

Carsten Hoffmann

List of Publications by Year in descending order

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Version: 2024-02-01

52
papers

3,132
citations

218677

26
h-index

223800

46
g-index

56
all docs

56
docs citations

56
times ranked

3536
citing authors

#	ARTICLE	IF	CITATIONS
1	GPCR kinase knockout cells reveal the impact of individual GRKs on arrestin binding and GPCR regulation. <i>Nature Communications</i> , 2022, 13, 540.	12.8	54
2	Suitability of GRK Antibodies for Individual Detection and Quantification of GRK Isoforms in Western Blots. <i>International Journal of Molecular Sciences</i> , 2022, 23, 1195.	4.1	4
3	Phosphorylation of the D ₁ Dopamine Receptor by G Protein-Coupled Receptor Kinases: phosphorylation site identification and linkage to functional effects. <i>FASEB Journal</i> , 2021, 35, .	0.5	0
4	G protein-coupled receptor kinase 2 can enhance β -arrestin recruitment to the D ₂ dopamine receptor in the absence of receptor phosphorylation. <i>FASEB Journal</i> , 2021, 35, .	0.5	0
5	Differential Regulation of GPCRs—Are GRK Expression Levels the Key?. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 687489.	3.7	32
6	Ligand-Specific Allosteric Coupling Controls G-Protein-Coupled Receptor Signaling. <i>ACS Pharmacology and Translational Science</i> , 2020, 3, 859-867.	4.9	15
7	Advanced fluorescence microscopy reveals disruption of dynamic CXCR4 dimerization by subpocket-specific inverse agonists. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 29144-29154.	7.1	42
8	Kinetic Analysis of the Early Signaling Steps of the Human Chemokine Receptor CXCR4. <i>Molecular Pharmacology</i> , 2020, 98, 72-87.	2.3	13
9	Modulation of CXCR4-Mediated Gi1 Activation by EGF Receptor and GRK2. <i>ACS Pharmacology and Translational Science</i> , 2020, 3, 627-634.	4.9	3
10	Structural insight into small molecule action on Frizzleds. <i>Nature Communications</i> , 2020, 11, 414.	12.8	38
11	Molecular determinants of the mechanosensitivity of G protein-coupled receptors. <i>FASEB Journal</i> , 2020, 34, 1-1.	0.5	0
12	The Role of Orthosteric Building Blocks of Bitopic Ligands for Muscarinic M1 Receptors. <i>ACS Omega</i> , 2020, 5, 31706-31715.	3.5	6
13	Chemokine Receptor Crystal Structures: What Can Be Learned from Them?. <i>Molecular Pharmacology</i> , 2019, 96, 765-777.	2.3	25
14	Context-Dependent Signaling of CXC Chemokine Receptor 4 and Atypical Chemokine Receptor 3. <i>Molecular Pharmacology</i> , 2019, 96, 778-793.	2.3	30
15	Dishevelled-3 conformation dynamics analyzed by FRET-based biosensors reveals a key role of casein kinase 1. <i>Nature Communications</i> , 2019, 10, 1804.	12.8	20
16	Helix 8 is the essential structural motif of mechanosensitive GPCRs. <i>Nature Communications</i> , 2019, 10, 5784.	12.8	79
17	To sense or not to sense—new insights from GPCR-based and arrestin-based biosensors. <i>Current Opinion in Cell Biology</i> , 2019, 57, 16-24.	5.4	19
18	Structure-based exploration and pharmacological evaluation of N-substituted piperidin-4-yl-methanamine CXCR4 chemokine receptor antagonists. <i>European Journal of Medicinal Chemistry</i> , 2019, 162, 631-649.	5.5	12

