Laura M Bohn

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#	Paper	IF	Citations
112	Enhanced morphine analgesia in mice lacking beta-arrestin 2. <i>Science</i> , 1999 , 286, 2495-8	33.3	815
111	Mu-opioid receptor desensitization by beta-arrestin-2 determines morphine tolerance but not dependence. <i>Nature</i> , 2000 , 408, 720-3	50.4	709
110	Desensitization of G protein-coupled receptors and neuronal functions. <i>Annual Review of Neuroscience</i> , 2004 , 27, 107-44	17	677
109	Morphine side effects in beta-arrestin 2 knockout mice. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2005 , 314, 1195-201	4.7	427
108	Mice lacking the norepinephrine transporter are supersensitive to psychostimulants. <i>Nature Neuroscience</i> , 2000 , 3, 465-71	25.5	383
107	Crystal Structure of the Human Cannabinoid Receptor CB. Cell, 2016, 167, 750-762.e14	56.2	323
106	Bias Factor and Therapeutic Window Correlate to Predict Safer Opioid Analgesics. <i>Cell</i> , 2017 , 171, 1165	-56725.	e 13 9
105	Crystal structures of agonist-bound human cannabinoid receptor CB. <i>Nature</i> , 2017 , 547, 468-471	50.4	270
104	Differential mechanisms of morphine antinociceptive tolerance revealed in (beta)arrestin-2 knock-out mice. <i>Journal of Neuroscience</i> , 2002 , 22, 10494-500	6.6	218
103	Functional selectivity at the Eppioid receptor: implications for understanding opioid analgesia and tolerance. <i>Pharmacological Reviews</i> , 2011 , 63, 1001-19	22.5	194
102	Dopaminergic supersensitivity in G protein-coupled receptor kinase 6-deficient mice. <i>Neuron</i> , 2003 , 38, 291-303	13.9	194
101	An opioid agonist that does not induce mu-opioid receptorarrestin interactions or receptor internalization. <i>Molecular Pharmacology</i> , 2007 , 71, 549-57	4.3	187
100	Crystal Structure of the Human Cannabinoid Receptor CB2. <i>Cell</i> , 2019 , 176, 459-467.e13	56.2	175
99	Muscarinic supersensitivity and impaired receptor desensitization in G protein-coupled receptor kinase 5-deficient mice. <i>Neuron</i> , 1999 , 24, 1029-36	13.9	168
98	Biased agonism: An emerging paradigm in GPCR drug discovery. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 241-250	2.9	167
97	Enhanced rewarding properties of morphine, but not cocaine, in beta(arrestin)-2 knock-out mice. <i>Journal of Neuroscience</i> , 2003 , 23, 10265-73	6.6	167
96	Agonist-directed signaling of the serotonin 2A receptor depends on beta-arrestin-2 interactions in vivo. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008 , 105, 1079-8-	4 ^{11.5}	161

(2000-2015)

