

THE CONCISE GUIDE TO PHARMACOLOGY 2015/16: G protein-coupled receptors

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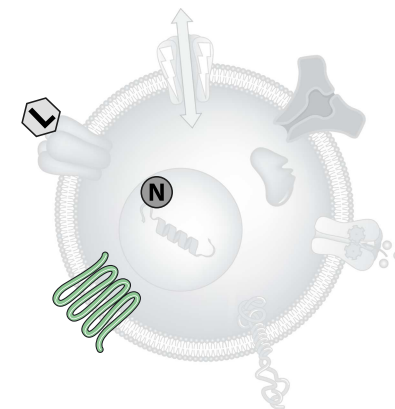
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Abstract

The Concise Guide to PHARMACOLOGY 2015/16 provides concise overviews of the key properties of over 1750 human drug targets with their pharmacology, plus links to an open access knowledgebase of drug targets and their ligands (www.guidetopharmacology.org), which provides more detailed views of target and ligand properties. The full contents can be found at <http://onlinelibrary.wiley.com/doi/10.1111/bph.13348/full>. G protein-coupled receptors are one of the eight major pharmacological targets into which the Guide is divided, with the others being: ligand-gated ion channels, voltage-gated ion channels, other ion channels, nuclear hormone receptors, catalytic receptors, enzymes and transporters. These are presented with nomenclature guidance and summary information on the best available pharmacological tools, alongside key references and suggestions for further reading. The Concise Guide is published in landscape format in order to facilitate comparison of related targets. It is a condensed version of material contemporary to late 2015, which is presented in greater detail and constantly updated on the website www.guidetopharmacology.org, superseding data presented in the previous Guides to Receptors & Channels and the Concise Guide to PHARMACOLOGY 2013/14. It is produced in conjunction with NC-IUPHAR and provides the official IUPHAR classification and nomenclature for human drug targets, where appropriate. It consolidates information previously curated and displayed separately in IUPHAR-DB and GRAC and provides a permanent, citable, point-in-time record that will survive database updates.

Conflict of interest

The authors state that there are no conflicts of interest to declare.

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Overview: G protein-coupled receptors (GPCRs) are the largest class of membrane proteins in the human genome. The term "7TM receptor" is commonly used interchangeably with "GPCR", although there are some receptors with seven transmembrane domains that do not signal through G proteins. GPCRs share a common architecture, each consisting of a single polypeptide with an extracellular N-terminus, an intracellular C-terminus and seven hydrophobic transmembrane domains (TM1-TM7) linked by three extracellular loops (ECL1-ECL3) and three intracellular

loops (ICL1-ICL3). About 800 GPCRs have been identified in man, of which about half have sensory functions, mediating olfaction (400), taste (33), light perception (10) and pheromone signalling (5) [1309]. The remaining 350 non-sensory GPCRs mediate intersignalling by ligands that range in size from small molecules to peptide to large proteins; they are the targets for the majority of drugs in clinical usage [1451, 1560], although only a minority of these receptors are exploited therapeutically. The first classification scheme to be proposed for GPCRs [984] di-

vided them, on the basis of sequence homology, into six classes. These classes and their prototype members were as follows: **Class A** (rhodopsin-like), **Class B** (secretin receptor family), **Class C** (metabotropic glutamate), **Class D** (fungal mating pheromone receptors), **Class E** (cyclic AMP receptors) and **Class F** (frizzled/smoothed). Of these, classes D and E are not found in vertebrates. An alternative classification scheme "GRAFS" [1666] divides vertebrate GPCRs into five classes, overlapping with the A-F nomenclature, viz:

Searchable database: <http://www.guidetopharmacology.org/index.jsp>

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Glutamate family (class C), which includes metabotropic glutamate receptors, a calcium-sensing receptor and GABA_B receptors, as well as three taste type 1 receptors [class C list] and a family of pheromone receptors (V2 receptors) that are abundant in rodents but absent in man [1309].

Rhodopsin family (class A), which includes receptors for a wide variety of small molecules, neurotransmitters, peptides and hormones, together with olfactory receptors, visual pigments, taste type 2 receptors and five pheromone receptors (V1 receptors). [Class A list]

Adhesion family GPCRs are phylogenetically related to class B receptors, from which they differ by possessing large extracellular N-termini that are autoproteolytically cleaved from their 7TM domains at a conserved "GPCR proteolysis site" (GPS) which lies within a much larger (320 residue) "GPCR autoproteolysis-inducing" (GAIN) domain, an evolutionary ancient motif also found in polycystic kidney disease 1 (PKD1)-like proteins, which has been suggested to be both required and sufficient for autoproteolysis [1538]. [Adhesion family list].

Frizzled family (class F) consists of 10 Frizzled proteins (FZD(1-10)) and Smoothed (SMO). [Frizzled family list]. The FZDs are activated by secreted lipoglycoproteins of the WNT family, whereas SMO is indirectly activated by the Hedgehog (HH) family of proteins acting on the transmembrane protein Patched (PTCH).

Secretin family (class B), encoded by 15 genes in humans. The ligands for receptors in this family are polypeptide hormones of 27-141 amino-acid residues; nine of the mammalian receptors respond to ligands that are structurally related to one another (glucagon, glucagon-like peptides (GLP-1, GLP-2), glucose-dependent insulinotropic polypeptide (GIP), secretin, vasoactive intestinal peptide (VIP), pituitary adenylate cyclase-activating polypeptide (PACAP) and growth-hormone-releasing hormone (GHRH) [703].

GPCR families

Family	Class A	Class B (Secretin)	Class C (Glutamate)	Adhesion	Frizzled
Receptors with known ligands	197 ^a	15	12	0	11
Orphans	87 (54) ^a	-	8 (1) ^a	26 (6) ^a	0
Sensory (olfaction)	390 ^{b,c}	-	-	-	-
Sensory (vision)	10 ^d opsins	-	-	-	-
Sensory (taste)	30 ^c taste 2	-	3 ^c taste 1	-	-
Sensory (pheromone)	5 ^c vomeronasal 1	-	-	-	-
Total	719	15	22	33	11

^aNumbers in brackets refer to orphan receptors for which an endogenous ligand has been proposed in at least one publication, see [396]; ^b[1443]; ^c[1309]; ^d[1866].

Much of our current understanding of the structure and function of GPCRs is the result of pioneering work on the visual pigment rhodopsin and on the β_2 adrenoceptor, the latter culminating in the award of the 2012 Nobel Prize in chemistry to Robert Lefkowitz and Brian Kobilka [975, 1073].

Family structure

5746	Orphan and other 7TM receptors	5780	Bradykinin receptors	5803	GABA _B receptors
5746	Class A Orphans	5781	Calcitonin receptors	5805	Galanin receptors
5756	Class C Orphans	5783	Calcium-sensing receptors	5806	Ghrelin receptor
5756	Taste 1 receptors	5784	Cannabinoid receptors	5807	Glucagon receptor family
5757	Taste 2 receptors	5785	Chemerin receptor	5809	Glycoprotein hormone receptors
5758	Other 7TM proteins	5785	Chemokine receptors	5810	Gonadotrophin-releasing hormone receptors
5759	5-Hydroxytryptamine receptors	5791	Cholecystokinin receptors	5811	GPR18, GPR55 and GPR119
5764	Acetylcholine receptors (muscarinic)	5792	Class Frizzled GPCRs	5812	Histamine receptors
5766	Adenosine receptors	5793	Complement peptide receptors	5814	Hydroxycarboxylic acid receptors
5768	Adhesion Class GPCRs	5795	Corticotropin-releasing factor receptors	5815	Kisspeptin receptor
5770	Adrenoceptors	5796	Dopamine receptors	5816	Leukotriene receptors
5774	Angiotensin receptors	5798	Endothelin receptors	5818	Lysophospholipid (LPA) receptors
5775	Apelin receptor	5799	G protein-coupled estrogen receptor	5819	Lysophospholipid (S1P) receptors
5777	Bile acid receptor	5800	Formylpeptide receptors	5820	Melanin-concentrating hormone receptors
5778	Bombesin receptors	5801	Free fatty acid receptors	5821	Melanocortin receptors

5822	Melatonin receptors	5835	Orexin receptors	5846	Relaxin family peptide receptors
5823	Metabotropic glutamate receptors	5836	Oxoglutarate receptor	5848	Somatostatin receptors
5826	Motilin receptor	5836	P2Y receptors	5850	Succinate receptor
5827	Neuromedin U receptors	5838	Parathyroid hormone receptors	5850	Tachykinin receptors
5828	Neuropeptide FF/neuropeptide AF receptors	5839	Platelet-activating factor receptor	5852	Thyrotropin-releasing hormone receptors
5829	Neuropeptide S receptor	5840	Prokineticin receptors	5852	Trace amine receptor
5829	Neuropeptide W/neuropeptide B receptors	5841	Prolactin-releasing peptide receptor	5854	Urotensin receptor
5830	Neuropeptide Y receptors	5842	Prostanoid receptors	5854	Vasopressin and oxytocin receptors
5832	Neurotensin receptors	5844	Proteinase-activated receptors	5856	VIP and PACAP receptors
5833	Opioid receptors	5846	QRFP receptor		

Orphan and other 7TM receptors

G protein-coupled receptors → Orphan and other 7TM receptors

Class A Orphans

G protein-coupled receptors → Orphan and other 7TM receptors → Class A Orphans

Overview: Table 1 lists a number of putative GPCRs identified by **NC-IUPHAR [530]**, for which preliminary evidence for an endogenous ligand has been published, or for which there exists a potential link to a disease, or disorder. These GPCRs have recently been reviewed in detail [396]. The GPCRs in Table 1 are all Class A, rhodopsin-like GPCRs. Class A orphan GPCRs not listed in Table 1 are putative GPCRs with as-yet unidentified endogenous ligands.

Table 1: Class A orphan GPCRs with putative endogenous ligands

<i>GPR1</i>	<i>GPR3</i>	<i>GPR4</i>	<i>GPR6</i>	<i>GPR12</i>	<i>GPR15</i>	<i>GPR17</i>	<i>GPR20</i>
<i>GPR22</i>	<i>GPR26</i>	<i>GPR31</i>	<i>GPR34</i>	<i>GPR35</i>	<i>GPR37</i>	<i>GPR39</i>	<i>GPR50</i>
<i>GPR63</i>	<i>GRP65</i>	<i>GPR68</i>	<i>GPR75</i>	<i>GPR84</i>	<i>GPR87</i>	<i>GPR88</i>	<i>GPR132</i>
<i>GPR149</i>	<i>GPR161</i>	<i>GPR183</i>	<i>LGR4</i>	<i>LGR5</i>	<i>LGR6</i>	<i>MAS1</i>	<i>MRGPRD</i>
<i>MRGPRX1</i>	<i>MRGPRX2</i>	<i>P2RY10</i>	<i>TAAR2</i>				

In addition the orphan receptors *GPR18*, *GPR55* and *GPR119* which are reported to respond to endogenous agents analogous to the endogenous cannabinoid ligands have been grouped together (*GPR18*, *GPR55* and *GPR119*).

Nomenclature	<i>GPR1</i>	<i>GPR3</i>	<i>GPR4</i>	<i>GPR6</i>
HGNC, UniProt	<i>GPR1</i> , P46091	<i>GPR3</i> , P46089	<i>GPR4</i> , P46093	<i>GPR6</i> , P46095
Endogenous ligand	–	–	Protons	–
Endogenous agonists	chemerin (<i>RARRES2</i> , Q99969) (p <i>K_d</i> 8.3) [95]	–	–	–
Agonists	–	diphenyleneiodonium chloride (p <i>EC</i> ₅₀ 6) [2091]	–	–

(continued)				
Nomenclature	GPR1	GPR3	GPR4	GPR6
Comments	Reported to act as a co-receptor for HIV [1724]. See review [396] for discussion of pairing with chemerin.	sphingosine 1-phosphate was reported to be an endogenous agonist [1921], but this finding was not replicated in subsequent studies [2093]. Reported to activate adenylyl cyclase constitutively through G _s [466]. Gene disruption results in premature ovarian ageing [1063], reduced β -amyloid deposition [1868] and hypersensitivity to thermal pain [1615] in mice. First small molecule inverse agonist [860] and agonists identified [2091].	An initial report suggesting activation by lysophosphatidylcholine and sphingosylphosphorylcholine [2131] has been retracted [2148]. GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [396, 1704]. Gene disruption is associated with increased perinatal mortality and impaired vascular proliferation [2085]. Negative allosteric modulators of GPR4 have been reported [1889].	An initial report that sphingosine 1-phosphate (S1P) was a high-affinity ligand (EC ₅₀ value of 39nM) [815, 1921] was not repeated by arrestin PathHunter[TM] assays [1785, 2093]. Reported to activate adenylyl cyclase constitutively through G _s and to be located intracellularly [1453]. GPR6-deficient mice showed reduced striatal cyclic AMP production <i>in vitro</i> and selected alterations in instrumental conditioning <i>in vivo</i> . [1134].
Nomenclature	GPR12	GPR15	GPR17	GPR19
HGNC, UniProt	GPR12 , P47775	GPR15 , P49685	GPR17 , Q13304	GPR19 , Q15760
Endogenous agonists	–	–	UDP-glucose (pEC ₅₀ 5.9–9.5) [130, 344], LTC₄ (pEC ₅₀ 7.8–9.5) [344], UDP-galactose (pEC ₅₀ 6–8.9) [130, 344], uridine diphosphate (pEC ₅₀ 6–8.8) [130, 344], LTD₄ (pEC ₅₀ 8.1–8.4) [344]	–
Comments	Reports that sphingosine 1-phosphate is a ligand of GPR12 [814, 1921] have not been replicated in arrestin-based assays [1785, 2093]. Gene disruption results in dyslipidemia and obesity [154].	Reported to act as a co-receptor for HIV [462]. In an infection-induced colitis model, <i>Gpr15</i> knockout mice were more prone to tissue damage and inflammatory cytokine expression [945].	Reported to be a dual leukotriene and uridine diphosphate receptor [344]. Another group instead proposed that GPR17 functions as a negative regulator of the CysLT ₁ receptor response to leukotriene D ₄ (LTD₄). For further discussion, see [396]. Reported to antagonize CysLT ₁ receptor signalling <i>in vivo</i> and <i>in vitro</i> [1175]. See reviews [250] and [396].	–

Nomenclature	GPR20	GPR21	GPR22	GPR25	GPR26
HGNC, UniProt	GPR20, Q99678	GPR21, Q99679	GPR22, Q99680	GPR25, O00155	GPR26, Q8NDV2
Comments	Reported to inhibit adenylyl cyclase constitutively through G _{i/o} [708]. GPR20 deficient mice exhibit hyperactivity characterised by increased total distance travelled in an open field test [207].	<i>Gpr21</i> knockout mice were resistant to diet-induced obesity, exhibiting an increase in glucose tolerance and insulin sensitivity, as well as a modest lean phenotype [1448].	Gene disruption results in increased severity of functional decompensation following aortic banding [10]. Identified as a susceptibility locus for osteoarthritis [494, 929, 1935].	–	Has been reported to activate adenylyl cyclase constitutively through G _s [880]. <i>Gpr26</i> knockout mice show increased levels of anxiety and depression-like behaviours [2117].

Nomenclature	GPR27	GPR31	GPR32	GPR33	GPR34
HGNC, UniProt	GPR27, Q9NS67	GPR31, O00270	GPR32, O75388	GPR33, Q49SQ1	GPR34, Q9UPCS
Rank order of potency	–	–	resolvin D1 > LXA ₄	–	–
Endogenous agonists	–	12S-HETE (Selective) (pEC ₅₀ 9.6) [665] – Mouse	resolvin D1 (pEC ₅₀ 11.1) [1006], LXA ₄ (pEC ₅₀ 9.7) [1006]	–	lysophosphatidylserine (Selective) (pEC ₅₀ 6.6–6.9) [960, 1817]
Labelled ligands	–	–	[³ H]resolvin D1 (Agonist) (pK _d 9.7) [1006]	–	–
Comments	Knockdown of Gpr27 reduces endogenous mouse insulin promotor activity and glucose-stimulated insulin secretion [1012].	See [396] for discussion of pairing.	resolvin D1 has been demonstrated to activate GPR32 in two publications [316, 1006]. The pairing was not replicated in a recent study based on arrestin recruitment [1785]. <i>GPR32</i> is a pseudogene in mice and rats. See reviews [250] and [396].	<i>GPR33</i> is a pseudogene in most individuals, containing a premature stop codon within the coding sequence of the second intracellular loop [1621].	Lysophosphatidylserine has been reported to be a ligand of GPR34 in several publications, but the pairing was not replicated in a recent study based on arrestin recruitment [1785]. Fails to respond to a variety of lipid-derived agents [2093]. Gene disruption results in an enhanced immune response [1102]. Characterization of agonists at this receptor is discussed in [819] and [396].

Nomenclature	GPR35	GPR37	GPR37L1	GPR39	GPR42
HGNC, UniProt	GPR35 , Q9HC97	GPR37 , O15354	GPR37L1 , O60883	GPR39 , O43194	GPR42 , O15529
Endogenous agonists	2-oleoyl-LPA (pEC ₅₀ 7.3–7.5) [1436], kynurenic acid (pEC ₅₀ 3.9–4.4) [1785 , 1980]	–	–	Zn²⁺ [775]	–
Agonists	–	neuropeptide head activator (pEC ₅₀ 8–8.5) [1578]	–	compound 1 [PMID: 24900608] (pEC ₅₀ 4.9–7.2) [166]	–
Comments	Several studies have shown that kynurenic acid is an agonist of GPR35 but it remains controversial whether the proposed endogenous ligand reaches sufficient tissue concentrations to activate the receptor [1015]. 2-oleoyl-LPA has also been proposed as an endogenous ligand [1436] but these results were not replicated in an arrestin assay [1785]. The phosphodiesterase inhibitor zaprinast [1863] has become widely used as a surrogate agonist to investigate GPR35 pharmacology and signalling [1863]. GPR35 is also activated by the pharmaceutical adjunct pamoic acid [2124]. See reviews [396] and [429].	Reported to associate and regulate the dopamine transporter [1207] and to be a substrate for parkin [1205]. Gene disruption results in altered striatal signalling [1206]. The peptides prosaptide and prosaposin are proposed as endogenous ligands for GPR37 and GPR37L1 [1264].	The peptides prosaptide and prosaposin are proposed as endogenous ligands for GPR37 and GPR37L1 [1264].	Zn²⁺ has been reported to be a potent and efficacious agonist of human, mouse and rat GPR39 [2089]. obestatin (GHRL , Q9UBU3), a fragment from the ghrelin precursor, was reported initially as an endogenous ligand, but subsequent studies failed to reproduce these findings. GPR39 has been reported to be down-regulated in adipose tissue in obesity-related diabetes [273]. Gene disruption results in obesity and altered adipocyte metabolism [1497]. Reviewed in [396].	–
Nomenclature	GPR45	GPR50	GPR52	GPR61	
HGNC, UniProt	GPR45 , Q9Y5Y3	GPR50 , Q13585	GPR52 , Q9Y2T5	GPR61 , Q9BZJ8	
Comments	–	GPR50 is structurally related to MT₁ and MT₂ melatonin receptors, with which it heterodimerises constitutively and specifically [1089]. Gpr50 knockout mice display abnormal thermoregulation and are much more likely than wild-type mice to enter fasting-induced torpor [111].	First small molecule agonist reported [1703].	GPR61 deficient mice exhibit obesity associated with hyperphagia [1363]. Although no endogenous ligands have been identified, 5-(nonyloxy)tryptamine has been reported to be a low affinity inverse agonist [1852].	

Nomenclature	GPR62	GPR63	GPR65	GPR68	GPR75
HGNC, UniProt	GPR62, Q9BZJ7	GPR63, Q9BZJ6	GPR65, Q8IYL9	GPR68, Q15743	GPR75, O95800
Endogenous ligand	–	–	Protons	Protons	–
Comments	–	sphingosine 1-phosphate and dioleoylphosphatidic acid have been reported to be low affinity agonists for GPR63 [1394] but this finding was not replicated in an arrestin-based assay [2093].	GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [396, 1704]. Reported to activate adenylyl cyclase; gene disruption leads to reduced eosinophilia in models of allergic airway disease [1000].	GPR68 was previously identified as a receptor for sphingosylphosphorylcholine (SPC) [2068], but the original publication has been retracted [2067]. GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [396, 1704]. A family of 3,5-disubstituted isoxazoles were identified as agonists of GPR68 [1617].	CCL5 (CCL5, P13501) was reported to be an agonist of GPR75 [816], but the pairing could not be repeated in an arrestin assay [1785].

Nomenclature	GPR78	GPR79	GPR82	GPR83	GPR84
HGNC, UniProt	GPR78, Q96P69	GPR79, –	GPR82, Q96P67	GPR83, Q9NYM4	GPR84, Q9NQ55
Agonists	–	–	–	Zn^{2+} (pEC ₅₀ 5) [1351] – Mouse	decanoic acid (pEC ₅₀ 5–5.4) [1785, 1981], undecanoic acid (pEC ₅₀ 5.1) [1981], lauric acid (pEC ₅₀ 5) [1981]
Comments	GPR78 has been reported to be constitutively active, coupled to elevated cAMP production [880].	–	Mice with <i>Gpr82</i> knockout have a lower body weight and body fat content associated with reduced food intake, decreased serum triglyceride levels, as well as higher insulin sensitivity and glucose tolerance [479].	One isoform has been implicated in the induction of CD4(+) CD25(+) regulatory T cells (Tregs) during inflammatory immune responses [696]. The extracellular N-terminal domain is reported as an intramolecular inverse agonist [1352].	Medium chain free fatty acids with carbon chain lengths of 9–14 activate GPR84 [1828, 1981]. A surrogate ligand for GPR84, 6-n-octylaminouracil has also been proposed [1828]. See review [396] for discussion of classification. Mutational analysis and molecular modelling of GPR84 has been reported [1397].

Nomenclature	GPR85	GPR87	GPR88	GPR101
HGNC, UniProt	GPR85, P60893	GPR87, Q9BY21	GPR88, Q9GZNO	GPR101, Q96P66
Endogenous agonists	–	LPA (pEC ₅₀ 7.4) [1344 , 1836]	–	–
Agonists	–	–	compound 2 [PMID: 24793972] (pEC ₅₀ 6.2) [868]	–
Comments	Proposed to regulate hippocampal neurogenesis in the adult, as well as neurogenesis-dependent learning and memory [303].	–	Gene disruption results in altered striatal signalling [1137]. Small molecule agonists have been reported [147].	Mutations in GPR101 have been linked to gigantism and acromegaly [1906].

Nomenclature	GPR132	GPR135	GPR139	GPR141	GPR142
HGNC, UniProt	GPR132, Q9UNW8	GPR135, Q8IZ08	GPR139, Q6DWJ6	GPR141, Q7Z602	GPR142, Q7Z601
Endogenous ligand	Protons	–	–	–	–
Agonists	–	–	compound 1a [PMID: 24900311] (pEC ₅₀ 7.4) [1721]	–	–
Comments	GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [396 , 1704]. Reported to respond to lysophosphatidylcholine [891], but later retracted [2038].	–	Peptide agonists have been reported [828].	–	Small molecule agonists have been reported [1890 , 2106].

Nomenclature	GPR146	GPR148	GPR149	GPR150	GPR151
HGNC, UniProt	GPR146, Q96CH1	GPR148, Q8TDV2	GPR149, Q86SP6	GPR150, Q8NGU9	GPR151, Q8TDV0
Comments	Yosten <i>et al.</i> demonstrated inhibition of proinsulin C-peptide (INS, P01308) -induced stimulation of cFos expression following knockdown of GPR146 in KATO III cells, suggesting proinsulin C-peptide as an endogenous ligand of the receptor [2103].	–	<i>Gpr149</i> knockout mice displayed increased fertility and enhanced ovulation, with increased levels of FSH receptor and cyclin D2 mRNA levels [463].	–	GPR151 responded to galanin with an EC ₅₀ value of 2 μM, suggesting that the endogenous ligand shares structural features with galanin (GAL, P22466) [813].

Nomenclature	GPR152	GPR153	GPR160	GPR161	GPR162
HGNC, UniProt	GPR152, Q8TDT2	GPR153, Q6NV75	GPR160, Q9UJ42	GPR161, Q8N6U8	GPR162, Q16538
Comments	–	–	–	A C-terminal truncation (deletion) mutation in Gpr161 causes congenital cataracts and neural tube defects in the vacuolated lens (vl) mouse mutant [1226]. The mutated receptor is associated with cataract, spina bifida and white belly spot phenotypes in mice [994]. Gene disruption is associated with a failure of asymmetric embryonic development in zebrafish [1085].	–

Nomenclature	GPR171	GPR173	GPR174	GPR176	GPR182
HGNC, UniProt	GPR171, O14626	GPR173, Q9NS66	GPR174, Q9BXC1	GPR176, Q14439	GPR182, O15218
Endogenous agonists	–	–	lysophosphatidylserine (pEC ₅₀ 7.1) [825]	–	–
Comments	GPR171 has been shown to be activated by the endogenous peptide BigLEN (Mouse). This receptor-peptide interaction is believed to be involved in regulating feeding and metabolism responses [621].	–	See [819] which discusses characterization of agonists at this receptor.	–	Rat GPR182 was first proposed as the adrenomedullin receptor [904]. However, it was later reported that rat and human GPR182 did not respond to adrenomedullin [927] and GPR182 is not currently considered to be a genuine adrenomedullin receptor [722].

Nomenclature	GPR183	LGR4	LGR5	LGR6	MAS1
HGNC, UniProt	GPR183 , P32249	LGR4 , Q9BXB1	LGR5 , O75473	LGR6 , Q9HBX8	MAS1 , P04201
Endogenous agonists	7α,25-dihydroxycholesterol (Selective) (pEC ₅₀ 8.1–9.8) [694 , 1125], 7α,27-dihydroxycholesterol (Selective) (pEC ₅₀ 8.9) [1125], 7β, 25-dihydroxycholesterol (Selective) (pEC ₅₀ 8.7) [1125], 7β, 27-dihydroxycholesterol (Selective) (pEC ₅₀ 7.3) [1125]	R-spondin-2 (RSPO2 , Q6UXX9) (pEC ₅₀ 12.5) [266], R-spondin-1 (RSPO1 , Q2MKA7) (pEC ₅₀ 10.7) [266], R-spondin-3 (RSPO3 , Q9BXY4) (pEC ₅₀ 10.7) [266], R-spondin-4 (RSPO4 , Q2IOM5) (pEC ₅₀ 10.1) [266]	R-spondin-2 (RSPO2 , Q6UXX9) (pEC ₅₀ 12) [266], R-spondin-1 (RSPO1 , Q2MKA7) (pEC ₅₀ 11.1) [266], R-spondin-3 (RSPO3 , Q9BXY4) (pEC ₅₀ 11) [266], R-spondin-4 (RSPO4 , Q2IOM5) (pEC ₅₀ 9.4) [266]	R-spondin-1 (RSPO1 , Q2MKA7) [266 , 2140], R-spondin-2 (RSPO2 , Q6UXX9) [266 , 2140], R-spondin-3 (RSPO3 , Q9BXY4) [266 , 2140], R-spondin-4 (RSPO4 , Q2IOM5) [266 , 2140]	–
Agonists	–	–	–	–	angiotensin-(1-7) (AGT , P01019) (pK _i 7.3) [612] – Mouse
Comments	Two independent publications have shown that 7α,25-dihydroxycholesterol is an agonist of GPR183 and have demonstrated by mass spectrometry that this oxysterol is present endogenously in tissues [694 , 1125]. Gpr183-deficient mice show a reduction in the early antibody response to a T-dependent antigen. GPR183-deficient B cells fail to migrate to the outer follicle and instead stay in the follicle centre [923 , 1488].	LGR4 does not couple to heterotrimeric G proteins or recruit arrestins when stimulated by the R-spondins, indicating a unique mechanism of action. R-spondins bind to LGR4, which specifically associates with Frizzled and LDL receptor-related proteins (LRPs) that are activated by the extracellular Wnt molecules and then trigger canonical Wnt signalling to increase gene expression [266 , 1612 , 2140]. Gene disruption leads to multiple developmental disorders [869 , 1154 , 1781 , 2005].	The four R-spondins can bind to LGR4, LGR5, and LGR6, which specifically associate with Frizzled and LDL receptor-related proteins (LRPs), proteins that are activated by extracellular Wnt molecules and which then trigger canonical Wnt signalling to increase gene expression [266 , 2140].	–	–

Nomenclature	MASTL	MRGPRD	MRGPRE	MRGPRF	MRGPRG
HGNC, UniProt	MASTL, P35410	MRGPRD, Q8TDS7	MRGPRE, Q86SM8	MRGPRF, Q96AM1	MRGPRG, Q86SM5
Endogenous agonists	–	β-alanine (pEC ₅₀ 4.8) [1729 , 1785]	–	–	–
Comments	–	An endogenous peptide with a high degree of sequence similarity to angiotensin-(1-7) (AGT , P01019), alamandine (AGT), was shown to promote NO release in MRGPRD-transfected cells. The binding of alamandine to MRGPRD to was shown to be blocked by D-Pro ⁷ -angiotensin-(1-7), β-alanine and PD123319 [1045]. Genetic ablation of MRGPRD+ neurons of adult mice decreased behavioural sensitivity to mechanical stimuli but not to thermal stimuli [278]. See reviews [396] and [1779].	See reviews [396] and [1779].	MRGPRF has been reported to respond to stimulation by angiotensin metabolites [589]. See reviews [396] and [1779].	See reviews [396] and [1779].

Nomenclature	MRGPRX1	MRGPRX2	MRGPRX3	MRGPRX4
HGNC, UniProt	MRGPRX1, Q96LB2	MRGPRX2, Q96LB1	MRGPRX3, Q96LB0	MRGPRX4, Q96LA9
Endogenous agonists	bovine adrenal medulla peptide 8-22 (PENK , P01210) (Selective) (pEC ₅₀ 5.3–7.8) [299 , 1080 , 1785]	PAMP-20 (ADM , P35318) (Selective) [899]	–	–
Agonists	–	cortistatin-14 {Mouse, Rat} (pEC ₅₀ 6.9–7.6) [899 , 1594 , 1785]	–	–
Selective agonists	–	PAMP-12 (human) (pEC ₅₀ 7.2–7.7) [899]	–	–
Comments	Reported to mediate the sensation of itch [1131 , 1739]. Reports that bovine adrenal medulla peptide 8-22 (PENK , P01210) was the most potent of a series of proenkephalin A-derived peptides as an agonist of MRGPRX1 in assays of calcium mobilisation and radioligand binding [1080] were replicated in an independent study using an arrestin recruitment assay [1785]. See reviews [396] and [1779].	A diverse range of substances has been reported to be agonists of MRGPRX2, with cortistatin 14 the highest potency agonist in assays of calcium mobilisation [1594], also confirmed in an independent study using an arrestin recruitment assay [1785]. See reviews [396] and [1779].	–	See reviews [396] and [1779].

Nomenclature	OPN3	OPN4	OPN5	P2RY8
HGNC, UniProt	OPN3, Q9H1Y3	OPN4, Q9UHM6	OPN5, Q6U736	P2RY8, Q86VZ1
Comments	–	–	Evidence indicates OPN5 triggers a UV-sensitive G _i -mediated signalling pathway in mammalian tissues [982].	–

Nomenclature	P2RY10	TAAR2	TAAR3	TAAR4P
HGNC, UniProt	P2RY10, O00398	TAAR2, Q9P1P5	TAAR3, Q9P1P4	TAAR4P, –
Rank order of potency	–	β-phenylethylamine > tryptamine [185]	–	–
Endogenous agonists	sphingosine 1-phosphate (Selective) (pEC ₅₀ 7.3) [1344], LPA (Selective) (pEC ₅₀ 6.9) [1344]	–	–	–
Comments	–	Probable pseudogene in 10-15% of Asians due to a polymorphism (rs8192646) producing a premature stop codon at amino acid 168 [396].	TAAR3 is thought to be a pseudogene in man though functional in rodents [396].	Pseudogene in man but functional in rodents [396].

Nomenclature	TAAR5	TAAR6	TAAR8	TAAR9
HGNC, UniProt	TAAR5, O14804	TAAR6, Q96R18	TAAR8, Q969N4	TAAR9, Q96R19
Comments	Trimethylamine is reported as an agonist [1974] and 3-iodothyronamine an inverse agonist [426].	–	–	TAAR9 appears to be functional in most individuals but has a polymorphic premature stop codon at amino acid 61 (rs2842899) with an allele frequency of 10-30% in different populations [1944].

Class C Orphans

G protein-coupled receptors → Orphan and other 7TM receptors → Class C Orphans

Nomenclature	<i>GPR156</i>	<i>GPR158</i>	<i>GPR179</i>	<i>GPRCSA</i>	<i>GPRCSB</i>	<i>GPRCSC</i>	<i>GPRCSD</i>
HGNC, UniProt	<i>GPR156</i> , Q8NFN8	<i>GPR158</i> , Q5T848	<i>GPR179</i> , Q6PRD1	<i>GPRCSA</i> , Q8NFJ5	<i>GPRCSB</i> , Q9NZH0	<i>GPRCSC</i> , Q9NQ84	<i>GPRCSD</i> , Q9NZD1

Taste 1 receptors

G protein-coupled receptors → Orphan and other 7TM receptors → Taste 1 receptors

Overview: Whilst the taste of acid and salty foods appear to be sensed by regulation of ion channel activity, bitter, sweet and umami tastes are sensed by specialised GPCR. Two classes of taste GPCR have been identified, T1R and T2R, which are similar in sequence and structure to Class C and Class A GPCR, respectively. Activation

of taste receptors appears to involve gustducin- (*Gαt3*) and *Gα14*-mediated signalling, although the precise mechanisms remain obscure. Gene disruption studies suggest the involvement of PLCβ2 [2122], TRPM5 [2122] and IP3 [764] receptors in post-receptor signalling of taste receptors. Although predominantly associated with

the oral cavity, taste receptors are also located elsewhere, including further down the gastrointestinal system, in the lungs and in the brain.

Sweet/Umami

T1R3 acts as an obligate partner in T1R1/T1R3 and T1R2/T1R3 heterodimers, which sense umami or sweet, respectively. T1R1/T1R3 heterodimers respond to *L*-glutamic acid and may be positively allosterically modulated by 5'-nucleoside monophosphates, such as 5'-GMP [1096]. T1R2/T1R3 heterodimers respond to sugars, such as sucrose, and artificial sweeteners, such as saccharin [1376].

Nomenclature	<i>TAS1R1</i>	<i>TAS1R2</i>	<i>TAS1R3</i>
HGNC, UniProt	<i>TAS1R1</i> , Q7RTX1	<i>TAS1R2</i> , Q8TE23	<i>TAS1R3</i> , Q7RTX0

Taste 2 receptors

G protein-coupled receptors → Orphan and other 7TM receptors → Taste 2 receptors

Bitter

The composition and stoichiometry of bitter taste receptors is not yet established. Bitter receptors appear to separate into two groups, with very restricted ligand specificity or much broader responsiveness. For example, T2R5 responded to [cycloheximide](#), but not 10 other bitter compounds [287], while T2R14 responded to at least eight different bitter tastants, including (-)- α -thujone and [picrotoxinin](#) [119].

Specialist database [BitterDB](#) contains additional information on bitter compounds and receptors [2023].

Nomenclature	TAS2R1	TAS2R3	TAS2R4	TAS2R5	TAS2R7	TAS2R8
HGNC, UniProt	TAS2R1, Q9NYW7	TAS2R3, Q9NYW6	TAS2R4, Q9NYW5	TAS2R5, Q9NYW4	TAS2R7, Q9NYW3	TAS2R8, Q9NYW2

Nomenclature	TAS2R9	TAS2R10	TAS2R13	TAS2R14	TAS2R16	TAS2R19
HGNC, UniProt	TAS2R9, Q9NYW1	TAS2R10, Q9NYW0	TAS2R13, Q9NYV9	TAS2R14, Q9NYV8	TAS2R16, Q9NYV7	TAS2R19, P59542

Nomenclature	TAS2R20	TAS2R30	TAS2R31	TAS2R38	TAS2R39	TAS2R40
HGNC, UniProt	TAS2R20, P59543	TAS2R30, P59541	TAS2R31, P59538	TAS2R38, P59533	TAS2R39, P59534	TAS2R40, P59535

Nomenclature	TAS2R41	TAS2R42	TAS2R43	TAS2R45	TAS2R46	TAS2R50	TAS2R60
HGNC, UniProt	TAS2R41, P59536	TAS2R42, Q7RTR8	TAS2R43, P59537	TAS2R45, P59539	TAS2R46, P59540	TAS2R50, P59544	TAS2R60, P59551

Other 7TM proteins

G protein-coupled receptors → Orphan and other 7TM receptors → Other 7TM proteins

Nomenclature	GPR107	GPR137	OR51E1	TPRA1	GPR143	GPR157
HGNC, UniProt	GPR107 , Q5VW38	GPR137 , Q96N19	OR51E1 , Q8TCB6	TPRA1 , Q86W33	GPR143 , P51810	GPR157 , Q5UAW9
Endogenous agonists	–	–	–	–	levodopa [1141]	–
Comments	GPR107 is a member of the LUSTR family of proteins found in both plants and animals, having similar topology to Gprotein-coupled receptors [461]	–	OR51E1 is a putative olfactory receptor.	TPRA1 shows no homology to known G protein-coupled receptors.	Loss-of-function mutations underlie ocular albinism type 1 [103].	GPR157 has ambiguous sequence similarities to several different GPCR families (class A, class B and the slime mould cyclic AMP receptor). Because of its distant relationship to other GPCRs, it cannot be readily classified.

5-Hydroxytryptamine receptors

G protein-coupled receptors → 5-Hydroxytryptamine receptors

Overview: 5-HT receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on 5-HT receptors [789] and subsequently revised [707]**) are, with the exception of the ionotropic 5-HT₃ class, GPCR receptors where the endogenous agonist is **5-hydroxytryptamine**. The diversity of metabotropic 5-HT

receptors is increased by alternative splicing that produces isoforms of the 5-HT_{2A} (non-functional), 5-HT_{2C} (non-functional), 5-HT₄, 5-HT₆ (non-functional) and 5-HT₇ receptors. Unique amongst the GPCRs, RNA editing produces 5-HT_{2C} receptor isoforms that differ in function, such as efficiency and specificity of coupling to G_{q/11}

and also pharmacology [164, 2011]. Most 5-HT receptors (except 5-HT_{1e} and 5-HT_{5a/5b}) play specific roles mediating functional responses in different tissues (reviewed by [1554, 1957]).

