THE CONCISE GUIDE TO PHARMACOLOGY 2017/18:
G protein-coupled receptors

Stephen PH Alexander1, Arthur Christopoulos2, Anthony P Davenport3, Eamonn Kelly4, Neil V Marrion4, John A Peters5, Elena Faccenda6, Simon D Harding6, Adam J Pawson6, Joanna L Sharman6, Christopher Southan6, Jamie A Davies6 and CGTP Collaborators

1 School of Life Sciences, University of Nottingham Medical School, Nottingham, NG7 2UH, UK
2 Monash Institute of Pharmaceutical Sciences and Department of Pharmacology, Monash University, Parkville, Victoria 3052, Australia
3 Clinical Pharmacology Unit, University of Cambridge, Cambridge, CB2 0QQ, UK
4 School of Physiology, Pharmacology and Neuroscience, University of Bristol, Bristol, BS8 1TD, UK
5 Neuroscience Division, Medical Education Institute, Ninewells Hospital and Medical School, University of Dundee, Dundee, DD1 9SY, UK
6 Centre for Integrative Physiology, University of Edinburgh, Edinburgh, EH8 9XD, UK

Abstract

The Concise Guide to PHARMACOLOGY 2017/18 provides concise overviews of the key properties of nearly 1800 human drug targets with an emphasis on selective pharmacology (where available), 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. Although the Concise Guide represents approximately 400 pages, the material presented is substantially reduced compared to information and links presented on the website. It provides a permanent, citable, point-in-time record that will survive database updates. The full contents of this section can be found at http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/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 landscape format of the Concise Guide is designed to facilitate comparison of related targets from material contemporary to mid-2017, and supersedes data presented in the 2015/16 and 2013/14 Concise Guides and previous Guides to Receptors and Channels. It is produced in close conjunction with the Nomenclature Committee of the Union of Basic and Clinical Pharmacology (NC-IUPHAR), therefore, providing official IUPHAR classification and nomenclature for human drug targets, where appropriate.

Conflict of interest

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

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Glutamate family (class C), which includes metabotropic glutamate receptors, a calcium-sensing receptor and GABAB receptors, as well as three taste type 1 receptors and a family of pheromone receptors (V2 receptors) that are abundant in rodents but absent in man [1362].

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).

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 [1609].

Frizzled family consists of 10 Frizzled proteins (FZD(1-10)) and Smoothened (SMO). 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)) [738].

GPCR families

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

*Numbers in brackets refer to orphan receptors for which an endogenous ligand has been proposed in at least one publication, see [414]; [1511]; [1362]; [1941].

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 β2 adrenoceptor, the latter culminating in the award of the 2012 Nobel Prize in chemistry to Robert Lefkowitz and Brian Kobilka [1021, 1137].
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 [557], 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 [414]. 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

| GPR1 | GPR3 | GPR4 | GPR6 | GPR12 | GPR15 | GPR17 | GPR20 |
|------|------|------|------|-------|-------|-------|-------|
| GPR22 | GPR26 | GPR31 | GPR34 | GPR35 | GPR37 | GPR39 | GPR50 |
| GPR63 | GPR65 | GPR68 | GPR75 | GPR84 | GPR87 | GPR88 | GPR132 |
| GPR149 | GPR161 | GPR183 | LGR4 | LGR5 | LGR6 | MAS1 | MRGPRD |
| MRGPRX1 | MRGPRX2 | P2RY10 | TAAR2 |

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 | GPR1 | GPR3 |
|--------------|------|------|
| HGNC, UniProt | GPR1, P46091 | GPR3, P46089 |
| Endogenous agonists | chemeerin (RARRES2, Q99969) [101] | – |
| Agonists | – | diphenyleiodonium chloride [2179] |
| Nomenclature | GPR1 | GPR3 |
|--------------|------|------|
| Comments     | Reported to act as a co-receptor for HIV [1791]. See review [414] for discussion of pairing with chemerin. | sphingosine 1-phosphate was reported to be an endogenous agonist [1997], but this finding was not replicated in subsequent studies [2182]. Reported to activate adenylyl cyclase constitutively through Go [494]. Gene disruption results in premature ovarian ageing [1128], reduced β-amyloid deposition [1943] and hypersensitivity to thermal pain [1689] in mice. First small molecule inverse agonist [903] and agonists identified [2179]. |

| Nomenclature | GPR4 | GPR6 | GPR42 |
|--------------|------|------|-------|
| HGNC, UniProt | GPR4, P46093 | GPR6, P46095 | GPR42, O15529 |
| Endogenous ligands | Protons | – | – |
| Comments     | An initial report suggesting activation by lysophosphatidylcholine and sphingosylphosphorylcholine [2225] has been retracted [1470]. GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [414, 1775]. Gene disruption is associated with increased perinatal mortality and impaired vascular proliferation [2173]. Negative allosteric modulators of GPR4 have been reported [1967]. | An initial report that sphingosine 1-phosphate (S1P) was a high-affinity ligand (EC50 value of 39nM) [855, 1997] was not repeated in arrestin-based assays [1854, 2182]. Reported to activate adenylyl cyclase constitutively through Go and to be located intracellularly [1521]. GPR6-deficient mice showed reduced striatal cyclic AMP production in vitro and selected alterations in instrumental conditioning in vivo [1200]. | – |

| Nomenclature | GPR12 | GPR15 | GPR17 |
|--------------|-------|-------|-------|
| HGNC, UniProt | GPR12, P47775 | GPR15, P49685 | GPR17, Q13304 |
| Endogenous agonists | – | – | UDP-glucose [134, 359], LTC4 [359], UDP-galactose [134, 359], uridine diphosphate [134, 359], uridine diphosphate [134, 359], LTD4 [359] |
| Comments     | Reports that sphingosine 1-phosphate is a ligand of GPR12 [854, 1997] have not been replicated in arrestin-based assays [1854, 2182]. Gene disruption results in dyslipidemia and obesity [158]. | Reported to act as a co-receptor for HIV [490]. In an infection-induced colitis model, Gpr15 knockout mice were more prone to tissue damage and inflammatory cytokine expression [991]. | Reported to be a dual leukotriene and uridine diphosphate receptor [359]. Another group instead proposed that GPR17 functions as a negative regulator of the CysLT1 receptor response to leukotriene D4 (LTD4). For further discussion, see [414]. Reported to antagonize CysLT1 receptor signalling in vivo and in vitro [1239]. See reviews [258] and [414]. |
### Nomenclature

| GPR19 | GPR20 | GPR21 | GPR22 | GPR25 | GPR26 | GPR27 |
|-------|-------|-------|-------|-------|-------|-------|
| **HGNC, UniProt** | GPR19, Q15760 | GPR20, Q99678 | GPR21, Q99679 | GPR22, Q99680 | GPR25, O00155 | GPR26, Q8NDV2 | GPR27, Q9NS67 |

### Comments

**GPR19**
- Reported to inhibit adenyl cyclase constitutively through Gi/o [743].

**GPR20**
- GPR20 deficient mice exhibit hyperactivity characterised by increased total distance travelled in an open field test [213].

**GPR21**
- Gpr21 knockout mice were resistant to diet-induced obesity, exhibiting an increase in glucose tolerance and insulin sensitivity, as well as a modest lean phenotype [1516].

**GPR22**
- Gene disruption results in increased severity of functional decompensation following aortic banding [10].

**GPR25**
- Identified as a susceptibility locus for osteoarthritis [520, 975, 2011].

**GPR26**
- Has been reported to activate adenyl cyclase constitutively through Gs [923].

**GPR27**
- Knockdown of Gpr27 reduces endogenous mouse insulin promotor activity and glucose-stimulated insulin secretion [1059].

### Nomenclature

| GPR31 | GPR32 | GPR33 | GPR34 |
|-------|-------|-------|-------|
| **HGNC, UniProt** | GPR31, O00270 | GPR32, Q75388 | GPR33, Q49SQ1 | GPR34, Q9UPC5 |

### Potency order of endogenous ligands

- 12S-HETE [700] – Mouse

### Endogenous agonists

- resolvin D1 > LXA4

### Labelled ligands

- [3H]resolvin D1 (Agonist) [1052]

### Comments

**GPR31**
- Has been demonstrated to activate GPR32 in two publications [331, 1052]. The pairing was not replicated in a recent study based on arrestin recruitment [1854].

**GPR32**
- GPR32 is a pseudogene in most individuals, containing a premature stop codon within the coding sequence of the second intracellular loop [1696].

**GPR33**
- 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 [1854]. Fails to respond to a variety of lipid-derived agents [2182]. Gene disruption results in an enhanced immune response [1168].

**GPR34**
- Characterization of agonists at this receptor is discussed in [859] and [414].
### Nomenclature
| GPR35 | GPR37 |
|---|---|
| HGNC, UniProt | GPR35, Q9HC97 | GPR37, O15354 |

### Endogenous agonists
| GPR35 | 2-oleoyl-LPA [1503], kynurenic acid [1854, 2066] |
|---|---|
| GPR37 | – |

### Agonists
- 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 [1061]. 2-oleoyl-LPA has also been proposed as an endogenous ligand [1503] but these results were not replicated in an arrestin assay [1854]. The phosphodiesterase inhibitor zaprinast [1937] has become widely used as a surrogate agonist to investigate GPR35 pharmacology and signalling [1937]. GPR35 is also activated by the pharmaceutical adjunct pamoic acid [2218]. See reviews [414] and [453].

### Comments
- Reported to associate and regulate the dopamine transporter [1269] and to be a substrate for parkin [1267]. Gene disruption results in altered striatal signalling [1268]. The peptides prosaptide and prosaposin are proposed as endogenous ligands for GPR37 and GPR37L1 [1324].

### Nomenclature
| GPR37L1 | GPR39 | GPR45 | GPR50 |
|---|---|---|---|
| HGNC, UniProt | GPR37L1, O60883 | GPR39, Q43194 | GPR45, Q9YSY3 |

### Endogenous agonists
| GPR37L1 | – |
|---|---|
| GPR39 | Zn$^{2+}$ [813] |

### Agonists
- Zn$^{2+}$ has been reported to be a potent and efficacious agonist of human, mouse and rat GPR39 [2176]. 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 [285]. Gene disruption results in obesity and altered adipocyte metabolism [1567]. Reviewed in [414].

### Comments
- GPR50 is structurally related to MT$_1$ and MT$_2$ melatonin receptors, with which it heterodimerises constitutively and specifically [1155]. Gpr50 knockout mice display abnormal thermoregulation and are much more likely than wild-type mice to enter fasting-induced torpor [117].

### Nomenclature
| GPR52 | GPR61 | GPR62 | GPR63 | GPR65 |
|---|---|---|---|---|
| HGNC, UniProt | GPR52, Q9Y2TS | GPR61, Q9BZJ8 | GPR62, Q9BZJ7 | GPR63, Q9BZJ6 |

### Endogenous ligands
| GPR52 | – |
|---|---|
| GPR61 | – |
| GPR62 | – |
| GPR63 | – |
| GPR65 | Protons |

### Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of Concise Guide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)**
### Nomenclature

| Gene Symbol | HGNC, UniProt Accession |
|-------------|-------------------------|
| GPR52       | GPR52, Q15743           |
| GPR61       | GPR61, Q95800           |
| GPR62       | GPR62, Q17509           |
| GPR63       | GPR63, Q96P69           |
| GPR65       | GPR65, Q96P67           |

### Endogenous Ligands

| Gene Symbol | Endogenous Ligands |
|-------------|--------------------|
| GPR52       | Protons            |
| GPR61       | sphingosine 1-phosphate and dioleoylphosphatidic acid |
| GPR62       | –                  |
| GPR63       | –                  |
| GPR65       | GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH. Reported to activate adenylyl cyclase; gene disruption leads to reduced eosinophilia in models of allergic asthma [414, 1775]. |

### Allosteric Modulators

| Gene Symbol | Allosteric Modulators |
|-------------|-----------------------|
| GPR52       | lorazepam (Positive)  |
| GPR61       | –                     |
| GPR62       | –                     |
| GPR63       | –                     |
| GPR65       | –                     |

### Comments

- **GPR52**: First small molecule agonist reported [1774].
- **GPR61**: GPR61 deficient mice exhibit obesity associated with hyperphagia [1422]. Although no endogenous ligands have been identified, 5-(nonyloxy)tryptamine has been reported to be a low affinity inverse agonist [1925].
- **GPR62**: sphingosine 1-phosphate and dioleoylphosphatidic acid have been reported to be low affinity agonists for GPR63 [1459] but this finding was not replicated in an arrestin-based assay [2182].
- **GPR65**: GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [414, 1775]. Reported to activate adenylyl cyclase; gene disruption leads to reduced eosinophilia in models of allergic asthma [1044].
- **GPR68**: was previously identified as a receptor for sphingosylphosphorylcholine (SPC) [2157], but the original publication has been retracted [2156]. GPR4, GPR65, GPR68 and GPR132 are now thought to function as proton-sensing receptors detecting acidic pH [414, 1775]. A family of 3,5-disubstituted isoxazoles were identified as agonists of GPR68 [1691].
- **GPR75**: CCL5 (CCL5, P13501) was reported to be an agonist of GPR75 [856], but the pairing could not be repeated in an arrestin assay [1854].
- **GPR78**: has been reported to be constitutively active, coupled to elevated cAMP production [923].
- **GPR79**: –
- **GPR82**: Mice with Gpr82 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 [507].

**Searchable database**: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)  
**Full Contents of Concise Guide**: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
### Nomenclature

**GPR83**

**GPR84**

**GPR85**

**GPR87**

**GPR88**

**GPR101**

**Nomenclature**

| Comments | Medium chain free fatty acids with carbon chain lengths of 9-14 activate GPR84 [1901, 2067]. A surrogate ligand for GPR84, 6-octylaminouracil has also been proposed [1901]. See review [414] for discussion of classification. Mutational analysis and molecular modelling of GPR84 has been reported [1463]. |

| Comments | Proposed to regulate hippocampal neurogenesis in the adult, as well as neurogenesis-dependent learning and memory [319]. |

| Comments | Gene disruption results in altered striatal signalling [1203]. Small molecule agonists have been reported [151]. |

| Comments | Mutations in GPR101 have been linked to gigantism and acromegaly [1982]. |

#### GPR132

**GPR132**

**GPR135**

**GPR139**

**GPR141**

**GPR142**

**GPR146**

**Endogenous ligands**

| Protons | – | – | Peptide agonists have been reported [867]. | – | Small molecule agonists have been reported [1968, 2196]. | – |

**Comments**

**GPR132** is now thought to function as a proton-sensing receptor detecting acidic pH [414, 1775]. Reported to respond to lysophosphatidylcholine [934], but later retracted [2126].

**GPR146** demonstrated inhibition of proinsulin C-peptide (INS, P01308)-induced stimulation of c-Fos expression following knockdown of GPR146 in KATO III cells, suggesting proinsulin C-peptide as an endogenous ligand of the receptor [2193].

#### GPR148

**GPR148**

**GPR149**

**GPR150**

**GPR151**

**GPR152**

**GPR153**

**GPR160**

**Endogenous ligands**

| – | – | – | – | – | – |

**Comments**

**Gpr149** knockout mice displayed increased fertility and enhanced ovulation, with increased levels of FSH receptor and cyclin D2 mRNA levels [491].

GPR151 responded to galanin with an EC50 value of 2 μM, suggesting that the endogenous ligand shares structural features with *galanin* (GAL, P22466) [853].

### Searchable database

- [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)
- [Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)

---

**S.P.H. Alexander et al. The Concise Guide to PHARMACOLOGY 2017/18: G protein-coupled receptors. British Journal of Pharmacology (2017) 174, S17–S129**
| Nomenclature | GPR161 | GPR162 | GPR171 | GPR173 | GPR174 | GPR176 | GPR182 |
|-------------|--------|--------|--------|--------|--------|--------|--------|
| HGNC, UniProt | GPR161, Q8N6UB | GPR162, O14626 | GPR171, Q9NS66 | GPR173, Q9BXC1 | GPR174, Q14439 | GPR176, Q15218 | GPR182, O15218 |
| Endogenous agonists | – | – | – | lysophosphatidylserine [864] | – | – | – |
| Comments | A C-terminal truncation (deletion) mutation in Gpr161 causes congenital cataracts and neural tube defects in the vacuolated lens (vl) mouse mutant [1289]. The mutated receptor is associated with cataract, spina bifida and white belly spot phenotypes in mice [1039]. Gene disruption is associated with a failure of asymmetric embryonic development in zebrafish [1151]. | – | 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 [654]. | – | See [859] which discusses characterization of agonists at this receptor. | – | Rat GPR182 was first proposed as the adrenomedullin receptor [947]. However, it was later reported that rat and human GPR182 did not respond to adrenomedullin [973] and GPR182 is not currently considered to be a genuine adrenomedullin receptor [756]. |

| Nomenclature | GPR183 | LGR4 |
|-------------|--------|------|
| HGNC, UniProt | GPR183, P32249 | LGR4, Q9BXB1 |
| Endogenous agonists | 7α,25-dihydroxycholesterol [729, 1191], 7α,27-dihydroxycholesterol [1191], 7β, 25-dihydroxycholesterol [1191], 7β, 27-dihydroxycholesterol [1191] | R-spondin-2 (RSPO2, Q6UXX9) [277], R-spondin-1 (RSPO1, Q2MKA7) [277], R-spondin-3 (RSPO3, Q9BXY4) [277], R-spondin-4 (RSPO4, Q2I0M5) [277] |
| 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 [729, 1191]. 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 [966, 1557]. | 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 [277, 426, 1686]. Gene disruption leads to multiple developmental disorders [911, 1219, 1849, 2092]. |
| Nomenclature | LGR5 | LGR6 | MAS1 | MAS1L |
|--------------|------|------|------|-------|
| HGNC, UniProt | LGR5, O75473 | LGR6, Q9HBX8 | MAS1, P04201 | MAS1L, P35410 |
| Endogenous agonists | R-spondin-2 (RSPO2, Q6UXX9) [277], R-spondin-1 (RSPO1, Q2MKA7) [277], R-spondin-3 (RSPO3, Q9BXY4) [277], R-spondin-4 (RSPO4, Q2I0M5) [277] | R-spondin-1 (RSPO1, Q2MKA7) [277, 426], R-spondin-2 (RSPO2, Q6UXX9) [277, 426], R-spondin-3 (RSPO3, Q9BXY4) [277, 426], R-spondin-4 (RSPO4, Q2I0M5) [277, 426] | – | – |
| Agonists | – | – | angiotensin-(1-7) (AGT, P01019) [645] – Mouse | – |
| Comments | 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 [277, 426]. | – | – | – |

| Nomenclature | MRGPRD | MRGPRE | MRGPRF | MRGPRG |
|--------------|--------|--------|--------|--------|
| HGNC, UniProt | MRGPRD, Q8TDS7 | MRGPRE, Q86SM8 | MRGPRF, Q96AM1 | MRGPRG, Q86SMS |
| Endogenous agonists | β-alanine [1797, 1854] | – | See reviews [414] and [1847]. | MRGPRF has been reported to respond to stimulation by angiotensin metabolites [620]. See reviews [414] and [1847]. |
| 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-Pro7-angiotensin-(1-7), β-alanine and PD123319 [1102]. Genetic ablation of MRGPRD+ neurons of adult mice decreased behavioural sensitivity to mechanical stimuli but not to thermal stimuli [292]. See reviews [414] and [1847]. | – | – | See reviews [414] and [1847]. |
| Nomenclature | MRGPRX1 | MRGPRX2 | MRGPRX3 | MRGPRX4 | OPN3 | OPN4 | OPNS |
|--------------|---------|---------|---------|---------|------|------|------|
| HGNC, UniProt| MRGPRX1, Q96LB2 | MRGPRX2, Q96LB1 | MRGPRX3, Q96LB0 | MRGPRX4, Q96LA9 | OPN3, Q9H1Y3 | OPN4, Q9UHM6 | OPNS, Q6U736 |
| Endogenous agonists | bovine adrenal medulla peptide 8-22 (PENK, P01210) [315, 1144, 1854] | PAMP-20 (ADM, P35318) [942] | – | – | – | – | – |
| Agonists | – | cortistatin-14 (Mouse, Rat) [942, 1667, 1854] | – | – | – | – | – |
| Selective agonists | – | PAMP-12 (human) [942] | – | – | – | – | – |
| Comments | Reported to mediate the sensation of itch [1196, 1808]. 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 [1144] were replicated in an independent study using an arrestin recruitment assay [1854]. See reviews [414] and [1847]. | 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 [1667], also confirmed in an independent study using an arrestin recruitment assay [1854]. See reviews [414] and [1847]. | – | See reviews [414] and [1847]. | – | – | – |

| Nomenclature | P2RY8 | P2RY10 | TAAR2 | TAAR3 | TAAR4P |
|--------------|-------|-------|-------|-------|--------|
| HGNC, UniProt | P2RY8, Q86VZ1 | P2RY10, O00398 | TAAR2, Q9P1P5 | TAAR3P, Q9P1P4 | TAAR4P, – |
| Potency order of endogenous ligands | – | – | β-phenylethylamine > tryptamine [189] | – | – |
| Endogenous agonists | – | sphingosine 1-phosphate [1401], LPA [1401] | – | – | – |
| Comments | – | – | Probable pseudogene in 10–15% of Asians due to a polymorphism (rs8192646) producing a premature stop codon at amino acid 168 [414]. | TAAR3 is thought to be a pseudogene in man though functional in rodents [414]. | Pseudogene in man but functional in rodents [414]. |
### Class C Orphans

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

| Nomenclature | TAAR5 | TAAR6 | TAAR8 | TAAR9 |
|---------------|-------|-------|-------|-------|
| HGNC, UniProt | TAAR5, O14804 | TAAR6, Q96R18 | TAAR8, Q96N4 | TAAR9, Q96R19 |
| Comments | Trimethylamine is reported as an agonist [2058] and 3-iodothyronamine an inverse agonist [450]. | – | – | – |

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 [2023].

### 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 Go14-mediated signalling, although the precise mechanisms remain obscure. Gene disruption studies suggest the involvement of PLCβ2 [2215], TRPM5 [2215] and IP3 [802] 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 [1162]. T1R2/T1R3 heterodimers respond to sugars, such as sucrose, and artificial sweeteners, such as saccharin [1440].

**GPRC6 receptor**

GPRC6 is a related Gq-coupled receptor which responds to basic amino acids [2090].

---

**Searchable database**: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide**: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Taste 2 receptors

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

**Overview:** 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 [302], while T2R14 responded to at least eight different bitter tastants, including (-)-α-thujone and picrotoxinin [124]. Specialist database BitterDB contains additional information on bitter compounds and receptors [2113].

**Nomenclature**

| Nomenclature | TAS1R1 | TAS1R2 | TAS1R3 |
|--------------|--------|--------|--------|
| HGNC, UniProt | TAS1R1, Q7RTX1 | TAS1R2, Q8TE23 | TAS1R3, Q7RTX0 |

**Nomenclature**

| Nomenclature | TAS2R1 | TAS2R3 | TAS2R4 | TAS2R5 | TAS2R7 | TAS2R8 | TAS2R9 |
|--------------|--------|--------|--------|--------|--------|--------|--------|
| HGNC, UniProt | TAS2R1, Q9NYW7 | TAS2R3, Q9NYW6 | TAS2R4, Q9NYW5 | TAS2R5, Q9NYW4 | TAS2R7, Q9NYW3 | TAS2R8, Q9NYW2 | TAS2R9, Q9NYW1 |

**Nomenclature**

| Nomenclature | TAS2R10 | TAS2R13 | TAS2R14 | TAS2R16 | TAS2R19 | TAS2R20 | TAS2R30 |
|--------------|---------|---------|---------|---------|---------|---------|---------|
| HGNC, UniProt | TAS2R10, Q9NYW0 | TAS2R13, Q9NY9 | TAS2R14, Q9NY8 | TAS2R16, Q9NY7 | TAS2R19, P59542 | TAS2R20, P59543 | TAS2R30, P59541 |

**Nomenclature**

| Nomenclature | TAS2R31 | TAS2R38 | TAS2R39 | TAS2R40 |
|--------------|---------|---------|---------|---------|
| HGNC, UniProt | TAS2R31, P59538 | TAS2R38, P59533 | TAS2R39, P59534 | TAS2R40, P59535 |
| Antagonists   | 6-methoxysakuranetin (pIC<sub>50</sub> 5.6) [1000], GIV3727 (pIC<sub>50</sub> 5.1–5.2) [1823] | – | – | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Other 7TM proteins

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

| Nomenclature | TAS2R41 | TAS2R42 | TAS2R43 | TAS2R45 | TAS2R46 | TAS2R50 | TAS2R60 |
|--------------|---------|---------|---------|---------|---------|---------|---------|
| HGNC, UniProt | P59536 | P59538 | P59537 | P59539 | P59540 | P59544 | P59551 |
| Nomenclature | GPR107 | GPR137 | OR51E1 | TPRA1   | GPR143 | GPR157 |
| HGNC, UniProt | Q5VW38 | Q96N19 | Q8TCB6 | Q86W33  | P51810 | QSUAW9 |
| Endogenous agonists | –      | –      | –      | –      | levodopa [1207] | –      |
| Comments | GPR107 is a member of the LUSTR family of proteins found in both plants and animals, having similar topology to G protein-coupled receptors [489]. | GPR137, Q96N19 | OR51E1 is a putative olfactory receptor. | TPRA1 shows no homology to known G protein-coupled receptors. | GPR143, P51810 | GPR157, QSUAW9 |
| Comments | GPR157 has ambiguous sequence similarities to several different GPCR families (class A, class B and the slime mold cyclic AMP receptor). Because of its distant relationship to other GPCRs, it cannot be readily classified. |

Further reading on Orphan and other 7TM receptors

Davenport AP et al. (2013) International Union of Basic and Clinical Pharmacology. LXXXVIII. G protein-coupled receptor list: recommendations for new pairings with cognate ligands. Pharmacol Rev 65: 967-86 [PMID:23686350]
Gilissen J et al. (2016) Insight into SUCNR1 (GPR91) structure and function. Pharmacology & Therapeutics 159: 56-65 [PMID:25118328]
Insel PA et al. (2015) G Protein-Coupled Receptor (GPCR) Expression in Native Cells: 'Novel' endoGPCRs as Physiologic Regulators and Therapeutic Targets. Molecular Pharmacology 88: 181-187 [PMID:25737495]
Khan MZ et al. (2017) Neuro-psychopharmacological perspective of Orphan receptors of Rhodopsin (class A) family of G protein-coupled receptors. Psychopharmacology (Berl) 234: 1181-1207 [PMID:28289782]
Mackenzie AE et al. (2017) The emerging pharmacology and function of GPR35 in the nervous system. Neuropharmacology 113: 661-671 [PMID:26232640]
Ngo T et al. (2016) Identifying ligands at orphan GPCRs: current status using structure-based approaches. Br J Pharmacol 173: 2934-2951 [PMID:26837045]
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 [828] and subsequently revised [742]) are, with the exception of the ionotropic 5-HT3 class, GPCRs 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-HT2A (non-functional), 5-HT2C (non-functional), 5-HT4, 5-HT6 (non-functional) and 5-HT7 receptors. Unique amongst the GPCRs, RNA editing produces 5-HT2C receptor isoforms that differ in function, such as efficiency and specificity of coupling to Gq/11 and also pharmacology [167, 2098]. Most 5-HT receptors (except 5-HT1E and 5-HT5B) play specific roles mediating functional responses in different tissues (reviewed by [1625, 2037]).

| Nomenclature | S-HT1A receptor | S-HT1B receptor |
|--------------|-----------------|-----------------|
| HGNC, UniProt | HTR1A, P08908   | HTR1B, P28222   |
| Agonists     | U92016A [1302], vilazodone (Partial agonist) [421], vortioxetine (Partial agonist) [96] | L-694,247 [671], naratriptan (Partial agonist) [1425], eletriptan [1425], frovatriptan [2158], zolmitriptan (Partial agonist) [1425], vortioxetine (Partial agonist) [96], rizatriptan (Partial agonist) [1425] |
| Selective agonists | 8-OH-DPAT [431, 720, 939, 1143, 1338, 1451, 1453, 1454], NLX-101 [1452] | CP94253 [1022] |
| Antagonists | (S)-UH 301 (pKᵢ 7.9) [1451] | SB 224289 (Inverse agonist) (pKᵢ 8.2–8.6) [614, 1449, 1768], SB236057 (Inverse agonist) (pKᵢ 8.2) [1331], GR-55562 (pKᵢ 7.4) [830] |
| Selective antagonists | WAY-100635 (pKᵢ 7.9–9.2) [1451, 1453], robalzotan (pKᵢ 9.2) [915] | – |
| Labelled ligands | [³H]robalzotan (Antagonist) (pKᵢ 9.8) [904], [³H]WAY100635 (Antagonist) (pKᵢ 9.5) [978], [³H]8-OH-DPAT (Agonist) [160, 939, 1450, 1453], [³H]NLX-112 (Agonist) [785], [¹¹C]WAY100635 (Antagonist) [1991], p-[¹⁸F]MPPF (Antagonist) [382] | [³H]N-methyl-AZ10419369 (Agonist, Partial agonist) [1245], [³H]GR 125,743 (Selective Antagonist) (pKᵢ 8.6–9.2) [671, 2150], [³H]alniditan (Agonist) [1150], [¹²⁵I]GTI (Agonist) [197, 237] – Rat, [³H]eletriptan (Agonist, Partial agonist) [1425], [³H]sumatriptan (Agonist, Partial agonist) [1425], [¹¹C]AZ10419369 (Agonist, Partial agonist) [2029] |

| Nomenclature | S-HT1D receptor | S-ht1e receptor | S-HT1F receptor |
|--------------|-----------------|-----------------|-----------------|
| HGNC, UniProt | HTR1D, P28221   | HTR1E, P28566   | HTR1F, P30939   |
| Agonists     | dihydroergotamine [719, 1150, 1157], ergotamine [648], L-694,247 [2140], naratriptan [457, 1425, 1651], zolmitriptan [1425], frovatriptan [2158], rizatriptan [1425] | BRL-54443 [232] | BRL-54443 [232], eletriptan [1425], sumatriptan [12, 13, 1425, 2052] |
| Selective agonists | PNU109291 [511] – Gorilla, eletriptan [1425] | – | lasmiditan [1439], LY334370 [2052], S-BODMT [1014], LY344864 [1572] |
| Selective antagonists | SB 714786 (pKᵢ 9.1) [2074] | – | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature  5-HT₁D receptor  5-HT₁E receptor  5-HT₁F receptor
Labelled ligands  [³H]eletriptan (Agonist) [1425], [³H]alniditan (Agonist) [1150], [¹²⁵I]GTI (Selective Agonist) [197, 237] – Rat, [³H]GR 125,743 (Selective Antagonist) (pKᵦ 8.6) [2150], [³H]sumatriptan (Agonist) [1425]  [³H]S-HT (Agonist) [1299, 1532]  [³H]LY34370 (Agonist) [2052], [¹²⁵I]LSD (Agonist) [45] – Mouse

Nomenclature  5-HT₂A receptor  5-HT₂B receptor
HGNC, UniProt  HTR2A, P28223  HTR2B, P41595
Agonists  DOI [210, 1438, 1825]  methysergide (Partial agonist) [1018, 1679, 2053], DOI [1077, 1438, 1730]  mianserin (pKᵦ 7.9–8.8) [184, 1018, 2053]
Selective agonists  risperidone (Inverse agonist) (pKᵦ 9.3–10) [1032, 1055, 1746], mianserin (pKᵦ 7.7–9.6) [1018, 1045, 1338], ziprasidone (pKᵦ 8.8–9.5) [1032, 1055, 1746, 1782], volinanserin (pIC₅₀ 6.5–9.3) [1018, 1208, 1640], blonanserin (pKᵦ 9.1) [1487], clozapine (Inverse agonist) (pKᵦ 7.6–9) [1018, 1055, 1335, 1746, 2022]  risperidone (Inverse agonist) (pKᵦ 9.3–10) [1077, 1438, 1730], Ro 60-0175 [1018]  mianserin (pKᵦ 7.9–8.8) [184, 1018, 2053]
Antagonists  ketanserin (pKᵦ 8.1–9.7) [241, 1018, 1630], pimavanserin (Inverse agonist) (pKᵦ 9.3) [603, 2022]  BF-1 (pKᵦ 10.1) [1742], RS-127445 (pKᵦ 9–9.5) [184, 1018], ECIS-7625 (pKᵦ 9) [1045]  ketanserin (pKᵦ 8.1–9.7) [241, 1018, 1630], pimavanserin (Inverse agonist) (pKᵦ 9.3) [603, 2022]  BF-1 (pKᵦ 10.1) [1742], RS-127445 (pKᵦ 9–9.5) [184, 1018], ECIS-7625 (pKᵦ 9) [1045]
Selectives antagonists  ketanserin (pKᵦ 8.1–9.7) [241, 1018, 1630], pimavanserin (Inverse agonist) (pKᵦ 9.3) [603, 2022]  ketanserin (pKᵦ 8.1–9.7) [241, 1018, 1630], pimavanserin (Inverse agonist) (pKᵦ 9.3) [603, 2022]  BF-1 (pKᵦ 10.1) [1742], RS-127445 (pKᵦ 9–9.5) [184, 1018], ECIS-7625 (pKᵦ 9) [1045]
Labelled ligands  [³H]fananserin (Antagonist) (pKᵦ 9.9) [1251] – Rat, [³H]ketanserin (Antagonist) (pKᵦ 8.6–9.7) [1018, 1630], [¹¹C]volinanserin (Antagonist) [712], [¹⁸F]altanserin (Antagonist) [1675]  [³H]LSD (Agonist) [1630], [³H]S-HT (Agonist) [2051] – Rat, [³H]mesulergine (Antagonist, Inverse agonist) (pKᵦ 7.9) [1018], [¹²⁵I]DOI (Agonist)