95	Fulfilling the Promise of "Biased" G Protein-Coupled Receptor Agonism. <i>Molecular Pharmacology</i> , 2015 , 88, 579-88	4.3	153
94	Relative opioid efficacy is determined by the complements of the G protein-coupled receptor desensitization machinery. <i>Molecular Pharmacology</i> , 2004 , 66, 106-12	4.3	138
93	Glutamatergic modulation of hyperactivity in mice lacking the dopamine transporter. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2001 , 98, 11047-54	11.5	135
92	The role of beta-arrestin2 in the severity of antinociceptive tolerance and physical dependence induced by different opioid pain therapeutics. <i>Neuropharmacology</i> , 2011 , 60, 58-65	5.5	134
91	Opioid receptor homo- and heterodimerization in living cells by quantitative bioluminescence resonance energy transfer. <i>Molecular Pharmacology</i> , 2005 , 67, 2173-84	4.3	131
90	Biased agonists of the kappa opioid receptor suppress pain and itch without causing sedation or dysphoria. <i>Science Signaling</i> , 2016 , 9, ra117	8.8	128
89	Sustained elevation of extracellular dopamine causes motor dysfunction and selective degeneration of striatal GABAergic neurons. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003 , 100, 11035-40	11.5	119
88	Requirement of receptor internalization for opioid stimulation of mitogen-activated protein kinase: biochemical and immunofluorescence confocal microscopic evidence. <i>Journal of Neuroscience</i> , 1999 , 19, 56-63	6.6	119
87	Serotonin, but not N-methyltryptamines, activates the serotonin 2A receptor via a Earrestin2/Src/Akt signaling complex in vivo. <i>Journal of Neuroscience</i> , 2010 , 30, 13513-24	6.6	112
86	Development of functionally selective, small molecule agonists at kappa opioid receptors. <i>Journal of Biological Chemistry</i> , 2013 , 288, 36703-16	5.4	109
85	Identification of Drosophila neuropeptide receptors by G protein-coupled receptors-beta-arrestin2 interactions. <i>Journal of Biological Chemistry</i> , 2003 , 278, 52172-8	5.4	103
84	Agonist-directed interactions with specific beta-arrestins determine mu-opioid receptor trafficking, ubiquitination, and dephosphorylation. <i>Journal of Biological Chemistry</i> , 2011 , 286, 31731-41	5.4	98
83	Potentiated opioid analgesia in norepinephrine transporter knock-out mice. <i>Journal of Neuroscience</i> , 2000 , 20, 9040-5	6.6	97
82	Activation and Signaling Mechanism Revealed by Cannabinoid Receptor-G Complex Structures. <i>Cell</i> , 2020 , 180, 655-665.e18	56.2	88
81	Drosophila CG8422 encodes a functional diuretic hormone receptor. <i>Journal of Experimental Biology</i> , 2004 , 207, 743-8	3	78
80	Enterestin2 regulates cannabinoid CB1 receptor signaling and adaptation in a central nervous system region-dependent manner. <i>Biological Psychiatry</i> , 2012 , 71, 714-24	7.9	76
79	Physiological and pharmacological implications of beta-arrestin regulation 2009 , 121, 285-93		76
78	Mitogenic signaling via endogenous kappa-opioid receptors in C6 glioma cells: evidence for the involvement of protein kinase C and the mitogen-activated protein kinase signaling cascade. Journal of Neurochemistry, 2000, 74, 564-73	6	75

77	Kappa opioids promote the proliferation of astrocytes via Gbetagamma and beta-arrestin 2-dependent MAPK-mediated pathways. <i>Journal of Neurochemistry</i> , 2008 , 107, 1753-65	6	72
76	Apparent loss-of-function mutant GPCRs revealed as constitutively desensitized receptors. <i>Biochemistry</i> , 2002 , 41, 11981-9	3.2	72
75	Mu opioid receptor regulation and opiate responsiveness. AAPS Journal, 2005, 7, E587-91	3.7	66
74	Functional selectivity of 6Sguanidinonaltrindole (6SGNTI) at Ebpioid receptors in striatal neurons. Journal of Biological Chemistry, 2013 , 288, 22387-98	5.4	65
73	A novel method for analyzing extremely biased agonism at G protein-coupled receptors. <i>Molecular Pharmacology</i> , 2015 , 87, 866-77	4.3	56
72	Enrestins: regulatory role and therapeutic potential in opioid and cannabinoid receptor-mediated analgesia. <i>Handbook of Experimental Pharmacology</i> , 2014 , 219, 427-43	3.2	54
71	Herkinorin analogues with differential beta-arrestin-2 interactions. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 2421-31	8.3	52
70	A comparison of noninternalizing (herkinorin) and internalizing (DAMGO) mu-opioid agonists on cellular markers related to opioid tolerance and dependence. <i>Synapse</i> , 2007 , 61, 166-75	2.4	51
69	Functional selectivity of GPCR signaling in animals. Current Opinion in Cell Biology, 2014, 27, 102-8	9	48
68	Antinociceptive effects of herkinorin, a MOP receptor agonist derived from salvinorin A in the formalin test in rats: new concepts in mu opioid receptor pharmacology: from a symposium on new concepts in mu-opioid pharmacology. <i>Drug and Alcohol Dependence</i> , 2012 , 121, 181-8	4.9	48
67	Recruitment of EArrestin into Neuronal Cilia Modulates Somatostatin Receptor Subtype 3 Ciliary Localization. <i>Molecular and Cellular Biology</i> , 2016 , 36, 223-35	4.8	45
66	Toward Directing Opioid Receptor Signaling to Refine Opioid Therapeutics. <i>Biological Psychiatry</i> , 2020 , 87, 15-21	7.9	44
65	Structure-activity relationship studies of functionally selective kappa opioid receptor agonists that modulate ERK 1/2 phosphorylation while preserving G protein over @rrestin2 signaling bias. ACS Chemical Neuroscience, 2015, 6, 1411-9	5.7	43
64	Synthesis of conolidine, a potent non-opioid analgesic for tonic and persistent pain. <i>Nature Chemistry</i> , 2011 , 3, 449-53	17.6	41
63	Mu-opioid agonist inhibition of kappa-opioid receptor-stimulated extracellular signal-regulated kinase phosphorylation is dynamin-dependent in C6 glioma cells. <i>Journal of Neurochemistry</i> , 2000 , 74, 574-81	6	40
62	Lubiprostone reverses the inhibitory action of morphine on intestinal secretion in guinea pig and mouse. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2010 , 334, 333-40	4.7	39
61	Optimization of a Series of Mu Opioid Receptor (MOR) Agonists with High G Protein Signaling Bias. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 8895-8907	8.3	38
60	Clozapine acts as an agonist at serotonin 2A receptors to counter MK-801-induced behaviors through a <code>Brrestin2-independent</code> activation of Akt. <i>Neuropsychopharmacology</i> , 2014 , 39, 1902-13	8.7	37