Nomenclature	5-HT _{1A} receptor	5-HT _{1B} receptor	5-HT _{1D} receptor	5-HT _{1e} receptor	5-HT _{1F} receptor
HGNC, UniProt	<i>HTR1A</i> , P08908	<i>HTR1B</i> , P28222	<i>HTR1D</i> , P28221	<i>HTR1E</i> , P28566	<i>HTR1F</i> , P30939
Agonists	U92016A (pK _i 9.7) [1240], vilazodone (Partial agonist) (pK _i 9.7) [402], vortioxetine (Partial agonist) (pK _i 7.8) [90]	L-694,247 (pK _i 9.2) [637], naratriptan (Partial agonist) (pK _i 8.1) [1365], eletriptan (pK _i 8) [1365], frovatriptan (pK _i 8) [2069], zolmitriptan (Partial agonist) (pK _i 7.7) [1365], vortioxetine (Partial agonist) (pK _i 7.5) [90], rizatriptan (Partial agonist) (pK _i 6.9) [1365]	dihydroergotamine (pK _i 9.2–9.9) [684, 1084, 1091], ergotamine (pK _i 9.1) [616], L-694,247 (pK _i 9) [2052], naratriptan (pK _i 8.4–9) [432, 1365, 1577], zolmitriptan (pK _i 8.9) [1365], frovatriptan (pK _i 8.4) [2069], rizatriptan (pK _i 7.9) [1365]	BRL-54443 (pK _i 8.7) [227]	BRL-54443 (pK _i 8.9) [227], eletriptan (pK _i 8) [1365], sumatriptan (pK _i 7.2–7.9) [11, 12, 1365, 1968]
Selective agonists	8-OH-DPAT (pK _i 8.4–9.4) [406, 685, 896, 1079, 1280, 1386, 1388, 1389], NLX-101 (pK _i 8.6) [1387]	CP94253 (pK _i 8.7) [976]	PNU109291 (pK _i 9.1) [483] – Gorilla, eletriptan (pK _i 8.9) [1365]	–	lasmiditan (pK _i 8.7) [1375], LY334370 (pK _i 8.7) [1968], 5-BODMT (pK _i 8.4) [966], LY344864 (pK _i 8.2) [1502]
Antagonists	(S)-UH 301 (pK _i 7.9) [1386]	–	–	–	–
Selective antagonists	WAY-100635 (pK _i 7.9–9.2) [1386, 1388], robalzotan (pK _i 9.2) [872]	SB 224289 (Inverse agonist) (pK _i 8.2–8.6) [583, 1384, 1696], SB236057 (Inverse agonist) (pK _i 8.2) [1272], GR-55562 (pK _B 7.4) [791]	SB 714786 (pK _i 9.1) [1987]	–	–

(continued)					
Nomenclature	5-HT _{1A} receptor	5-HT _{1B} receptor	5-HT _{1D} receptor	5-HT _{1E} receptor	5-HT _{1F} receptor
Labelled ligands	<p>³H]robalzotan (Antagonist) (pK_d 9.8) [861],</p> <p>³H]WAY100635 (Antagonist) (pK_d 9.5) [933],</p> <p>³H]8-OH-DPAT (Agonist) (pK_d 6–9.4) [156, 896, 1385, 1388],</p> <p>³H]NLX-112 (Agonist) (pK_d 8.9) [748], [¹¹C]WAY100635 (Antagonist) [1915],</p> <p>p-[¹⁸F]MPPF (Antagonist) [368]</p>	<p>³H]N-methyl-AZ10419369 (Agonist, Partial agonist) (pK_d 9.4) [1182], ³H]GR 125,743 (Selective Antagonist) (pK_d 8.6–9.2) [637, 2061],</p> <p>³H]alniditan (Agonist) (pK_d 8.6–9) [1084], [¹²⁵I]GTI (Agonist) (pK_d 8.9) [193, 232] – Rat, ³H]eletriptan (Agonist, Partial agonist) (pK_d 8.5) [1365], ³H]sumatriptan (Agonist, Partial agonist) (pK_d 8) [1365], [¹¹C]AZ10419369 (Agonist, Partial agonist) [1950]</p>	<p>³H]eletriptan (Agonist) (pK_d 9.1) [1365], ³H]alniditan (Agonist) (pK_d 8.8–8.9) [1084], [¹²⁵I]GTI (Selective Agonist) (pK_d 8.9) [193, 232] – Rat, ³H]GR 125,743 (Selective Antagonist) (pK_d 8.6) [2061], ³H]sumatriptan (Agonist) (pK_d 8.2) [1365]</p>	<p>³H]5-HT (Agonist) (pK_d 8.1–8.2) [1237, 1463]</p>	<p>³H]LY334370 (Agonist) (pK_d 9.4) [1968], [¹²⁵I]LSD (Agonist) (pK_d 9) [44] – Mouse</p>
Comments	–	Wang <i>et al.</i> (2013) report X-ray structures which reveal the binding modality of ergotamine and dihydroergotamine to the 5-HT _{1B} receptor in comparison with the structure of the 5-HT _{2B} receptor [1978].	–	–	–

	5-HT _{2A} receptor	5-HT _{2B} receptor	5-HT _{2C} receptor	5-HT ₄ receptor
Nomenclature	5-HT _{2A} receptor	5-HT _{2B} receptor	5-HT _{2C} receptor	5-HT ₄ receptor
HGNC, UniProt	<i>HTR2A</i> , P28223	<i>HTR2B</i> , P41595	<i>HTR2C</i> , P28335	<i>HTR4</i> , Q13639
Agonists	DOI (pK _i 7.4–9.2) [204, 1374, 1755]	methysergide (Partial agonist) (pK _i 8–9.4) [970, 1605, 1969], DOI (pK _i 7.6–7.7) [1025, 1374, 1659]	DOI (pK _i 7.2–8.6) [465, 1374, 1659], Ro 60-0175 (pK _i 7.7–8.2) [953, 970]	cisapride (Partial agonist) (pK _i 6.4–7.4) [77, 128, 597, 1266, 1267, 1941]
Selective agonists	–	BW723C86 (pK _i 7.3–8.6) [108, 970, 1659], Ro 60-0175 (pK _i 8.3) [970]	WAY-163909 (pK _i 6.7–8) [454], lorcaserin (pK _i 7.8) [1878]	TD-8954 (pK _i 9.4) [1250], ML 10302 (Partial agonist) (pK _i 7.9–9) [136, 160, 1266, 1267, 1268], RS67506 (pEC ₅₀ 8.8) [731] – Rat, relenopride (Partial agonist) (pK _i 8.3) [607], velusetrag (pK _i 7.7) [1139, 1763], BIMU 8 (pK _i 7.3) [347]
Antagonists	risperidone (Inverse agonist) (pK _i 9.3–10) [986, 1008, 1675], mianserin (pK _i 7.7–9.6) [970, 1001, 1280], ziprasidone (pK _i 8.8–9.5) [986, 1008, 1675, 1711], volinsarin (pIC ₅₀ 6.5–9.3) [970, 1142, 1568], blonsarin (pK _i 9.1) [1421], clozapine (Inverse agonist) (pK _i 7.6–9) [970, 1008, 1277, 1675, 1943], olanzapine (pK _i 8.6–8.9) [986, 1008, 1675, 1711], nefazodone (pK _i 8.2) [1698], chlorpromazine (Inverse agonist) (pK _i 8.1) [1008], loxapine (Inverse agonist) (pK _i 8.1) [1008], trifluoperazine (pK _i 7.9) [1008], pimozide (pK _i 7.1–7.7) [986, 1008], trazodone (pK _i 7.4) [970], haloperidol (pK _i 6.7–7.3) [1008, 1277, 1675, 1711, 1943], mesoridazine (pK _i 7.3) [326], mirtazapine (pK _i 7.2) [513], mirtazapine (pK _i 7.2) [513], quetiapine (pK _i 6.4–7) [986, 1008], molindone (pK _i 6.5) [1008]	mianserin (pK _i 7.9–8.8) [180, 970, 1969]	mianserin (Inverse agonist) (pK _i 8.3–9.2) [524, 970, 1280], methysergide (pK _i 8.6–9.1) [465, 970], ziprasidone (Inverse agonist) (pK _i 7.9–9) [743, 1008, 1711], olanzapine (Inverse agonist) (pK _i 8.1–8.4) [743, 1008, 1711], loxapine (Inverse agonist) (pK _i 7.8–8) [743, 1008], mirtazapine (pK _i 7.4) [513], mirtazapine (pK _i 7.4) [513], trazodone (pK _i 6.6) [970], trifluoperazine (pK _i 6.4) [1008], agomelatine (pK _i 6.2) [1276]	–

(continued)				
Nomenclature	5-HT_{2A} receptor	5-HT_{2B} receptor	5-HT_{2C} receptor	5-HT₄ receptor
Selective antagonists	ketanserin (pK _i 8.1–9.7) [234, 970, 1559], pimavanserin (Inverse agonist) (pK _i 9.3) [572, 1943]	BF-1 (pK _i 10.1) [1671], RS-127445 (pK _i 9–9.5) [180, 970], EGIS-7625 (pK _i 9) [1001]	FR260010 (pK _i 9) [700], SB 242084 (pK _i 8.2–9) [928, 970], RS-102221 (pK _i 8.3–8.4) [181, 970]	RS 100235 (pK _i 8.7–12.2) [347, 1589], SB 204070 (pK _i 9.8–10.4) [128, 1266, 1267, 1941], GR 113808 (pK _i 9.3–10.3) [77, 128, 160, 347, 1267, 1589, 1941]
Labelled ligands	[³H]fananserin (Antagonist) (pK _d 9.9) [1188] – Rat, [³H]ketanserin (Antagonist) (pK _d 8.6–9.7) [970, 1559], [¹¹C]volinanserin (Antagonist) [676], [¹⁸F]altanserin (Antagonist) [1601]	[³H]LSD (Agonist) (pK _d 8.7) [1559], [³H]5-HT (Agonist) (pK _d 8.1) [1967] – Rat, [³H]mesulergine (Antagonist, Inverse agonist) (pK _d 7.9) [970], [¹²⁵I]DOI (Agonist) (pK _d 7.7–7.6)	[¹²⁵I]DOI (Agonist) (pK _d 8.7–9) [524], [³H]mesulergine (Antagonist, Inverse agonist) (pK _d 9.3–8.7) [524, 1559], [³H]LSD (Agonist)	[¹²³I]SB 207710 (Antagonist) (pK _d 10.1) [228] – Pig, [³H]GR 113808 (Antagonist) (pK _d 10.3–9.7) [77, 128, 1268, 1941], [³H]RS 57639 (Selective Antagonist) (pK _d 9.7) [179] – Guinea pig, [¹¹C]SB207145 (Antagonist) (pK _d 8.6) [1169]
Comments	–	LSD (lysergic acid) and ergotamine show a strong preference for arrestin recruitment over G protein coupling at the 5-HT _{2B} receptor, with no such preference evident at 5-HT _{1B} receptors, and they also antagonise 5-HT _{7A} receptors [1963]. DHE (dihydroergocryptine), pergolide and cabergoline also show significant preference for arrestin recruitment over G protein coupling at 5-HT _{2B} receptors [1963].	The serotonin antagonist mesulergine was key to the discovery of the 5-HT _{2C} receptor [1479].	–

Nomenclature	5-ht_{5a} receptor	5-ht_{5b} receptor	5-HT₆ receptor	5-HT₇ receptor
HGNC, UniProt	HTR5A, P47898	HTR5BP, –	HTR6, P50406	HTR7, P34969
Selective agonists	–	–	WAY-181187 (pK _i 8.7) [1663], E6801 (Partial agonist) (pK _i 8.7) [769], WAY-208466 (pK _i 8.3) [135], EMD-386088 (pIC ₅₀ 8.1) [1228]	LP-12 (pK _i 9.9) [1082], LP-44 (pK _i 9.7) [1082], LP-211 (pK _i 9.2) [1083] – Rat, AS-19 (pK _i 9.2) [947], E55888 (pK _i 8.6) [206]

(continued)	5-HT _{5A} receptor	5-HT _{5B} receptor	5-HT ₆ receptor	5-HT ₇ receptor
Nomenclature				
Antagonists	–	–	–	lurasidone (pK _i 9.3) [829], pimozone (pK _i 9.3) [1604] – Rat, vortioxetine (pK _i 6.3) [90]
Selective antagonists	SB 699551 (pK _i 8.2) [366]	–	SB399885 (pK _i 9) [763], SB 271046 (pK _i 8.9) [224], cerlapirdine (pK _i 8.9) [358], SB357134 (pK _i 8.5) [225], Ro 63-0563 (pK _i 7.9–8.4) [168, 1754]	SB269970 (pK _i 8.6–8.9) [1874], SB656104 (pK _i 8.7) [531], DR-4004 (pK _i 8.7) [615, 938], JNJ-18038683 (pK _i 8.2) [177], SB 258719 (Inverse agonist) (pK _i 7.5) [1875]
Labelled ligands	[¹²⁵ I]LSD (Agonist) (pK _d 9.7) [636], [³ H]5-CT (Agonist) (pK _d 8.6) [636]	[¹²⁵ I]LSD (Agonist) (pK _d 9.3) [1227] – Mouse, [³ H]5-CT (Agonist) [1965] – Mouse	[¹¹ C]GSK215083 (Antagonist) (pK _i 9.8) [1462], [¹²⁵ I]SB258585 (Selective Antagonist) (pK _d 9) [763], [³ H]LSD (Agonist) (pK _d 8.7) [167], [³ H]Ro 63-0563 (Antagonist) (pK _d 8.3) [168], [³ H]5-CT (Agonist)	[³ H]5-CT (Agonist) (pK _d 9.4) [1874], [³ H]5-HT (Agonist) (pK _d 8.1–9) [93, 1793], [³ H]SB269970 (Selective Antagonist) (pK _d 8.9) [1874], [³ H]LSD (Agonist) (pK _d 8.5–8.6) [1793]

Comments: Tabulated pK_i and K_D values refer to binding to human 5-HT receptors unless indicated otherwise. The nomenclature of 5-HT_{1B}/5-HT_{1D} receptors has been revised [707]. Only the non-rodent form of the receptor was previously called 5-HT_{1D}: the human 5-HT_{1B} receptor (tabulated) displays a different pharmacology to the rodent forms of the receptor due to Thr335 of the human

sequence being replaced by Asn in rodent receptors. NAS181 is a selective antagonist of the rodent 5-HT_{1B} receptor. Fananserlin and ketanserlin bind with high affinity to dopamine D4 and histamine H₁ receptors respectively, and ketanserlin is a potent α₁ adrenoceptor antagonist, in addition to blocking 5-HT_{2A} receptors. The human 5-HT_{5A} receptor has been claimed to couple to several signal transduction pathways when stably expressed in C6 glioma cells [1404]. The

human orthologue of the mouse 5-HT_{5B} receptor is non-functional due to interruption of the gene by stop codons. The 5-HT_{1E} receptor appears not to have been cloned from mouse, or rat, impeding definition of its function. In addition to the receptors listed in the table, an 'orphan' receptor, unofficially termed 5-HT_{1P}, has been described [600].

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Acetylcholine receptors (muscarinic)

G protein-coupled receptors → Acetylcholine receptors (muscarinic)

Overview: Muscarinic acetylcholine receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Muscarinic Acetylcholine Receptors [275]**) are GPCRs of the Class A, rhodopsin-like family where the endogenous agonist is **acetylcholine**. In addition to the agents listed in the table,

AC-42, its structural analogues **AC-260584** and **77-LH-28-1**, **N-desmethylozapine**, **TBPB** and **LuAE51090** have been described as functionally selective agonists of the M₁ receptor subtype *via* binding in a mode distinct from that utilized by non-selective agonists [71, 878, 1040, 1041, 1232, 1635, 1786, 1787, 1825]. There are

two pharmacologically characterised allosteric sites on muscarinic receptors, one defined by it binding **gallamine**, **strychnine** and **brucine**, and the other defined by the binding of **KT 5720**, **WIN 62,577**, **WIN 51,708** and **staurosporine** [1052, 1053].

Nomenclature	M ₁ receptor	M ₂ receptor
HGNC, UniProt	<i>CHRM1</i> , P11229	<i>CHRM2</i> , P08172
Agonists	carbachol (pK _i 3.2–5.3) [334, 846, 2040], pilocarpine (Partial agonist) (pK _i 5.1) [846], bethanechol (pK _i 4) [846]	bethanechol (pK _i 4) [846]
Antagonists	glycopyrrolate (pIC ₅₀ 9.9) [1801], umeclidinium (pK _i 9.8) [1035, 1632], AE9C90CB (pK _i 9.7) [1749], propranolol (pK _i 9.7) [797], atropine (pK _i 8.5–9.6) [334, 552, 759, 797, 1486, 1762], tiotropium (pK _i 9.6) [428], 4-DAMP (pK _i 9.2) [458], dicyclomine (pK _i 9.1) [68], scopolamine (pK _i 9) [797], trihexyphenidyl (pK _i 8.9) [68], tripitramine (pK _i 8.8) [1176], UH-AH 37 (pK _i 8.6–8.7) [609, 2012], tolterodine (pK _i 8.5–8.7) [609, 1749], oxybutynin (pK _i 8.6) [410, 818, 1749], darifenacin (pK _i 7.5–8.3) [609, 730, 759, 818, 1749], pirenzepine (pK _i 7.8–8.3) [238, 458, 730, 797, 875, 2012], solifenacin (pK _i 7.6) [818, 1749], AFDX384 (pK _i 7.5) [458], AQ-RA 741 (pK _i 7.2–7.5) [458, 609], methoctramine (pK _i 6.6–7.3) [458, 493, 730, 1762], himbacine (pK _i 6.7–7.1) [458, 875, 1286], muscarinic toxin 3 (pK _i 7.1) [875], otenzepad (pK _d 6.2) [493]	tiotropium (pK _i 9.9) [428], umeclidinium (pK _i 9.8) [1035, 1632], propranolol (pK _i 9.5) [797], glycopyrrolate (Full agonist) (pIC ₅₀ 9.3) [1801], atropine (pK _i 7.8–9.2) [238, 310, 759, 797, 1002, 1373, 1486], AE9C90CB (pK _i 8.6) [1749], tolterodine (Inverse agonist) (pK _i 8.4–8.6) [609, 1373, 1749], AQ-RA 741 (pK _i 8.4) [458, 609], himbacine (pK _i 7.9–8.4) [458, 875, 1002, 1286], methoctramine (pK _i 7.3–8.4) [238, 458, 493, 730, 1002, 1373], 4-DAMP (pK _i 8.3) [1002], AFDX384 (pK _i 8.2) [458], biperiden (pK _d 8.2) [173], oxybutynin (pK _i 7.7–8.1) [818, 1749], darifenacin (Inverse agonist) (pK _i 7–7.6) [609, 730, 759, 818, 1373, 1749], UH-AH 37 (pK _i 7.3–7.4) [609, 2012], otenzepad (pK _i 6.7–7.2) [238, 1002], solifenacin (pK _i 6.9–7.1) [818, 1749], pirenzepine (pK _i 6–6.7) [238, 458, 730, 797, 875, 1002, 1373, 2012], VU0255035 (pK _i 6.2) [1717], muscarinic toxin 3 (pK _i <6) [875], guanylpirenzepine (pK _i 5.3) [1966] – Rat, muscarinic toxin 7 (pK _i <5) [1414]
Selective antagonists	biperiden (pK _d 9.3) [173], VU0255035 (pK _i 7.8) [1717], guanylpirenzepine (pK _i 7.3–7.6) [23, 1966] – Rat	tripitramine (pK _i 9.6) [1176]
Allosteric modulators	muscarinic toxin 7 (Negative) (pK _i 11–11.1) [1414], benzoquinazolinone 12 (Positive) (pK _B 6.6) [4], KT 5720 (Positive) (pK _d 6.4) [1052], brucine (Positive) (pK _d 4.5–5.8) [846, 1051], BQCA (Positive) (pK _B 4–4.8) [4, 5, 261, 1161], VU0029767 (Positive) [1208], VU0090157 (Positive) [1208]	W-84 (Negative) (pK _d 6–7.5) [1299, 1908], C₇/3-phth (Negative) (pK _d 7.1) [335], alcuronium (Negative) (pK _d 6.1–6.9) [846, 1908], gallamine (Negative) (pK _d 5.9–6.3) [348, 1049], LY2119620 (Positive) (pK _d 5.7) [383, 1010], LY2033298 (Positive) (pK _d 4.4) [1933]
Labelled ligands	[³H]QNB (Antagonist) (pK _d 10.6–10.8) [336, 1486], Cy3B-telenzepine (Antagonist) (pK _d 10.5) [742], [³H]N-methyl scopolamine (Antagonist) (pK _d 9.4–10.3) [280, 334, 336, 759, 846, 847, 875, 932, 1049], [³H](+)-telenzepine (Antagonist) (pK _i 9.4) [500] – Rat, Alexa-488-telenzepine (Antagonist) (pK _d 9.3) [742], [³H]pirenzepine (Antagonist) (pK _d 7.9) [1995], BODIPY-pirenzepine (Antagonist) (pK _i 7) [820], [¹¹C]butylthio-TZTP (Agonist) [504], [¹¹C]xanomeline (Agonist) [504], [¹⁸F](R,R)-quinuclidinyl-4-fluoromethyl-benzilate (Antagonist) [935] – Rat	[³H]QNB (Antagonist) (pK _d 10.1–10.6) [1486], Cy3B-telenzepine (Antagonist) (pK _i 10.4) [1380], [³H]tiotropium (Antagonist) (pK _d 10.3) [1632], [³H]N-methyl scopolamine (Antagonist) (pK _d 9.3–9.9) [280, 310, 759, 846, 847, 875, 932, 1049, 1985], Alexa-488-telenzepine (Antagonist) (pK _i 8.8) [1380], [³H]acetylcholine (Agonist) (pK _d 8.8) [1050], [³H]oxotremorine-M (Agonist) (pK _d 8.7) [137], [³H]dimethyl-W84 (Allosteric modulator, Positive) (pK _d 8.5) [1908], [¹⁸F]FP-TZTP (Agonist) [845] – Mouse

Nomenclature	M₃ receptor	M₄ receptor	M₅ receptor
HGNC, UniProt	CHRM3, P20309	CHRM4, P08173	CHRM5, P08912
Agonists	pilocarpine (Partial agonist) (pK _i 5.1) [846], carbachol (pK _i 4–4.4) [310, 846, 2040], bethanechol (pK _i 4.2) [846]	pilocarpine (Partial agonist) (pK _i 5.2) [846], carbachol (pK _i 4.3–4.9) [846, 2040], bethanechol (pK _i 4) [846]	pilocarpine (Partial agonist) (pK _i 5) [639], carbachol (pK _i 4.9) [2040]
Antagonists	tiotropium (pK _i 9.5–11.1) [428, 442], umeclidinium (pK _i 10.2) [1035, 1632], proprantheline (pK _i 10) [797], AE9C90CB (pK _i 9.9) [1749], atropine (pK _i 8.9–9.8) [238, 442, 759, 797, 1486, 1762], ipratropium (pK _i 9.3–9.8) [442, 759], acridinium (pIC ₅₀ 9.8) [1530], clidinium (pK _i 9.6) [442], glycopyrrolate (pIC ₅₀ 9.6) [1801], 4-DAMP (pK _i 9.3) [458], darifenacin (pK _i 8.6–9.1) [609, 730, 759, 818, 1749], dicyclomine (pK _i 9) [68], oxybutynin (pK _i 8.8–8.9) [410, 818, 1749], tolterodine (pK _i 8.4–8.5) [609, 1749], biperiden (pK _d 8.4) [173], UH-AH 37 (pK _i 8.1–8.2) [609, 2012], solifenacin (pK _i 7.7–8) [818, 1749], tropicamide (pK _i 7.5) [68], AQ-RA 741 (pK _i 7.2–7.3) [458, 609], AFDX384 (pK _i 7.2) [458], himbacine (pK _i 6.9–7.2) [458, 875, 1286], methoctramine (pK _i 6.1–6.9) [238, 458, 493, 730, 1762], tripitramine (pK _i 6.8) [1288], pirenzepine (pK _i 6.5–6.8) [238, 458, 730, 797, 875, 2012], guanylpirenzepine (pK _i 6.2) [1966] – Rat, VU0255035 (pK _i 6.1) [1717], otenzepad (pK _i 6.1) [238], muscarinic toxin 3 (pK _i <6) [875], muscarinic toxin 7 (pK _i <5) [1414]	umeclidinium (pK _i 10.3) [1632], glycopyrrolate (pIC ₅₀ 9.8) [1801], AE9C90CB (pK _i 9.5) [1749], 4-DAMP (pK _i 8.9) [458], oxybutynin (pK _i 8.7) [1749], biperiden (pK _d 8.6) [173], UH-AH 37 (pK _i 8.3–8.4) [609, 2012], tolterodine (pK _i 8.3–8.4) [609, 1749], AQ-RA 741 (pK _i 7.8–8.2) [458, 609], himbacine (pK _i 7.9–8.2) [458, 875, 1286], darifenacin (pK _i 7.3–8.1) [609, 730, 759, 1749], AFDX384 (pK _i 8) [458], tripitramine (pK _i 7.9) [1288], pirenzepine (pK _i 7–7.6) [458, 730, 797, 875, 2012], methoctramine (pK _i 6.6–7.5) [238, 458, 493, 730], otenzepad (pK _d 7) [493], solifenacin (pK _i 6.8) [1749], guanylpirenzepine (pK _i 6.2) [1966] – Rat, VU0255035 (pK _i 5.9) [1717], muscarinic toxin 7 (pK _i <5) [1414]	umeclidinium (pK _i 9.9) [1632], glycopyrrolate (pIC ₅₀ 9.7) [1801], AE9C90CB (pK _i 9.5) [1749], 4-DAMP (pK _i 9) [458], tolterodine (pK _i 8.5–8.8) [609, 1749], darifenacin (pK _i 7.9–8.6) [609, 730, 759, 1749], UH-AH 37 (pK _i 8.3) [609, 2012], biperiden (pK _d 8.2) [173], oxybutynin (pK _i 7.9) [1749], AQ-RA 741 (pK _i 6.1–7.8) [458, 609], tripitramine (pK _i 7.5) [1176], methoctramine (pK _i 6.3–7.2) [238, 458, 493, 730], solifenacin (pK _i 7.2) [1749], pirenzepine (pK _i 6.8–7.1) [730, 875, 2012], guanylpirenzepine (pK _d 6.8) [514] – Unknown, himbacine (pK _i 5.4–6.5) [458, 875, 1286], AFDX384 (pK _i 6.3) [458], muscarinic toxin 3 (pK _i <6) [875], otenzepad (pK _i 5.6) [238], muscarinic toxin 7 (pK _i <5) [1414]
Selective antagonists	–	–	ML381 (pK _i 6.3) [593]
Allosteric modulators	WIN 62,577 (Positive) (pK _d 5.1) [1053], N-chloromethyl-brucine (Positive) (pK _d 3.3) [1051]	muscarinic toxin 3 (Negative) (pK _i 8.7) [875, 1444], LY2119620 (Positive) (pK _d 5.7) [383], thiochrome (Positive) (pK _d 4) [1050], LY2033298 (Positive) [286], VU0152099 (Positive) [201], VU0152100 (Positive) [201]	ML380 (Positive) (pEC ₅₀ 6.7) [595]
Selective allosteric modulators	–	–	ML375 (Negative) (pIC ₅₀ 6.5) [594]
Labelled ligands	[³H]tiotropium (Antagonist) (pK _d 10.7) [1632], [³H]QNB (Antagonist) (pK _d 10.4) [1486], [³H]N-methyl scopolamine (Antagonist) (pK _d 9.7–10.2) [280, 310, 759, 797, 846, 875, 932, 1049], [³H]darifenacin (Antagonist) (pK _d 9.5) [1762]	[³H]QNB (Antagonist) (pK _d 9.7–10.5) [336, 1486], [³H]N-methyl scopolamine (Antagonist) (pK _d 9.9–10.2) [280, 310, 336, 759, 846, 875, 932, 1049, 1444, 1985], [³H]acetylcholine (Agonist) (pK _d 8.2) [1050]	[³H]QNB (Antagonist) (pK _d 10.2–10.7), [³H]N-methyl scopolamine (Antagonist) (pK _d 9.3–9.7) [280, 310, 759, 875, 932, 1985]

Comments: LY2033298 and BQCA have also been shown to directly activate the M₄ and M₁ receptors, respectively, via an allosteric site [1059, 1060, 1366, 1367]. The allosteric site for gallamine and strychnine on M₂ receptors can be labelled by [³H]dimethyl-W84 [1908]. McN-A-343 is a functionally selective partial agonist that appears to interact in a bitopic mode with both the orthosteric and an allosteric site on the M₂ muscarinic recep-

tor [1934]. THR160209, hybrid 1 and hybrid 2, are multivalent (bitopic) ligands that also achieve selectivity for M₂ receptors by binding both to the orthosteric and a nearby allosteric site [52, 1796].

Although numerous ligands for muscarinic acetylcholine receptors have been described, relatively few selective antagonists have been described, so it is common to assess the rank order of affinity

of a number of antagonists of limited selectivity (*e.g.* 4-DAMP, darifenacin, pirenzepine) in order to identify the involvement of particular subtypes. It should be noted that the measured affinities of antagonists (and agonists) in radioligand binding studies are sensitive to ionic strength and can increase over 10-fold at low ionic strength compared to its value at physiological ionic strengths [151].

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Adenosine receptors

G protein-coupled receptors → Adenosine receptors

Overview: Adenosine receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Adenosine Receptors** [541]) are activated by the endogenous ligand adenosine (potentially inosine also at A₃ receptors). Crystal structures for the antagonist-bound and agonist-bound A_{2A} adenosine receptors have been described [835, 2065].

Nomenclature	A ₁ receptor	A _{2A} receptor	A _{2B} receptor	A ₃ receptor
HGNC, UniProt	<i>ADORA1</i> , P30542	<i>ADORA2A</i> , P29274	<i>ADORA2B</i> , P29275	<i>ADORA3</i> , PODMS8
Agonists	cyclopentyladenosine (pK _i 6.5–9.4) [388, 570, 736, 839, 870, 1592, 2141]	–	–	–
(Sub)family-selective agonists	NECA (pK _i 5.3–8.2) [570, 870, 1592, 1903, 2077]	NECA (pK _i 6.9–8.7) [189, 427, 570, 943, 1021, 2077]	NECA (pK _i 5.7–6.9) [146, 189, 862, 1113, 1798, 1945, 2077]	NECA (pK _i 7.5–8.4) [189, 570, 840, 1634, 1946, 2077]
Selective agonists	5-Cl-5-deoxy-(±)-ENBA (pK _i 9.3) [536], GR79236 (pK _i 8.5) [839] – Rat, CCPA (pK _i 7.7–8.1) [839, 1419]	apadenoson (pK _i 9.3) [1481], UK-432,097 (pK _i 8.3) [2065], CGS 21680 (pK _i 6.7–8.1) [189, 427, 570, 839, 943, 968, 1021, 1419], regadenoson (pK _i 6.5) [839]	BAY 60-6583 (pK _i 8–8.5) [460]	IB-MECA (pK _i 8.7–9.2) [511, 561, 968, 1946], CI-IB-MECA (pK _i 8–8.9) [202, 840, 941], MRS5698 (pK _i 8.5) [1898]
Antagonists	caffeine (pK _i 4.3–5) [6, 409, 838]	SCH 58261 (pK _i 8.3–9.2) [427, 1021, 1445], theophylline (pK _i 5.2–5.8) [427, 838, 968, 1945], caffeine (pK _i 4.6–5.6) [6, 838, 1021]	theophylline (pK _i 4.1–5) [141, 511, 944, 1945], caffeine (pK _i 4.5–5) [141, 188, 944]	caffeine (pK _i 4.9) [838]
(Sub)family-selective antagonists	CGS 15943 (pK _i 8.5) [1445], xanthine amine congener (pK _d 7.5) [536]	CGS 15943 (pK _i 7.7–9.4) [427, 943, 968, 1445], xanthine amine congener (pK _i 8.4–9) [427, 968]	xanthine amine congener (pK _i 6.9–8.8) [146, 862, 863, 968, 1113, 1798], CGS 15943 (pK _i 6–8.1) [65, 862, 863, 968, 1445, 1798]	CGS 15943 (pK _i 7–7.9) [949, 968, 1445, 1946], xanthine amine congener (pK _i 7–7.4) [968, 1634, 1946]
Selective antagonists	PSB36 (pK _i 9.9) [6] – Rat, DPCPX (pK _i 7.4–9.2) [826, 1419, 1592, 2015, 2141], derenofylline (pK _i 9) [897], WRC-0571 (pK _i 8.8) [1210]	SCH442416 (pK _i 8.4–10.3) [1728, 1891], ZM-241385 (pK _i 8.8–9.1) [1445]	PSB-0788 (pK _i 9.4) [188], PSB603 (pK _i 9.3) [188], MRS1754 (pK _i 8.8) [862, 948], PSB1115 (pK _i 7.3) [723]	MRS1220 (pK _i 8.2–9.2) [840, 949, 1818, 2090], VUF5574 (pK _i 8.4) [2143], MRS1523 (pK _i 7.7) [1092], MRS1191 (pK _i 7.5) [840, 866, 1097]
Labelled ligands	[³ H]CCPA (Agonist) (pK _d 9.2) [968, 1592], [³ H]DPCPX (Antagonist) (pK _d 8.4–9.2) [388, 511, 968, 1445, 1592, 1903]	[³ H]ZM 241385 (Antagonist) (pK _d 8.7–9.1) [36, 569], [³ H]CGS 21680 (Agonist) (pK _d 7.7–7.8) [852, 1976]	[³ H]MRS1754 (Antagonist) (pK _d 8.8) [862]	[¹²⁵ I]AB-MECA (Agonist) (pK _d 9–9.1) [1445, 1946]

Comments: Adenosine inhibits many intracellular ATP-utilising enzymes, including adenylyl cyclase (P-site). A pseudogene exists for the A_{2B} adenosine receptor (*ADORA2BP1*) with 79% identity to the A_{2B} adenosine receptor cDNA coding sequence, but which is unable to encode a functional receptor [841]. DPCPX also exhibits antagonism at A_{2B} receptors (pK_i ca. 7, [34, 968]). Antagonists at A₃ recep-

tors exhibit marked species differences, such that only MRS1523 and MRS1191 are selective at the rat A₃ receptor. In the absence of other adenosine receptors, [³H]DPCPX and [³H]ZM 241385 can also be used to label A_{2B} receptors (K_D ca. 30 and 60 nM respectively). [¹²⁵I]AB-MECA also binds to A₁ receptors [968]. [³H]CGS 21680 is relatively selective for A_{2A} receptors, but may also bind to other sites

in cerebral cortex [384, 871]. [³H]NECA binds to other non-receptor elements, which also recognise adenosine [1143]. XAC-BY630 has been described as a fluorescent antagonist for labelling A₁ adenosine receptors in living cells, although activity at other adenosine receptors was not examined [212].

Further Reading

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Adhesion Class GPCRs

G protein-coupled receptors → Adhesion Class GPCRs

Overview: Adhesion GPCRs are structurally identified on the basis of a large extracellular region, similar to the Class B GPCR, but which is linked to the 7TM region by a "stalk" motif containing a GPCR proteolytic site. The N-terminus often shares structural homology with proteins such as lectins and immunoglobulins, leading to the term adhesion GPCR [543, 2097]. **The nomenclature of these receptors was revised in 2015 as recommended by NC-IUPHAR and the Adhesion GPCR Consortium [683].**

Nomenclature	ADGRA1	ADGRA2	ADGRA3	ADGRB1	ADGRB2
HGNC, UniProt	ADGRA1, Q86SQ6	ADGRA2, Q96PE1	ADGRA3, Q8IWK6	ADGRB1, O14514	ADGRB2, O60241
Endogenous agonists	–	–	–	phosphatidylserine [1461]	–
Comments	–	–	–	ADGRB1 is reported to respond to phosphatidylserine [1461].	–

Nomenclature	ADGRB3	CELSR1	CELSR2	CELSR3	ADGRD1
HGNC, UniProt	ADGRB3, O60242	CELSR1, Q9NYQ6	CELSR2, Q9HCU4	CELSR3, Q9NYQ7	ADGRD1, Q6QNK2

Nomenclature	ADGRD2	ADGRE1	ADGRE2	ADGRE3	ADGRE4P
HGNC, UniProt	ADGRD2, Q7Z7M1	ADGRE1, Q14246	ADGRE2, Q9UHX3	ADGRE3, Q9BY15	ADGRE4P, Q86SQ3

Nomenclature	ADGRES	ADGRF1	ADGRF2	ADGRF3	ADGRF4
HGNC, UniProt	ADGRES , P48960	ADGRF1 , Q5T601	ADGRF2 , Q8IZF7	ADGRF3 , Q8IZF5	ADGRF4 , Q8IZF3

Nomenclature	ADGRF5	ADGRG1	ADGRG2	ADGRG3	ADGRG4
HGNC, UniProt	ADGRF5 , Q8IZF2	ADGRG1 , Q9Y653	ADGRG2 , Q8IZP9	ADGRG3 , Q86Y34	ADGRG4 , Q8IZF6
Comments	–	Reported to bind tissue transglutaminase 2 [2066] and collagen, which activates the G _{12/13} pathway [1155].	–	–	–

Nomenclature	ADGRG5	ADGRG6	ADGRG7	ADGRL1
HGNC, UniProt	ADGRG5 , Q8IZF4	ADGRG6 , Q86SQ4	ADGRG7 , Q96K78	ADGRL1 , O94910

Nomenclature	ADGRL2	ADGRL3	ADGRL4	ADGRV1
HGNC, UniProt	ADGRL2 , O95490	ADGRL3 , Q9HAR2	ADGRL4 , Q9HBW9	ADGRV1 , Q8WXG9
Comments	–	–	–	Loss-of-function mutations are associated with Usher syndrome, a sensory deficit disorder [843].

Adrenoceptors

G protein-coupled receptors → Adrenoceptors

Overview:

Adrenoceptors, α_1

α_1 -Adrenoceptors (nomenclature as agreed by the **NC-IUPHAR Subcommittee on Adrenoceptors** [248], see also [752]) are activated by the endogenous agonists (-)-adrenaline and (-)-noradrenaline with equal potency. Phenylephrine, methoxamine and cirazoline are agonists selective for α_1 -adrenoceptors relative to α_2 -adrenoceptors, while prazosin (8.5–10.5) and corynanthine (6.5–7.5) are antagonists con-

sidered selective for α_1 -adrenoceptors relative to α_2 -adrenoceptors. [³H]prazosin (0.25 nM) and [¹²⁵I]HEAT (0.1 nM; also known as BE2254) are relatively selective radioligands. The α_{1A} -adrenoceptor antagonist S(+)-niguldipine also has high affinity for L-type Ca²⁺ channels. The conotoxin rho-TIA acts as a negative allosteric modulator at the α_{1B} -adrenoceptor [1716], while the snake toxin ρ -Da1a acts as a selective non-competitive antagonist at the α_{1A} -adrenoceptor [1236, 1548]. Fluorescent derivatives of prazosin (Bodipy PL-prazosin-QAPB) are increasingly used to examine cellular localisation of α_1 -adrenoceptors. The vasoconstrictor effects of

selective α_1 -adrenoceptor agonists have led to their use as nasal decongestants; antagonists are used to treat hypertension (doxazosin, prazosin) and benign prostatic hyperplasia (alfuzosin, tamsulosin). The combined α_1 - and β_2 -adrenoceptor antagonist carvedilol is widely used to treat congestive heart failure, although the contribution of α_1 -adrenoceptor blockade to the therapeutic effect is unclear. Several anti-depressants and anti-psychotic drugs possess α_1 -adrenoceptor blocking properties that are believed to contribute to side effects such as orthostatic hypotension and extrapyramidal effects.

	α_{1A} -adrenoceptor	α_{1B} -adrenoceptor	α_{1D} -adrenoceptor
Nomenclature	α_{1A} -adrenoceptor	α_{1B} -adrenoceptor	α_{1D} -adrenoceptor
HGNC, UniProt	ADRA1A, P35348	ADRA1B, P35368	ADRA1D, P25100
Endogenous agonists	–	–	(-)-adrenaline (pK _i 7.2) [1722]
Agonists	oxymetazoline (pK _i 8–8.2) [780, 1420, 1722, 1862], phenylephrine (pK _i 5.2–5.4) [1862], methoxamine (pK _i 5–5.2) [1722, 1862]	phenylephrine (pIC ₅₀ 6.3–7.5) [532, 1289]	–
Selective agonists	A61603 (pIC ₅₀ 7.8–8.4) [532, 969], dabuzalgron (pK _i 7.4) [162]	–	–
Antagonists	prazosin (Inverse agonist) (pK _i 9–9.9) [289, 389, 532, 1722, 2029], doxazosin (pK _i 9.3) [689], terazosin (pK _i 8.7) [1263], phentolamine (pK _i 8.6) [1722], alfuzosin (pK _i 8.1) [750]	prazosin (Inverse agonist) (pK _i 9.6–9.9) [532, 1722, 2029], tamsulosin (Inverse agonist) (pK _i 9.5–9.7) [532, 1722, 2029], doxazosin (pK _i 9.1) [689], alfuzosin (pK _i 8.6) [751], terazosin (pK _i 8.6) [1263], phentolamine (pK _i 7.5) [1722]	prazosin (Inverse agonist) (pK _i 9.5–10.2) [532, 1722, 2029], tamsulosin (pK _i 9.8–10.2) [532, 1722, 2029], doxazosin (pK _i 9.1) [689], terazosin (pK _i 9.1) [1263], alfuzosin (pK _i 8.4) [750], dapiprazole (pK _i 8.4) [68], phentolamine (Inverse agonist) (pK _i 8.2) [1722], RS-100329 (pK _i 7.9) [2029], labetalol (pK _i 6.6) [68]
Selective antagonists	tamsulosin (pK _i 10–10.7) [289, 389, 532, 1722, 2029], silodosin (pK _i 10.4) [1722], S(+)-niguldipine (pK _i 9.1–10) [532, 1722], RS-100329 (pK _i 9.6) [2029], SNAP5089 (pK _i 8.8–9.4) [750, 1081, 2014], ρ -Da1a (pK _i 9.2–9.3) [1236, 1548], RS-17053 (pK _i 9.2–9.3) [289, 389, 529, 532]	Rec 15/2615 (pK _i 9.5) [1867], L-765314 (pK _i 7.7) [1470], AH 11110 (pK _i 7.5) [1652]	BMY-7378 (pK _i 8.7–9.1) [268, 2102]

Adrenoceptors, α_2

α_2 -Adrenoceptors (**nomenclature as agreed by NC-IUPHAR Subcommittee on Adrenoceptors, [248]**) are activated by endogenous agonists with a relative potency of (-)-adrenaline > (-)-noradrenaline. **Brimonidine** and **talipexole** are agonists selective for α_2 -adrenoceptors relative to α_1 -adrenoceptors, **rauwolscine** and **yohimbine** are antagonists selective for α_2 -adrenoceptors relative to α_1 -adrenoceptors. [³H]rauwolscine (1 nM), [³H]brimonidine (5 nM) and [³H]RX821002 (0.5 nM and 0.1 nM at α_{2C}) are relatively selective radioligands. There is species variation in the pharmacology of the α_{2A} -adrenoceptor; for example, **yohimbine**, **rauwolscine** and **oxymetazoline** have an ~20-fold higher affinity for the human α_{2A} -adrenoceptor compared to the rat, mouse and bovine recep-

tor. These α_{2A} orthologues are sometimes referred to as α_{2D} -adrenoceptors. Multiple mutations of α_2 -adrenoceptors have been described, some of which are associated with alterations in function. Presynaptic α_2 -adrenoceptors are widespread in the nervous system and regulate many functions, hence the multiplicity of actions. The effects of classical (not subtype selective) α_2 -adrenoceptor agonists such as **clonidine**, **guanabenz** and **brimonidine** on central baroreflex control (hypotension and bradycardia), as well as their ability to induce hypnotic effects and analgesia, and their ability to modulate seizure activity and platelet aggregation are mediated by α_{2A} -adrenoceptors. **Clonidine** has been used as an anti-hypertensive and also to counteract opioid withdrawal. Actions on imidazoline recognition sites may contribute to the pharmacological effects of cloni-

dine. α_2 -Adrenoceptor agonists such as **dexmedetomidine** have been widely used as sedatives and analgesics in veterinary medicine (also **xylazine**) and are now used frequently in humans. **Dexmedetomidine** also has analgesic, sympatholytic and anxiolytic properties but is notable for the production of sedation without respiratory depression. α_2 -Adrenoceptor antagonists are relatively little used therapeutically although **yohimbine** has been used to treat erectile dysfunction and several anti-depressants (e.g. **Mirtazapine**) that block α_2 -adrenoceptors may work through this mechanism. The roles of α_{2B} and α_{2C} -adrenoceptors are less clear but the α_{2B} subtype appears to be involved in neurotransmission in the spinal cord and α_{2C} in regulating catecholamine release from adrenal chromaffin cells.

Nomenclature	α_{2A} -adrenoceptor	α_{2B} -adrenoceptor	α_{2C} -adrenoceptor
HGNC, UniProt	ADRA2A, P08913	ADRA2B, P18089	ADRA2C, P18825
Endogenous agonists	(-)-adrenaline (pK _i 5.6–8.3) [854, 1503]	(-)-noradrenaline (Partial agonist) (pK _i 5.6–9.1) [854, 1484, 1503]	(-)-noradrenaline (pK _i 5.9–8.7) [854, 1484, 1503]
Agonists	dexmedetomidine (Partial agonist) (pK _i 7.6–9.6) [854, 1163, 1484, 1503], clonidine (Partial agonist) (pK _i 7.2–9.2) [854, 1484, 1503], brimonidine (pK _i 6.7–8.7) [854, 1163, 1484, 1503], apraclonidine (pK _i 8.5) [1343], guanabenz (pIC ₅₀ 7.4) [68], guanfacine (Partial agonist) (pK _i 7.1–7.3) [854, 1165]	dexmedetomidine (pK _i 7.5–9.7) [854, 1163, 1484, 1503], clonidine (Partial agonist) (pK _i 6.7–9.5) [854, 1484, 1503], brimonidine (Partial agonist) (pK _i 6–8.3) [854, 1484, 1503], guanabenz (pIC ₅₀ 6.8) [68], guanfacine (pK _i 5.8–6.5) [854]	dexmedetomidine (pK _i 7–9.3) [854, 1484, 1503], brimonidine (Partial agonist) (pK _i 5.7–7.6) [854, 1163, 1484, 1503], apraclonidine (pK _i 7.5) [1343], guanfacine (Partial agonist) (pK _i 5.4–6.2) [854], guanabenz (pIC ₅₀ 6) [68]
Selective agonists	oxymetazoline (Partial agonist) (pK _i 8–8.6) [854, 1163, 1922]	–	–
Antagonists	yohimbine (pK _i 8.5–9.5) [247, 416, 1922], WB 4101 (pK _i 8.4–9.4) [247, 416, 1922], spiroxatrine (pK _i 9) [1922], mirtazapine (pK _i 7.7) [513], tolazoline (pK _i 5.4) [854]	yohimbine (pK _i 8.4–9.2) [247, 416, 1922]	yohimbine (pK _i 7.9–8.9) [247, 416, 1922], phenoxybenzamine (pK _i 8.5) [2002], tolazoline (pK _i 5.5) [854]
Selective antagonists	BRL 44408 (pK _i 8.2–8.8) [1922, 2104]	imiloxan (pK _i 7.3) [1269] – Rat	JP1302 (pK _B 7.8) [1631]
Labelled ligands	–	–	[³ H]MK-912 (Antagonist) (pK _d 10.1) [1922]

Adrenoceptors, β

β -Adrenoceptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Adrenoceptors, [248]**) are activated by the endogenous agonists (-)-adrenaline and (-)-noradrenaline. **Isoprenaline** is a synthetic agonist selective for β -adrenoceptors relative to α_1 - and α_2 -adrenoceptors, while for β_1 and β_2 adrenoceptors, **propranolol** (pK_i 8.2–9.2) and **cyanopindolol** (pK_i 10.0–11.0) are relatively selective antagonists. (-)-noradrenaline, **xamoterol** and (-)-Ro 363 are agonists that show selectivity for β_1 -

relative to β_2 -adrenoceptors. Pharmacological differences exist between human and mouse β_3 -adrenoceptors, and the 'rodent selective' agonists **BRL 37344** and **CL316243** have low efficacy at the human β_3 -adrenoceptor whereas **CGP 12177** and **L 755507** activate human β_3 -adrenoceptors [1649]. β_3 -Adrenoceptors are relatively resistant to blockade by **propranolol** (pK_i 5.8–7.0), but can be blocked by high concentrations of **bupranolol** (pK_i 8.65 [1650]). **SR59230A** has reasonably high affinity at β_3 -adrenoceptors [1197],

but does not discriminate well between the three β -adrenoceptor subtypes [262] and has been reported to have lower affinity for the β_3 -adrenoceptor in some circumstances [913]. **L-748337** is the most selective antagonist for β_3 adrenoceptors. [¹²⁵I]-cyanopindolol, [¹²⁵I]-hydroxybenzylpindolol and [³H]-alprenolol are high affinity radioligands widely used to label β_1 - and β_2 -adrenoceptors and β_3 -adrenoceptors can be labelled with higher concentrations (nM)

of [¹²⁵I]-cyanopindolol in the presence of appropriate concentrations of β₁- and β₂-adrenoceptor antagonists. [³H]L-748337 is a β₃-selective radioligand. Fluorescent ligands such as BODIPY-TMR-CGP12177 are also increasingly being used to track β-adrenoceptors at the cellular level [86]. Somewhat selective β₁-adrenoceptor selective agonists (**denopamine**, **dobutamine**) are used short-term to

treat cardiogenic shock but, in the longer term, reduce survival. β₁-Adrenoceptor-preferring antagonists are used to treat hypertension (**atenolol**, **betaxolol**, **bisoprolol**, **metoprolol** and **nebivolol**), cardiac arrhythmias (**atenolol**, **bisoprolol**, **esmolol**) and cardiac failure (**metoprolol**, **nebivolol**). Cardiac failure is also successfully treated with **carvedilol** which blocks both β₁- and β₂-adrenoceptors, as well as α₁-adrenoceptors. β₂-Adrenoceptor-selective agonists are

powerful bronchodilators widely used to treat respiratory disorders. There are both short (**salbutamol**, **terbutaline**) and long acting drugs (**formoterol**, **salmeterol**). Although many first generation β-adrenoceptor antagonists (**propranolol**) block both β₁- and β₂-adrenoceptors there are no β₂-adrenoceptor-selective antagonists used therapeutically. The β₃-adrenoceptor agonist **mirabegron** is used to control overactive bladder syndrome.