Nomenclature  5-HT₂C receptor  5-HT₄ receptor
HGNC, UniProt  HTR2C, P28335  HTR4, Q13639
Agonists  DOI [493, 1438, 1730], Ro 60-0175 [999, 1018]  cisapride (Partial agonist) [80, 132, 631, 1326, 1327, 2013]  TD-8954 [1312], ML 10302 (Partial agonist) [140, 164, 1326, 1327, 1328], RS67506 [765] – Rat, relenopride (Partial agonist) [641], velusetrag [1205, 1832], BIMU 8 [362]  cisapride (Partial agonist) [80, 132, 631, 1326, 1327, 2013]  TD-8954 [1312], ML 10302 (Partial agonist) [140, 164, 1326, 1327, 1328], RS67506 [765] – Rat, relenopride (Partial agonist) [641], velusetrag [1205, 1832], BIMU 8 [362]
Selective agonists  WAY-163909 [482], lorcanerin [1955]  FR260010 (pKᵦ 9) [735], SB 242084 (pKᵦ 8.2–9) [974, 1018], RS-102221 (pKᵦ 8.3–8.4) [185, 1018]  RS 100235 (pKᵦ 8.7–12.2) [362, 1663], SB 204070 (pKᵦ 9.8–10.4) [132, 1326, 1327, 2013], CR 113808 (pKᵦ 9.3–10.3) [80, 132, 164, 362, 1327, 1663, 2013]  FR260010 (pKᵦ 9) [735], SB 242084 (pKᵦ 8.2–9) [974, 1018], RS-102221 (pKᵦ 8.3–8.4) [185, 1018]  RS 100235 (pKᵦ 8.7–12.2) [362, 1663], SB 204070 (pKᵦ 9.8–10.4) [132, 1326, 1327, 2013], CR 113808 (pKᵦ 9.3–10.3) [80, 132, 164, 362, 1327, 1663, 2013]
Antagonists  mianserin (Inverse agonist) (pKᵦ 8.3–9.2) [551, 1018, 1338], methysergide (pKᵦ 8.6–9.1) [493, 1018], ziprasidone (Inverse agonist) (pKᵦ 7.9–9) [779, 1055, 1782], olanzapine (Inverse agonist) (pKᵦ 8.1–8.4) [779, 1055, 1782], loxapine (Inverse agonist) (pKᵦ 7.8–8) [779, 1055]  –
Selectives antagonists  FR260010 (pKᵦ 9) [735], SB 242084 (pKᵦ 8.2–9) [974, 1018], RS-102221 (pKᵦ 8.3–8.4) [185, 1018]  FR260010 (pKᵦ 9) [735], SB 242084 (pKᵦ 8.2–9) [974, 1018], RS-102221 (pKᵦ 8.3–8.4) [185, 1018]  FR260010 (pKᵦ 9) [735], SB 242084 (pKᵦ 8.2–9) [974, 1018], RS-102221 (pKᵦ 8.3–8.4) [185, 1018]
### Nomenclature

| 5-HT$_{2C}$ receptor | 5-HT$_4$ receptor |
|----------------------|------------------|
| [H$^3$]mesulergine (Antagonist, Inverse agonist) (pK$_d$ 8.7–9.3) [551, 1630], [H$^{125}$I]DOI (Agonist) [551], [H$^3$]LSD (Agonist) | [H$^3$]GR 113808 (Antagonist) (pK$_d$ 9.7–10.3) [80, 132, 1328, 2013], [H$^{125}$I]SB 207710 (Antagonist) (pK$_d$ 10.1) [233] – Pig, [H$^3$]RS 57639 (Selective Antagonist) (pK$_d$ 9.7) [183] – Guinea pig, [H$^{11}$C]SB207145 (Antagonist) (pK$_d$ 8.6) [1233] |

### Table

| Nomenclature | 5-HT$_{3A}$ receptor | 5-HT$_{3B}$/5-HT$_{3C}$ receptor | 5-HT$_6$ receptor | 5-HT$_7$ receptor |
|--------------|----------------------|-------------------------------|----------------|-----------------|
| HGNC, UniProt| HTR5A, P47898        | HTR5BP, –                      | HTR6, P05046   | HTR7, P34969    |
| Selective agonists | WAY-181187 [1734], E6801 (Partial agonist) [808], WAY-208466 [139], EMD-386088 [1291] | – | – | – |
| Antagonists | – | WAY-181187, WAY-208466 [139], EMD-386088 [1291] | – | – |
| Selective antagonists | SB 699551 (pK$_i$ 8.2) [380] | – | SB399885 (pK$_i$ 9) [801], SB 271046 (pK$_i$ 8.9) [229], cerlapidine (pK$_i$ 8.9) [371], SB357134 (pK$_i$ 8.5) [230], Ro 63-0563 (pK$_i$ 7.9–8.4) [170, 1824] | SB269970 (pK$_i$ 8.6–8.9) [1949], SB656104 (pK$_i$ 8.7) [558], DR-4004 (pK$_i$ 8.7) [647, 985], JNJ-18038683 (pK$_i$ 8.2) [181], SB 258719 (Inverse agonist) (pK$_i$ 7.5) [1950] | – |

### Labelled ligands

| 5-HT$_{3A}$ receptor | 5-HT$_{3B}$/5-HT$_{3C}$ receptor | 5-HT$_6$ receptor | 5-HT$_7$ receptor |
|----------------------|-------------------------------|----------------|----------------|
| [H$^{125}$]LSD (Agonist) [670], [H$^{11}$C]-S-CT (Agonist) [670] – Mouse, [H$^{125}$]LSD (Agonist) [1290] – Mouse, [H$^{11}$C]-S-CT (Agonist) [2049] – Mouse | [H$^{11}$C]-GLS215803 (Antagonist) (pK$_i$ 9.8) [1531], [H$^{125}$]SB258585 (Selective Antagonist) (pK$_d$ 9) [801], [H$^{125}$]LSD (Agonist) [169], [H$^{11}$]Ro 63-0563 (Antagonist) (pK$_d$ 8.3) [170], [H$^{11}$C]-S-CT (Agonist) | [H$^{11}$C]-S-CT (Agonist) [1949], [H$^{11}$C]-S-HT (Agonist) [99, 1864], SB269970 (Selective Antagonist) (pK$_d$ 8.9) [1949], [H$^{11}$C]-LSD (Agonist) [1864] | – |

### 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 [742]. 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. Wang et al. (2013) report X-ray structures which reveal the binding modality of ergotamine and dihydroergotamine to the 5-HT$_{2B}$ receptor in comparison with the structure of the 5-HT$_{2A}$ receptor [2064]. NAS181 is a selective antagonist of the rodent 5-HT$_{1B}$ receptor. Fananserin and ketanserin bind with high affinity to dopamine D4 and histamine H1 receptors respectively, and ketanserin is a potent α1 adrenoceptor antagonist, in addition to blocking 5-HT$_{2A}$ receptors. Lysergic acid (LSD) 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-HT7A receptors [2047]. DHE (dihydroergocryptine), pergolide and cabergoline also show significant preference for arrestin recruitment over G protein coupling at 5-HT$_{2B}$ receptors [2047]. The serotonin antagonist mesulergine was key to the discovery of the 5-HT$_{2C}$ receptor [1546]. The human 5-HT$_{5A}$ receptor has been claimed to couple to several signal transduction pathways when stably expressed in C6 glioma cells [1472] and electrophysiological recordings from mice and rat prefrontal cortex (layer V pyramidal neurons) demonstrate 5-HT-elicted outward currents mediated via the 5-HT$_{5A}$ receptor [660]. 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 [635].
Further reading on 5-Hydroxytryptamine receptors

Bockaert J et al. (2011) 5-HT(4) receptors, a place in the sun: act two. Curr Opin Pharmacol 11: 87-93 [PMID:21342787]

Hayes DJ et al. (2011) 5-HT receptors and reward-related behaviour: a review. Neurosci Biobehav Rev 35: 1419-49 [PMID:21402098]

Hoyer D et al. (1994) International Union of Pharmacology classification of receptors for 5-hydroxytryptamine (Serotonin). Pharmacol. Rev. 46: 157-203 [PMID:7938165]

Leopoldo M et al. (2011) Serotonin 5-HT7 receptor agents: Structure-activity relationships and potential therapeutic applications in central nervous system disorders. Pharmacol. Ther. 129: 120-48 [PMID:20923682]

Meltzer HY et al. (2011) The role of serotonin receptors in the action of atypical antipsychotic drugs. Curr Opin Pharmacol 11: 59-67 [PMID:21420906]

Roberts AJ et al. (2012) The 5-HT(7) receptor in learning and memory. Hippocampus 22: 762-71 [PMID:21484935]

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 [288]) 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-desmethyllozapine, TBPBand 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 [74, 921, 1097, 1294, 1708, 1857, 1858, 1898]. 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 [1110, 1111].

| Nomenclature | M1 receptor | M2 receptor |
|--------------|-------------|-------------|
| HGNC, UniProt| CHRM1, P11229 | CHRM2, P08172 |
| Agonists     | carbachol [350, 888, 2129], pilocarpine (Partial agonist) [888], bethanechol [888] | bethanechol [888] |
| Antagonists  | glycopyrrolate (pIC_{50} 9.9) [1874], umeclidinium (pK_{i} 9.8) [1090, 1705], AE9C90CB (pK_{i} 9.7) [1818], atropine (pK_{i} 8.5–9.6) [350, 888, 1225], tiotropium (pK_{i} 9.6) [452], 4-DAMP (pK_{i} 9.2) [486] | tiotropium (pK_{i} 9.9) [452], umeclidinium (pK_{i} 9.8) [1090, 1705], propantheline (pK_{i} 9.5) [837], glycopyrrolate (Full agonist) (pIC_{50} 9.3) [1874], atropine (pK_{i} 7.8–9.2) [245, 325, 797, 837, 1046, 1437, 1555], AE9C90CB (pK_{i} 8.6) [1818], tolterodine (Inverse agonist) (pK_{i} 8.4–8.6) [642, 1437, 1818] |
| Selective antagonists | biperiden (pK_{d} 9.3) [175], VU0255035 (pK_{i} 7.8) [1786], guanlypirenzepine (pK_{i} 7.3–7.6) [23, 2050] – Rat | tiotropium (pK_{i} 9.6) [1240] |
| Allosteric modulators | muscarinic toxin 7 (Negative) (pK_{d} 11–11.1) [1480], benzoquinazolinone 12 (Positive) (pK_{d} 6.6) [4], KT 5720 (Positive) (pK_{d} 6.4) [1100], brucine (Positive) (pK_{d} 4.5–5.8) [888, 1109], BQCA (Positive) (pK_{d} 4–4.8) [4, 5, 271, 1225], VU0029767 (Positive) [1270], VU0090157 (Positive) [1270] | W-84 (Negative) (pK_{d} 7.1) [351], alcuronium (Negative) (pK_{d} 6.1–6.9) [888, 1983], gallamine (Negative) (pK_{d} 5.9–6.3) [363, 1107], LY2119620 (Positive) (pK_{d} 5.7) [399, 1057], LY2033298 (Positive) (pK_{d} 4.4) [2009] |

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

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
### M1 receptor

**Nomenclature**
- CHRM1, P20309

**Agonists**
- pilocarpine (Partial agonist) [888], carbachol [325, 888, 2129], bethanechol [888]

**Antagonists**
- tiotropium (pKᵢ 9.5–11.1) [452, 469], umeclidinium (pKᵢ 10.2) [1090, 1705], propantheline (pKᵢ 10) [837], AE9C90CB (pKᵢ 9.9) [1818], atropine (pKᵢ 8.9–9.8) [245, 469, 797, 837, 1555, 1831], ipratropium (pKᵢ 9.3–9.8) [469, 797], acridinium (pIC₅₀ 9.8) [1601]

**Selective antagonists**
- WIN 62,577 (Positive) (pKᵢ 5.1) [1111], N-chloromethyl-brucine (Positive) (pKᵢ 3.3) [1109]

**Allosteric modulators**
- muscarinic toxin 3 (Negative) (pKᵢ 8.7) [918, 1512], VU0152100 (Positive) (pEC₅₀ 6.4) [207] – Rat, VU0152099 (Positive) (pEC₅₀ 6.4) [207] – Rat, LY2119620 (Positive) (pKᵢ 5.7) [399], thiochrome (Positive) (pKᵢ 4) [1108], LY2033298 (Positive) [301]

**Selective allosteric modulators**
- [³H]tiotropium (Antagonist) (pKᵢ 10.7) [1705], [³H]QNB (Antagonist) (pKᵢ 10.4) [1555], [³H]N-methyl scopolamine (Antagonist) (pKᵢ 9.7–10.2) [294, 325, 797, 837, 888, 918, 977, 1107], [³H]darifenacin (Antagonist) (pKᵢ 9.5) [1831]

**Labelled ligands**
- [³H]QNB (Antagonist) (pKᵢ 10.1–10.6) [1555], Cy3B-telenzepine (Antagonist) (pKᵢ 9.4–10.3) [294, 350, 352, 797, 888, 889, 918, 977, 1107], [³H]telenzepine (Antagonist) (pKᵢ 9.4) [526] – Rat, Alexa-488-telenzepine (Antagonist) (pKᵢ 9.3) [777], [³H]pirenzepine (Antagonist) (pKᵢ 10.5) [777], [³H]N-methyl scopolamine (Antagonist) (pKᵢ 9.3–9.9) [294, 325, 797, 888, 918, 977, 1107, 2072], Alexa-488-telenzepine (Antagonist) (pKᵢ 8.8) [1444], [³H]acetylcholine (Agonist) [888], [³H]oxotremorine-M (Agonist) [141], [³H]N-methyl-DL-scopolamine (Antagonist) (pKᵢ 9.7–10.2) [294, 325, 797, 888, 918, 977, 1107, 2072], [³H]Darifenacin (Antagonist) [1831] – Mouse

### M2 receptor

**Nomenclature**
- CHRM2, P08173

**Agonists**
- pilocarpine (Partial agonist) [888], carbachol [888, 2129], bethanechol [888]

**Antagonists**
- tiotropium (pKᵢ 9.5–11.1) [452, 469], umeclidinium (pKᵢ 10.2) [1090, 1705], propantheline (pKᵢ 10) [837], AE9C90CB (pKᵢ 9.9) [1818], atropine (pKᵢ 8.9–9.8) [245, 469, 797, 837, 1555, 1831], ipratropium (pKᵢ 8.3–8.6) [469, 797], aclidinium (pIC₅₀ 9.8) [1601]

**Selective antagonists**
- –

**Allosteric modulators**
- ML381 (pKᵢ 6.3) [625], ML380 (Positive) (pEC₅₀ 6.7) [627]

**Selective allosteric modulators**
- –

**Labelled ligands**
- [³H]QNB (Antagonist) (pKᵢ 9.7–10.5) [352, 1555], [³H]N-methyl scopolamine (Antagonist) (pKᵢ 9.9–10.2) [294, 325, 352, 797, 888, 918, 977, 1107, 1512, 2072], [³H]Acetylcholine (Agonist) [1108]

### M3 receptor

**Nomenclature**
- CHRM3, P20309

**Agonists**
- pilocarpine (Partial agonist) [888], carbachol [888, 2129], bethanechol [888]

**Antagonists**
- tiotropium (pKᵢ 9.5–11.1) [452, 469], umeclidinium (pKᵢ 10.2) [1090, 1705], propantheline (pKᵢ 10) [837], AE9C90CB (pKᵢ 9.9) [1818], atropine (pKᵢ 8.9–9.8) [245, 469, 797, 837, 1555, 1831], ipratropium (pKᵢ 9.3–9.8) [469, 797], aclidinium (pIC₅₀ 9.8) [1601]

**Selective antagonists**
- –

**Allosteric modulators**
- ML381 (pKᵢ 6.3) [625], ML380 (Positive) (pEC₅₀ 6.7) [627]

**Selective allosteric modulators**
- –

**Labelled ligands**
- [³H]QNB (Antagonist) (pKᵢ 9.7–10.5) [352, 1555], [³H]N-methyl scopolamine (Antagonist) (pKᵢ 9.9–10.2) [294, 325, 352, 797, 888, 918, 977, 1107, 1512, 2072], [³H]Acetylcholine (Agonist) [1108]
Comments: LY2033298 and BQCA have also been shown to directly activate the M4 and M1 receptors, respectively, via an allosteric site [1119, 1121, 1427, 1428]. The allosteric site for gallamine and strychnine on M2 receptors can be labelled by [3H]dimethyl-W84 [1983]. 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 receptor [2010]. THRX160209, 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 [55, 1866]. 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 their values at physiological ionic strengths [155].

Further reading on Acetylcholine receptors (muscarinic)

Caulfield MP et al. (1998) International Union of Pharmacology. XVII. Classification of muscarinic acetylcholine receptors. Pharmacol. Rev. 50: 279-290 [PMID:9647869]
Eglen RM. (2012) Overview of muscarinic receptor subtypes. Handb Exp Pharmacol 3-28 [PMID:22222692]
Gregory KJ et al. (2007) Allosteric modulation of muscarinic acetylcholine receptors. Curr Neuropsychopharmacol 5: 157-67 [PMID:19305798]

Adenosine receptors
G protein-coupled receptors → Adenosine receptors

Overview: Adenosine receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Adenosine Receptors [569]) are activated by the endogenous ligand adenosine (potentially inosine also at A3 receptors). Crystal structures for the antagonist-bound [374, 876, 1198, 1765], agonist-bound [1126, 1127, 2154] and G protein-bound A2A adenosine receptors [193, 451, 602, 880, 990, 1016, 1071, 1485] have been described.

| Nomenclature | A1 receptor | A2A receptor | A2B receptor | A3 receptor |
|--------------|-------------|--------------|--------------|-------------|
| HGNC, UniProt| ADORA1, P30542 | ADORA2A, P29274 | ADORA2B, P29275 | ADORA3, P0DMS8 |
| Sub/family-selective agonists | NECA [602, 913, 1665, 1980, 2166] | NECA [193, 451, 602, 990, 1071, 2166] | NECA [148, 193, 905, 1179, 1870, 2024, 2166] | NECA [193, 905, 906, 1016, 1513, 1870] |
| Selective agonists | cyclopentyladenosine [404, 428, 602, 771, 880, 913, 1665], 5-Ci-3-deoxy-(±)-ENBA [565], TCPA [150], CCPA [880, 1485] | apadenoson [1548], UK-432,097 [2154], compound 4g [374], CGS 21680 [193, 451, 602, 880, 990, 1016, 1071, 1485], regadenoson [880] | BAY 60-6583 [488] | piclidenoson [537, 592, 1016, 2025], Cl-IB-MECA [208, 883, 987], MRS5698 [1977] |
| Sub/family-selective antagonists | CGS 15943 (pK9 8.5) [1513], xanthine amine congener (pK9 7.5) [565] | CGS 15943 (pK9 7.7–9.4) [451, 990, 1016, 1513], xanthine amine congener (pK9 8.4–9) [451, 1016] | xanthine amine congener (pK9 6.9–8.8) [148, 905, 906, 1016, 1179, 1870], CGS 15943 (pK9 6–8.1) [68, 905, 906, 1016, 1513, 1870] | CGS 15943 (pK9 7–7.9) [995, 1016, 1513, 2025], xanthine amine congener (pK9 7–7.4) [1016, 1707, 2166] |
(continued)

| Nomenclature | $A_1$ receptor | $A_{2A}$ receptor | $A_{2B}$ receptor | $A_3$ receptor |
|--------------|----------------|------------------|------------------|---------------|
| Selective antagonists | PSB36 ($pK_i$ 9.9) [6] – Rat, DPCPX ($pK_i$ 7.4–9.2) [428, 865, 1485, 1665, 2102], 8-phenyltheophylline ($pK_i$ 9) [939], WRC-0571 ($pK_i$ 8.8) [1272], DU172 ($pK_i$ 7.4) [649] | SCH442416 ($pK_i$ 8.4–10.3) [1796, 1969], ZM-241385 ($pK_i$ 8.8–9.1) [1513] | PSB-0788 ($pK_i$ 9.4) [192], PSB603 ($pK_i$ 9.3) [192], MRS1754 ($pK_i$ 8.8) [905, 994], PSB1115 ($pK_i$ 7.3) [757] | MRS1220 ($pK_i$ 8.2–9.2) [883, 995, 1892, 2177], VUF5574 ($pK_i$ 8.4) [2016], MRS1523 ($pK_i$ 7.7) [1158], MRS1191 ($pK_i$ 7.5) [883, 909, 1163] |
| Allosteric modulators | PD81723 (Positive) [239] | – | – | LUF6000 (Positive) [701], LUF6096 (Positive) [770] |
| Labelled ligands | $[^{3}H]$CCPA (Agonist) [1016, 1665], $[^{3}H]$DPCPX (Antagonist) ($pK_d$ 8.4–9.2) [404, 537, 1016, 1513, 1665, 1980] | $[^{3}H]$ZM 241385 (Antagonist) ($pK_i$ 8.7–9.1) [36, 600], $[^{3}H]$CGS 21680 (Agonist) [894, 2062] | $[^{3}H]$MR15754 (Antagonist) ($pK_i$ 9.8) [905] | $[^{125}I]$AB-MECA (Agonist) [1513, 2025] |

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 [884]. DPCPX also exhibits antagonism at $A_{2B}$ receptors ($pK_i$ ca. 7, [34, 1016]). Antagonists at $A_3$ receptors exhibit marked species differences, such that only MRS1523 and MRS1191 are selective at the rat $A_3$ receptor. In the absence of other adenosine receptors, $[^{3}H]$DPCPX and $[^{3}H]$ZM 241385 can also be used to label $A_{2B}$ receptors ($K_i$ ca. 30 and 60 nM respectively). $[^{125}I]$AB-MECA also binds to $A_1$ receptors [1016]. $[^{3}H]$CGS 21680 is relatively selective for $A_{2A}$ receptors, but may also bind to other sites in cerebral cortex [400, 914]. $[^{3}H]$NECA binds to other non-receptor elements, which also recognise adenosine [1209]. XAC-BY630 has been described as a fluorescent antagonist for labelling $A_1$ adenosine receptors in living cells, although activity at other adenosine receptors was not examined [217].

Further reading on Adenosine receptors

Fredholm BB et al. (2011) International Union of Basic and Clinical Pharmacology. LXXXI. Nomenclature and classification of adenosine receptors—an update. Pharmacol. Rev. 63:1-34 [PMID:21303899]
Guo D et al. (2017) Kinetic Aspects of the Interaction between Ligand and G Protein-Coupled Receptor: The Case of the Adenosine Receptors. Chem. Rev. 117:38-66 [PMID:27088232]
Göblyös A et al. (2011) Allosteric modulation of adenosine receptors. Biochim. Biophys. Acta 1808:1309-18 [PMID:20599682]
Headrick JP et al. (2011) Adenosine and its receptors in the heart: regulation, retaliation and adaptation. Biochim. Biophys. Acta 1808:1413-28 [PMID:21094127]
Lasley RD. (2011) Adenosine receptors and membrane microdomains. Biochim. Biophys. Acta 1808:1284-9 [PMID:20888790]
Mundell S et al. (2011) Adenosine receptor desensitization and trafficking. Biochim. Biophys. Acta 1808:1319-28 [PMID:20550943]
Wei CJ et al. (2011) Normal and abnormal functions of adenosine receptors in the central nervous system revealed by genetic knockout studies. Biochim. Biophys. Acta 1808:1358-79 [PMID:21185258]

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 GPCR autophosphorylation-inducing (GAIN) domain [56] 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 [571, 2187]. The nomenclature of these receptors was revised in 2015 as recommended by NC-IUPHAR and the Adhesion GPCR Consortium [718].

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

Adhesion Class GPCRs S37
| Nomenclature | HGNC, UniProt | Comments |
|--------------|--------------|----------|
| Nomenclature | HGNC, UniProt | Comments |
| ADGRA1 | ADGRA1, Q86SQ6 | – |
| ADGRA2 | ADGRA2, Q96PE1 | – |
| ADGRA3 | ADGRA3, Q8IWK6 | – |
| ADGRB1 | ADGRB1, O14514 | – |
| ADGRB2 | ADGRB2, O60241 | – |
| ADGRB3 | ADGRB3, O60242 | – |
| CELSR1 | CELSR1, Q9NYQ6 | – |
| Endogenous agonists | – | – |
| Comments | – | – |
| HGNC, UniProt | CELSR2, Q9HCU4 | – |
| CELSR3 | CELSR3, Q9NYQ7 | – |
| ADGRD1 | ADGRD1, Q6QNK2 | – |
| ADGRD2 | ADGRD2, Q7Z7M1 | – |
| ADGRE1 | ADGRE1, Q14246 | – |
| ADGRE2 | ADGRE2, Q9UHX3 | A mutation destabilizing the GAIN domain sensitizes mast cells to IgE-independent vibration-induced degranulation [202]. |
| ADGRE3 | ADGRE3, Q9BY15 | – |
| ADGRE4P | ADGRE4P, Q86SQ3 | – |
| ADGRES | ADGRES, P48960 | – |
| ADGRF1 | ADGRF1, QST601 | – |
| ADGRF2 | ADGRF2, Q8IZF7 | – |
| ADGRF3 | ADGRF3, Q8IZF5 | – |
| ADGRF4 | ADGRF4, Q8IZF3 | – |
| ADGRF5 | ADGRF5, Q8IZF2 | – |
| ADGRG1 | ADGRG1, Q9Y653 | Reported to bind tissue transglutaminase 2 [2155] and collagen, which activates the G12/13 pathway [1220]. |
| ADGRG2 | ADGRG2, Q8IZP9 | – |
| ADGRG3 | ADGRG3, Q86Y34 | – |
| ADGRG4 | ADGRG4, Q8IZF6 | – |
| ADGRG5 | ADGRG5, Q8IZF4 | – |

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)
Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Adrenoceptors

G protein-coupled receptors → Adrenoceptors

Overview: The nomenclature of the Adrenoceptors has been agreed by the NC-IUPHAR Subcommittee on Adrenoceptors [256], see also [789].

Adrenoceptors, \( \alpha \)

\( \alpha \)-Adrenoceptors are activated by the endogenous agonists (-)-adrenaline and (-)-noradrenaline. Phenylephrine, methoxamine and cirazoline are agonists and prazosin and cirazoline antagonists considered selective for \( \alpha_1 \)-relative to \( \alpha_2 \)-adrenoceptors. \([^{3}H]\)prazosin and \([^{125}I]\)HEAT (BE2254) are relatively selective radioligands. \( \text{S(+)-niguldipine} \) also has high affinity for L-type Ca\(^{2+} \) channels. Fluorescent derivatives of prazosin (Bodipy PLprazosin- QAPB) are used to examine cellular localisation of \( \alpha_3 \)-adrenoceptors. Selective \( \alpha_1 \)-adrenoceptor agonists are used as nasal decongestants; antagonists to treat hypertension (doxazosin, prazosin) and benign prostatic hyperplasia (alfuzosin, tamsulosin). The \( \alpha_1 \)- and \( \beta_2 \)-adrenoceptor antagonist carvedilol is used to treat congestive heart failure, although the contribution of \( \alpha_1 \)-adrenoceptor blockade to the therapeutic effect is unclear. Several anti-depressants and anti-psychotic drugs are \( \alpha_1 \)-adrenoceptor antagonists contributing to side effects such as orthostatic hypotension and extrapyramidal effects.

### Nomenclature

| Nomenclature | ADGRG6 | ADGRG7 | ADGRL1 | ADGRL2 | ADGRL3 | ADGRL4 | ADGRV1 |
|--------------|--------|--------|--------|--------|--------|--------|--------|
| HGNC, UniProt| ADGRG6, Q86SQ4| ADGRG7, Q96K78| ADGRL1, O94910| ADGRL2, O95490| ADGRL3, Q9HAR2| ADGRL4, Q9HBW9| ADGRV1, Q8WXG9|

Comments: Loss-of-function mutations are associated with Usher syndrome, a sensory deficit disorder [885].

---

Further reading on Adhesion Class GPCRs

Hamann J et al. (2015) International Union of Basic and Clinical Pharmacology. XCIV. Adhesion G protein-coupled receptors. Pharmacol. Rev. 67: 338-67 [PMID:25713288]

Yona S et al. (2008) Adhesion-GPCRs: emerging roles for novel receptors. Trends Biochem. Sci. 33: 491-500 [PMID:18789697]
### Adrenoceptors, $\alpha_2$

$\alpha_2$-Adrenoceptors are activated by (-)-adrenaline and with lower potency by (-)-noradrenaline. Brimonidine and talipexole are agonists and rauwolscine and yohimbine antagonists selective for $\alpha_2$-relative to $\alpha_1$-adrenoceptors. [3H]rauwolscine, [3H]brimonidine and [3H]RX821002 are relatively selective radioligands. There is species variation in the pharmacology of the $\alpha_{2A}$-adrenoceptor. Multiple mutations of $\alpha_2$-adrenoceptors have been described, some associated with alterations in function. Presynaptic $\alpha_2$-adrenoceptors regulate many functions in the nervous system. The $\alpha_2$-adrenoceptor agonists clonidine, guanabenz and brimonidine affect central baroreflex control (hypotension and bradycardia), induce hypnotic effects and analgesia, and modulate seizure activity and platelet aggregation. Clonidine is an anti-hypertensive and counteracts opioid withdrawal. Dexmedetomidine (also xylazine) is used as a sedative and analgesic in human and veterinary medicine with sympatholytic and anxiolytic properties. The $\alpha_2$-adrenoceptor antagonist yohimbine has been used to treat erectile dysfunction and mirtazapine as an anti-depressant. The $\alpha_{2B}$ subtype appears to be involved in neurotransmission in the spinal cord and $\alpha_{2C}$ in regulating catecholamine release from adrenal chromaffin cells.

| Nomenclature | $\alpha_{2A}$-adrenoceptor | $\alpha_{2B}$-adrenoceptor | $\alpha_{2C}$-adrenoceptor |
|--------------|-----------------------------|-----------------------------|-----------------------------|
| HGNC, UniProt| ADRA2A, P08913              | ADRA2B, P18089              | ADRA2C, P18825              |
| Endogenous agonists | (-)-adrenaline [896, 1573], (-)-noradrenaline [896, 1573] | (-)-noradrenaline (Partial agonist) [896, 1573], (-)-adrenaline [896] | (-)-noradrenaline [896, 1573], (-)-adrenaline [896] |
| Agonists | dexmedetomidine (Partial agonist) [896, 1228, 1552, 1573], clonidine (Partial agonist) [896, 1552, 1573], brimonidine [896, 1228, 1552, 1573], apraclonidine [1399], guanfacine (Partial agonist) [896, 1231] | dexmedetomidine [896, 1228, 1552, 1573], clonidine (Partial agonist) [896, 1552, 1573], brimonidine (Partial agonist) [896, 1552, 1573], guanfacine [896] | dexmedetomidine [896, 1552, 1573], brimonidine (Partial agonist) [896, 1228, 1552, 1573], apraclonidine [1399], guanfacine (Partial agonist) [896], guanfacine [896] |
| Selective agonists | oxymetazoline (Partial agonist) [896, 1228, 1998] | - | - |
| Antagonists | yohimbine (pK$_i$ 8.4–9.2) [255, 440, 1998] | yohimbine (pK$_i$ 7.9–8.9) [255, 440, 1998], phenoxybenzamine (pK$_i$ 8.5) [2088], tolazoline (pK$_i$ 5.5) [896] | yohimbine (pK$_i$ 8.5–9.5) [255, 440, 1998], WB 4101 (pK$_i$ 8.4–9.4) [255, 440, 1998], spiroxatrine (pK$_i$ 9) [1998], mirtazapine (pK$_i$ 7.7) [539], tolazoline (pK$_i$ 5.4) [896] |
| Selective antagonists | BRL 44408 (pK$_i$ 8.2–8.8) [1998, 2194] | imiloxan (pK$_i$ 7.3) [1329] – Rat | JP1302 (pK$_i$ 7.8) [1704] |
| Labelled ligands | - | - | [3H]MK-912 (Antagonist) (pK$_d$ 10.1) [1998] |
Adrenoceptors, β

β-Adrenoceptors are activated by the endogenous agonists (-)-adrenaline and (-)-noradrenaline. Isoproterenol is selective for β-1-adrenoceptors relative to α1- and α2-adrenoceptors, while propranolol (pKᵢ 8.2–9.2) and cyanopindolol (pKᵢ 10.0–11.0) are relatively β₁ and β₂ adrenoceptor-selective antagonists. (-)-Noradrenaline, xamoterol and (-)-Ro 363 show selectivity for β₁-adrenoceptors. Pharmacological differences exist between human and mouse β₂-adrenoceptors. Antagonists carvedilol (pKᵢ 9.5–9.9) [272], propranolol (pKᵢ 7.3–9) [272, 1210], levo-bunolol (pKᵢ 8.4) [71], labetalol (pKᵢ 8.2) [71], metoprolol (pKᵢ 7–7.8) [87, 272, 806, 1210], esmolol (pKᵢ 6.9) [71], nadolol (pKᵢ 6.9) [272], practolol (pKᵢ 6.1–6.8) [87, 1210], propranolol (pKᵢ 6.7) [71], sotalol (pKᵢ 6.1) [71] and (-)-adrenaline (pKᵢ 7.3) [272], timolol (pKᵢ 9.7) [87], propranolol (pKᵢ 9.1–9.5) [87, 90, 870, 1210], levo-bunolol (pKᵢ 9.3) [71], propranolol (pKᵢ 8.3–9.1) [272, 1210], alprenolol (pKᵢ 9) [87], nadolol (pKᵢ 7–8.6) [87, 272], labetalol (pKᵢ 8) [71], propranolol (pKᵢ 7.4) [71], sotalol (pKᵢ 6.5) [71] and carazolol [1318] are high affinity radioligands that label β₁- and β₂-adrenoceptors and β₁-adrenoceptors can be labelled with higher concentrations (nM) of [¹²⁵I]-cyanopindolol together with β₁- and β₂-adrenoceptor antagonists. [¹²⁵I]-L-748377 is a β₁-selective radioligand [2020]. Fluorescent ligands such as BODIPY-TMR-CGP12177 can be used to track β₁-adrenoceptors at the cellular level [8]. Somewhat selective β₁-adrenoceptor agonists (denopamine, dutabutamine) are used short term to treat cardiogenic shock but, chronically, reduce survival. β₁-Adrenoceptor-prefering antagonists are used to treat hypertension (atenolol, betaxolol, bisoprolol, metoprolol and nebivolol), cardiac arhythmias (atenolol, bisoprolol, esmolol) and cardiac failure (metoprolol, nebivolol). Cardiac failure is also treated with carvedilol that blocks β₁- and β₂-adrenoceptors, as well as α₁-adrenoceptors. Short (salbutamol, terbutaline) and long (formoterol, salmeterol) acting β₂-adrenoceptor-selective agonists are powerful bronchodilators used to treat respiratory disorders. Many first generation β-adrenoceptor antagonists (propranolol) block both β₁- and β₂-adrenoceptors and there are no β₂-adrenoceptor-selective antagonists used therapeutically. The β₂-adrenoceptor agonist mirabegron is used to control overactive bladder syndrome.
Physiological actions of adrenoceptors

Adrenoceptors, α<sub>1</sub>

The α<sub>1C</sub>-adrenoceptor corresponds to the pharmacologically defined α<sub>1A</sub>-adrenoceptor [789]. Some tissues possess α<sub>1A</sub>-adrenoceptors (α<sub>1</sub>-adrenoceptors [559, 1382]) that display relatively low affinity in functional and binding assays for prazosin indicative of different receptor states or locations. α<sub>1A</sub>-adrenoceptor C-terminal splice variants form homo- and heterodimers, but fail to generate a functional α<sub>1L</sub>-adrenoceptor [1628]. α<sub>1D</sub>-Adrenoceptors form heterodimers with α<sub>1B</sub>- or β<sub>2</sub>-adrenoceptors that show increased cell-surface expression [1993]. Recombinant α<sub>1D</sub>-adrenoceptors have been shown in some heterologous systems to be mainly located intracellularly but cell-surface localization is encouraged by truncation of the N-terminus, or by co-expression of α<sub>1A</sub>- or β<sub>2</sub>-adrenoceptors [706, 1993]. In blood vessels all three α<sub>1</sub>-adrenoceptor subtypes are located on the surface and intracellularly [1320, 1321]. Signalling is predominantly via G<sub>q</sub>/11 but α<sub>1</sub>-adrenoceptors also couple to G<sub>i/o</sub>, G<sub>8</sub> and G<sub>12/13</sub>. Several α<sub>1A</sub>-adrenoceptor agonists display ligand directed signalling bias relative to noradrenaline [521]. There are also differences between subtypes in coupling efficiency to different pathways. In vascular smooth muscle, the potency of agonists is differences between subtypes in coupling efficiency to different GM-coupled receptors. In vascular smooth muscle, the potency of agonists is differences between subtypes in coupling efficiency to different GM-coupled receptors. 