(2017-2010)

59	Serotonin receptor signaling and regulation via Earrestins. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2010 , 45, 555-66	8.7	37
58	Discovery of Small Molecule Kappa Opioid Receptor Agonist and Antagonist Chemotypes through a HTS and Hit Refinement Strategy. <i>ACS Chemical Neuroscience</i> , 2012 , 3, 221-236	5.7	36
57	A G protein signaling-biased agonist at the Eppioid receptor reverses morphine tolerance while preventing morphine withdrawal. <i>Neuropsychopharmacology</i> , 2020 , 45, 416-425	8.7	36
56	Opioid receptor signaling: relevance for gastrointestinal therapy. <i>Current Opinion in Pharmacology</i> , 2006 , 6, 559-63	5.1	35
55	GIRK channel modulation by assembly with allosterically regulated RGS proteins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012 , 109, 19977-82	11.5	32
54	Arrestin recruitment to dopamine D2 receptor mediates locomotion but not incentive motivation. <i>Molecular Psychiatry</i> , 2020 , 25, 2086-2100	15.1	32
53	Morphine-induced physiological and behavioral responses in mice lacking G protein-coupled receptor kinase 6. <i>Drug and Alcohol Dependence</i> , 2009 , 104, 187-96	4.9	31
52	5-HT1A Autoreceptors in the Dorsal Raphe Nucleus Convey Vulnerability to Compulsive Cocaine Seeking. <i>Neuropsychopharmacology</i> , 2016 , 41, 1210-22	8.7	30
51	G protein-coupled receptor kinase/beta-arrestin systems and drugs of abuse: psychostimulant and opiate studies in knockout mice. <i>NeuroMolecular Medicine</i> , 2004 , 5, 41-50	4.6	30
50	Characterization of conditioned place preference to cocaine in congenic dopamine transporter knockout female mice. <i>Psychopharmacology</i> , 2005 , 180, 408-13	4.7	29
49	Investigation of the role of Brrestin2 in kappa opioid receptor modulation in a mouse model of pruritus. <i>Neuropharmacology</i> , 2015 , 99, 600-9	5.5	28
48	Lubiprostone reverses the inhibitory action of morphine on mucosal secretion in human small intestine. <i>Digestive Diseases and Sciences</i> , 2011 , 56, 330-8	4	26
47	Stimulation of mucosal secretion by lubiprostone (SPI-0211) in guinea pig small intestine and colon. <i>American Journal of Physiology - Renal Physiology</i> , 2009 , 296, G823-32	5.1	26
46	RasGRP1 promotes amphetamine-induced motor behavior through a Rhes interaction network ("Rhesactome") in the striatum. <i>Science Signaling</i> , 2016 , 9, ra111	8.8	25
45	G protein signaling-biased agonism at the Eppioid receptor is maintained in striatal neurons. <i>Science Signaling</i> , 2018 , 11,	8.8	25
44	Brain opioid receptor adaptation and expression after prenatal exposure to buprenorphine. <i>Developmental Brain Research</i> , 1998 , 111, 35-42		24
43	Dynamic Strategic Bond Analysis Yields a Ten-Step Synthesis of 20-nor-Salvinorin A, a Potent EOR Agonist. <i>ACS Central Science</i> , 2017 , 3, 1329-1336	16.8	23
42	Seeking (and Finding) Biased Ligands of the Kappa Opioid Receptor. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 694-700	4.3	23