Nomenclature	β ₁ -adrenoceptor	β ₂ -adrenoceptor	β ₃ -adrenoceptor
HGNC, UniProt	ADRB1 , P08588	ADRB2 , P07550	ADRB3 , P13945
Rank order of potency	(-)-noradrenaline > (-)-adrenaline	(-)-adrenaline > (-)-noradrenaline	(-)-noradrenaline = (-)-adrenaline
Endogenous agonists	noradrenaline (pK _i 6) [549]	–	–
Agonists	pindolol (Partial agonist) (pK _i 9.3) [1011], isoprenaline (pK _i 6.6–7) [549, 1651], dobutamine (Partial agonist) (pK _i 5.5) [831]	pindolol (Partial agonist) (pK _i 9.4) [1011], arformoterol (pK _i 8.6) [37], isoprenaline (pK _i 6.4) [1651], dobutamine (Partial agonist) (pK _i 6.2) [1100], ephedrine (Partial agonist) (pK _i 5.6) [851]	carazolol (pK _i 8.7) [1256]
Selective agonists	(-)-Ro 363 (pK _i 8) [1301], xamoterol (Partial agonist) (pK _i 7) [831], denopamine (Partial agonist) (pK _i 5.8) [831, 1830]	formoterol (pEC ₅₀ 10.1) [81], salmeterol (pEC ₅₀ 9.9) [81], zinterol (pEC ₅₀ 9.5) [81], vilanterol (pEC ₅₀ 9.4) [1534], procaterol (pEC ₅₀ 8.4) [81], indacaterol (pK _i 7.8) [107], fenoterol (pK _i 6.9) [55], salbutamol (Partial agonist) (pK _i 5.8–6.1) [83, 831], terbutaline (Partial agonist) (pK _i 5.6) [83], orciprenaline (pK _d 5.3) [1784]	L 755507 (pEC ₅₀ 10.1) [81], L742791 (pEC ₅₀ 8.8) [2000], mirabegron (pEC ₅₀ 7.7) [1849], CGP 12177 (Partial agonist) (pK _i 6.1–7.3) [159, 1144, 1256, 1301], SB251023 (pEC ₅₀ 7.1) [810] – Mouse, BRL 37344 (pK _i 6.4–7) [159, 431, 768, 1256], CL316243 (pK _i 5.2) [2080]
Antagonists	carvedilol (pK _i 9.5) [262], bupranolol (pK _i 7.3–9) [262, 1144], levobunolol (pK _i 8.4) [68], labetalol (pK _i 8.2) [68], metoprolol (pK _i 7–7.6) [83, 262, 768, 1144], esmolol (pK _i 6.9) [68], nadolol (pK _i 6.9) [262], practolol (pK _i 6.1–6.8) [83, 1144], propafenone (pK _i 6.7) [68], sotalol (pK _i 6.1) [68]	carvedilol (pK _i 9.4–9.9) [83, 262], timolol (pK _i 9.7) [83], propranolol (pK _i 9.1–9.5) [83, 86, 831, 1144], levobunolol (pK _i 9.3) [68], bupranolol (pK _i 8.3–9.1) [262, 1144], alprenolol (pK _i 9) [83], nadolol (pK _i 7–8.6) [83, 262], labetalol (pK _i 8) [68], propafenone (pK _i 7.4) [68], sotalol (pK _i 6.5) [68]	carvedilol (pK _i 9.4) [262], bupranolol (pK _i 6.8–7.3) [159, 262, 1144, 1256], propranolol (pK _i 6.3–7.2) [1144, 1517], levobunolol (pK _i 6.8) [1517]
Selective antagonists	CGP 20712A (pK _i 8.5–9.2) [83, 262, 1144], levobetaxolol (pK _i 9.1) [1715], betaxolol (pK _i 8.8) [1144], nebivolol (pK _i 8.1–8.7) [1476] – Rabbit, atenolol (pK _i 6.7–7.6) [83, 886, 1144], acebutolol (pK _i 6.4) [68]	ICI 118551 (Inverse agonist) (pK _i 9.2–9.5) [83, 86, 1144]	L-748337 (pK _i 8.4) [262], SR59230A (pK _i 6.9–8.4) [262, 405, 768], L748328 (pK _i 8.4) [262]
Labelled ligands	[¹²⁵ I]CYP (Selective Antagonist) (pK _d 10.4–11.3) [831, 1144, 1651]	[¹²⁵ I]CYP (Antagonist) (pK _d 11.1) [1144, 1651]	[¹²⁵ I]CYP (Agonist, Partial agonist) (pK _d 9.2–9.8) [1144, 1301, 1517, 1651, 1806]
Comments	The agonists indicated have less than two orders of magnitude selectivity [81].	–	Agonist SB251023 has a pEC ₅₀ of 6.9 for the splice variant of the mouse β ₃ receptor, β _{3b} [810].

Comments:

Adrenoceptors, α₁

The clone originally called the α_{1C}-adrenoceptor corresponds to the pharmacologically defined α_{1A}-adrenoceptor [752]. Some tis-

sues possess α_{1A}-adrenoceptors (termed α_{1L}-adrenoceptors [532, 1329]) that display relatively low affinity in functional and binding assays for **prazosin** (pK_i < 9) indicative of different receptor states or locations. α_{1A}-adrenoceptor C-terminal splice variants form homo- and heterodimers, but fail to generate a functional α_{1L}-

adrenoceptor [1557]. α_{1D}-Adrenoceptors form heterodimers with α_{1B}- or β₂-adrenoceptors that show increased cell-surface expression [1917]. Recombinant α_{1D}-adrenoceptors have been shown in some heterologous systems to be mainly located intracellularly but cell-surface localization is attained by truncation of the N-terminus,

or by co-expression of $\alpha_1\text{B}$ - or β_2 -adrenoceptors to form heterodimers [670, 1917]. In smooth muscle of native blood vessels all three α_1 -adrenoceptor subtypes are located on the surface and intracellularly [1260, 1261].

Signalling is predominantly via $G_{q/11}$ but α_1 -adrenoceptors also couple to $G_{i/o}$, G_s and $G_{12/13}$. Several ligands activating $\alpha_1\text{A}$ -adrenoceptors display ligand directed signalling bias relative to noradrenaline. For example, **oxymetazoline** is a full agonist for extracellular acidification rate (ECAR) and a partial agonist for Ca^{2+} release but does not stimulate cAMP production. **Phenylephrine** is biased toward ECAR versus Ca^{2+} release or cAMP accumulation but not between Ca^{2+} release and cAMP accumulation [495]. There are also differences between subtypes in coupling efficiency to different pathways *e.g.* in some systems coupling efficiency to Ca^{2+} signalling is $\alpha_1\text{A} > \alpha_1\text{B} > \alpha_1\text{D}$, but for MAP kinase signalling is $\alpha_1\text{D} > \alpha_1\text{A} > \alpha_1\text{B}$. In vascular smooth muscle, the potency of agonists is related to the predominant subtype, $\alpha_1\text{D}$ -conveying greater agonist sensitivity than $\alpha_1\text{A}$ -adrenoceptors [526].

Adrenoceptors, α_2

ARC-239 (pK_i 8.0) and **prazosin** (pK_i 7.5) show selectivity for $\alpha_2\text{B}$ - and $\alpha_2\text{C}$ -adrenoceptors over $\alpha_2\text{A}$ -adrenoceptors. **Oxymetazoline** is a reduced efficacy agonist and is one of many α_2 -adrenoceptor agonists that are imidazolines or closely related compounds. Other binding sites for imidazolines, distinct from α_2 -adrenoceptors, and structurally distinct from the 7TM adrenoceptors, have been identified and classified as I_1 , I_2 and I_3 sites [390]; catecholamines have a low affinity, while rilmenidine and moxonidine are selective ligands for these sites, evoking hypotensive effects *in vivo*. I_1 -imidazoline receptors are involved in central inhibition of sympathetic tone, I_2 -imidazoline receptors are an allosteric binding site on monoamine oxidase B, and I_3 -imidazoline receptors regulate insulin secretion from pancreatic β -cells. $\alpha_2\text{A}$ -adrenoceptor stimulation reduces insulin

secretion from β -islets [2083], with a polymorphism in the 5'-UTR of the ADRA2A gene being associated with increased receptor expression in β -islets and heightened susceptibility to diabetes [1599]. $\alpha_2\text{A}$ - and $\alpha_2\text{C}$ -adrenoceptors form homodimers [1758]. Heterodimers between $\alpha_2\text{A}$ - and either the $\alpha_2\text{C}$ -adrenoceptor or μ opioid peptide receptor exhibit altered signalling and trafficking properties compared to the individual receptors [1758, 1858, 1956]. Signalling by α_2 -adrenoceptors is primarily via $G_{i/o}$, however the $\alpha_2\text{A}$ -adrenoceptor also couples to G_s [459]. Imidazoline compounds display bias relative to each other at the $\alpha_2\text{A}$ -adrenoceptor when assayed by [^{35}S] GTP γS binding compared to inhibition of cAMP accumulation [1477]. The noradrenaline reuptake inhibitor desipramine acts directly on the $\alpha_2\text{A}$ -adrenoceptor, promoting internalisation via recruitment of arrestin without activating G proteins [371].

Adrenoceptors, β

Radioligand binding with [^{125}I]ICYP can be used to define β_1 - or β_2 -adrenoceptors when conducted in the presence of a 'saturating' concentration of either a β_1 - or β_2 -adrenoceptor-selective antagonist. [^3H]CGP12177 or [^3H]dihydroalprenolol can be used in place of [^{125}I]ICYP. Binding of a fluorescent analogue of CGP 12177 to β_2 -adrenoceptors in living cells has been described [84]. [^{125}I]ICYP at higher (nM) concentrations can be used to label β_3 -adrenoceptors in systems where there are few if any other β -adrenoceptor subtypes. Pharmacological differences exist between human and mouse β_3 -adrenoceptors, and the 'rodent selective' agonists **BRL 37344** and **CL316243** are partial agonists at the human β_3 -adrenoceptor whereas **CGP 12177** and **L 755507** activate human β_3 -adrenoceptors with greater potency [1650]. The β_3 -adrenoceptor has an intron in the coding region, but splice variants have only been described for the mouse [496], where the isoforms display different signalling characteristics [810]. There are 3 β -adrenoceptors in turkey (termed the $\text{t}\beta$, $\text{t}\beta_3\text{c}$ and $\text{t}\beta_4\text{c}$) that have a pharmacology that differs from the human β -adrenoceptors

[82]. Numerous polymorphisms have been described for the three β -adrenoceptors; some are associated with alterations in agonist-evoked signalling and trafficking, altered susceptibility to disease and/or altered responses to pharmacotherapy [1103].

All β -adrenoceptors couple to G_s (activating adenylyl cyclase and elevating cAMP levels), but it is also clear that they activate other G proteins such as G_i and many other G protein-independent signalling pathways, including arrestin-mediated signalling, which may in turn lead to activation of MAP kinases. Many antagonists at β_1 - and β_2 -adrenoceptors are agonists at β_3 -adrenoceptors (**CL316243**, **CGP 12177** and **carazolol**). Many 'antagonists' that block agonist-stimulated cAMP accumulation, for example **carvedilol** and **bucindolol**, are able to activate MAP kinase pathways [85, 497, 559, 560, 1649, 1650] and thus display 'protean agonism'. **Bupranolol** appears to act as a neutral antagonist in most systems so far examined. Agonists also display biased signalling at the β_2 -adrenoceptor via G_s or arrestins [443].

The X-ray crystal structures have been described of the agonist bound [1988] and antagonist bound forms of the β_1 - [1989], agonist-bound [313] and antagonist-bound forms of the β_2 -adrenoceptor [1561, 1598], as well as a fully active agonist-bound, G_s protein-coupled β_2 -adrenoceptor [1562]. **Carvedilol** and **bucindolol** bind to an extended site of the β_1 -adrenoceptor involving contacts in TM2, 3, and 7 and extracellular loop 2 that may facilitate coupling to arrestins [1989]. Compounds displaying arrestin-biased signalling at the β_2 -adrenoceptor also have a greater effect on the conformation of TM7, whereas full agonists for G_s coupling promote movement of TM5 and TM6 [1127]. Recent studies using NMR spectroscopy have demonstrated significant conformational flexibility in the β_2 -adrenoceptor which is stabilized by both agonist and G proteins highlighting the dynamic nature of interactions with both ligand and downstream signalling partners [946, 1199, 1413]. Such flexibility will likely have consequences for our understanding of biased agonism, and for the future therapeutic exploitation of this phenomenon.

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Angiotensin receptors

G protein-coupled receptors → Angiotensin receptors

Overview: The actions of angiotensin II (AGT, P01019) (Ang II) are mediated by AT₁ and AT₂ receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Angiotensin Receptors [2137]), which have around 30% sequence similarity. Endogenous ligands are angiotensin II (AGT, P01019) and angiotensin III (AGT, P01019) (Ang III), while angiotensin I (AGT, P01019) is weakly active in some systems.

Nomenclature	AT ₁ receptor	AT ₂ receptor
HGNC, UniProt	AGTR1, P30556	AGTR2, P50052
Selective agonists	L-162,313 (pIC ₅₀ 7.8–7.9) [1490]	CGP42112 (pIC ₅₀ 9.6) [190], [p-aminoPhe6]ang II (pK _d 9.1–9.4) [1789, 2139] – Rat
Antagonists	telmisartan (pIC ₅₀ 8.4) [1241], olmesartan (pIC ₅₀ 8.1) [981]	–
Selective antagonists	candesartan (pIC ₅₀ 9.5–9.7) [1942], EXP3174 (pIC ₅₀ 7.4–9.5) [1887, 1942], eprosartan (pIC ₅₀ 8.4–8.8) [464], irbesartan (pIC ₅₀ 8.7–8.8) [1942], losartan (pIC ₅₀ 7.4–8.7) [1887, 2139], valsartan (pIC ₅₀ 8.6) [2138], azilsartan (pIC ₅₀ 8.1–8.1) [1551, 1844]	PD123177 (pIC ₅₀ 8.5–9.5) [291, 321, 450] – Rat, EMA401 (pIC ₅₀ 8.5–9.3) [518, 1582, 1767], PD123319 (pK _d 8.7–9.2) [449, 2025, 2139]
Labelled ligands	[³ H]A81988 (Antagonist) (pK _d 9.2) [690] – Rat, [³ H]L158809 (Antagonist) (pK _d 9.2) [305] – Rat, [³ H]eprosartan (Antagonist) (pK _d 9.1) [21] – Rat, [³ H]valsartan (Antagonist) (pIC ₅₀ 8.8–9) [1954], [¹²⁵ I]EXP985 (Antagonist) (pK _d 8.8) [322] – Rat, [³ H]losartan (Antagonist) (pK _d 8.2) [294] – Rat	[¹²⁵ I]CGP42112 (Agonist) (pK _d 10.6) [2017, 2018, 2139]

Comments: AT₁ receptors are predominantly coupled to Gq₁₁, however they are also linked to arrestin recruitment and stimulate G protein-independent arrestin signalling [1156]. Most species express a single *AGTR1* gene, but two related *agtr1a* and *agtr1b* receptor genes are expressed in rodents. The AT₂ receptor counteracts several of the growth responses initiated by the AT₁ receptors. The AT₂ receptor is much less abundant than the AT₁ receptor in adult tissues

and is upregulated in pathological conditions. AT₁ receptor antagonists bearing substituted 4-phenylquinoline moieties have been synthesized, which bind to AT₁ receptors with nanomolar affinity and are slightly more potent than losartan in functional studies [264]. The antagonist activity of CGP42112 at the AT₂ receptor has also been reported [2147].

The AT₁ and bradykinin B2 receptors have been proposed to form a heterodimeric complex [3].

There is also evidence for an AT₄ receptor that specifically binds angiotensin IV (AGT, P01019) and is located in the brain and kidney. An additional putative endogenous ligand for the AT₄ receptor has been described (LVV-hemorphin (HBB, P68871), a globin decapeptide) [1296].

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Apelin receptor

G protein-coupled receptors → Apelin receptor

Overview: The apelin receptor (nomenclature as agreed by the **NC-IUPHAR Subcommittee on the apelin receptor [1510]**) responds to apelin, a 36 amino-acid peptide derived initially from bovine stomach. Apelin-36 (*APLN*, Q9ULZ1), apelin-13 (*APLN*, Q9ULZ1) and [Pyr¹]apelin-13 (*APLN*, Q9ULZ1) are the predominant endogenous ligands which are cleaved from a 77 amino-acid precursor peptide (*APLN*, Q9ULZ1) by a so far unidentified enzymatic pathway [1864]. A second family of peptides discovered independently and named Elabela [323] or Toddler, that has little sequence similarity to apelin, has been proposed as a second endogenous apelin receptor ligand [1475].

Nomenclature	apelin receptor
HGNC, UniProt	APLNR , P35414
Rank order of potency	[Pyr¹]apelin-13 (APLN, Q9ULZ1) ≥ apelin-13 (APLN, Q9ULZ1) > apelin-36 (APLN, Q9ULZ1) [503, 1864]
Endogenous agonists	apelin-13 (APLN, Q9ULZ1) (Selective) (pIC ₅₀ 8.8–9.5) [503, 785, 1254], apelin receptor early endogenous ligand (APELA, PODMC3) (Selective) (pK _d 9.3) [412], apelin-17 (APLN, Q9ULZ1) (Selective) (pIC ₅₀ 7.9–9) [468, 1254], [Pyr¹]apelin-13 (APLN, Q9ULZ1) (Selective) (pIC ₅₀ 7–8.8) [918, 1254], Elabela/Toddler-21 (APELA, PODMC3) (pIC ₅₀ 8.7) [2086], Elabela/Toddler-32 (APELA, PODMC3) (pIC ₅₀ 8.7) [2086], apelin-36 (APLN, Q9ULZ1) (Selective) (pIC ₅₀ 8.2–8.6) [503, 785, 918, 1254], Elabela/Toddler-11 (APELA, PODMC3) (pIC ₅₀ 7.2) [2086]
Selective agonists	MM07 (Biased agonist) (pEC ₅₀ 9.5) [203]
Antagonists	MM54 (pK _i 8.2) [1166]
Labelled ligands	[¹²⁵I][Nle⁷⁵,Tyr⁷⁷]apelin-36 (human) (Agonist) (pK _d 11.2) [918], [¹²⁵I][Glp⁶⁵Nle⁷⁵,Tyr⁷⁷]apelin-13 (Agonist) (pK _d 10.7) [785], [¹²⁵I](Pyr¹)apelin-13 (Agonist) (pK _d 9.5) [911], [¹²⁵I]apelin-13 (Agonist) (pK _d 9.2) [503], [³H](Pyr¹)[Met(0)11]-apelin-13 (Agonist) (pK _d 8.6) [1254]

Comments: Potency order determined for heterologously expressed human apelin receptor (pD₂ values range from 9.5 to 8.6). The apelin receptor may also act as a co-receptor with CD4 for isolates of human immunodeficiency virus, with apelin blocking this function [279]. A modified apelin-13 peptide, [apelin-13\(F13A\)](#) was reported to block the hypotensive response to apelin in rat *in vivo* [1067], however, this peptide exhibits agonist activity in HEK293 cells stably expressing the recombinant apelin receptor [503].

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Bile acid receptor

G protein-coupled receptors → Bile acid receptor

Overview: The bile acid receptor (GPBA) responds to bile acids produced during the liver metabolism of [cholesterol](#). Selective agonists are promising drugs for the treatment of metabolic disorders, such as type II diabetes, obesity and atherosclerosis.

Nomenclature	GPBA receptor
HGNC, UniProt	GPBAR1 , Q8TDU6
Rank order of potency	lithocholic acid > deoxycholic acid > chenodeoxycholic acid , cholic acid (Unknown) [917 , 1214]
Selective agonists	betulinic acid (pEC ₅₀ 6) [590], oleanolic acid (pEC ₅₀ 5.7) [1648]

Comments: The triterpenoid natural product [betulinic acid](#) has also been reported to inhibit inflammatory signalling through the NFκB pathway [[1842](#)]. Disruption of GPBA expression is reported to protect from cholesterol gallstone formation [[1951](#)]. A new series of 5-phenoxy-1,3-dimethyl-1H-pyrazole-4-carboxamides have been reported as highly potent agonists [[1138](#)].

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Bombesin receptors

G protein-coupled receptors → Bombesin receptors

Overview: Bombesin receptors (**nomenclature recommended by the NC-IUPHAR Subcommittee on bombesin receptors, [857]**) are activated by the endogenous ligands [gastrin-releasing peptide \(GRP, P07492\)](#) (GRP), [neuromedin B \(NMB, P08949\)](#) (NMB) and [GRP-\(18-27\) \(GRP, P07492\)](#) (previously named neuromedin C). **Bombesin** is a tetradecapeptide, originally derived from amphibians, and is an agonist at BB₁ and BB₂ receptors. These

receptors couple primarily to the G_{q/11} family of G proteins (but see also [857]). Each of these receptors is widely distributed in the CNS and peripheral tissues [625, 857, 1556, 1642]. Activation of BB₁ and BB₂ receptors causes a wide range of physiological actions, including the stimulation of normal and neoplastic tissue growth, smooth-muscle contraction, appetite and feeding behavior, secretion and many central nervous system effects [857, 858, 859, 1185,

1317, 1556]. A physiological role for the BB₃ receptor has yet to be fully defined although recently studies using receptor knockout mice and newly described agonists/antagonists suggest an important role in glucose and insulin regulation, metabolic homeostasis, feeding and other CNS behaviors and growth of normal/neoplastic tissues [625, 1186, 1430].

	BB ₁ receptor	BB ₂ receptor	BB ₃ receptor
Nomenclature	BB ₁ receptor	BB ₂ receptor	BB ₃ receptor
HGNC, UniProt	NMBR, P28336	GRPR, P30550	BR53, P32247
Endogenous agonists	neuromedin B (NMB, P08949) (Selective) (pK _i 8.1–10.3) [857, 1556, 1919]	neuromedin C (pIC ₅₀ 9.9) [1919], gastrin releasing peptide(14-27) (human) (Selective) (pIC ₅₀ 9.7–9.8) [1919]	–
Selective agonists	–	–	compound 8a [PMID: 24900283] (pIC ₅₀ 8.9) [1129], compound 9g [PMID: 24412111] (pEC ₅₀ 8.8) [1220], MK-7725 (pIC ₅₀ 8.5) [324], MK-5046 (pK _i 7.7–8.4) [1321, 1689], [D-Tyr⁶,Apa-4Cl¹¹,Phe¹³,Nle¹⁴]bombesin-(6-14) (pK _i 8.1) [1202], compound 17c [PMID: 25497965] (pEC ₅₀ 7.9) [1219], compound 9f [PMID: 24412111] (pEC ₅₀ 7.8) [1220], bag-1 (pIC ₅₀ 7.7) [659], compound 22e [PMID: 20167483] (pIC ₅₀ 7.6) [727], bag-2 (pIC ₅₀ 7) [659]
Antagonists	D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Nal-NH₂ (pIC ₅₀ 6.2–6.6) [624]	–	–

(continued)			
Nomenclature	BB₁ receptor	BB₂ receptor	BB₃ receptor
Selective antagonists	PD 176252 (pIC ₅₀ 9.3–9.8) [624], PD 168368 (pIC ₅₀ 9.3–9.6) [624], dNal-cyc(Cys-Tyr-dTrp-Orn-Val)-Nal-NH ₂	[D-Phe ⁶ , Leu ¹³ , Cpa ¹⁴ , ψ ¹³⁻¹⁴]bombesin-(6-14) (pK _i 9.8) [624], JMV641 (pIC ₅₀ 9.3) [1892] – Mouse, [(3-Ph-Pr ⁶), His ⁷ , D-Ala ¹¹ , D-Pro ¹³ , ψ ¹³⁻¹⁴ , Phe ¹⁴] bombesin-(6-14) (pIC ₅₀ 9.2) [624, 1062], [D-Tpi ⁶ , Leu ¹³ ψ(CH ₂ NH)-Leu ¹⁴]bombesin-(6-14) (pIC ₅₀ 8.9) [624], Ac-GRP-(20-26)-methylester (pIC ₅₀ 8.7) [624], JMV594 (pIC ₅₀ 8.7–8.7) [1133, 1892] – Mouse	bantag-1 (pIC ₅₀ 8.6–8.7) [659, 1321], ML-18 (pIC ₅₀ 5.3) [1316]
Labelled ligands	[¹²⁵ I]BH-NMB (human, mouse, rat) (Agonist), [¹²⁵ I][Tyr ⁴]bombesin (Agonist)	[¹²⁵ I][D-Tyr ⁶]bombesin-(6-13)-methyl ester (Selective Antagonist) (pK _d 9.3) [1201] – Mouse, [¹²⁵ I][Tyr ⁴]bombesin (Agonist) (pK _d 8.2) [131], [¹²⁵ I]GRP (human) (Agonist)	[³ H]bag-2 (Agonist) (pK _d 8.6) [659] – Mouse, [¹²⁵ I][D-Tyr ⁶ , β-Ala ¹¹ , Phe ¹³ , Nle ¹⁴]bombesin-(6-14) (Agonist) (pK _d 8–8.4) [1203, 1321]

Comments: All three subtypes may be activated by [D-Phe⁶, β-Ala¹¹, Phe¹³, Nle¹⁴]bombesin-(6-14) [1203]. [D-Tyr⁶, Apa-4Cl¹¹, Phe¹³, Nle¹⁴]bombesin-(6-14) has more than 200-fold selectivity for BB₃ receptors over BB₁ and BB₂ [1202].

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Bradykinin receptors

G protein-coupled receptors → Bradykinin receptors

Overview: Bradykinin (or kinin) receptors (**nomenclature as agreed by the NC-IUPHAR subcommittee on Bradykinin (kinin) Receptors [1072]**) are activated by the endogenous peptides bradykinin (*KNG1*, P01042) (BK), [des-Arg⁹]bradykinin (*KNG1*, P01042), Lys-BK (kallidin (*KNG1*, P01042)), [des-Arg¹⁰]kallidin (*KNG1*, P01042), T-kinin (*KNG1*, P01042) (Ile-Ser-BK), [Hyp³]bradykinin (*KNG1*, P01042) and Lys-[Hyp³]-bradykinin (*KNG1*, P01042). The variation in affinity or inactivity of B₂ receptor antagonists could reflect the existence of species homologues of B₂ receptors.

Nomenclature	B ₁ receptor	B ₂ receptor
HGNC, UniProt	<i>BDKRB1</i> , P46663	<i>BDKRB2</i> , P30411
Rank order of potency	[des-Arg ¹⁰]kallidin (<i>KNG1</i> , P01042) > [des-Arg ⁹]bradykinin (<i>KNG1</i> , P01042) = kallidin (<i>KNG1</i> , P01042) > bradykinin (<i>KNG1</i> , P01042)	kallidin (<i>KNG1</i> , P01042) > bradykinin (<i>KNG1</i> , P01042) ≫ [des-Arg ⁹]bradykinin (<i>KNG1</i> , P01042), [des-Arg ¹⁰]kallidin (<i>KNG1</i> , P01042)
Endogenous agonists	[des-Arg ¹⁰]kallidin (<i>KNG1</i> , P01042) (Selective) (pK _i 9.6–10) [69, 104, 876]	–
Selective agonists	[Sar,D-Phe ⁸ ,des-Arg ⁹]bradykinin (pK _i 5.7) [876]	[Hyp ³ ,Tyr(Me) ⁸]BK, [Phe ⁸ ,ψ(CH ₂ -NH)Arg ⁹]BK
Antagonists	[Leu ⁹ ,des-Arg ¹⁰]kallidin (pK _i 9.1–9.3) [69, 104]	–
Selective antagonists	B-9958 (pK _i 9.2–10.3) [596, 1570], R-914 (pA ₂ 8.6) [617], R-715 (pA ₂ 8.5) [618]	icatibant (pK _i 10.2) [39], FR173657 (pA ₂ 8.2) [1593], anantibant (pK _i 8.2) [1537]
Labelled ligands	[¹²⁵ I]Hpp-desArg ¹⁰ HOE140 (pK _d 10), [³ H]Lys-[des-Arg ⁹]BK (Agonist) (pK _d 9.4), [³ H]Lys-[Leu ⁸][des-Arg ⁹]BK (Antagonist)	[³ H]BK (human, mouse, rat) (Agonist) (pK _d 9.4) [2034] – Mouse, [³ H]NPC17731 (Antagonist) (pK _d 9.1–9.4) [2119, 2120], [¹²⁵ I][Tyr ⁸]bradykinin (Agonist)

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Calcitonin receptors

G protein-coupled receptors → Calcitonin receptors

Overview: This receptor family comprises a group of receptors for the calcitonin/CGRP family of peptides. The calcitonin (CT), amylin (AMY), calcitonin gene-related peptide (CGRP) and adrenomedullin (AM) receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on CGRP, AM, AMY, and CT receptors [721, 1528]**) are generated by the genes *CALCR* (which codes for the CT receptor (CTR)) and *CALCL* (which codes for the calcitonin receptor-like receptor, CLR, previously known as CRLR). Their function and pharmacology are altered in the presence of RAMPs (receptor activity-modifying proteins), which are single TM domain pro-

teins of ca. 130 amino acids, identified as a family of three members; RAMP1, RAMP2 and RAMP3. There are splice variants of CTR; these in turn produce variants of the AMY receptor [1528], some of which can be potentially activated by CGRP. The endogenous agonists are the peptides calcitonin (*CALCA*, P01258), α -CGRP (*CALCA*, P06881) (formerly known as CGRP-I), β -CGRP (*CALCB*, P10092) (formerly known as CGRP-II), amylin (*IAPP*, P10997) (occasionally called islet-amyloid polypeptide, diabetes-associated polypeptide), adrenomedullin (*ADM*, P35318) and adrenomedullin 2/intermedin (*ADM2*, Q7Z4H4). There are species differences in peptide se-

quences, particularly for the CTs. **CTR-stimulating peptide** {Pig} (CRSP) is another member of the family with selectivity for the CTR but it is not expressed in humans [907]. **Olcegepant** (also known as BIBN4096BS, pKi 10.5) and **telcagepant** (also known as MK0974, pKi 9) are the most selective antagonists available, having a high selectivity for CGRP receptors, with a particular preference for those of primate origin.

CLR by itself binds no known endogenous ligand, but in the presence of RAMPs it gives receptors for CGRP, adrenomedullin and adrenomedullin 2/intermedin.

	CT receptor	AMY ₁ receptor	AMY ₂ receptor	AMY ₃ receptor
Nomenclature	CT receptor	AMY ₁ receptor	AMY ₂ receptor	AMY ₃ receptor
HGNC, UniProt	CALCR , P30988	–	–	–
Subunits	–	RAMP1 (Accessory protein), CT receptor	CT receptor, RAMP2 (Accessory protein)	CT receptor, RAMP3 (Accessory protein)
Rank order of potency	calcitonin (salmon) ≥ calcitonin (<i>CALCA</i> , P01258) ≥ amylin (<i>IAPP</i> , P10997), α -CGRP (<i>CALCA</i> , P06881) > adrenomedullin (<i>ADM</i> , P35318), adrenomedullin 2/intermedin (<i>ADM2</i> , Q7Z4H4)	calcitonin (salmon) ≥ amylin (<i>IAPP</i> , P10997) ≥ α -CGRP (<i>CALCA</i> , P06881) > adrenomedullin 2/intermedin (<i>ADM2</i> , Q7Z4H4) ≥ calcitonin (<i>CALCA</i> , P01258) > adrenomedullin (<i>ADM</i> , P35318)	Poorly defined	calcitonin (salmon) ≥ amylin (<i>IAPP</i> , P10997) > α -CGRP (<i>CALCA</i> , P06881) ≥ adrenomedullin 2/intermedin (<i>ADM2</i> , Q7Z4H4) ≥ calcitonin (<i>CALCA</i> , P01258) > adrenomedullin (<i>ADM</i> , P35318)
Endogenous agonists	calcitonin (<i>CALCA</i> , P01258) (Selective) (pEC ₅₀ 9–11.2) [32, 58, 718, 1027, 1087, 1341]	amylin (<i>IAPP</i> , P10997) (pEC ₅₀ 9–9.7) [610]	amylin (<i>IAPP</i> , P10997) (pEC ₅₀ 8.3–9.1) [610]	amylin (<i>IAPP</i> , P10997) (pEC ₅₀ 8.9–9.6) [610]
Labelled ligands	[¹²⁵ I]CT (human) (Agonist) (pK _d 9–10), [¹²⁵ I]CT (salmon) (Agonist) (pK _d 10)	[¹²⁵ I]BH-AMY (rat, mouse) (Agonist) (pK _d 9–10)	[¹²⁵ I]BH-AMY (rat, mouse) (Agonist) (pK _d 9–10)	[¹²⁵ I]BH-AMY (rat, mouse) (Agonist) (pK _d 9–10)

	calcitonin receptor-like receptor	CGRP receptor	AM ₁ receptor	AM ₂ receptor
Nomenclature	CALCRL , Q16602	–	–	–
HGNC, UniProt	–	–	–	–
Subunits	–	calcitonin receptor-like receptor, RAMP1 (Accessory protein)	calcitonin receptor-like receptor, RAMP2 (Accessory protein)	calcitonin receptor-like receptor, RAMP3 (Accessory protein)
Rank order of potency	–	α -CGRP (CALCA , P06881) > adrenomedullin (ADM , P35318) \geq adrenomedullin 2/intermedin (ADM2 , Q7Z4H4) > amylin (IAPP , P10997) \geq calcitonin (salmon)	adrenomedullin (ADM , P35318) > adrenomedullin 2/intermedin (ADM2 , Q7Z4H4) > α -CGRP (CALCA , P06881), amylin (IAPP , P10997) > calcitonin (salmon)	adrenomedullin (ADM , P35318) \geq adrenomedullin 2/intermedin (ADM2 , Q7Z4H4) \geq α -CGRP (CALCA , P06881) > amylin (IAPP , P10997) > calcitonin (salmon)
Endogenous agonists	–	β -CGRP (CALCB , P10092) (pK _i 9.9–11) [20 , 1251], α -CGRP (CALCA , P06881) (pK _i 9.7–10) [20 , 1251]	adrenomedullin (ADM , P35318) (pK _i 8.3–9.2) [20 , 1251]	adrenomedullin (ADM , P35318) (pK _i 8.3–9) [20 , 539]
Antagonists	–	olcegepant (pK _i 10.2–10.7) [435 , 719 , 720 , 1194], telcagepant (pK _i 9.1) [1633]	–	–
Selective antagonists	–	–	AM-(22-52) (human) (pK _i 7–7.8) [20 , 720 , 1251]	–
Labelled ligands	–	[¹²⁵ I] α CGRP (human) (Agonist) (pK _d 10), [¹²⁵ I] α CGRP (mouse, rat) (Agonist)	[¹²⁵ I]AM (rat) (Agonist) (pK _d 10–9)	[¹²⁵ I]AM (rat) (Agonist) (pK _d 9–10)

Comments: It is important to note that a complication with the interpretation of pharmacological studies with AMY receptors in transfected cells is that most of this work has likely used a mixed population of receptors, encompassing RAMP-coupled CTR as well as CTR alone. This means that although in binding assays human calcitonin ([CALCA](#), [P01258](#)) has low affinity for [¹²⁵I]-AMY binding sites, cells transfected with CTR and RAMPs can display potent CT functional responses. Transfection of human CTR with any RAMP can generate receptors with a high affinity for both salmon CT and AMY and varying affinity for different antagonists [[337](#), [718](#), [719](#)]. The major human CTR splice variant (hCT_(a), which does not contain an insert) with RAMP1 (*i.e.* the AMY_{1(a)} receptor) has a high affinity for CGRP, unlike hCT_(a)-RAMP3 (*i.e.* AMY_{3(a)} receptor) [[337](#), [718](#)]. However, the AMY receptor phenotype is RAMP-type, splice variant and cell-line-dependent [[1886](#)]. In particular, CGRP is a more potent agonist than amylin ([IAPP](#), [P10997](#)) at increasing cAMP at the delta 47 hCT(a) receptor, when transfected with RAMP1 (to give the corresponding AMY1(a) receptor) in Cos 7 cells [[1543](#)].

The ligands described represent the best available but their selectivity is limited. For example, adrenomedullin has appreciable affinity for CGRP receptors. CGRP can show significant cross-reactivity at AMY receptors and AM₂ receptors. Adrenomedullin 2/intermedin also has high affinity for the AM₂ receptor [[779](#)]. CGRP-(8-37) acts as an antagonist of CGRP (pK_i 8) and inhibits some AM and AMY responses (pK_i 6-7). It is weak at CT receptors. Salmon CT-(8-32) is an antagonist at both AMY and CT receptors. AC187, a salmon CT analogue, is also an antagonist at AMY and CT receptors. Human AM-(22-52) has some selectivity towards AM receptors, but with modest potency (pK_i 7), limiting its use [[720](#)]. AM-(22-52) is slightly more effective at AM₁ than AM₂ receptors but this difference is not sufficient for this peptide to be a useful discriminator of the AM receptor subtypes. Olcegepant shows the greatest selectivity between receptors but still has significant affinity for AMY₁ receptors [[1973](#)].

Ligand responsiveness at CT and AMY receptors can be affected by receptor splice variation and can depend on the pathway being measured. Particularly for AMY receptors, relative potency can vary with

the type and level of RAMP present and can be influenced by other factors such as G proteins [[1324](#), [1886](#)].

G_s is a prominent route for effector coupling for CLR and CTR but other pathways (*e.g.* Ca²⁺, ERK, Akt), and G proteins can be activated [[1972](#)]. There is evidence that CGRP-RCP (a 148 amino-acid hydrophilic protein, [ASL](#) ([P04424](#))) is important for the coupling of CLR to adenylyl cyclase [[498](#)].

[¹²⁵I]-Salmon CT is the most common radioligand for CT receptors but it has high affinity for AMY receptors and is also poorly reversible. [¹²⁵I]-Tyr⁰-CGRP is widely used as a radioligand for CGRP receptors.

Some early literature distinguished between CGRP₁ and CGRP₂ receptors. It is now clear that the complex of [CALCRL](#) and [RAMP1](#) represents the CGRP₁ subtype and is now known simply as the CGRP receptor [[721](#)]. The CGRP₂ receptor is now considered to have arisen from the actions of CGRP at AM₂ and AMY receptors. This term should not be used [[721](#)].

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Calcium-sensing receptors

G protein-coupled receptors → Calcium-sensing receptors

Overview: The calcium-sensing receptor (CaS, **provisional nomenclature as recommended by NC-IUPHAR [530]**) responds to extracellular calcium and magnesium in the millimolar range and to gadolinium and some polycations in the micromolar range [229]. The sensitivity of CaS to primary agonists can be increased by aromatic L-amino acids [362] and also by elevated extracellular pH [1544] or decreased extracellular ionic strength [1545]. This receptor bears no sequence or structural relation to the plant calcium receptor, also called CaS.

Nomenclature	CaS receptor	GPRC6 receptor
HGNC, UniProt	CASR, P41180	GPRC6A, Q5T6X5
Amino-acid rank order of potency	L-phenylalanine, L-tryptophan, L-histidine > L-alanine > L-serine, L-proline, L-glutamic acid > L-aspartic acid (not L-lysine, L-arginine, L-leucine and L-isoleucine) [362]	–
Cation rank order of potency	Gd ³⁺ > Ca ²⁺ > Mg ²⁺ [229]	–
Polyamine rank order of potency	spermine > spermidine > putrescine [1546]	–
Allosteric modulators	AC265347 (Positive) (pEC ₅₀ 7.6–8.1) [1160], NPS 2143 (Negative) (pIC ₅₀ 7.1–7.4) [1377, 2087], cinacalcet (Positive) (pEC ₅₀ 7.3) [1378], calindol (Positive) (pEC ₅₀ 6.5) [1499], calindol (Positive) (pK _d 6–6.5) [930], tecalcet (Positive) (pK _d 6.5) [1379], calhex 231 (Negative) (pIC ₅₀ 6.4) [1500]	–
Comments	2-benzylpyrrolidine derivatives of NPS 2143 are also negative allosteric modulators of the calcium sensing receptor [2087]. etelcalcetide is a novel peptide agonist of the receptor [1975].	GPRC6 is a related G _q -coupled receptor which responds to basic amino acids [2004].

Comments: Positive allosteric modulators of CaS are termed Type II calcimimetics and can suppress parathyroid hormone (PTH (PTH, P01270)) secretion [1379]. Negative allosteric modulators are called calcilytics and can act to increase PTH (PTH, P01270) secretion [1377].

The central role of CaS in the maintenance of extracellular calcium homeostasis is seen most clearly in patients with loss-of-function CaS mutations who develop familial hypocalcaemic hypercalcaemia (heterozygous mutation) or neonatal severe hyperparathyroidism (homozygous mutation) and in CaS null mice [293, 765], which exhibit

similar increases in PTH secretion and blood Ca²⁺ levels. A gain-of-function mutation in the CaS gene is associated with autosomal dominant hypocalcaemia.

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Cannabinoid receptors

G protein-coupled receptors → [Cannabinoid receptors](#)

Overview: Cannabinoid receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Cannabinoid Receptors [1494]**) are activated by endogenous ligands that include N-arachidonylethanolamine (**anandamide**), N-homo- γ -linolenylethanolamine, N-docosatetra-7,10,13,16-enylethanolamine and 2-arachidonoylglycerol. Potency determinations of endogenous agonists at these receptors are complicated by the possibility of differential susceptibility of endogenous ligands to enzymatic conversion [35].

Nomenclature	CB ₁ receptor	CB ₂ receptor
HGNC, UniProt	CNR1, P21554	CNR2, P34972
(Sub)family-selective agonists	HU-210 (pK _i 9.1–10.2) [509, 1733], CP55940 (pK _i 8.3–9.2) [509, 1602, 1733], WIN55212-2 (pK _i 6.9–8.7) [509, 1730, 1733], Δ^9 -tetrahydrocannabinol (Partial agonist) (pK _i 7.3–7.4) [509, 1733]	HU-210 (pK _i 9.3–9.8) [509, 1579, 1733], WIN55212-2 (pK _i 8.4–9.6) [509, 1730, 1733], CP55940 (pK _i 8.6–9.2) [509, 1602, 1733], Δ^9 -tetrahydrocannabinol (Partial agonist) (pK _i 7.1–7.5) [106, 509, 1579, 1733]
Selective agonists	arachidonyl-2-chloroethylamide (pK _i 8.9) [755] – Rat, arachidonylcyclopropylamide (pK _i 8.7) [755] – Rat, O-1812 (pK _i 8.5) [420] – Rat, R-(+)-methanandamide (pK _i 7.7) [931] – Rat	JWH-133 (pK _i 8.5) [804, 1493], L-759,633 (pK _i 7.7–8.2) [576, 1602], AM1241 (pK _i 8.1) [2088], L-759,656 (pK _i 7.7–7.9) [576, 1602], HU-308 (pK _i 7.6) [699]
Selective antagonists	rimonabant (pK _i 7.9–8.7) [508, 509, 1586, 1613, 1733], AM251 (pK _i 8.1) [1038] – Rat, AM281 (pK _i 7.9) [1037] – Rat, LY320135 (pK _i 6.9) [508]	SR144528 (pK _i 8.3–9.2) [1587, 1602], AM-630 (pK _i 7.5) [1602]
Labelled ligands	[³ H]rimonabant (Antagonist) (pK _d 8.9–10) [205, 761, 889, 1498, 1588, 1742, 1873] – Rat	–

Comments: Both CB₁ and CB₂ receptors may be labelled with [³H]CP55940 (0.5 nM; [1733]) and [³H]WIN55212-2 (2–2.4 nM; [1756, 1783]). Anandamide is also an agonist at vanilloid receptors (TRPV1) and PPARs [1418, 2135]. There is evidence for an allosteric site on the CB₁ receptor [1532]. All of the compounds listed as antagonists behave as inverse agonists in some bioassay systems [1494]. Moreover, GPR18, GPR55 and GPR119, although showing little structural similarity to CB₁ and CB₂ receptors, respond to endogenous agents that are structurally similar to the endogenous cannabinoid ligands [1494].

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Chemerin receptor

G protein-coupled receptors → Chemerin receptor

Overview: The chemerin receptor (**nomenclature as recommended by NC-IUPHAR [396]**) is activated by chemerin [1148, 1253, 2108] and the lipid-derived, anti-inflammatory ligand **resolvin E1** (RvE1), which is the result of sequential metabolism of EPA by aspirin-modified cyclooxygenase and lipoxygenase [56, 57]. In addition, two GPCRs for **resolvin D1** (RvD1) have been identified, FPR2/ALX, the lipoxin A₄ receptor, and GPR32, an orphan receptor [1006].

Nomenclature	chemerin receptor
HGNC, UniProt	CMKLR1, Q99788
Rank order of potency	resolvin E1 > chemerin C-terminal peptide > 18R-HEPE > EPA [56]
Selective agonists	resolvin E1
Labelled ligands	[³ H]resolvin E1 (Agonist) (pK _d 8) [56, 57]

Chemokine receptors

G protein-coupled receptors → Chemokine receptors

Overview: Chemokine receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Chemokine Receptors [78, 1346, 1347]**) comprise a large subfamily of 7TM proteins that bind one or more chemokines, a large family of small cytokines typically possessing chemotactic activity for leukocytes. Chemokine re-

ceptors can be divided by function into two main groups: G protein-coupled chemokine receptors, which mediate leukocyte trafficking, and "Atypical chemokine receptors", which may signal through non-G protein-coupled mechanisms and act as chemokine scavengers to downregulate inflammation or shape chemokine gradients [78].

