sensing mechanisms. Molecular Metabolism, 2018, 12, 62-75.	6.5	124
34	Enteroendocrine cell types revisited. Current Opinion in Pharmacology, 2013, 13, 912-921.	3.5	123
35	Neurotensin Is Coexpressed, Coreleased, and Acts Together With GLP-1 and PYY in Enteroendocrine Control of Metabolism. Endocrinology, 2016, 157, 176-194.	2.8	119
36	Effects of PYY_{3â€“36} and GLP-1 on energy intake, energy expenditure, and appetite in overweight men. American Journal of Physiology - Endocrinology and Metabolism, 2014, 306, E1248-E1256.	3.5	114

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37	Conversion of agonist site to metal-ion chelator site in the beta 2-adrenergic receptor. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 12322-12327.	7.1	110
38	Differentiation between binding sites for angiotensin II and nonpeptide antagonists on the angiotensin II type 1 receptors.. Proceedings of the National Academy of Sciences of the United States of America, 1994, 91, 7046-7050.	7.1	109
39	Molecular Mechanism of Action of Monocyclam Versus Bicyclam Non-peptide Antagonists in the CXCR4 Chemokine Receptor. Journal of Biological Chemistry, 2007, 282, 27354-27365.	3.4	104
40	Septide and neurokinin A are high-affinity ligands on the NK-1 receptor: evidence from homologous versus heterologous binding analysis. FEBS Letters, 1996, 399, 264-266.	2.8	99
41	Molecular Pharmacological Phenotyping of EBI2. Journal of Biological Chemistry, 2006, 281, 13199-13208.	3.4	98
42	Virally encoded 7TM receptors. Oncogene, 2001, 20, 1582-1593.	5.9	92
43	Ghrelin receptor mutations -- too little height and too much hunger. Journal of Clinical Investigation, 2006, 116, 637-641.	8.2	92
44	Mutational Evidence for a Common Î² Antagonist Binding Pocket in the Wild-Type Î² and Mutant Î¼ [K303E] Opioid Receptorsâ€. Journal of Medicinal Chemistry, 1998, 41, 4911-4914.	6.4	91
45	Metal Ion Site Engineering Indicates a Global Toggle Switch Model for Seven-transmembrane Receptor Activation. Journal of Biological Chemistry, 2006, 281, 17337-17346.	3.4	88
46	Two Active Molecular Phenotypes of the Tachykinin NK1 Receptor Revealed by G-protein Fusions and Mutagenesis. Journal of Biological Chemistry, 2001, 276, 19793-19799.	3.4	82
47	G Protein-Coupled Receptor 39 Deficiency Is Associated with Pancreatic Islet Dysfunction. Endocrinology, 2009, 150, 2577-2585.	2.8	82
48	Construction of a High Affinity Zinc Switch in the k-Opioid Receptor. Journal of Biological Chemistry, 1996, 271, 7875-7878.	3.4	79
49	Ago-Allosteric Modulation and Other Types of Allostery in Dimeric 7TM Receptors. Journal of Receptor and Signal Transduction Research, 2006, 26, 107-128.	2.5	77
50	GPR119, a Major Enteroendocrine Sensor of Dietary Triglyceride Metabolites Coacting in Synergy With FFA1 (GPR40). Endocrinology, 2016, 157, 4561-4569.	2.8	77
51	Overlapping Binding Site for the Endogenous Agonist, Small-Molecule Agonists, and Ago-allosteric Modulators on the Ghrelin Receptor. Molecular Pharmacology, 2009, 75, 44-59.	2.3	66
52	Isolation and biogenesis of a new peptide from pancreatic islets. Nature, 1981, 294, 589-591.	27.8	65
53	Steric Hindrance Mutagenesis versus Alanine Scan in Mapping of Ligand Binding Sites in the Tachykinin NK1 Receptor. Molecular Pharmacology, 1998, 53, 166-175.	2.3	63
54	Partial Agonism through a Zinc-Ion Switch Constructed between Transmembrane Domains III and VII in the Tachykinin NK1 Receptor. Molecular Pharmacology, 2000, 58, 263-270.	2.3	61