Adrenoceptors, β

ABC-239 and prazosin show selectivity for α<sub>2B</sub>- and α<sub>2C</sub>-adrenoceptors over α<sub>1</sub>-adrenoceptors. Oxymetazoline is a re-duced efficacy imidazole agonist but also binds to non-GPCR binding sites for imidazolines, classified as I<sub>1</sub>, I<sub>2</sub> and I<sub>3</sub> sites [406]; catecholamines have a low affinity, while rilmenidine and moxonidine are selective ligands evoking hypotensive effects in vivo. I<sub>1</sub>-imidazoline receptors cause central inhibition of sympathetic tone, I<sub>2</sub>-imidazoline receptors are an allosteric binding site on monoamine oxidase B, and I<sub>3</sub>-imidazoline receptors regulate insulin secretion from pancreatic β-cells. α<sub>2A</sub>-adrenoceptor stimulation reduces insulin secretion from β- islets [2171], 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 [1673]. α<sub>2A</sub>- and α<sub>2C</sub>-adrenoceptors form homodimers [1829]. Heterodimers between α<sub>2A</sub>- and either the α<sub>2C</sub>-adrenoceptor or μ opioid peptide receptor exhibit altered signalling and trafficking properties compared to the individual receptors [1829, 1931, 2036]. Signalling by α<sub>2</sub>-adrenoceptors is primarily via G<sub>i</sub>/<sub>o</sub>, although the α<sub>2A</sub>-adrenoceptor also couples to G<sub>8</sub> [487]. Imidazoline compounds display bias relative to each other at the α<sub>2A</sub>-adrenoceptor [1544]. The noradrenaline reuptake inhibitor desipramine acts directly on the α<sub>2A</sub>-adrenoceptor to promote internalisation via recruitment of arrestin [385].

Adrenoceptors, β

[125I]ICYP can be used to define β<sub>1</sub>- or β<sub>2</sub>-adrenoceptors when conducted in the presence of a β<sub>1</sub>- or β<sub>2</sub>-adrenoceptor-selective antagonist. A fluorescent analogue of CGP 12177 can be used to study β<sub>2</sub>-adrenoceptors in living cells [88]. [125I]ICYP at higher (nM) concentrations can be used to label β<sub>1</sub>-adrenoceptors in systems with few if any other β-adrenoceptor subtypes. The β<sub>1</sub>-adrenoceptor has an intron in the coding region, but splice variants have only been described for the mouse [522], where the isoforms display different signalling characteristics [850].

There are 3 β-adrenoceptors in turkey (termed the tβ, tβ3c and tβ4c) that have a pharmacology that differs from the human β-adrenoceptors [86]. Numerous polymorphisms have been described for the β-adrenoceptors; some are associated with signalling and trafficking, altered susceptibility to disease and/or altered responses to pharmacotherapy [1169]. All β-adrenoceptors couple to G<sub>i</sub> (activating adenylyl cyclase and elevating cAMP levels), but also activate G<sub>q</sub> and β-arrestin-mediated signalling. Many β<sub>1</sub>- and β<sub>2</sub>-adrenoceptor antagonists are agonists at β<sub>3</sub>-adrenoceptors (CL316243, CGP 12177 and carazolol). Many ‘antagonists’ of cAMP accumulation, for example carvedilol and bucindolol, weakly activate MAP kinase pathways [89, 523, 589, 590, 1721, 1722] and thus display ‘protean agonism.’ Bupranolol acts as a neutral antagonist in most systems so far examined. Agonists also display biased signalling at the β<sub>2</sub>-adrenoceptor via G<sub>i</sub> or arrests [470]. X-ray crystal structures have been described of the agonist bound [2075] and antagonist bound forms of the β<sub>1</sub>- [2076], agonist-bound [328] and antagonist-bound forms of the β<sub>2</sub>-adrenoceptor [1632, 1672], as well as a fully active agonist-bound, G<sub>i</sub> protein-coupled β<sub>2</sub>-adrenoceptor [1633]. Carvedilol and bucindolol bind to a site on the β<sub>1</sub>-adrenoceptor involving contacts in TM2, 3, and 7 and extracellular loop 2 that may facilitate coupling to arrestins [2076]. Compounds displaying arrestin-biased signalling at the β<sub>2</sub>-adrenoceptor have a greater effect on the conformation of TM7, whereas full agonists for G<sub>i</sub> coupling promote movement of TM5 and TM6 [1192]. Recent studies using NMR spectroscopy demonstrate significant conformational flexibility in the β<sub>3</sub>-adrenoceptor that is stabilized by both agonist and G proteins highlighting the dynamic nature of interactions with both ligand and downstream signalling partners [992, 1260, 1479]. Such flexibility likely has consequences for our understanding of biased agonism, and for the future therapeutic exploitation of this phenomenon.
Angiotensin receptors
G protein-coupled receptors \(\rightarrow\) Angiotensin receptors

**Overview:** The actions of angiotensin II (AGT, P01019) (Ang II) are mediated by AT\(_1\) and AT\(_2\) receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Angiotensin receptors [423, 950]), which have around 30% sequence similarity. The decapeptide angiotensin I (AGT, P01019), the octapeptide angiotensin II (AGT, P01019) and the heptapeptide angiotensin III (AGT, P01019) are endogenous ligands. Losartan, candesartan, telmisartan, etc. are clinically used AT\(_1\) receptor blockers.

| Nomenclature | AT\(_1\) receptor | AT\(_2\) receptor |
|--------------|-------------------|-------------------|
| HGNC, UniProt | AGTR1, P30556     | AGTR2, P50052     |
| Endogenous agonists | angiotensin II (AGT, P01019) [425, 2021], angiotensin III (AGT, P01019) [425] | angiotensin III (AGT, P01019) [394, 425, 2105], angiotensin II (AGT, P01019) [425, 1838, 2105], angiotensin-(1-7) (AGT, P01019) [194] |
| Selective agonists | L-162,313 [1559] | CGP42112 [194], [p-aminoPhe6]ang II [425, 1860] – Rat |
| Antagonists | telmisartan (pIC\(_{50}\) 8.4) [1303], olmesartan (pIC\(_{50}\) 8.1) [1027] | telmisartan (pIC\(_{50}\) 8.4–8.7) [2021], EXP3174 (pIC\(_{50}\) 7.4–9.5) [1965, 2021], eprosartan (pIC\(_{50}\) 8.4–8.8) [492], irbesartan (pIC\(_{50}\) 8.7–8.8) [2021], losartan (pIC\(_{50}\) 7.4–8.7) [425, 1965], valsartan (pIC\(_{50}\) 8.6) [424], azilsartan (pIC\(_{50}\) 8.1–8.1) [1623, 1917] | PD123177 (pIC\(_{50}\) 8.5–9.5) [305, 336, 478] – Rat, EMA401 (pIC\(_{50}\) 8.5–9.3) [543, 1656, 1836], PD123319 (pK\(_{d}\) 8.7–9.2) [425, 477, 2115] |
| Selective antagonists | candesartan (pIC\(_{50}\) 9.5–9.7) [2021], EXP3174 (pIC\(_{50}\) 7.4–9.5) [1965, 2021], eprosartan (pIC\(_{50}\) 8.4–8.8) [492], irbesartan (pIC\(_{50}\) 8.7–8.8) [2021], losartan (pIC\(_{50}\) 7.4–8.7) [425, 1965], valsartan (pIC\(_{50}\) 8.6) [424], azilsartan (pIC\(_{50}\) 8.1–8.1) [1623, 1917] | PD123177 (pIC\(_{50}\) 8.5–9.5) [305, 336, 478] – Rat, EMA401 (pIC\(_{50}\) 8.5–9.3) [543, 1656, 1836], PD123319 (pK\(_{d}\) 8.7–9.2) [425, 477, 2115] |
| Labelled ligands | \([^{3}H]\)A81988 (Antagonist) (pK\(_{d}\) 9.2) [725] – Rat, \([^{3}H]\)L158809 (Antagonist) (pK\(_{d}\) 9.2) [320] – Rat, \([^{3}H]\)Eprosartan (Antagonist) (pK\(_{d}\) 9.1) [22] – Rat, \([^{3}H]\)Valsartan (Antagonist) (pIC\(_{50}\) 8.8–9) [2034], \([^{125}I]\)EXP985 (Antagonist) (pK\(_{d}\) 8.8) [337] – Rat, \([^{3}H]\)Losartan (Antagonist) (pK\(_{d}\) 8.2) [309] – Rat | \([^{125}I]\)CGP42112 (A agonist) [425, 2105, 2106] |
| Comments | telmisartan and candesartan are also reported to be agonists of PPAR\(_{\gamma}\) [1877] | – |

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)
Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
**Comments:** AT₁ receptors are predominantly coupled to Gq/11, however they are also linked to arrestin recruitment and stimulate G protein-independent arrestin signalling [1221]. 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 [275]. The antagonist activity of CGP42112 at the AT₂ receptor has also been reported [1469]. 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) [1351].

**Further reading on Angiotensin receptors**

de Gasparo M et al. (2000) International Union of Pharmacology. XXIII. The angiotensin II receptors. Pharmacol. Rev. 52: 415-472 [PMID:10977869]
Karnik SS et al. (2015) International Union of Basic and Clinical Pharmacology. XCIX. Angiotensin Receptors: Interpreters of Pathophysiological Angiotensinergic Stimuli [corrected]. Pharmacol. Rev. 67: 754-819 [PMID:26315714]
Zhang H et al. (2015) Structural Basis for Ligand Recognition and Functional Selectivity at Angiotensin Receptor. J. Biol. Chem. 290: 29127-39 [PMID:26420482]
Zhang H et al. (2015) Structure of the Angiotensin receptor revealed by serial femtosecond crystallography. Cell 161: 833-44 [PMID:25913193]

**Apelin receptor**

**G protein-coupled receptors → Apelin receptor**

**Overview:** The apelin receptor (nomenclature as agreed by the NC-IUPHAR Subcommittee on the apelin receptor [1582]) 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 [1938]. A second family of peptides discovered independently and named Elabela [338] or Toddler, that has little sequence similarity to apelin, has been proposed as a second endogenous apelin receptor ligand [1542]. Structure-activity relationship Elabela analogues have been described [1406].

**Nomenclature**
apelin receptor

**HGNC, UniProt**
APLNR, P35414

**Potency order of endogenous ligands**
[Pyr₁]apelin-13 (APLN, Q9ULZ1) ≥ apelin-13 (APLN, Q9ULZ1) > apelin-36 (APLN, Q9ULZ1) [529, 1938]

**Endogenous agonists**
apelin-13 (APLN, Q9ULZ1) [529, 824, 1315], apelin receptor early endogenous ligand (APELA, P0DMC3) [436], apelin-17 (APLN, Q9ULZ1) [496, 1315], [Pyr₁]apelin-13 (APLN, Q9ULZ1) [961, 1315], Elabela/Toddler-21 (APELA, P0DMC3) [2174], Elabela/Toddler-32 (APELA, P0DMC3) [2174], apelin-36 (APLN, Q9ULZ1) [529, 824, 961, 1315], Elabela/Toddler-11 (APELA, P0DMC3) [2174]

**Selective agonists**
CMF-019 (Biased agonist) [1639], MM07 (Biased agonist) [209]

**Antagonists**
MM54 (pKi 8.2) [1227]

**Labelled ligands**
[¹²⁵I][Nle⁷⁵,Tyr⁷⁷]apelin-36 (human) (Agonist) [961], [¹²⁵I][Glp⁶⁵,Nle⁷⁵,Tyr⁷⁷]apelin-13 (Agonist) [824], [¹²⁵I][Pyr₁]apelin-13 (Agonist) [955], [¹²⁵I]apelin-13 (Agonist) [529], [³H][Pyr₁][Met(0)₁₁]-apelin-13 (Agonist) [1315]
**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 [293]. A modified apelin-13 peptide, apelin-13(F13A) was reported to block the hypotensive response to apelin in rat in vivo [1132], however, this peptide exhibits agonist activity in HEK293 cells stably expressing the recombinant apelin receptor [529].

**Further reading on Apelin receptor**

Cheng B et al. (2012) Neuroprotection of apelin and its signaling pathway. *Peptides* **37**: 171-3 [PMID:22820556]

Langelaan DN et al. (2009) Structural insight into G-protein coupled receptor binding by apelin. *Biochemistry* **48**: 537-48 [PMID:19123778]

O’Carroll AM et al. (2013) The apelin receptor APJ: journey from an orphan to a multifaceted regulator of homeostasis. *J. Endocrinol.* **219**: R13-35 [PMID:23943882]

Pitkin SL et al. (2010) International Union of Basic and Clinical Pharmacology. LXXIV. Apelin receptor nomenclature, distribution, pharmacology, and function. *Pharmacol. Rev.* **62**: 331-42 [PMID:20605969]

Yang P et al. (2015) Apelin, Elabela/Toddler, and biased agonists as novel therapeutic agents in the cardiovascular system. *Trends Pharmacol. Sci.* [PMID:26143239]

---

**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**

| HGNC, UniProt | GPBA receptor |
|--------------|---------------|

**Potency order of endogenous ligands**

| Lithocholic acid | Deoxycholic acid | Chenodeoxycholic acid, cholic acid [960, 1278] |

**Selective agonists**

- S-EMCA [1550] – Mouse, betulinic acid [621], oleanolic acid [1720]

**Comments**: The triterpenoid natural product betulinic acid has also been reported to inhibit inflammatory signalling through the NFκB pathway [1916]. Disruption of GPBA expression is reported to protect from cholesterol gallstone formation [2031]. A new series of 5-phenoxy-1,3-dimethyl-1H-pyrazole-4-carboxamides have been reported as highly potent agonists [1204].

**Further reading on Bile acid receptor**

Duboc H et al. (2014) The bile acid TGR5 membrane receptor: from basic research to clinical application. *Dig Liver Dis* **46**: 302-12 [PMID:24411485]

Lefebvre P et al. (2009) Role of bile acids and bile acid receptors in metabolic regulation. *Physiol. Rev.* **89**: 147-91 [PMID:19126757]

Lieu T et al. (2014) GPBA: a GPCR for bile acids and an emerging therapeutic target for disorders of digestion and sensation. *Br. J. Pharmacol.* **171**: 1156-66 [PMID:24111923]

van Nierop FS et al. (2016) Clinical relevance of the bile acid receptor TGR5 in metabolism. *Lancet Diabetes Endocrinol* [PMID:27639537]

---

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Bombesin receptors

G protein-coupled receptors → Bombesin receptors

**Overview:** Mammalian bombesin (Bn) receptors comprise 3 subtypes: BB₁, BB₂, BB₃ (nomenclature recommended by the NC-IUPHAR Subcommittee on bombesin receptors, [900]). BB₁ and BB₂ are activated by the endogenous ligands gastrin-releasing peptide (GRP, P07492) (GRP), neuropeptide B (NMB, P08949) (NMB) and GRP-(18-27) (GRP, P07492) (previously named neuropeptide C). Bombesin is a tetradecapeptide, originally derived from amphibians. The three Bn receptor subtypes couple primarily to the G₁₁ and G₁₂ family of G proteins [900] (but see also [908, 1995]). Each of these receptors is widely distributed in the CNS and peripheral tissues [659, 900, 1590, 1626, 1627, 1715, 1715, 2208]. 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 [900, 901, 902, 1248, 1371, 1626, 1627].

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, regulation of body temperature and other CNS behaviors, obesity, diabetes mellitus and growth of normal/neoplastic tissues [659, 1249, 1249, 1496, 1496, 2145].

| Nomenclature | BB₁ receptor | BB₂ receptor | BB₃ receptor |
|--------------|-------------|-------------|-------------|
| HGNC, UniProt | NMBR, P28336 | GRPR, P30550 | BRS3, P32247 |
| Endogenous agonists | neuropeptide B (NMB, P08949) [900, 1626, 1993] | neuropeptide C [1995], gastrin releasing peptide (14-27) (human) [1995] | |
| Selective agonists | – | – | compound 8a [1194], compound 9g [1284], MK-7725 [339], MK-5046 [1375, 1759], [D-Tyr⁶,Apa-⁴Cl₁₁,Phe₁₃,Nle₁₄]bombesin-(6-14) [1263], compound 17c [1283], compound 9f [1284], bag-1 [692], compound 22e [761], bag-2 [692] |
| Antagonists | D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Nal-NH₂ (pIC₅₀ 6.2–6.6) [658] | – | bantag-1 (pIC₅₀ 6.8–6.7) [692, 1375], ML-18 (pIC₅₀ 5.3) [1370] |
| Selective antagonists | PD 176252 (pIC₅₀ 9.3–9.8) [658], PD 168368 (pIC₅₀ 9.3–9.6) [658], dNal-cyc(Cys-Tyr-dTrp-Orn-Val)-Nal-NH₂ (pIC₅₀ 7.6–8.9) [658] | [D-Phe⁶, Leu¹³, Cpa¹⁴, ψ 13-14]bombesin-(6-14) (pKᵢ 9.8) [658], JMV641 (pIC₅₀ 9.3) [1970] – Mouse, (3-Ph-Pr⁶), His⁷,D-Ala¹¹,D-Pro¹³, ψ 13-14,Phe¹⁴]bombesin-(6-14) (pIC₅₀ 8.9) [1199, 1970] – Mouse, [D-Tpi⁶, Leu¹³, ψ(CH₂NH)-Leu¹⁴]bombesin-(6-14) (pIC₅₀ 8.9) [658], Ac-GRP-(20-26)-methylester (pIC₅₀ 8.7) [658] | |
| Labelled ligands | [¹²⁵I]BH-NMB (human, mouse, rat) (Agonist), [¹²⁵I][Tyr⁷]bombesin (Agonist) | [¹²⁵I][D-Tyr⁶]bombesin-(6-13)-methyl ester (Selective Antagonist) (pKᵢ 9.3) [1262] – Mouse, [¹²⁵I][Tyr⁷]bombesin (Agonist) [135], [¹²⁵I]GRP (human) (Agonist) | [³H]bag-2 (Agonist) [692] – Mouse, [¹²⁵I][D-Tyr⁶, ψ-Ala¹¹,Phe¹³,Nle¹⁴]bombesin-(6-14) (Agonist) [1264, 1375] |

**Comments:** All three human subtypes may be activated by [D-Phe⁶,β-Ala¹¹,Phe¹³,Nle¹⁴]bombesin-(6-14) [1264]. [D-Tyr⁶, Apa-⁴Cl₁₁,Phe¹³,Nle¹⁴] bombesin-(6-14) has more than 200-fold selectivity for BB₃ receptors over BB₁ and BB₂ [1263, 1264, 1626, 1627].

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp) Bombesin receptors S46
Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Further reading on Bombesin receptors

Ferreira CA et al. (2017) Radiolabeled bombesin derivatives for preclinical oncological imaging. *Biomed. Pharmacother.* **87**: 58-72 [PMID:28040598]
González N et al. (2015) Bombesin receptor subtype 3 as a potential target for obesity and diabetes. *Expert Opin. Ther. Targets* 1-18 [PMID:26066663]
Jensen RT et al. (2008) International Union of Pharmacology. LXVIII. Mammalian bombesin receptors: nomenclature, distribution, pharmacology, signaling, and functions in normal and disease states. *Pharmacol. Rev.* **60**: 1-42 [PMID:18055507]
Ramos-Álvarez I et al. (2013) Bombesin Peptides (Cancer). In *Handbook of Biologically Active Peptides. 2nd Revised edition.* Edited by Kastin AJ: Elsevier: 506-511 [ISBN: 9780123850959]
Sayegh AI. (2013) The role of bombesin and bombesin-related peptides in the short-term control of food intake. *Prog Mol Biol Transl Sci* **114**: 343-70 [PMID:23317790]

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 [1136]) 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 | B1 receptor | B2 receptor |
|--------------|-------------|-------------|
| HGNC, UniProt | *BDKRB1*, P46663 | *BDKRB2*, P30411 |

| Potency order of endogenous ligands | [des-Arg10]kallidin (KNG1, P01042) > [des-Arg9]bradykinin (KNG1, P01042) = kallidin (KNG1, P01042) > bradykinin (KNG1, P01042) |
|-------------------------------|----------------------------------------------------------------------------------------------------------------------------------|

| Endogenous agonists | [des-Arg10]kallidin (KNG1, P01042) [72, 110, 919] |
|---------------------|--------------------------------------------------|
| Selective agonists  | [Sar,D-Phe8,des-Arg9]bradykinin [919] |
| Antagonists         | [Leu9,des-Arg10]kallidin (pKi 9.1–9.3) [72, 110] |
| Selective antagonists| B-9958 (pK8 9.2–10.3) [630, 1642], R-914 (pA2 8.6) [650], R-715 (pA2 8.5) [651] |
| Labelled ligands    | [125I]Hpp-desArg10HOE140 (pKd 10), [3H]Lys-[des-Arg9]BK (Agonist), [3H]Lys-[Leu8][des-Arg9]BK (Antagonist) |

Further reading on Bradykinin receptors

Campos MM et al. (2006) Non-peptide antagonists for kinin B1 receptors: new insights into their therapeutic potential for the management of inflammation and pain. *Trends Pharmacol. Sci.* **27**: 646-51 [PMID:17056130]
Duchene J et al. (2009) The kinin B1 receptor and inflammation: new therapeutic target for cardiovascular disease. *Curr Opin Pharmacol* **9**: 125-31 [PMID:19124274]
Marceau F et al. (2004) Bradykinin receptor ligands: therapeutic perspectives. *Nat Rev Drug Discov* **3**: 845-52 [PMID:15459675]
Paquet JL et al. (1999) Pharmacological characterization of the bradykinin B2 receptor: inter-species variability and dissociation between binding and functional responses. *Br. J. Pharmacol.* **126**: 1083-90 [PMID:10204994]
Thornton E et al. (2010) Kinin receptor antagonists as potential neuroprotective agents in central nervous system injury. *Molecules* **15**: 6598-618 [PMID:20877247]

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)
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 [755, 1600]) are generated by the genes CALCR (which codes for the CT receptor) and CALCRL (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 proteins of ca. 130 amino acids, identified as a family of three members; RAMP1, RAMP2 and RAMP3. There are splice variants of the CT receptor, these in turn produce variants of the AMY receptor [1600], some of which can be potently activated by CGRP. The endogenous agonists are the peptides calcitonin (CALCA, P01258), a-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 sequences, particularly for the CTs. CTR-stimulating peptide{Pig} (CRSP) is another member of the family with selectivity for the CT receptor but it is not expressed in humans [952]. Olcegepant (also known as BIBN4096BS, pKi ~10.5) and telcagepant (also known as MK0974, pKi ~9) are the most selective antagonists available, showing selectivity for CGRP receptors, with a particular preference for those of primate origin. CLR (calcitonin receptor-like receptor) by itself binds no known endogenous ligand, but in the presence of RAMPs it gives receptors for CGRP, adrenomedullin and adrenomedullin 2/intermedin.

| Nomenclature          | CT receptor | AMY<sub>1</sub> receptor | AMY<sub>2</sub> receptor | AMY<sub>3</sub> receptor |
|-----------------------|-------------|--------------------------|--------------------------|--------------------------|
| HGNC, UniProt         | CALCR, P30988 | –                        | –                        | –                        |
| Subunits              | –           | CT receptor, RAMP1 (Accessory protein) | CT receptor, RAMP2 (Accessory protein) | CT receptor, RAMP3 (Accessory protein) |
| Potency order of endogenous ligands | calcitonin (salmon) > calcitonin (CALCA, P01258) > amylin (IAPP, P10997) > α-CGRP (CALCA, P06881) > adrenomedullin 2/intermedin (ADM2, Q7Z4H4) > calcitonin (CALCA, P01258) > adrenomedullin (ADM, P35318) | Poorly defined | calcitonin (salmon) > amylin (IAPP, P10997) > α-CGRP (CALCA, P06881) > adrenomedullin 2/intermedin (ADM2, Q7Z4H4) > calcitonin (CALCA, P01258) > adrenomedullin (ADM, P35318) |
| Endogenous agonists   | calcitonin (CALCA, P01258) [32, 62, 752, 1080, 1153, 1396] | α-CGRP (CALCA, P06881) [752, 1079, 1080, 1153, 2057], amylin (IAPP, P10997) [643] | amylin (IAPP, P10997) [643] | amylin (IAPP, P10997) [643] |
| Sub/family-selective agonists | pramlintide [643] | pramlintide [643] | pramlintide [643] | pramlintide [643] |
| Sub/family-selective antagonists | CT-(8-32) (salmon) (pK<sub>D</sub> 9) [793], AC187 (pK<sub>i</sub> 7.2) [752] | AC187 (pK<sub>i</sub> 8) [752], CT-(8-32) (salmon) (pK<sub>i</sub> 7.7) [752] | CT-(8-32) (salmon) (pK<sub>i</sub> 7.9) [752], AC187 (pK<sub>i</sub> 7.7) [752] | CT-(8-32) (salmon) (pK<sub>i</sub> 7.9) [752], AC187 (pK<sub>i</sub> 7.7) [752] |
| Labelled ligands      | [<sup>125</sup>I]CT (human) (Agonist), [<sup>125</sup>I]CT (salmon) (Agonist) | [<sup>125</sup>I]αCGRP (human) (Agonist), [<sup>125</sup>I]BH-AMY (rat, mouse) (Agonist) | [<sup>125</sup>I]BH-AMY (rat, mouse) (Agonist) | [125]BH-AMY (rat, mouse) (Agonist) |

**Searchable database:** http://www.guidetopharmacology.org/index.jsp

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

Calcitonin receptors  S48
**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 [353, 752, 753]. The major human CTR splice variant (hCT(a), which does not contain an insert) with RAMP1 (i.e. the AMY1(a) receptor) has a high affinity for CGRP [2057], unlike hCT(α2)-RAMP3 (i.e. AMY3(a) receptor) [353, 752]. Actions of CGRP at AMY (and the AM2) receptors led to proposals for a CGRP2 receptor in early literature [755]. However, the AMY receptor phenotype is RAMP-type, splice variant and cell-line-dependent [1376, 1614, 1964].

The ligands described have limited selectivity. Adrenomedullin has appreciable affinity for CCRP receptors. CGRP can show significant cross-reactivity at AMY receptors and AM2 receptors. Adrenomedullin 2/intermedin also has high affinity for the AM2 receptor [818]. CGRP-(8-37) acts as an antagonist of CGRP (pKz ≥ 8) and inhibits some AM and AMY responses (pKz 6.7). It is weak at CT receptors. Human AM-(22-52) has some selectivity towards AM receptors, but with modest potency (pKz 7), limiting its use [754]. Olcegepant shows the greatest selectivity between receptors but still has significant affinity for AMY1 receptors [2057].

**Further reading on Calcitonin receptors**

Booe JM et al. (2015) Structural Basis for Receptor Activity-Modifying Protein-Dependent Selective Peptide Recognition by a G Protein-Coupled Receptor. *Mol. Cell* 58: 1040-52 [PMID:25982113]

Hay DL et al. (2015) Amylin: Pharmacology, Physiology, and Clinical Potential. *Pharmacol. Rev.* 67: 564-600 [PMID:26071095]

Hay DL et al. (2016) Receptor Activity-Modifying Proteins (RAMPs): New Insights and Roles. *Annu. Rev. Pharmacol. Toxicol.* 56: 469-87 [PMID:26514202]

Kato J et al. (2015) Bench-to-bedside pharmacology of adrenomedullin. *Eur. J. Pharmacol.* 764: 140-8 [PMID:26144371]

Russell FA et al. (2014) Calcitonin gene-related peptide: physiology and pathophysiology. *Physiol. Rev.* 94: 1099-142 [PMID:25287861]

Russo AF. (2015) Calcitonin gene-related peptide (CGRP): a new target for migraine. *Annu. Rev. Pharmacol. Toxicol.* 55: S33-52 [PMID:25340934]

---

**Nomenclature**

- **HGNC, UniProt:**
- **Subunits:**
- **Potency order of endogenous ligands:**
- **Antagonists:**
- **Sub/family-selective antagonists:**
- **Labelled ligands:**

**CGRP receptor**

- **AM1 receptor**
- **AM2 receptor**

| Nomenclature | calcitonin receptor-like receptor | CGRP receptor | AM1 receptor | AM2 receptor |
|--------------|---------------------------------|---------------|--------------|--------------|
| HGNC, UniProt | CALCRL, Q16602                  |               |              |              |
| Subunits     | calcitonin receptor-like receptor, RAMP1 (Accessory protein) | calcitonin receptor-like receptor, RAMP2 (Accessory protein) | calcitonin receptor-like receptor, RAMP3 (Accessory protein) |
| Potency order of endogenous ligands | α-CGRP (CALCA, P06881) > adrenomedullin (ADM, P35318) > adrenomedullin 2/intermedin (ADM2, Q74H4) > amylin (IAPP, P10997) ≥ calcitonin (salmon) | adrenomedullin (ADM, P35318) > adrenomedullin 2/intermedin (ADM2, Q74H4) > α-CGRP (CALCA, P06881), amylin (IAPP, P10997) ≥ calcitonin (salmon) | adrenomedullin (ADM, P35318) ≥ adrenomedullin 2/intermedin (ADM2, Q74H4) ≥ α-CGRP (CALCA, P06881) > amylin (IAPP, P10997) > calcitonin (salmon) |
| Endogenous agonists | α-CGRP (CALCA, P10092) [21, 1313], α-CGRP (CALCA, P06881) [21, 1313] | adrenomedullin (ADM, P35318) [21, 1313] | adrenomedullin (ADM, P35318) [21, 1313] | adrenomedullin (ADM, P35318) [21, 568] |
| Antagonists | olcegepant (pKz 10.7-11) [462, 753, 754, 929, 1256], telcagepant (pKz 9.1) [1706] | AM-(22-52) (human) (pKz 7-7.8) [754] | – | – |
| Sub/family-selective antagonists | – | – | – | – |
| Labelled ligands | [125I]αCGRP (human) (Agonist), [125I]αCGRP (mouse, rat) (Agonist) | [125I]JAM (rat) (Agonist) | [125I]JAM (rat) (Agonist) | [125I]JAM (rat) (Agonist) |
Calcium-sensing receptor
G protein-coupled receptors → Calcium-sensing receptor

Overview: The calcium-sensing receptor (CaS, provisional nomenclature as recommended by NC-IUPHAR [557]) responds to multiple endogenous ligands, including extracellular calcium and other divalent/trivalent cations, polyamines and polycationic peptides, L-amino acids (particularly L-Trp and L-Phe), glutathione and various peptide analogues, ionic strength and extracellular pH (reviewed in [1122]). While divalent/trivalent cations, polyamines and polycations are CaS receptor agonists [234, 1618], L-amino acids, glutamyl peptides, ionic strength and pH are allosteric modulators of agonist function [375, 557, 803, 1616, 1617]. Indeed, L-amino acids have been identified as 'co-agonists', with both concomitant calcium and L-amino acid binding required for full receptor activation [623, 2205]. The sensitivity of the CaS receptor to primary agonists is increased by elevated extracellular pH [270] or decreased extracellular ionic strength [1617]. This receptor bears no sequence or structural relation to the plant calcium receptor, also called CaS.

| Nomenclature | CaS receptor |
|--------------|-------------|
| HGNC, UniProt | CASR, P41180 |
| 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) [375] |
| Cation rank order of potency | Gd3+ > Ca2+ > Mg2+ [234] |
| Glutamyl peptide rank order of potency | S-methylglutathione ≈ γGlu-Val-Gly > glutathione > γGlu-Cys [226, 1498, 2068] |
| Polyamine rank order of potency | spermine > spermidine > putrescine [1618] |
| Allosteric modulators | ATF 936 (Negative) (pIC50 8.9) [2109], encaleret (Negative) (pIC50 7.9) [1795], SB-423562 (Negative) (pIC50 7.1) [1074], ronacaleret (Negative) (pIC50 6.5–6.8) [92], NPS 2143 (Negative) (pKB 6.2–6.7) [418, 1120, 1123], cinacalcet (Positive) (pKB 5.9–6.6) [378, 418, 1120, 1123], tecalcet (Positive) (pKB 6.2–6.6) [378, 418], AC265347 (Positive) (pKB 6.3) [378, 1120], calhex 231 (Negative) (pIC50 6.4) [1569], calindol (Positive) (pKB 6.3) [378] |

Comments: The CaS receptor has a number of physiological functions, but it is best known for its central role in parathyroid and renal regulation of extracellular calcium homeostasis [728]. This is seen most clearly in patients with loss-of-function CaS receptor mutations who develop familial hypocalciuric hypercalcaemia (heterozygous mutations) or neonatal severe hyperparathyroidism (heterozygous, compound heterozygous or homozygous mutations) [728] and in Casr null mice [307, 803], which exhibit similar increases in PTH secretion and blood calcium levels. Gain-of-function CaS mutations are associated with autosomal dominant hypocalcaemia and Bartter syndrome type V [728]. The CaS receptor primarily couples to G13(11), G12(13) and Gi(0) [418, 634, 836, 1954], but in some cell types can couple to G5 [1258]. However, the CaS receptor can form heteromers with Class C GABAB [308, 327] and mGlu1/5 receptors [595], which may introduce further complexity in its signalling capabilities. Multiple other small molecule chemotypes are positive and negative allosteric modulators of the CaS receptor [980, 1441]. Further, etelcalcetide is a novel peptide agonist of the receptor [2059]. Agonists and positive allosteric modulators of the CaS receptor are termed Type I and II calcimimetics, respectively, and can suppress parathyroid hormone (PTH (PTH, P01270)) secretion [1443]. Negative allosteric modulators are called calcilytics and can act to increase PTH (PTH, P01270) secretion [1442].