Chemokines in turn can be divided by structure into four subclasses by the number and arrangement of conserved cysteines. CC (also known as β -chemokines; $n=28$), CXC (also known as α -chemokines; $n=17$) and CX₃C ($n=1$) chemokines all have four conserved cysteines, with zero, one and three amino acids separating the first two

cysteines respectively. C chemokines ($n=2$) have only the second and fourth cysteines found in other chemokines. Chemokines can also be classified by function into homeostatic and inflammatory subgroups. Most chemokine receptors are able to bind multiple high-affinity chemokine ligands, but the ligands for a given receptor are almost always restricted to the same structural subclass. Most chemokines bind to more than one receptor subtype. Receptors for inflammatory chemokines are typically highly promiscuous with regard to ligand specificity, and may lack a selective endogenous ligand. G protein-

coupled chemokine receptors are named according to the class of chemokines bound, whereas ACKR is the root acronym for atypical chemokine receptors [79]. Listed are those human agonists with EC_{50} values <50 nM in either Ca^{2+} flux or chemotaxis assays at human recombinant G protein-coupled chemokine receptors expressed in mammalian cell lines. There can be substantial cross-species differences in the sequences of both chemokines and chemokine receptors, and in the pharmacology and biology of chemokine recep-

tors. Endogenous and microbial non-chemokine ligands have also been identified for chemokine receptors. Many chemokine receptors function as HIV co-receptors, but CCR5 is the only one demonstrated to play an essential role in HIV/AIDS pathogenesis. The tables include both standard chemokine receptor names [2101] and the most commonly used aliases. Numerical data quoted are typically pK_i or pIC_{50} values from radioligand binding to heterologously expressed receptors.

	CCR1	CCR2	CCR3
Nomenclature	CCR1	CCR2	CCR3
HGNC, UniProt	CCR1, P32246	CCR2, P41597	CCR3, P51677
Endogenous agonists	CCL3 (CCL3, P10147) (pK_i 7.8–10.2) [328, 357, 747, 2134], CCL23 (CCL23, P55773) (Selective) (pK_i 8.9) [328], CCL5 (CCL5, P13501) (pK_i 6.8–8.2) [357, 747], CCL7 (CCL7, P80098) (pK_i 8.1) [328, 667], CCL15 (CCL15, Q16663) (Selective) (pIC_{50} 7.9) [373], CCL14 (CCL14, Q16627) (pK_i 7.4) [328], CCL13 (CCL13, Q99616), CCL8 (CCL8, P80075)	CCL2 (CCL2, P13500) (pIC_{50} 9.3–10.2) [373, 1159, 1291, 1465, 1920], CCL13 (CCL13, Q99616) (pIC_{50} 8.6–8.7) [1159, 1920], CCL7 (CCL7, P80098) (pIC_{50} 8.4–8.7) [373, 1159, 1920], CCL11 (CCL11, P51671) (Partial agonist) (pIC_{50} 7.1–7.7) [1159, 1465], CCL16 (CCL16, O15467)	CCL13 (CCL13, Q99616) (pIC_{50} 8.7–10.3) [1332, 1920], CCL24 (CCL24, O00175) (Selective) (pIC_{50} 8–9.4) [1332, 1465], CCL5 (CCL5, P13501) (pK_i 8.5–9.3) [391], CCL7 (CCL7, P80098) (pK_i 8.6–9.2) [391], CCL11 (CCL11, P51671) (Selective) (pIC_{50} 8.7–9) [452, 961, 1332, 1625, 1920], CCL26 (CCL26, Q9Y258) (Selective) (pIC_{50} 7.9–8.9) [961, 1332, 1465], CCL15 (CCL15, Q16663) (pIC_{50} 8.6) [373], CCL28 (CCL28, Q9NRJ3), CCL8 (CCL8, P80075)
Agonists	–	–	CCL11 (Mouse) (pK_i 9.5–10) [391]
Endogenous antagonists	CCL4 (CCL4, P13236) (Selective) (pK_i 7.1–7.8) [328, 357]	CCL26 (CCL26, Q9Y258) (Selective) (pIC_{50} 8.5) [1465]	CXCL10 (CXCL10, P02778) (Selective), CXCL11 (CXCL11, O14625) (Selective), CXCL9 (CXCL9, Q07325) (Selective)
Antagonists	–	–	–
Selective antagonists	BX 471 (pK_i 8.2–9) [1098], compound 2b-1 [PMID: 12614873] (pIC_{50} 8.7) [1368], CP-481,715 (pK_d 8) [614], UCB35625 (pIC_{50} 8) [1625]	GSK Compound 34 (pK_i 7.6)	banyu (I) (Inverse agonist) (pK_i 8.5) [1977], SB328437 (pK_i 8.4), BMS compound 87b (pK_i 8.1) [1964]
Antibodies	–	–	–
Labelled ligands	[^{125}I]CCL7 (human) (Agonist) (pK_d 9.2) [127], [^{125}I]CCL3 (human) (Agonist) (pK_d 8–8.8) [127, 623, 1646], [^{125}I]CCL5 (human) (Agonist) (pK_d 8.2) [1646]	[^{125}I]CCL2 (human) (Agonist), [^{125}I]CCL7 (human) (Agonist)	[^{125}I]CCL11 (human) (Antagonist) (pK_d 8.3) [1977], [^{125}I]CCL5 (human) (Agonist), [^{125}I]CCL7 (human) (Agonist)

Nomenclature	CCR4	CCR5
HGNC, UniProt	CCR4, P51679	CCR5, P51681
Endogenous agonists	CCL22 (CCL22, O00626) (Selective) (pIC ₅₀ 9.2) [822], CCL17 (CCL17, Q92583) (Selective) (pIC ₅₀ 8.7) [822]	CCL5 (CCL5, P13501) (pK _i 9.2–9.7) [75, 1364, 1611], CCL4 (CCL4, P13236) (Selective) (pK _i 9.4–9.6) [1364, 1611], CCL8 (CCL8, P80075) (pK _i 9.3) [1611], CCL3 (CCL3, P10147) (pK _i 8–8.9) [1364, 1611, 2134], CCL11 (CCL11, P51671) (pIC ₅₀ 7.7) [157], CCL2 (CCL2, P13500) (pK _i 7.5) [1364], CCL14 (CCL14, Q16627) (pK _i 7.2) [1364], CCL16 (CCL16, O15467)
Agonists	vMIP-III	R5-HIV-1 gp120
Endogenous antagonists	–	CCL7 (CCL7, P80098) (Selective) (pK _i 7.5) [1364]
Antagonists	–	vicriviroc (pK _i 9.1) [1805], ancriviroc (pK _i 7.8–8.7) [1173, 1455, 1805]
Selective antagonists	–	E913 (pIC ₅₀ 8.7) [1174], aplaviroc (pK _i 8.5) [1173], maraviroc (pIC ₅₀ 8.1) [1364], TAK-779 (pK _i 7.5) [1173], MRK-1 [1023] – Rat
Antibodies	mogamulizumab (Inhibition) [51, 1731]	–
Labelled ligands	[¹²⁵I]CCL17 (human) (Agonist), [¹²⁵I]CCL27 (human) (Agonist)	[¹²⁵I]CCL4 (human) (Agonist) (pK _d 9.6) [1364], [¹²⁵I]CCL3 (human) (Agonist), [¹²⁵I]CCL5 (human) (Agonist), [¹²⁵I]CCL8 (human) (Agonist)

Nomenclature	CCR6	CCR7	CCR8	CCR9	CCR10
HGNC, UniProt	CCR6, P51684	CCR7, P32248	CCR8, P51685	CCR9, P51686	CCR10, P46092
Endogenous agonists	CCL20 (CCL20, P78556) (pIC ₅₀ 7.9–8.5) [18, 74, 1526], beta-defensin 4A (DEFB4A DEFB4B, O15263) (Selective) [2081]	CCL21 (CCL21, O00585) (Selective) (pIC ₅₀ 9.3) [2099], CCL19 (CCL19, Q99731) (Selective) (pIC ₅₀ 7.7–9, median 8.6) [1449, 2098, 2099]	CCL1 (CCL1, P22362) (Selective) (pIC ₅₀ 8.5–9.8) [387, 710, 824], CCL8 {Mouse} – Mouse	CCL25 (CCL25, O15444) (Selective)	CCL27 (CCL27, Q9Y4X3) (Selective), CCL28 (CCL28, Q9NRJ3) (Selective)
Agonists	–	–	vMIP-1 (pIC ₅₀ 8.9–9.9) [387, 824]	–	–
Selective antagonists	–	–	vMCC-1 (pIC ₅₀ 9.4) [387]	–	–
Labelled ligands	[¹²⁵I]CCL20 (human) (Agonist) (pK _d ~10) [641]	[¹²⁵I]CCL19 (human) (Agonist), [¹²⁵I]CCL21 (human) (Agonist) [856]	[¹²⁵I]CCL1 (human) (Agonist) (pK _d 8.9–9.7) [824, 1597]	[¹²⁵I]CCL25 (human) (Agonist)	–

	CXCR1	CXCR2	CXCR3
Nomenclature	CXCR1	CXCR2	CXCR3
HGNC, UniProt	CXCR1, P25024	CXCR2, P25025	CXCR3, P49682
Endogenous agonists	CXCL8 (CXCL8, P10145) (pK _i 8.8–9.5) [142, 675, 1068, 2032, 2049], CXCL6 (CXCL6, P80162) (pK _i 7) [2053]	CXCL1 (CXCL1, P09341) (Selective) (pK _i 8.4–9.7) [675, 1068, 2049], CXCL8 (CXCL8, P10145) (pK _i 8.8–9.5) [142, 675, 1068, 2032, 2049], CXCL7 (PPBP, P02775) (Selective) (pIC ₅₀ 6.3–9.3) [16], CXCL3 (CXCL3, P19876) (Selective) (pIC ₅₀ 7.8–9.2) [16], CXCL2 (CXCL2, P19875) (Selective) (pIC ₅₀ 7–9.1) [16], CXCL5 (CXCL5, P42830) (Selective) (pIC ₅₀ 6.9–9) [16], CXCL6 (CXCL6, P80162) (pK _d 7) [2053]	CXCL11 (CXCL11, O14625) (Selective) (pK _i 10.4–10.5) [734], CXCL10 (CXCL10, P02778) (Selective) (pK _i 7.8–9.8) [734, 2006], CXCL9 (CXCL9, Q07325) (Selective) (pK _i 7.3–8.3) [734, 2006]
Agonists	vCXCL1 (pIC ₅₀ 7.4) [1158], HIV-1 matrix protein p17 (pK _d 5.7) [602]	vCXCL1 (pIC ₅₀ 8.2) [1158], HIV-1 matrix protein p17 (pK _d 6.9) [602]	–
Selective agonists	–	–	–
Endogenous antagonists	–	–	CCL11 (CCL11, P51671) (Selective) (pK _i 7.2) [2006], CCL7 (CCL7, P80098) (Selective) (pK _i 6.6) [2006]
Antagonists	–	–	–
Selective antagonists	–	navarixin (pIC ₅₀ 10.3) [78, 456], danirixin (pIC ₅₀ 7.9) [1285], SB 225002 (pIC ₅₀ 7.7) [2016], elubirixin (pIC ₅₀ 7.7) [78], SX-517 (pIC ₅₀ 7.2) [1172]	–
Allosteric modulators	reparixin (Negative) (pIC ₅₀ 9) [142]	reparixin (Negative) (pIC ₅₀ 6.4) [142]	–
Labelled ligands	[¹²⁵ I]CXCL8 (human) (Agonist) (pK _d 8.9–9.6) [675, 1584]	[¹²⁵ I]CXCL8 (human) (Agonist) (pK _d 9–9.4) [675, 1584], [¹²⁵ I]CXCL1 (human) (Agonist), [¹²⁵ I]CXCL5 (human) (Agonist), [¹²⁵ I]CXCL7 (human) (Agonist)	[¹²⁵ I]CXCL10 (human) (Agonist), [¹²⁵ I]CXCL11 (human) (Agonist)
Comments	Reports on the expression of native CXCR1 by mouse leukocytes are not conclusive. There are reports on the existence of mouse <i>Cxcr1</i> and on <i>Cxcr1</i> knockout mice, but the distinct function of the gene and of its knockout phenotype are unclear [118, 351, 1297, 1628, 1794].	–	–

Nomenclature	CXCR4	CXCR5	CXCR6
HGNC, UniProt	CXCR4 , P61073	CXCR5 , P32302	CXCR6 , O00574
Endogenous agonists	CXCL12α (CXCL12 , P48061) (Selective) (p <i>K</i> _d 7.7–8.2) [746, 1136], CXCL12β (CXCL12 , P48061) (Selective) (p <i>K</i> _d 7.9) [746]	CXCL13 (CXCL13 , O43927) (Selective) (p <i>K</i> _d 7.3) [97]	CXCL16 (CXCL16 , Q9H2A7) (Selective) (p <i>K</i> _d 9) [2026]
Agonists	–	–	–
Selective agonists	ALX40-4C (Partial agonist) (p <i>C</i> ₅₀ 6.1) [2121], X4-HIV-1 gp120	–	–
Endogenous antagonists	–	–	–
Antagonists	plerixafor (p <i>K</i> _i 7) [2121]	–	–
Selective antagonists	T134 (p <i>C</i> ₅₀ 8.4) [1856], AMD070 (p <i>C</i> ₅₀ 7.9) [1750], HIV-Tat	–	–
Allosteric modulators	–	–	–
Labelled ligands	[¹²⁵ I] CXCL12α (human) (Agonist) (p <i>K</i> _d 8.1–8.4) [421, 746]	[¹²⁵ I] CXCL13 (mouse) (Agonist) [222] – Mouse	[¹²⁵ I] CXCL16 (human) (Agonist)
Comments	–	–	–

Nomenclature	CX₃CR1	XCR1	ACKR1	ACKR2
HGNC, UniProt	CX3CR1 , P49238	XCR1 , P46094	ACKR1 , Q16570	ACKR2 , O00590
Endogenous ligands	–	–	CXCL5 (CXCL5 , P42830), CXCL6 (CXCL6 , P80162), CXCL8 (CXCL8 , P10145), CXCL11 (CXCL11 , O14625), CCL2 (CCL2 , P13500), CCL5 (CCL5 , P13501), CCL7 (CCL7 , P80098), CCL11 (CCL11 , P51671), CCL14 (CCL14 , Q16627), CCL17 (CCL17 , Q92583)	CCL2 (CCL2 , P13500), CCL3 (CCL3 , P10147), CCL4 (CCL4 , P13236), CCL5 (CCL5 , P13501), CCL7 (CCL7 , P80098), CCL8 (CCL8 , P80075), CCL11 (CCL11 , P51671), CCL13 (CCL13 , Q99616), CCL14 (CCL14 , Q16627), CCL17 (CCL17 , Q92583), CCL22 (CCL22 , O00626)
Endogenous agonists	CX₃CL1 (CX3CL1 , P78423) (Selective) (p <i>C</i> ₅₀ 8.9) [577]	XCL1 (XCL1 , P47992) (Selective), XCL2 (XCL2 , Q9UBD3) (Selective)	–	–
Labelled ligands	[¹²⁵ I] CX₃CL1 (human) (Agonist)	–	–	–
Comments	–	When fused with secreted alkaline phosphatase (SEAP), XCL1 functions as a probe at XCR1	ACKR1 is used by <i>Plasmodium vivax</i> and <i>Plasmodium knowlesi</i> for entering erythrocytes.	–

Nomenclature	ACKR3	ACKR4	CCRL2
HGNC, UniProt	ACKR3, P25106	ACKR4, Q9NPB9	CCRL2, O00421
Endogenous ligands	–	–	chemerin C-terminal peptide, CCL19 (CCL19, Q99731) [95]
Endogenous agonists	CXCL12α (CXCL12, P48061) (pEC ₅₀ 7.5–7.9) [640, 1785], CXCL11 (CXCL11, O14625) , adrenomedullin {Mouse} [965] – Mouse	CCL19 (CCL19, Q99731) (pK _i 8.4) [1997], CCL25 (CCL25, O15444) (pK _i 7.6) [1997], CCL21 (CCL21, O00585) (pK _i 6.9) [1997]	–

Comments: Mouse Cxcr binds iodinated mouse KC (CXCL1 {Mouse}) and mouse MIP-2 (CXCL2 {Mouse}) with high affinity (mouse KC and MIP-2 are homologues of human CXCL1 (CXCL1, P09341), CXCL2 (CXCL2, P19875) and CXCL3 (CXCL3, P19876)), but shows low affinity for human IL-8 (CXCL8 (CXCL8, P10145)).

Specific chemokine receptors facilitate cell entry by microbes, such as ACKR1 for *Plasmodium vivax*, and CCR5 for HIV-1. Virally encoded chemokine receptors are known (e.g. US28, a homologue of CCR1 from human cytomegalovirus and ORF74, which encodes a homolog of CXCR2 in *Herpesvirus saimiri* and Herpesvirus-68), but their role in viral life cycles is not established. Viruses can exploit or subvert the chemokine system by producing chemokine antagonists and scavengers.

The CC chemokine family (CCL1–28) includes I309 (CCL1 (CCL1, P22362)), MCP-1 (CCL2 (CCL2, P13500)), MIP-1 α (CCL3 (CCL3, P10147)), MIP-1 β (CCL4 (CCL4, P13236)), RANTES (CCL5 (CCL5,

P13501)), MCP-3 (CCL7 (CCL7, P80098)), MCP-2 (CCL8 (CCL8, P80075)), eotaxin (CCL11 (CCL11, P51671)), MCP-4 (CCL13 (CCL13, Q99616)), HCC-1 (CCL14 (CCL14, Q16627)), Lkn-1/HCC-2 (CCL15 (CCL15, Q16663)), TARC (CCL17 (CCL17, Q92583)), ELC (CCL19 (CCL19, Q99731)), LARC (CCL20 (CCL20, P78556)), SLC (CCL21 (CCL21, O00585)), MDC (CCL22 (CCL22, O00626)), MPIF-1 (CCL23 (CCL23, P55773)), eotaxin-2 (CCL24 (CCL24, O00175)), TECK (CCL25 (CCL25, O15444)), eotaxin (CCL26 (CCL26, Q9Y258)), eskine/CTACK (CCL27 (CCL27, Q9Y4X3)) and MEC (CCL28 (CCL28, Q9NRJ3)). The CXC chemokine family (CXCL1–17) includes GRO α (CXCL1 (CXCL1, P09341)), GRO β (CXCL2 (CXCL2, P19875)), GRO γ (CXCL3 (CXCL3, P19876)), platelet factor 4 (CXCL4 (PF4, P02776)), ENA78 (CXCL5 (CXCL5, P42830)), GCP-2 (CXCL6 (CXCL6, P80162)), NAP-2 (CXCL7 (PPBP, P02775)), IL-8 (CXCL8 (CXCL8, P10145)), MIG (CXCL9 (CXCL9, Q07325)), IP10 (CXCL10 (CXCL10, P02778)), I-TAC (CXCL11 (CXCL11, O14625)), SDF-1 (CXCL12, i.e. CXCL12 α (CXCL12, P48061) and CXCL12 β (CXCL12, P48061)), BLC (CXCL13

(CXCL13, O43927)), BRAK (CXCL14 (CXCL14, O95715)), mouse lungkine (CXCL15 {Mouse}) SR-PSOX (CXCL16 (CXCL16, Q9H2A7)) and CXCL17 (CXCL17, Q6UXB2). The CX₃C chemokine (CX₃CL1 (CX3CL1, P78423)) is also known as fractalkine (neurotactin in the mouse). Like CXCL16 (CXCL16, Q9H2A7), and unlike other chemokines, CX₃CL1 (CX3CL1, P78423) is multimodular containing a chemokine domain, an elongated mucin-like stalk, a transmembrane domain and a cytoplasmic tail. Both plasma membrane-associated and shed forms have been identified. The C chemokine (XCL1 (XCL1, P47992)) is also known as lymphotactin.

Two chemokine receptor antagonists have now been approved by the FDA: the CCR5 antagonist **maraviroc** (Pfizer) for treatment of HIV/AIDS in patients with CCR5-using strains; and the CXCR4 antagonist **plerixafor** (Sanofi) for hematopoietic stem cell mobilization with **G-CSF (CSF3, P09919)** in patients undergoing transplantation in the context of chemotherapy for lymphoma and multiple myeloma.

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Cholecystokinin receptors

G protein-coupled receptors → Cholecystokinin receptors

Overview: Cholecystokinin receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on CCK receptors [1403]**) are activated by the endogenous peptides cholecystokinin-8 (CCK-8 (CCK, P06307)), CCK-33 (CCK, P06307), CCK-58 (CCK, P06307) and gastrin (gastrin-17 (GAST, P01350)).

There are only two distinct subtypes of CCK receptors, CCK₁ and CCK₂ receptors [992, 1986], with some alternatively spliced forms most often identified in neoplastic cells. The CCK receptor subtypes are distinguished by their peptide selectivity, with the CCK₁ receptor requiring the carboxyl-terminal heptapeptide-amide that includes a

sulfated tyrosine for high affinity and potency, while the CCK₂ receptor requires only the carboxyl-terminal tetrapeptide shared by both CCK and gastrin peptides. These receptors have characteristic and distinct distributions, with both present in both the central nervous system and peripheral tissues.

Nomenclature	CCK ₁ receptor	CCK ₂ receptor
HGNC, UniProt	CCKAR, P32238	CCKBR, P32239
Rank order of potency	CCK-8 (CCK, P06307) ≫ gastrin-17 (GAST, P01350), desulfated cholecystokinin-8 > CCK-4 (CCK, P06307)	CCK-8 (CCK, P06307) ≥ gastrin-17 (GAST, P01350), desulfated cholecystokinin-8, CCK-4 (CCK, P06307)
Endogenous agonists	–	desulfated cholecystokinin-8 (pIC ₅₀ 8.3–8.7) [1071], gastrin-17 (GAST, P01350) (Selective) (pIC ₅₀ 8.3) [805] – Mouse, CCK-4 (CCK, P06307) (pIC ₅₀ 7.5) [832], desulfated gastrin-14 (GAST, P01350), desulfated gastrin-17 (GAST, P01350), desulfated gastrin-34 (GAST, P01350), desulfated gastrin-71 (GAST, P01350), gastrin-14 (GAST, P01350), gastrin-34 (GAST, P01350), gastrin-71 (GAST, P01350)
Selective agonists	A-71623 (pIC ₅₀ 8.4) [63] – Rat, JMV180 (pIC ₅₀ 8.3) [926], GW-5823 (pIC ₅₀ 7.6) [737]	RB-400 (pK _i 9.1) [123] – Rat, PBC-264 (pIC ₅₀ 9.1) [844] – Rat
Antagonists	linitript (pIC ₅₀ 8.3) [632]	–
Selective antagonists	devazepide (pIC ₅₀ 9.7) [805] – Rat, T-0632 (pIC ₅₀ 9.6) [1861] – Rat, PD-140548 (pIC ₅₀ 8.6) [1748] – Rat, lorglumide (pIC ₅₀ 6.7–8.2) [805, 834] – Rat	YF-476 (pIC ₅₀ 9.7) [196, 1854], GV150013 (pIC ₅₀ 9.4) [1930], L-740093 (pIC ₅₀ 9.2) [1398], YM-022 (pIC ₅₀ 9.2) [1398], JNJ-26070109 (pIC ₅₀ 8.5) [1336], L-365260 (pIC ₅₀ 8.4) [1071], RP73870 (pIC ₅₀ 8) [1115] – Rat, LY262691 (pIC ₅₀ 7.5) [1561] – Rat
Labelled ligands	[³ H]devazepide (Antagonist) (pK _d 9.7) [292], [¹²⁵ I]DTyr-Gly-[(Nle28,31)CCK-26-33 (Agonist) (pIC ₅₀ 9) [1527]	[³ H]PD140376 (Antagonist) (pK _i 9.7–10) [809] – Guinea pig, [¹²⁵ I]PD142308 (Antagonist) (pK _d 9.6) [781] – Guinea pig, [¹²⁵ I]DTyr-Gly-[(Nle28,31)CCK-26-33 (Agonist) (pIC ₅₀ 9) [1527], [¹²⁵ I]gastrin (Agonist) (pIC ₅₀ 9), [³ H]gastrin (Agonist) (pIC ₅₀ 9), [³ H]L365260 (Antagonist) (pK _d 8.2–8.5) [1398], [¹²⁵ I]-BDZ ₂ (Antagonist) (pK _i 8.4) [25]

Comments: While a cancer-specific CCK receptor has been postulated to exist, which also might be responsive to incompletely processed forms of CCK (Gly-extended forms), this has never been isolated. An alternatively spliced form of the CCK₂ receptor in which intron 4 is retained, adding 69 amino acids to the intracellular loop 3

(ICL3) region, has been described to be present particularly in certain neoplasms where mRNA mis-splicing has been commonly observed [1764], but it is not clear that this receptor splice form plays a special role in carcinogenesis. Another alternative splicing event for the CCK₂ receptor was reported [1782], with alternative donor sites in

exon 4 resulting in long (452 amino acids) and short (447 amino acids) forms of the receptor differing by five residues in ICL3, however, no clear functional differences have been observed.

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Class Frizzled GPCRs

G protein-coupled receptors → Class Frizzled GPCRs

Overview: Receptors of the Class Frizzled (FZD, **nomenclature as agreed by the NC-IUPHAR subcommittee on the Class Frizzled GPCRs [1676]**), are GPCRs originally identified in *Drosophila* [285], which are highly conserved across species. FZDs are activated by WNTs, which are cysteine-rich lipoglycoproteins with fundamental functions in ontogeny and tissue homeostasis. FZD signalling was initially divided into two pathways, being either dependent on the accumulation of the transcription regulator β -catenin (*CTNNB1*, P35222) or being β -catenin-independent (often referred to as canonical vs. non-canonical WNT/FZD signalling, respectively). WNT stimulation of FZDs can, in cooperation

with the low density lipoprotein receptors *LRP5* (O75197) and *LRP6* (O75581), lead to the inhibition of a constitutively active destruction complex, which results in the accumulation of β -catenin and subsequently its translocation to the nucleus. β -Catenin, in turn, modifies gene transcription by interacting with TCF/LEF transcription factors. β -Catenin-independent FZD signalling is far more complex with regard to the diversity of the activated pathways. WNT/FZD signalling can lead to the activation of pertussis toxin-sensitive heterotrimeric G proteins [939], the elevation of intracellular calcium [1757], activation of cGMP-specific PDE6 [17] and elevation of cAMP as well as RAC-1, JNK, Rho and Rho kinase signalling [695]. Fur-

thermore, the phosphoprotein Dishevelled constitutes a key player in WNT/FZD signalling. As with other GPCRs, members of the Frizzled family are functionally dependent on the arrestin scaffolding protein for internalization [306], as well as for β -catenin-dependent [235] and -independent [236, 940] signalling. The pattern of cell signalling is complicated by the presence of additional ligands, which can enhance or inhibit FZD signalling (secreted Frizzled-related proteins (sFRP), *Wnt-inhibitory factor* (*WIF1*, Q9Y5W5) (WIF), *sclerostin* (*SOST*, Q9BQB4) or Dickkopf (DKK)), as well as modulatory (co)-receptors with *Ryk*, *ROR1*, *ROR2* and Kremen, which may also function as independent signalling proteins.

Nomenclature	FZD ₁	FZD ₂	FZD ₃	FZD ₄	FZD ₅
HGNC, UniProt	FZD1, Q9UP38	FZD2, Q14332	FZD3, Q9NPG1	FZD4, Q9ULV1	FZD5, Q13467

Nomenclature	FZD ₆	FZD ₇	FZD ₈	FZD ₉	FZD ₁₀
HGNC, UniProt	FZD6, O60353	FZD7, O75084	FZD8, Q9H461	FZD9, O00144	FZD10, Q9ULW2

Nomenclature	SMO
HGNC, UniProt	SMO, Q99835
Antagonists	saridegib (pIC ₅₀ 8.9) [1904], glasdegib (pIC ₅₀ 8.3) [1342], erismodegib (pK _i 8.2) [1979]
Selective antagonists	vismodegib (pK _i 7.8) [1979]

Comments: There is limited knowledge about WNT/FZD specificity and which molecular entities determine the signalling outcome of a specific WNT/FZD pair. Understanding of the coupling to G proteins is incomplete (see [423]). There is also a scarcity of information on basic pharmacological characteristics of FZDs, such as binding constants, ligand specificity or concentration-response relationships [937].

Ligands associated with FZD signalling

WNTs: Wnt-1 (WNT1, P04628), Wnt-2 (WNT2, P09544) (also known as Int-1-related protein), Wnt-2b (WNT2B, Q93097) (also

known as WNT-13), Wnt-3 (WNT3, P56703), Wnt-3a (WNT3A, P56704), Wnt-4 (WNT4, P56705), Wnt-5a (WNT5A, P41221), Wnt-5b (WNT5B, Q9H1J7), Wnt-6 (WNT6, Q9Y6F9), Wnt-7a (WNT7A, O00755), Wnt-7b (WNT7B, P56706), Wnt-8a (WNT8A, Q9H1J5), Wnt-8b (WNT8B, Q93098), Wnt-9a (WNT9A, O14904) (also known as WNT-14), Wnt-9b (WNT9B, O14905) (also known as WNT-15 or WNT-14b), Wnt-10a (WNT10A, Q9GZT5), Wnt-10b (WNT10B, O00744) (also known as WNT-12), Wnt-11 (WNT11, O96014) and Wnt-16 (WNT16, Q9UBV4).

Extracellular proteins that interact with FZDs: norrin (NDP, Q00604), R-spondin-1 (RSPO1, Q2MKA7), R-spondin-2

(RSPO2, Q6UXX9), R-spondin-3 (RSPO3, Q9BXY4), R-spondin-4 (RSPO4, Q2I0M5), sFRP-1 (SFRP1, Q8N474), sFRP-2 (SFRP2, Q96HF1), sFRP-3 (FRZB, Q92765), sFRP-4 (SFRP4, Q6FHJ7), sFRP-5 (SFRP5, Q6FHJ7).

Extracellular proteins that interact with WNTs or LRP5: Dickkopf 1 (DKK1, O94907), WIF1 (Q9Y5W5), sclerostin (SOST, Q9BQB4), kremen 1 (KREMEN1, Q96MU8) and kremen 2 (KREMEN2, Q8NCW0)

Small exogenous ligands: Foxy-5 [1835], Box-5, UM206 [1031], and XWnt8 (P28026) also known as mini-Wnt8.

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Complement peptide receptors

G protein-coupled receptors → Complement peptide receptors

Overview: Complement peptide receptors (nomenclature as agreed by the **NC-IUPHAR subcommittee on Complement peptide receptors** [967]) are activated by the endogenous 75 amino-acid anaphylatoxin polypeptides C3a (C3, P01024), C4a (C4A, P0C0L4) and C5a (C5, P01031), generated upon stimulation of the complement cascade.

Nomenclature	C3a receptor	C5a₁ receptor	C5a₂ receptor
HGNC, UniProt	C3AR1 , Q16581	C5AR1 , P21730	C5AR2 , Q9P296
Rank order of potency	C3a (C3 , P01024) > C5a (C5 , P01031) [41]	C5a (C5 , P01031), C5a des-Arg (C5) > C3a (C3 , P01024) [41]	–
Endogenous agonists	–	ribosomal protein S19 (RPS19 , P39019) [2071]	–
Agonists	E7 (pEC ₅₀ 8.7) [43], compound 21 [PMID: 25259874] (pEC ₅₀ 7.7) [1571], SQ007-5 (Partial agonist) (pEC ₅₀ 6.7) [124], Ac-RHYPLWR (pEC ₅₀ 6) [672]	N-methyl-Phe-Lys-Pro-D-Cha-Cha-D-Arg-CO₂H (pIC ₅₀ 7.6) [916 , 989]	–
Antagonists	SB290157 (pIC ₅₀ 7.6) [40], compound 4 [PMID: 25259874] (pIC ₅₀ 5.9) [1571]	CHIPS (pK _d 9) [1522], W54011 (pK _i 8.7) [1819], AcPhe-Orn-Pro-D-Cha-Trp-Arg (pIC ₅₀ 7.9) [2039], N-methyl-Phe-Lys-Pro-D-Cha-Trp-D-Arg-CO₂H (pIC ₅₀ 7.2) [989]	–
Labelled ligands	[¹²⁵I]C3a (human) (Agonist) (pK _d 8.4) [296]	[¹²⁵I]C5a (human) (Agonist) (pK _d 8.7) [803]	[¹²⁵I]C5a (human) (Agonist)
Comments	–	–	Binds C5a complement factor, but appears to lack G protein signalling and has been termed a decoy receptor [1684].

Comments: [SB290157](#) has also been reported to have agonist properties at the C3a receptor [[1218](#)]. The putative chemoattractant receptor termed C5a₂ (also known as GPR77, C5L2) binds [¹²⁵I]C5a with no clear signalling function, but has a putative role opposing inflammatory responses [[257](#), [568](#), [585](#)]. Binding to this site may be

displaced with the rank order [C5a des-Arg](#) ([C5](#)) > [C5a](#) ([C5](#), [P01031](#)) [[257](#), [1440](#)] while there is controversy over the ability of [C3a](#) ([C3](#), [P01024](#)) and [C3a des Arg](#) ([C3](#), [P01024](#)) to compete [[778](#), [894](#), [895](#), [1440](#)]. C5a₂ appears to lack G protein signalling and has been termed a decoy receptor [[1684](#)]. However, C5a₂ does recruit arrestin after ligand binding, which might provide a signaling pathway

for this receptor [[89](#), [1937](#)], and forms heteromers with C5a₁. C5a, but not C5a-des Arg, induces upregulation of heteromer formation between complement C5a receptors C5aR and C5L2 [[380](#)]. There are also reports of pro-inflammatory activity of C5a₂, mediated by HMGB1, but the signaling pathway that underlies this is currently unclear (reviewed in [[1095](#)]).

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Corticotropin-releasing factor receptors

G protein-coupled receptors → Corticotropin-releasing factor receptors

Overview: Corticotropin-releasing factor (CRF, **nomenclature as agreed by the NC-IUPHAR subcommittee on Corticotropin-releasing Factor Receptors [716]**) receptors are activated by the endogenous peptides **corticotrophin-releasing hormone (CRH, P06850)**, a 41 amino-acid

peptide, **urocortin 1 (UCN, P55089)**, 40 amino-acids, **urocortin 2 (UCN2, Q96RP3)**, 38 amino-acids and **urocortin 3 (UCN3, Q969E3)**, 38 amino-acids. CRF₁ and CRF₂ receptors are activated non-selectively by **corticotrophin-releasing hormone (CRH, P06850)** and **urocortin 1 (UCN, P55089)**. Binding to CRF receptors can be

conducted using [¹²⁵I]Tyr⁰-CRF or [¹²⁵I]Tyr⁰-sauvagine with K_d values of 0.1–0.4 nM. CRF₁ and CRF₂ receptors are non-selectively antagonized by **α-helical CRF, D-Phe-CRF-(12-41)** and **astressin**.

Nomenclature	CRF ₁ receptor	CRF ₂ receptor
HGNC, UniProt	CRHR1, P34998	CRHR2, Q13324
Endogenous agonists	–	urocortin 2 (UCN2, Q96RP3) (Selective) (pK _d 8.5–8.6) [392], urocortin 3 (UCN3, Q969E3) (Selective) (pK _d 7.9–8) [392]
Antagonists	SSR125543A (pK _i 8.7) [663]	–
Selective antagonists	CP 154,526 (pIC ₅₀ 9.3–10.4) [1153] – Rat, DMP696 (pK _i 8.3–9) [726], NBI27914 (pK _i 8.3–9) [298], R121919 (pK _i 8.3–9) [2133], antalarmin (pK _i 8.3–9) [2001], CP376395 (pIC ₅₀ 8.3) [307] – Rat, CRA1000 (pIC ₅₀ 6.4–7.1) [284]	antisauvagine (pK _d 8.8–9.6) [394], K41498 (pK _i 9.2) [1048], K31440 (pK _i 8.7–8.8) [1622]

Comments: A CRF binding protein has been identified ([CRHBP, P24387](#)) to which both **corticotrophin-releasing hormone (CRH, P06850)** and **urocortin 1 (UCN, P55089)** bind with high affinities, which has been suggested to bind and inactivate circulating **corticotrophin-releasing hormone (CRH, P06850)** [[1489](#)].

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Dopamine receptors

G protein-coupled receptors → Dopamine receptors

Overview: Dopamine receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Dopamine Receptors [1677]**) are commonly divided into D₁-like (D₁ and D₅) and D₂-like (D₂, D₃ and D₄) families, where the endogenous agonist is [dopamine](#).

Nomenclature	D₁ receptor	D₂ receptor
HGNC, UniProt	DRD1 , P21728	DRD2 , P14416
Endogenous agonists	dopamine (pK _i 4.3–5.6) [1823 , 1884]	dopamine (pK _i 4.7–7.2) [245 , 545 , 1653]
Agonists	fenoldopam (pK _i 6.5–7.9) [1884]	rotigotine (pK _i 10.2) [424], cabergoline (Partial agonist) (pK _i 9–9.2) [1279], aripiprazole (Partial agonist) (pK _i 9.1) [2111], bromocriptine (pK _i 7.3–8.3) [545 , 1279 , 1653], MLS1547 (Biased agonist) (pK _i 8.2) [544], ropinirole (pK _i 8.1) [732], apomorphine (Partial agonist) (pK _i 5.7–7.5) [245 , 545 , 1279 , 1653 , 1776], pramipexole (pK _i 5.1–7.4) [1273 , 1653], benzquinamide (pK _i 5.4) [643]
(Sub)family-selective agonists	A68930 (pEC ₅₀ 6.8) [1381], SKF-38393 (Partial agonist) (pK _i 6.2–6.8) [1823 , 1884]	quinpirole (pK _i 4.9–7.7) [245 , 1273 , 1473 , 1776 , 1778 , 1940]
Selective agonists	SKF-83959 (Biased agonist) (pEC ₅₀ 9.7) [364], SKF-81297 (pK _i 8.7) [46] – Rat	sumanirole (pK _i 8.1) [1239]
Antagonists	flupentixol (pK _i 7–8.4) [1823 , 1884]	blonanserin (pK _i 9.9) [1421], pipotiazine (pK _i 9.7) [1777], perphenazine (pK _i 8.9–9.6) [1008 , 1691], risperidone (pK _i 9.4) [60], perospirone (pK _i 9.2) [1692], trifluoperazine (pK _i 8.9–9) [1008 , 1693], asenapine (pK _i 8.9) [1711], sertindole (pK _i 8–8.9) [986 , 1008 , 1691], fluphenazine (pK _i 8.8) [1647], flupentixol (pK _i 8.8) [545], pimozide (pK _i 7–8.8) [545 , 1776], olanzapine (pK _i 8.7) [60], mesoridazine (pK _i 8.7) [326], ziprasidone (pK _i 8.6) [60], prochlorperazine (pK _i 8.4) [68], loxapine (pK _i 7.9–8.3) [1008 , 1693], (-)- sulpiride (pK _i 6.3–8) [545 , 1776 , 1860], amisulpride (pK _i 7.9–8) [1195 , 1776], metoclopramide (pK _i 7.5) [1221] – Mouse, quetiapine (pK _i 7.2) [60], trans-flupenthixol (pK _i 6.9) [545], clozapine (pK _i 5.8–6.9) [545 , 1164 , 1711 , 1776 , 1860], promazine (pK _i 6.5) [246]
(Sub)family-selective antagonists	SCH-23390 (pK _i 7.4–9.5) [1823 , 1884], SKF-83556 (pK _i 9.5) [1823], ecopipam (pK _i 8.3) [1885]	haloperidol (pK _i 7.4–8.8) [545 , 1164 , 1273 , 1776 , 1885]
Selective antagonists	–	L-741,626 (pK _i 7.9–8.5) [655 , 1020], domperidone (pK _i 7.9–8.4) [545 , 1776], raclopride (pK _i 8) [1281], ML321 (pK _i 7) [2058 , 2059]
Labelled ligands	[³H]SCH-23390 (Antagonist) (pK _d 9.5) [2127]	[³H]spiperone (Antagonist) (pK _d 10.2) [239 , 767 , 2125] – Rat
Labelled ligands	[¹²⁵I]SCH23982 (Antagonist) (pK _d 9.5) [408]	[³H]raclopride (Antagonist) (pK _d 8.9) [1028] – Rat

Nomenclature	D ₃ receptor	D ₄ receptor	D ₅ receptor
HGNC, UniProt	<i>DRD3</i> , P35462	<i>DRD4</i> , P21917	<i>DRD5</i> , P21918
Endogenous agonists	dopamine (pK _i 6.4–7.3) [245, 545, 1653, 1778]	dopamine (pK _i 7.6) [1940]	dopamine (pK _i 6.6) [1823]
Agonists	pramipexole (pK _i 8.4–8.7) [1273, 1653], bromocriptine (Partial agonist) (pK _i 7.1–8.2) [545, 1279, 1653], ropinirole (pK _i 7.7) [732], apomorphine (Partial agonist) (pK _i 6.1–7.6) [245, 545, 1279, 1653, 1776]	apomorphine (Partial agonist) (pK _i 8.4) [1279]	–
(Sub)family-selective agonists	quinpirole (pK _i 6.4–8) [245, 1273, 1281, 1473, 1653, 1776, 1778, 1940]	quinpirole (pK _i 7.5) [1279, 1473, 1940]	A68930 (pEC ₅₀ 6.6) [1381]
Selective agonists	PD 128907 (pK _i 7.6–7.7) [1539, 1653]	PD168,077 (Partial agonist) (pK _i 8.8) [995] – Rat, A412997 (pK _i 8.1) [1319] – Rat, A412997 (pK _i 8.1) [1319]	–
Antagonists	perospirone (pK _i 9.6) [1776], sertindole (pK _i 8–8.8) [60, 1675, 1691], prochlorperazine (pK _i 8.4) [68], (-)-sulpiride (pK _i 6.7–7.7) [545, 1776, 1860], loxapine (pK _i 7.7) [1691], domperidone (pK _i 7.1–7.6) [545, 1776], promazine (pK _i 6.8) [246]	perospirone (pK _i 10.1) [1694], sertindole (pK _i 7.8–9.1) [246, 1691, 1693, 1694], sonepiprazole (pK _i 8.9) [1668], loxapine (pK _i 8.1) [1693]	–
(Sub)family-selective antagonists	haloperidol (pK _i 7.5–8.6) [545, 1711, 1776, 1885]	haloperidol (pK _i 8.7–8.8) [1033, 1711, 1885]	SCH-23390 (pK _i 7.5–9.5) [1823], SKF-83556 (pK _i 9.4) [1823], ecopipam (pK _i 8.3) [1823]
Selective antagonists	S33084 (pK _i 9.6) [1278], nafadotride (pK _i 9.5) [1654], PG01037 (pK _i 9.2) [656], NGB 2904 (pK _i 8.8) [2055], SB 277011-A (pK _i 8) [1569], (+)-S-14297 (pK _i 6.9–7.9) [1275, 1281]	L745870 (pK _i 9.4) [1020], A-381393 (pK _i 8.8) [1361], L741742 (pK _i 8.5) [1609], ML398 (pK _i 7.4) [138]	–
Selective allosteric modulators	SB269652 (Negative) (pK _i ~9) [558]	–	–
Labelled ligands	–	[³ H]spiperone (Antagonist) (pK _d 9.5) [749, 1940]	[³ H]SCH-23390 (Antagonist) (pK _d 9.2) [1580]
Labelled ligands	[³ H]spiperone (Antagonist) (pK _d 9.9) [767, 2125] – Rat, [³ H]7-OH-DPAT (Agonist) (pK _d 9.6) [1581], [³ H]PD128907 (Agonist) (pK _d 9) [27]	[¹²⁵ I]L750667 (Antagonist) (pK _d 9.8) [1473], [³ H]NGD941 (Antagonist) (pK _d 8.3) [1533]	[¹²⁵ I]SCH23982 (Antagonist) (pK _d 9.1) – Unknown

Comments: The selectivity of many of these agents is less than two orders of magnitude. [³H]raclopride exhibits similar high affinity for D₂ and D₃ receptors (low affinity for D₄), but has been used to label D₂ receptors in the presence of a D₃-selective antagonist.

[³H]7-OH-DPAT has similar affinity for D₂ and D₃ receptors, but labels only D₃ receptors in the absence of divalent cations. The pharmacological profile of the D₅ receptor is similar to, yet distinct from, that of the D₁ receptor. The splice variants of the D₂ receptor are

commonly termed D_{2S} and D_{2L} (short and long). The *DRD4* gene encoding the D₄ receptor is highly polymorphic in humans, with allelic variations of the protein from amino acid 387 to 515.