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55	The adhesion G protein-coupled receptor G2 (ADGRG2/GPR64) constitutively activates SRE and NFÎºB and is involved in cell adhesion and migration. Cellular Signalling, 2015, 27, 2579-2588.	3.6	61
56	Extracellular succinate hyperpolarizes M2 macrophages through SUCNR1/GPR91-mediated Gq signaling. Cell Reports, 2021, 35, 109246.	6.4	61
57	Molecular mechanism of Zn²⁺ agonism in the extracellular domain of GPR39. FEBS Letters, 2008, 582, 2583-2588.	2.8	60
58	GluVII:06 - A Highly Conserved and Selective Anchor Point for Non-Peptide Ligands in Chemokine Receptors. Current Topics in Medicinal Chemistry, 2006, 6, 1319-1333.	2.1	60
59	Two nonpeptide tachykinin antagonists act through epitopes on corresponding segments of the NK1 and NK2 receptors.. Proceedings of the National Academy of Sciences of the United States of America, 1993, 90, 6194-6198.	7.1	59
60	Transcriptional and Functional Characterization of the G Protein-Coupled Receptor Repertoire of Gastric Somatostatin Cells. Endocrinology, 2015, 156, 3909-3923.	2.8	56
61	L-Cell Differentiation Is Induced by Bile Acids Through GPBAR1 and Paracrine GLP-1 and Serotonin Signaling. Diabetes, 2020, 69, 614-623.	0.6	54
62	Microbial regulation of the L cell transcriptome. Scientific Reports, 2018, 8, 1207.	3.3	52
63	Mapping Substance P Binding Sites on the Neurokinin-1 Receptor Using Genetic Incorporation of a Photoreactive Amino Acid. Journal of Biological Chemistry, 2014, 289, 18045-18054.	3.4	49
64	Binding of Norbinaltorphimine (norBNI) Congeners to Wild-Type and Mutant Mu and Kappa Opioid Receptors:â€‰ Molecular Recognition Loci for the Pharmacophore and Address Components of Kappa Antagonists. Journal of Medicinal Chemistry, 2000, 43, 1573-1576.	6.4	48
65	Molecular Interaction of a Potent Nonpeptide Agonist with the Chemokine Receptor CCR8. Molecular Pharmacology, 2007, 72, 327-340.	2.3	47
66	Conserved Water-mediated Hydrogen Bond Network between TM-I, -II, -VI, and -VII in 7TM Receptor Activation. Journal of Biological Chemistry, 2010, 285, 19625-19636.	3.4	45
67	Gq and Gs signaling acting in synergy to control GLP-1 secretion. Molecular and Cellular Endocrinology, 2017, 449, 64-73.	3.2	45
68	Human substance P receptor binding mode of the antagonist drug aprepitant by NMR and crystallography. Nature Communications, 2019, 10, 638.	12.8	43
69	The aromatic amino acid sensor GPR142 controls metabolism through balanced regulation of pancreatic and gut hormones. Molecular Metabolism, 2019, 19, 49-64.	6.5	43
70	Unique Interaction Pattern for a Functionally Biased Ghrelin Receptor Agonist. Journal of Biological Chemistry, 2011, 286, 20845-20860.	3.4	42
71	An Aromatic Region To Induce a Switch between Agonism and Inverse Agonism at the Ghrelin Receptor. Journal of Medicinal Chemistry, 2012, 55, 7437-7449.	6.4	42
72	Ligand Modulation of the Epstein-Barr Virus-induced Seven-transmembrane Receptor EBI2. Journal of Biological Chemistry, 2011, 286, 29292-29302.	3.4	41

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73	Inhibiting RHOA Signaling in Mice Increases Glucose Tolerance and Numbers of Enteroendocrine and Other Secretory Cells in the Intestine. Gastroenterology, 2018, 155, 1164-1176.e2.	1.3	41
74	Activation of the CXCR3 Chemokine Receptor through Anchoring of a Small Molecule Chelator Ligand between TM-III, -IV, and -VI. Molecular Pharmacology, 2007, 71, 930-941.	2.3	40
75	Biased signaling of lipids and allosteric actions of synthetic molecules for GPR119. Biochemical Pharmacology, 2016, 119, 66-75.	4.4	40