41	Differential signaling properties at the kappa opioid receptor of 12-epi-salvinorin A and its analogues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 1023-6	2.9	22
40	Molecular-interaction and signaling profiles of AM3677, a novel covalent agonist selective for the cannabinoid 1 receptor. <i>ACS Chemical Neuroscience</i> , 2015 , 6, 1400-10	5.7	19
39	Post-activation-mediated changes in opioid receptors detected by N-terminal antibodies. <i>Journal of Biological Chemistry</i> , 2008 , 283, 10735-44	5.4	19
38	Evidence for kappa- and mu-opioid receptor expression in C6 glioma cells. <i>Journal of Neurochemistry</i> , 1998 , 70, 1819-25	6	17
37	Characterization of kappa opioid receptor mediated, dynorphin-stimulated [35S]GTPB binding in mouse striatum for the evaluation of selective KOR ligands in an endogenous setting. Neuropharmacology, 2015, 99, 131-41	5.5	16
36	Seeking Ligand Bias: Assessing GPCR Coupling to Beta-Arrestins for Drug Discovery. <i>Drug Discovery Today: Technologies</i> , 2010 , 7, e37-e42	7.1	14
35	Pharmacological characterization of a selective agonist for bombesin receptor subtype-3. Biochemical and Biophysical Research Communications, 2009 , 387, 283-8	3.4	13
34	Stolonidiol: Synthesis, Target Identification, and Mechanism for Choline Acetyltransferase Activation. <i>Journal of the American Chemical Society</i> , 2017 , 139, 5865-5869	16.4	12
33	Constitutive traffickingmore than just running in circles?. <i>Molecular Pharmacology</i> , 2007 , 71, 957-8	4.3	10
32	Community Guidelines for GPCR Ligand Bias: IUPHAR Review XX <i>British Journal of Pharmacology</i> , 2022 ,	8.6	10
31	Approaches to Assess Functional Selectivity in GPCRs: Evaluating G Protein Signaling in an Endogenous Environment. <i>Methods in Molecular Biology</i> , 2015 , 1335, 177-89	1.4	9
30	O6C-20-nor-salvinorin A is a stable and potent KOR agonist. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 2770-2772	2.9	8
29	The effect of quinine in two bottle choice procedures in C57BL6 mice: Opioid preference, somatic withdrawal, and pharmacokinetic outcomes. <i>Drug and Alcohol Dependence</i> , 2018 , 191, 195-202	4.9	8
28	Repurposing of a Nucleoside Scaffold from Adenosine Receptor Agonists to Opioid Receptor Antagonists. <i>ACS Omega</i> , 2018 , 3, 12658-12678	3.9	8
27	Low Intrinsic Efficacy Alone Cannot Explain the Improved Side Effect Profiles of New Opioid Agonists. <i>Biochemistry</i> , 2021 ,	3.2	8
26	Comparison of morphine, oxycodone and the biased MOR agonist SR-17018 for tolerance and efficacy in mouse models of pain. <i>Neuropharmacology</i> , 2021 , 185, 108439	5.5	7
25	Potency enhancement of the Eppioid receptor antagonist probe ML140 through sulfonamide constraint utilizing a tetrahydroisoquinoline motif. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 3948-5	∂·4	6
24	Kappa-opioid receptor binding varies inversely with tumor grade in human gliomas. <i>Cancer</i> , 1998 , 83, 2561-6	6.4	6