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Endothelin receptors

G protein-coupled receptors → Endothelin receptors

Overview: Endothelin receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Endothelin Receptors [395]**) are activated by the endogenous 21 amino-acid peptides endothelins 1-3 (endothelin-1 (*EDN1*, P05305), endothelin-2 (*EDN2*, P20800) and endothelin-3 (*EDN3*, P14138)).

Nomenclature	ET _A receptor	ET _B receptor
HGNC, UniProt	<i>EDNRA</i> , P25101	<i>EDNRB</i> , P24530
Family selective agonists	endothelin-1 (<i>EDN1</i> , P05305) = endothelin-2 (<i>EDN2</i> , P20800) > endothelin-3 (<i>EDN3</i> , P14138) [1178]	endothelin-1 (<i>EDN1</i> , P05305) = endothelin-2 (<i>EDN2</i> , P20800), endothelin-3 (<i>EDN3</i> , P14138)
Selective agonists	–	sarafotoxin S6c (pK _d 8.8–9.8) [1016, 1616], BQ 3020 (pK _i 9.7) [1576], [Ala ^{1,3,11,15}]ET-1 (pK _d 8.7–9.2) [1300], IRL 1620 (pK _i 8.7) [1991]
(Sub)family-selective antagonists	SB209670 (pK _B 9.4) [474] – Rat, TAK 044 (pA ₂ 8.4) [1993] – Rat, bosentan (pA ₂ 7.2) [354] – Rat	SB209670 (pK _B 9.4) [474] – Rat, TAK 044 (pA ₂ 8.4) [1993] – Rat, bosentan (pK _i 7.1) [1349]
Selective antagonists	atrasentan (pA ₂ 9.2–10.5) [1446], PD-156707 (pK _d 9–9.8) [1180], macitentan (pIC ₅₀ 9.3) [174], sitaxsentan (pA ₂ 8) [2047], FR139317 (Inverse agonist) (pIC ₅₀ 7.3–7.9) [1178], ambrisentan (pIC ₅₀ 7.7) [175], BQ123 (pA ₂ 6.9–7.4) [1178], avosentan (pIC ₅₀ 7.3) [210], ambrisentan (pA ₂ 7.1) [175]	A192621 (pK _d 8.1) [2145], BQ788 (pK _d 7.9–8) [1616], IRL 2500 (pK _d 7.2) [1616], Ro 46-8443 (pIC ₅₀ 7.2) [209]
Labelled ligands	[¹²⁵ I]PD164333 (Antagonist) (pK _d 9.6–9.8) [398], [³ H]S0139 (Antagonist) (pK _d 9.2), [¹²⁵ I]PD151242 (Antagonist) (pK _d 9–9.1) [399], [³ H]BQ123 (Antagonist) (pK _d 8.5) [817]	[¹²⁵ I]IRL1620 (Agonist) (pK _d 9.9–10.1) [1362], [¹²⁵ I]BQ3020 (Agonist) (pK _d 8.3–10) [702, 1300, 1495], [¹²⁵ I][Ala ^{1,3,11,15}]ET-1 (Agonist) (pK _d 9.7) [1300]

Comments: Splice variants of the ET_A receptor have been identified in rat pituitary cells; one of these, ET_AR-C13, appeared to show loss of function with comparable plasma membrane expression to wild type receptor [713]. Subtypes of the ET_B receptor have been proposed, although gene disruption studies in mice suggest that only a single gene product exists [1295].

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G protein-coupled estrogen receptor

G protein-coupled receptors → G protein-coupled estrogen receptor

Overview: The G protein-coupled estrogen receptor (GPER, **nomenclature as agreed by the NC-IUPHAR Subcommittee on the G protein-coupled estrogen receptor [1536]**) was identified following observations of estrogen-evoked cyclic AMP signalling in breast cancer cells [61], which mirrored the differential expression of an orphan 7-transmembrane receptor GPR30 [265]. There are observations of both cell-surface and intracellular expression of the GPER receptor [1573, 1877].

Nomenclature	GPER
HGNC, UniProt	GPER1, Q99527
Selective agonists	G1 (pK _i 8) [176]
Selective antagonists	G36 (pIC ₅₀ 6.8–6.9) [414], G15 (pIC ₅₀ 6.7) [413]
Labelled ligands	[³ H]17β-estradiol (Agonist) (pK _d 8.5–8.6) [1877]

Comments: Antagonists at the nuclear estrogen receptor, such as fulvestrant and tamoxifen [515], as well as the flavonoid ‘phytoestrogens’ genistein and quercetin [1177], are agonists at GPER receptors.

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Formylpeptide receptors

G protein-coupled receptors → Formylpeptide receptors

Overview: The formylpeptide receptors (nomenclature agreed by the **NC-IUPHAR Subcommittee on the formyl peptide receptor family [2092]**) respond to exogenous ligands such as the bacterial product **fMet-Leu-Phe** (fMLP) and endogenous ligands such as **annexin I** (*ANXA1*, **P04083**), **cathepsin G** (*CTSG*, **P08311**), amyloid β 42, serum amyloid A and **spinorphin**, derived from **β -haemoglobin** (*HBB*, **P68871**).

Nomenclature	FPR1	FPR2/ALX	FPR3
HGNC, UniProt	<i>FPR1</i> , P21462	<i>FPR2</i> , P25090	<i>FPR3</i> , P25089
Rank order of potency	fMet-Leu-Phe > cathepsin G (<i>CTSG</i> , P08311) > annexin I (<i>ANXA1</i> , P04083) [1058 , 1821]	LXA ₄ =aspirin triggered lipoxin A4=ATLa2>LTC ₄ =LTD ₄ » 15-deoxy-LXA ₄ »fMet-Leu-Phe [352 , 519 , 521 , 651 , 1846]	–
Endogenous agonists	–	LXA ₄ (Selective) (pEC ₅₀ ~12) [1006], resolvin D1 (Selective) (pEC ₅₀ ~11.9) [1006], aspirin triggered lipoxin A4 (Selective)	F2L (<i>HEBP1</i> , Q9NRV9) (Selective) (pEC ₅₀ 8–8.2) [1274]
Agonists	fMet-Leu-Phe (pEC ₅₀ 10.1–10.2) [546 , 1734]	–	–
Selective agonists	–	ATLa2 [662]	–
Endogenous antagonists	spinorphin (Selective) (pIC ₅₀ 4.3) [1099 , 1348]	–	–
Antagonists	t-Boc-FLFLF (pK _i 6–6.5) [2008]	–	–
Selective antagonists	cyclosporin H (pK _i 6.1–7.1) [2008 , 2078]	–	–
Labelled ligands	[³ H]fMet-Leu-Phe (Agonist) (pK _d 7.6–9.3) [990]	[³ H]LXA ₄ (Agonist) (pK _d 9.2–9.3) [519 , 520]	–
Comments	A FITC-conjugated fMLP analogue has been used for binding to the mouse recombinant receptor [724].	The agonist activity of the lipid mediators described has been questioned [697 , 1513], which may derive from batch-to-batch differences, partial agonism or biased agonism. Recent results from Cooray <i>et al.</i> (2013) [365] have addressed this issue and the role of homodimers and heterodimers in the intracellular signaling .	–

Comments: Note that the data for FPR2/ALX are also reproduced on the [leukotriene](#) receptor page.

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Free fatty acid receptors

G protein-coupled receptors → Free fatty acid receptors

Overview: Free fatty acid receptors (FFA, **nomenclature as agreed by the NC-IUPHAR Subcommittee on free fatty acid receptors [396, 1803]**) are activated by free fatty acids. Long-chain saturated and unsaturated fatty acids (C14:0 (*myristic acid*), C16:0 (*palmitic acid*), C18:1 (*oleic acid*), C18:2

(*linoleic acid*), C18:3, (*α-linolenic acid*), C20:4 (*arachidonic acid*), C20:5,n-3 (*EPA*), C22:6,n-3 (*docosahexaenoic acid*)) activate FFA1 [218, 833, 998] and FFA4 receptors [757, 812, 1427], while short chain fatty acids (C2 (*acetic acid*), C3 (*propanoic acid*), C4 (*butyric acid*) and C5 (*pentanoic acid*)) activate FFA2 [226, 1057,

1399] and FFA3 [226, 1057] receptors. In addition, thiazolidinedione PPAR γ agonists such as *rosiglitazone* activate FFA1 (pEC₅₀ 5.2; [999, 1768, 1802]) and small molecule allosteric modulators, such as *4-CMTB*, have recently been characterised for FFA2 [801, 1070, 1769].

	FFA1 receptor	FFA2 receptor	FFA3 receptor	FFA4 receptor	GPR42
Nomenclature	FFA1 receptor	FFA2 receptor	FFA3 receptor	FFA4 receptor	GPR42
HGNC, UniProt	<i>FFAR1</i> , O14842	<i>FFAR2</i> , O15552	<i>FFAR3</i> , O14843	<i>FFAR4</i> , Q5NUL3	<i>GPR42</i> , O15529
Endogenous agonists	<i>docosahexaenoic acid</i> (pEC ₅₀ 5.4–6) [218, 833]	–	–	<i>α-linolenic acid</i> (pEC ₅₀ 5.5) [1727]	–
Agonists	<i>fasiglifam</i> (pEC ₅₀ 7.1) [893, 1791, 1909]	–	–	–	–
(Sub)family-selective agonists	<i>α-linolenic acid</i> (pEC ₅₀ 4.6–5.7) [218, 833, 998], <i>oleic acid</i> (pEC ₅₀ 3.9–5.7) [218, 833, 998], <i>myristic acid</i> (pEC ₅₀ 4.5–5.1) [218, 833, 998]	<i>propanoic acid</i> (pEC ₅₀ 3–4.9) [226, 1057, 1399, 1670], <i>acetic acid</i> (pEC ₅₀ 3.1–4.6) [226, 1057, 1399, 1670], <i>butyric acid</i> (pEC ₅₀ 2.9–4.6) [226, 1057, 1399, 1670], <i>trans-2-methylcrotonic acid</i> (pEC ₅₀ 3.8) [1670], <i>1-methylcyclopropanecarboxylic acid</i> (pEC ₅₀ 2.6) [1670]	<i>propanoic acid</i> (pEC ₅₀ 3.9–5.7) [226, 1057, 1670, 2063], <i>butyric acid</i> (pEC ₅₀ 3.8–4.9) [226, 1057, 1670, 2063], <i>1-methylcyclopropanecarboxylic acid</i> (pEC ₅₀ 3.9) [1670], <i>acetic acid</i> (pEC ₅₀ 2.8–3.9) [226, 1057, 1670, 2063]	<i>myristic acid</i> (pEC ₅₀ 5.2) [1996], <i>oleic acid</i> (pEC ₅₀ 4.7) [1996]	–
Selective agonists	<i>AMG-837</i> (pEC ₅₀ 8.5) [1110], <i>TUG-770</i> (pEC ₅₀ 8.2) [332], <i>GW9508</i> (pEC ₅₀ 7.3) [217], <i>linoleic acid</i> (pEC ₅₀ 4.4–5.7) [218, 833, 998]	<i>compound 1</i> [PMID: 23589301] (pEC ₅₀ 7.1) [800] – Rat, <i>(S)-4-CMTB</i> (pEC ₅₀ 6.4) [801, 1070]	–	<i>compound A</i> [PMID 24997608] (pEC ₅₀ 7.6) [1428], <i>TUG-891</i> (pEC ₅₀ 7) [1727] – Unknown, <i>NCG21</i> (pEC ₅₀ 5.9) [1829]	–

(continued)	FFA1 receptor	FFA2 receptor	FFA3 receptor	FFA4 receptor	GPR42
Nomenclature	FFA1 receptor	FFA2 receptor	FFA3 receptor	FFA4 receptor	GPR42
Selective antagonists	GW1100 (pIC ₅₀ 6) [217]	GLPG0974 (pIC ₅₀ 8.1) [1512], CATPB (pIC ₅₀ 6.5) [801]	–	–	–
Comments	Antagonist GW1100 has been shown to reduce [³⁵ S]GTPγS binding in FFAR1-expressing cells [1802]. GW1100 is also an oxytocin receptor antagonist [217]. TUG-770 and GW9508 are both approximately 100 fold selective for FFA1 over FFA4 [217, 332]. AMG-837 and the related analogue AM6331 have been suggested to have an allosteric mechanism of action at FFA1, with respect to the orthosteric fatty acid binding site [1110, 2064].	–	Beta-hydroxybutyrate has been reported to antagonise FFA3 responses to short chain fatty acids [951]. A range of FFA3 selective molecules with agonist and antagonist properties, but which bind at sites distinct from the short chain fatty acid binding site, have recently been described [799].	compound A [PMID 24997608] exhibits more than 1000 fold selectivity [1428], and TUG-891 50-1000 fold selectivity for FFA4 over FFA1 [1727], dependent on the assay. NGC21 exhibits approximately 15 fold selectivity for FFA4 over FFA1 [1820].	–

Comments: Short (361 amino acids) and long (377 amino acids) splice variants of human FFA4 have been reported [1318], which differ by a 16 amino acid insertion in intracellular loop 3, and exhibit differences in intracellular signalling properties in recombinant sys-

tems [1996]. The long FFA4 splice variant has not been identified in other primates or rodents to date [757, 1318].

GPR42 was originally described as a pseudogene within the family (ENSM0025000002583), but the recent discovery of several poly-

morphisms suggests that some versions of GPR42 may be functional [1101]. GPR84 is a structurally-unrelated G protein-coupled receptor which has been found to respond to medium chain fatty acids [1981].

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GABA_B receptors

G protein-coupled receptors → GABA_B receptors

Overview: Functional GABA_B receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on GABA_B receptors** [194, 1507]) are formed from the heterodimerization of two similar 7TM subunits termed GABA_{B1} and GABA_{B2} [194, 478, 1506, 1507, 1926]. GABA_B receptors are widespread in the CNS and regulate both pre- and postsynaptic activity. The GABA_{B1} subunit, when expressed alone, binds both antagonists and agonists, but the affinity of the latter is generally 10–100-fold less than for the native receptor. The GABA_{B1} subunit when expressed alone is not transported to the cell membrane and is non-functional. However, Richer *et al.* (2008) report that GABA_{B1} alone can control ERK/MAPK pathway activity [1585]. Co-expression of GABA_{B1} and GABA_{B2} subunits allows transport of GABA_{B1} to the cell surface and generates a functional receptor that can couple to signal transduction pathways such as high-voltage-activated Ca²⁺ channels (Cav2.1, Cav2.2), or inwardly rectifying potassium channels (Kir3) [144, 194, 195]. The GABA_{B2} subunit also determines the rate of internalisation of the dimeric GABA_B receptor [693]. The GABA_{B1} subunit har-

bours the GABA (orthosteric)-binding site within an extracellular domain (ECD) venus flytrap module (VTM), whereas the GABA_{B2} subunit mediates G protein-coupled signalling [194, 591, 592, 1506]. The two subunits interact by direct allosteric coupling [1313], such that GABA_{B2} increases the affinity of GABA_{B1} for agonists and reciprocally GABA_{B1} facilitates the coupling of GABA_{B2} to G proteins [591, 1013, 1506]. GABA_{B1} and GABA_{B2} subunits assemble in a 1:1 stoichiometry by means of a coiled-coil interaction between α -helices within their carboxy-termini that masks an endoplasmic reticulum retention motif (RXRR) within the GABA_{B1} subunit but other domains of the proteins also contribute to their heteromerization [144, 243, 1506]. Recent evidence indicates that higher order assemblies of GABA_B receptor comprising dimers of heterodimers occur in recombinant expression systems and *in vivo* and that such complexes exhibit negative functional cooperativity between heterodimers [361, 1505]. Adding further complexity, KCTD (potassium channel tetramerization proteins) 8, 12, 12b and 16 associate as tetramers with the carboxy terminus of the GABA_{B2} subunit to impart altered signalling kinetics and agonist potency to the receptor

complex [102, 1680, 1914] and reviewed by [1508]. Four isoforms of the human GABA_{B1} subunit have been cloned. The predominant GABA_{B1(a)} and GABA_{B1(b)} isoforms, which are most prevalent in neonatal and adult brain tissue respectively, differ in their ECD sequences as a result of the use of alternative transcription initiation sites. GABA_{B1(a)}-containing heterodimers localise to distal axons and mediate inhibition of glutamate release in the CA3-CA1 terminals, and GABA release onto the layer 5 pyramidal neurons, whereas GABA_{B1(b)}-containing receptors occur within dendritic spines and mediate slow postsynaptic inhibition [1541, 1955]. Isoforms generated by alternative splicing are GABA_{B1(c)} that differs in the ECD, and GABA_{B1(e)}, which is a truncated protein that can heterodimerize with the GABA_{B2} subunit but does not constitute a functional receptor. Only the 1a and 1b variants are identified as components of native receptors [194]. Additional GABA_{B1} subunit isoforms have been described in rodents and humans [1065] and reviewed by [144].

Nomenclature	GABA _B receptor
Subunits	kcdt12b (Accessory protein), KCTD16 (Accessory protein), KCTD12 (Accessory protein), GABA _{B2} , GABA _{B1} , KCTD8 (Accessory protein)
Agonists	CGP 44532 (pIC ₅₀ 8.6) [551] – Rat, (-)-baclofen (pIC ₅₀ 8.5) [551] – Rat, 3-APPA (pK _i 5.2–7.2) [762], baclofen (pK _i 4.3–6.2) [762, 2041], 3-APMPA (pK _i 5.1) [2041]
Antagonists	CGP 62349 (pK _i 8.5–8.9) [762, 2041], CGP 55845 (pK _i 7.8) [2041], SCH 50911 (pK _i 5.5–6) [762, 2041], CGP 35348 (pK _i 4.4) [2041], 2-hydroxy-saclofen (pIC ₅₀ 4.1) [914] – Rat
Labelled ligands	[³ H]CGP 54626 (Antagonist) (pK _i 9.1) [879] – Rat, [³ H]CGP 62349 (Antagonist) (pK _d 9.1) [922] – Rat, [¹²⁵ I]CGP 64213 (Antagonist) (pK _d 9) [563] – Rat, [¹²⁵ I]CGP 71872 (Antagonist) (pK _d 9) [914] – Rat, [³ H](R)-(-)-baclofen (Agonist)

Subunits

Nomenclature	GABA _{B1}	GABA _{B2}
HGNC, UniProt	GABBR1, Q9UBSS	GABBR2, O75899

Comments: Potencies of agonists and antagonists listed in the table, quantified as IC₅₀ values for the inhibition of [³H]CGP27492 binding to rat cerebral cortex membranes, are from [194, 550, 551]. Radioligand K_D values relate to binding to rat brain membranes. CGP 71872 is a photoaffinity ligand for the GABA_{B1} subunit [122]. CGP27492 (3-APPA), CGP35024 (3-APMPA) and CGP 44532 act as antagonists at human GABA_A ρ1 receptors, with potencies in the low micromolar range [550]. In addition to the ligands listed in the table, Ca²⁺ binds to the VTM of the GABA_{B1} subunit to act as a positive

allosteric modulator of GABA [563]. In cerebellar Purkinje neurones, the interaction of Ca²⁺ with the GABA_B receptor enhances the activity of mGlu₁, through functional cross-talk involving G-protein Gβγ subunits [1590, 1837]. Synthetic positive allosteric modulators with low, or no, intrinsic activity include CGP7930, GS39783, BHF-177 and (+)-BHFF [9, 144, 150, 550]. The site of action of CGP7930 and GS39783 appears to be on the heptahelical domain of the GABA_{B2} subunit [455, 1506]. In the presence of CGP7930 or GS39783, CGP 35348 and 2-hydroxy-saclofen behave as partial agonists [550]. A negative allosteric modulator of GABA_B activity

has been reported [302]. Knock-out of the GABA_{B1} subunit in C57B mice causes the development of severe tonic-clonic convulsions that prove fatal within a month of birth, whereas GABA_{B1}^{-/-} BALB/c mice, although also displaying spontaneous epileptiform activity, are viable. The phenotype of the latter animals additionally includes hyperalgesia, hyperlocomotion (in a novel, but not familiar, environment), hyperdopaminergia, memory impairment and behaviours indicative of anxiety [482, 1932]. A similar phenotype has been found for GABA_{B2}^{-/-} BALB/c mice [582].

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Galanin receptors

G protein-coupled receptors → Galanin receptors

Overview: Galanin receptors (**provisional nomenclature as recommended by NC-IUPHAR [530]**) are activated by the endogenous peptides **galanin** (*GAL*, P22466) and **galanin-like peptide** (*GALP*, Q9UBC7). Human **galanin** (*GAL*, P22466) is a 30 amino-acid non-amidated peptide [499]; in other species, it is 29 amino acids long and C-terminally amidated. Amino acids 1-14 of galanin are highly conserved in mammals, birds, reptiles, amphibia and fish. Shorter peptide species (e.g. human galanin-1-19 [139] and porcine galanin-5-29 [1740]) and N-terminally extended forms (e.g. N-terminally seven and nine residue elongated forms of porcine galanin [140, 1740]) have been reported.

Nomenclature	<i>GAL</i> ₁ receptor	<i>GAL</i> ₂ receptor	<i>GAL</i> ₃ receptor
HGNC, UniProt	<i>GALR1</i> , P47211	<i>GALR2</i> , O43603	<i>GALR3</i> , O60755
Rank order of potency	galanin (<i>GAL</i> , P22466) > galanin-like peptide (<i>GALP</i> , Q9UBC7) [1433]	galanin-like peptide (<i>GALP</i> , Q9UBC7) ≥ galanin (<i>GAL</i> , P22466) [1433]	galanin-like peptide (<i>GALP</i> , Q9UBC7) > galanin (<i>GAL</i> , P22466) [1039]
Agonists	–	galanin(2-29) (rat/mouse) (p <i>K</i> _i 7.2–8.7) [1457, 1982, 1983, 1984] – Rat	–
Selective agonists	–	[D-Trp ²]galanin-(1-29) (p <i>K</i> _i 8.1) [1765] – Rat	–
Selective antagonists	2,3-dihydro-1,4-dithiin-1,1,4,4-tetroxide (p <i>C</i> ₅₀ 5.6) [1688]	M871 (p <i>K</i> _i 7.9) [1780]	SNAP 398299 (p <i>K</i> _i 8.3) [987, 988, 1833], SNAP 37889 (p <i>K</i> _i 7.8–7.8) [987, 988, 1833]
Selective allosteric modulators	–	CYM2503 (Positive) (p <i>E</i> <i>C</i> ₅₀ 9.2) [1147] – Rat	–
Labelled ligands	[¹²⁵ I][Tyr ²⁶]galanin (human) (Agonist) (p <i>K</i> _d 10.3) [525], [¹²⁵ I][Tyr ²⁶]galanin (human) (Agonist) (p <i>K</i> _d 7.8) [525]	[¹²⁵ I][Tyr ²⁶]galanin (human) (Agonist) (p <i>K</i> _d 9.2) [1983] – Rat	[¹²⁵ I][Tyr ²⁶]galanin (pig) (Agonist) (p <i>K</i> _d 8.6) [187, 1766]
Comments	–	The CYM2503 PAM potentiates the anticonvulsant activity of endogenous galanin in mouse seizure models [1147].	–

Comments: **galanin-(1-11)** is a high-affinity agonist at *GAL*₁/*GAL*₂ (p*K*_i 9), and **galanin(2-11)** is selective for *GAL*₂ and *GAL*₃ compared with *GAL*₁ [1146]. [¹²⁵I][Tyr²⁶]galanin binds to all three subtypes with *K*_d values generally reported to range from 0.05 to 1 nM, depending on the assay conditions used [525, 1752, 1765, 1766, 1983]. Porcine galanin-(3-29) does not bind to cloned *GAL*₁, *GAL*₂ or *GAL*₃ receptors, but a receptor that is functionally activated by porcine galanin-(3-29) has been reported in pituitary and

gastric smooth muscle cells [658, 2054]. Additional galanin receptor subtypes are also suggested from studies with chimeric peptides (e.g. M15, M35 and M40), which act as antagonists in functional assays in the cardiovascular system [1924], spinal cord [2024], locus coeruleus, hippocampus [100] and hypothalamus [101, 1078], but exhibit agonist activity at some peripheral sites [101, 658]. The chimeric peptides M15, M32, M35, M40 and C7 are agonists at *GAL*₁ receptors expressed endogenously in Bowes human

melanoma cells [1433], and at heterologously expressed recombinant *GAL*₁, *GAL*₂ and *GAL*₃ receptors [525, 1765, 1766]. Recent studies have described the synthesis of a series of novel, systemically-active, galanin analogues, with modest preferential binding at the *GAL*₂ receptor. Specific chemical modifications to the galanin backbone increased brain levels of these peptides after *i.v.* injection and several of these peptides exerted a potent antidepressant-like effect in mouse models of depression [1623].

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Ghrelin receptor

G protein-coupled receptors → Ghrelin receptor

Overview: The ghrelin receptor (**nomenclature as agreed by the NC-IUPHAR Subcommittee for the Ghrelin receptor [397]**) is activated by a 28 amino-acid peptide originally isolated from rat stomach, where it is cleaved from a 117 amino-acid precursor (*GHRL*, *Q9UBU3*). The human gene encoding the precursor peptide has 83% sequence homology to rat prepro-ghrelin, although the mature peptides from rat and human differ by only two amino acids [1222]. Alternative splicing results in the formation of a second

peptide, [*des-Gln*¹⁴]ghrelin (*GHRL*, *Q9UBU3*) with equipotent biological activity [783]. A unique post-translational modification (octanoylation of Ser³, catalysed by ghrelin *O*-acyltransferase (*MBOAT4*, *Q96T53*) [2082] occurs in both peptides, essential for full activity in binding to ghrelin receptors in the hypothalamus and pituitary, and for the release of growth hormone from the pituitary [983]. Structure activity studies showed the first five N-terminal amino acids to

be the minimum required for binding [116], and receptor mutagenesis has indicated overlap of the ghrelin binding site with those for small molecule agonists and allosteric modulators of ghrelin (*GHRL*, *Q9UBU3*) function [776]. In cell systems, the ghrelin receptor is constitutively active [777], but this is abolished by a naturally occurring mutation (A204E) that results in decreased cell surface receptor expression and is associated with familial short stature [1458].

Nomenclature	ghrelin receptor
HGNC, UniProt	<i>GHSR</i> , Q92847
Rank order of potency	ghrelin (<i>GHRL</i> , <i>Q9UBU3</i>) = [<i>des-Gln</i> ¹⁴]ghrelin (<i>GHRL</i> , <i>Q9UBU3</i>) [115, 1222]
Selective antagonists	GSK1614343 (pIC ₅₀ 8.4) [1624], GSK1614343 (pK _B 8) [1487] – Rat
Labelled ligands	[¹²⁵ I][His ⁹]ghrelin (human) (Agonist) (pK _d 9.4) [912], [¹²⁵ I][Tyr ⁴]ghrelin (human) (Agonist) (pK _d 9.4) [1339]

Comments: [*des-octanoyl*]ghrelin (*GHRL*, *Q9UBU3*) has been shown to bind (as [¹²⁵I]Tyr⁴-*des-octanoyl*-ghrelin) and have effects in the cardiovascular system [115], which raises the possible existence of different receptor subtypes in peripheral tissues and the central nervous system. A potent inverse agonist has been identified

([D-Arg¹,D-Phe⁵,D-Trp^{7,9},Leu¹¹]substance P, pD₂ 8.3; [774]). Ulimorelin, described as a ghrelin receptor agonist (pK_i 7.8 and pD₂ 7.5 at human recombinant ghrelin receptors), has been shown to stimulate ghrelin receptor mediated food intake and gastric emptying but not elicit release of growth hormone, or modify ghrelin stimulated growth hormone release, thus pharmacologically discrim-

inating the orexigenic and gastrointestinal actions of ghrelin (*GHRL*, *Q9UBU3*) from the release of growth hormone [538]. A number of selective antagonists have been reported, including peptidomimetic [1338] and non-peptide small molecules including GSK1614343 [1487, 1624].

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Glucagon receptor family

G protein-coupled receptors → Glucagon receptor family

Overview: The glucagon family of receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on the Glucagon receptor family [1234]**) are activated by the endogenous peptide (27-44 aa) hormones **glucagon** (GCG, P01275), **glucagon-like peptide 1** (GCG, P01275), **glucagon-like peptide 2** (GCG, P01275), glucose-dependent insulinotropic polypeptide (also known as **gastric inhibitory polypeptide** (GIP, P09681)), **GHRH** (GHRH, P01286) and **secretin** (SCT, P09683). One common precursor (GCG) generates **glucagon** (GCG, P01275), **glucagon-like peptide 1** (GCG, P01275) and **glucagon-like peptide 2** (GCG, P01275) peptides [827].

Nomenclature	GHRH receptor	GIP receptor	GLP-1 receptor
HGNC, UniProt	<i>GHRHR</i> , Q02643	<i>GIPR</i> , P48546	<i>GLP1R</i> , P43220
Endogenous agonists	–	gastric inhibitory polypeptide (GIP, P09681) (Selective) (pK _d 8.7) [1961]	glucagon-like peptide 1-(7-36) amide (GCG, P01275) (Selective) (pK _i 9.2) [885], glucagon-like peptide 1-(7-37) (GCG, P01275) (Selective) [425]
Agonists	Jl-38 [255], sermorelin	–	liraglutide (pEC ₅₀ 10.2) [972], lixisenatide (pK _i 8.9) [2010], WB4-24 (pA ₂ 4.9) [502]
Selective agonists	BIM28011 [379], tesamorelin	–	exendin-4 (pIC ₅₀ 9.2) [1290], exendin-4 (pK _i 8.7–9) [885], exendin-3 (P20394) [1564]
Selective antagonists	JV-1-36 (pK _i 10.1–10.4) [1662, 1947, 1948] – Rat, JV-1-38 (pK _i 10.1) [1662, 1947, 1948] – Rat	[Pro³]GIP [584] – Mouse	exendin-(9-39) (pK _i 8.1) [885], GLP-1-(9-36) (pIC ₅₀ 6.9) [1314] – Rat, T-0632 (pIC ₅₀ 4.7) [1883]
Labelled ligands	[¹²⁵I]GHRH (human) (Agonist) (pK _d 7.6) [192] – Rat	[¹²⁵I]GIP (human) (Agonist) (pK _d 8.6) [562] – Rat	[¹²⁵I]GLP-1-(7-36)-amide (Agonist) (pK _d 9.3) [885], [¹²⁵I]exendin-(9-39) (Antagonist) (pK _d 8.3) [885], [¹²⁵I]GLP-1-(7-37) (human) (Agonist)

Nomenclature	GLP-2 receptor	glucagon receptor	secretin receptor
HGNC, UniProt	GLP2R, O95838	GCCR, P47871	SCTR, P47872
Endogenous agonists	glucagon-like peptide 2 (GCG, P01275) (Selective) (pIC ₅₀ 8.5) [1880]	glucagon (GCG, P01275) (Selective) (pEC ₅₀ 9) [1515]	secretin (SCT, P09683) (Selective) (pEC ₅₀ 9.7) [329]
Agonists	teduglutide [1248]	–	–
Selective antagonists	–	L-168,049 (pIC ₅₀ 8.4) [269], des-His ¹ -[Glu ⁹]glucagon-NH ₂ (pA ₂ 7.2) [1928 , 1929] – Rat, NNC 92-1687 (pK _i 5) [1170], BAY27-9955 [1496]	[(CH ₂ NH) ^{4,5}]secretin (pK _i 5.3) [668]
Labelled ligands	–	[¹²⁵ I]glucagon (human, mouse, rat) (Agonist)	[¹²⁵ I](Tyr ¹⁰)secretin-27 (rat) (Agonist) [1925] – Rat

Comments: The glucagon receptor has been reported to interact with receptor activity modifying proteins (RAMPs), specifically [RAMP2](#), in heterologous expression systems [[333](#)], although the physiological significance of this has yet to be established.

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Glycoprotein hormone receptors

G protein-coupled receptors → Glycoprotein hormone receptors

Overview: Glycoprotein hormone receptors (**provisional nomenclature** [530]) are activated by a non-covalent heterodimeric glycoprotein made up of a common α chain (glycoprotein hormone common alpha subunit (CGA, P01215) CGA,

P01215), with a unique β chain that confers the biological specificity to FSH (CGA FSHB, P01215 P01225), LH (CGA LHB, P01215 P01229), hCG (CGA CGB, P01215 P01233) or TSH (CGA TSHB, P01215 P01222). There is binding cross-reactivity across the en-

dogenous agonists for each of the glycoprotein hormone receptors. The deglycosylated hormones appear to exhibit reduced efficacy at these receptors [1626].

Nomenclature	FSH receptor	LH receptor	TSH receptor
HGNC, UniProt	FSHR, P23945	LHCGR, P22888	TSHR, P16473
Endogenous agonists	–	hCG (CGA CGB, P01215 P01233) (Selective) (pK_d 9.9–11.8) [864, 1353], LH (CGA LHB, P01215 P01229) (Selective) ($pI_{C_{50}}$ 9.9–10.9) [864, 1353]	–
Antagonists	FSH deglycosylated α/β (pK_d 10) [527, 921]	–	–
Labelled ligands	[¹²⁵ I]FSH (human) (Agonist)	[¹²⁵ I]LH (Agonist), [¹²⁵ I]chorionic gonadotropin (human) (Agonist)	[¹²⁵ I]TSH (human) (Agonist)
Comments	Animal follitropins are less potent than the human hormone as agonists at the human FSH receptor. Gain- and loss-of-function mutations of the FSH receptor are associated with human reproductive disorders [19, 109, 650, 1900]. The rat FSH receptor also stimulates phosphoinositide turnover through an unidentified G protein [1547].	Loss-of-function mutations of the LH receptor are associated with Leydig cell hypoplasia and gain-of-function mutations are associated with male-limited gonadotropin-independent precocious puberty (<i>e.g.</i> [1044, 1720]) and Leydig cell tumours [1126].	Autoimmune antibodies that act as agonists of the TSH receptor are found in patients with Graves' disease (<i>e.g.</i> [1558]). Mutants of the TSH receptor exhibiting constitutive activity underlie hyperfunctioning thyroid adenomas [1464] and congenital hyperthyroidism [993]. TSH receptor loss-of-function mutations are associated with TSH resistance [1824].

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Gonadotrophin-releasing hormone receptors

G protein-coupled receptors → Gonadotrophin-releasing hormone receptors

Overview: GnRH₁ and GnRH₂ receptors (**provisonal nomenclature** [530], also called Type I and Type II GnRH receptor, respectively [1284]) have been cloned from numerous species, most of which express two or three types of GnRH receptor [1283, 1284, 1741]. GnRH I (*GNRH1*, P01148) (p-Glu-His-Trp-Ser-Tyr-Gly-Leu-Arg-Pro-Gly-NH₂) is a hypothalamic decapeptide also known as luteinizing hormone-releasing hormone, gonadoliberin, luliberin, gonadorelin or simply as GnRH. It is a member of a family of similar peptides found in many species [1283, 1284, 1741] including GnRH II (*GNRH2*, O43555) (pGlu-His-Trp-Ser-His-Gly-Trp-Tyr-Pro-Gly-NH₂ (which is also known as chicken GnRH-II)). Receptors

for three forms of GnRH exist in some species but only GnRH I and GnRH II and their cognate receptors have been found in mammals [1283, 1284, 1741]. GnRH₁ receptors are expressed primarily by pituitary gonadotrophs, and mediate central control of mammalian reproduction. They are selectively activated by GnRH I and all lack the COOH-terminal tails found in other GPCRs. GnRH₂ receptors do have COOH-terminal tails and (where tested) are selective for GnRH II over GnRH I. GnRH₂ receptors are expressed by some primates but are thought not to be expressed by humans because the human *GNRHR2* gene contains a frame shift and an internal stop codon [1325]. An alternative phylogenetic classification divides GnRH receptors into

three classes and includes both GnRH I-selective mammalian and GnRH II-selective non-mammalian GnRH receptors as GnRH₁ receptors [1284]. A more recent phylogenetic classification groups vertebrate GnRH receptors into five subfamilies [2028] and highlights examples of gene loss through evolution, with humans notably retaining only one ancient gene. Although thousands of peptide analogues of GnRH I have been synthesized and several (agonists and antagonists) are used therapeutically [934], the potency of most of these peptides at GnRH₂ receptors is unknown.

Nomenclature	GnRH ₁ receptor	GnRH ₂ receptor
HGNC, UniProt	<i>GNRHR</i> , P30968	<i>GNRHR2</i> , Q96P88
Rank order of potency	GnRH I (<i>GNRH1</i> , P01148) > GnRH II (<i>GNRH2</i> , O43555) [1284]	GnRH II (<i>GNRH2</i> , O43555) > GnRH I (<i>GNRH1</i> , P01148) (Monkey) [1282]
Endogenous agonists	–	GnRH II (<i>GNRH2</i> , O43555) (pIC ₅₀ 9) [1282] – Monkey, GnRH I (<i>GNRH1</i> , P01148) (pIC ₅₀ 7.4) [1282] – Monkey
Selective agonists	triptorelin (pK _i 9.3–9.5) [112], leuprolide (pK _i 8.5–9.1) [1807], busarelin, goserelin, histrelin, nafarelin	–
Antagonists	itrelax (pK _i 9.5) [1591]	–
Selective antagonists	cetorelix (pK _i 9.3–10) [113, 114, 1807], abarelix (pK _i 9.1–9.5) [1807], degarelix (pK _i 8.8) [1938], ganirelix	trptorelix-1 [1183] – Monkey
Labelled ligands	[¹²⁵ I]busarelin (Agonist) (pK _d 7.4) [1024] – Rat, [¹²⁵ I]GnRH I (human, mouse, rat) (Agonist)	–
Comments	–	Probable transcribed pseudogene in man [1284].

Comments: GnRH₁ and GnRH₂ receptors couple primarily to G_{q/11} [653] but coupling to G_s and G_i is evident in some systems [1009, 1024]. GnRH₂ receptors may also mediate (heterotrimeric) G protein-independent signalling to protein kinases [276]. There is increasing evidence for expression of GnRH receptors on hormone-dependent cancer cells where they can exert antiproliferative and/or

proapoptotic effects and mediate effects of cytotoxins conjugated to GnRH analogues [309, 706, 1108, 1661]. In some human cancer cell models GnRH II (*GNRH2*, O43555) is more potent than GnRH I (*GNRH1*, P01148), implying mediation by GnRH₂ receptors [657]. However, GnRH₂ receptors that are expressed by some primates are probably not expressed in humans because the human *GNRHR2* gene contains a frame shift and internal stop codon [1325]. The

possibility remains that this gene generates GnRH₂ receptor-related proteins (other than the full-length receptor) that mediate responses to GnRH II (*GNRH2*, O43555) (see [1372]). Alternatively, there is evidence for multiple active GnRH receptor conformations [276, 277, 516, 1231, 1284] raising the possibility that GnRH₁ receptor-mediated proliferation inhibition in hormone-dependent cancer cells is dependent upon different conformations (with different ligand

specificity and ligand biased signalling) than effects on G_{q/11} in pituitary cells [277, 1231]. Loss-of-function mutations in the GnRH₁ receptor and deficiency of GnRH1 (*GNRH1*, P01148) are associated with hypogonadotropic hypogonadism although some ‘loss of

function’ mutations may actually prevent trafficking of ‘functional’ GnRH₁ receptors to the cell surface, as evidenced by recovery of function by nonpeptide antagonists [1061]. Human GnRH₁ receptors appear to be poorly expressed at the cell surface because of failure to meet structural quality control criteria for endoplasmic reticu-

lum exit [517, 1061]. This may increase susceptibility to point mutations that further impair trafficking and also increase effects of non-peptide antagonists on GnRH₁ receptor trafficking to the plasma membrane [517, 1061]. GnRH receptor signalling may be dependent upon receptor oligomerisation [363, 1007].

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GPR18, GPR55 and GPR119

G protein-coupled receptors → GPR18, GPR55 and GPR119

Overview: GPR18, GPR55 and GPR119 (**provisional nomenclature**), although showing little structural similarity to CB₁ and CB₂ cannabinoid receptors, respond to endogenous agents analogous to the endogenous cannabinoid ligands, as well as some natural/synthetic cannabinoid receptor ligands [1494]. Although there are multiple reports to indicate that GPR18, GPR55 and GPR119 can be activated *in vitro* by N-arachidonoylglycine, lysophosphatidylinositol and N-oleoylethanolamide, respectively, there is a lack of evidence for activation by these lipid messengers *in vivo*. As such, therefore, these receptors retain their orphan status.

Nomenclature	<i>GPR18</i>	<i>GPR55</i>	<i>GPR119</i>
HGNC, UniProt	<i>GPR18</i> , Q14330	<i>GPR55</i> , Q9Y2T6	<i>GPR119</i> , Q8TDV5
Rank order of potency	–	–	N-oleoylethanolamide, N-palmitoylethanolamine > SEA (anandamide is ineffective) [1452]
Endogenous agonists	N-arachidonoylglycine [980]	lysophosphatidylinositol (pEC ₅₀ 5.5–7.3) [738, 1435, 1785], 2-arachidonoylglycerolphosphoinositol (Selective) [1437]	N-oleoylethanolamide (pEC ₅₀ 5.4–6.3) [338, 1452, 1785], N-palmitoylethanolamine, SEA
Selective agonists	–	AM251 (pEC ₅₀ 5–7.4) [738, 905, 1620]	AS1269574 (pEC ₅₀ 5.6) [2100], PSN632408 (pEC ₅₀ 5.3) [1452], PSN375963 (pEC ₅₀ 5.1) [1452]
Selective antagonists	–	CID16020046 (apparent pA ₂) (pA ₂ 7.3) [906]	–
Comments	The pairing of N-arachidonoylglycine with GPR18 was not replicated in two studies based on arrestin assays [1785, 2093]. See [396] for discussion.	See reviews [396] and [1732].	In addition to those shown above, further small molecule agonists have been reported [687].

Comments: GPR18 failed to respond to a variety of lipid-derived agents in an *in vitro* screen [2093], but has been reported to be activated by Δ^9 -tetrahydrocannabinol [1246]. GPR55 responds to AM251 and rimonabant at micromolar concentrations, compared to their nanomolar affinity as CB₁ re-

ceptor antagonists/inverse agonists [1494]. It has been reported that lysophosphatidylinositol acts at other sites in addition to GPR55 [2075]. N-Arachidonoylserine has been suggested to act as a low efficacy agonist/antagonist at GPR18 *in vitro* [1244]. It has also been suggested oleoyl-lysophosphatidylcholine acts, at

least in part, through GPR119 [1400]. Although PSN375963 and PSN632408 produce GPR119-dependent responses in heterologous expression systems, comparison with N-oleylethanolamide-mediated responses suggests additional mechanisms of action [1400].

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Histamine receptors

G protein-coupled receptors → Histamine receptors

Overview: Histamine receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Histamine Receptors [754, 1459]) are activated by the endogenous ligand histamine. Marked species differences exist between histamine receptor orthologues [754].

Nomenclature	H ₁ receptor	H ₂ receptor	H ₃ receptor	H ₄ receptor
HGNC, UniProt	<i>HRH1</i> , P35367	<i>HRH2</i> , P25021	<i>HRH3</i> , Q9Y5N1	<i>HRH4</i> , Q9H3N8
Selective agonists	methylhistaprodifen (pK _i 6.4) [1695], histaprodifen (pK _i 5.7) [1107]	amthamine (pEC ₅₀ 6.4) [1003]	immethridine (pK _i 9.1) [963], methimepip (pK _i 9) [962], MK-0249 (Inverse agonist) (pK _i 8.8) [1354]	clobenpropit (Partial agonist) (pK _i 7.4–8.3) [490, 1107, 1122, 1123, 1335], 4-methylhistamine (pK _i 7.3–8.2) [586, 1107], VUF 8430 (pK _i 7.5) [1106]
Antagonists	ciproheptadine (pK _i 10.2) [1298], promethazine (pK _i 9.6) [601], pyrilamine (Inverse agonist) (pK _i 8.7–9) [184, 1563], hydroxyzine (pK _i 8.7) [608], ketotifen (pK _i 8.6) [1014], cetirizine (Inverse agonist) (pK _i 8.2) [1298], diphenhydramine (pK _i 7.9) [184]	–	iodophenpropit (pK _i 8.2–8.7) [2022, 2051], thioperamide (pK _i 7.1–7.7) [355, 489, 490, 1104, 1145, 2022, 2051]	–
Selective antagonists	clemastine (pK _i 10.3) [68], desloratadine (pK _i 9) [1090], triprolidine (pK _i 8.5–9) [184, 1298], azelastine (pK _i 8.9) [1535], astemizole (pK _i 8.5) [1480], cyclizine (pK _i 8.4) [68], chlorpheniramine (pK _i 8.1) [1535], fexofenadine (pK _i 7.6) [64], loratadine (pK _i 7.4) [850], terfenadine (pK _i 7.4) [64], tripeleminamine (pK _i 7.4) [635]	tiotidine (pK _i 7.5) [145] – Rat, ranitidine (pK _i 7.1) [1086], cimetidine (pK _i 6.8) [263]	clobenpropit (pK _i 8.4–9.4) [355, 490, 1104, 1122, 1145, 2022, 2051], A331440 (pK _i 8.5) [688]	JNJ 7777120 (pK _i 7.8–8.3) [1107, 1771, 1881]
Labelled ligands	[³ H]pyrilamine (Antagonist, Inverse agonist) (pK _d 8.4–9.1) [403, 1298, 1675, 1695], [¹¹ C]doxepin (Antagonist) (pK _d 9) [830], [¹¹ C]pyrilamine (Antagonist, Inverse agonist)	[¹²⁵ I]iodoaminopotentidine (Antagonist) (pK _d 8.7) [1029] – Rat, [³ H]tiotidine (Antagonist) (pK _d 7.7–8.7) [1310]	[¹²³ I]iodoproxyfan (Antagonist) (pK _d 10.2) [1104], [¹²⁵ I]iodophenpropit (Antagonist) (pK _d 9.2) [849] – Rat, [³ H](R)-α-methylhistamine (Agonist) (pK _d 9.2) [1122], N-[³ H]α-methylhistamine (Agonist) (pK _d 9) [301] – Mouse	[³ H]JNJ 7777120 (Antagonist) (pK _d 8.4) [1881]

Comments: histaprodifen and methylhistaprodifen are reduced efficacy agonists. The H₄ receptor appears to exhibit broadly similar pharmacology to the H₃ receptor for imidazole-containing ligands, although (R)-α-methylhistamine and N-α-methylhistamine are

less potent, while clobenpropit acts as a reduced efficacy agonist at the H₄ receptor and an antagonist at the H₃ receptor [1122, 1360, 1390, 1422, 2132]. Moreover, 4-methylhistamine is identified as a high affinity, full agonist for the human H₄ receptor [1107].