Where functional pKB values are provided for allosteric modulators, this refers to ligand affinity determined in an assay that measures a functional readout of receptor activity (i.e. a receptor signalling assay), as opposed to affinity determined in a radioligand binding assay. The functional pKB may differ depending on the signalling pathway studied. Consult the ‘More detailed page’ for the assay description, as well as other functional readouts.

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Further reading on Calcium-sensing receptor

Breitwieser GE. (2012) Minireview: the intimate link between calcium sensing receptor trafficking and signaling: implications for disorders of calcium homeostasis. Mol. Endocrinol. 26: 1482-95 [PMID:22745192]
Brown EM. (2013) Role of the calcium-sensing receptor in extracellular calcium homeostasis. Best Pract. Res. Clin. Endocrinol. Metab. 27: 333-43 [PMID:23856263]

Conigrave AD et al. (2013) Calcium-sensing receptor (CaSR): pharmacological properties and signaling pathways. Best Pract. Res. Clin. Endocrinol. Metab. 27: 315-31 [PMID:23856262]
Nemeth EF et al. (2013) Calcimimetic and calcilytic drugs for treating bone and mineral-related disorders. Best Pract. Res. Clin. Endocrinol. Metab. 27: 373-84 [PMID:23856266]

Cannabinoid receptors → Cannabinoid receptors

Overview: Cannabinoid receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Cannabinoid Receptors [1564]) are activated by endogenous ligands that include N-arachidonoylethanolamine (anandamide), N-homo-γ-linolenoyl ethanolamine, N-docosatetra-7,10,13,16-enoyl ethanolamine 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]. There are currently three licenced cannabinoid medicines each of which contains a compound that can activate CB1 and CB2 receptors [1562]. Two of these medicines were developed to suppress nausea and vomiting produced by chemotherapy. These are nabilone (Cesamet®), a synthetic CB1/CB2 receptor agonist, and synthetic Δ⁹-tetrahydrocannabinol (Marinol®; dronabinol), which can also be used as an appetite stimulant. The third medicine, Sativex®, contains mainly Δ⁹-tetrahydrocannabinol and cannabidiol, both extracted from cannabis, and is used to treat multiple sclerosis and cancer pain.

| Nomenclature | CB₁ receptor | CB₂ receptor |
|--------------|--------------|--------------|
| HGNC, UniProt| CNR1, P21554 | CNR2, P34972 |
| Agonists     | cannabinol (Partial agonist) [535, 1801] | cannabinol (Partial agonist) [535, 1801] |
| Sub/family-selective agonists | HU-210 [535, 1801], CP55940 [535, 1676, 1801], WIN55212-2 [535, 1798, 1801], Δ⁹-tetrahydrocannabinol (Partial agonist) [535, 1801] | HU-210 [535, 1653, 1801], WIN55212-2 [535, 1798, 1801], CP55940 [535, 1676, 1801], Δ⁹-tetrahydrocannabinol (Partial agonist) [113, 535, 1653, 1801] |
| Selective agonists | arachidonyl-2-chloroethylamide [791] – Rat, arachidonylcyclopentylamide [791] – Rat, O-1812 [443] – Rat, R-(+)-methanandamide [976] – Rat | [WIN-133 [844, 1563], L-759,633 [607, 1676], AM1241 [2175], L-759,656 [607, 1676], HU-308 [734] |
| Selective antagonists | rimonabant (pKᵢ 7.9–8.7) [534, 535, 1660, 1687, 1801], AM251 (pKᵢ 8.1) [1094] – Rat, AM281 (pKᵢ 7.9) [1093] – Rat, LY320135 (pKᵢ 6.9) [534] | SR144528 (pKᵢ 8.3–9.2) [1661, 1676], AM-630 (pKᵢ 7.5) [1676] |
| Allosteric modulators | GAT100 (Negative) (pEC₅₀ 7.7) [1100], ZCZ011 (Positive) (pEC₅₀ 6.3) [1100] – Mouse, cannabinoids (Negative) [1100] | – |
| Labelled ligands | [³H]rimonabant (Antagonist) (pKᵢ 8.9–10) [211, 799, 932, 1568, 1662, 1811, 1948] – Rat | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Comments: Both CB1 and CB2 receptors may be labelled with $[^{3}H]CP55940$ (0.5 nM; [1801]) and $[^{3}H]WIN55212-2$ (2–2.4 nM; [1826, 1852]). Anandamide is also an agonist at vanilloid receptors (TRPV1) and PPARs [1484]. There is evidence for an allosteric site on the CB1 receptor [1603]. All of the compounds listed as antagonists behave as inverse agonists in some bioassay systems [1564]. For some cannabinoid receptor ligands, additional pharmacological targets that include GPR55 and GPR119 have been identified [1564]. 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 [1564].

Further reading on Cannabinoid receptors

Howlett AC et al. (2002) International Union of Pharmacology. XXVII. Classification of cannabinoid receptors. Pharmacol. Rev. 54: 161-202 [PMID:12037135]
Pertwee RG. (2010) Receptors and channels targeted by synthetic cannabinoid receptor agonists and antagonists. Curr. Med. Chem. 17: 1360-81 [PMID:20169277]
Pertwee RG et al. (2010) International Union of Basic and Clinical Pharmacology. LXXIX. Cannabinoid receptors and their ligands: beyond CB1 and CB2. Pharmacol. Rev. 62: 588-631 [PMID:21079038]

Chemerin receptor

G protein-coupled receptors → Chemerin receptor

Overview: The chemerin receptor (nomenclature as recommended by NC-IUPHAR [414]) is activated by the lipid-derived, anti-inflammatory ligand resolvin E1 (RvE1), which is the result of sequential metabolism of EPA by aspirin-modified cyclooxygenase and lipoxygenase [60, 61]. In addition, two GPCRs for resolvin D1 (RvD1) have been identified, FPR2/ALX, the lipoxin A4 receptor, and GPR32, an orphan receptor [1052].

| Nomenclature | chemerin receptor |
|--------------|------------------|
| HGNC, UniProt | CMKLR1, Q99788   |
| Potency order of endogenous ligands | resolvin E1 > chemerin C-terminal peptide > 18R-HEPE > EPA [60] |
| Selective agonists | resolvin E1 |
| Labelled ligands | $[^{3}H]$resolvin E1 (Agonist) [60, 61] |

Comments: CCX832 (structure not disclosed) is a selective antagonist, pKi=9.2 [969].
Overview: Chemokine receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Chemokine Receptors [81, 1402, 1403]) 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 receptors 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 [81].

Chemokines in turn can be divided by structure into four sub-classes by the number and arrangement of conserved cysteines. CC (also known as β-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 [82]. Listed are those human agonists with EC<50 values <50nM 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 receptors. 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 [2191] and aliases. Numerical data quoted are typically pK<sub>i</sub> or pIC<sub>50</sub> values from radioligand binding to heterologously expressed receptors.

### Nomenclature

| CCR1 | CCR2 | CCR3 |
|------|------|------|
| HGNC, UniProt | CCR1, P32246 | CCR2, P41597 |
| Endogenous agonists | CCL3 (CCL3, P10147) [342, 370, 783, 2228], CCL23 (CCL23, P55773) [342], CCL5 (CCL5, P13501) [370, 783], CCL7 (CCL7, P80098) [342, 703], CCL15 (CCL15, Q16663) [387], CCL14 (CCL14, Q16627) [342], CCL13 (CCL13, Q99616), CCL8 (CCL8, P80075) | CCL2 (CCL2, P13500) [387, 1224, 1347, 1533, 1996], CCL13 (CCL13, Q99616) [1224, 1996], CCL7 (CCL7, P80098) [387, 1224, 1996], CCL11 (CCL11, P51671) (Partial agonist) [1224, 1533], CCL16 (CCL16, O15467) |
| Agonists | – | – |
| Endogenous antagonists | CCL4 (CCL4, P13236) (pK<sub>i</sub> 7.1–7.8) [342, 370] | CCL26 (CCL26, Q9Y258) (pIC<sub>50</sub> 8.5) [1533] |
| Selective antagonists | BX 471 (pK<sub>i</sub> 8.2–9) [1164], compound 2b (pIC<sub>50</sub> 8.7) [1429], UCB36526 (pIC<sub>50</sub> 8) [1700], CP-481,715 (pK<sub>i</sub> 7.6) [646] | GSK Compound 34 (pK<sub>i</sub> 7.6) |
| Labelled ligands | [125I]CCL7 (human) (Agonist) [131], [125I]CCL3 (human) (Agonist) [131, 656, 1719], [125I]CCL5 (human) (Agonist) | [125I]CCL2 (human) (Agonist), [125I]CCL7 (human) (Agonist) |
| | | [125I]CCL11 (human) (Agonist) (pK<sub>d</sub> 8.3) [2063], [125I]CCL5 (human) (Agonist), [125I]CCL7 (human) (Agonist) |

**Searchable database:** http://www.guidetopharmacology.org/index.jsp

**Full Contents of ConciseGuide:** http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

**Chemokine receptors**

S.P.H. Alexander et al. The Concise Guide to PHARMACOLOGY 2017/18: G protein-coupled receptors. British Journal of Pharmacology (2017) 174, S17–S129
| Nomenclature | CCR4 | CCR5 | CCR6 | CCR7 | CCR8 | CCR9 | CCR10 |
|--------------|------|------|------|------|------|------|-------|
| HGNC, UniProt | CCR4, P51679 | CCR5, P51681 | CCR6, P51684 | CCR7, P32248 | CCR8, P51685 | CCR9, P51686 | CCR10, P46092 |
| Endogenous agonists | CCL22 (CCL22, O00626) [862], CCL17 (CCL17, Q92583) [862] | CCL5 (CCL5, P13501 ) [78, 1424, 1685], CCL4 (CCL4, P13236) [1424, 1685], CCL8 (CCL8, P80075) [1685], CCL3 (CCL3, P10147) [1424, 1685, 2228], CCL11 (CCL11, P51671) [161], CCL2 (CCL2, P13500) [1424], CCL14 (CCL14, Q16627) [1424], CCL16 (CCL16, O15467) | CCL20 (CCL20, P78556) [20, 77, 1598], beta-defensin 4A (DEFB4A, DEFB48, O15263) [2169] | CCL21 (CCL21, O00585) [2189], CCL19 (CCL19, Q99731) [1517, 2188, 2189] | CCL1 (CCL1, P22362) [403, 745, 863], CCL8 (Mouse) – Mouse | CCL25 (CCL25, O15444) | CCL27 (CCL27, Q9Y4X3) [816], CCL28 (CCL28, Q9NR3) |
| Agonists | vMIP-III | RS-HIV-1 gp120 | – | – | vMIP-I [403, 863] | – | – |
| Endogenous antagonists | – | CCL7 (CCL7, P80098) (pKᵢ 7.5) [1424] | – | – | – | – | – |
| Antagonists | – | vicriviroc (pKᵢ 9.1) [1879], ancriviroc (pKᵢ 7.8–8.7) [1237, 1523, 1879] | – | – | – | – | – |
| Selective antagonists | compound 8ic (pIC₅₀ 7.7) [2186], plerixafor (pIC₅₀ 6.2) [577] | E913 (pIC₅₀ 8.7) [1238], aplaviroc (pKᵢ 8.5) [1237], maraviroc (pIC₅₀ 8.1) [1424], TAK-779 (pKᵢ 7.5) [1237], MRK-1 [1073] – Rat | – | – | vMCC-I (pIC₅₀ 9.4) [403] | – | – |
| Selective allosteric modulators | – | – | – | – | – | vercirnon (Antagonist) (pIC₅₀ 8.2) [2060] | – |
| Antibodies | mogamulizumab (Inhibition) [54, 1799] | – | – | – | – | – | – |
| Labelled ligands | [¹²⁵I]CCL17 (human) (Agonist), [¹²⁵I]CCL27 (human) (Agonist) | [¹²⁵I]CCL4 (human) (Agonist) [1424], [¹²⁵I]CCL3 (human) (Agonist), [¹²⁵I]CCL5 (human) (Agonist), [¹²⁵I]CCL8 (human) (Agonist) | [¹²⁵I]CCL20 (human) (Agonist) [675] | [¹²⁵I]CCL19 (human) (Agonist), [¹²⁵I]CCL21 (human) (Agonist) [899] | [¹²⁵I]CCL1 (human) (Agonist) [863, 1671] | [¹²⁵I]CCL25 (human) (Agonist) |

Searchable database: http://www.guidetopharmacology.org/index.jsp  
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
| Nomenclature | HGN/C, UniProt | Endogenous agonists | Agonists | Selective agonists | Endogenous antagonists | Antagonists | Selective antagonists | Allosteric modulators | Labelled ligands |
|--------------|---------------|---------------------|----------|-------------------|------------------------|------------|----------------------|---------------------|-----------------|
| CXCR1        | CXCR1, P25024 | CXCL8 (CXCL8, P01045) [145, 711, 1133, 2121, 2137], CXCL6 (CXCL6, P80162) [2141] | vCXCL1 [1223], HIV-1 matrix protein p17 [637] | – | – | – | – | – |
| CXCR2        | CXCR2, P25025 | CXCL1 (CXCL1, P09241) [711, 1133, 2137], CXCL8 (CXCL8, P01045) [145, 711, 1133, 2121, 2137], CXCL7 (PPBP, P002775) [18], CXCL3 (CXCL3, P19876) [18], CXCL2 (CXCL2, P19875) [18], CXCL5 (CXCL5, P42830) [18], CXCL6 (CXCL6, P80162) [2141] | vCXCL1 [1223], HIV-1 matrix protein p17 [637] | – | – | – | – | – |
| CXCR3        | CXCR3, P49682 | CXCL11 (CXCL11, P14625) [768], CXCL10 (CXCL10, P002775) [18], CXCL3 (CXCL3, P19876) [18], CXCL5 (CXCL5, P42830) [18], CXCL6 (CXCL6, P80162) [2141] | – | – | – | – | – | – |
| CXCR4        | CXCR4, P61073 | CXCL12α (CXCL12, P48061) [782, 1202], CXCL12β (CXCL12, P48061) [782] | ALX40-4C (Partial agonist) [2213], X4-HIV-1 gp120 | – | – | – | – | – |
| CXCR5        | CXCR5, P32302 | – | – | – | – | – | – | – |
| CXCR6        | CXCR6, O00574 | – | – | – | – | – | – | – |
| CX3CR1       | CX3CR1, P49238 | – | – | – | – | – | – | – |

**Endogenous agonists**

- CXCL8 (CXCL8, P01045) [145, 711, 1133, 2121, 2137], CXCL6 (CXCL6, P80162) [2141]
- CXCL1 (CXCL1, P09241) [711, 1133, 2137], CXCL8 (CXCL8, P01045) [145, 711, 1133, 2121, 2137]
- CXCL7 (PPBP, P002775) [18], CXCL3 (CXCL3, P19876) [18], CXCL2 (CXCL2, P19875) [18]
- CXCL5 (CXCL5, P42830) [18], CXCL6 (CXCL6, P80162) [2141]

**Endogenous antagonists**

- CCL11 (CCL11, P51671) (pKᵢ 7.2) [2093], CCL7 (CCL7, P80098) (pKᵢ 6.6) [2093]

**Antagonists**

- Navarixin (pIC₅₀ 10.3) [81, 484], danirixin (pIC₅₀ 7.9) [1343], SB 225002 (pIC₅₀ 7.7) [2103]
- Elubirixin (pIC₅₀ 7.7) [81], SX-517 (pIC₅₀ 7.2) [1236]

**Selective antagonists**

- T134 (pIC₅₀ 8.4) [1929], X4P-001 (pIC₅₀ 7.9) [1819], HIV-Tat

**Allosteric modulators**

- Reparixin (Negative) (pIC₅₀ 9) [145]
- Reparixin (Negative) (pIC₅₀ 6.4) [145]

**Labelled ligands**

- [¹²⁵I]CXCL8 (human) (Agonist) [711, 1658]
- [¹²⁵I]CXCL8 (human) (Agonist) [711, 1658], [¹²⁵I]CXCL1 (human) (Agonist) [125],[¹²⁵I]CXCL5 (human) (Agonist) [125],[¹²⁵I]CXCL11 (human) (Agonist) [125], [¹²⁵I]CXCL12α (human) (Agonist) [444, 782], [¹²⁵I]CXCL13 (mouse) (Agonist) [227] – Mouse, [¹²⁵I]CXCL16 (human) (Agonist) [125], [¹²⁵I]CX3CL1 (human) (Agonist)
### Nomenclature

| Name   | HGNC, UniProt | Endogenous ligands | Endogenous agonists | Comments |
|--------|---------------|---------------------|---------------------|----------|
| XCR1   | XR1, P46094   | 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) | XCL1 (XCL1, P47992) [564], XCL2 (XCL2, Q9UBD3) [564] | XCL1 cannot be iodinated, but a secreted alkaline phosphatase (SEAP)-XCL1 fusion peptide can be used as a probe at XCR1. ACKR1 is used by *Plasmodium vivax* and *Plasmodium knowlesi* for entering erythrocytes. Several lines of evidence have suggested that adrenomedullin is a ligand for ACKR3; however, classical direct binding to the receptor has not yet been convincingly demonstrated. |
| ACKR1  | ACKR1, Q16570 | –                   | –                   | ACKR1 is used for treating 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 Hodgkin's Disease and multiple myeloma. |
| ACKR2  | ACKR2, O00590 | –                   | CCL2 (CCL2, P13500), CCL3 (CCL3, P10147), CCL4 (CCL4, P13236), CCL5 (CCL5, P13501), CCL7 (CCL7, P80098), CCL8 (CCL8, P80098), CCL11 (CCL11, P51671), CCL13 (CCL13, Q99616), CCL14 (CCL14, Q16627), CCL17 (CCL17, Q92583), CCL22 (CCL22, O00626) | – |
| ACKR3  | ACKR3, P25106 | –                   | –                   | – |
| ACKR4  | ACKR4, Q9NPB9 | –                   | –                   | – |
| CCLR2  | CCLR2, O00421 | –                   | –                   | – |

**Comments**: Specific chemokine receptors facilitate cell entry by microbes, such as ACKR1 for *Plasmodium vivax*, and CCR5 and CXCR4 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 gamma-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. 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 Hodgkin's Disease and multiple myeloma.

### Further reading on Chemokine receptors

- Bachelerie F *et al.* (2015) An atypical addition to the chemokine receptor nomenclature: IUPHAR Review “15”. *Br. J. Pharmacol.* [PMID:25958743]
- Koelink PJ *et al.* (2012) Targeting chemokine receptors in chronic inflammatory diseases: an extensive review. *Pharmacol. Ther.* 133: 1-18 [PMID:21839114]
- Murphy PM. (2002) International Union of Pharmacology. XXX. Update on chemokine receptor nomenclature. *Pharmacol. Rev.* 54: 227-9 [PMID:12037138]
- Murphy PM *et al.* (2000) International Union of Pharmacology. XXII. Nomenclature for chemokine receptors. *Pharmacol. Rev.* 52: 145-176 [PMID:10699158]
- Scholten DJ *et al.* (2012) Pharmacological modulation of chemokine receptor function. *Br. J. Pharmacol.* 165: 1617-43 [PMID:21699506]

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Cholecystokinin receptors

Overview: Cholecystokinin receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on CCK receptors [1471]) 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 [1038, 2073], 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 each CCK and gastrin peptides. These receptors have characteristic and distinct distributions, with both present in both the central nervous system and peripheral tissues.

| Nomenclature | CCK1 receptor | CCK2 receptor |
|--------------|---------------|---------------|
| HGNC, UniProt | CCKAR, P32238 | CCKBR, P32239 |
| Potency order of endogenous ligands | 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 [1135], gastrin-17 (GAST, P01350) [845], desulfated cholecystokinin-8, CCK-4 (CCK, P06307) |
| Selective agonists | A-71623 [67] – Rat, JMV180 [971], GW-5823 [722] | RB-400 [129] – Rat, PBC-264 [886] – Rat |
| Antagonists | lintitript (pIC$_{50}$ 8.3) [667] | – |
| Selective antagonists | devazepide (pIC$_{50}$ 9.7) [845] – Rat, T-0632 (pIC$_{50}$ 9.6) [1935] – Rat, PD-140548 (pIC$_{50}$ 8.8) [1817] – Rat, lorglumide (pIC$_{50}$ 6.7–8.2) [845, 875] – Rat | YF-476 (pIC$_{50}$ 9.7) [201, 1927], GV150013 (pIC$_{50}$ 9.4) [2006], L-740093 (pIC$_{50}$ 9.2) [1464], YM-022 (pIC$_{50}$ 9.2) [1464], Nj-26070109 (pIC$_{50}$ 8.5) [1390], L-365260 (pIC$_{50}$ 8.4) [1135], RP73870 (pIC$_{50}$ 8.3) [1181] – Rat, LY262691 (pIC$_{50}$ 7.5) [1632] – Rat |
| Labelled ligands | $^{3}$H]ldevazepide (Antagonist) (pK$_{d}$ 9.7) [306], $^{[125]}$IDTyr-Gly-[Nle28,31]CCK-26-33 (Agonist) [1599] | $^{[3]}$H]PD140376 (Antagonist) (pK$_{i}$ 9.7–10) [849] – Guinea pig, $^{[125]}$JPD142308 (Antagonist) (pK$_{d}$ 9.6) [820] – Guinea pig, $^{[125]}$IDTyr-Gly-[Nle28,31]CCK-26-33 (Agonist) [1599], $^{[125]}$Jgastrin (Agonist), $^{[3]}$H]gastrin (Agonist), $^{[3]}$H]365260 (Antagonist) (pK$_{d}$ 8.2–8.5) [1464], $^{[125]}$BDZ2 (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 [1833], 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 [1850], 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.

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Further reading on Cholecystokinin receptors

Cawston EE et al. (2010) Therapeutic potential for novel drugs targeting the type 1 cholecystokinin receptor. Br. J. Pharmacol. 159: 1009-21 [PMID:19922535]

Dockray GJ. (2009) Cholecystokinin and gut-brain signalling. Regul. Pept. 155: 6-10 [PMID:19345244]

Dufresne M et al. (2006) Cholecystokinin and gastrin receptors. Physiol. Rev. 86: 805-47 [PMID:16816139]

Miller LJ et al. (2008) Structural basis of cholecystokinin receptor binding and regulation. Pharmacol. Ther. 119: 83-95 [PMID:1858433]

Class Frizzled GPCRs

Overview: Receptors of the Class Frizzled (FZD, nomenclature as agreed by the NC-IUPHAR subcommittee on the Class Frizzled GPCRs [1747]), are GPCRs originally identified in Drosophila [300], which are highly conserved across species. While SMO shows structural resemblance to the 10 FZDs, it is functionally separated as it mediates effects in the Hedgehog signalling pathway [1747]. 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 LRPS (O75197) and LRPS6 (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 heterotrimeric G proteins [447], the elevation of intracellular calcium [1828], activation of cGMP-specific PDE6 [19] and elevation of cAMP as well as RAC-1, JNK, Rho and Rho kinase signalling [730]. Furthermore, 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 [321], as well as for β-catenin-dependent [242] and -independent [243,986] 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 FZD1 FZD2 FZD3 FZD4 FZD5 FZD6 FZD7
HGNC, UniProt FZD1, Q9UP38 FZD2, Q14332 FZD3, Q9NPG1 FZD4, Q9ULV1 FZD5, Q13467 FZD6, Q60335 FZD7, Q75084

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

Class Frizzled GPCRs S58
**Nomenclature**

|            | FZD<sub>8</sub> | FZD<sub>9</sub> | FZD<sub>10</sub> | SMO |
|------------|----------------|----------------|----------------|-----|
| HGNC, UniProt | FZD8, Q9H461  | FZD9, O00144  | FZD10, Q9ULW2 | SMO, Q99835 |

**Antagonists**
- –
- –
- saridegib (pIC<sub>50</sub> 8.9) [1981], glasdegib (pIC<sub>50</sub> 8.3) [1398], sonidegib (pKi 8.2) [2065]

**Selective antagonists**
- –
- –
- vismodegib (pK<sub>i</sub> 7.8) [2065]

**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 [447]). There is also a scarcity of information on basic pharmacological characteristics of FZDs, such as binding constants, ligand specificity or concentration-response relationships [984].

**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, Q9H17), 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, Q19405) (also known as WNT-15 or WNT-14b), Wnt-10a (WNT10A, Q9GZT5), Wnt-10b (WNT10B, O00744) (also known as WNT-12), Wnt-11 (WNT11, Q96014) and Wnt-16 (WNT16, Q9UBV4).

**Extracellular proteins that interact with FZDs**: norrin (NDP, Q90604), R-spondin-1 (RSPO1, Q2MKA7), R-spondin-2 (RSPO2, Q6UXX9), R-spondin-3 (RSPO3, Q9BXY4), R-spondin-4 (RSPO4, Q210MS), 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 LRP**: Dickkopf 1 (DKK1, Q94907), WIF1 (Q9Y5W5), sclerostin (SOST, Q99QB4), kremen 1 (KREMEN1, Q96MU8) and kremen 2 (KREMEN2, Q8NCW0)

**Small exogenous ligands**: Foxy-5 [1910], Box-5, UM206 [1086], and XWnt8 (P28026) also known as mini-Wnt8.

**Further reading on Class Frizzled GPCRs**

Angers S et al. (2009) Proximal events in Wnt signal transduction. Nat. Rev. Mol. Cell Biol. 10: 468-77 [PMID:19536106]

Schulte G. (2015) Frizzleds and WNT/β-catenin signaling—The black box of ligand-receptor selectivity, complex stoichiometry and activation kinetics. Eur. J. Pharmacol. 763: 191-5 [PMID:26003275]

van Amerongen R. (2012) Alternative Wnt pathways and receptors. Cold Spring Harb Perspect Biol 4: [PMID:22935094]

Wang Y et al. (2016) Frizzled Receptors in Development and Disease. Curr. Top. Dev. Biol. 117: 113-39 [PMID:26969975]

---

**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 [1015]) are activated by the endogenous ~75 amino-acid anaphylatoxin polypeptides C3a (C3, P01024) and C5a (C5, P01031), generated upon stimulation of the complement cascade.

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

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature C3a receptor C5a receptor C5a2 receptor HGNC, UniProt C3AR1, Q16581 CSAR1, P21730 CSAR2, Q9P296 Potency order of endogenous ligands C3a (C3, P01024) > C5a (C5, P01031) [41] C5a (C5, P01031), C5a des-Arg (C5) > C3a (C3, P01024) [41] C5a (C5, P01031), C5a des-Arg (C5) [125I] C5a (C5, P01031), C5a des-Arg (C5) [2160] Endogenous agonists ribosomal protein S19 [RP57], P39019) [2160] N-methyl-Phe-Lys-Pro-D-Cha-Cha-D-Arg-CO2H [125I]C5a (human) (Agonist) [310] 

Agonists E7 [43], compound 17 [1644], compound 21 [1643], Ac-RHYPLWR [707] N-methyl-Phe-Lys-Pro-D-Cha-D-Arg-CO2H [959, 1035] N-methyl-Phe-Lys-Pro-D-Cha-Trp-D-Arg-CO2H [1035] 

Selective agonists–– P59 (Biased agonist) [396], P32 (Biased agonist) [396]–

Antagonists SB290157 (pIC50 7.6) [40], compound 4 (pIC50 5.9) [1643] avacopan (pIC50 9.7) [125], WS5011 (pKi 8.7) [1893], DF2593A (pIC50 8.3) [2128], N-methyl-Phe-Lys-Pro-D-Cha-D-Arg-CO2H (pIC50 7.2) [1035]–

Labelled ligands [125I]C3a (human) (Agonist) [310] [125I]CSa (human) (Agonist) [843] [125I]C5a (human) (Agonist)Antagonists SB290157 (pIC50 7.6) [40], compound 4 (pIC50 5.9) [1643] avacopan (pIC50 9.7) [125], WS5011 (pKi 8.7) [1893], DF2593A (pIC50 8.3) [2128], N-methyl-Phe-Lys-Pro-D-Cha-D-Arg-CO2H (pIC50 7.2) [1035]–

Labelled ligands [125I]C3a (human) (Agonist) [310] [125I]CSa (human) (Agonist) [843] [125I]C5a (human) (Agonist)

**Comments:** SB290157 has also been reported to have agonist properties at the C3a receptor [1282]. The putative chemoattractant receptor termed C5a2 (also known as GPR77, C5L2) binds 125I C3a with no clear signalling function, but has a putative role opposing inflammatory responses [267, 599, 616]. Binding to this site may be displaced with the rank order C5a des-Arg (C5) > C5a (C5, P01031) while there is controversy over the ability of C3a (C3, P01024) and C3a des Arg (C3, P01024) to compete [817, 936, 937, 1508]. C5a2 appears to lack G protein signalling and has been termed a decoy receptor [1753]. However, C5a2 does recruit arrestin after ligand binding, which might provide a signalling pathway for this receptor [94, 2015], and forms heteromers with C5a1. C5a, but not C5a-des Arg, induces upregulation of heteromer formation between complement C5a receptors C5a1 and C5a2 [395]. There are also reports of pro-inflammatory activity of C5a2, mediated by HMGB1, but the signalling pathway that underlies this is currently unclear (reviewed in [1161]). More recently, work in T cells has shown that C5a2 and C5a2 act in opposition to each other and that altering the equilibrium between the two receptors, by differential expression or production of C5a-des Arg (which favours C5a2), can affect the final cellular response [57].