76	Receptor structure-based discovery of non-metabolite agonists for the succinate receptor GPR91. Molecular Metabolism, 2017, 6, 1585-1596.	6.5	40
77	G proteinâ€“coupled receptor modulation with pepducins: moving closer to the clinic. Annals of the New York Academy of Sciences, 2011, 1226, 34-49.	3.8	39
78	Research Resource: A Chromogranin A Reporter for Serotonin and Histamine Secreting Enteroendocrine Cells. Molecular Endocrinology, 2015, 29, 1658-1671.	3.7	39
79	Opposite Regulation of Ghrelin and Glucagon-like Peptide-1 by Metabolite G-Protein-Coupled Receptors. Trends in Endocrinology and Metabolism, 2016, 27, 665-675.	7.1	39
80	NK1 Receptor Fused to Î²-Arrestin Displays a Single-Component, High-Affinity Molecular Phenotype. Molecular Pharmacology, 2002, 62, 30-37.	2.3	38
81	GLP1- and GIP-producing cells rarely overlap and differ by bombesin receptor-2 expression and responsiveness. Journal of Endocrinology, 2016, 228, 39-48.	2.6	35
82	Molecular dynamics-guided discovery of an ago-allosteric modulator for GPR40/FFAR1. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 7123-7128.	7.1	35
83	Stable expression of high affinity NK1(substance P) and NK2(neurokinin A) receptors but low affinity NK3(neurokinin B) receptors in transfected CHO cells. FEBS Letters, 1992, 296, 241-244.	2.8	34
84	Conserved HisVI-17of the NK-1 receptor is involved in binding of non-peptide antagonists but not substance P. FEBS Letters, 1993, 336, 506-510.	2.8	34
85	Simultaneous recording of the gastro-entero-pancreatic hormonal peptide response to food in man. Metabolism: Clinical and Experimental, 1980, 29, 777-779.	3.4	33
86	A moving story of receptors. Nature, 2008, 455, 473-474.	27.8	33
87	Modulation of Constitutive Activity and Signaling Bias of the Ghrelin Receptor by Conformational Constraint in the Second Extracellular Loop. Journal of Biological Chemistry, 2012, 287, 33488-33502.	3.4	33
88	Evidence that a deviation in the kynurenine pathway aggravates atherosclerotic disease in humans. Journal of Internal Medicine, 2021, 289, 53-68.	6.0	33
89	Impaired Î±-carboxyamidation of gastrin in vitamin C-deficient guinea pigs. FEBS Letters, 1986, 196, 151-154.	2.8	32
90	Paracrine crosstalk between intestinal L- and D-cells controls secretion of glucagon-like peptide-1 in mice. American Journal of Physiology - Endocrinology and Metabolism, 2019, 317, E1081-E1093.	3.5	32

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91	PheVI:09 (Phe6.44) as a Sliding Microswitch in Seven-transmembrane (7TM) G Protein-coupled Receptor Activation. Journal of Biological Chemistry, 2012, 287, 43516-43526.	3.4	31
92	Biased Gs Versus Gq Proteins and Î²-Arrestin Signaling in the NK1 Receptor Determined by Interactions in the Water Hydrogen Bond Network. Journal of Biological Chemistry, 2015, 290, 24495-24508.	3.4	31
93	Why Warburg Works: Lactate Controls Immune Evasion through GPR81. Cell Metabolism, 2020, 31, 666-668.	16.2	31
94	The Arginine of the DRY Motif in Transmembrane Segment III Functions as a Balancing Micro-switch in the Activation of the Î²2-Adrenergic Receptor. Journal of Biological Chemistry, 2012, 287, 31973-31982.	3.4	30
95	The MicroRNA Repertoire in Enteroendocrine Cells: Identification of miR-375 as a Potential Regulator of the Enteroendocrine Lineage. Endocrinology, 2015, 156, 3971-3983.	2.8	29
96	Model-Based Discovery of Synthetic Agonists for the Zn²⁺-Sensing G-Protein-Coupled Receptor 39 (GPR39) Reveals Novel Biological Functions. Journal of Medicinal Chemistry, 2017, 60, 886-898.	6.4	29