[³H]histamine has been used to label the H₄ receptor in heterologous expression systems.

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Hydroxycarboxylic acid receptors

G protein-coupled receptors → Hydroxycarboxylic acid receptors

Overview: The hydroxycarboxylic acid family of receptors (ENFSM00500000271913, nomenclature as agreed by the **NC-IUPHAR Subcommittee on Hydroxycarboxylic acid receptors** [396, 1424]) respond to organic acids, including the endogenous hydroxy carboxylic acids 3-hydroxy butyric acid and L-lactic acid, as well as the lipid lowering agents nicotinic acid (niacin), acipimox and acifran [1774, 1913, 2036]. These receptors were provisionally described as nicotinic acid receptors, although nicotinic acid shows submicromolar potency at HCA₂ receptors only and is unlikely to be the natural ligand [1913, 2036].

	HCA ₁ receptor	HCA ₂ receptor	HCA ₃ receptor
Nomenclature	HCA ₁ receptor	HCA ₂ receptor	HCA ₃ receptor
HGNC, UniProt	HCAR1, Q9BXC0	HCAR2, Q8TDS4	HCAR3, P49019
Endogenous agonists	L-lactic acid (Selective) (pEC ₅₀ 1.3–2.9) [14, 256, 1124, 1785]	β-D-hydroxybutyric acid (pEC ₅₀ 3.1) [1838]	3-hydroxyoctanoic acid (pEC ₅₀ 5.1) [13]
Agonists	compound 2 [PMID: 24486398] (pEC ₅₀ 7.2) [1630], 3,5-dihydroxybenzoic acid (pEC ₅₀ 3.7) [1121]	SCH 900271 (pEC ₅₀ 8.7) [1454], GSK256073 (pEC ₅₀ 7.5) [1790]	–
Selective agonists	–	MK 6892 (pEC ₅₀ 7.8) [1719], MK 1903 (pEC ₅₀ 7.6) [163], nicotinic acid (pEC ₅₀ 6–7.2) [1774, 1913, 2036], acipimox (pEC ₅₀ 5.2–5.6) [1774, 2036], monomethylfumarate (pEC ₅₀ 5) [1859]	compound 6o [PMID: 19524438] (pEC ₅₀ 8.5) [1751], IBC 293 (pEC ₅₀ 6.4) [1697]
Labelled ligands	–	[³ H]nicotinic acid (Agonist) (pK _d 7–7.3) [1774, 1913, 2036]	–

Comments: Further closely-related GPCRs include the 5-oxoeicosanoid receptor (OXER1, Q8TDS5) and GPR31 (O00270).

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Kisspeptin receptor

G protein-coupled receptors → Kisspeptin receptor

Overview: The kisspeptin receptor (**nomenclature as agreed by the NC-IUPHAR Subcommittee on the kisspeptin receptor [958]**), like neuropeptide FF (NPFF), prolactin-releasing peptide (PrP) and QRFP receptors (provisional nomenclature) responds to endogenous peptides with an arginine-phenylalanine-amide (RFamide) motif. **Kisspeptin-54 (KISS1, Q15726)** (KP54, originally named metastin), **kisspeptin-13 (KISS1, Q15726)** (KP13) and **kisspeptin-10 (KISS1)** (KP10) are biologically-active peptides cleaved from the **KISS1 (Q15726)** gene product.

Nomenclature	kisspeptin receptor
HGNC, UniProt	KISS1R , Q969F8
Endogenous agonists	kisspeptin-10 (KISS1) (Selective) (pK _i 8.6–10.4) [996 , 1434], kisspeptin-54 (KISS1, Q15726) (Selective) (pK _i 8.8–9.5) [996 , 1434], kisspeptin-14 (KISS1, Q15726) (pK _i 8.8) [996], kisspeptin-13 (KISS1, Q15726) (Selective) (pK _i 8.4) [996]
Selective agonists	4-fluorobenzoyl-FGLRW-NH ₂ (pEC ₅₀ 9.2) [1894], [dY] ¹ KP-10 (pIC ₅₀ 8.4) [385] – Mouse
Selective antagonists	peptide 234 [1600]
Labelled ligands	[¹²⁵ I]Tyr ⁴⁵ -kisspeptin-15 (Agonist) (pK _d 10) [1434], [¹²⁵ I]kisspeptin-13 (human) (Agonist) (pK _d 9.7) [1252], [¹²⁵ I]kisspeptin-10 (human) (Agonist) (pK _d 8.7) [996], [¹²⁵ I]kisspeptin-14 (human) (Agonist) [1252]

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Searchable database: <http://www.guidetopharmacology.org/index.jsp>

Full Contents of ConciseGuide: <http://onlinelibrary.wiley.com/doi/10.1111/bph.13348/full>

Kisspeptin receptor 5815

Leukotriene receptors

G protein-coupled receptors → Leukotriene receptors

Overview: Leukotriene receptors (**nomenclature as agreed by the NC-IUPHAR subcommittee on Leukotriene Receptors [249, 250]**) is activated by the endogenous ligands leukotrienes (LT), synthesized from lipoxygenase metabolism of arachidonic acid.

The human BLT₁ receptor is the high affinity LTB₄ receptor whereas the BLT₂ receptor in addition to being a low-affinity LTB₄ receptor also binds several other lipoxygenase-products, such as 12S-HETE, 12S-HPETE, 15S-HETE, and the thromboxane synthase

product 12-hydroxyheptadecatrienoic acid. The BLT receptors mediate chemotaxis and immunomodulation in several leukocyte populations and are in addition expressed on non-myeloid cells, such as vascular smooth muscle and endothelial cells. In addition to BLT receptors, LTB₄ has been reported to bind to the peroxisome proliferator activated receptor (PPAR) α [1112] and the vanilloid TRPV1 ligand-gated nonselective cation channel [1245].

The receptors for the cysteinyl-leukotrienes (*i.e.* LTC₄, LTD₄ and LTE₄) are termed CysLT₁ and CysLT₂ and exhibit distinct expression

patterns in human tissues, mediating for example smooth muscle cell contraction, regulation of vascular permeability, and leukocyte activation. There is also evidence in the literature for additional CysLT receptor subtypes, derived from functional in vitro studies, radioligand binding and in mice lacking both CysLT₁ and CysLT₂ receptors [250]. Cysteinyl-leukotrienes have also been suggested to signal through the P2Y₁₂ receptor [542, 1407, 1466], GPR17 [344] and GPR99 [900].

	BLT ₁ receptor	BLT ₂ receptor	CysLT ₁ receptor	CysLT ₂ receptor
Nomenclature	BLT ₁ receptor	BLT ₂ receptor	CysLT ₁ receptor	CysLT ₂ receptor
HGNC, UniProt	LTBR, Q15722	LTBR2, Q9NPC1	CYSLTR1, Q9Y271	CYSLTR2, Q9NS75
Rank order of potency	LTB ₄ > 20-hydroxy-LTB ₄ >> 12R-HETE [2096]	12-hydroxyheptadecatrienoic acid > LTB ₄ > 12S-HETE = 12S-HPETE > 15S-HETE > 12R-HETE > 20-hydroxy-LTB ₄ [1442, 2096]	LTD ₄ > LTC ₄ > LTE ₄ [1157, 1643]	LTC ₄ ≥ LTD ₄ >> LTE ₄ [733, 1411, 1847]
Endogenous agonists	–	12S-HETE (Partial agonist) (pEC ₅₀ < 7.5) [2096]	–	–
Antagonists	–	–	ICI198615 (- 8.4–8.6)	BAYu9773 (against LTC ₄ and LTD ₄ induced contraction in smooth muscle preparation) (pA ₂ 6.8–7.7) [1912] – Rat
Selective antagonists	BIIL 260 (pK _i 8.8) [152, 457], CP105696 (pIC ₅₀ 8.1) [1735], U75302 (pK _i 6.4) [172]	LY255283 (pIC ₅₀ 6–7.1) [744, 2096]	zafirlukast (against [³ H]LTD ₄ in COS-7 or HEK-293 cells) (pIC ₅₀ 8.6–8.7) [1157, 1643], SR2640 (pK _i 8.7), montelukast (against [³ H]LTD ₄ in COS-7 or HEK-293 cells) (pIC ₅₀ 8.3–8.6) [1157, 1643], sulukast (pK _i 8.3), pobilukast (against [³ H]LTD ₄ in HEK-293) (pIC ₅₀ 7.5) [1643]	BayCysLT ₂ (against 30–300nM LTD ₄ β-arrestin assay in C2C12 myofibroblasts) (pIC ₅₀ 6.6–7.3) [1391]
Labelled ligands	[³ H]LTB ₄ (Agonist) (pK _d 9.8) [2095], [³ H]CGS23131 (Antagonist) (pK _d 7.9) [836]	[³ H]LTB ₄ (pK _d 7.6–9.7)	[³ H]LTD ₄ (Agonist) (pK _d 8–10.7), [³ H]ICI-198615 (Antagonist) (pK _d 10.6) [1608]	[³ H]LTD ₄ (Agonist) (pK _d 7.3–9.4) [733]

Nomenclature	FPR2/ALX	OXE receptor
HGNC, UniProt	FPR2, P25090	OXER1, Q8TDSS
Rank order of potency	LXA ₄ =aspirin triggered lipoxin A4 =ATLa2>LTC ₄ =LTD ₄ >>15-deoxy-LXA ₄ >> fMet-Leu-Phe [352, 519, 521, 651, 1846]	5-oxo-EETE, 5-oxo-C20:3, 5-oxo-ODE > 5-oxo-15-HETE > 5S-HPETE > 5S-HETE [784, 877, 1472]
Endogenous agonists	LXA ₄ (Selective) (pEC ₅₀ ~12) [1006], resolvin D1 (Selective) (pEC ₅₀ ~11.9) [1006], aspirin triggered lipoxin A4 (Selective)	5-oxo-EETE (Selective) (pEC ₅₀ 8.3–8.5) [638, 1417, 1472, 1525, 1681]
Selective agonists	ATLa2 [662]	–
Endogenous antagonists	–	5-oxo-12-HETE (Selective) (pIC ₅₀ 6.3) [1524]
Antagonists	–	–
Selective antagonists	–	–
Labelled ligands	[³ H]LXA ₄ (Agonist) (pK _d 9.2–9.3) [519, 520]	[³ H]5-oxo-EETE (Agonist) (pK _d 8.4) [1417]
Comments	The agonist activity of the lipid mediators described has been questioned [697, 1513], which may derive from batch-to-batch differences, partial agonism or biased agonism. Recent results from Cooray <i>et al.</i> (2013) [365] have addressed this issue and the role of homodimers and heterodimers in the intracellular signaling .	–

Comments: The FPR2/ALX receptor (**nomenclature as agreed by the NC-IUPHAR subcommittee on Leukotriene and Lipoxin Receptors [250]**) is activated by the endogenous lipid-derived, anti-inflammatory ligands lipoxin A₄ (LXA₄) and 15-epi-LXA₄ (aspirin triggered lipoxin A4, ATL). The FPR2/ALX receptor also interacts with endogenous peptide and protein ligands, such as MHC binding peptide [315] as well as annexin I (ANXA1, P04083) (ANXA1) and its N-terminal peptides [365, 1491]. In addition, a

soluble hydrolytic product of protease action on the urokinase-type plasminogen activator receptor has been reported to activate the FPR2/ALX receptor [1572]. Furthermore, FPR2/ALX has been suggested to act as a receptor mediating the proinflammatory actions of the acute-phase reactant, serum amyloid A [1772, 1809]. A receptor selective for LXB₄ has been suggested from functional studies [53, 1168, 1596]. Note that the data for FPR2/ALX are also reproduced on the [Formylpeptide receptor pages](#).

Oxoecicosanoid receptors (**OXE, nomenclature agreed by the NC-IUPHAR subcommittee on Oxoecicosanoid Receptors [214]**) are activated by endogenous chemotactic eicosanoid ligands oxidised at the C-5 position, with 5-oxo-EETE the most potent agonist identified for this receptor. Initial characterization of the heterologously expressed OXE receptor suggested that polyunsaturated fatty acids, such as docosahexaenoic acid and EPA, acted as receptor antagonists [784]

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Lysophospholipid (LPA) receptors

G protein-coupled receptors → Lysophospholipid (LPA) receptors

Overview: Lysophosphatidic acid (LPA) receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Lysophospholipid Receptors** [396, 936]) are activated by the endogenous phospholipid metabolite LPA. The first receptor, LPA₁, was identified as *ventricular zone gene-1* (*vzg-1*), leading to deorphanisation of members of the endothelial differentiation gene (*edg*) family as other LPA receptors along with sphingosine 1-phosphate (S1P) receptors. Additional LPA receptor GPCRs were later identified. Gene names have been codified as *LPAR1*, etc. to reflect the receptor function of proteins. The crystal structure of LPA₁ was re-

cently solved and demonstrates ligand access characteristics that allows for extracellular LPA binding [331]; these studies have also implicated cross-talk with endocannabinoids *via* phosphorylated intermediates that can activate this receptor. The identified receptors can account for most, although not all, LPA-induced phenomena in the literature, indicating that a majority of LPA-dependent phenomena are receptor-mediated. Radioligand binding has been conducted in heterologous expression systems using [³H]LPA (*e.g.* [556]). In native systems, analysis of binding data is complicated by metabolism and high levels of nonspecific binding, and therefore the relation-

ship between recombinant and endogenously expressed receptors is unclear. Targeted deletion of LPA receptors has clarified signalling pathways and identified physiological and pathophysiological roles. Independent validation by multiple groups has been reported in the peer-reviewed literature for all six LPA receptors described in the tables, including further validation using a distinct read-out via a novel TGFα “shedding” assay [825]. LPA has also been described as an agonist at other orphan GPCRs (PSP24, GPR87 and GPR35), as well as at the nuclear hormone PPARγ receptors [1247, 1743], although the physiological significance of these observations remain unclear.

Nomenclature	LPA ₁ receptor	LPA ₂ receptor	LPA ₃ receptor
HGNC, UniProt	<i>LPAR1</i> , Q92633	<i>LPAR2</i> , Q9HBW0	<i>LPAR3</i> , Q9UBY5
Selective agonists	–	dodecylphosphate (pEC ₅₀ 6.2) [1958], decyl dihydrogen phosphate (pEC ₅₀ 5.4) [1958], GRI977143 (pEC ₅₀ 4.5) [959]	OMPT (pEC ₅₀ 7.2) [709]
Selective antagonists	AM966 (pIC ₅₀ 6.7–7.8) [1832]	–	dioctanoylglycerol pyrophosphate (pK _i 5.5–7) [522, 1432]
Comments	–	Virtual screening experiments have shown H2L5186303 to be a potent antagonist of LPA ₂ [510]. dodecylphosphate is also an antagonist at LPA ₃ receptors [1958].	–

Nomenclature	LPA ₄ receptor	LPA ₅ receptor	LPA ₆ receptor
HGNC, UniProt	<i>LPAR4</i> , Q99677	<i>LPAR5</i> , Q9H1C0	<i>LPAR6</i> , P43657

Comments: Ki16425 [1432], VPC12249 [735] and VPC32179 [729] have antagonist activity at LPA₁ and LPA₃ receptors. There is growing evidence for *in vivo* efficacy of these chemical antagonists in several disorders, including fetal hydrocephalus [2107], lung fibrosis [1429], and systemic sclerosis [1429].

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Lysophospholipid (S1P) receptors

G protein-coupled receptors → Lysophospholipid (S1P) receptors

Overview: Sphingosine 1-phosphate (S1P) receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Lysophospholipid receptors [936]**) are activated by the endogenous lipid sphingosine 1-phosphate (S1P) and with lower apparent affinity, sphingosylphosphorylcholine (SPC). Originally cloned as orphan members of the endothelial differentiation gene (*edg*) family, deorphanisation as lysophospholipid receptors for S1P was based on sequence homology to LPA receptors. Current gene names have been codified as S1PR1, etc. to reflect the receptor function of these proteins. Most cellular phenomena ascribed to S1P can

be explained by receptor-mediated mechanisms; S1P has also been described to act at intracellular sites [1841], and awaits precise definition. Previously-proposed SPC (or lysophosphosphatidylcholine) receptors- G2A, TDAG8, OGR1 and GPR4 - continue to lack confirmation of these roles [396]. The relationship between recombinant and endogenously expressed receptors is unclear. Radioligand binding has been conducted in heterologous expression systems using [³²P]S1P (*e.g* [1438]). In native systems, analysis of binding data is complicated by metabolism and high levels of nonspecific binding. Targeted deletion of several S1P receptors and key enzymes involved

in S1P biosynthesis or degradation has clarified signalling pathways and physiological roles. A crystal structure of an S1P₁-T4 fusion protein has been described [698].

The S1P receptor modulator, fingolimod (FTY720, Gilenya), has received world-wide approval as the first oral therapy for relapsing forms of multiple sclerosis. This drug has a novel mechanism of action involving modulation of S1P receptors in both the immune and nervous systems [325, 356, 654], although the precise nature of its interaction requires clarification.

Nomenclature	S1P ₁ receptor	S1P ₂ receptor	S1P ₃ receptor	S1P ₄ receptor	S1P ₅ receptor
HGNC, UniProt	S1PR1, P21453	S1PR2, O95136	S1PR3, Q99500	S1PR4, O95977	S1PR5, Q9H228
Rank order of potency	sphingosine 1-phosphate > dihydro sphingosine-1-phosphate > sphingosylphosphorylcholine [45, 1438]	sphingosine 1-phosphate > dihydro sphingosine-1-phosphate > sphingosylphosphorylcholine [45, 1438]	sphingosine 1-phosphate > dihydro sphingosine-1-phosphate > sphingosylphosphorylcholine [1438]	sphingosine 1-phosphate > dihydro sphingosine-1-phosphate > sphingosylphosphorylcholine [1936]	sphingosine 1-phosphate > dihydro sphingosine-1-phosphate > sphingosylphosphorylcholine [821]
Agonists	SEW2871 (pK _i 5.5–7.7) [1640]	–	–	–	–

(continued)	S1P ₁ receptor	S1P ₂ receptor	S1P ₃ receptor	S1P ₄ receptor	S1P ₅ receptor
Nomenclature	S1P ₁ receptor	S1P ₂ receptor	S1P ₃ receptor	S1P ₄ receptor	S1P ₅ receptor
Antagonists	VPC44116 (pK _i 8.5) [533], VPC23019 (pK _i 7.9) [400]	–	VPC44116 (pK _i 6.5) [533], VPC23019 (pK _i 5.9) [400]	–	–
Selective antagonists	W146 (pK _i 7.1) [1641]	JTE-013 (pIC ₅₀ 7.8) [1447]	–	–	–
Selective agonists	AUY954 (pEC ₅₀ 8.9) [1456]	–	–	–	–

Comments: The approved immunomodulator drug [fingolimod](#) can be phosphorylated *in vivo* [31] to generate a relatively potent agonist with activity at S1P₁, S1P₃, S1P₄ and S1P₅ receptors [215, 1198], although its biological activity appears to involve an element of functional antagonism [339, 356, 1405].

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Melanin-concentrating hormone receptors

G protein-coupled receptors → Melanin-concentrating hormone receptors

Overview: Melanin-concentrating hormone (MCH) receptors (**provisional nomenclature as recommended by NC-IUPHAR [530]**) are activated by an endogenous nonadecameric cyclic peptide identical in humans and rats (DFDMLRCMLGRVYRPCWQV) generated from a precursor (*PMCH*, P20382), which also produces [neuropeptide EI](#) (*PMCH*, P20382) and [neuropeptide GE](#) (*PMCH*, P20382).

Nomenclature	MCH ₁ receptor	MCH ₂ receptor
HGNC, UniProt	MCHR1 , Q99705	MCHR2 , Q969V1
Rank order of potency	melanin-concentrating hormone (<i>PMCH</i> , P20382) > MCH (salmon)	melanin-concentrating hormone (<i>PMCH</i> , P20382) = MCH (salmon) [753]
Selective antagonists	GW803430 (pIC ₅₀ 9.3) [745], SNAP-7941 (pA ₂ 9.2) [186], T-226296 (pIC ₅₀ 8.3) [1853], ATC0175 (pIC ₅₀ 7.9–8.1) [283]	–
Labelled ligands	[¹²⁵ I]S36057 (Antagonist) (pK _d 9.2–9.5) [66], [¹²⁵ I][Phe ¹³ , Tyr ¹⁹]MCH (Agonist) (pK _d 9.2) [242], [³ H]MCH (human, mouse, rat) (Agonist) [242]	–

Comments: The MCH₂ receptor appears to be a non-functional pseudogene in rodents [1857].

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Melanocortin receptors

G protein-coupled receptors → Melanocortin receptors

Overview: Melanocortin receptors (**provisional nomenclature as recommended by NC-IUPHAR [530]**) are activated by members of the melanocortin family (α -MSH (*POMC*, P01189), β -MSH (*POMC*, P01189) and γ -MSH (*POMC*, P01189) forms; $-\delta$ form is not found in mammals) and adrenocorticotrophin (*ACTH* (*POMC*, P01189)). Endogenous antagonists include *agouti* (*ASIP*, P42127) and *agouti-related protein* (*AGRP*, O00253).

Nomenclature	MC ₁ receptor	MC ₂ receptor	MC ₃ receptor	MC ₄ receptor	MC ₅ receptor
HGNC, UniProt	<i>MC1R</i> , Q01726	<i>MC2R</i> , Q01718	<i>MC3R</i> , P41968	<i>MC4R</i> , P32245	<i>MC5R</i> , P33032
Rank order of potency	α -MSH (<i>POMC</i> , P01189) > β -MSH (<i>POMC</i> , P01189) > ACTH (<i>POMC</i> , P01189), γ -MSH (<i>POMC</i> , P01189)	ACTH (<i>POMC</i> , P01189)	γ -MSH (<i>POMC</i> , P01189), β -MSH (<i>POMC</i> , P01189) > ACTH (<i>POMC</i> , P01189), α -MSH (<i>POMC</i> , P01189)	β -MSH (<i>POMC</i> , P01189) > α -MSH (<i>POMC</i> , P01189), ACTH (<i>POMC</i> , P01189) > γ -MSH (<i>POMC</i> , P01189)	α -MSH (<i>POMC</i> , P01189) > β -MSH (<i>POMC</i> , P01189) > ACTH (<i>POMC</i> , P01189) > γ -MSH (<i>POMC</i> , P01189)
Selective agonists	–	corticotropin zinc hydroxide	[D-Trp ⁸] γ -MSH (pIC ₅₀ 8.2) [645]	THIQ (pIC ₅₀ 8.9) [1690]	–
Antagonists	–	–	PG-106 (pIC ₅₀ 6.7) [646]	–	–
Selective antagonists	–	–	–	MBP10 (pIC ₅₀ 10) [117], HS014 (pK _i 8.5) [1667]	–
Labelled ligands	[¹²⁵ I]NDP-MSH (Agonist) (pK _d 9.5) [991]	[¹²⁵ I]ACTH-(1-24) (Agonist)	[¹²⁵ I]NDP-MSH (Agonist) (pK _d 9.7) [991], [¹²⁵ I]SHU9119 (Antagonist) [1392]	[¹²⁵ I]SHU9119 (Antagonist) (pK _d 9.2) [1392], [¹²⁵ I]NDP-MSH (Agonist) (pK _d 8.4–8.9) [991, 1665]	[¹²⁵ I]NDP-MSH (Agonist) (pK _d 8.6) [991]

Comments: Polymorphisms of the MC1 receptor have been linked to variations in skin pigmentation. Defects of the MC2 receptor underlie familial glucocorticoid deficiency. Polymorphisms of the MC4 receptor have been linked to obesity [282, 505].

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Melatonin receptors

G protein-coupled receptors → Melatonin receptors

Overview: Melatonin receptors (nomenclature as agreed by the [NC-IUPHAR Subcommittee on Melatonin Receptors \[446\]](#)) are activated by the endogenous ligands [melatonin](#) and [N-acetylserotonin](#).

Nomenclature	MT₁ receptor	MT₂ receptor
HGNC, UniProt	MTNR1A , P48039	MTNR1B , P49286
Endogenous agonists	melatonin (pK _i 9.1–9.7) [67 , 445 , 447]	melatonin (pK _i 9.4–9.8) [67 , 445 , 447]
Agonists	ramelteon (pK _i 10.9) [909], agomelatine (pK _i 10–10.4) [67 , 132]	agomelatine (pK _i 9.9–10.5) [67 , 132], ramelteon (pK _i 10) [909 , 1565]
Selective agonists	–	llok7 (pK _i 10.3) [506 , 1814], 5-methoxy-luzindole (Partial agonist) (pK _i 9.6) [447]
Selective antagonists	–	4P-PDOT (pK _i 8.8–9.4) [67 , 447 , 448], K185 (pK _i 9.3) [506 , 1814], DH97 (pK _i 8) [1865]
Labelled ligands	[¹²⁵I]SD6 (Agonist) (pK _d 10.9) [1074], 2-[¹²⁵I]melatonin (Agonist) (pK _d 9.9–10.7) [67 , 447], [³H]melatonin (Agonist) (pK _d 9.4–9.9) [230]	[¹²⁵I]SD6 (Agonist) (pK _d 10.2) [1074], 2-[¹²⁵I]melatonin (Agonist) (pK _d 9.7–10) [67 , 447], [¹²⁵I]DIV880 (Agonist, Partial agonist) (pK _d 9.7) [1074], [³H]melatonin (Agonist) (pK _d 9–9.6) [230]

Comments: [melatonin](#), [2-iodo-melatonin](#), [agomelatine](#), [GR 196429](#), [LY 156735](#) and [ramelteon](#) [[909](#)] are nonselective agonists for MT₁ and MT₂ receptors. (-)-AMMTC displays an ~400-fold greater agonist potency than (+)-AMMTC at rat MT₁ receptors (see [AMMTC](#) for structure) [[1888](#)]. [Luzindole](#) is an MT₁/MT₂ melatonin

receptor-selective competitive antagonist with some selectivity for the MT₂ receptor [[448](#)]. MT₁/MT₂ heterodimers present different pharmacological profiles from MT₁ and MT₂ receptors [[72](#)].

The MT₃ binding site of hamster brain and peripheral tissues such

as kidney and testis, also termed the ML₂ receptor, binds selectively [2-iodo-\[¹²⁵I\]SMCA-NAT](#) [[1302](#)]. Pharmacological investigations of MT₃ binding sites have primarily been conducted in hamster tissues. At this site, [N-acetylserotonin](#) [[467](#), [1149](#), [1302](#), [1516](#)] and

5MCA-NAT [1516] appear to function as agonists, while prazosin [1149] functions as an antagonist. The MT_3 binding site of hamster kidney was also identified as the hamster homologue of human quinone reductase 2 (NQO2, P16083 [1408, 1409]). The MT_3 bind-

ing site activated by 5MCA-NAT in eye ciliary body is positively coupled to adenylyl cyclase and regulates chloride secretion [802]. *Xenopus* melanophores and chick brain express a distinct receptor (x420, P49219; c346, P49288, initially termed Mel $_{1C}$) coupled to the $G_{i/o}$

family of G proteins, for which GPR50 has recently been suggested to be a mammalian counterpart [451] although melatonin does not bind to GPR50 receptors.

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Metabotropic glutamate receptors

G protein-coupled receptors → Metabotropic glutamate receptors

Overview: Metabotropic glutamate (mGlu) receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Metabotropic Glutamate Receptors** [1672]) are activated by the endogenous ligands L-glutamic acid, L-serine-O-phosphate, N-acetylaspartylglutamate (NAAG) and L-cysteine sulphinic acid. Examples of agonists selective for mGlu receptors compared with ionotropic glutamate receptors are (1S,3R)-ACPD and L-CCG-I, which show limited selectivity for Group-II receptors. An example of an antagonist selective for mGlu receptors is LY341495, which blocks mGlu $_2$ and mGlu $_3$ at low nanomolar concentrations, mGlu $_8$ at high nanomolar concentrations, and mGlu $_4$, mGlu $_5$, and mGlu $_7$ in the micromolar range [955]. Three groups of native receptors are distinguishable on the bases of similarities of agonist pharmacology, pri-

mary sequence and G protein coupling to effector: Group-I (mGlu $_1$ and mGlu $_5$), Group-II (mGlu $_2$ and mGlu $_3$) and Group-III (mGlu $_4$, mGlu $_6$, mGlu $_7$ and mGlu $_8$) (see Further reading). Group-I mGlu receptors may be activated by 3,5-DHPG and (S)-3HPC [198] and antagonized by (S)-hexylhomobotenic acid [1171]. Group-II mGlu receptors may be activated by LY389795 [1311], LY379268 [1311], eglumegad [1673, 2050], DCG-IV and (2R,3R)-APDC [1674], and antagonised by eGlu (4.3, [848] and LY307452 [491, 2009]. Group-III mGlu receptors may be activated by L-AP4 and (R,S)-4-PPG [579].

In addition to orthosteric ligands that directly interact with the glutamate recognition site directly, allosteric modulators have been described. Negative allosteric modulators are listed separately. The positive allosteric modulators most often act as 'potentiators' of an

orthosteric agonist response, without significantly activating the receptor in the absence of agonist.

Although mGlu receptors have been thought to only form homodimers, recent studies revealed the possible formation of heterodimers between either group-I receptors, or within and between group-II and -III receptors [441]. Although well characterized in transfected cells, co-localization and specific pharmacological properties also suggest the existence of such heterodimers in the brain [2094].

The structure of the 7 transmembrane (TM) domains of both mGlu $_1$ and mGlu $_5$ have been solved, and confirm a general helical organization similar to that of other GPCRs, although the helices appear more compacted [438, 2048].

	mGlu₁ receptor	mGlu₂ receptor	mGlu₃ receptor	mGlu₄ receptor
Nomenclature	mGlu₁ receptor	mGlu₂ receptor	mGlu₃ receptor	mGlu₄ receptor
HGNC, UniProt	GRM1, Q13255	GRM2, Q14416	GRM3, Q14832	GRM4, Q14833
Endogenous agonists	–	–	NAAG (Selective) (pK _i 4.7) [1679]	–
Agonists	–	–	–	L-AP4 (pEC ₅₀ 6.5) [2050], L-serine-O-phosphate (pEC ₅₀ 5.9) [2050]
Selective agonists	–	–	–	LSP4-2022 (pEC ₅₀ 7) [631]
Antagonists	LY367385 (pIC ₅₀ 5.1) [349]	–	–	MAP4 (pK _i 4.6) [686] – Rat
Selective antagonists	3-MATIDA (pIC ₅₀ 5.2) [1333] – Rat, (S)-(+)-CBPG (pIC ₅₀ 4.2) [1200] – Rat, (S)-TBPG (pIC ₅₀ 4.2) [367] – Rat, AIDA (pA ₂ 4.2) [1334]	PCCG-4 (pIC ₅₀ 5.1) [1483] – Rat	–	–
Allosteric modulators	YM298198 (Negative) (pIC ₅₀ 7.8) [979] – Rat	CBiPES (Positive) (pEC ₅₀ 7) [874], 4-MPPTS (Positive) (pIC ₅₀ 5.8) [94, 873, 874, 1660]	–	SIB-1893 (Positive) (pEC ₅₀ 6.3–6.8) [1217], MPEP (Positive) (pEC ₅₀ 6.3–6.6) [1217], PHCCC (Positive) (pEC ₅₀ 4.5) [1184]
Selective allosteric modulators	BAY 367620 (Negative) (pK _i 9.5) [267] – Rat, JNJ16259685 (Negative) (pIC ₅₀ 8.9) [1047], A-841720 (Negative) (pIC ₅₀ 8) [2126], Ro67-7476 (Positive) (pK _i 7.5–7.9) [971] – Rat, 3,5-dimethyl PPP (Negative) (pIC ₅₀ 7.8) [1271] – Rat, EM-TBPC (Negative) (pK _i 7.8) [1191] – Rat, Ro01-6128 (Positive) (pK _i 7.5–7.7) [971] – Rat, LY456236 (Negative) (pIC ₅₀ 6.9) [1094], CPCCOEt (Negative) (pIC ₅₀ 5.2–5.8) [1116], Ro67-4853 (Positive) (pK _i 5.1) [971] – Rat, PHCCC (Positive)	Ro64-5229 (Negative) (pIC ₅₀ 7) [985] – Rat, biphenylindanone A (Positive) (pEC ₅₀ 7) [183]	–	VU0361737 (Positive) (pEC ₅₀ 6.6) [480], VU0155041 (Positive) (pEC ₅₀ 6.1) [1402]
Comments	–	–	–	pEC ₅₀ values for MPEP and SIB-1893 were obtained in the presence of L-AP4 [1217].

Nomenclature	mGlu ₅ receptor	mGlu ₆ receptor	mGlu ₇ receptor	mGlu ₈ receptor
HGNC, UniProt	<i>GRM5</i> , P41594	<i>GRM6</i> , O15303	<i>GRM7</i> , Q14831	<i>GRM8</i> , O00222
Endogenous agonists	–	–	–	L-serine-O-phosphate (pIC ₅₀ 6.2–7.2) [1192, 2050]
Agonists	–	–	LSP4-2022 (pEC ₅₀ 4.9) [631], L-serine-O-phosphate (pEC ₅₀ 4.5) [2050], L-AP4 (pEC ₅₀ 3.8) [2050]	(S)-3,4-DCEPG (pEC ₅₀ 7.5) [1876], L-AP4 (pIC ₅₀ 7–7.2) [1192]
Selective agonists	(S)-(+)-CBPG (Partial agonist) (pEC ₅₀ 4.3) [1200] – Rat, CHPG (pIC ₅₀ 3.4) [1350]	1-benzyl-APDC (pEC ₅₀ 4.7) [1911] – Rat, homo-AMPA (pEC ₅₀ 4.1) [237] [1350]	–	–
Antagonists	–	MAP4 (pIC ₅₀ 3.5) [1504] – Rat, THPG [1879] – Unknown	–	MPPG (pIC ₅₀ 4.3) [2050]
Selective antagonists	ACDPP (pIC ₅₀ 6.9) [182]	–	–	–
Allosteric modulators	3,3'-difluorobenzaldazine (Positive) (pIC ₅₀ 5.6–8.5) [1415, 1416], alloswitch-1 (Negative) (pIC ₅₀ 8.1) [1511] – Rat, CDPPB (Positive) (pEC ₅₀ 7.6–8) [956, 1114], MTEP (Negative) (pK _i 7.8) [223], MPEP (Negative) (pIC ₅₀ 7.4–7.7) [578, 580], fenobam (Negative) (pIC ₅₀ 7.2) [1519], SIB-1893 (Negative) (pIC ₅₀ 5.9–6.5) [578, 1949], SIB-1757 (Negative) (pIC ₅₀ 6–6.4) [578, 1949], CPPHA (Positive) (pIC ₅₀ 6.3) [1416]	–	MMPIP (Negative) (pIC ₅₀ 6.1–7.6) [1401, 1827] – Rat, AMN082 (Positive) (pEC ₅₀ 6.5–6.8) [1293], XAP044 (Negative) (pIC ₅₀ 5.6) [587]	–
Selective allosteric modulators	VU-1545 (Positive) (pEC ₅₀ 8) [2142]	–	–	–

Comments: The activity of NAAG as an agonist at mGlu₃ receptors was questioned on the basis of contamination with glutamate [327, 547], but this has been refuted [1369].

Radioligand binding using a variety of radioligands has been conducted on recombinant receptors (for example, [³H]R214127 [1046] and [³H]YM298198 [979] at mGlu₁ receptors and [³H]M-MPEP [578] and [³H]methoxymethyl-MTEP [47] at mGlu₅ receptors). Although a number of radioligands have been used to examine binding in native tissues, correlation with individual subtypes is limited. Many pharmacological agents have not been fully tested

across all known subtypes of mGlu receptors. Potential differences linked to the species (*e.g.* human *versus* rat or mouse) of the receptors and the receptor splice variants are generally not known. The influence of receptor expression level on pharmacology and selectivity has not been controlled for in most studies, particularly those involving functional assays of receptor coupling.

(S)-(+)-CBPG is an antagonist at mGlu₁, but is an agonist (albeit of reduced efficacy) at mGlu₅ receptors. DCG-IV also exhibits agonist activity at NMDA glutamate receptors [1931], and is an antagonist at all group-III mGluRs with an IC₅₀ of 30 μM. A potential novel metabotropic glutamate receptor coupled to phospho-

inositide turnover has been observed in rat brain; it is activated by 4-methylhomobutenoic acid (ineffective as an agonist at recombinant Group I metabotropic glutamate receptors), but resistant to LY341495 [341]. There are also reports of a distinct metabotropic glutamate receptor coupled to phospholipase D in rat brain, which does not readily fit into the current classification [964, 1482]

A related class C receptor composed of two distinct subunits, T1R1 + T1R3 is also activated by glutamate and is responsible for umami taste detection.

All selective antagonists at metabotropic glutamate receptors are competitive.

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Motilin receptor

G protein-coupled receptors → Motilin receptor

Overview: Motilin receptors (**provisional nomenclature**) are activated by a 22 amino-acid peptide derived from a precursor (*MLN*, P12872), which may also generate a **motilin-associated peptide** (*MLN*, P12872). These receptors are also suggested to be responsible for the gastrointestinal prokinetic effects of certain macrolide antibiotics (often called motilides; *e.g.* erythromycin), although for many of these molecules the evidence is sparse.

Nomenclature	motilin receptor
HGNC, UniProt	MLNR , O43193
Endogenous agonists	motilin (MLN , P12872) (pK_i 8.4–8.7) [372 , 1223 , 1224 , 1225]
Agonists	alemcinal (pIC_{50} 7.2) [1872], erythromycin-A (pIC_{50} 5.5–6.5) [507 , 1872], azithromycin (pEC_{50} 5.5) [220]
Selective agonists	camicinal (pEC_{50} 7.9) [99 , 1639], mitemcinal (pEC_{50} 7.5–7.8) [977 , 1845] – Rabbit
Selective antagonists	MA-2029 (pA_2 9.2) [1811], GM-109 (pIC_{50} 8) [701] – Pig
Labelled ligands	[¹²⁵ I]motilin (human) (Agonist) (pK_d 10) [507]

Comments: In laboratory rodents, the gene encoding the motilin precursor appears to be absent, while the receptor appears to be a pseudogene [[725](#), [1637](#)]. Functions of [motilin](#) ([MLN](#), [P12872](#)) are not usually detected in rodents, although brain and other responses to motilin and the macrolide [alemcinal](#) have been reported and the mechanism of these actions are obscure [[1249](#), [1396](#)]. Marked dif-

ferences in ligand affinities for the motilin receptor in dogs and humans may be explained by significant differences in receptor structure [[1638](#)]. Note that for the complex macrolide structures, selectivity of action has often not been rigorously examined and other actions are possible (*e.g.* P2X inhibition by erythromycin; [[2123](#)]). Small molecule motilin receptor agonists are now described [[1093](#),

[1639](#), [2013](#)]. The motilin receptor does not appear to have constitutive activity [[774](#)]. Although not proven, the existence of biased agonism at the receptor has been suggested [[1225](#), [1292](#), [1636](#)]. A truncated 5-transmembrane structure has been identified but this is without activity when transfected into a host cell [[507](#)].

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Neuromedin U receptors

G protein-coupled receptors → Neuromedin U receptors

Overview: Neuromedin U receptors (**provisional nomenclature as recommended by NC-IUPHAR [530]**) are activated by the endogenous 25 amino acid peptide neuromedin U (**neuromedin U-25 (NMU, P48645)**), NmU-25, a peptide originally isolated from pig spinal cord [1287]. In humans, NmU-25 appears to be the sole product of a precursor gene (**NMU, P48645**) showing a broad tissue distribution, but which is expressed at highest lev-

els in the upper gastrointestinal tract, CNS, bone marrow and fetal liver. Much shorter versions of NmU are found in some species, but not in human, and are derived at least in some instances from the proteolytic cleavage of the longer NmU. Despite species differences in NmU structure, the C-terminal region (particularly the C-terminal pentapeptide) is highly conserved and contains biological activity. Neuromedin S (**neuromedin S-33 (NMS, Q5H8A3)**) has also been

identified as an endogenous agonist [1326]. NmS-33 is, as its name suggests, a 33 amino-acid product of a precursor protein derived from a single gene and contains an amidated C-terminal heptapeptide identical to NmU. NmS-33 appears to activate NMU receptors with equivalent potency to NmU-25.

Nomenclature	NMU1 receptor	NMU2 receptor
HGNC, UniProt	NMUR1, Q9HB89	NMUR2, Q9GZQ4
Antagonists	–	R-PSOP (pK _B 7) [1128]

Comments: NMU1 and NMU2 couple predominantly to G_{q/11} although there is evidence of good coupling to G_{i/o} [213, 786, 794]. NMU1 and NMU2 can be labelled with [¹²⁵I]-NmU and [¹²⁵I]-NmS (of various species, *e.g.* [1259]), BODIPY[®] TMR-NMU or Cy3B-NMU-8 [213]. A range of radiolabelled (¹²⁵I-), fluorescently labelled (*e.g.* Cy3, Cy5, rhodamine and FAM) and biotin labelled versions of **neuromedin U-25 (NMU, P48645)** and **neuromedin S-33 (NMS, Q5H8A3)** are now commercially available.

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Neuropeptide FF/neuropeptide AF receptors

G protein-coupled receptors → Neuropeptide FF/neuropeptide AF receptors

Overview: The Neuropeptide FF receptor family contains two subtypes, NPFF1 and NPFF2 (**provisional nomenclature [530]**), which exhibit high affinities for neuropeptide FF (*NPFF*, O15130) and RFamide related peptides (RFRP: precursor gene symbol *NPVF*, Q9HCQ7). NPFF1 is broadly distributed in the central nervous system with the highest levels found in the limbic system and the hypothalamus. NPFF2 is present in high density in the superficial layers of the mammalian spinal cord where it is involved in nociception and modulation of opioid functions.