Further reading on Complement peptide receptors

- Arbore G et al. (2016) A novel “complement-metabolism-inflammasome axis” as a key regulator of immune cell effector function. *Eur. J. Immunol.* 46: 1563-73 [PMID:27184294]
- Klos A et al. (2013) International Union of Pharmacology. LXXXVII. Complement peptide C5a, C4a, and C3a receptors. *Pharmacol. Rev.* 65: 500-43 [PMID:23383423]
- Monk PN et al. (2007) Function, structure and therapeutic potential of complement C5a receptors. *Br. J. Pharmacol.* 152: 429-48 [PMID:17603557]

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 [750]) receptors are activated by the endogenous peptides corticotrophin-releasing hormone (CRH, P06850), a 41 amino-acid peptide, urocaritin 1 (UCN, P55089), 40 amino-acids, urocaritin 2 (UCN2, Q96RP3), 38 amino-acids and urocaritin 3 (UCN3, Q969P3), 38 amino-acids. CRF1 and CRF2 receptors are activated non-selectively by corticotrophin-releasing hormone (CRH, P06850) and urocaritin 1 (UCN, P55089). Binding to CRF receptors can be conducted using 125ITyr0-CRF or 125ITyr1-sauvagine with Kd 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: CRF₁ receptor
HGNC, UniProt: CRHR1, P34998
Endogenous agonists: –
Antagonists: SSR125543A (pKᵢ 8.7) [698]
Selective antagonists: CP 154,526 (pIC₅₀ 9.3–10.4) [1218] – Rat, DMP696 (pKᵢ 8.3–9) [760], NBI27914 (pKᵢ 8.3–9) [314], R121919 (pKᵢ 8.3–9) [2227], antalarmin (pKᵢ 8.3–9) [2087], CP376395 (pIC₅₀ 8.3) [322] – Rat, CRA1000 (pIC₅₀ 6.4–7.1) [298]

Comments: A CRF binding protein has been identified (CRHBP, P24387) to which both corticotrophin-releasing hormone (CRH, P06850) and urocaritin 1 (UCN, P55089) bind with high affinities, which has been suggested to bind and inactivate circulating corticotrophin-releasing hormone (CRH, P06850) [1558].

Further reading on Corticotropin-releasing factor receptors
Grammatopoulos DK. (2012) Insights into mechanisms of corticotropin-releasing hormone receptor signal transduction. Br. J. Pharmacol. 166: 85-97 [PMID:21883143]
Gysling K. (2012) Relevance of both type-1 and type-2 corticotropin releasing factor receptors in stress-induced relapse to cocaine seeking behaviour. Biochem. Pharmacol. 83:1 - 5 [PMID:21843515]
Hauger RL et al. (2003) International Union of Pharmacology. XXXVI. Current status of the nomenclature for receptors for corticotropin-releasing factor and their ligands. Pharmacol Rev. 55: 21-26 [PMID:12615952]

Valentino RJ et al. (2013) Sex-biased stress signaling: the corticotropin-releasing factor receptor as a model. Mol. Pharmacol. 83: 737-45 [PMID:23239826]
Zhu H et al. (2011) Corticotropin-releasing factor family and its receptors: pro-inflammatory or anti-inflammatory targets in the periphery? Inflamm. Res. 60: 715-21 [PMID:21476084]

Dopamine receptors
G protein-coupled receptors → Dopamine receptors

Overview: Dopamine receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Dopamine Receptors [1748]) 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
HGNC, UniProt: DRD1, P21728
Sub/family-selective labelled ligands: [¹²⁵I]SCH23982 (Antagonist) (pKᵦ 9.5) [433], [³H]SCH-23390 (Antagonist) (pKᵦ 9.5) [2221]
Endogenous agonists: dopamine [1897, 1962]
Agonists: fenoldopam [1962]

D₂ receptor
HGNC, UniProt: DRD2, P14416
Sub/family-selective labelled ligands: [³H]spiperone (Antagonist) (pKᵦ 10.2) [246, 805, 2219] – Rat
Endogenous agonists: dopamine [252, 573, 1725]
Agonists: rotigotine [448], cabergoline (Partial agonist) [1337], aripiprazole (Partial agonist) [2199], bromocriptine [573, 1337, 1725], MLS1547 (Biased agonist) [572], ropinirole [766], apomorphine (Partial agonist) [252, 573, 1337, 1725, 1844], pramipexole [1332, 1725], benzquinamide [677]

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature

Sub/family-selective agonists
- D₁ receptor: A68930 [1445], SKF-38393 (Partial agonist) [1897, 1962]
- D₂ receptor: quinpirole [252, 1332, 1539, 1844, 1846, 2019]

Selective agonists
- SKF-83959 (Biased agonist) [377], SKF-81297 (Partial agonist) [47] – Rat

Antagonists
- flupentixol (pKᵢ 7–8.4) [1897, 1962]
- haloperidol (pKᵢ 7.4–8.8) [573, 1230, 1332, 1844, 1963]

Sub/family-selective antagonists
- SCH-23390 (pKᵢ 7.4–9.5) [1897, 1962], SKF-83566 (pKᵢ 9.5) [1897], ecopipam (pKᵢ 8.3) [1963]

Selective antagonists
- L-741,626 (pKᵢ 7.9–8.5) [688, 1069], domperidone (pKᵢ 7.1–7.6) [573, 1844], ML321 (pKᵢ 7) [2147, 2148]

Labelled ligands
- [³H]raclopride (Antagonist) (pKᵢ 8.9) [1081] – Rat

Nomenclature

HGNC, UniProt
- D₃ receptor: DRD3, P35462
- D₄ receptor: DRD4, P21917
- D₅ receptor: DRD5, P21918

Endogenous agonists
- dopamine [252, 573, 1725, 1846]

Agonists
- pramipexole [1332, 1725], bromocriptine (Partial agonist) [573, 1337, 1725], ropinirole [766], apomorphine (Partial agonist) [252, 573, 1337, 1725, 1844]

Sub/family-selective agonists
- quinpirole [252, 1332, 1539, 1725, 1844, 1846, 2019]

Selective antagonists
- PD 128907 [1610, 1725] – Rat, A412997 [1373] – Rat, A412997 [1373]

Antagonists
- perospirone (pKᵢ 10.1) [1764], sertindole (pKᵢ 7.8–9.1) [253, 1761, 1763, 1764], sonepiprazole (pKᵢ 8.9) [1739], loxapine (pKᵢ 8.1) [1763]

Sub/family-selective antagonists
- haloperidol (pKᵢ 7.5–8.8) [573, 1782, 1844, 1963]

Selective antagonists
- S33084 (pKᵢ 9.6) [1336], nafadotride (pKᵢ 9.5) [1726], PG01037 (pKᵢ 9.2) [689], NCGR 2904 (pKᵢ 8.8) [2143], SB 277011-A (pKᵢ 8) [1641], (+)-5-14297 (pKᵢ 6.9–7.9) [1334, 1339], L745870 (pKᵢ 9.4) [1069], A-381393 (pKᵢ 8.8) [1420], L741742 (pKᵢ 8.5) [1683], ML398 (pKᵢ 7.4) [142]
Nomenclature | D₃ receptor | D₄ receptor | D₅ receptor
--- | --- | --- | ---
Selective allosteric modulators | SB269652 (Negative) (pKᵢ ~ 9) [588] | – | –
Labelled ligands | [³H]spiperone (Antagonist) (pKᵢ 9.9) [805, 2219] – Rat, [³H]7-OH-DPAT (Agonist) [1655], [³H]PD128907 (Agonist) [27] | [¹²⁵]JL750667 (Antagonist) (pKᵢ 9.8) [1539], [¹²⁵]JNGD941 (Antagonist) (pKᵢ 8.3) [1604] | [¹²⁵]J SCH23982 (Antagonist) (pKᵢ 9.1)

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 D₂S and D₂L (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.

Further reading on Dopamine receptors
Beaulieu JM et al. (2015) Dopamine receptors - IUPHAR Review 13. Br. J. Pharmacol. 172:1 - 23 [PMID:25671228]
Beaulieu JM et al. (2011) The physiology, signaling, and pharmacology of dopamine receptors. Pharmacol. Rev. 63:182-217 [PMID:21303898]
Cumming F. (2011) Absolute abundances and affinity states of dopamine receptors in mammalian brain: A review. Synapse 65:892-909 [PMID:21308799]
Maggio R et al. (2010) Dopamine D2-D3 receptor heteromers: pharmacological properties and therapeutic significance. Curr Opin Pharmacol 10:100-7 [PMID:19896900]
Ptácek R et al. (2011) Dopamine D4 receptor gene DRD4 and its association with psychiatric disorders. Med. Sci. Monit. 17:RA215-20 [PMID:21873960]
Schwartz J-C et al. (1998) Dopamine Receptors. In The IUPHAR Compendium of Receptor Characterization and Classification Edited by Girdlestone D: IUPHAR Media: 141-151
Undieh AS. (2010) Pharmacology of signaling induced by dopamine D(1)-like receptor activation. Pharmacol. Ther. 128:37-60 [PMID:20547182]

Endothelin receptors
G protein-coupled receptors → Endothelin receptors

Overview: Endothelin receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Endothelin Receptors [413]) 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)).
(continued)

**Nomenclature**

| Sub/family-selective antagonists | ET<sub>A</sub> receptor | ET<sub>B</sub> receptor |
|---------------------------------|------------------------|------------------------|
| SB209670 (pK<sub>B</sub> 9.4) [502] – Rat, TAK 044 (pA<sub>2</sub> 8.4) [2081] – Rat, bosentan (pA<sub>2</sub> 7.2) [367] – Rat | SB209670 (pK<sub>B</sub> 9.4) [502] – Rat, TAK 044 (pA<sub>2</sub> 8.4) [2081] – Rat, bosentan (pK<sub>B</sub> 7.1) [1405] |

**Selective antagonists**

| macitentan (pIC<sub>50</sub> 9.3) [177], sitaxsentan (pA<sub>2</sub> 8) [2135], FR139317 (Inverse agonist) (pIC<sub>50</sub> 7.3–7.9) [1242], BQ123 (pA<sub>2</sub> 6.9–7.4) [1242], ambrisentan (pA<sub>2</sub> 7.1) [178] | A192621 (pK<sub>d</sub> 8.1) [2043], BQ788 (pK<sub>d</sub> 7.9–8) [1690], IRL 2500 (pK<sub>d</sub> 7.2) [1690], Ro 46-8443 (pIC<sub>50</sub> 7.2) [215] |

**Labelled ligands**

| [125I]PD164333 (Antagonist) (pK<sub>d</sub> 9.6–9.8) [416], [3H]S0139 (Antagonist) (pK<sub>d</sub> 9.2), [125I]PD151242 (Antagonist) (pK<sub>d</sub> 9–9.1) [417], [3H]BQ123 (Antagonist) (pK<sub>d</sub> 8.5) [858] | [125I]IRL1620 (Agonist) [1421], [125I]BQ3020 (Agonist) [737, 1354, 1565], [125I]Ala<sup>1</sup>,3<sup>11</sup>,15<sup>15</sup>ET-1 (Agonist) [1354] |

**Comments**: Splice variants of the ET<sub>A</sub> receptor have been identified in rat pituitary cells; one of these, ET<sub>A</sub>R-C13, appeared to show loss of function with comparable plasma membrane expression to wild type receptor [748]. Subtypes of the ET<sub>B</sub> receptor have been proposed, although gene disruption studies in mice suggest that only a single gene product exists [1350].

**Further reading on Endothelin receptors**

Clozel M et al. (2013) Endothelin receptor antagonists. *Handb Exp Pharmacol* **218**: 199-227 [PMID:24092342]

Davenport AP. (2002) International Union of Pharmacology. XXIX. Update on endothelin receptor nomenclature. *Pharmacol. Rev.* **54**: 219-26 [PMID:12037137]

Davenport AP et al. (2016) Endothelin. *Pharmacol. Rev.* **68**: 357-418 [PMID:26956245]

Maguire JI et al. (2014) Endothelin®25 - new agonists, antagonists, inhibitors and emerging research frontiers: IUPHAR Review 12. *Br. J. Pharmacol.* **171**: 5555-72 [PMID:25131455]

---

**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** [1607]) was identified following observations of estrogen-evoked cyclic AMP signalling in breast cancer cells [65], which mirrored the differential expression of an orphan 7-transmembrane receptor GPR30 [276]. There are observations of both cell-surface and intracellular expression of the GPER receptor [1647, 1953].

**Nomenclature**

| Nomenclature | GPER |
|--------------|------|
| HGNC, UniProt | GPER1, Q99527 |

**Agonists**

| raloxifene [1570] |

**Selective agonists**

| G1 [179] |

**Selective antagonists**

| G36 (pIC<sub>50</sub> 6.8–6.9) [438], G15 (pIC<sub>50</sub> 6.7) [437] |

**Labelled ligands**

| [3H]17β-estradiol (Agonist) [1953] |

---

**Searchable database**: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide**: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Comments: Antagonists at the nuclear estrogen receptor, such as fulvestrant, tamoxifen [540] and raloxifene [1570], as well as the flavonoid ‘phytoestrogens’ genistein and quercetin [1241], are agonists at GPER receptors. A complete review of GPER pharmacology has been recently published [1607].

Further reading on G protein-coupled estrogen receptor
Prossnitz ER et al. (2015) International Union of Basic and Clinical Pharmacology. XCVII. G Protein-Coupled Estrogen Receptor and Its Pharmacologic Modulators. Pharmacol. Rev. 67: 505-40 [PMID:26023144]

Prossnitz ER et al. (2015) What have we learned about GPER function in physiology and disease from knockout mice? J. Steroid Biochem. Mol. Biol. 153: 114-26 [PMID:26189910]

Formylpeptide receptors
G protein-coupled receptors → Formylpeptide receptors

Overview: The formylpeptide receptors (nomenclature agreed by the NC-IUPHAR Subcommittee on the formylpeptide receptor family [2180]) 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 β42, serum amyloid A and spinorphin, derived from β-haemoglobin (HBB, P68871).

| Nomenclature | FPR1 | FPR2/ALX | FPR3 |
|--------------|------|----------|------|
| HGNC, UniProt | FPR1, P21462 | FPR2, P25090 | FPR3, P25089 |

Potency order of endogenous ligands
fMet-Leu-Phe > cathepsin G (CTSG, P08311) > annexin I (ANXA1, P04083) [1118, 1895]

Endogenous agonists

Endogenous antagonists
spinorphin (pIC50 4.3) [1165, 1404]

Antagonists
t-Boc-FLFLF (pKᵢ 6–6.5) [2095]

Selective antagonists
cyclosporin H (pKᵢ 6.1–7.1) [2095, 2167]

Labelled ligands
[3H]fMet-Leu-Phe (Agonist) [1036]

Comments: A FITC-conjugated fMLP analogue has been used for binding to the mouse recombinant receptor [758].

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Further reading on Formylpeptide receptors

Dorward DA et al. (2015) The Role of Formylated Peptides and Formyl Peptide Receptor 1 in Governing Neutrophil Function during Acute Inflammation. *Am. J. Pathol.* **185**:1172-1184 [PMID:25791526]

Dufton N et al. (2010) Therapeutic anti-inflammatory potential of formyl-peptide receptor agonists. *Pharmacol. Ther.* **127**:175-88 [PMID:20546777]

Liu M et al. (2012) G protein-coupled receptor FPR1 as a pharmacologic target in inflammation and human glioblastoma. *Int. Immunopharmacol.* **14**:283-8 [PMID:22863814]

Rabiet MJ et al. (2011) N-formyl peptide receptor 3 (FPR3) departs from the homologous FPR2/ALX receptor with regard to the major processes governing chemoattractant receptor regulation, expression at the cell surface, and phosphorylation. *J. Biol. Chem.* **286**:26718-31 [PMID:21543323]

Yazid S et al. (2012) Anti-inflammatory drugs, eicosanoids and the annexin A1/FPR2 anti-inflammatory system. *Prostaglandins Other Lipid Mediat.* **98**:94-100 [PMID:222173264]

Ye RD et al. (2009) International Union of Basic and Clinical Pharmacology. LXXIII. Nomenclature for the formyl peptide receptor (FPR) family. *Pharmacol. Rev.* **61**:119-61 [PMID:19498085]

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 [414, 1876]) 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) and C22:6,n-3 (docosahexaenoic acid)) activate FFA1 [223, 872, 1043] and FFA4 receptors [795, 852, 1494], while short chain fatty acids (C2 (acetic acid), C3 (propanoic acid), C4 (butyric acid) and C5 (pentanoic acid)) activate FFA2 [231, 1117, 1465] and FFA3 [231, 1117] receptors. The crystal structure for agonist bound FFA1 has been described [1862].

| Nomenclature       | FFA1 receptor | FFA2 receptor |
|--------------------|---------------|---------------|
| HGNC, UniProt      | FFAR1, O14842 | FFAR2, O15552 |

**Endogenous agonists**
docosahexaenoic acid [223, 872], α-linolenic acid [223, 872, 1043], oleic acid [223, 872, 1043], myristic acid [223, 872, 1043]

**Selective agonists**
AMG-837 [1176], compound 4 [347], TUG-770 [346], TUG-905 [345], GW9508 (Partial agonist) [222], fasiglifam [935, 1434, 1862, 1985]

**Selective antagonists**
GW1100 (pIC<sub>50</sub> 6) [222, 1875]

**Comments**
Antagonist GW1100 is also an oxytocin receptor antagonist [222], Fasiglifam, TUG-770 and GW9508 are approximately 100 fold selective for FFA1 over FFA4 [222, 346, 1434]. 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 [1176, 2153].

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Nomenclature | FFA3 receptor | FFA4 receptor | GPR42
--- | --- | --- | ---
HGNC, UniProt | FFA3, O14843 | FFA4, Q5NUL3 | GPR42, O15529
Endogenous agonists | propanoic acid [231, 1117, 1741, 2152], butyric acid [231, 1117, 1741, 2152], 1-methylcyclopropanecarboxylic acid [1741] | α-linolenic acid [1794], myristic acid [2084], α-linolenic acid [1932] – Rat, oleic acid [2084] | –
Agonists | acetic acid [231, 1117, 1741, 2152] | – | –
Selective agonists | – | compound A [1493], TUG-891 [1794], NCG21 [1902] | –
Comments | Beta-hydroxybutyrate has been reported to antagonise FFA3 responses to short chain fatty acids [997]. 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 (i.e. allosteric modulators), have recently been described [180, 839, 1226]. | A wide range of both saturated and unsaturated fatty acids containing from 6 to 22 carbons have been shown to act as agonists at FFA4 [348] with a small subset listed above. Compound A [PMID 24997608] exhibits more than 1000 fold selectivity [1493], and TUG-891 50-1000 fold selectivity for FFA4 over FFA1 [1794], dependent on the assay. NGC21 exhibits approximately 15 fold selectivity for FFA4 over FFA1 [1894]. | –

**Comments:** Short (361 amino acids) and long (377 amino acids) splice variants of human FFA4 have been reported [1372], which differ by a 16 amino acid insertion in intracellular loop 3, and exhibit differences in intracellular signalling properties in recombinant systems [2084]. The long FFA4 splice variant has not been identified in other primates or rodents to date [795, 1372]. GPR42 was originally described as a pseudogene within the family (ENSFM00250000002583), but the discovery of several polymorphisms suggests that some versions of GPR42 may be functional [1167]. GPR84 is a structurally-unrelated G protein-coupled receptor which has been found to respond to medium chain fatty acids [2067].

### Further reading on Free fatty acid receptors

Bolognini D et al. (2016) The Pharmacology and Function of Receptors for Short-Chain Fatty Acids. *Mol. Pharmacol.* **89**: 388-98 [PMID:26719580]

Mancini AD et al. (2013) The fatty acid receptor FFA1/GPR40 a decade later: how much do we know? *Trends Endocrinol. Metab.* **24**: 398-407 [PMID:23631851]

Moniri NH. (2016) Free-fatty acid receptor-4 (GPR120): Cellular and molecular function and its role in metabolic disorders. *Biochem. Pharmacol.* **110-111**: 1-15 [PMID:26827942]

Stoddart LA et al. (2008) International Union of Pharmacology. LXXI. Free fatty acid receptors FFA1, -2, and -3: pharmacology and pathophysiological functions. *Pharmacol. Rev.* **60**: 405-17 [PMID:19047536]

Talukdar S et al. (2011) Targeting GPR120 and other fatty acid-sensing GPCRs ameliorates insulin resistance and inflammatory diseases. *Trends Pharmacol. Sci.* **32**: 543-50 [PMID:21663979]

Watterson KR et al. (2014) Treatment of type 2 diabetes by free Fatty Acid receptor agonists. *Front Endocrinol (Lausanne)* **5**: 137 [PMID:25221541]

### GABA<sub>B</sub> receptors

G protein-coupled receptors → GABA<sub>B</sub> receptors

**Overview:** Functional GABA<sub>B</sub> receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on GABA<sub>B</sub> receptors [199, 1579]) are formed from the heterodimerization of two similar 7TM subunits termed GABA<sub>B1</sub> and GABA<sub>B2</sub> [199, 506, 1578, 1579, 2002]. GABA<sub>B</sub> receptors are widespread in the CNS and regulate both pre- and postsynaptic activity. The GABA<sub>B1</sub> 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. Co-expression of GABA<sub>B1</sub> and GABA<sub>B2</sub> subunits allows transport of GABA<sub>B1</sub> to the cell surface and generates a functional receptor that can couple to signal transduction pathways such as high-voltage-activated Ca<sup>2+</sup> channels (Ca<sub>v</sub>2.1, Ca<sub>v</sub>2.2), or inwardly rectifying potassium channels (Kir3) [147, 199, 200]. The GABA<sub>B1</sub> subunit harbours the GABA (orthosteric)-binding site within an extracellular domain (ECD)

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
venus flytrap module (VTM), whereas the GABAB2 subunit mediates G protein-coupled signalling [199, 622, 624, 1578]. The two subunits interact by direct allosteric coupling [1367], such that GABAB2 increases the affinity of GABAB1 for agonists and reciprocally GABAB1 facilitates the coupling of GABAB2 to G proteins [622, 1060, 1578]. GABAB1 and GABAB2 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 [147, 250, 1578]. Recent evidence indicates that higher order assemblies of GABAB receptor comprising dimers of heterodimers occur in recombinant expression systems and in vivo, and that such complexes exhibit negative functional cooperativity between heterodimers [373, 1577]. Adding further complexity, KCTD (potassium channel tetramerization proteins) 8, 12, 12b and 16 associate as tetramers with the carboxy terminus of the GABAB2 subunit to impart altered signalling kinetics and agonist potency to the receptor complex [108, 1751, 1990] and are reviewed by [1580]. The molecular complexity of GABAB receptors is further increased through association with trafficking and effector proteins [Schwenk et al., 2016, Nature Neuroscience 19(2): 233-42] and reviewed by [1576]. Four isoforms of the human GABAB1 subunit have been cloned. The predominant GABAB1a and GABAB1b 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. GABAB1a-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 GABAB1b-containing receptors occur within dendritic spines and mediate slow postsynaptic inhibition [1613, 2035]. Only the 1a and 1b variants are identified as components of native receptors [199]. Additional GABAB1 subunit isoforms have been described in rodents and humans [1130] and reviewed by [147].

| Subunits | Nomenclature | KGNC, UniProt |
|----------|--------------|---------------|
| GABAB1   | GABBR1, Q9UBS5 |                |
| GABAB2   | 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 [199, 580, 581]. Radioligand Kp values relate to binding to rat brain membranes. CGP 71872 is a photoaffinity ligand for the GABAB1 subunit [128]. CGP27492 (3-APPA), CGP35024 (3-APMPA) and CGP44532 act as antagonists at human GABAB1 receptors, with potencies in the low micromolar range [580]. In addition to the ligands listed in the table, Ca2+ binds to the VTM of the GABAB1 subunit to act as a positive allosteric modulator of GABA [594]. Synthetic positive allosteric modulators with low, or no, intrinsic activity include CGP7930, GS39783, BHE-177 [2040] and (+)-BHFF [9, 147, 154, 580]. The site of action of CGP7930 and GS39783 appears to be on the heptahelical domain of the GABAB2 subunit [483, 1578]. In the presence of CGP7930 or GS39783, CGP 35348 and 2-hydroxy-saclofen behave as partial agonists [580]. A negative allosteric modulator of GABAB activity has been reported [318]. Knock-out of the GABAB1 subunit in C57B mice causes the development of severe tonic-clonic convulsions that prove fatal within a month of birth, whereas GABAB1-/- 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 [510, 2008]. A similar phenotype has been found for GABAB2-/- BALB/c mice [613].
Further reading on GABAB receptors

Bowery NG et al. (2002) International Union of Pharmacology. XXXIII. Mammalian gamma-aminobutyric acid(B) receptors: structure and function. Pharmaco1 Rev. 54: 247-264 [PMID:12037141]
Frostl W. (2011) An historical perspective on GABAergic drugs. Future Med Chem 3: 163-75
Gassmann M et al. (2012) Regulation of neuronal GABA(B) receptor functions by subunit composition. Nat. Rev. Neurosci. 13: 380-94 [PMID:22595784]
Pin JP et al. (2016) Organization and functions of mGlu and GABAB receptor complexes. Nature 540: 60-68 [PMID:27905440]

Galanin receptors

G protein-coupled receptors → Galanin receptors

Overview: Galanin receptors (provisional nomenclature as recommended by NC-IUPHAR [557]) 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 [525]; 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 [143] and porcine galanin-5–29 [1809]) and N-terminally extended forms (e.g. N-terminally seven and nine residue elongated forms of porcine galanin [144, 1809]) have been reported.

| Nomenclature | GAL1 receptor | GAL2 receptor | GAL3 receptor |
|--------------|---------------|---------------|---------------|
| HGNC, UniProt | GALR1, P47211 | GALR2, O43603 | GALR3, O60755 |
| Potency order of endogenous ligands | galanin (GAL, P22466) > galanin-like peptide (GALP, Q9UBC7) [1500] | galanin-like peptide (GALP, Q9UBC7) ≥ galanin (GAL, P22466) [1500] | galanin-like peptide (GALP, Q9UBC7) > galanin (GAL, P22466) [1095] |
| Agonists | – | galanin(2-29) (rat/mouse) [1526, 2069, 2070, 2071] – Rat | – |
| Selective agonists | – | [D-Trp²]galanin-(1-29) [1834] – Rat | – |
| Selective antagonists | 2,3-dihydro-1,4-dithiin-1,1,4,4-tetroxide (pIC₅₀ 5.6) [1758] | M871 (pKᵢ 7.9) [1848] | SNAP 398299 (pKᵢ 8.3) [1033, 1034, 1906], SNAP 37889 (pKᵢ 7.8–7.8) [1033, 1034, 1906] |
| Selective allosteric modulators | – | CYM2503 (Positive) (pEC₅₀ 9.2) [1213] – Rat | – |
| Labelled ligands | [¹²⁵I][Tyr²⁶]galanin (human) (Agonist) [552], [¹²⁵I][Tyr²⁶]galanin (human) (Agonist) [552] | [¹²⁵I][Tyr²⁶]galanin (human) (Agonist) [2070] – Rat | [¹²⁵I][Tyr²⁶]galanin (pig) (Agonist) [191, 1835] |
| Comments | – | The CYM2503 PAM potentiates the anticonvulsant activity of endogenous galanin in mouse seizure models [1213]. | – |
**Comments:** galanin-(1-11) is a high-affinity agonist at GAL1/GAL2 (pK<sub>i</sub> 9), and galanin-(2-11) is selective for GAL2 and GAL3 compared with GAL1 [1212]. [125I]Tyr<sup>26</sup>galanin binds to all three subtypes with K<sub>i</sub> values generally reported to range from 0.05 to 1 nM, depending on the assay conditions used [552, 1821, 1834, 1835, 2070]. Porcine galanin-(3-29) does not bind to cloned GAL1, GAL2 or GAL3 receptors, but a receptor that is functionally activated by porcine galanin-(3–29) has been reported in pituitary and gastric smooth muscle cells [691, 2142]. 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 [2000], spinal cord [2114], locus coeruleus, hippocampus [106] and hypothalamus [107, 1142], but exhibit agonist activity at some peripheral sites [107, 691]. The chimeric peptides M15, M32, M35, M40 and C7 are agonists at GAL1 receptors expressed endogenously in Bowes human melanoma cells [1500], and at heterologously expressed recombinant GAL1, GAL2 and GAL3 receptors [552, 1834, 1835]. Recent studies have described the synthesis of a series of novel, systemically-active, galanin analogues, with modest preferential binding at the GAL2 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 [1698].

**Further reading on Galanin receptors**

Foord SM et al. (2005) International Union of Pharmacology. XLVI. G protein-coupled receptor list. Pharmacol Rev 57: 279-288 [PMID:15914470]

Lang R et al. (2015) Physiology, signaling, and pharmacology of galanin peptides and receptors: three decades of emerging diversity. Pharmacol. Rev. 67: 118-75 [PMID:25428932]

Lawrence C et al. (2011) Galanin-like peptide (GALP) is a hypothalamic regulator of energy homeostasis and reproduction. Front Neuroendocrinol 32: 1-9 [PMID:20558195]

Webling KE et al. (2012) Galanin receptors and ligands. Front Endocrinol (Lausanne) 3: 146 [PMID:23233848]

---

**Ghrelin receptor**

**G protein-coupled receptors → Ghrelin receptor**

**Overview:** The ghrelin receptor (nomenclature as agreed by the NC-IUPHAR Subcommittee for the Ghrelin receptor [415]) 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 [1285]. Alternative splicing results in the formation of a second peptide, [des-Gln<sup>14</sup>]ghrelin (GHRL, Q9UBU3) with equipotent biological activity [822]. A unique post-translational modification (octanoylation of Ser<sup>3</sup>, catalysed by ghrelin O-acyltransferase (MBOAT4, Q96T53) [2170] 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 [1029]. Structure activity studies showed the first five N-terminal amino acids to be the minimum required for binding [122], 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 [814]. In cell systems, the ghrelin receptor is constitutively active [815], 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 [1527].

**Nomenclature**

| HGNC, UniProt | ghrelin receptor | GHSR, Q92847 |
|-------------|-----------------|--------------|

**Potency order of endogenous ligands**

| ghrelin (GHRL, Q9UBU3) = [des-Gln<sup>14</sup>]ghrelin (GHRL, Q9UBU3) [121, 1285] |

**Selective antagonists**

| GSK1614343 (pIC<sub>50</sub> 8.4) [1699], GSK1614343 (pK<sub>B</sub> 8) [1556] – Rat |

**Labelled ligands**

| [125I][His<sup>9</sup>]ghrelin (human) (Agonist) [956], [125I][Tyr<sup>4</sup>]ghrelin (human) (Agonist) [1394] |

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Comments: [des-octanoyl]ghrelin (GHRL, Q9UBU3) has been shown to bind (as [125I]Tyr4-des-octanoyl-ghrelin) and have effects in the cardiovascular system [121], 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-Trp7, 9, Leu11]substance P, pD2 8.3; [812]). Ulimorelin, described as a ghrelin receptor agonist (pKi 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 discriminating the orexigenic and gastrointestinal actions of ghrelin (GHRL, Q9UBU3) from the release of growth hormone [567]. A number of selective antagonists have been reported, including peptidomimetic [1393] and non-peptide small molecules including GSK1614343 [1556, 1699].

Further reading on Ghrelin receptor

Andrews ZB. (2011) The extra-hypothalamic actions of ghrelin on neuronal function. Trends Neurosci. 34: 31-40 [PMID:21035199]

Angelidis G et al. (2010) Current and potential roles of ghrelin in clinical practice. J. Endocrinol. Invest. 33: 823-38 [PMID:21293171]

Briggs DI et al. (2011) Metabolic status regulates ghrelin function on energy homeostasis. Neuroendocrinology 93: 48-57 [PMID:21124019]

Callaghan B et al. (2014) Novel and conventional receptors for ghrelin, desacyl-ghrelin, and pharmacologically related compounds. Pharmacol. Rev. 66: 984-1001 [PMID:25107984]

Davenport AP et al. (2005) International Union of Pharmacology. LVI. Ghrelin receptor nomenclature, distribution, and function. Pharmacol. Rev. 57: 541-6 [PMID:16382107]

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 [1296]) 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 [866].

| Nomenclature       | GHRH receptor | GIP receptor | GLP-1 receptor |
|--------------------|---------------|--------------|---------------|
| HGNC, UniProt      | GHRHR, Q02643 | GIPR, P48546 | GLP1R, P43220 |
| Endogenous agonists| –             | gastric inhibitory polypeptide (GIP, P09681) [2042] | glucagon-like peptide 1-(7-36) amide (GCG, P01275) [927], glucagon-like peptide 1-(7-37) (GCG, P01275) [449] |
| Agonists           | JJ-38 [265], sermorelin | – | liraglutide [1020], lixisenatide [2097], WB4-24 [528] |
| Selective agonists | BIM28011 [393], tesamorelin | – | exendin-4 [1346], exendin-4 [927], exendin-3 (P20394) [1635] |
| Selective antagonists | JV-1-36 (pKᵢ 10.1–10.4) [1733, 2026, 2027] – Rat, JV-1-38 (pKᵢ 10.1) [1733, 2026, 2027] – Rat | [Pro³]GIP [615] – Mouse | exendin-(9-39) (pKᵢ 8.1) [927], GLP-1-(9-36) (pIC₅₀ 6.9) [1368] – Rat, T-0632 (pIC₅₀ 4.7) [1961] |
| Labelled ligands   | [125I]GHRH (human) (Agonist) [196] – Rat | [125I]GIP (human) (Agonist) [593] – Rat | [125I]GLP-1-(7-36)-amide (Agonist) [927], [125I]exendin-(9-39) (Antagonist) (pKᵢ 8.3) [927], [125I]GLP-1-(7-37) (human) (Agonist) |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature | GLP-2 receptor | glucagon receptor | secretin receptor  
--- | --- | --- | ---  
HGNC, UniProt | GLP2R, P095838 | GCR, P47871 | SCTR, P47872  
Endogenous agonists | glucagon-like peptide 2 (GCG, P01275) [1958] | glucagon (GCG, P01275) [1587] | secretin (SCT, P09683) [343]  
Agonists | teduglutide [1310] | |  
Selective antagonists | – | L-168,049 (pIC50 8.4) [282], adomeglivant (pK8 8.2) [963, 967], des-His1-[Glu9]glucagon-NH2 (pA2 7.2) [2004, 2005] – Rat, NNC 92-1687 (pK5 1234), BAY27-9955 (pK5 1566) | [(CH2NH)4,5]secretin (pKi 5.3) [704]  
Labelled ligands | – | [125I]glucagon (human, mouse, rat) (Agonist) | [125I][Tyr10]secretin-27 (rat) (Agonist) [2001] – Rat

**Comments:** The glucagon receptor has been reported to interact with receptor activity modifying proteins (RAMPs), specifically RAMP2, in heterologous expression systems [349], although the physiological significance of this has yet to be established.