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19	Muscarinic receptors promote pacemaker fate at the expense of secondary conduction system tissue in zebrafish. <i>JCI Insight</i> , 2019, 4, .	5.0	9
20	Lack of beta-arrestin signaling in the absence of active G proteins. <i>Nature Communications</i> , 2018, 9, 341.	12.8	297
21	Enhanced Fluorescence Resonance Energy Transfer in G-Protein-Coupled Receptor Probes on Nanocoated Microscopy Coverslips. <i>ACS Photonics</i> , 2018, 5, 2225-2233.	6.6	7
22	Intramolecular and Intermolecular FRET Sensors for GPCRs â€“ Monitoring Conformational Changes and Beyond. <i>Trends in Pharmacological Sciences</i> , 2018, 39, 123-135.	8.7	53
23	A split luciferase-based probe for quantitative proximal determination of GÎ±q signalling in live cells. <i>Scientific Reports</i> , 2018, 8, 17179.	3.3	16
24	FZD ₅ is a GÎ±q-coupled receptor that exhibits the functional hallmarks of prototypical GPCRs. <i>Science Signaling</i> , 2018, 11, .	3.6	46
25	FRET Studies of Quinolone-Based Bitopic Ligands and Their Structural Analogues at the Muscarinic M ₁ Receptor. <i>ACS Chemical Biology</i> , 2017, 12, 833-843.	3.4	17
26	A Photoswitchable Dualsteric Ligand Controlling Receptor Efficacy. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 7282-7287.	13.8	61
27	Ein photoschaltbarer Ligand zur Regulierung der Rezeptoraktivierung. <i>Angewandte Chemie</i> , 2017, 129, 7388-7393.	2.0	14
28	Functional and structural characterization of axonal opioid receptors as targets for analgesia. <i>Molecular Pain</i> , 2016, 12, 174480691662873.	2.1	22
29	Optical probes based on G proteinâ€“coupled receptors â€“ added work or added value?. <i>British Journal of Pharmacology</i> , 2016, 173, 255-266.	5.4	24
30	Î²-Arrestin biosensors reveal a rapid, receptor-dependent activation/deactivation cycle. <i>Nature</i> , 2016, 531, 661-664.	27.8	190
31	A New Generation of FRET Sensors for Robust Measurement of GÎ±1, GÎ±2 and GÎ±3 Activation Kinetics in Single Cells. <i>PLoS ONE</i> , 2016, 11, e0146789.	2.5	50
32	A Perspective on Studying G-Proteinâ€“Coupled Receptor Signaling with Resonance Energy Transfer Biosensors in Living Organisms. <i>Molecular Pharmacology</i> , 2015, 88, 589-595.	2.3	28
33	Ligand Residence Time at G-proteinâ€“Coupled Receptorsâ€“Why We Should Take Our Time To Study It. <i>Molecular Pharmacology</i> , 2015, 88, 552-560.	2.3	66
34	Arrestin Interactions with G Protein-Coupled Receptors. <i>Handbook of Experimental Pharmacology</i> , 2014, 219, 15-56.	1.8	62
35	Dynamic ligand binding dictates partial agonism at a G proteinâ€“coupled receptor. <i>Nature Chemical Biology</i> , 2014, 10, 18-20.	8.0	45
36	Comparison of the Activation Kinetics of the M ₃ Acetylcholine Receptor and a Constitutively Active Mutant Receptor in Living Cells. <i>Molecular Pharmacology</i> , 2012, 82, 236-245.	2.3	30

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37	The allosteric vestibule of a seven transmembrane helical receptor controls G-protein coupling. <i>Nature Communications</i> , 2012, 3, 1044.	12.8	117
38	Fluorescence/Bioluminescence Resonance Energy Transfer Techniques to Study G-Protein-Coupled Receptor Activation and Signaling. <i>Pharmacological Reviews</i> , 2012, 64, 299-336.	16.0	279
39	FRET-based sensors for the human M1-, M3-, and M5-acetylcholine receptors. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 1048-1054.	3.0	79
40	G Protein-Coupled Receptor Activation: Amino Acid Movements Caught Infrâ€Redâ€Handed. <i>ChemBioChem</i> , 2010, 11, 2247-2249.	2.6	0
41	Fluorescent labeling of tetracysteine-tagged proteins in intact cells. <i>Nature Protocols</i> , 2010, 5, 1666-1677.	12.0	192
42	A Fluorescence Resonance Energy Transfer-based M2 Muscarinic Receptor Sensor Reveals Rapid Kinetics of Allosteric Modulation. <i>Journal of Biological Chemistry</i> , 2010, 285, 8793-8800.	3.4	66
43	Differential Signaling of the Endogenous Agonists at the β 2-Adrenergic Receptor. <i>Journal of Biological Chemistry</i> , 2010, 285, 36188-36198.	3.4	101
44	Minireview: GPCR and G Proteins: Drug Efficacy and Activation in Live Cells. <i>Molecular Endocrinology</i> , 2009, 23, 590-599.	3.7	73
45	Fluorescence Resonance Energy Transfer Analysis of β 2-Adrenergic Receptor Activation Reveals Distinct Agonist-Specific Conformational Changes. <i>Molecular Pharmacology</i> , 2009, 75, 534-541.	2.3	103
46	Contribution of Fluorophores to Protein Kinase C FRET Probe Performance. <i>ChemBioChem</i> , 2008, 9, 1379-1384.	2.6	26
47	Conformational changes in G-protein-coupled receptorsâ€”the quest for functionally selective conformations is open. <i>British Journal of Pharmacology</i> , 2008, 153, S358-66.	5.4	68
48	Optical techniques to analyze real-time activation and signaling of G-protein-coupled receptors. <i>Trends in Pharmacological Sciences</i> , 2008, 29, 159-165.	8.7	119
49	Gq-coupled Receptor signaling â€” A kinetic analysis in living cells. <i>FASEB Journal</i> , 2008, 22, 722.1.	0.5	0
50	Direct Measurement Of Receptor/Gq Interaction. <i>FASEB Journal</i> , 2007, 21, A429.	0.5	0
51	A FLAsH-based FRET approach to determine G protein-coupled receptor activation in living cells. <i>Nature Methods</i> , 2005, 2, 171-176.	19.0	471
52	Using Intramolecular Fluorescence Resonance Energy Transfer to Study Receptor Conformation. , 0, , 133-146.		0