97	Microbial fermentation of flaxseed fibers modulates the transcriptome of GPR41-expressing enteroendocrine cells and protects mice against diet-induced obesity. American Journal of Physiology - Endocrinology and Metabolism, 2019, 316, E453-E463.	3.5	29
98	Partial processing of the neuropeptide Y precursor in transfected CHO cells. FEBS Letters, 1990, 261, 101-105.	2.8	28
99	Disulfide Bridge Engineering in the Tachykinin NK1 Receptor. Biochemistry, 2000, 39, 667-675.	2.5	28
100	Nervous control of pancreatic endocrine secretion in pigs. Acta Physiologica Scandinavica, 1981, 111, 15-22.	2.2	26
101	Split-receptors in the tachykinin neurokinin-1 system. Mutational analysis of intracellular loop 3. FEBS Journal, 1998, 251, 217-226.	0.2	25
102	Mutational analysis of the interaction of the N- and C-terminal ends of angiotensin II with the rat AT1A receptor. British Journal of Pharmacology, 2000, 130, 1263-1268.	5.4	25
103	N -acyl taurines are endogenous lipid messengers that improve glucose homeostasis. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 24770-24778.	7.1	25
104	Molecular adaptations in human subcutaneous adipose tissue after ten weeks of endurance exercise training in healthy males. Journal of Applied Physiology, 2019, 126, 569-577.	2.5	25
105	Structural basis for GPCR signaling by small polar versus largeÂlipid metabolitesâ€"discovery of non-metabolite ligands. Current Opinion in Cell Biology, 2020, 63, 38-48.	5.4	24
106	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.	5.4	23
107	Oral 2â€oleyl glyceryl ether improves glucose tolerance in mice through the GPR119 receptor. BioFactors, 2016, 42, 665-673.	5.4	23
108	Mutations in transmembrane segment VIJ of the AT₁ receptor differentiate between closely related insurmountable and competitive angiotensin antagonists. British Journal of Pharmacology, 1994, 113, 331-333.	5.4	22

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109	Functional and genetic epidemiological characterisation of theFFAR4(GPR120) p.R270H variant in the Danish population. Journal of Medical Genetics, 2016, 53, 616-623.	3.2	20
110	Bidirectional GPR119 Agonism Requires Peptide YY and Glucose for Activity in Mouse and Human Colon Mucosa. Endocrinology, 2018, 159, 1704-1717.	2.8	19
111	Bioorthogonal Labeling of Ghrelin Receptor to Facilitate Studies of Ligand-Dependent Conformational Dynamics. Chemistry and Biology, 2015, 22, 1431-1436.	6.0	17
112	Getting from A to Bâ€"exploring the activation motifs of the class B adhesion G proteinâ€coupled receptor subfamily G member 4/GPR112. FASEB Journal, 2016, 30, 1836-1848.	0.5	17
113	Molecular Interaction of a Potent Nonpeptide Agonist with the Chemokine Receptor CCR8. Molecular Pharmacology, 2007, 72, 327-340.	2.3	17
114	Snapshot of a signalling complex. Nature, 2011, 477, 540-541.	27.8	16
115	Autocrine negative feedback regulation of lipolysis through sensing of NEFAs by FFAR4/GPR120 in WAT. Molecular Metabolism, 2020, 42, 101103.	6.5	16
116	EBI2 overexpression in mice leads to B1 B-cell expansion and chronic lymphocytic leukemiaâ€"like B-cell malignancies. Blood, 2017, 129, 866-878.	1.4	14
117	Full monty of family B GPCRs. Nature Chemical Biology, 2017, 13, 819-821.	8.0	14
118	Isolation of ovine pancreatic icosapeptide: a peptide product containing one cysteine residue. FEBS Letters, 1984, 168, 293-298.	2.8	13
119	High molecular weight PEGylation of human pancreatic polypeptide at position 22 improves stability and reduces food intake in mice. British Journal of Pharmacology, 2016, 173, 3208-3221.	5.4	13