**Further reading on Glucagon receptor family**

Ahrén B. (2015) Glucagon–Early breakthroughs and recent discoveries. *Peptides* **67**: 74-81 [PMID:25814364]

Campbell JE et al. (2013) Pharmacology, physiology, and mechanisms of incretin hormone action. *Cell Metab.* **17**: 819-37 [PMID:23684623]

Donnelly D. (2012) The structure and function of the glucagon-like peptide-1 receptor and its ligands. *Br. J. Pharmacol.* **166**: 27-41 [PMID:21950636]

Kazda CM et al. (2016) Evaluation of Efficacy and Safety of the Glucagon Receptor Antagonist LY2409021 in Patients With Type 2 Diabetes: 12- and 24-Week Phase 2 Studies. *Diabetes Care* **39**: 1241-9 [PMID:26681715]

Mayo KE et al. (2003) International Union of Pharmacology. XXXV. The glucagon receptor family. *Pharmacol. Rev.* **55**: 167-94 [PMID:12615957]

Trujillo JM et al. (2014) GLP-1 receptor agonists for type 2 diabetes mellitus: recent developments and emerging agents. *Pharmacotherapy* **34**: 1174-86 [PMID:25382096]

**Glycoprotein hormone receptors**

**G protein-coupled receptors → Glycoprotein hormone receptors**

**Overview:** Glycoprotein hormone receptors (provisional nomenclature [557]) are activated by a non-covalent heterodimeric glycoprotein made up of a common α chain (glycoprotein hormone common alpha subunit (CGA, P01215)) and a unique β chain that confers the biological specificity to FSH (CGA FSHB, P01215 P01225), LH (CGA LHB, P01215 P01229), hCG (CGA CGB3, P01215 P01233) or TSH (CGA TSHB, P01215 P01222). There is binding cross-reactivity across the endogenous agonists for each of the glycoprotein hormone receptors. The deglycosylated hormones appear to exhibit reduced efficacy at these receptors [1701].

[Searchable database: http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

[Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Gonadotrophin-releasing hormone receptors

**Overview:** GnRH₁ and GnRH₂ receptors (provisional nomenclature [557]), also called Type I and Type II GnRH receptor, respectively [1342]) have been cloned from numerous species, most of which express two or three types of GnRH receptor [1341, 1342, 1810]. 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 [1341, 1342, 1810] 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 [1341, 1342, 1810]. GnRH₁ receptors are expressed by pituitary gonadotrophs, where they mediate the effects of GnRH on gonadotropin hormone synthesis and secretion that underpin central control of mammalian reproduction. GnRH analogues are used in assisted reproduction and to treat steroid hormone-dependent conditions [981]. Notably, agonists cause desensitization of GnRH-stimulated gonadotropin secretion and the consequent reduction in circulating sex steroids is exploited to treat hormone-dependent cancers of the breast, ovary and prostate [981]. GnRH₁ receptors 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 not by humans [1377]. Phylogenetic classifications divide GnRH receptors into three [1342] or five groups [2117] and highlight examples of gene loss through evolution, with humans retaining only one ancient gene.

Further reading on Glycoprotein hormone receptors

Jiang X *et al.* (2012) Structure of follicle-stimulating hormone in complex with the entire ectodomain of its receptor. *Proc. Natl. Acad. Sci. U.S.A.* **109:** 12491-6 [PMID:22802634]

Kleinau G *et al.* TSH receptor mutations and disease. Accessed on 2017-02-23. Thyroid Disease Manager.

Tao YX *et al.* (2009) Follicle stimulating hormone receptor mutations and reproductive disorders. *Prog Mol Biol Transl Sci* **89:** 115-31 [PMID:20374735]

Troppmann B *et al.* (2013) Structural and functional plasticity of the luteinizing hormone/choriogonadotrophin receptor. *Hum. Reprod. Update* **19:** 583-602 [PMID:23686864]
Nomenclature

HGNC,

| Nomenclature       | GnRH₁ receptor | GnRH₂ receptor |
|--------------------|----------------|----------------|
| HGNC, UniProt      | GNRH₁, P30968  | GNRH₂, Q96P88  |

Potency order of endogenous ligands

- GnRH I (GNRH₁, P01148) > GnRH II (GNRH₂, O43555) [1342]
- GnRH II (GNRH₂, O43555) > GnRH I (GNRH₁, P01148) (Monkey) [1340]

Endogenous agonists

- GnRH I (GNRH₁, P01148) [1214], GnRH II (GNRH₂, O43555) [550, 1214, 1869]
- GnRH II (GNRH₂, O43555) [1340] – Monkey, GnRH I (GNRH₁, P01148) [1340] – Monkey

Selective agonists

- Buserelin [1432], buserelin [1431], triptorelin [118], leuprolide [1881], goserelin, histrelin, nafarelin

Selective antagonists

- Cetrorelix (pKᵢ 9.3–10) [119, 120, 1881], abarelix (pKᵢ 9.1–9.5) [1881], degarelix (pKᵢ 8.8) [2017], ganirelix

Labelled ligands

- [¹²⁵I]Cetrorelix (Antagonist) (pKᵢ 9.7) [807], [¹²⁵I]Triptorelin (Agonist) [435] – Rat, [¹²⁵I]Buserelin (Agonist) [1076] – Rat, [¹²⁵I]GnRH I (human, mouse, rat) (Agonist)

Comments:

- GnRH₁ and GnRH₂ receptors couple primarily to Gq/11 [686] but coupling to Gs and Gi is evident in some systems [1056, 1076]. GnRH₂ receptors may also mediate (heterotrimetric) G protein-independent signalling to protein kinases [289]. 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 [324, 741, 1174, 1732]. In some human cancer cell models GnRH II (GNRH₂, O43555) is more potent than GnRH I (GNRH₁, P01148), implying mediation by GnRH₂ receptors [690], but GnRH₂ receptors are not expressed by humans because the human GNRHR2 gene contains a frame shift and internal stop codon [1377]. The possibility remains that this gene generates GnRH receptor-related proteins (other than the full-length receptor) that mediate responses to GnRH II (GNRH₂, O43555) (see [1436]). Alternatively, evidence for multiple active GnRH receptor conformations [289, 290, 541, 1293, 1342] raises the possibility that GnRH₁ receptor-mediated proliferation inhibition in hormone-dependent cancer cells is dependent upon different conformations than effects on Gq/11 in pituitary cells [290, 1293]. Loss-of-function mutations in the GnRH₁ receptor and deficiency of GnRH I (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 [1124]. Human GnRH₁ receptors are poorly expressed at the cell surface because of failure to meet structural quality control criteria for endoplasmic reticulum exit [542, 1124], and this increase susceptibility to point mutations that further impair trafficking [542, 1124]. GnRH receptor signalling may require receptor oligomerisation [376, 1054].

Further reading on Gonadotrophin-releasing hormone receptors

Limonta P et al. (2012) GnRH receptors in cancer: from cell biology to novel targeted therapeutic strategies. Endocr. Rev. 33: 784-811 [PMID:22778172]

McArdle CA and Roberson MS. (2015) Gonadotropes and gonadotropin-releasing hormone signaling. In Knobil and Neill’s Physiology of Reproduction (4th edition). Edited by Plant TM and Zeleznik AJ.: Elsevier Inc.: [ISBN: 9780123971753]

Millar RP et al. (2004) Gonadotropin-releasing hormone receptors. Endocr Rev 25: 235-275 [PMID:15082521]

Tao YX et al. (2014) Chaperoning G protein-coupled receptors: from cell biology to therapeutics. Endocr. Rev. 35: 602-47 [PMID:24661201]
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 [1564]. Although there are multiple reports to indicate that GPR18, GPR55 and GPR119 can be activated in vitro by N-arachidonoylglycine, lysophosphatidylinositol and N-oleylethanolamide, 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 | GPR18 | GPR55 | GPR119 |
|--------------|-------|-------|--------|
| HGNC, UniProt | GPR18, Q14330 | GPR55, Q9Y2T6 | GPR119, Q8TDVS |
| Potency order of endogenous ligands | – | – | N-oleylethanolamide, N-palmitoylethanolamine > SEA (anandamide is ineffective) [1520] |
| Endogenous agonists | N-arachidonoylglycine [1026] | lysophosphatidylinositol [773, 1502, 1854], 2-arachidonoylglycerophosphoinositol [1504] | N-oleylethanolamide [354, 1520, 1854], N-palmitoylethanolamine, SEA |
| Selective agonists | – | AM251 [773, 948, 1695] | AS1269574 [2190], PSN632408 [1520], PSN375963 [1520] |
| Selective antagonists | – | CID16020046 (p<sub>A</sub> 2 7.3) [949] | – |
| Comments | The pairing of N-arachidonoylglycine with GPR18 was not replicated in two studies based on arrestin assays [1854, 2182]. See [414] for discussion. | See reviews [414] and [1800]. | In addition to those shown above, further small molecule agonists have been reported [722]. |

**Comments:** GPR18 failed to respond to a variety of lipid-derived agents in an in vitro screen [2182], but has been reported to be activated by Δ⁷-tetrahydrocannabinol [1308]. GPR55 responds to AM251 and rimonabant at micromolar concentrations, compared to their nanomolar affinity as CB₁ receptor antagonists/inverse agonists [1564]. It has been reported that lysophosphatidylinositol acts at other sites in addition to GPR55 [2164]. N-Arachidonoyleritine has been suggested to act as a low efficacy agonist/antagonist at GPR18 in vitro [1306]. It has also been suggested that oleoyl-lysophosphatidylcholine acts, at least in part, through GPR119 [1466]. Although PSN375963 and PSN632408 produce GPR119-dependent responses in heterologous expression systems, comparison with N-oleylethanolamine-mediated responses suggests additional mechanisms of action [1466].

**Further reading on GPR18, GPR55 and GPR119**

Davenport AP et al. (2013) International Union of Basic and Clinical Pharmacology. LXXXVIII. G protein-coupled receptor list: recommendations for new pairings with cognate ligands. Pharmacol. Rev. 65: 967-86 [PMID:23686380]

Hassing HA et al. (2016) Biased signaling of lipids and allosteric actions of synthetic molecules for GPR119. Biochem Pharmacol 119: 66-75 [PMID:27569424]

Liu B et al. (2015) GPR55: from orphan to metabolic regulator? Pharmacol Ther 145: 35-42 [PMID:24972076]

Pertwee RG et al. (2010) International Union of Basic and Clinical Pharmacology. LXXIX. Cannabinoid receptors and their ligands: beyond CB₁ and CB₂. Pharmacol. Rev. 62: 588-631 [PMID:21079038]

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Histamine receptors
G protein-coupled receptors → Histamine receptors

Overview: Histamine receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Histamine Receptors [790, 1528]) are activated by the endogenous ligand histamine. Marked species differences exist between histamine receptor orthologues [790]. The human and rat H3 receptor genes are subject to significant splice variance [91]. The potency order of histamine at histamine receptor subtypes is $H_3 = H_4 > H_2 > H_1$ [1528]. Some agonists at the human H3 receptor display significant ligand bias [1659]. Antagonists of all 4 histamine receptors have clinical uses: H1 antagonists for allergies (e.g. cetirizine), H2 antagonists for acid-reflux diseases (e.g. ranitidine), H3 antagonists for narcolepsy (e.g. pitolisant/WAKIX; Registered) and H4 antagonists for atopic dermatitis (e.g. ZPL-3893787; Phase IIa) [1528].

| Nomenclature | H1 receptor | H2 receptor | H3 receptor | H4 receptor |
|--------------|-------------|-------------|-------------|-------------|
| HGNC, UniProt | HRH1, P35367 | HRH2, P25021 | HRH3, Q9YSN1 | HRH4, Q9H3N8 |
| Selective agonists | methylhistaprodifen [1766], histaprodifen [1173] | amthamine [1048] | immethridine [1011], methimepip [1010], MK-0249 (Inverse agonist) [1413] | clobenpropit (Partial agonist) [517, 1173, 1188, 1189, 1389], 4-methylhistamine [617, 1173], ST-1006 [1528], VUF 8430 [1172] |
| Antagonists | cyproheptadine (pKi 10.2) [1352], promethazine (pKi 9.6) [636], pyrilamine (Inverse agonist) (pKi 8.7–9) [188, 1634], cetirizine (Inverse agonist) (pKi 8.2) [1352], diphenhydramine (pKi 7.9) [188] | iodophenpropit (pKi 8.2–8.7) [2112, 2139] | thioperamide (Selective for $H_3/H_4$ compared to $H_1$ and $H_3$) (pKi 7.1–7.7) [368, 516, 517, 1211, 2112, 2139] | thioperamide (Selective for $H_3/H_4$ compared to $H_1$ and $H_3$) (pKi 6.3–7.6) [516, 517, 1188, 1189, 1389, 2226] |
| Sub/family-selective agonists | – | – | – | – |
| Selective agonists | clemastine (pKi 10.3) [71], desloratadine (pKi 9) [1156], tripolidine (pKi 8.5–9) [188, 1352], azelastine (pKi 8.9) [1606], astemizole (pKi 8.5) [1547], tiotidine (pKi 7.5) [149] – Rat, ranitidine (pKi 7.1) [1152], cimetidine (pKi 6.8) [274] | pitolisant (pKi 8.1–8.6) [1528, 2254], A331440 (pKi 8.5) [723], conessine (pKi 8.3) [1528], MK-0249 (pKi 8.2) [1528], ciproxifan (pKi 6.7–7.3) [368, 516, 517, 1170, 1528, 2139] | ZPL-3893787 (pKi 8.3) [1528], INCB-38579 (pKi 8.3) [1528], JNJ 7777120 (pKi 7.8–8.3) [1173, 1839, 1959], JNJ-39758979 (pKi 7.9) [1528, 1727] | – |
| Labelled ligands | [3H]pyrilamine (Antagonist, Inverse agonist) (pKD 8.4–9.1) [422, 1352, 1746, 1766], [11C]doxepin (Antagonist) (pKi 9) [869], [11C]pyrilamine (Antagonist, Inverse agonist) | [125I]iodoaminopotentidine (Antagonist) (pKD 8.7) [1082] – Rat, [3H]tiotidine (Antagonist) (pKD 7.7–8.7) [1363] | [123I]iodoxyproxyfan (Antagonist) (pKD 10.2) [1170], [125I]iodophenpropit (Antagonist) (pKD 9.2) [891] – Rat, [3H](R)-α-methylhistamine (Agonist) [1188], N-[3H](S)-α-methylhistamine (Agonist) [317] – Mouse | [3H]JNJ 7777120 (Antagonist) (pKD 8.4) [1599] |
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 [1188, 1419, 1455, 1489, 2226]. Moreover, 4-methylhistamine is identified as a high affinity, full agonist for the human H₄ receptor [1173]. [³H]histamine has been used to label the H₄ receptor in heterologous expression systems.

Further reading on Histamine receptors

Gbahou F et al. (2012) The histamine autoreceptor is a short isoform of the H₃ receptor. Br. J. Pharmacol. 166: 1860-71 [PMID:22356432]
Nieto-Alamilla G et al. (2016) The Histamine H3 Receptor: Structure, Pharmacology, and Function. Mol. Pharmacol. 90: 649-673 [PMID:27563055]
Panula P et al. (2015) International Union of Basic and Clinical Pharmacology. XCVIII. Histamine Receptors. Pharmacol. Rev. 67: 601-55 [PMID:26084539]
van Rijn RM et al. (2008) Cloning and characterization of dominant negative splice variants of the human histamine H4 receptor. Biochem. J. 414: 121-31 [PMID:18452403]

Hydroxycarboxylic acid receptors

G protein-coupled receptors → Hydroxycarboxylic acid receptors

Overview: The hydroxycarboxylic acid family of receptors (ENSFM005000000271913, nomenclature as agreed by the NC-IUPHAR Subcommittee on Hydroxycarboxylic acid receptors [414, 1491]) 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 [1842, 1989, 2125]. 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 [1989, 2125].

| Nomenclature | HCA₁ receptor | HCA₂ receptor | HCA₃ receptor |
|--------------|---------------|---------------|---------------|
| HGNC, UniProt | HCA₁, Q9BXC0 | HCA₂, Q8TDS4 | HCA₃, P49019 |
| Potency order of endogenous ligands | – | β-D-hydroxybutyric acid > butyric acid | – |
| Endogenous agonists | L-lactic acid [16, 266, 1190, 1854] | β-D-hydroxybutyric acid [1912], butyric acid | 3-hydroxyoctanoic acid [15] |
| Agonists | 3,5-dihydroxybenzoic acid [1187] | SCH 900271 [1522], GSK256073 [1861] | compound 6o [1820], IBC 293 [1769] |
| Selective agonists | – | MK 6892 [1788], MK 1903 [166], nicotinic acid [1842, 1989, 2125], acipimox [1842, 2125], monomethylfumarate [1933] | – |
| Labelled ligands | – | [³H]nicotinic acid (Agonist) [1842, 1989, 2125] | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
**Comments:** Further closely-related GPCRs include the 5-oxoicosanoid receptor (OXER1, Q8TDS5) and GPR31 (Q600270). Lactate activates HCA1 on adipocytes in an autocrine manner. It inhibits lipolysis and thereby promotes anabolic effects. HCA2 and HCA3 regulate adipocyte lipolysis and immune functions under conditions of increased FFA formation through lipolysis (e.g., during fasting). HCA2 agonists acting mainly through the receptor on immune cells exert antiatherogenic and anti-inflammatory effects. HCA2 is also a receptor for butyrate and mediates some of the beneficial effects of short-chain fatty acids produced by gut microbiota.

**Further reading on Hydroxycarboxylic acid receptors**

Boatman PD et al. (2008) Nicotinic acid receptor agonists. *J. Med. Chem.* 51: 7653-62 [PMID:18983141]

Graff EC et al. (2016) Anti-inflammatory effects of the hydroxycarboxylic acid receptor 2. *Metab. Clin. Exp.* 65: 102-13 [PMID:26773933]

Kamanna VS et al. (2013) Recent advances in niacin and lipid metabolism. *Curr. Opin. Lipidol.* 24: 239-45 [PMID:23619367]

Offermanns S. (2017) Hydroxy-Carboxylic Acid Receptor Actions in Metabolism. *Trends Endocrinol. Metab.* [PMID:28087125]

Offermanns S et al. (2011) International Union of Basic and Clinical Pharmacology. LXXXII: Nomenclature and Classification of Hydroxy-carboxylic Acid Receptors (GPR81, GPR109A, and GPR109B). *Pharmacol. Rev.* 63: 269-90 [PMID:21454438]

Offermanns S et al. (2015) Nutritional or pharmacological activation of HCA(2) ameliorates neuroinflammation. *Trends Mol Med* 21: 245-55 [PMID:25766751]

**Kisspeptin receptor**

G protein-coupled receptors → Kisspeptin receptor

**Overview:** The kisspeptin receptor (nomenclature as agreed by the NC-IUPHAR Subcommittee on the kisspeptin receptor [1004]), 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. Kisspeptins have roles in, for example, cancer metastasis, fertility/puberty regulation and glucose homeostasis.

| Nomenclature       | kispeptin receptor |
|--------------------|--------------------|
| HGNC, UniProt     | KISS1R, Q969F8 |
| Endogenous agonists| kisspeptin-10 (KISS1) [1041, 1501], kisspeptin-54 (KISS1, Q15726) [1041, 1501], kisspeptin-14 (KISS1, Q15726) [1041], kisspeptin-13 (KISS1, Q15726) [1041] |
| Selective agonists | 4-fluorobenzoyl-FGLRW-NH2 [1973], [dY]1KP-10 [401] – Mouse |
| Selective antagonists | peptide 234 [1674] |
| Labelled ligands   | [123I]Ile-29-kisspeptin-15 (Agonist) [1501], [125I]kisspeptin-13 (human) (Agonist) [1314], [125I]kisspeptin-10 (human) (Agonist) [1041], [125I]kisspeptin-14 (human) (Agonist) [1041] |

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Further reading on Kisspeptin receptor

Kanda S et al. (2013) Structure, synthesis, and phylogeny of kisspeptin and its receptor. *Adv. Exp. Med. Biol.* **784**: 9-26 [PMID:23550000]

Kirby HR et al. (2010) International Union of Basic and Clinical Pharmacology. LXXVII. Kisspeptin receptor nomenclature, distribution, and function. *Pharmacol. Rev.* **62**: 565-78 [PMID:21079036]

Millar RP et al. (2010) Kisspeptin antagonists: unraveling the role of kisspeptin in reproductive physiology. *Brain Res.* **1364**: 81-9 [PMID:20858467]

Oakley AE et al. (2009) Kisspeptin signaling in the brain. *Endocr. Rev.* **30**: 713-43 [PMID:19770291]

Pasquier J et al. (2014) Molecular evolution of GPCRs: Kisspeptin/kisspeptin receptors. *J. Mol. Endocrinol.* **52**: T101-17 [PMID:24577719]

Leukotriene receptors

G protein-coupled receptors → Leukotriene receptors

**Overview**: The leukotriene receptors (nomenclature as agreed by the NC-IUPHAR subcommittee on Leukotriene Receptors [257, 258]) are activated by the endogenous ligands leukotrienes (LT), synthesized from lipoxygenase metabolism of arachidonic acid. The human BLT1 receptor is the high affinity LT4 receptor whereas the BLT2 receptor in addition to being a low-affinity LT4 receptor also binds several other lipoxygenase-products, such as 12S-HETE, 12S-HPETE, 15S-HETE, and the thromboxane synthesize 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, LT4 has been reported to bind to the peroxisome proliferator activated receptor (PPAR) α [1178] and the vanilloid TRPV1 ligand-gated nonselective cation channel [1307]. 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 [258]. Cysteinyl-leukotrienes have also been suggested to signal through the P2Y12 receptor [570, 1473, 1534], GPR17 [359] and GPR99 [943].

| Nomenclature | BLT1 receptor | BLT2 receptor | CysLT1 receptor | CysLT2 receptor |
|--------------|--------------|--------------|-----------------|---------------|
| HGNC, UniProt | LTB4R, Q15722 | LTB4R2, Q9NPC1 | CYSLTR1, Q9Y271 | CYSLTR2, Q9NS75 |
| Potency order of endogenous ligands | LTB4 > 20-hydroxy-LTB4 \( \gg 12R\)-HETE [2185] | 12-hydroxyheptadecatrienoic acid > LTB4 \( \gg 12R\)-HETE = 12S-HPETE > 15S-HETE > 20-hydroxy-LTB4 [1510, 2185] | LTD4 > LTC4 > LTE4 [1222, 1716] | LTD4 > LTD4 > LTE4 [767, 1477, 1920] |
| Endogenous agonists | – | 12S-HETE (Partial agonist) [2185] | – | – |
| Antagonists | – | – | ICI198615 (pK<sub>i</sub> 9.7) [591] – Guinea pig zafirlukast (pIC<sub>50</sub> 8.6–8.7) [1222, 1716], SR2640 (pK<sub>i</sub> 8.7), montelukast (pIC<sub>50</sub> 8.3–8.6) [1222, 1716], sulukast (pK<sub>i</sub> 8.3), pobilukast (pIC<sub>50</sub> 7.5) [1716] | BayCysLT2 (pIC<sub>50</sub> 6.6–7.3) [1456] |
| Selective antagonists | BILL 260 (pK<sub>i</sub> 8.8) [156, 485], CP105696 (pIC<sub>50</sub> 8.1) [1803], U75302 (pK<sub>i</sub> 6.4) [174] | LY255283 (pIC<sub>50</sub> 6–7.1) [780, 2185] | ICI198615 (pK<sub>i</sub> 9.7) [591] – Guinea pig zafirlukast (pIC<sub>50</sub> 8.6–8.7) [1222, 1716], SR2640 (pK<sub>i</sub> 8.7), montelukast (pIC<sub>50</sub> 8.3–8.6) [1222, 1716], sulukast (pK<sub>i</sub> 8.3), pobilukast (pIC<sub>50</sub> 7.5) [1716] | – |
| Labelled ligands | \(^{3}\)H]LTB4 (Agonist) [2184], \(^{3}\)H]CGS23131 (Antagonist) (pK<sub>d</sub> 7.9) [877] | \(^{3}\)H]LTB4 (pK<sub>d</sub> 7.6–9.7) | \(^{3}\)H]LTC4 (Agonist) [1870], \(^{3}\)H]CI-198615 (Antagonist) (pK<sub>d</sub> 10.6) [1682] | \(^{3}\)H]LTD4 (Agonist) [767] |

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Nomenclature | OXE receptor | FPR2/ALX
---|---|---
HGNC, UniProt | OXER1, Q8TDS5 | FPR2, P25090
Potency order of endogenous ligands | 5-oxo-ETE, 5-oxo-C20:3, 5-oxo-ODE > 5-oxo-15-HETE > SS-HPETE > SS-HETE | LXA4 = aspirin triggered lipoxin A4 = ATLα2 = resolvins D1 > LTC4 = LTD4 >> 15-deoxy-LXA4 >> fMet-Leu-Phe [365, 544, 546, 684, 1919]
Endogenous agonists | 5-oxo-ETE [672, 1483, 1538, 1597, 1752] | LXA4 [1052], resolvins D1 [1052], aspirin-triggered resolvins D1 [1051], aspirin triggered lipoxin A4
Selective agonists | – | ATLα2 [697]
Endogenous antagonists | 5-oxo-12-HETE (pIC\textsubscript{50} 6.3) [1596] | –
Selective antagonists | – | WRWWWWW (pIC\textsubscript{50} 6.6) [83], t-Boc-FLFLF (pIC\textsubscript{50} 4.3–6) [574, 1867, 2061]
Labelled ligands | [\textsuperscript{3}H]5-oxo-ETE (Agonist) [1483] | [\textsuperscript{3}H]LXA\textsubscript{4} (Agonist) [544, 545]

**Comments:** The FPR2/ALX receptor (nomenclature as agreed by the NC-IUPHAR subcommittee on Leukotriene and Lipoxin Receptors [258]) is activated by the endogenous lipid-derived, anti-inflammatory ligands lipoxin A\textsubscript{4} (LXA\textsubscript{4}) and 15-epi-LXA\textsubscript{4} (aspirin triggered lipoxin A\textsubscript{4}, ATL). The FPR2/ALX receptor also interacts with endogenous peptide and protein ligands, such as MHC binding peptide [330] as well as annexin I (ANXA1, P04083) (ANXA1) and its N-terminal peptides [379, 1560]. 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 [1646]. Furthermore, FPR2/ALX has been suggested to act as a receptor mediating the proinflammatory actions of the acute-phase reactant, serum amyloid A [1840, 1883]. The agonist activity of the lipid mediators described has been questioned [732, 1585], which may derive from batch-to-batch differences, partial agonism or biased agonism. Recent results from Cooray et al. (2013) [379] have addressed this issue and the role of homodimers and heterodimers in intracellular signaling. A receptor selective for LXB\textsubscript{4} has been suggested from functional studies [58, 1232, 1670]. Note that the data for FPR2/ALX are also reproduced on the Formylpeptide receptor pages.

Oxoeicosanoid receptors (OXE, nomenclature agreed by the NC-IUPHAR subcommittee on Oxoeicosanoid Receptors [219]) are activated by endogenous chemotactic eicosanoid ligands oxidised at the C-5 position, with 5-oxo-ETE the most potent agonist identified for this receptor. Initial characterization of the heterologously expressed OXE receptor suggested that polyunsaturated fatty acids, such as docosahexanoic acid and EPA, acted as receptor antagonists [823].

**Further reading on Leukotriene receptors**

Bäck M et al. (2011) International Union of Basic and Clinical Pharmacology. LXXXIV: leukotriene receptor nomenclature, distribution, and pathophysiological functions. Pharmacol. Rev. 63: 539-84 [PMID:21771892]
Bäck M et al. (2014) Update on leukotriene, lipoxin and oxoeicosanoid receptors: IUPHAR Review 7. Br. J. Pharmacol. 171: 3551-74 [PMID:24588652]

Brink C et al. (2004) International Union of Pharmacology XLIV. Nomenclature for the Oxoeicosanoid Receptor. Pharmacol. Rev. 56: 149-157 [PMID:15001665]
Brink C et al. (2003) International Union of Pharmacology XXXVII. Nomenclature for leukotriene and lipoxin receptors. Pharmacol. Rev. 55: 195-227 [PMID:12615958]
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 [414, 983]) 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 recently solved and demonstrates extracellular LPA access to the binding pocket, consistent with proposed delivery via autotaxin. These studies have also implicated cross-talk with endocannabinoids via phosphorylated intermediates that can also activate these receptors. 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. [586]). In native systems, analysis of binding data is complicated by metabolism and high levels of nonspecific binding, and therefore the relationship 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 [864]. LPA has also been described as an agonist for the transient receptor potential (Trp) ion channel TRPV1 [1461] and TRPA1 [1012]. In addition, orphan GPCRs (PSP24 [547] and GPR87 [1488]) are proposed as LPA receptors. LPA was originally proposed to be a ligand for GPCR35, but recent data shows that in fact it is a receptor for CXCL17 (CXCL17, Q6UXB2) [1266]. Further, the nuclear hormone receptor PPARγ [1309, 1812], has been reported as an LPA receptor. All of these proposed entities require confirmation and are not currently recognized as bona fide LPA receptors.

| Nomenclature | LPA₁ receptor | LPA₂ receptor | LPA₃ receptor | LPA₄ receptor | LPA₅ receptor | LPA₆ receptor |
|--------------|--------------|--------------|--------------|--------------|--------------|--------------|
| HGNC, UniProt | LPAR1, Q92633 | LPAR2, Q9HBW0 | LPAR3, Q9UBYS | LPAR4, Q99677 | LPAR5, Q9H1C0 | LPAR6, P43657 |
| Selective agonists | – | dodecylphosphate [2038], decyl dihydrogen phosphate [2038], GRI977143 [1007] | – | – | – | – |
| Sub/family-selective antagonists | Ki16425 (pIC₅₀ 6.6–6.9) [1499] – Mouse, VPC12249 (pKᵢ 5.2–6.9) [769] – Mouse, VPC32179 [763] | – | Ki16425 (pKᵢ 6.4) [1499], VPC12249 (pKᵢ 6.4) [769], VPC32179 [763] | – | – | – |
| Selective antagonists | BMS-986020 (pIC₀ 6.7–7.8) [1905], ONO-7300243 (pIC₀ 6.8) [1905], AM0995 (pIC₀ 6.6–6.1) [1905] | – | dioctanoylglycerol pyrophosphate (pKᵢ 5.5–7) [548, 1499] | – | TCLA5 (pIC₀ 6.1) [1047] | – |

Comments: Ki16425 [1499], VPC12249 [769] and VPC32179 [763] have dual 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 [2197], fetal hypoxia [778], lung fibrosis [1495], systemic sclerosis [1495] and atherosclerosis progression [1053]. Virtual screening experiments have shown H2L5186303 to be a potent antagonist of LPA₂ [536]. Dodecylphosphate is also an antagonist at LPA₃ receptors [2038].

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of Concise Guide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Further reading on Lysophospholipid (LPA) receptors

Chun J et al. (2010) International Union of Basic and Clinical Pharmacology. LXXVIII. Lysophospholipid receptor nomenclature. Pharmacol. Rev. 62: 579-87 [PMID:21079037]
Kiara Y et al. (2014) Lysophospholipid receptor nomenclature review: IUPHAR Review 8. Br. J. Pharmacol. 171: 3575-94 [PMID:24602016]

Yung YC et al. (2014) LPA receptor signaling; pharmacology, physiology, and pathophysiology. J. Lipid Res. 55: 1192-1214 [PMID:24643338]
Yung YC et al. (2015) Lysophosphatidic Acid signaling in the nervous system. Neuron 85: 669-82 [PMID:25695267]

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 [983]) 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 S1P1R, 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 [915], and awaits precise definition. Previously-proposed SPC (or lysophosphatidylcholine) receptors- G2A, TDAG8, OGR1 and GPR4- continue to lack confirmation of these roles [414]. The relationship between recombinant and endogenously expressed receptors is unclear. Radioligand binding has been conducted in heterologous expression systems using [32P]S1P [e.g 1505]. 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 [733].