Nomenclature	NPFF1 receptor	NPFF2 receptor
HGNC, UniProt	<i>NPFFR1</i> , Q9GZQ6	<i>NPFFR2</i> , Q9Y5X5
Rank order of potency	RFRP-1 (<i>NPVF</i> , Q9HCQ7) > RFRP-3 (<i>NPVF</i> , Q9HCQ7) > FMRFneuropeptide FF (<i>NPFF</i> , O15130) > neuropeptide AF (<i>NPFF</i> , O15130) > neuropeptide SF (<i>NPFF</i> , O15130), QRFP43 (<i>QRFP</i> , P83859), PrRP-31 (<i>PRLH</i> , P81277) [628]	neuropeptide AF (<i>NPFF</i> , O15130), neuropeptide FF (<i>NPFF</i> , O15130) > PrRP-31 (<i>PRLH</i> , P81277) > FMRF, QRFP43 (<i>QRFP</i> , P83859) > neuropeptide SF (<i>NPFF</i> , O15130) [628]
Endogenous agonists	neuropeptide FF (<i>NPFF</i> , O15130) (Selective) (pK _i : 8.5–9.9) [628, 629, 1306], RFRP-3 (<i>NPVF</i> , Q9HCQ7) (Selective) (pK _i : 9.2–9.3) [629, 630, 1306]	neuropeptide FF (<i>NPFF</i> , O15130) (Selective) (pK _i : 9.7) [629, 1305]
Selective agonists	–	dNPA (pK _i : 10.6) [1607], AC263093 (pEC ₅₀ : 5.2–5.9) [1036]
Antagonists	RF9 (pK _i : 7.2) [1745]	–
Selective antagonists	AC262620 (pK _i : 7.7–8.1) [1036], AC262970 (pK _i : 7.4–8.1) [1036]	–
Labelled ligands	[¹²⁵ I]Y-RFRP-3 (Agonist) (pK _d : 9.7) [629], [³ H]NPVF (Agonist) (pK _d : 8.6) [1855], [¹²⁵ I]NPFF (Agonist) [628]	[¹²⁵ I]EYF (Agonist) (pK _d : 10.2) [1306], [³ H]EYF (Agonist) (pK _d : 9.3) [1855], [¹²⁵ I]NPFF (Agonist) [628]

Comments: An orphan receptor *GPR83* (Q9NYM4) shows sequence similarities with NPFF1, NPFF2, PrRP and QRFP receptors. The antagonist RF9 is selective for NPFF receptors, but does not distinguish between the NPFF1 and NPFF2 subtypes (pK_i: 7.1 and 7.2, respectively, [1745]).

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Neuropeptide S receptor

G protein-coupled receptors → Neuropeptide S receptor

Overview: The neuropeptide S receptor (NPS, **provisional nomenclature** [530]) responds to the 20 amino-acid peptide neuropeptide S derived from the precursor (NPS, POCOP6).

Nomenclature	NPS receptor
HGNC, UniProt	NPSR1, Q6W5P4
Endogenous agonists	neuropeptide S (NPS, POCOP6) (pEC ₅₀ 8) [2070]
Labelled ligands	[¹²⁵ I]Tyr ¹⁰ NPS (human) (Agonist) (pK _d 9.5) [2070]

Comments: Polymorphisms in the NPS receptor have been suggested to be associated with asthma [1953] and irritable bowel syndrome [386].

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Neuropeptide W/neuropeptide B receptors

G protein-coupled receptors → Neuropeptide W/neuropeptide B receptors

Overview: The neuropeptide BW receptor 1 (NPBW1, **provisional nomenclature** [530]) is activated by two 23-amino-acid peptides, neuropeptide W (neuropeptide W-23 (NPW, Q8N729)) and neuropeptide B (neuropeptide B-23 (NPB, Q8NG41)) [554, 1725]. C-terminally extended forms of the peptides (neuropeptide W-30 (NPW, Q8N729)) and neuropeptide B-29 (NPB,

Q8NG41)) also activate NPBW1 [211]. Unique to both forms of neuropeptide B is the N-terminal bromination of the first tryptophan residue, and it is from this post-translational modification that the nomenclature NPB is derived. These peptides were first identified from bovine hypothalamus and therefore are classed as neuropeptides. Endogenous variants of the peptides without the N-

terminal bromination, des-Br-neuropeptide B-23 (NPB, Q8NG41) and des-Br-neuropeptide B-29 (NPB, Q8NG41), were not found to be major components of bovine hypothalamic tissue extracts. The NPBW2 receptor is activated by the short and C-terminal extended forms of neuropeptide W and neuropeptide B [211].

Nomenclature	NPBW1 receptor	NPBW2 receptor
HGNC, UniProt	NPBWR1, P48145	NPBWR2, P48146
Rank order of potency	neuropeptide B-29 (NPB, Q8NG41) > neuropeptide B-23 (NPB, Q8NG41) > neuropeptide W-23 (NPW, Q8N729) > neuropeptide W-30 (NPW, Q8N729) [211]	neuropeptide W-23 (NPW, Q8N729) > neuropeptide W-30 (NPW, Q8N729) > neuropeptide B-29 (NPB, Q8NG41) > neuropeptide B-23 (NPB, Q8NG41) [211]
Selective agonists	Ava3 (pK _i 9.4–9.4) [902], Ava5 (pK _i 8.8–9) [902]	–
Labelled ligands	[¹²⁵I]NPW-23 (human) (Agonist) (pK _d 9.4) [1747]	[¹²⁵I]NPW-23 (human) (Agonist) (pK _d 7.7) [1725]

Comments: Potency measurements were conducted with heterologously-expressed receptors with a range of 0.14–0.57 nM (NPBW1) and 0.98–21 nM (NPBW2).

NPBW1^{-/-} mice show changes in social behavior, suggesting that the NPBW1 pathway may have an important role in the emotional responses of social interaction [1355].

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Neuropeptide Y receptors

[G protein-coupled receptors](#) → [Neuropeptide Y receptors](#)

Overview: Neuropeptide Y (NPY) receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Neuropeptide Y Receptors** [1270]) are activated by the endogenous peptides [neuropeptide Y \(NPY, P01303\)](#), [neuropeptide Y-\(3-36\), peptide YY \(PYY, P10082\)](#), [PYY-\(3-36\)](#) and [pancreatic polypeptide \(PPY, P01298\)](#) (PP). The receptor originally identified as the Y3 receptor has been identified as the [CXCR4 chemokine receptor](#) (orig-

inally named LESTR, [1135]). The y6 receptor is a functional gene product in mouse, absent in rat, but contains a frame-shift mutation in primates producing a truncated non-functional gene [642]. Many of the agonists exhibit differing degrees of selectivity dependent on the species examined. For example, the potency of PP is greater at the rat Y₄ receptor than at the human receptor [485]. In addition, many agonists lack selectivity for individual subtypes, but can exhibit comparable potency against pairs of NPY receptor subtypes, or have not been examined for activity at all subtypes. [¹²⁵I]-PYY or [¹²⁵I]-NPY can be used to label Y₁, Y₂, Y₅ and Y₆ subtypes non-selectively, while [¹²⁵I][cPP(1-7), NPY(19-23), Ala³¹, Aib³², Gln³⁴]hPP may be used to label Y₅ receptors preferentially (note that cPP denotes chicken peptide sequence and hPP is the human sequence).

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Nomenclature	Y₁ receptor	Y₂ receptor	Y₄ receptor	Y₅ receptor	y₆ receptor
HGNC, UniProt	NPY1R, P25929	NPY2R, P49146	NPY4R, P50391	NPY5R, Q15761	NPY6R, Q99463
Rank order of potency	neuropeptide Y > peptide YY >> pancreatic polypeptide	neuropeptide Y > peptide YY >> pancreatic polypeptide	pancreatic polypeptide > neuropeptide Y = peptide YY	neuropeptide Y > peptide YY > pancreatic polypeptide	neuropeptide Y = peptide YY > pancreatic polypeptide
Endogenous agonists	neuropeptide Y (NPY, P01303) , peptide YY (PYY, P10082)	PYY-(3-36) (PYY, P10082) (pK _i 9.2–9.7) [588 , 599], neuropeptide Y (NPY, P01303) , neuropeptide Y-(3-36) (NPY, P01303) , peptide YY (PYY, P10082)	pancreatic polypeptide (PPY, P01298) (pK _i 8.7–10.9) [92 , 1152 , 1899 , 2076]	–	–
Selective agonists	[Leu³¹, Pro³⁴]NPY (pEC ₅₀ 7.1) [378], [Leu³¹, Pro³⁴]PYY (human), [Pro³⁴]NPY , [Pro³⁴]PYY (human)	–	–	[Ala³¹, Aib³²]NPY (pig) (pIC ₅₀ 8.2) [254]	–
Selective antagonists	BIBO3304 (pIC ₅₀ 9.5) [2020], BIBP3226 (pK _i 8.1–9.3) [436 , 2021]	BIIE0246 (pIC ₅₀ 8.5) [434], JNJ-5207787 (pIC ₅₀ 6.9–7.1) [178]	–	L-152,804 (pK _i 7.6) [901]	–
Labelled ligands	[³H]BIBP3226 (Antagonist) (pK _d 8.7), [¹²⁵I][Leu³¹, Pro³⁴]NPY (Agonist)	[¹²⁵I]PYY-(3-36) (human) (Agonist)	[¹²⁵I]PP (human) (Agonist)	[¹²⁵I][cPP(1-7), NPY(19-23), Ala³¹, Aib³², Gln³⁴]hPP (Agonist) (pK _d 9.2–9.3) [453] – Rat	–
Comments	Note that Pro ³⁴ -containing NPY and PYY can also bind Y ₄ and Y ₅ receptors, so strictly speaking are not selective, but are the 'preferred' agonists.	–	–	–	The y ₆ receptor is a pseudogene in humans, but is functional in mouse, rabbit and some other mammals.

Comments: The Y₁ agonists indicated are selective relative to Y₂ receptors. [BIBP3226](#) is selective relative to Y₂, Y₄ and Y₅ receptors [[598](#)]. [NPY-\(1-36\)](#) is Y₂ selective relative to Y₁ and Y₅ receptors. [PYY-\(3-36\)](#) is Y₂ selective relative to Y₁ receptors.

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Neurotensin receptors

G protein-coupled receptors → Neurotensin receptors

Overview: Neurotensin receptors (**nomenclature as recommended by NC-IUPHAR [530]**) are activated by the endogenous tridecapeptide neurotensin (pGlu-Leu-Tyr-Glu-Asn-Lys-Pro-Arg-Arg-Pro-Tyr-Ile-Leu) derived from a precursor (*NTS*, 30990), which also generates neuromedin N, an agonist at the NTS₂ receptor. A nonpeptide antagonist, **SRI42948A**, shows high affinity (pK_i ~9) at both NTS₁ and NTS₂ receptors [664]. [³H]neurotensin (human, mouse, rat) and [¹²⁵I]neurotensin (human, mouse, rat) may be used to label NTS₁ and NTS₂ receptors at 0.1-0.3 and 3-5 nM concentrations respectively.

Nomenclature	NTS ₁ receptor	NTS ₂ receptor
HGNC, UniProt	<i>NTSR1</i> , P30989	<i>NTSR2</i> , O95665
Rank order of potency	neurotensin (<i>NTS</i> , P30990) > neuromedin N {Mouse, Rat} [741]	neurotensin (<i>NTS</i> , P30990) = neuromedin N {Mouse, Rat} [1235]
Selective agonists	JMV449 (pK _i 10) [1753] – Rat	levocabastine (pK _i 6.8) [1235, 1583]
Antagonists	meclizant (pIC ₅₀ 7.5–8.2) [664]	–
Labelled ligands	[³ H]meclizant (Antagonist) (pK _d 8.5) [1030] – Rat	–
Comments	–	A splice variant of the NTS ₂ receptor bearing 5 transmembrane domains has been identified in mouse [191] and later in rat [1492].

Comments: neurotensin (*NTS*, P30990) appears to be a low-efficacy agonist at the NTS₂ receptor [1959], while the NTS₁ receptor antagonist meclizant is an agonist at NTS₂ receptors [1959]. An additional protein, provisionally termed NTS₃ (also known as NTR3, gp95 and sortilin; *ENSG00000134243*), has been suggested to bind lipoprotein lipase and mediate its degradation [1395]. It has been reported to interact with the NTS₁ receptor [1211] and has been implicated in hormone trafficking and/or neurotensin uptake.

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Opioid receptors

G protein-coupled receptors → Opioid receptors

Overview: Opioid and opioid-like receptors are activated by a variety of endogenous peptides including [Met]enkephalin (*PENK*, P01210) (met), [Leu]enkephalin (*PENK*, P01210) (leu), β -endorphin (*POMC*, P01189) (β -end), α -neodynorphin (*PDYN*, P01213), dynorphin A (*PDYN*, P01213) (dynA), dynorphin B

(*PDYN*, P01213) (dynB), big dynorphin (*PDYN*, P01213) (Big dyn), nociceptin/orphanin FQ (*PNOC*, Q13519) (N/OFQ); endomorphin-1 and endomorphin-2 are also potential endogenous peptides. The Greek letter nomenclature for the opioid receptors, μ , δ and κ , is well established, and **NC-IUPHAR** considers this nomenclature most

appropriate [376, 417, 530]. The human N/OFQ receptor is considered 'opioid-related' rather than opioid because while it exhibits a high degree of structural homology with the conventional opioid receptors [1308], it displays a distinct pharmacology.

Nomenclature	δ receptor	κ receptor	μ receptor	NOP receptor
HGNC, UniProt	<i>OPRD1</i> , P41143	<i>OPRK1</i> , P41145	<i>OPRM1</i> , P35372	<i>OPRL1</i> , P41146
Principal endogenous agonists	β -endorphin (<i>POMC</i> , P01189), [Leu]enkephalin (<i>PENK</i> , P01210), [Met]enkephalin (<i>PENK</i> , P01210)	big dynorphin (<i>PDYN</i> , P01213), dynorphin A (<i>PDYN</i> , P01213)	β -endorphin (<i>POMC</i> , P01189), [Met]enkephalin (<i>PENK</i> , P01210), [Leu]enkephalin (<i>PENK</i> , P01210), endomorphin-1, endomorphin-2	–
Endogenous agonists	–	–	endomorphin-2 (Selective) (pK _i 8.5) [2109] – Rat, endomorphin-1 (Selective) (pK _i 8.3) [622, 2109]	nociceptin/orphanin FQ (<i>PNOC</i> , Q13519) (Selective) (pK _i 9.7–10.4) [149, 1242, 1303, 1307, 1439]
Agonists	–	–	levorphanol (pIC ₅₀ 9.9) [692], hydromorphone (pK _i 9.6) [2007], fentanyl (pK _i 9.2) [1893], buprenorphine (Partial agonist) (pK _i 8.8) [1893], methadone (pIC ₅₀ 8.4) [1523], codeine (pK _i 6.9) [1893], tapentadol (pK _i 6.8) [1916], meperidine (pIC ₅₀ 6.5) [1523]	–
Selective agonists	[D-Ala ²]deltorphin I (pK _d 9.4) [487, 1795], DPDPE (pK _i 8.8) [1337, 1893], [D-Ala ²]deltorphin II (pK _i 8.8) [488], SNC80 (pK _i 7.2) [258, 1549]	U50488 (pK _i 7.8–9.7) [297, 1478, 1744, 1893, 1962, 2128, 2130], enadoline (pK _i 9.6) [808, 1383], U69593 (pK _i 9.5) [1034, 1893], salvinorin A (pK _i 7.8–8.7) [251, 1603]	sufentanil (pK _i 9.9) [1960], DAMGO (pK _i 9.3) [691, 1893], loperamide (pK _i 9.3) [308], morphine (pK _i 9) [620, 1893], PL017 (pK _i 8.2) [290, 1893]	N/OFQ-(1-13)-NH ₂ (pK _i 10.1–10.4) [149, 661, 1242, 1439], Ro64-6198 (pK _i 9.6) [855]

(continued)	δ receptor	κ receptor	μ receptor	NOP receptor
Nomenclature	δ receptor	κ receptor	μ receptor	NOP receptor
Antagonists	naltrexone (pK _i 8) [1893], naloxone (pK _i 7.2) [1893]	buprenorphine (pK _i 9.1–10.2) [1893, 2130], nalmefene (pK _i 9.5) [1893], naltrexone (pK _i 8.4–9.4) [1478, 1744, 1893], naloxone (pK _i 7.6–8.6) [1478, 1744, 1893, 2128, 2130]	naltrexone (pK _i 9.7) [1893], nalmefene (pK _i 9.5) [1893], nalorphine (pK _i 8.9) [1893], methylnaltrexone (pK _i 8.7) [2007]	–
Selective antagonists	naltriben (pK _i 10) [1773, 1893], naltrindole (pK _i 9.7) [1521, 1893], TIPP ψ (Inverse agonist) (pK _i 9) [1664, 1893]	nor-binaltorphimine (pK _i 8.9–11) [1478, 1520, 1744, 1893, 2128, 2130], 5'-guanidinonaltrindole (pK _i 9.7–9.9) [882, 1478, 1797]	alvimopan (pK _i 9.3) [1056], levallorphan (pK _i 8.8–9.3) [1187], CTAP (pK _i 8.6) [290, 1893]	UFP-101 (pK _i 10.2) [259], Banyu Compound-24 (pK _i 9.6) [523], SB 612111 (pK _i 9.5) [2112], J-113397 (pI _{C50} 8.3) [919]
Labelled ligands	[³ H]naltrindole (Antagonist) (pK _d 10.4) [2072] – Rat, [³ H]DPDPE (Agonist) [26], [³ H]deltorphan II (Agonist) [252], [³ H]naltriben (Antagonist) [1088]	[³ H]U69593 (Agonist) (pK _d 8.7–8.8) [1034, 1478, 1744], [³ H]enadoline (Agonist) [1746]	[³ H]DAMGO (Agonist) (pK _d 9.2) [1567] – Rat, [³ H]PL017 (Agonist) [717] – Rat	[³ H]N/OFQ (Agonist) (pK _d 10.2) [437, 1307]

Comments: Three naloxone-sensitive opioid receptor genes have been identified in humans, and while the μ -receptor in particular may be subject to extensive alternative splicing [1468], these putative isoforms have not been correlated with any of the subtypes of receptor proposed in years past. Opioid receptors may heterodimerize with each other or with other 7TM receptors [884], and give rise to complexes with a unique pharmacology, however, evidence for such heterodimers in native cells is equivocal and the consequences this heterodimerization for signalling remains largely unknown. For μ -opioid receptors at least, dimerization does not seem to be required for signalling [1026]. A distinct met-enkephalin receptor lacking structural resemblance to the opioid receptors listed has been

identified (*OGFR*, *9NZT2*) and termed an opioid growth factor receptor [2110].

Endomorphin-1 and **endomorphin-2** have been identified as highly selective, putative endogenous agonists for the μ -opioid receptor. At present, however, the mechanisms for endomorphin synthesis *in vivo* have not been established, and there is no gene identified that encodes for either. Thus, the status of these peptides as endogenous ligands remains unproven.

Two areas of increasing importance in defining opioid receptor function are the presence of functionally relevant single nucleotide polymorphisms in human μ -receptors [1423] and the identification of bi-

ased signalling by opioid receptor ligands, in particular, compounds previously characterized as antagonists [231]. Pathway bias for agonists makes general rank orders of potency and efficacy somewhat obsolete, so these do not appear in the table. As ever, the mechanisms underlying the acute and long term regulation of opioid receptor function are the subject of intense investigation and debate.

The richness of opioid receptor pharmacology has been enhanced with the recent discovery of allosteric modulators of MOPr and DOPr, notably the positive allosteric modulators and silent allosteric "antagonists" outlined in [240, 241]. Negative allosteric modulation of opioid receptors has been previously suggested [908], whether all compounds are acting at a similar site remains to be established.

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Orexin receptors

G protein-coupled receptors → Orexin receptors

Overview: Orexin receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Orexin receptors [627]**) are activated by the endogenous polypeptides orexin-A (*HCRT*, O43612) and orexin-B (*HCRT*, O43612) (also known as hypocretin-1 and -2; 33 and 28 aa) derived from a common precursor, **preproorexin or orexin precursor**, by proteolytic cleavage [1629]. Binding to both receptors may be accomplished with [¹²⁵I]orexin A (human, mouse, rat) [773].

Nomenclature	OX ₁ receptor	OX ₂ receptor
HGNC, UniProt	<i>HCRT1</i> , O43613	<i>HCRT2</i> , O43614
Rank order of potency	orexin-A (<i>HCRT</i> , O43612) > orexin-B (<i>HCRT</i> , O43612)	orexin-A (<i>HCRT</i> , O43612) = orexin-B (<i>HCRT</i> , O43612)
Selective agonists	–	[Ala ¹¹ , D-Leu ¹⁵]orexin-B (pEC ₅₀ 9.9) [62]
(Sub)family-selective antagonists	suvorexant (pK _i 9.3) [377], SB-649868 (pK _i 9.1) [419], filorexant (pK _i 8.6) [2035], almorexant (pIC ₅₀ 7.9) [216]	filorexant (pK _i 9.5) [2035], suvorexant (pK _i 9.5) [377], SB-649868 (pK _i 8.9) [419], almorexant (pIC ₅₀ 8.1) [216]
Selective antagonists	SB-408124 (pK _i 7.2–7.6) [1042, 1190], SB-334867 (pK _i 7.4–7.5) [1190, 1518]	EMPA (pK _i 9) [1189], JNJ 10397049 (pK _i 7.9–8.6) [1238], TCS-OX2-29 (pK _i 7.4) [760]
Labelled ligands	[³ H]SB-674042 (Antagonist) (pK _d 8.3–9.1) [1042, 1190, 1193]	–

Comments: The primary coupling of orexin receptors to G_{q/11} proteins is rather speculative and based on the strong activation of phospholipase C. Coupling of both receptors to G_{i/o} and G_s has also been reported [1019, 1555]; for most cellular responses observed, the G protein pathway is unknown. The rank order of endogenous agonist potency may depend on the cellular signal transduction machinery. The synthetic [Ala¹¹, D-Leu¹⁵]orexin-B may show poor OX₂ receptor selectivity [1540].

Loss-of-function mutations in the gene encoding the OX₂ receptor underlie canine hereditary narcolepsy [1111].

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Oxoglutarate receptor

G protein-coupled receptors → Oxoglutarate receptor

Overview: Nomenclature as recommended by NC-IUPHAR [396].

Nomenclature	oxoglutarate receptor
HGNC, UniProt	<i>OXGR1</i> , Q96P68
Endogenous agonists	α -ketoglutaric acid (pEC ₅₀ 3.3–4.5) [728, 1785]

P2Y receptors

G protein-coupled receptors → P2Y receptors

Overview: P2Y receptors (nomenclature as agreed by the **NC-IUPHAR Subcommittee on P2Y Receptors [1, 2]**) are activated by the endogenous ligands ATP, adenosine diphosphate, uridine triphosphate, uridine diphosphate and UDP-glucose. The relationship of many of the cloned receptors to endogenously expressed receptors is not yet established and so it might be appropriate to use wording such as 'uridine triphosphate-preferring (or ATP-, etc.) P2Y receptor' or 'P2Y₁-like', etc., until further, as yet undefined, corroborative criteria can be applied [244, 486, 837, 2003, 2146].

Nomenclature	P2Y ₁ receptor	P2Y ₂ receptor	P2Y ₄ receptor	P2Y ₆ receptor
HGNC, UniProt	<i>P2RY1</i> , P47900	<i>P2RY2</i> , P41231	<i>P2RY4</i> , P51582	<i>P2RY6</i> , Q15077
Rank order of potency	adenosine diphosphate>ATP	uridine triphosphate=ATP	uridine triphosphate>ATP (at rat recombinant receptors, UTP = ATP)	uridine diphosphate>> uridine triphosphate>ATP
Endogenous agonists	–	–	–	uridine diphosphate (Selective) (pEC ₅₀ 6.5) [359]
Agonists	ADP β S (pEC ₅₀ 7.3) [1848], 2MeSADP (pIC ₅₀ 5.4–7) [1658, 1970]	–	–	Rp-5-OMe-UDP α B (pEC ₅₀ 8.1) [611, 666]
Selective agonists	MRS2365 (pEC ₅₀ 9.4) [314], 2-CI-ADP(α -BH ₃) (pEC ₅₀ 8.1) [73]	2-thioUTP (pEC ₅₀ 7.3) [470], PSB1114 (EC ₅₀ value determined using an IP ₃ functional assay) (pEC ₅₀ 6.9) [471], Ap ₄ A (pEC ₅₀ 6.1) [270, 1471], UTP γ S (pEC ₅₀ 5.8) [1054], MRS2768 (EC ₅₀ value determined using an IP ₃ functional assay) (pEC ₅₀ 5.7) [973]	MRS4062 (pEC ₅₀ 7.6) [1213], UTP γ S [1055] – Unknown	MRS2957 (pEC ₅₀ 7.9) [1212], MRS2693 (pEC ₅₀ 7.8) [143], 3-phenacyl-UDP (pEC ₅₀ 7.2) [470]

(continued)				
Nomenclature	P2Y ₁ receptor	P2Y ₂ receptor	P2Y ₄ receptor	P2Y ₆ receptor
Antagonists	–	–	ATP (pK _d 6.2) [925]	–
Selective antagonists	MRS2500 (pK _i 8.8–9.1) [274, 942], BMS compound 16 [PMID:23368907] (pK _i 8.2) [295, 2115], MRS2279 (pK _i 7.9) [1970], MRS2179 (pK _i 7–7.1) [197, 1970], 2,2'-pyridylisatogen tosylate (pK _i 6.8) [570]	AR-C118925XX (pIC ₅₀ ~6) [924]	–	MRS2578 (pIC ₅₀ 7.4) [1196]
Labelled ligands	[³ H]MRS2279 (Antagonist) (pK _d 8.1) [1970], [³ H]2MeSADP (Agonist) (pK _d 7.3) [1848], [³⁵ S]ADPβS (Agonist) – Unknown	–	–	–
Nomenclature	P2Y ₁₁ receptor	P2Y ₁₂ receptor	P2Y ₁₃ receptor	P2Y ₁₄ receptor
HGNC, UniProt	P2RY11, Q96G91	P2RY12, Q9H244	P2RY13, Q9BPV8	P2RY14, Q15391
Rank order of potency	ATP>uridine triphosphate	–	adenosine diphosphate≫ATP	–
Rank order of potency Human	–	–	–	uridine diphosphate ≥ UDP-glucose
Endogenous agonists	–	adenosine diphosphate (Selective) (pK _i 5.9) [740]	–	–
Agonists	–	2MeSADP (pK _i 9.2) [740]	–	MRS2690 (pEC ₅₀ 6.6–7.3) [571, 974]
Selective agonists	AR-C67085 (pEC ₅₀ 8.5) [88, 360], NF546 (pEC ₅₀ 6.3) [1255], NAADP [1322], NAD [1323]	–	–	–
Antagonists	NF340 (pIC ₅₀ 6.4–7.1) [1255]	PSB-0739 (pK _i 7.6) [91]	–	–
Selective antagonists	NF157 (pK _i 7.3) [1923]	AZD1283 (pK _i 8) [76, 2116], ARL66096 (pIC ₅₀ 7.9) [806, 807], ticagrelor (pK _i 7.8) [2113]	MRS2211 (pIC ₅₀ 6) [950]	PPTN (pK _i 10.1) [96]
Labelled ligands	–	[³ H]2MeSADP (Agonist) (pIC ₅₀ 7.5–9.6) [1848], [³ H]PSB-0413 (Antagonist) (pK _d 8.3–8.5) [469, 1431]	–	–

Comments: [cangrelor](#) shows selectivity for P2Y₁₂ and P2Y₁₃ receptors compared with other P2Y receptors [1209, 1848]. [NF157](#) also has antagonist activity at P2X₁ receptors [1923]. [Uridine diphosphate](#) has been reported to be an antagonist at the P2Y₁₄ receptor [548]. [³⁵S]ATP α S has been used to label P2Y receptors in rat synaptosomal membranes [1682, 1683].

An orphan GPCR suggested to be a 'P2Y₁₅' receptor [823] appears not to be a genuine nucleotide receptor [2], but rather responds to dicarboxylic acids [728]. Further P2Y-like receptors have been cloned from non-mammalian sources; a clone from chick brain, termed a p2y₃ receptor ([ENSGALG00000017327](#)), couples to the G_{q/11} family of G proteins and shows the rank order

of potency [adenosine diphosphate](#) > [uridine triphosphate](#) > [ATP](#) = [uridine diphosphate](#) [1998]. In addition, human sources have yielded a clone with a preliminary identification of p2y5 ([LPA6](#), [P43657](#)) and contradictory evidence of responses to [ATP](#) [954, 1999]. This protein is now classified as [LPA₆](#), a receptor for lysophosphatidic acid ([LPA](#)) [1467, 2079]. The clone termed p2y9 ([LPA4](#), [Q99677](#)) is also a receptor for lysophosphatidic acid, [LPA₄](#) [1406]. The clone p2y7 ([NOP9](#), [Q86U38](#)), originally suggested to be a P2Y receptor [22], has been shown to encode a leukotriene receptor [2095]. A P2Y receptor that was initially termed a p2y8 receptor ([P79928](#)) has been cloned from *Xenopus laevis*; it shows the rank order of potency [ADP \$\beta\$ S](#) > [ATP](#) = [uridine triphosphate](#) = [guanosine-5'-triphosphate](#) = [CTP](#) = [TTP](#) = [ITP](#) > [ATP \$\gamma\$ S](#) and elic-

its a Ca²⁺-dependent Cl⁻ current in *Xenopus* oocytes [169]. The p2y10 clone ([P2RY10](#), [O00398](#)) lacks functional data. Diadenosine polyphosphates also have effects on as yet uncloned P2Y-like receptors with the rank order of potency of [Ap₄A](#) > [Ap_{5a}](#) > [Ap_{3a}](#), coupling via G_{q/11} [270]. P2Y-like receptors have recently been described on mitochondria [126]. CysLT1 and CysLT2 leukotriene receptors respond to nanomolar concentrations of [uridine diphosphate](#), although they are activated principally by leukotrienes [LTC₄](#) and [LTD₄](#) [1257, 1258]. Human [GPR17](#) (13304) and rat GPR17, which are structurally related to CysLT and P2Y receptors, are also activated by leukotrienes [1542] as well as [uridine diphosphate](#) and [UDP-glucose](#) [344, 540]. Activity at the rat GPR17 is inhibited by submicromolar concentrations of [MRS2179](#) and [cangrelor](#) [344].

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Parathyroid hormone receptors

G protein-coupled receptors → Parathyroid hormone receptors

Overview: The parathyroid hormone receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Parathyroid Hormone Receptors [575]**) are family B G protein-coupled receptors. The parathyroid hormone (PTH)/parathyroid hormone-related peptide (PTHrP) receptor (PTH1 receptor) is activated by precursor-derived peptides: [PTH](#) ([PTH](#), [P01270](#)) (84 amino acids), and [PTHrP](#) ([PTHrP](#), [P12272](#)) (141 amino-acids) and related peptides (PTH-(1-34), [PTHrP-\(1-36\)](#) ([PTHrP](#), [P12272](#))). The parathyroid hormone 2 receptor (PTH2 receptor) is activated by the precursor-derived peptide [TIP39](#) ([PTH2](#), [Q96A98](#)) (39 amino acids). [¹²⁵I]PTH may be used to label both PTH1 and PTH2 receptors.

Nomenclature	PTH1 receptor	PTH2 receptor
HGNC, UniProt	PTH1R, Q03431	PTH2R, P49190
Rank order of potency	PTH (PTH, P01270) = PTHrP (PTHLH, P12272)	TIP39 (PTH2, Q96A98), PTH (PTH, P01270) ≫ PTHrP (PTHLH, P12272)
Endogenous agonists	–	TIP39 (PTH2, Q96A98) (pIC ₅₀ 7.6–9.2) [626, 766]
Agonists	teriparatide (pIC ₅₀ 7.4) [573]	–
Selective agonists	PTHrP-(1-34) (human) (pIC ₅₀ 7.8–8.1) [574] – Rat	–

Comments: Although [PTH](#) ([PTH, P01270](#)) is an agonist at human PTH2 receptors, it fails to activate the rodent orthologues. [TIP39](#) ([PTH2, Q96A98](#)) is a weak antagonist at PTH1 receptors [[883](#)].

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Platelet-activating factor receptor

G protein-coupled receptors → Platelet-activating factor receptor

Overview: Platelet-activating factor (PAF, 1-O-alkyl-2-acetyl-sn-glycero-3-phosphocholine) is an ether phospholipid mediator associated with platelet coagulation, but also subserves inflammatory roles. The PAF receptor (**provisional nomenclature recommended by NC-IUPHAR** [[530](#)]) is activated by PAF and other suggested endogenous ligands are oxidized phosphatidylcholine [[1204](#)] and lysophosphatidylcholine [[1425](#)]. It may also be activated by bacterial lipopolysaccharide [[1358](#)].

Nomenclature	PAF receptor
HGNC, UniProt	PTAFR, P25105
Selective agonists	methylcarbamyl PAF – Unknown
Selective antagonists	foropafant (pK _i 10.3) [739], ABT-491 (pK _i 9.2) [30], CV-6209 (pIC ₅₀ 8.1–8.3) [619, 1357], L659989 (pK _i 7.8) [811], apafant (pK _i 5.2–7.5) [1460, 1831]
Labelled ligands	[³H]PAF (Agonist) (pK _d 8.8–8.9) [555, 1357]

Comments: Note that a previously recommended radioligand ($[^3\text{H}]$ apafant; K_d 44.6 nM) is currently unavailable.

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Prokineticin receptors

G protein-coupled receptors → Prokineticin receptors

Overview: Prokineticin receptors, PKR₁ and PKR₂ (**provisional nomenclature as recommended by NC-IUPHAR [530]**) respond to the cysteine-rich 81–86 amino-acid peptides **prokineticin-1 (PROK1, Q9HC23)** (also known as endocrine gland-derived vascular

endothelial growth factor, mambakine) and **prokineticin-2 (PROK2, Q9HC23)** (protein Bv8 homologue). An orthologue of PROK1 from black mamba (*Dendroaspis polylepis*) venom, mamba intestinal toxin 1 (MIT1, [1678]) is a potent, non-selective agonist at prokineticin

receptors [1215], while **Bv8**, an orthologue of PROK2 from amphibians (*Bombina sp.*, [1304]), is equipotent at recombinant PKR₁ and PKR₂ [1371], and has high potency in macrophage chemotaxis assays, which are lost in PKR₁-null mice.

Nomenclature	PKR ₁	PKR ₂
HGNC, UniProt	PROKR1, Q8TCW9	PROKR2, Q8NFJ6
Rank order of potency	prokineticin-2 (PROK2, Q9HC23) > prokineticin-1 (PROK1, Q9HC23) > prokineticin-2β (PROK2) [1109, 1215, 1775]	prokineticin-2 (PROK2, Q9HC23) > prokineticin-1 (PROK1, Q9HC23) > prokineticin-2β (PROK2) [1109, 1215, 1775]
Endogenous agonists	prokineticin-2 (PROK2, Q9HC23) (pIC ₅₀ 8.2–8.4) [300, 1215], prokineticin-1 (PROK1, Q9HC23) (pIC ₅₀ 6.6–7.6) [300, 1215], prokineticin-2β (PROK2) (pIC ₅₀ 7.5) [300]	prokineticin-2 (PROK2, Q9HC23) (pIC ₅₀ 8.1–8.2) [300, 1215], prokineticin-1 (PROK1, Q9HC23) (pIC ₅₀ 7.1–7.3) [300, 1215], prokineticin-2β (PROK2) (pIC ₅₀ <6) [300]
Agonists	MIT1 (pIC ₅₀ 8.4) [1215]	MIT1 (pIC ₅₀ 9.2) [1215]
Selective agonists	IS20 (pEC ₅₀ 7.4) [581], IS1 (pEC ₅₀ 5.6) [581]	–
Selective antagonists	triazine compound PC1 (pK _i 7.7) [87], triazine compound PC7 (pIC ₅₀ 7.5) [842, 1552], triazine compound PC10 (pIC ₅₀ 7) [842]	PKR-A (pIC ₅₀ 7.3–7.4) [312]
Labelled ligands	[¹²⁵ I]BH-MIT1 (Agonist) (pIC ₅₀ 8.4) [1215]	[¹²⁵ I]BH-MIT1 (Agonist) (pIC ₅₀ 9.2) [1215]

Comments: Genetic mutations in *PROKR1* are associated with Hirschsprung's disease [1614], while genetic mutations in *PROKR2* are associated with hypogonadotropic hypogonadism with anosmia [430], hypopituitarism with pituitary stalk interruption [1575] and Hirschsprung's disease [1614].

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Prolactin-releasing peptide receptor

G protein-coupled receptors → Prolactin-releasing peptide receptor

Overview: The precursor (*PRLH*, P81277) for PrRP generates 31 and 20-amino-acid versions. QRFP43 (*QRFP*, P83859) (named after a pyroglutamylated arginine-phenylalanine-amide peptide) is a 43 amino acid peptide derived from QRFP (P83859) and is also known as P518 or 26RFa. RFRP is an RF amide-related peptide [756] derived from a FMRFamide-related peptide precursor (*NPVF*, Q9HCQ7), which is cleaved to generate neuropeptide SF (*NPFF*, O15130), neuropeptide RFRP-1 (*NPVF*, Q9HCQ7), neuropeptide RFRP-2 (*NPVF*, Q9HCQ7) and neuropeptide RFRP-3 (*NPVF*, Q9HCQ7) (neuropeptide NPVF).

Nomenclature	PrRP receptor
HGNC, UniProt	<i>PRLHR</i> , P49683
Rank order of potency	PrRP-20 (<i>PRLH</i> , P81277), PrRP-31 (<i>PRLH</i> , P81277) [1043]
Endogenous agonists	PrRP-20 (<i>PRLH</i> , P81277) (Selective) (pK _i 9–9.6) [481, 1043], PrRP-31 (<i>PRLH</i> , P81277) (Selective) (pK _i 9–9.2) [481, 1043]
Endogenous antagonists	neuropeptide Y (<i>NPY</i> , P01303) (Selective) (pK _i 5.4) [1032]
Labelled ligands	[¹²⁵ I]PrRP-20 (human) (Agonist) (pK _d 9.2–10.6) [1043], [¹²⁵ I]PrRP31 (Agonist) [473]

Comments: The orphan receptor *GPR83* (Q9NYM4) shows sequence similarities with NPFF1, NPFF2, PrRP and QRFP receptors.

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Prostanoid receptors

G protein-coupled receptors → Prostanoid receptors

Overview: Prostanoid receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Prostanoid Receptors [2043]**) are activated by the endogenous ligands prostaglandins PGD_2 , PGE_2 , $\text{PGF}_{2\alpha}$, PGH_2 , prostacyclin [PGI_2] and **thromboxane A_2** . Measurement of the potency of PGI_2 and **thromboxane A_2** is hampered by their instability in physiological salt solution; they are often replaced by **cicaprost** and **U46619**, respectively, in receptor characterization studies.

Nomenclature	DP ₁ receptor	DP ₂ receptor	IP receptor	FP receptor	TP receptor
HGNC, UniProt	<i>PTGDR</i> , Q13258	<i>PTGDR2</i> , Q9YSY4	<i>PTGIR</i> , P43119	<i>PTGFR</i> , P43088	<i>TBXA2R</i> , P21731
Rank order of potency	$\text{PGD}_2 \gg \text{PGE}_2 > \text{PGF}_{2\alpha} > \text{PGI}_2, \text{thromboxane } A_2$	–	$\text{PGI}_2 \gg \text{PGD}_2, \text{PGE}_2, \text{PGF}_{2\alpha} > \text{thromboxane } A_2$	$\text{PGF}_{2\alpha} > \text{PGD}_2 > \text{PGE}_2 > \text{PGI}_2, \text{thromboxane } A_2$	$\text{thromboxane } A_2 = \text{PGH}_2 \gg \text{PGD}_2, \text{PGE}_2, \text{PGF}_{2\alpha}, \text{PGI}_2$
Rank order of potency	–	$\text{PGD}_2 \gg \text{PGF}_{2\alpha}, \text{PGE}_2 > \text{PGI}_2, \text{thromboxane } A_2$	–	–	–
Agonists	–	13,14-dihydro-15-keto-PGD ₂ (pK _i 7.4–8.5) [712, 1656, 1815]	iloprost (pK _i 7.5–8) [7, 2030], treprostinil (pK _i 7.5) [2019]	bimatoprost (pIC ₅₀ 5.3) [2044]	–
Selective agonists	BW 245C (pK _i 8.4–9.4) [171, 2045, 2046], L-644,698 (pK _i 9–9.3) [2045, 2046], SQ-27986 (pK _i 8) [1712], RS 93520 (Partial agonist) (pK _i 7.5) [1712], ZK118182 (pK _i 7.3) [1712]	15(R)-15-methyl-PGD ₂ (pK _i 8.9) [712, 1312, 1815]	AFP-07 (pIC ₅₀ 8.5) [288], BMY 45778 (pIC ₅₀ 8) [881], esuberaprost (pK _d 7.9) [892], cicaprost (pK _i 7.8) [7]	fluprostenol (pK _i 8.6) [7], latanoprost (free acid form) (pK _i 8.6) [7], AL12180 (pEC ₅₀ 7.7–7.9) [1714], tafluprost [1843]	I-BOP (pK _d 8.9–9.3) [1233], U46619 (pK _i 7.5) [7], STA ₂ (pIC ₅₀ 6.4–7.1) [59]
Antagonists	–	ramatroban (pK _i 7.4) [1815]	–	–	ramatroban (pK _i 8) [1869]
Selective antagonists	laropiprant (pK _i 10.1) [1808] – Unknown, BWA868C (pK _i 8.6–9.3) [171, 606, 2045], S-5751 (pK _i 8.8) [54], ONO-AE3-237 (pK _i 7.7) [758, 1895, 1897]	CAY 10471 (pIC ₅₀ 8.9) [1610, 1927], AZD1981 (pIC ₅₀ 8.4) [1150]	RO1138452 (pK _i 8.7) [158], RO3244794 (pA ₂ 8.5) [158]	AS604872 (pK _i 7.5) [346]	ifetroban (pK _i 8.4–10) [1426], vapiroprost (pK _i 8.3–9.4) [59, 1151], SQ-29548 (pK _i 8.1–9.1) [7, 1834, 2030], ONO-3708 (pK _i 7.4–8.9) [910]
Labelled ligands	[³ H]PGD ₂ (Agonist) (pK _d 7.9–9.5) [2030, 2045]	[³ H]PGD ₂ (Agonist) (pK _d 7.8–8.2) [1216, 1723]	[³ H]iloprost (Agonist) (pK _d 7.7–9) [7, 170, 2030]	[³ H]PGF _{2α} (Agonist) (pK _d 8.1–9) [7, 8, 2030], [³ H](+)-fluprostenol (Agonist) (pK _d 7.5) – Unknown	[¹²⁵ I]SAP (Antagonist) (pK _d 7.7–9.3) [1356], [¹²⁵ I]BOP (Agonist) (pK _d 8.7) [1328], [³ H]SQ-29548 (Antagonist) (pK _d 7.4–8.2) [7, 2030]

	EP ₁ receptor	EP ₂ receptor	EP ₃ receptor	EP ₄ receptor
Nomenclature	EP ₁ receptor	EP ₂ receptor	EP ₃ receptor	EP ₄ receptor
HGNC, UniProt	<i>PTGER1</i> , P34995	<i>PTGER2</i> , P43116	<i>PTGER3</i> , P43115	<i>PTGER4</i> , P35408
Rank order of potency	PGE ₂ > PGF _{2α} , PGI ₂ > PGD ₂ , thromboxane A ₂	PGE ₂ > PGF _{2α} , PGI ₂ > PGD ₂ , thromboxane A ₂	PGE ₂ > PGF _{2α} , PGI ₂ > PGD ₂ , thromboxane A ₂	PGE ₂ > PGF _{2α} , PGI ₂ > PGD ₂ , thromboxane A ₂
Endogenous agonists	PGE ₁ (pK _i 6.8) [1713], PGI ₂ (pK _i 4.8) [1713]	PGE ₂ (pK _i 7.5–8.3) [7, 1799, 2030]	–	–
Agonists	17-phenyl-ω-trinor-PGE ₂ (pK _i 8.1) [1713]	evatanepag (pIC ₅₀ 7.3) [260] – Rat	misoprostol (methyl ester) (EP ₃ -III isoform) (pK _i 6.5) [7]	–
Selective agonists	ONO-DI-004 (pK _i 6.8) [1826] – Mouse	ONO-AE1-259 (pK _i 8.5) [1826] – Mouse, butaprost (free acid form) (pK _i 5.9–7) [7, 1799]	SC46275 (pEC ₅₀ 10.4) [1655] – Guinea pig, MB-28767 (EP ₃ -III isoform) (pK _i 9.9) [7], ONO-AE-248 (pEC ₅₀ 5.6–6.7) [534, 1140]	L902688 (pEC ₅₀ 8.1–10.3) [535, 1064], ONO-AE1-437 (pK _i 9.1) [1294] – Mouse, CP734432 (pIC ₅₀ 8.7) [1529], ONO-AE1-329 (pEC ₅₀ 7.7–7.8) [534, 535]
Antagonists	–	–	–	evatanepag (pK _i 8.6) [1345],
Selective antagonists	ONO-8711 (pK _i 9.2) [1992], GW848687X (pIC ₅₀ 8.6) [605], SC-51322 (pK _i 7.9) [7]	TG4-155 (TG4-155 also has affinity for the human DP1 receptor (pK _B = 7.8)) (pK _B 8.6) [865], TG7-171 (pK _B 8.6) [567], PF-04852946 (pK _B 8.4–8.5) [920], PF-04418948 (PF-04418948 has weaker affinity at the EP2-receptor in guinea-pigs) (pK _B 8.3) [153, 2136]	L-798,106 (EP ₃ -III isoform) (pK _i 7.8–9.7) [888, 890, 1810], L-826266 (EP ₃ -III isoform (pK _i =8.04 in the presence of HSA)) (pK _i 9.1) [890], ONO-AE3-240 (pIC ₅₀ 8.8) [38] – Mouse, DG-041 (pK _i 8.4) [888]	MK-2894 (pK _i 9.2) [7, 161, 350], ONO-AE3-208 (pK _i 8.5), BGC201531 (pK _i 7.9) [1230], ER819762 (pIC ₅₀ 7.2) [304], GW 627368 (pK _i 7–7.1) [2030, 2031]
Labelled ligands	[³ H]PGE ₂ (Agonist) (pK _d 7.6–7.9) [7, 1713, 2030]	[³ H]PGE ₂ (Agonist) (pK _d 7.7–7.9) [7, 2030]	[³ H]PGE ₂ (Agonist) (pK _d 8.2–9.5) [7, 2030]	[³ H]PGE ₂ (Agonist) (pK _d 7.6–9.5) [7, 401, 2019, 2030]

Comments: ramatroban is an antagonist at both DP₂ and TP receptors. Whilst cicaprost is selective for IP receptors, it does exhibit moderate agonist potency at EP₄ receptors [7]. Apart from IP receptors, iloprost also binds to other prostanoid receptors such as EP₁ receptors. The TP receptor exists in α and β isoforms due to alternative splicing of the cytoplasmic tail [1566]. The IP receptor agonist treprostinil binds also to human EP₂ and DP₁ receptors with high affinity (pK_i 8.44 and 8.36, respectively).