Selectivity for G Protein or Arrestin-Mediated Signaling 2009, 71-85 6 23 Synthesis of Kappa Opioid Antagonists Based On Pyrrolo[1,2-#quinoxalinones Using an 22 N-Arylation/Condensation/Oxidation Reaction Sequence. Journal of Organic Chemistry, **2016**, 81, 10538- 10 550 6 Synthetic Ligands of Cannabinoid Receptors Affect Dauer Formation in the Nematode 21 3.2 5 Caenorhabditis elegans. G3: Genes, Genomes, Genetics, 2016, 6, 1695-705 A Genetically Encoded F-19 NMR Probe Reveals the Allosteric Modulation Mechanism of 20 16.4 Cannabinoid Receptor 1. Journal of the American Chemical Society, 2021, 143, 16320-16325 Approaches to Assess Biased Signaling at the CB1R Receptor. Methods in Enzymology, 2017, 593, 259-279.7 19 4 Probing the CB Cannabinoid Receptor Binding Pocket with AM6538, a High-Affinity Irreversible 18 4.3 Antagonist. Molecular Pharmacology, **2019**, 96, 619-628 Novel Functionalized Cannabinoid Receptor Probes: Development of Exceptionally Potent 8.3 17 3 Agonists. Journal of Medicinal Chemistry, 2021, 64, 3870-3884 Quantitating Ligand Bias Using the Competitive Model of Ligand Activity. Methods in Molecular 16 1.4 Biology, 2019, 1957, 235-247 Exploring the Biology of G Protein-Coupled Receptors from In Vitro to In Vivo. Molecular 15 2 4.3 Pharmacology, **2015**, 88, 534-5 Low intrinsic efficacy alone cannot explain the improved side effect profiles of new opioid agonists 14 Arrestins: Ligand-Directed Regulators of 5-HT2A Receptor Trafficking and Signaling Events 2018, 31-55 13 1 G protein signaling-biased mu opioid receptor agonists that produce sustained G protein activation are noncompetitive agonists. Proceedings of the National Academy of Sciences of the United States 11.5 of America, 2021, 118, Beta-arrestin2 Contributes to the Development of Opioid-Induced Constipation. FASEB Journal, 11 0.9 1 2012, 26, 1123.6 Some effects of putative G-protein biased mu-opioid receptor agonists in male rhesus monkeys. 10 2.4 Behavioural Pharmacology, 2021, 32, 453-458 Improved cyclobutyl nabilone analogs as potent CB1 receptor agonists.. European Journal of 6.8 9 Medicinal Chemistry, **2021**, 230, 114027 How specific are "target-specific" drugs? Celecoxib as a case in point. Molecular Interventions: 8 Pharmacological Perspectives From Biology, Chemistry and Genomics, 2006, 6, 196-8 Mu Opioid Receptor Regulation and Opiate Responsiveness 2008, 617-624 Biased agonism and opioid receptor-mediated analgesia. Proceedings for Annual Meeting of the Japanese Pharmacological Society, 2018, WCP2018, SY85-4

5	Expression of different adenylyl cyclase isoforms impacts ligand bias downstream of the kappa opioid receptor. <i>FASEB Journal</i> , 2019 , 33, 503.11	0.9
4	Detecting the Role of Arrestins in G Protein-Coupled Receptor Regulation. <i>Neuromethods</i> , 2011 , 347-3	5&.4
3	The role of beta-arrestin2 in clozapines actions at the serotonin 2A receptor. <i>FASEB Journal</i> , 2012 , 26, lb573	0.9
2	Decaf or regular? Energizing the caffeine receptor. <i>Cell</i> , 2021 , 184, 1659-1660	56.2
1	Structure activity relationship investigation of triazole-based kappa opioid receptor agonists. Medicinal Chemistry Research, 2021, 30, 1386-1396	2.2