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 [340, 369, 687], although the precise nature of its interaction requires clarification.

| Nomenclature | S1P1 receptor | S1P2 receptor | S1P3 receptor | S1P4 receptor | S1P3 receptor |
|-------------|---------------|---------------|---------------|---------------|---------------|
| HGNC, UniProt | S1P1R, P21453 | S1P2R, O95136 | S1P3R, Q99500 | S1P4R, Q95977 | S1P3R, Q9H228 |
| Potency order of endogenous ligands | sphingosine 1-phosphate > dihydrosphingosine 1-phosphate > sphingosylphosphorylcholine [46, 1505] | sphingosine 1-phosphate > dihydrosphingosine 1-phosphate > sphingosylphosphorylcholine [46, 1505] | sphingosine 1-phosphate > dihydrosphingosine 1-phosphate > sphingosylphosphorylcholine [1050] | sphingosine 1-phosphate > dihydrosphingosine 1-phosphate > sphingosylphosphorylcholine [2012] | sphingosine 1-phosphate > dihydrosphingosine 1-phosphate > sphingosylphosphorylcholine [861] |
| Agonists | amiselimod phosphate [1887], FTY720-phosphate [220, 560, 1525], siponimod [1524], AUY924 [1525], AFD(2) [220], etrasimod [254], fingolimod [708] | – | – | – | – |
| Selective agonists | ozaninmod [657, 1274, 1757], ponesimod [176], KRP 203-phosphate [1851] – Mouse, CYMS181 [657], SEW2871 [1713] – Mouse | – | – | – | – |
| Antagonists | VPC23019 (pK<sub>i</sub> 7.9) [419], VPC03090-P (pK<sub>i</sub> 7.6–7.7) [972], VPC44116 (pIC<sub>50</sub> 7.6) [561] | – | VPC44116 (pK<sub>i</sub> 6.5) [561], VPC23019 (pK<sub>i</sub> 5.9) [419] | – | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature  | S1P₁ receptor | S1P₂ receptor | S1P₃ receptor | S1P₄ receptor | S1P₅ receptor
--- | --- | --- | --- | --- | ---
Selective antagonists  | NIBR-0213 (pIC₅₀ 8.6) [1615], W146 (pKᵢ 7.1) [1714]  | JTE-013 (pIC₅₀ 7.8) [1515]  | –  | –  | –

**Comments**: The FDA-approved immunomodulator fingolimod (FTY720) can be phosphorylated in vivo [31] to generate a relatively potent agonist with activity at S1P₁, S1P₃, S1P₄ and S1P₅ receptors [220, 1259]. The physiological consequences of FTY720-phosphate administration, as well as those of other S1P₁ agonists, may involve functional antagonism via ubiquitination and subsequent degradation of S1P₁ [1514].

**Further reading on Lysophospholipid (S1P) receptors**

Chew WS et al. (2016) To fingolimod and beyond: The rich pipeline of drug candidates that target S1P signaling. *Pharmacol. Res.* **113**: 521-532 [PMID:27663260]

Chun J et al. (2010) International Union of Basic and Clinical Pharmacology. LXXVIII. Lysophospholipid receptor nomenclature. *Pharmacol. Rev.* **62**: 579-87 [PMID:21079037]

Pyne NJ et al. (2017) Sphingosine 1-Phosphate Receptor 1 Signaling in Mammalian Cells. *Molecules* **22**: [PMID:28241498]

Rosen H et al. (2013) Sphingosine-1-phosphate and its receptors: structure, signaling, and influence. *Annu. Rev. Biochem.* **82**: 637-62 [PMID:23527695]

**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 [557]) are activated by an endogenous nonadecameric cyclic peptide identical in humans and rats (DFDMLRCMLGRVYRPCWQV) generated from a precursor (*PMCH, P20382*), which also produces neuropeptide E1 (*PMCH, P20382*) and neuropeptide GE (*PMCH, P20382*).

| Nomenclature | MCH₁ receptor | MCH₂ receptor |
| --- | --- | --- |
| HGNC, UniProt | MCHR1, Q99705 | MCHR2, Q969V1 |
| Selective antagonists | GW803430 (pIC₅₀ 9.3) [781], SNAP-7941 (pA₂ 9.2) [190], T-226296 (pIC₅₀ 8.3) [1926], ATC0175 (pIC₅₀ 7.9–8.1) [297] | – |
| Labelled ligands | [¹²⁵I]S36057 (Antagonist) (pKᵦ 9.2–9.5) [69], [¹²⁵I][Phe¹³,Tyr¹⁹]MCH (Agonist) [249], [³H]MCH (human, mouse, rat) (Agonist) [249] | – |

**Comments**: The MCH₂ receptor appears to be a non-functional pseudogene in rodents [1930].

**Searchable database**: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide**: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Further reading on Melanin-concentrating hormone receptors

Chung S et al. (2011) Recent updates on the melanin-concentrating hormone (MCH) and its receptor system: lessons from MCH1R antagonists. J. Mol. Neurosci. 43: 115-21 [PMID:20582487]
Eberle AN et al. (2010) Cellular models for the study of the pharmacology and signaling of melanin-concentrating hormone receptors. J. Recept. Signal Transduct. Res. 30: 385-402 [PMID:21083507]

Melanocortin receptors

Overview: Melanocortin receptors (provisional nomenclature as recommended by NC-IUPHAR [557]) are activated by members of the melanocortin family (α-MSH (POMC, P01189), β-MSH (POMC, P01189) and γ-MSH (POMC, P01189) forms; δ form is not found in mammals) and adrenocorticotrophin (ACTH (POMC, P01189)). Endogenous antagonists include agouti (ASIP, P42127) and agouti-related protein (AGRP, O00253). ACTH(1-24) was approved by the US FDA as a diagnostic agent for adrenal function test. At least 2 synthetic melanocortin receptor agonists are under clinical development as of 2017.

| Nomenclature | MC1 receptor | MC2 receptor | MC3 receptor | MC4 receptor | MC5 receptor |
|--------------|--------------|--------------|--------------|--------------|--------------|
| HGNC, UniProt | MC1R, Q01726 | MC2R, Q01718 | MC3R, P41968 | MC4R, P32245 | MC5R, P33032 |
| Potency order of endogenous ligands | α-MSH (POMC, P01189) > β-MSH (POMC, P01189) > ACTH (POMC, P01189), γ-MSH (POMC, P01189) | ACTH (POMC, P01189) | γ-MSH (POMC, P01189), β-MSH (POMC, P01189) > ACTH (POMC, P01189), α-MSH (POMC, P01189) | β-MSH (POMC, P01189) > α-MSH (POMC, P01189) > ACTH (POMC, P01189) > γ-MSH (POMC, P01189) | α-MSH (POMC, P01189) > β-MSH (POMC, P01189) > ACTH (POMC, P01189) > γ-MSH (POMC, P01189) |
| Selective agonists | − | corticotropic zinc hydroxide | [D-Trp8]γ-MSH [679] | THIQ [1760] | − |
| Antagonists | − | − | PG-106 (pIC50 6.7) [680] | − | − |
| Selective antagonists | − | − | − | MBP10 (pIC50 10) [123], HS014 (pKi 8.5) [1738] | − |
| Labelled ligands | [125]I NDP-MSH (Agonist) [1037] | [125]I ACTH-(1-24) (Agonist) | [125]I NDP-MSH (Agonist) [1037], [125]I SHU9119 (Agonist) [1457] | [125]I SHU9119 (Agonist) (pKD 9.2) [1457], [125]I NDP-MSH (Agonist) [1037, 1736] | [125]I NDP-MSH (Agonist) [1037] |

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 [296, 531].
## Melatonin receptors

**G protein-coupled receptors → Melatonin receptors**

**Overview:** Melatonin receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Melatonin Receptors [474]) are activated by the endogenous ligands melatonin and clinically used drugs like ramelteon and agomelatine.

| Nomenclature | MT₁ receptor | MT₂ receptor |
|--------------|--------------|--------------|
| HGCN, UniProt | MTRNR1A, P48039 | MTRNR1B, P49286 |
| Endogenous agonists | melatonin [70, 473, 475] | melatonin [70, 473, 475] |
| Agonists | ramelteon [954], agomelatine [70, 136] | agomelatine [70, 136], ramelteon [954, 1636] |
| Selective agonists | – | UCM1014 [1855], IIG7 [532, 1888], 5-methoxy-luzindole (Partial agonist) [475] |
| Selective antagonists | – | 4P-PDOT (pKᵢ 8.8–9.4) [70, 475, 476], K185 (pKᵢ 9.3) [532, 1888], DH97 (pKᵢ 8) [1939] |
| Labelled ligands | [¹²⁵I]SD6 (Agonist) [1138], 2-[¹²⁵I]melatonin (Agonist) [70, 475], [³H]melatonin (Agonist) [235] | [¹²⁵I]SD6 (Agonist) [1138], 2-[¹²⁵I]melatonin (Agonist) [70, 475], [¹²⁵I]DIV880 (Agonist, Partial agonist) [1138], [³H]melatonin (Agonist) [235] |

**Comments:** melatonin, 2-iodo-melatonin, agomelatine, GR 196429, 17 156735 and ramelteon [954] are nonselective agonists for MT₁ and MT₂ receptors. (+)-AMMTTC displays an ~400-fold greater agonist potency than (-)-AMMTTC at rat MT₂ receptors (see AMMTTC for structure) [1996]. Luzindole is an MT₁/MT₂ non-selective competitive melatonin receptor antagonist with about 15-25 fold selectivity for the MT₂ receptor [476]. MT₁/MT₂ heterodimers present different pharmacological profiles from MT₁ and MT₂ receptors [75].

The MT₂ 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 [1356]. Pharmacological investigations of MT₂ binding sites have primarily been conducted in hamster tissues. At this site, The endogenous ligand N-acetylsalicylic acid [495, 1215, 1356, 1588] and SMCA-NAT [1588] appear to function as agonists, while prazosin [1215] functions as an antagonist. The MT₂ binding site of hamster kidney was also identified as the hamster homologue of human quinone reductase 2 (NQO2, P16083 [1474, 1475]). The MT₃ binding site activated by SMCA-NAT in eye ciliary body is positively coupled to adenyl cyclase and regulates chloride secretion [842]. Xenopus melanophores and chick brain express a distinct receptor (x420, P49219; c346, P49288, initially termed Mel₁ argues coupled to the Gᵢ/o family of G proteins, for which GPR50 has recently been suggested to be a mammalian counterpart [479] although melatonin does not bind to GPR50 receptors. Several variants of the MTNR1B gene have been associated with increased type 2 diabetes risk.

---

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
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 [1743]) are a family of G-protein coupled receptors activated by the neurotransmitter glutamate. The mGlu family is composed of eight members (named mGlu1 to mGlu8) which are divided in three groups based on similarities of agonist pharmacology, primary sequence and G protein coupling to effector: Group-I (mGlu1 and mGlu5), Group-II (mGlu2 and mGlu3) and Group-III (mGlu4, mGlu6, mGlu7 and mGlu8) (see Further reading).

Structurally, mGlu are composed of three juxtaposed domains: a core G-protein-activating seven-transmembrane domain (TM), common to all GPCRs, is linked via a rigid cysteine-rich domain (CRD) to the Venus Flytrap domain (VFTD), a large bi-lobed extracellular domain where glutamate binds. The structures of the VFTD of mGlu1, mGlu2, mGlu3, mGlu5 and mGlu7 have been solved [1075, 1364, 1408, 1984]. The structure of the 7 transmembrane (TM) domains of both mGlu1 and mGlu5 have been solved, and confirm a general helical organization similar to that of other GPCRs, although the helices appear more compacted [465, 2136]. mGlu form constitutive dimers crosslinked by a disul-fide bridge. 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 [468]. Although well characterized in transfected cells, co-localization and specific pharmacological properties also suggest the existence of such heterodimers in the brain [2183]. The endogenous ligands of mGlu are L-glutamic acid, L-serine-O-phosphate, N-acetylaspartylglutamate (NAAG) and L-cysteine sulphinic acid. Group-I mGlu receptors may be activated by 3,5-DHPG and (S)-3HPG [204] and antagonized by (S)-hexylhomolbotenic acid [1235]. Group-II mGlu receptors may be activated by LY389795 [1365, LY379268 [1365], eglumegad [1744, 2138], DCG-IV and (2R,3R)-APDC [1745], and antagonised by eGlu [890] and LY307452 [518, 2096]. Group-III mGlu receptors may be activated by L-AP4 and (R,S)-4-PPG [610]. An example of an antagonist selective for mGlu receptors is LY341495, which blocks mGlu2 and mGlu4 at low nanomolar concentrations, mGlu6 at high nanomolar concentrations, and mGlu4, mGlu5, and mGlu7 in the micromolar range [1001]. In addition to orthosteric ligands that directly interact with the glutamate recognition site, allosteric modulators that bind within the TM domain 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.

| Nomenclature | mGlu1 receptor | mGlu2 receptor | mGlu3 receptor | mGlu4 receptor | mGlu5 receptor |
|--------------|----------------|----------------|----------------|----------------|----------------|
| HGNC, UniProt | GRM1, Q13255   | GRM2, Q14416   | GRM3, Q14832   | GRM4, Q14833   | GRM5, P41594   |
| Endogenous agonists | L-glutamic acid [1574] | L-glutamic acid [1574] | L-glutamic acid [1574], NAAG [1750] | L-glutamic acid [1574] | L-glutamic acid [1574] |
| Agonists | – | – | – | L-AP4 [2138], L-serine-O-phosphate [2138] | – |
| Selective agonists | – | – | – | LSP4-2022 [666] | (S,)-CBPG (Partial agonist) [1261] |
| Antagonists | LY367385 (pIC50 5.1) [364] | – | – | MAP4 (pK1 4.6) [721] – Rat, CHPG [1407] | – |

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

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
### Nomenclature

| Receptor   | mGlu₁ receptor | mGlu₂ receptor | mGlu₃ receptor | mGlu₄ receptor | mGlu₅ receptor |
|------------|----------------|----------------|----------------|----------------|----------------|
| **Selective antagonists** | 3-MATIDA (pIC₅₀ 5.2) [1386] – Rat, (S)–(-)–CBPG (pIC₅₀ 4.2) [1261] – Rat, (S)–TBPG (pIC₅₀ 4.2) [381] – Rat, AIDA (pA₂ 4.2) [1387] | PCCG-4 (pIC₅₀ 5.1) [1551] – Rat | – | – | ACDPP (pIC₅₀ 6.9) [186] |
| **Selective allosteric modulators** | BAY 367620 (Negative) (pKᵢ 9.5) [279] – Rat, IN16259685 (Negative) (pIC₅₀ 8.9) [1104], Ro01-6128 (Positive) (pKᵢ 7.5–7.7) [1019] – Rat, LY456236 (Negative) (pIC₅₀ 6.9) [1160], CPCCC0Et (Negative) (pIC₅₀ 5.2–5.8) [1183] | Ro64-5229 (Negative) (pIC₅₀ 7) [1031] – Rat, biphenylindanone A (Positive) (pEC₅₀ 7) [187] | – | VU0361737 (Positive) (pEC₅₀ 6.6) [508], VU0155041 (Positive) (pEC₅₀ 6.1) [1468] | VU-1545 (Positive) (pEC₅₀ 8) [429] |

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)  
**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Comments: The activity of NAAG as an agonist at mGlu3 receptors was questioned on the basis of contamination with glutamate [341, 576], but this has been refuted [1430].

Radioligand binding using a variety of radioligands has been conducted on recombinant receptors (for example, [3H]R214127 [1103] and [3H]YM298198 [1025] at mGlu3 receptors and [3H]M-MPEP [609] and [3H]methoxymethyl-MTEP [48] 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 [2007], and is an antagonist at all Group-III mGluRs with an IC50 of 30 μM.

A potential novel metabotropic glutamate receptor coupled to phosphoinositide turnover has been observed in rat brain; it is activated by 4-methylhomoibotenic acid (ineffective as an agonist at recombinant Group I metabotropic glutamate receptors), but is resistant to LY341495 [356]. 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 [1013, 1549].

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.

Further reading on Metabotropic glutamate receptors

Conn PJ et al. (1997) Pharmacology and functions of metabotropic glutamate receptors. Annu. Rev. Pharmacol. Toxicol. 37: 205-237 [PMID:9131252]
Ferraguti F et al. (2006) Metabotropic glutamate receptors. Cell Tissue Res. 326: 483-504 [PMID:16847639]
Nicoletti F et al. (2011) Metabotropic glutamate receptors: from the workbench to the bedside. Neuropsychopharmacology 60: 1017-41 [PMID:21036182]
Niswender CM et al. (2010) Metabotropic glutamate receptors: physiology, pharmacology, and disease. Annu. Rev. Pharmacol. Toxicol. 50: 295-322 [PMID:20055706]

Motilin receptor

G protein-coupled receptors → Motilin receptor

Overview: Motilin receptors (provisional nomenclature) are activated by motilin (MLN, P12872), a 22 amino-acid peptide derived from a precursor (MLN, P12872), which may also generate a motilin-associated peptide (MLN, P12872). These receptors promote gastrointestinal motility and are 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) [386, 1286, 1287, 1288]
Agonists alemcinal [1947], erythromycin-A [533, 1947], azithromycin [225]
Selective agonists camicinal [105, 1712], mitemcinal [1023, 1918] – Rabbit
Selective antagonists MA-2029 (pA2 9.2) [1884], GM-109 (pIC50 8) [736] – Rabbit
Labelled ligands [125I]motilin (human) (Agonist) [533]

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
**Comments**: In terms of structure, the motilin receptor has closest homology with the ghrelin receptor. Thus, the human motilin receptor shares 52% overall amino acid identity with the ghrelin receptor and 86% in the transmembrane regions [759, 1918, 1947]. However, differences between the N-terminus regions of these receptors means that their cognate peptide ligands do not readily activate each other [408, 1712]. In laboratory rodents, the gene encoding the motilin precursor appears to be absent, while the receptor appears to be a pseudogene [759, 1710]. 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 is obscure [1311, 1462]. Notably, in some non-laboratory rodents (e.g. the North American kangaroo rat (Dipodomys) and mouse (Microdipodops) a functional form of motilin may exist but the motilin receptor is non-functional [1159]. Marked differences in ligand affinities for the motilin receptor in dogs and humans may be explained by significant differences in receptor structure [1711]. 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; [2216]). Small molecule motilin receptor agonists are now described [1159, 1712, 2100]. The motilin receptor does not appear to have constitutive activity [812]. Although not proven, the existence of biased agonism at the receptor has been suggested [1288, 1348, 1709]. A truncated 5-transmembrane structure has been identified but this is without activity when transfected into a host cell [533]. Receptor dimerisation has not been reported.

**Further reading on Motilin receptor**

De Smet B et al. (2009) Motilin and ghrelin as prokinetic drug targets. *Pharmacol. Ther.* **123**: 207-23 [PMID:19427331]

Sanger GJ et al. (2016) Ghrelin and motilin receptors as drug targets for gastrointestinal disorders. *Nat Rev Gastroenterol Hepatol* **13**: 38-48 [PMID:26392067]

---

**Neuromedin U receptors**

**G protein-coupled receptors → Neuromedin U receptors**

**Overview**: Neuromedin U receptors (provisional nomenclature as recommended by NC-IUPHAR [557]) 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 [1344]. 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 levels 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 (NM'S, Q5H8A3)) has also been identified as an endogenous agonist [1378]. 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, Q9CZQ4 |
| Antagonists   | –             | R-PSOP (pK₈ 7) [1193] |

**Comments**: NMU1 and NMU2 couple predominantly to Gq/11, although there is evidence of good coupling to Gs[41]. NMU1 and NMU2 can be labelled with [¹²⁵I]-NmU and [¹²⁵I]-NmS (of various species, e.g. [1319]), BODIPY® TMR-NMU or Cy3B-NMU-8 [218]. 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 (NM'S, Q5H8A3) are now commercially available.
Further reading on Neuromedin U receptors

Brighton PJ et al. (2004) Neuromedin U and its receptors: structure, function, and physiological roles. Pharmacol. Rev. 56: 231-48 [PMID:15169928]
Budhiraja S et al. (2009) Neuromedin U: physiology, pharmacology and therapeutic potential. Fundam Clin Pharmacol 23: 149-57 [PMID:19645813]

Mitchell JD et al. (2009) Emerging pharmacology and physiology of neuromedin U and the structurally related peptide neuromedin S. Br. J. Pharmacol. 158: 87-103 [PMID:19519756]
Novak CM. (2009) Neuromedin S and U. Endocrinology 150: 2985-7 [PMID:19549882]

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 [557]), 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 | NPFFR1, Q9GZQ6 | NPFFR2, Q9YSX5 |

Potency order of endogenous ligands

RFRP-1 (NPVF, Q9HCQ7) > RFRP-3 (NPVF, Q9HCQ7) > FMRFneuropeptide FF (NPFF, O15130) > neuropeptide FF (NPFF, O15130) > neuropeptide SF (NPFF, O15130), QRFP43 (QRFP, P83859), PrRP-31 (PRLH, P81277) [663]

Endogenous agonists

neuropeptide FF (NPFF, O15130) [663, 664, 1359], RFRP-3 (NPVF, Q9HCQ7) [664, 665, 1359]

Selective agonists

–

Antagonists

RF9 (pK_i 7.2) [1814]

Selective antagonists

AC262620 (pK_i 7.7–8.1) [1092], AC262970 (pK_i 7.4–8.1) [1092]

Labelled ligands

[^125I]Y-RFRP-3 (Agonist) [664], [3H]NPVF (Agonist) [1928], [^125I]NPFF (Agonist) [663], [^125I]EYF (Agonist) [1359], [3H]EYF (Agonist) [1928], [^125I]NPFF (Agonist) [663]

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, [1814]).
Further reading on Neuropeptide FF/neuropeptide AF receptors

Moulédous L et al. (2010) Opioid-modulating properties of the neuropeptide FF system. *Biofactors* **36**: 423-9 [PMID:20803521]
Vyas N et al. (2006) Structure-activity relationships of neuropeptide FF and related peptidic and non-peptidic derivatives. *Peptides* **27**: 990-6 [PMID:16490282]

Yang HY et al. (2008) Modulatory role of neuropeptide FF system in nociception and opiate analgesia. *Neuropeptides* **42**: 1-18 [PMID:17854890]

Neuropeptide S receptor

**G protein-coupled receptors → Neuropeptide S receptor**

**Overview:** The neuropeptide S receptor (NPS, provisional nomenclature [557]) responds to the 20 amino-acid peptide neuropeptide S derived from a precursor (NPS, P0C0P6).

| Nomenclature | NPS receptor |
|-------------|-------------|
| HGNC, UniProt | NPSR1, Q6W5P4 |
| Endogenous agonists | neuropeptide S (NPS, P0C0P6) [2159] |
| Selective agonists | PWT1-NPS [1692] – Mouse |
| Selective antagonists | NCGC 84 (pA2 9) [1957], SHA 68 (pA2 8.1) [1693] – Mouse, RTI-118 [2214] |
| Labeled ligands | [125I]Tyr10NPS (human) (Agonist) [2159] |

**Comments:** Multiple single-nucleotide polymorphisms (SNP) and several splice variants have been identified in the human NPS receptor. The most interesting of these is an Asn-Ile exchange at position 107 (Asn107Ile). The human NPS receptor Asn107Ile displayed similar binding affinity but higher NPS potency (by approx. 10-fold) than human NPS receptor Asn107 [1645]. Several epidemiological studies reported an association between Asn107Ile receptor variant and susceptibility to panic disorders [458, 460, 1506, 1621]. The SNP Asn107Ile has also been linked to sleep behavior [662], inflammatory bowel disease [402], schizophrenia [1145], increased impulsivity and ADHD symptoms [1083]. Interestingly, a carboxy-terminal splice variant of human NPS receptor was found to be overexpressed in asthmatic patients [1091].

Further reading on Neuropeptide S receptor

Guerrini R et al. (2010) Neurobiology, pharmacology, and medicinal chemistry of neuropeptide S and its receptor. *Med Res Rev* **30**: 751-77 [PMID:19824051]

Xu YL et al. (2004) Neuropeptide S: a neuropeptide promoting arousal and anxiolytic-like effects. *Neuron* **43**: 487-497 [PMID:15312648]

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Neuropeptide W/neuropeptide B receptors

Overview: The neuropeptide BW receptor 1 (NPBW1, provisional nomenclature [587]) is activated by two 23-amino-acid peptides, neuropeptide W (neuropeptide W-23 (NPW, Q8N729)) and neuropeptide B (neuropeptide B-23 (NPB, Q8NG41)) [584, 1792]. C-terminally extended forms of the peptides (neuropeptide W-30 (NPW, Q8N729) and neuropeptide B-29 (NPB, Q8NG41)) also activate NPBW1 [216]. 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 [216].

| Nomenclature | Potency order of endogenous ligands | Selective agonists | Labeled ligands |
|--------------|-----------------------------------|-------------------|----------------|
| NPBW1 receptor | neuropeptide B-29 (NPB, Q8NG41) > neuropeptide B-23 (NPB, Q8NG41) > neuropeptide W-23 (NPW, Q8N729) (NPW, Q8N729) [216] | Ava3 [945], Ava5 [945] | [125I]NPW-23 (human) (Agonist) [1816] |
| NPBW2 receptor | neuropeptide W-30 (NPW, Q8N729) > neuropeptide W-23 (NPW, Q8N729) > neuropeptide B-29 (NPB, Q8NG41) > neuropeptide B-23 (NPB, Q8NG41) [216] | – | [125I]NPW-23 (human) (Agonist) [1792] |

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 [1414]. For a review of the contribution of neuropeptide W/B to social dominance, see [2080]. It has been reported that neuropeptide W may have a key role in the gating of stressful stimuli when mice are exposed to novel environments [1392]. Two antagonists have been discovered and reported to have affinity for NPBW1, ML181 and ML250, the latter exhibiting improved selectivity (~ 100 fold) for NPBW1 compared to MCH1 receptors [694, 695]. Computational insights into the binding of antagonists to this receptor have also been described [1541].

Further reading on Neuropeptide W/neuropeptide B receptors

Sakurai T. (2013) NPBWR1 and NPBWR2: Implications in Energy Homeostasis, Pain, and Emotion. Front Endocrinol (Lausanne) 4, 23 [PMID:23515889]

Singh G et al. (2006) Neuropeptide B and W: neurotransmitters in an emerging G-protein-coupled receptor system. Br. J. Pharmacol. 148: 1033-41 [PMID:16847439]
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 [1330]) are activated by the endogenous peptides neuropeptide Y (NPY, P01303), neuropeptide Y-(3-36), peptide YY (PYY, P10082), PYY-(3-36) and pancreatic polypeptide (PP, P01298) (PP). The receptor originally identified as the Y3 receptor has been identified as the CXCR4 chemokine receptor (originally named LESTR, [1201]). The y6 receptor is a functional gene product in mouse, absent in rat, but contains a frameshift mutation in primates producing a truncated non-functional gene [676]. 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 [513]. 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).

| Nomenclature | Y1 receptor | Y2 receptor | Y4 receptor | Y5 receptor | Y6 receptor |
|--------------|-------------|-------------|-------------|-------------|-------------|
| HGNC, UniProt| NPY1R, P25929 | NPY2R, P49146 | NPY4R, P50391 | NPY5R, Q15761 | NPY6R, Q99463 |
| Potency order of endogenous ligands| neuropeptide Y = peptide YY >> pancreatic polypeptide | peptide YY = peptide YY(3-36) = neuropeptide Y = neuropeptide Y-(3-36) >> 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) [619, 633], neuropeptide Y (NPY, P01303), neuropeptide Y-(3-36) (NPY, P01303), peptide YY (PYY, P10082) | pancreatic polypeptide (PP, P10298) [98, 1217, 1978, 2165] | – | – |
| Agonists| [Leu31,Pro34]NPY [392], [Leu31,Pro34]PYY (human), [Pro34]PYY (human) | – | – | – | – |
| Selective agonists| – | – | – | [Ala31,Aib32]NPY (pig) [264] | – |
| Selective antagonists| BIBO3304 (pIC50 9.5) [2110], BIBP3226 (pK1 6.9–7.1) [182] | BIIE0246 (pIC50 8.5) [461], JNJ-5207787 (pIC50 6.9–7.1) [182] | – | L-152,804 (pK1 7.6) [944] | – |
| Selective allosteric modulators| – | – | niclosamide (Positive) [1827] | – | – |
| Labelled ligands| [3H]BIBP3226 (Antagonist) (pKd 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) [481] | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature

Comments: The Y_1 agonists indicated are selective relative to Y_2 receptors. BIBP3226 is selective relative to Y_2, Y_4 and Y_5 receptors [632]. NPY-(13-36) is Y_2 selective relative to Y_1 and Y_5 receptors. PYY-(3-36) is Y_2 selective relative to Y_1 receptors. Note that Pro34-containing NPY and PYY can also bind Y_4 and Y_5, thus they are selective only relative to Y_2. The y_6 receptor is a pseudogene in humans, but is functional in mouse, rabbit and some other mammals.

Further reading on Neuropeptide Y receptors

Bowers ME et al. (2012) Neuropeptide regulation of fear and anxiety: Implications of cholecystokinin, endogenous opioids, and neuropeptide Y. *Physiol. Behav.* 107: 699-710 [PMID:22429904]

Michel MC et al. (1998) XVI. International Union of Pharmacology recommendations for the nomenclature of neuropeptide Y, peptide YY and pancreatic polypeptide receptors. *Pharmacol. Rev.* 50: 143-150 [PMID:9549761]

Pedragosa-Badia X et al. (2013) Neuropeptide Y receptors: how to get subtype selectivity. *Front Endocrinol (Lausanne)* 4: 5 [PMID:23382728]

Zhang L et al. (2011) The neuropeptide Y system: pathophysiological and therapeutic implications in obesity and cancer. *Pharmacol. Ther.* 131: 91-113 [PMID:21439311]

Neurotensin receptors

G protein-coupled receptors → Neurotensin receptors

Overview: Neurotensin receptors (nomenclature as recommended by NC-IUPHAR [557]) 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 NTS_2 receptor. [^H]neurotensin (human, mouse, rat) and [^{125}I]neurotensin (human, mouse, rat) may be used to label NTS_1 and NTS_2 receptors at 0.1-0.3 and 3-5 nM concentrations respectively.

| Nomenclature | NTS_1 receptor | NTS_2 receptor |
|--------------|----------------|----------------|
| HGNC, UniProt | NTSR1, P30989  | NTSR2, O95665  |
| Potency order of endogenous ligands | neurotensin (NTS, 30990) > neuromedin N (Mouse, Rat) [776] | neurotensin (NTS, P30990) = neuromedin N (Mouse, Rat) [1297] |
| Selective agonists | JMV449 [1822] – Rat | levocabastine [1297, 1657] |
| Selective antagonists | meclinertant (pIC_{50} 7.5–8.2) [699] | – |
| Labelled ligands | [^{3}H]meclinertant (Antagonist) (pK_{d} 8.5) [1085] – Rat | – |

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

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
**Comments:** neurotensin (NTS, P30990) appears to be a low-efficacy agonist at the NTS$_2$ receptor [2039], while the NTS$_1$ receptor antagonist meclintant is an agonist at NTS$_2$ receptors [2039]. An additional protein, provisionally termed NTS$_3$ (also known as NTR3, gp95 and sortilin; ENSG00000134243), has been suggested to bind lipoprotein lipase and mediate its degradation [1460]. It has been reported to interact with the NTS$_1$ receptor [1273] and the NTS$_2$ receptor [260], and has been implicated in hormone trafficking and/or neurotensin uptake. A splice variant of the NTS$_2$ receptor bearing 5 transmembrane domains has been identified in mouse [195] and later in rat [1561].

**Further reading on Neurotensin receptors**

Boules M et al. (2013) Diverse roles of neurotensin agonists in the central nervous system. *Front Endocrinol (Lausanne)* 4: 36 [PMID:23526754]

Mazella J et al. (2012) Neurotensin and its receptors in the control of glucose homeostasis. *Front Endocrinol (Lausanne)* 3: 143 [PMID:23230428]

Myers RM et al. (2009) Cancer, chemistry, and the cell: molecules that interact with the neurotensin receptors. *ACS Chem. Biol.* 4: 503-25 [PMID:19462983]

Tanganelli S et al. (2012) Relevance of dopamine D(2)/neurotensin NTS1 and NMDA/neurotensin NTS1 receptor interaction in psychiatric and neurodegenerative disorders. *Curr. Med. Chem.* 19: 304-16 [PMID:22335510]

**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-neoendorphin (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 appropriate, along with the symbols spelled out (mu, delta, and kappa), and the acronyms, MOP, DOP, and KOP. [390, 441, 557]. The human N/OFQ receptor, NOP, is considered ‘opioid-related’ rather than opioid because, while it exhibits a high degree of structural homology with the conventional opioid receptors [1361], it displays a distinct pharmacology. Currently there are numerous clinically used drugs, such as morphine and many other opioid analgesics, as well as antagonists such as naloxone, however only for the μ receptor.