120	Expression of human pancreatic polypeptide precursors from a dicistronic mRNA in mammalian cells. FEBS Letters, 1987, 219, 181-188.	2.8	11
121	Structure-Activity Investigations and Optimisations of Non-metabolite Agonists for the Succinate Receptor 1. Scientific Reports, 2018, 8, 10010.	3.3	11
122	Activation of metabolite receptor GPR91 promotes platelet aggregation and transcellular biosynthesis of leukotriene C4. Journal of Thrombosis and Haemostasis, 2020, 18, 976-984.	3.8	11
123	Adhesion receptor ADGRG2/GPR64 is in the GI-tract selectively expressed in mature intestinal tuft cells. Molecular Metabolism, 2021, 51, 101231.	6.5	11
124	Vagal afferent cholecystokinin receptor activation is required for glucagonâ€like peptideâ€1â€"induced satiation. Diabetes, Obesity and Metabolism, 2022, 24, 268-280.	4.4	11
125	Hyperinsulinemia Is Highly Associated With Markers of Hepatocytic Senescence in Two Independent Cohorts. Diabetes, 2022, 71, 1929-1936.	0.6	11
126	Implications of replacing peptide bonds in the COOHâ€terminal B chain domain of insulin by the Î¨(CH₂â€NH) linker. International Journal of Peptide and Protein Research, 1993, 42, 578-584.	0.1	10

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127	An Enteroendocrine Full Package Solution. Cell Metabolism, 2010, 11, 445-447.	16.2	10
128	Downregulation of 5-HT₇ Serotonin Receptors by the Atypical Antipsychotics Clozapine and Olanzapine. Role of Motifs in the C-Terminal Domain and Interaction with GASP-1. ACS Chemical Neuroscience, 2015, 6, 1206-1218.	3.5	10
129	Structureâ€based discovery of novel US28 small molecule ligands with different modes of action. Chemical Biology and Drug Design, 2017, 89, 289-296.	3.2	10
130	The HETE Is on FFAR1 and Pancreatic Islet Cells. Cell Metabolism, 2018, 27, 273-275.	16.2	10
131	Synthetic G proteinâ€coupled bile acid receptor agonists and bile acids act via basolateral receptors in ileal and colonic mucosa. Neurogastroenterology and Motility, 2020, 32, e13943.	3.0	10
132	Beta-Hydroxybutyrate Suppresses Hepatic Production of the Ghrelin Receptor Antagonist LEAP2. Endocrinology, 2022, 163, .	2.8	10
133	Protective succinate-SUCNR1 metabolic stress signaling gone bad. Cell Metabolism, 2021, 33, 1276-1278.	16.2	9
134	Preassociation between the 5â€HT ₇ serotonin receptor and G protein G _s : molecular determinants and association with low potency activation of adenylyl cyclase. FASEB Journal, 2019, 33, 3870-3886.	0.5	8
135	Activation of succinate receptor 1 boosts human mast cell reactivity and allergic bronchoconstriction. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 2677-2687.	5.7	7
136	Mutation-Guided Unbiased Modeling of the Fat Sensor GPR119 for High-Yield Agonist Screening. Structure, 2015, 23, 2377-2386.	3.3	6
137	Disruption of GPR35 Signaling in Bone Marrow-Derived Cells Does Not Influence Vascular Inflammation and Atherosclerosis in Hyperlipidemic Mice. Metabolites, 2021, 11, 411.	2.9	6
138	The Molecular Diversity of Vagal Afferents Revealed. Trends in Neurosciences, 2019, 42, 663-666.	8.6	5
139	Post-oral fat-induced satiation is mediated by endogenous CCK and GLP-1 in a fat self-administration mouse model. Physiology and Behavior, 2021, 234, 113315.	2.1	4
140	Construction of covalently coupled, concatameric dimers of 7TM receptors. Journal of Receptor and Signal Transduction Research, 2009, 29, 235-245.	2.5	3
141	The first PP supper--Camelot at Bispebjerg. Scandinavian Journal of Clinical and Laboratory Investigation, Supplement, 2001, 234, 109-21.	2.7	0