The EP₁ agonist 17-phenyl-ω-trinor-PGE₂ also shows agonist activity at EP₃ receptors. Butaprost and SC46275 may require de-esterification within tissues to attain full agonist potency. There is evidence for subtypes of FP [1105], IP [1851, 2037] and TP [1005] receptors. mRNA for the EP₁ and EP₃ receptors undergo alternative

splicing to produce two [1441] and at least six variants, respectively, which can interfere with signalling [1441] or generate complex patterns of G-protein (G_{i/o}, G_{q/11}, G_s and G_{12,13}) coupling (e.g. [997, 1370]). The number of EP₃ receptor (protein) variants are variable depending on species, with five in human, three in rat and three in mouse. The possibility of additional receptors for the isoprostanes has been suggested [1531]. Putative receptor(s) for prostamide F (which as yet lack molecular correlates) and which preferentially recognize PGF₂-1-ethanolamide and its analogues (e.g. Bimatoprost) have been identified, together with moderate-potency antagonists (e.g. AGN 211334) [2042].

The free acid form of AL-12182, AL12180, used in *in vitro* studies, has a EC₅₀ value of 15nM which is the concentration of the compound

giving half-maximal stimulation of inositol phosphate turnover in HEK-293 cells expressing the human FP receptor [1714].

References given alongside the TP receptor agonists I-BOP [1233] and STA₂ [59] use human platelets as the source of TP receptors for competition radio-ligand binding assays to determine the indicated activity values.

Pharmacological evidence for a second IP receptor, denoted IP₂, in the central nervous system [1851, 1994] and in the BEAS-2B human airway epithelial cell line [2033] is available. This receptor is selectively activated by 15R-17,18,19,20-tetranor-16-m-tolyl-isocarbacyclin (15R-TIC) and 15R-Deoxy 17,18,19,20-tetranor-16-m-tolyl-isocarbacyclin (15-deoxy-TIC). However, molecular biological evidence for the IP₂ subtype is currently lacking.

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Proteinase-activated receptors

G protein-coupled receptors → Proteinase-activated receptors

Overview: Proteinase-activated receptors (PARs, **nomenclature as agreed by the NC-IUPHAR Subcommittee on Proteinase-activated Receptors [770]**) are unique members of the GPCR superfamily activated by proteolytic cleavage of their amino terminal exodomains. Agonist proteinase-induced hydrolysis unmasks a tethered ligand (TL) at the exposed amino terminus, which acts intramolecularly at the binding site in the body of

the receptor to effect transmembrane signalling. TL sequences at human PAR1-4 are SFLLRN-NH₂, SLIGKV-NH₂, TFRGAP-NH₂ and GYPGQV-NH₂, respectively. With the exception of PAR3, these synthetic peptide sequences (as carboxyl terminal amides) are able to act as agonists at their respective receptors. Several proteinases, including neutrophil elastase, cathepsin G and chymotrypsin can have inhibitory effects at PAR1 and PAR2 such that they cleave the ex-

odomain of the receptor without inducing activation of Gαq-coupled calcium signalling, thereby preventing activation by activating proteinases but not by agonist peptides. Neutrophil elastase cleavage of PAR2 can however activate MAP kinase signaling by exposing a TL that is different from the one revealed by trypsin [1553]. The role of such an action *in vivo* is unclear.

Nomenclature	PAR1	PAR2	PAR3	PAR4
HGNC, UniProt	F2R, P25116	F2RL1, P55085	F2RL2, O00254	F2RL3, Q96R10
Agonist proteases	thrombin (F2, P00734), activated protein C (PROC, P04070), matrix metalloproteinase 1 (MMP1, P45452), matrix metalloproteinase 13 (MMP13, P45452) [70]	Trypsin, tryptase, TF/VIIa, Xa	thrombin (F2, P00734)	thrombin (F2, P00734), trypsin, cathepsin G (CTSG, P08311)
Selective agonists	TFLLR-NH ₂ (pEC ₅₀ 5.4) [340]	GB110 (pEC ₅₀ 6.5) [98], 2-furoyl-LIGRLO-amide (pK _i 5.4) [1243], SLIGKV-NH ₂ [1069], SLIGRL-NH ₂ [1069]	–	AYPGKF-NH ₂ , GYPGKF-NH ₂ , GYPGQV-NH ₂

(continued)	PAR1	PAR2	PAR3	PAR4
Nomenclature	PAR1	PAR2	PAR3	PAR4
Selective antagonists	vorapaxar (pK _i 8.1) [281], atopaxar (pIC ₅₀ 7.7) [978], RWJ-56110 (pIC ₅₀ 6.4) [48]	GB88 (pIC ₅₀ 5.7) [1813], P2pal18s [1705]	–	–
Labelled ligands	[³ H]haTRAP (Agonist) (pK _d 7.8) [15]	2-furoyl-LIGRL[N-(Alexa Fluor 594)-O]-NH ₂ (Agonist) [771], 2-furoyl-LIGRL[N- ³ H]propionyl]-O-NH ₂ (Agonist) [771], [³ H]2-furoyl-LIGRL-NH ₂ (Selective Agonist) [903], trans-cinnamoyl-LIGRLO [N- ³ H]propionyl]-NH ₂ (Agonist) [28]	–	–
Comments	TFLLR-NH ₂ is selective relative to the PAR ₂ receptor [155, 915].	2-Furoyl-LIGRLO-NH ₂ activity was measured via calcium mobilisation in HEK 293 cells which constitutively coexpress human PAR ₁ and PAR ₂ .	–	–

Comments: thrombin (F2, P00734) is inactive at the PAR₂ receptor.

Endogenous serine proteases (EC 3.4.21.) active at the proteinase-activated receptors include: thrombin (F2, P00734), generated by the action of Factor X (F10, P00742) on liver-derived prothrombin (F2, P00734); trypsin, generated by the action of enterokinase (TMPRSS15, P98073) on pancreatic-derived trypsinogen (PRSS1, P07477); tryptase, a family of enzymes (α/β 1 TPSAB1, Q15661 ; γ 1 TPSG1, Q9NRR2; δ 1 TPSD1, Q9BZJ3) secreted from mast cells; cathepsin G (CTSG, P08311) generated from leukocytes; liver-derived protein C (PROC, P04070) generated in plasma by thrombin (F2, P00734) and matrix metalloproteinase 1 (MMP1, P45452).

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QRFP receptor

G protein-coupled receptors → QRFP receptor

Overview: The human gene encoding the QRFP receptor (QRFP, also known as the peptide P518 receptor), previously designated as an orphan GPCR receptor was identified in 2001 by Lee *et al.* from a hypothalamus cDNA library [1066]. However, the reported cDNA (AF411117) is a chimera with bases 1-127 derived from chromosome 1 and bases 155-1368 derived from chromosome 4. When corrected, QRFP (also referred to as SP9155 or AQ27) encodes a 431 amino acid protein that shares sequence similarities in the transmembrane spanning regions with other peptide receptors. These include neuropeptide FF2 (38%), neuropeptide Y2 (37%) and galanin GalR1 (35%) receptors.

Nomenclature	QRFP receptor
HGNC, UniProt	QRFP, Q96P65
Endogenous agonists	QRFP43 (QRFP, P83859) (pIC ₅₀ 7.8–9.3) [557, 1850] – Rat, QRFP26 (QRFP) (pEC ₅₀ 8.2) [867]
Labelled ligands	[¹²⁵ I]QRFP43 (human) (Agonist) (pK _d 7.8–10.3) [557, 1017, 1850]

Comments: The orphan receptor *GPR83* (9NYM4) shows sequence similarities with the QRFP receptor, as well as with the NPFF1, NPFF2, and PrRP receptors.

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Relaxin family peptide receptors

G protein-coupled receptors → Relaxin family peptide receptors

Overview: Relaxin family peptide receptors (RXFP, **nomenclature as agreed by the NC-IUPHAR Subcommittee on Relaxin family peptide receptors** [105, 677]) may be divided into two pairs, RXFP1/2 and RXFP3/4. Endogenous agonists at these receptors are a number of heterodimeric peptide hormones analogous to insulin: relaxin-1 (RLN1, P04808), relaxin (RLN2, P04090), relaxin-3 (RLN3, Q8WXF3) (also known as INSL7), insulin-like pep-

ptide 3 (INSL3 (INSL3, P51460)) and INSL5 (INSL5, Q9Y5Q6).

Species homologues of relaxin have distinct pharmacology - relaxin (RLN2, P04090) interacts with RXFP1, RXFP2 and RXFP3, whereas mouse and rat relaxin selectively bind to and activate RXFP1 [1686] and porcine relaxin may have a higher efficacy than human relaxin (RLN2, P04090) [678]. Relaxin-3 (RLN3, Q8WXF3) has differential affinity for RXFP2 receptors between species; mouse and rat RXFP2

have a higher affinity for relaxin-3 (RLN3, Q8WXF3) [1685]. At least two binding sites have been identified on the RXFP1 and RXFP2 receptors: a high-affinity site in the leucine-rich repeat region of the ectodomain and a somewhat lower-affinity site located in the surface loops of the transmembrane domain [678, 1812]. The unique N-terminal LDLa module of RXFP1 and RXFP2 is essential for receptor signalling [1687].

Nomenclature	RXFP1 receptor	RXFP2 receptor	RXFP3 receptor	RXFP4 receptor
HGNC, UniProt	<i>RXFP1</i> , Q9HBX9	<i>RXFP2</i> , Q8WXD0	<i>RXFP3</i> , Q9NSD7	<i>RXFP4</i> , Q8TDU9
Rank order of potency	relaxin (<i>RLN2</i> , P04090) = relaxin-1 (<i>RLN1</i> , P04808) > relaxin-3 (<i>RLN3</i> , Q8WXF3) [1812]	INSL3 (<i>INSL3</i> , P51460) > relaxin (<i>RLN2</i> , P04090) ≫ relaxin-3 (<i>RLN3</i> , Q8WXF3) [1022, 1812]	relaxin-3 (<i>RLN3</i> , Q8WXF3) > relaxin-3 (B chain) (<i>RLN3</i> , Q8WXF3) > relaxin (<i>RLN2</i> , P04090) [1119]	INSL5 (<i>INSL5</i> , Q9Y5Q6) = relaxin-3 (<i>RLN3</i> , Q8WXF3) > relaxin-3 (B chain) (<i>RLN3</i> , Q8WXF3) [1117, 1118]
Endogenous antagonists	–	–	INSL5 (<i>INSL5</i> , Q9Y5Q6) (p <i>K_i</i> 7) [2129]	–
Antagonists	B-R13/17K H2 relaxin (pEC ₅₀ 5.7–6.7) [788, 1382], LGR7-truncate [1687]	–	R3(BΔ23-27)R/I5 chimeric peptide (pIC ₅₀ 9.2) [1018]	R3(BΔ23-27)R/I5 chimeric peptide (pIC ₅₀ 8–8.6) [714, 1018]
Selective antagonists	–	A(9-26)INSL3 (p <i>K_i</i> 9.1) [787], A(10-24)INSL3 (p <i>K_i</i> 8.7) [787], A(C10/15S)INSL3 (p <i>K_i</i> 8.6) [2118], INSL3 B chain dimer analogue 8 (p <i>K_i</i> 8.5) [1710], A(Δ10/15C)INSL3 (p <i>K_i</i> 8.3) [2118], cyclic INSL3 B-chain analogue 6 (p <i>K_i</i> 6.7) [1708], INSL3 B-chain analogue (p <i>K_i</i> 5.1) [411], (des 1-8) A-chain INSL3 analogue [253]	minimised relaxin-3 analogue 3 (p <i>K_i</i> 7.6) [1706], R3-B1-22R (pIC ₅₀ 7.4) [714]	minimised relaxin-3 analogue 3 (pIC ₅₀ 6.6) [1706]
Selective allosteric modulators	ML290 (Agonist) (pEC ₅₀ 7) [2057, 2060]	–	–	–
Labelled ligands	[³³ P]relaxin (human) (Agonist) (p <i>K_d</i> 9.3–9.7) [678, 1812], [¹²⁵ I]relaxin (human) (Agonist)	[¹²⁵ I]INSL3 (human) (Agonist) (p <i>K_d</i> 10) [1340], [³³ P]relaxin (human) (Agonist) (p <i>K_d</i> 9–9.2) [678, 1812]	[¹²⁵ I]relaxin-3 (human) (Agonist) (p <i>K_d</i> 9.5) [1119], [¹²⁵ I]relaxin-3-B/INSL5 A chimera (Agonist) (p <i>K_d</i> 9.3) [1117]	[¹²⁵ I]relaxin-3 (human) (Agonist) (p <i>K_d</i> 8.7–9.7) [1118], [¹²⁵ I]relaxin-3-B/INSL5 A chimera (Agonist) (p <i>K_d</i> 8.9) [1117], europium-labelled INSL5 (p <i>K_d</i> 8.3) [714]
Comments	europium-labelled relaxin is a fluorescent ligand for this receptor (<i>K_d</i> =0.5nM) [1707].	europium-labelled INSL3 is a fluorescent ligand for this receptor (<i>K_d</i> =1nM) [1709].	europium-labelled relaxin-3-B/INSL5 A chimera and R3-B1-22R are fluorescent ligands for this receptor (<i>K_d</i> =5nM and 28nM) [714, 715].	europium-labelled relaxin-3-B/INSL5 A chimera is a fluorescent probe at this receptor (<i>K_d</i> =5nM) [714]. europium-labelled mouse INSL5 is a fluorescent ligand at this receptor (<i>K_d</i> =5nM) [120].

Comments: Relaxin has recently successfully completed a Phase III clinical trial for the treatment of acute heart failure. 48 hr infusion of relaxin reduced dyspnoea and 180 day mortality [1262]. Small molecule agonists active at RXFP1 receptors have been developed [1718, 2060], and one of these (ML290) is an allosteric agonist at

RXFP1 [2060]. The antifibrotic actions of relaxin are dependent on the angiotensin receptor AT₂, are absent in AT₂ knockout mice, and are associated with heterodimer formation between RXFP1 and AT₂ [330]. Mutations in *INSL3* and *LGR8* (RXFP2) have been reported in populations of patients with cryptorchidism [512]. Numerous splice

variants of the human RXFP1 and RXFP2 receptors have been identified, most of which do not bind relaxin family peptides [1340]. Splice variants of RXFP1 encoding the N-terminal LDLa module act as antagonists of RXFP1 signalling [1685, 1687]. cAMP elevation appears to be a major signalling pathway for RXFP1 and RXFP2 [795, 796],

but RXFP1 also activates MAP kinases, nitric oxide signalling, tyrosine kinase phosphorylation and relaxin can interact with glucocorticoid receptors [681]. RXFP1 signalling involves lipid rafts, residues in the C-terminus of the receptor and activation of phosphatidylinositol-3-kinase [682]. More recent studies provide evidence that RXFP1 is pre-assembled in signalosomes with other signalling proteins including G_{α_s} , $G_{\beta\gamma}$ and adenylyl cyclase 2 that display constitutive activity and are exquisitely sensitive to sub-picomolar concentrations of relaxin [679]. The cyclic AMP signalling pattern is highly dependent on the cell type in which RXFP1 is expressed [680].

The receptor expression profiles suggested that RXFP3 was a neuropeptide receptor and RXFP4 a gut hormone receptor. Studies in rats and mice (including wildtype, and relaxin-3 and RXFP3 gene-deletion strains [671, 782, 1759, 1971] have revealed putative roles for the relaxin-3/RXFP3 system in the modulation of feeding [564, 566, 714, 1706, 1760], anxiety [1618, 2114], and reward and motivated, goal-directed behaviours [782, 1619, 1971], particularly in relation to the integration of stress and corticotrophin-releasing factor signalling [1162], with implications for the therapeutic treatment

of clinical anxiety, depression, eating disorders and addiction (see [565, 1761] for review). Relaxin-3 (*RLN3*, *Q8WXF3*) acts as an agonist at both RXFP3 and RXFP4 whereas *INSL5* (*INSL5*, *Q9YSQ6*) is an agonist at RXFP4 and a weak antagonist at RXFP3. Unlike RXFP1 and RXFP2 both RXFP3 and RXFP4 are encoded by a single exon and therefore no splice variants exist. The rat RXFP3 sequence has two potential start codons that encode RXFP3L and RXFP3S with the longer variant having an additional 7 amino-acids at the N-terminus. It is not known which variant is expressed. Rat and dog RXFP4 sequences are pseudogenes [2027]. Recent studies suggest that *INSL5* is an incretin secreted from enteroendocrine L cells and that the *INSL5*/RXFP4 system has roles in controlling food intake and glucose homeostasis [652]. RXFP3 couples to $G_{i/o}$ and inhibits adenylyl cyclase [1119, 2144], and also causes Erk1/2 phosphorylation [2144]. Relatively little is known about RXFP4 signalling but like RXFP3 it couples to inhibitory $G_{i/o}$ G-proteins [1120]. Recent studies suggest that relaxin (*RLN2*, *P04090*) also interacts with RXFP3 to cause a pattern of activation of signalling pathways that are a subset of those activated by relaxin-3 (*RLN3*, *Q8WXF3*). The

two patterns of signaling observed in several cell types expressing RXFP3 are strong inhibition of forskolin-stimulated cyclic AMP accumulation, ERK1/2 activation and nuclear factor $\text{NF}\kappa\text{-B}$ reporter gene activation with relaxin-3 (*RLN3*, *Q8WXF3*), and weaker activity with relaxin (*RLN2*, *P04090*), porcine relaxin, or insulin-like peptide 3 (*INSL3* (*INSL3*, *P51460*)) and a strong stimulation of activator protein (AP)-1 reporter genes with relaxin (*RLN2*, *P04090*), and weaker activation with relaxin-3 (*RLN3*, *Q8WXF3*) or porcine relaxin [2144]. Thus at RXFP3, relaxin (*RLN2*, *P04090*) is a biased ligand compared to the cognate ligand relaxin-3 (*RLN3*, *Q8WXF3*). Two pharmacologically distinct ligand binding sites were also identified on RXFP3-expressing cells using [125 I]relaxin-3-B/*INSL5* A chimera which binds with high affinity and displays competition by relaxin-3 (*RLN3*, *Q8WXF3*) or a relaxin-3 (B chain) (*RLN3*, *Q8WXF3*) peptide, and [125 I]relaxin (human) which displays competition by relaxin (*RLN2*, *P04090*), relaxin-3 (*RLN3*, *Q8WXF3*), or *INSL3* (*INSL3*, *P51460*) and weakly by porcine relaxin.

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Somatostatin receptors

G protein-coupled receptors → Somatostatin receptors

Overview: Somatostatin (somatotropin release inhibiting factor) is an abundant neuropeptide, which acts on five subtypes of somatostatin receptor (*sst*₁₋₅; **nomenclature as agreed by the NC-IUPHAR Subcommittee on Somatostatin Receptors** [790]). Activation of these receptors produces a wide range

of physiological effects throughout the body including the inhibition of secretion of many hormones. The relationship of the cloned receptors to endogenously expressed receptors is not yet well established in some cases. Endogenous ligands for these receptors are somatostatin-14 (*SRIF-14* (*SST*, *P61278*)) and somatostatin-28

(*SRIF-28* (*SST*, *P61278*)). *Cortistatin-14* {Mouse, Rat} has also been suggested to be an endogenous ligand for somatostatin receptors [404].

Nomenclature	sst₁ receptor	sst₂ receptor	sst₃ receptor	sst₄ receptor	sst₅ receptor
HGNC, UniProt	SSTR1, P30872	SSTR2, P30874	SSTR3, P32745	SSTR4, P31391	SSTR5, P35346
Agonists	pasireotide (pIC ₅₀ 8) [1669]	vapreotide (pK _i 8.3–10.1) [233 , 1474], pasireotide (pIC ₅₀ 9) [1669]	pasireotide (pIC ₅₀ 8.8) [1669], vapreotide (pK _i 7.4–7.9) [233 , 1474 , 1738]	NNC269100 (pK _i 8.2) [1132]	pasireotide (pIC ₅₀ 9.8) [1669], vapreotide (pK _i 7.3–9.2) [233 , 1265 , 1474 , 1736 , 1737 , 1738]
Selective agonists	L-797,591 (pK _i 8.8) [1595], Des-Ala^{1,2,5}-[D-Trp⁸, IAmp⁹]SRIF (pIC ₅₀ 7.5) [484]	L-054,522 (pK _i 11) [2084], BIM 23027 (pIC ₅₀ 10.9) [271], seglitide (pK _i 8.8–10.3) [233 , 1474 , 1736 , 1737 , 1738 , 2084], octreotide (pK _i 8.7–9.9) [233 , 1474 , 1736 , 1737 , 1738 , 2084]	L-796,778 (pK _i 7.6) [1595]	L-803,087 (pK _i 9.2) [1595]	BIM 23052 (pK _i 7.4–9.6) [1265 , 1736 , 1737 , 1738], L-817,818 (pK _i 9.4) [1595], BIM 23268 (pK _i 8.7) [1265]
Selective antagonists	SRA880 (pK _d 8–8.1) [792]	[D-Tyr⁸]CYN 154806 (pK _d 8.1–8.9) [1412]	NVP ACQ090 (pK _i 7.9) [793]	–	–
Labelled ligands	–	[¹²⁵I]Tyr³ SMS 201-995 (Agonist) (pK _d 9.9) [1736 , 1737], [¹²⁵I]BIM23027 (Agonist) (pIC ₅₀ 9.7) [772] – Rat	–	–	[¹²⁵I]Tyr³ SMS 201-995 (Agonist) (pK _d 9.6) [1736 , 1737]
Comments	–	–	Troxler <i>et al.</i> (2010) describe the identification of non-peptidic, subtype-selective sst ₃ receptor antagonists [1907].	–	–

Comments: [¹²⁵I]Tyr¹¹-SRIF-14, [¹²⁵I]LTT-SRIF-28, [¹²⁵I]CGP 23996 and [¹²⁵I]Tyr¹⁰-CST14 may be used to label somatostatin receptors nonselectively. A number of nonpeptide subtype-selective agonists have been synthesised [[1595](#)]. A novel peptide somatostatin analogue, [somatoprim](#), has affinity for sst₂, sst₄ and sst₅ receptors and is a potent inhibitor of GH secretion [[1514](#), [1726](#)].

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Succinate receptor

G protein-coupled receptors → Succinate receptor

Overview: Nomenclature as recommended by NC-IUPHAR [396].

Nomenclature	succinate receptor
HGNC, UniProt	<i>SUCNR1</i> , <i>Q9BXA5</i>
Endogenous agonists	succinic acid (pEC ₅₀ 3.1–4.7) [728, 1785]

Tachykinin receptors

G protein-coupled receptors → Tachykinin receptors

Overview: Tachykinin receptors (**provisional nomenclature as recommended by NC-IUPHAR [530]**) are activated by the endogenous peptides substance P (*TAC1*, P20366) (SP), neurokinin A (*TAC1*, P20366) (NKA; previously known as substance K, neurokinin α , neuromedin L), neurokinin B (*TAC3*, Q9UHF0) (NKB; previously

known as neurokinin β , neuromedin K), neuropeptide K (*TAC1*, P20366) and neuropeptide γ (*TAC1*, P20366) (N-terminally extended forms of neurokinin A). The neurokinins (A and B) are mammalian members of the tachykinin family, which includes peptides of mammalian and nonmammalian origin containing the consensus se-

quence: Phe-x-Gly-Leu-Met. Marked species differences in *in vitro* pharmacology exist for all three receptors, in the context of nonpeptide ligands.

Nomenclature	NK ₁ receptor	NK ₂ receptor	NK ₃ receptor
HGNC, UniProt	<i>TAC1</i> , P25103	<i>TACR2</i> , P21452	<i>TACR3</i> , P29371
Rank order of potency	substance P (<i>TAC1</i> , P20366) > neurokinin A (<i>TAC1</i> , P20366) > neurokinin B (<i>TAC3</i> , Q9UHF0)	neurokinin A (<i>TAC1</i> , P20366) > neurokinin B (<i>TAC3</i> , Q9UHF0) >> substance P (<i>TAC1</i> , P20366)	neurokinin B (<i>TAC3</i> , Q9UHF0) > neurokinin A (<i>TAC1</i> , P20366) > substance P (<i>TAC1</i> , P20366)
Agonists	substance P-OMe (pIC ₅₀ 7.4–7.5) [1882]	–	–
Selective agonists	[Sar ⁹ ,Met(O ₂) ¹¹]SP (pIC ₅₀ 9.7–9.9) [1882], septide (pK _i 7–9.3) [125, 711], [Pro ⁹]SP (pIC ₅₀ 8.6) [1896] – Rat	[Lys ⁵ ,Me-Leu ⁹ ,Nle ¹⁰]NKA-(4-10) (pIC ₅₀ 8.8–9.4) [1229] – Rat, GR64349 (pEC ₅₀ 8.4) [407] – Rat, [β Ala ⁸]neurokinin A-(4-10) (pK _d 6) [477]	[Phe(Me) ⁷]neurokinin B (pK _i 8.7–9.6) [1644, 1645], senktide (pK _i 7.1–8.6) [1644, 1645, 1882]

(continued)	NK ₁ receptor	NK ₂ receptor	NK ₃ receptor
Nomenclature			
Selective antagonists	aprepitant (pK _i 10.1) [673, 674], lanepitant (pK _i 9.8–10) [613], lanepitant (pI _C ₅₀ 9.8) [798], CP 99994 (pK _i 9.3–9.7) [50, 1645], casopitant (pK _i 9.4) [798, 1905], vestipitant (pK _i 9.4) [221, 418], nolpitanium (pI _C ₅₀ 8.9–9) [1882], RP67580 (pI _C ₅₀ 7.7) [528]	GR94800 (pK _i 9.8) [200], saredutant (pK _i 9.4–9.7) [50, 477, 1645], GR 159897 (pK _d 7.8–9.5) [133, 477, 1770], MEN10627 (pK _i 9.2) [603], nepadutant (pK _i 8.5–8.7) [272, 343]	osanetant (pK _i 8.4–9.7) [50, 110, 342, 476, 898, 1450, 1644, 1645, 1882], talnetant (pK _i 7.4–9) [129, 604, 1644, 1645], PD157672 (pI _C ₅₀ 7.8–7.9) [165, 1882]
Labelled ligands	[¹²⁵ I]L703,606 (Antagonist) (pK _d 9.5) [537], [¹²⁵ I]BH-[Sar ⁹ ,Met(O ₂) ¹¹]SP (Agonist) (pK _d 9) [1901] – Rat, [³ H]BH-[Sar ⁹ ,Met(O ₂) ¹¹]SP (Agonist) (pK _d 8.7) [1902] – Rat, [³ H]SP (human, mouse, rat) (Agonist) (pK _d 8.6) [80], [¹²⁵ I]SP (human, mouse, rat) (Agonist), [¹⁸ F]SPA-RQ (Antagonist) [317]	[³ H]saredutant (Antagonist) (pK _d 9.7) [649] – Rat, [¹²⁵ I]NKA (human, mouse, rat) (Agonist) (pK _d 9.3) [1990], [³ H]GR100679 (Antagonist) (pK _d 9.2) [669]	[³ H]osanetant (Antagonist) (pK _d 9.9), [³ H]senktide (Agonist) (pK _d 8.1–8.7) [660] – Guinea pig, [¹²⁵ I][MePhe ⁷]NKB (Agonist)

Comments: The NK₁ receptor has also been described to couple to other G proteins [1606]. The hexapeptide agonist septide appears to bind to an overlapping but non-identical site to substance P (TAC1, P20366) on the NK₁ receptor. There are suggestions for additional subtypes of tachykinin receptor; an orphan receptor (SwissProt P30098) with structural similarities to the NK₃ receptor was found to respond to NKB when expressed in *Xenopus* oocytes or Chinese hamster ovary cells [433, 1004].

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Thyrotropin-releasing hormone receptors

G protein-coupled receptors → Thyrotropin-releasing hormone receptors

Overview: Thyrotropin-releasing hormone (TRH) receptors (**provisional nomenclature as recommended by NC-IUPHAR [530]**) are activated by the endogenous tripeptide TRH (*TRH*, P20396) (pGlu-His-ProNH₂). TRH (*TRH*, P20396) and TRH analogues fail to distinguish TRH₁ and TRH₂ receptors [1822]. [³H]TRH (human, mouse, rat) is able to label both TRH₁ and TRH₂ receptors with K_d values of 13 and 9 nM respectively.

Nomenclature	TRH ₁ receptor	TRH ₂ receptor
HGNC, UniProt	<i>TRHR</i> , P34981	–
Antagonists	diazepam (pK _i 5.2) [444] – Rat	–
Selective antagonists	midazolam (pK _i 5.5) [444] – Rat, chlordiazepoxide (pK _i 4.8) [444] – Rat, chlordiazepoxide (pK _i 4.7) [1804] – Mouse	–
Comments	–	A class A G protein-coupled receptor: not present in man

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Trace amine receptor

G protein-coupled receptors → Trace amine receptor

Overview: Trace amine-associated receptors were initially discovered as a result of a search for novel 5-HT receptors [185], where 15 mammalian orthologues were identified and divided into two families. The TA₁ receptor (**nomenclature as agreed by the NC-IUPHAR Subcommittee for the Trace amine recep-**

tor [1181]) has been shown to have affinity for the endogenous trace amines tyramine, β-phenylethylamine and octopamine in addition to the classical amine dopamine [185]. Emerging evidence suggests that TA₁ is a modulator of monoaminergic activity in the brain [2062] with TA₁ and dopamine D₂ receptors shown to form

constitutive heterodimers when co-expressed [492]. In addition to trace amines, receptors can be activated by amphetamine-like psychostimulants, and endogenous thyronamines such as thyronamine and 3-iodothyronamine.

Nomenclature	TA ₁ receptor
HGNC, UniProt	TAAR1, Q96RJO
Rank order of potency	tyramine > β-phenylethylamine > octopamine = dopamine [185]
Agonists	RO5166017 (pEC ₅₀ 7.3) [1574]
Antagonists	EPPTB (Inverse agonist) (pIC ₅₀ 5.1) [199]
Labelled ligands	[³ H]tyramine (Agonist) (pK _d 7.7) [185]

Comments: In addition to TA₁, analysis has shown that in man there are up to 5 functional TAAR genes (TAAR2,5,6,8,9). See [185] for detailed discussion. The product of the gene TAAR2 (also known as GPR58) appears to respond to β-phenylethylamine > tyramine and to couple through G_S [185].

TAAR3, in some individuals, and TAAR4 are pseudogenes in man, although functional in rodents. The signalling characteristics and pharmacology of TAA₅ (PNR, Putative Neurotransmitter Receptor: TAAR5, O14804), TAA₆ (Trace amine receptor 4, TaR-4: TAAR6, 96RI8), TAA₈ (Trace amine receptor 5, GPR102: TAAR8, Q969N4) and TAA₉ (trace amine associated receptor 9: TAAR9, 96RI9) are lacking. The thy-

ronamines, endogenous derivatives of thyroid hormone, have been shown to have affinity for rodent cloned trace amine receptors, including TA₁ [1657]. An antagonist EPPTB has recently been described that has a pK_i of 9.1 at the mouse TA₁ but less than 5.3 for human TA₁ [1792].

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Urotensin receptor

G protein-coupled receptors → Urotensin receptor

Overview: The urotensin-II (U-II) receptor (UT, **nomenclature as agreed by the NC-IUPHAR Subcommittee on the Urotensin receptor [439, 530, 1952]**) is activated by the endogenous dodecapeptide urotensin-II (*UTS2*, *O95399*), originally isolated from the urophysis, the endocrine organ of the caudal neurosecretory system of teleost fish [134]. Several structural forms of U-II exist in fish and amphibians. The Goby orthologue was used to

identify U-II as the cognate ligand for the predicted receptor encoded by the rat gene *gpr14* [375, 1130, 1327, 1410]. Human urotensin-II (*UTS2*, *O95399*), an 11-amino-acid peptide [375], retains the cyclohexapeptide sequence of goby U-II that is thought to be important in ligand binding [219, 957]. This sequence is also conserved in the deduced amino-acid sequence of rat urotensin-II {Rat} (14 amino-acids) and mouse urotensin-II {Mouse} (14 amino-acids), although

the N-terminal is more divergent from the human sequence [374]. A second endogenous ligand for UT has been discovered in rat [1816]. This is the urotensin II-related peptide (*UTS2B*, *Q76510*), an octapeptide that is derived from a different gene, but shares the C-terminal sequence (CFWKYCV) common to U-II from other species. Identical sequences to rat urotensin II-related peptide (*UTS2B*, *Q76510*) are predicted for the mature mouse and human peptides.

Nomenclature	UT receptor
HGNC, UniProt	<i>UTS2R</i> , <i>Q9UKP6</i>
Endogenous agonists	urotensin II-related peptide (<i>UTS2B</i> , <i>Q76510</i>) (pK _d 9.6) [1179], urotensin-II (<i>UTS2</i> , <i>O95399</i>) (pK _i 8.6) [440, 475, 647]
Selective agonists	[Pen5]-U (4-11) (human) (pK _i 9.7) [647], U-II-(4-11) (human) (pK _i 9.6) [647], <i>FL104</i> (pEC ₅₀ 5.8–7.5) [1075, 1077], <i>AC-7954</i> (pK _i 6.6) [382, 1076]
Selective antagonists	urantide (pK _i 8.3) [1469], <i>SB-706375</i> (pK _i 8) [440], palosuran (pIC ₅₀ 7.1) [353], <i>SB-611812</i> (pK _i 6.6) [1550]
Labelled ligands	[¹²⁵ I]U-II (human) (Agonist) (pK _d 9.4–9.6) [42, 1179]

Comments: In human vasculature, human urotensin-II (*UTS2*, *O95399*) elicits both vasoconstrictor (pD₂ 9.3-10.1, [1179]) and vasodilator (pIC₅₀ 10.3-10.4, [1800]) responses.

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Vasopressin and oxytocin receptors

G protein-coupled receptors → Vasopressin and oxytocin receptors

Overview: Vasopressin (AVP) and oxytocin (OT) receptors (**nomenclature as recommended by NC-IUPHAR [530]**) are activated by the endogenous cyclic nonapeptides vasopressin (*AVP*, *P01185*) and oxytocin (*OXT*, *P01178*). These peptides are derived from precursors which also produce neurophysins (neurophysin I for oxytocin; neurophysin II for vasopressin).

Searchable database: <http://www.guidetopharmacology.org/index.jsp>

Full Contents of ConciseGuide: <http://onlinelibrary.wiley.com/doi/10.1111/bph.13348/full>

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Nomenclature	V _{1A} receptor	V _{1B} receptor	V ₂ receptor	OT receptor
HGNC, UniProt	AVPR1A, P37288	AVPR1B, P47901	AVPR2, P30518	OXTR, P30559
Rank order of potency	vasopressin (AVP, P01185) > oxytocin (OXT, P01178)	vasopressin (AVP, P01185) > oxytocin (OXT, P01178)	vasopressin (AVP, P01185) > oxytocin (OXT, P01178)	oxytocin (OXT, P01178) > vasopressin (AVP, P01185)
Endogenous agonists	vasopressin (AVP, P01185) (pK _i 8.5–9.3) [24, 311, 369, 415, 1359, 1501, 1627, 1839, 1840, 1870, 1871, 2073]	vasopressin (AVP, P01185) (pK _i 9–9.5) [24, 311, 415, 648, 1359, 1627, 1839, 1840, 1871, 2073]	vasopressin (AVP, P01185) (pK _i 7.9–9.1) [24, 311, 319, 415, 1359, 1627, 1700, 1839, 1840, 1871, 2073]	oxytocin (OXT, P01178) (pK _i 8.2–9.6) [24, 319, 320, 345, 648, 853]
Selective agonists	F180 (pK _d 7.9–8.3) [49, 369]	d[Leu ⁴]LVP (pK _i 9.8) [1485], d[Cha ⁴]AVP (pK _i 9–9.7) [415, 648]	VNA932 (pIC ₅₀ 7.1) [501], OPC-51803 (pK _i 7) [1359], d[Val ⁴ , DArg ⁸]VP	[Thr ⁴ , Gly ⁷]OT (pK _i 8.2–8.4) [320, 472, 853]
Antagonists	conivaptan (pK _i 8.2–8.4) [1839, 1840]	nelivaptan (pK _i 8.4–9.3) [644, 648, 1702]	–	L-371,257 (pK _i 8.8) [648]
Selective antagonists	relcovaptan (pK _i 8.1–9.3) [24, 369, 648, 1501, 1700, 1839, 1870, 1871, 1910], d(CH ₂) ₅ [Tyr(Me) ² , Arg ⁸]VP (pK _i 9)	–	conivaptan (pK _i 9.4) [381], tolvaptan (pK _i 9.4) [2073], satavaptan (pK _i 8.4–9.3) [24, 369, 370, 1699, 1700, 1839, 1910], lixivaptan (Inverse agonist) (pK _i 8.9–9.2) [33, 1700], d(CH ₂) ₅ [D-Ile ² , Ile ⁴]AVP (pK _i 6.9–8.4) [1700], mozavaptan (Inverse agonist) (pK _i 7.4–8.1) [370, 1700, 1839, 1871, 2073, 2074]	SSR126768A (pK _i 8.8–9.1) [1701], desGlyNH ₂ -d(CH ₂) ₅ [Tyr(Me) ² , Thr ⁴ , Orn ⁸]OT (pK _i 8.5), L-372662 (pK _i 8.4) [121]
Labelled ligands	[¹²⁵ I]OH-LVA (Antagonist) (pK _d 10.3–10.4) [319, 369, 1501], [³ H]AVP (human, mouse, rat) (Agonist) (pK _d 8.6–10.2) [208, 319, 369, 370, 1359, 1501, 1627, 1839, 1840, 1870, 1871, 1910, 2073], [³ H]d(CH ₂) ₅ [Tyr(Me) ²]AVP (Antagonist) (pK _d 9)	[³ H]AVP (human, mouse, rat) (Agonist) (pK _d 8.6–9.6) [208, 319, 369, 370, 1359, 1501, 1627, 1839, 1840, 1870, 1871, 1910, 2073]	[³ H]AVP (human, mouse, rat) (Agonist) (pK _d 8.4–9.4) [319, 369, 370, 1359, 1627, 1839, 1840, 1871, 1910, 2073], [³ H]dDAVP (Agonist) (pK _d 7.2–9.1) [319, 370, 1871], [³ H]desGly-NH ₂ [D-Ile ² , Ile ⁴]VP (pK _d 8.6)	[¹²⁵ I]d(CH ₂) ₅ [Tyr(Me) ² , Thr ⁴ , Orn ⁸ , Tyr-NH ₂ ⁹]OVT (Antagonist) (pK _d 10), [³ H]OT (human, mouse, rat) (Agonist) (pK _d 8.2–9.5) [319, 553, 853, 952], [¹¹¹ In]DOTA-dLVT (pK _d 8.3) [318]

Comments: The V₂ receptor exhibits marked species differences, such that many ligands (d(CH₂)₅[D-Ile², Ile⁴]AVP and [³H]desGly-NH₂[D-Ile², Ile⁴]VP) exhibit low affinity at human V₂ receptors [29]. Similarly, [³H]d[D-Arg⁸]VP is V₂ selective in the rat, not in the human [1627]. The gene encoding the V₂ receptor is polymorphic in man, underlying nephrogenic diabetes insipidus [148]. D[Cha⁴]AVP is selective only for the human and bovine V_{1b} receptors [415], while d[Leu⁴]LVP has high affinity for the rat V_{1b} receptor [1485].

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VIP and PACAP receptors

G protein-coupled receptors → VIP and PACAP receptors

Overview: Vasoactive intestinal peptide (VIP) and pituitary adenylate cyclase-activating peptide (PACAP) receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Vasoactive Intestinal Peptide Receptors [704, 705]**) are activated by the endogenous peptides VIP (VIP, P01282), PACAP-38 (ADCYAP1, P18509), PACAP-27 (ADCYAP1, P18509), peptide histidine isoleucineamide (PHI {Mouse, Rat}), peptide histidine methionineamide (PHM (VIP, P01282)) and peptide histidine valine (PHV (VIP, P01282)). VPAC₁ and VPAC₂ receptors display compa-

parable affinity for the PACAP peptides, PACAP-27 (ADCYAP1, P18509) and PACAP-38 (ADCYAP1, P18509), and VIP (VIP, P01282), whereas PACAP-27 (ADCYAP1, P18509) and PACAP-38 (ADCYAP1, P18509) are >100 fold more potent than VIP (VIP, P01282) as agonists of most isoforms of the PAC₁ receptor. However, one splice variant of the human PAC₁ receptor has been reported to respond to PACAP-38 (ADCYAP1, P18509), PACAP-27 (ADCYAP1, P18509) and VIP (VIP, P01282) with comparable affinity [393]. PG 99-465 [1320] has been used as a selective VPAC₂ receptor antagonist in a number of phys-

iological studies, but has been reported to have significant activity at VPAC₁ and PAC₁ receptors [422]. The selective PAC₁ receptor agonist maxadilan, was extracted from the salivary glands of sand flies (*Lutzomyia longipalpis*) and has no sequence homology to VIP (VIP, P01282) or the PACAP peptides [1330]. Two deletion variants of maxadilan, M65 [1918] and Max.d.4 [1331] have been reported to be PAC₁ receptor antagonists, but these peptides have not been extensively characterised.

	PAC ₁ receptor	VPAC ₁ receptor	VPAC ₂ receptor
Nomenclature	PAC ₁ receptor	VPAC ₁ receptor	VPAC ₂ receptor
HGNC, UniProt	ADCYAP1R1, P41586	VIPR1, P32241	VIPR2, P41587
Rank order of potency	PACAP-27 (ADCYAP1, P18509), PACAP-38 (ADCYAP1, P18509) ≫ VIP (VIP, P01282)	VIP (VIP, P01282), PACAP-27 (ADCYAP1, P18509), PACAP-38 (ADCYAP1, P18509) ≫ GHRH (GHRH, P01286), PHI {Pig}, secretin (SCT, P09683)	VIP (VIP, P01282), PACAP-38 (ADCYAP1, P18509), PACAP-27 (ADCYAP1, P18509) > PHI {Pig} ≫ GHRH (GHRH, P01286), secretin (SCT, P09683)
Selective agonists	maxadilan (pEC ₅₀ 10.3) [422], maxadilan (pEC ₅₀ 6.2) [422]	[Lys ¹⁵ ,Arg ¹⁶ ,Leu ²⁷]VIP-(1-7)/GRF-(8-27)-NH ₂ (pEC ₅₀ 8.3) [1315], [Ala ¹¹ ,22,28]VIP (pK _i 8.1) [1393]	Ro 25-1553 (pIC ₅₀ 7.8–9.5) [634, 887, 1315], Ro 25-1392 (pK _i 8) [2056]
Selective antagonists	–	PG 97-269 (pIC ₅₀ 8.7) [633, 887]	–
Labelled ligands	[¹²⁵ I]PACAP-27 (Agonist) (pK _d 9.1) [1509]	[¹²⁵ I]VIP (human, mouse, rat) (Agonist) (pK _d 9.4) [1393], [¹²⁵ I]PACAP-27 (Agonist)	[¹²⁵ I]VIP (human, mouse, rat) (Agonist) (pK _d 9.2) [1393], [¹²⁵ I]PACAP-27 (Agonist)

Comments: Subtypes of PAC₁ receptors have been proposed based on tissue differences in the potencies of PACAP-27 (ADCYAP1, P18509) and PACAP-38 (ADCYAP1, P18509); these might result from differences in G protein coupling and second messenger mechanisms [1939], or from alternative splicing of PAC₁ receptor mRNA [1788].

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