**Nomenclature**

| Nomenclature | δ receptor | κ receptor | μ receptor | NOP receptor |
|--------------|------------|------------|------------|--------------|
| HGNC, UniProt | OPRD1, P41143 | OPRK1, P41145 | OPRM1, P35372 | OPR1, P41146 |
| Principal endogenous agonists | β-endorphin (POMC, P01189), [Leu]enkephalin (PENK, P01210), [Met]enkephalin (PENK, P01210) | big dynorphin (PDYN, P01213), dynorphin A (PDYN, P01213) | β-endorphin (POMC, P01189), [Met]enkephalin (PENK, P01210), [Leu]enkephalin (PENK, P01210) | nociceptin/orphanin FQ (PNOC, Q13519) |
| Potential endogenous agonists | – | – | endomorphin-1, endomorphin-2 | – |
| Agonists | DADLE [1972], etorphine [1972], ethylketocyclazocine [1972] | – | levorphanol [727], hydromorphone [2094], fentanyl [1972], buprenorphine (Partial agonist) [1972], methadone [1595], codeine [1972], tapentadol [1992], pethidine [1595] | – |

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
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 [1535], 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 [926], and give rise to complexes with a unique pharmacology, however, evidence for such heterodimers in native cells is equivocal and the consequences of this heterodimerization for signalling remains largely unknown. For µ-opioid receptors at least, dimerization does not seem to be required for signalling [1078]. 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 [2198]. 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 [1490] and the identification of biased signalling by opioid receptor ligands, in particular, compounds previously characterized as antagonists [236]. Pathway bias for agonists makes general rank orders of potency and efficacy somewhat obsolete, so these do not appear in

| Nomenclature | δ receptor | κ receptor | µ receptor | NOP receptor |
|--------------|------------|------------|------------|-------------|
| Sub/family-selective agonists | BU08028 (Partial agonist) [979] | BU08028 [979] | BU08028 (Partial agonist) [979] | cebranopadol [1182], BU08028 (Partial agonist) [979] |
| Selective agonists | UFP-512 [2033], BW373U86 [1115], ADLS859 [1115], DPDPE [1391, 1972], [D-Ala2]deltorphin II [515], ADLS747 [1116], SNC80 [268, 1620] | US0488 [313, 1545, 1813, 1972, 2046, 2222, 2224], enadoline [848, 1447], U69593 [1089, 1972], salvinorin A [259, 1677] | sufentanil [2041], DAMGO [726, 1972], loperamide [323], morphine [653, 1972], PL017 [304, 1972] | N/OFQ-(1-13)-NH2 [153, 696, 1304, 1507], Ac-RRyRy-WKH-NH2 (Partial agonist) [464, 1304], SCH221510 [2030], Ro64-6198 [898, 2108] |
| Antagonists | naltrexone (pKi 8) [1972], naloxone (pKi 7.2) [1972] | buprenorphine (pKi 9.1–10.2) [1972, 2224], nalmefene (pKi 9.5) [1972], naltrexone (pKi 8.4–9.4) [1545, 1813, 1972], naloxone (pKi 7.6–8.6) [1545, 1813, 1972, 2222, 2224] | naltrexone (pKi 9.1–9.7) [965, 1972], nalmefene (pKi 9.5) [1972], nalorphine (pKi 8.9) [1972], naloxone (pKi 8.9) [1972], methylnaltrexone (pKi 8.7) [2094] | – |
| Sub/family-selective antagonists | AT-076 (pKi 7.7) [1972, 2201] | AT-076 (pKi 8.9) [1972, 2202] | AT-076 (pKi 8.8) [1972, 2202] | – |
| Selective antagonists | naltriben (pKi 10) [1841, 1972], naltrindole (pKi 9.7) [1594, 1972], TIPP (Inverse agonist) (pKi 9) [1735, 1972] | nor-binaltorphimine (pKi 8.9–11) [1545, 1593, 1813, 1972, 2222, 2224], S′-guanidinotalnindole (pKi 9.7–9.9) [924, 1545, 1868], µTic (pKi 9–9.4) [1400, 1951, 2202] | alvimopan (pKi 9.3) [1114], levallorphan (pKi 8.8–9.3) [1250], CTAP (pKi 8.6) [304, 1972] | BMS-986123 (Neutral) (pKi 6) [247], BMS-986121 (Positive) (pKi 5.7) [247], BMS-986124 (Neutral) (pKi 5.7) [247], BMS-986122 (Positive) (pKi 6) [247] |
| Allosteric modulators | – | – | BMS-986122 (Neutral) (pKi 6) [247], BMS-986121 (Positive) (pKi 5.7) [247], BMS-986124 (Neutral) (pKi 5.7) [247], BMS-986122 (Positive) (pKi 6) [247] | – |
| Labelled ligands | [3H]Naltrindole (Antagonist) (pKd 10.4) [2161] – Rat, [3H][D-Ala2]deltorphin I (Selective Agonist) [1865], [3H]diprenorphine (Agonist) [52, 1972], [3H]DPDPE (Agonist) [26], [3H]deltorphin II (Agonist) [261], [3H]Naltriben (Antagonist) [1154] | [3H]Diprenorphine (Antagonist) (pKd 9.1) [52, 1813], [3H]U69593 (Agonist) [1089, 1545, 1813], [3H]enadoline (Agonist) [1813] | [3H]Diprenorphine (Antagonist) (pKd 10.1) [1638] – Mouse, [3H]DAMGO (Agonist) [1638] – Rat, [3H]PL017 (Agonist) [751] – Rat | [3H]N/OFQ (Agonist) [464, 1360] |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

Opioid receptors 996
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 \( \mu \) and \( \delta \) receptors, notably the positive allosteric modulators and silent allosteric "antagonists" outlined in [247, 248]. Negative allosteric modulation of opioid receptors has been previously suggested [953], whether all compounds are acting at a similar site remains to be established.

**Further reading on Opioid receptors**

Butelman ER et al. (2012) \( \kappa \)-opioid receptor/dynorphin system: genetic and pharmacotherapeutic implications for addiction. Trends Neurosci. 35: 587-96 [PMID:22709632]

Cox BM et al. (2015) Challenges for opioid receptor nomenclature: IUPHAR Review 9. Br. J. Pharmacol. 172: 317-23 [PMID:24528283]

Pradhan AA et al. (2011) The delta opioid receptor: an evolving target for the treatment of brain disorders. Trends Pharmacol. Sci. 32: 581-90 [PMID:21925742]

Williams JT et al. (2013) Regulation of \( \mu \)-opioid receptors: desensitization, phosphorylation, internalization, and tolerance. Pharmacol. Rev. 65: 223-54 [PMID:23321159]

**Orexin receptors**

G protein-coupled receptors → Orexin receptors

**Overview:** Orexin receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Orexin receptors [557]) 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 [1703].

| Nomenclature | OX1 receptor | OX2 receptor |
|--------------|--------------|--------------|
| HGNC, UniProt | HCRT1, O43613 | HCRT2, O43614 |
| Potency order of endogenous ligands | orexin-A (HCRT, O43612) > orexin-B (HCRT, O43612) | orexin-A (HCRT, O43612) = orexin-B (HCRT, O43612) |
| Selective agonists | – | [Ala11, D-Leu15]orexin-B [66, 1612] |
| Selective antagonists | suvorexant (pK\(_\text{d} \) 9.3) [391], SB-649868 (pK\(_\text{d} \) 9.1) [442], SB-674042 (pK\(_\text{d} \) 8.7–9.1) [1098, 1253, 1255], filorexant (pK\(_\text{d} \) 8.6) [2124], almorexant (pIC\(_{50} \) 7.9) [221], SB-408124 (pK\(_\text{d} \) 7.2–7.6) [1098, 1253], SB-334867 (pK\(_\text{d} \) 7.4–7.5) [1253, 1591] | filorexant (pK\(_\text{d} \) 9.5) [2124], suvorexant (pK\(_\text{d} \) 9.5) [391], EMPA (pK\(_\text{d} \) 9) [1252], SB-649868 (pK\(_\text{d} \) 8.9) [442], JNJ-10397049 (pK\(_\text{d} \) 8.6–8.8) [1300], almorexant (pIC\(_{50} \) 8.1) [221], TCS-OX2-29 (pK\(_\text{d} \) 7.4) [798] |
| Labelled ligands | – | [\(^{3}\text{H}\)]almorexant (Selective Antagonist) (pK\(_\text{d} \) 8.9–9.8) [1253, 1255], [\(^{3}\text{H}\)]Cp-1 (Selective Antagonist) (pK\(_\text{d} \) 9.2–9.4) [1253], [\(^{3}\text{H}\)]EMPA (Selective Antagonist) (pK\(_\text{d} \) 8.6–9) [1252, 1255], [\(^{125}\text{I}\)]orexin-A (Agonist) [1066, 1611, 1703] |

**Comments:** The primary coupling of orexin receptors to G\(_{q/11}\) proteins is rather speculative and based on the strong activation of phospholipase C, though recent studies in recombinant CHO cells also stress the importance of G\(_{q/11}\) [1065]. Coupling of both receptors to G\(_{10}\) and G\(_{i}\) has also been reported [951, 1068, 1146, 1629] for most cellular responses observed, the G protein pathway is unknown. The potency order of endogenous ligands may depend on the cellular signal transduction machinery. Most of the OX2 receptor selective antagonists listed are weakly selective (±10-fold), or selectivity may be less than 100-fold or not unequivocally determined. [Ala\(_{11}\), D-Leu\(_{15}\)]orexin-B may show poor OX2 receptor selectivity [1612].

Orexin receptors have been reported to be able to form complexes with each other and some other GPCRs as well as CRF receptors [1067, 1426], which might affect the signaling and

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
pharmacology. Recently a promising synthetic orexin receptor ligand (compound 26) has been reported but not thoroughly characterized [1412]. Loss-of-function mutations in the gene encoding the OX2 receptor underlie canine hereditary narcolepsy [1177]. Antagonists of the orexin receptors are the focus of major drug discovery effort for their potential to treat insomnia and other disorders of wakefulness [1668], while agonists would likely be useful in human narcolepsy.

**Further reading on Orexin receptors**

Baimel C et al. (2015) Orexin/hypocretin role in reward: implications for opioid and other addictions. Br. J. Pharmacol. 172: 334-48 [PMID:24641197]

Kukkonen JP. (2013) Physiology of the orexinergic/hypocretinergic system: a revisit in 2012. Am. J. Physiol., Cell Physiol. 304: C2-32 [PMID:23034387]

Li SB et al. (2016) Hypocretins, Neural Systems, Physiology, and Psychiatric Disorders. Curr Psychiatry Rep 18: 7 [PMID:26733323]

Mahler SV et al. (2014) Motivational activation: a unifying hypothesis of orexin/hypocretin function. Nat. Neurosci. 17: 1298-303 [PMID:25254979]

---

**Oxoglutarate receptor**

*G protein-coupled receptors → Oxoglutarate receptor*

**Overview:** Nomenclature as recommended by NC-IUPHAR [414].

| Nomenclature | oxoglutarate receptor |
|--------------|-----------------------|
| HGNC, UniProt | OXGR1, Q96P68         |
| Endogenous agonists | α-ketoglutaric acid [762, 1854] |

---

**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, ADP, 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 [251, 514, 878, 2044, 2089].

Clinically used drugs acting on these receptors include the dinucleoside polyphosphate diquafosol, agonist of the P2Y2 receptor subtype, approved in Japan for the management of dry eye disease [1101], and the P2Y12 receptor antagonists prasugrel, ticagrelor and cangrelor, all approved as antiplatelet drugs [273, 1602].

---

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
| Nomenclature                  | P2Y<sub>1</sub> receptor | P2Y<sub>2</sub> receptor | P2Y<sub>4</sub> receptor | P2Y<sub>6</sub> receptor |
|------------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| HGNC, UniProt                | P2RY1, P47900            | P2RY2, P41231            | P2RY4, P51582            | P2RY6, Q15077            |
| Potency order of endogenous ligands | ADP > ATP              | uridine triphosphate > ATP [1112] | uridine triphosphate > ATP (at rat recombinant receptors, UTP = ATP) | uridine diphosphate >> uridine triphosphate > ADP |
| Endogenous agonists          | –                        | uridine triphosphate [989, 1112] | –                        | –                        |
| Agonists                     | ADPβS [1921], 2MeSADP [1729, 2054] | –                        | –                        | –                        |
| Sub/family-selective agonists | –                        | diquafosol [1554], denufosol [1113, 1554, 2181], UTPβS [1112] | diquafosol [240], denufosol [2181], UTPβS [1113] | –                        |
| Selective agonists           | MRS2365 [329], 2-CI-ADP(α-BH₃) [76] | MRS2698 [874], 2-thioUTP [498], PSB1114 (EC₅₀ value determined using an IP₃ functional assay) [498, 499, 873] | MRS4062 [1276], MRS2927 [1276], (N)methanocarba-UTP [989] | Rp-5-OMe-UDPαB [644, 702], MRS2957 [1275], MRS2693 [146] |
| Antagonists                  | suramin (p𝐾ᵢ 5.3) [2054], PPADS (p𝐾ᵢ 5.2) [2054] | –                        | ATP (p𝐾ᵢ 6.2) [970] | –                        |
| Sub/family-selective antagonists | –                      | reactive blue-2 (pIC₅₀ 6) [892], suramin (pIC₅₀ 4.3) [892, 1729] | PPADS (pEC₅₀ 2–5) [881], reactive blue-2 (pIC₅₀ 4.7) [171] – Rat | reactive blue-2 (p𝐾ᵢ 6) [2045], PPADS (p𝐾ᵢ 4) [2045], suramin (p𝐾ᵢ 4) [2045] |
| Selective antagonists         | MRS2500 (p𝐾ᵢ 8.8–9.1) [286, 988], MRS2279 (p𝐾ᵢ 7.9) [2054], MRS2179 (p𝐾ᵢ 7–7.1) [203, 2054] | AR-C118925XX (pIC₅₀ ~6) [968], AR-C126313 (pEC₅₀ 6) [874], PSB-416 (pIC₅₀ 4.7) [792] | ATP (p𝐾ᵢ 6.2) [970] | MRS2578 (pIC₅₀ 7.4) [1257], MRS2567 (pIC₅₀ 6.9) [1257] |
| Allosteric modulators         | 2,2'-pyridylisatogen tosylate (Negative) (pIC₅₀ 7.8) [601] | –                        | –                        | –                        |
| Selective allosteric modulators | BMS compound 16 (Negative) (p𝐾ᵢ 6.9) [2206] | –                        | –                        | –                        |
| Labelled ligands              | [³H]MRS2279 (Antagonist) (p𝐾ᵢ 8.1) [2054], [³H]2MeSADP (Agonist) [1921], [³⁵S]ADPβS (Agonist) | –                        | –                        | MRS4162 (Selective Antagonist) (pEC₅₀ 7.6) [897] |

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)
Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
**Nomenclature**

|     | P2Y$_{11}$ receptor | P2Y$_{12}$ receptor | P2Y$_{13}$ receptor | P2Y$_{14}$ receptor |
|-----|---------------------|---------------------|---------------------|---------------------|
| HGNC, UniProt | P2RY11, Q96C91 | P2RY12, Q9H244 | P2RY13, Q9BPV8 | P2RY14, Q15391 |

**Potency order of endogenous ligands**

- ATP
- ADP

**Sub/family-selective agonists**

- ATP
- ADP

**Selective agonists**

- AR-C67085 [93, 372], ATP$_5$ [372]
- PSB-0739 (pK$_i$ 7.6) [97]
- AZD1283 (pK$_i$ 8) [79, 2207], ARL66096 (pK$_i$ 7.8) [2203]

**Antagonists**

- Suramin (pIC$_{50}$ 4.8–6) [372], reactive blue-2 (pIC$_{50}$ 5) [372]
- Cangrelor (pIC$_{50}$ 9.4) [882], Ap4A (pIC$_{50}$ 6) [1271], 2MeSAMP (pIC$_{50}$ 5.4) [1921]
- Cangrelor (pIC$_{50}$ 8.3) [1271], Ap4A (pIC$_{50}$ 6.7) [1271], 2MeSAMP (pIC$_{50}$ 5.6) [1271]

**Sub/family-selective antagonists**

- Suramin (pIC$_{50}$ 4.8–6) [372], reactive blue-2 (pIC$_{50}$ 5) [372]
- Cangrelor (pIC$_{50}$ 9.4) [882], Ap4A (pIC$_{50}$ 6) [1271], 2MeSAMP (pIC$_{50}$ 5.4) [1921]
- Cangrelor (pIC$_{50}$ 8.3) [1271], Ap4A (pIC$_{50}$ 6.7) [1271], 2MeSAMP (pIC$_{50}$ 5.6) [1271]

**Selective antagonists**

- Suramin (pIC$_{50}$ 4.8–6) [372], reactive blue-2 (pIC$_{50}$ 5) [372]
- Cangrelor (pIC$_{50}$ 9.4) [882], Ap4A (pIC$_{50}$ 6) [1271], 2MeSAMP (pIC$_{50}$ 5.4) [1921]
- Cangrelor (pIC$_{50}$ 8.3) [1271], Ap4A (pIC$_{50}$ 6.7) [1271], 2MeSAMP (pIC$_{50}$ 5.6) [1271]

**Labelled ligands**

- [3H]2MeSADP (Agonist) [1921], [3H]PSB-0413 (Antagonist) (pK$_d$ 8.3–8.5) [497, 1497]
- [33P]2MeSADP (Agonist) [1271]
- [33P]2MeSADP (Agonist) [1271]
- [33P]2MeSADP (Agonist) [1271]

**Comments**

A series of 4-alkyloxyimino derivatives of uridine-5'-triphosphate which could be useful for derivatization as fluorescent P2Y$_{2/4/6}$ receptor probes has been recently synthesized [897]. Single nucleotide polymorphisms of the P2Y$_{1}$ gene are associated with different platelet reactivity to ADP ADP [784]. Three frequent nonsynonymous P2Y$_{2}$ receptor polymorphisms have been identified, one of which was significantly more common in cystic fibrosis patients. This polymorphism is linked to increases in Ca$^{2+}$ influx in transfected cells, and might therefore play a role in disease development [263]. Although uridine triphosphate (UTP) was also shown to be a biased agonist at P2Y$_{11}$, this is still under debate [1388, 2104]. A group of single nucleotide polymorphisms in the P2Y$_{12}$ gene, forming the so called P2Y$_{12}$ H2 haplotype, has been associated with increased platelet responsiveness to ADP, increased risk of peripheral arterial disease and with coronary artery disease [291]. The platelet-type bleeding disorder due to P2Y$_{12}$ receptor defects is an autosomal recessive condition characterized by mild to moderate mucocutaneous bleeding and excessive bleeding after surgery or trauma. The defect is due to the inability of ADP to induce platelet aggregation [287]. The P2Y$_{13}$ receptor Met-158-Thr polymorphism, which is in linkage disequilibrium with the P2Y$_{12}$ locus, is not associated with acute myocardial infarction, diabetes mellitus or related risk factors [44]. The P2Y$_{14}$ receptor was previously considered to exclusively bind sugar nucleotides such as UDP-glucose and UDP-galactose [299]. However, more recent evidence with several cell lines has demonstrated that uridine diphosphate (UDP) is 5-fold more potent than UDP-glucose [281]. UDP was also shown to competitively antagonise the UDP-glucose response at the human recombinant P2Y$_{14}$ receptor [578].

**Further reading on P2Y receptors**

- Abbracchio MP et al. (2006) International Union of Pharmacology LVIII: update on the P2Y G protein-coupled nucleotide receptors: from molecular mechanisms and pathophysiology to therapy. *Pharmacol. Rev.* **58**: 281-341 [PMID:16968944]
- Jacobson KA et al. (2015) Nucleotides Acting at P2Y Receptors: Connecting Structure and Function. *Mol. Pharmacol.* **88**: 220-30 [PMID:25837834]
- von Kügelgen I et al. (2016) Pharmacology and structure of P2Y receptors. *Neuropharmacology* **104**: 50-61 [PMID:26519900]
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 [606]) 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 (PTHLH, P12272) (141 amino-acids) and related peptides (PTH-(1-34), PTHrP-(1-36) (PTHLH, 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 |
| Potency order of endogenous ligands | PTH (PTH, P01270) = PTHrP (PTHLH, P12272) | TIP39 (PTH2, Q96A98), PTH (PTH, P01270) ≫ PTHrP (PTHLH, P12272) |
| Agonists             | teriparatide [604] | TIP39 (PTH2, Q96A98) [661, 804] |
| Selective agonists   | PTHrP-(1-34) (human) [605] – Rat | – |

Comments: The parathyroid hormone type 1 receptor (PTHR) is the canonical GPCR for PTH and PTHrP. It is coupled to Gs and Gq and regulates the development of bone, heart, mammary glands and other tissues in response to PTHrP, and blood concentrations of calcium and phosphate ions, as well as vitamin D, in response to PTH. Another important action of the PTH/PTHR system is to stimulate bone formation when the hormone is intermittently administered (daily injection). 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 [925].

Further reading on Parathyroid hormone receptors
Cheloha RW et al. (2015) PTH receptor-1 signalling-mechanistic insights and therapeutic prospects. Nat Rev Endocrinol [PMID:26303600]
Gardella TJ et al. (2015) International Union of Basic and Clinical Pharmacology. XCIII. The Parathyroid Hormone Receptors-Family B G Protein-Coupled Receptors. Pharmacol. Rev. 67: 310-37 [PMID:25713287]

Vilardaga JP et al. (2014) Endosomal generation of cAMP in GPCR signaling. Nat. Chem. Biol. 10: 700-6 [PMID:25271346]

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 [557]) is activated by PAF and other suggested endogenous ligands are oxidized phosphatidylycholine [1265] and lysophosphatidylycholine [1492]. It may also be activated by bacterial lipopolysaccharide [1417].

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature: PAF receptor
HGNC, UniProt: PTAFR, P25105
Selective agonists: methylcarbamyl PAF
Selective antagonists: foropafant (pKᵢ 10.3) [774], ABT-491 (pKᵢ 9.2) [30], CV-6209 (pIC₅₀ 8.1–8.3) [652, 1416], L659989 (pKᵢ 7.8) [851], apafant (pKᵢ 5.2–7.5) [1529, 1904]
Labelled ligands: [³H]PAF (Agonist) [585, 1416]

Comments: Note that a previously recommended radioligand ([³H]apafant; Kᵢ 44.6 nM) is currently unavailable.

Further reading on Platelet-activating factor receptor
Foord SM et al. (2005) International Union of Pharmacology. XLVI. G protein-coupled receptor list. Pharmacol Rev 57: 279-288 [PMID:15914470]
Ishii S et al. (2000) Platelet-activating factor (PAF) receptor and genetically engineered PAF receptor mutant mice. Prog. Lipid Res. 39: 41-82 [PMID:10729607]
Prescott SM et al. (2000) Platelet-activating factor and related lipid mediators. Annu. Rev. Biochem. 69: 419-45 [PMID:10966465]

Prokineticin receptors
G protein-coupled receptors → Prokineticin receptors

Overview: Prokineticin receptors, PKR₁ and PKR₂ (provisional nomenclature as recommended by NC-IUPHAR [557]) 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, [1749]) is a potent, non-selective agonist at prokineticin receptors [1279], while Bv8, an orthologue of PROK₂ from amphibians (Bombina sp., [1357]), is equipotent at recombinant PKR₁ and PKR₂ [1435], and has high potency in macrophage chemotaxis assays, which are lost in PKR₁-null mice.

Nomenclature: PKR₁, Q8TCW9
HGNC, UniProt: PROKR1
Potency order of endogenous ligands: prokineticin-2 (PROK₂, Q9HC23) > prokineticin-1 (PROK₁, Q9HC23) > prokineticin-2 β (PROK₂) [1175, 1279, 1843]
Endogenous agonists: prokineticin-2 (PROK₂, Q9HC23) [316, 1279], prokineticin-1 (PROK₁, Q9HC23) [316, 1279], prokineticin-2 β (PROK₂) [316]
Agonists: MIT₁ [1279]

Nomenclature: PKR₂
HGNC, UniProt: PROKR2, Q8NFJ6
Potency order of endogenous ligands: prokineticin-2 (PROK₂, Q9HC23) > prokineticin-1 (PROK₁, Q9HC23) > prokineticin-2 β (PROK₂) [1175, 1279, 1843]
Endogenous agonists: prokineticin-2 (PROK₂, Q9HC23) [316, 1279], prokineticin-1 (PROK₁, Q9HC23) [316, 1279], prokineticin-2 β (PROK₂) [316]
Agonists: MIT₁ [1279]

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature PKR1
Selective agonists IS20 [612], IS1 [612]
Labelled ligands [125I]BH-MIT1 (Agonist) [1279]

Comments: Genetic mutations in PROKR1 are associated with Hirschsprung’s disease [1688], while genetic mutations in PROKR2 are associated with hypogonadotropic hypogonadism with anosmia [455], hypopituitarism with pituitary stalk interruption [1649] and Hirschsprung’s disease [1688].

Further reading on Prokineticin receptors

Boulberdaa M et al. (2011) Prokineticin receptor 1 (PKR1) signalling in cardiovascular and kidney functions. Cardiovasc. Res. 92: 191-8 [PMID:21856786]
Negri L et al. (2012) Bv8/PK2 and prokineticin receptors: a druggable pronociceptive system. Curr Opin Pharmacol 12: 62-6 [PMID:22136937]
Ngan ES et al. (2008) Prokineticin-signaling pathway. Int. J. Biochem. Cell Biol. 40: 1679-84 [PMID:18440852]

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 [794] derived from a FMRFamide-related peptide precursor (NPVF, Q9HCQ7), which is cleaved to generate neuropeptide SF (NPFF, Q9HCQ7), neuropeptide RFRP-1 (NPVF, Q9HCQ7), neuropeptide RFRP-2 (NPVF, Q9HCQ7) and neuropeptide RFRP-3 (NPVF, Q9HCQ7) (neuropeptide NPFF).

Nomenclature PrRP receptor
HGNC, UniProt PRLHR, P49683
Potency order of endogenous ligands PrRP-20 (PRLH, P81277) = PrRP-31 (PRLH, P81277) [1099]
Endogenous agonists PrRP-20 (PRLH, P81277) [509, 1099], PrRP-31 (PRLH, P81277) [509, 1099]
Endogenous antagonists neuropeptide Y (NPY, P01303) (pK_i 5.4) [1087]
Labelled ligands [125I]PrRP-20 (human) (Agonist) [1099], [125I]PrRP31 (Agonist) [501]

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

**G protein-coupled receptors → Prostanoid receptors**

**Overview:** Prostanoid receptors (nomenclature as agreed by the NC-IUPHAR Subcommittee on Prostanoid Receptors [2132]) are activated by the endogenous ligands prostaglandins PGD$_2$, PGE$_1$, PGE$_2$, 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 |
|-----------------------|-----------------|-----------------|
| HGNC, UniProt         | PTGDR, Q13258   | PTGDR2, Q9Y5Y4  |
| Potency order of endogenous ligands | PGD$_2$ > PGE$_1$ > PGE$_2$ > PGF$_{2\alpha}$ > PGI$_2$, thromboxane A$_2$ | PGD$_2$ > PGF$_{2\alpha}$, PGE$_2$ > PGI$_2$, thromboxane A$_2$ |
| Selective agonists    | BW 245C [173, 2133, 2134], L-644,698 [2133, 2134] | 15(8)-TS-methyl-PGD$_2$ [747, 1366, 1889] |
| Antagonists           | –               | fevipiprant (pK$_d$ 9) [1908, 1909], ramatroban (pK$_i$ 7.4) [1889] |
| Selective antagonists | laropiprant (pK$_i$ 10.1) [1882], BWA868C (pK$_i$ 8.6–9.3) [173, 640, 2133], ONO-AE3-237 (pK$_i$ 7.7) [796, 1974, 1976] | CAY 10471 (pIC$_{50}$ 8.9) [1684, 2003] |
| Labelled ligands      | [³H]PGD$_2$ (Agonist) [2119, 2133] | [³H]PGD$_2$ (Agonist) [1280, 1790] |

| Nomenclature          | EP$_1$ receptor | EP$_2$ receptor | EP$_3$ receptor | EP$_4$ receptor |
|-----------------------|-----------------|-----------------|-----------------|-----------------|
| HGNC, UniProt         | PTGER1, P34995  | PTGER2, P43116  | PTGER3, P43115  | PTGER4, P35408  |
| Potency order of endogenous ligands | PGE$_2$ > PGE$_1$ > PGF$_{2\alpha}$, PGI$_2$ > PGD$_2$, thromboxane A$_2$ | PGE$_2$ = PGE$_1$ > PGF$_{2\alpha}$, PGI$_2$ > PGD$_2$, thromboxane A$_2$ | PGE$_2$, PGE$_1$ > PGF$_{2\alpha}$, PGI$_2$ > PGD$_2$, thromboxane A$_2$ | PGE$_2$ = PGE$_1$ > PGF$_{2\alpha}$, PGI$_2$ > PGD$_2$, thromboxane A$_2$ |
| Endogenous agonists   | –               | PGE$_2$ [7, 1871, 2119] | PGE$_2$ (EP$_3$-III isoform) [7] | –               |
| Agonists              | 17-phenyl-ω-trinor-PGE$_2$ [1783] | PGE$_1$ [111] | misoprostol (methyl ester) (EP$_3$-III isoform) [7] | –               |
| Selective agonists    | ONO-DI-004 [1899] – Mouse | ONO-AE1-259 [1899] – Mouse, butaproteren (free acid form) [7, 1871] | sulprostone (EP$_3$-III isoform) [7], ONO-AE-248 [562, 1206] | –               |
| Antagonists           | –               | –               | –               | EP$_4$ (pK$_i$ 7.6–8.5) [1229, 2195] |

Searchable database: [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

Full Contents of ConciseGuide: [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
Nomenclature | EP<sub>1</sub> receptor | EP<sub>2</sub> receptor | EP<sub>3</sub> receptor | EP<sub>4</sub> receptor
--- | --- | --- | --- | ---
Selective antagonists | ONO-8711 (pK<sub>i</sub> 9.2) [2079], SC-51322 (pK<sub>i</sub> 7.9) [7] | PF-04418948 (PF-04418948 has weaker affinity at the EP2-receptor in guinea-pigs) (pK<sub>i</sub> 8.3) [14, 157], TG6-129 (pK<sub>i</sub> 8.1) [598] | L-826266 (EP<sub>1</sub>-III isofrm (pK<sub>i</sub>=8.04 in the presence of HSA)) (pK<sub>i</sub> 9.1) [933], ONO-AE3-240 (pC50 8.8) [38] | ONO-AE3-208 (pK<sub>i</sub> 8.5), GW 627368 (pK<sub>i</sub> 7–7.1) [2119, 2120]
Labelled ligands | [³H]PGE<sub>2</sub> (Agonist) [7, 1783, 2119] | [³H]PGE<sub>2</sub> (Agonist) [7, 2119] | [³H]PGE<sub>2</sub> (Agonist) [7, 2119] | [³H]PGE<sub>2</sub> (Agonist) [7, 420, 2107, 2119]

**Comments**: Whilst cicaprost is selective for IP receptors, it does exhibit moderate agonist potency at EP<sub>4</sub> receptors [7]. Apart from IP receptors, iloprost also binds to EP<sub>1</sub> receptors. The IP receptor agonist treprostinil binds also to human EP<sub>2</sub> and DP<sub>1</sub> receptors with high affinity (pK<sub>i</sub> 8.44 and 8.36, respectively) [2107]. The EP<sub>1</sub> agonist 17-phenyl-ω-trinor-PGE<sub>2</sub> also shows agonist activity at EP<sub>3</sub> receptors. Butaprost and SC46275 may require de-esterification within tissues to attain full agonist potency. There is evidence for subtypes of FP [1171] and TP receptors [1050, 1637]. mRNA for the EP<sub>3</sub> receptor undergoes alternative splicing to produce variants which can interfere with signalling [1509] or generate complex patterns of G-protein (G<sub>i10</sub>, G<sub>q11</sub>, G<sub>q</sub> and G<sub>12,13</sub>) coupling (e.g. [1042, 1433]). The number of EP<sub>2</sub> receptor (protein) variants are variable depending on species, with five in human, three in rat and three in mouse. Putative receptor(s) for prostamide F (which as yet lack molecular correlates) and which preferentially recognize PGF<sub>2</sub>-ethanolamide and its analogues (e.g. Bimatoprost) have been identified, together with moderate-potency antagonists (e.g. AGN 211334) [2131]. The free acid form of AL-12182, AL12180, used in in vitro studies, has an EC<sub>50</sub> 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 [1784]. References given alongside the TP receptor agonists I-BOP [1295] and STA<sub>2</sub> [63] use human platelets as the source of TP receptors for competition radioligand binding assays to determine the indicated activity values. Pharmacological evidence for a second IP receptor, denoted IP<sub>2</sub>, in the central nervous system [1924, 2082] and in the BEAS-2B human airway epithelial cell line [2122] is available. This receptor is selectively activated by 15R-TIC and 15-deoxy-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 an IP<sub>2</sub> subtype is currently lacking.
Further reading on Prostanoid receptors

Woodward DF et al. (2011) International union of basic and clinical pharmacology. LXXXIII: classification of prostanoid receptors, updating 15 years of progress. Pharmacol. Rev. 63: 471-538 [PMID:21752876]

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 [809]) 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-NH$_2$, SLIGKV-NH$_2$, TFRGAP-NH$_2$ and GYPGQV-NH$_2$, 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 exodomain of the receptor without inducing activation of G-$\alpha_q$ coupled calcium signalling, thereby preventing activation by activating proteinases but not by agonist peptides. Neutrophil elastase (NE) cleavage of PAR1 and PAR2 can however activate MAP kinase signalling by exposing a TL that is different from the one revealed by trypsin [1624]. PAR2 activation by NE regulates inflammation and pain responses [1397, 2217] and triggers mucin secretion from airway epithelial cells [2220].

| 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) [73] | Trypsin, tryptase, TF/VIIa, Xa | thrombin (F2, P00734) | trypsin, cathepsin G (CTSG, P08311) |
| Agonists | F16357 | – | – | – |
| Selective agonists | TFLLR-NH$_2$ [355] | AC264613 [1767], AY77 [2178], AC-55541 [1767], CB110 [104], 2-furoyl-LIGRL-amide [1305], SLIGKV-NH$_2$ [1134], SLIGRL-NH$_2$ [1134] | – | – |
| Selective antagonists | vorapaxar (pK$_i$ 8.1) [295], atopaxar (pIC$_{50}$ 7.7) [1024], RWJ-56110 (pIC$_{50}$ 6.4) [49] | G888 (pIC$_{50}$ 5.7) [1886], P2pal18s [1776] | – | YD-3 (pIC$_{50}$ 6.9) [2091], ML354 (pIC$_{50}$ 6.8) [2091] |
| Labelled ligands | $[^{3}H]$haTRAP (Agonist) [17] | 2-furoyl-LIGRL[N-(Alexa Fluor 594)-O]-NH$_2$ (Agonist) [810], 2-furoyl-LIGRL[N-$[^{3}H]$propionyl]-O-NH$_2$ (Agonist) [810], $[^{3}H]$2-furoyl-LIGRL-NH$_2$ (Selective Agonist) [946], trans-cinnamoyl-LIGRLo [N-$[^{3}H]$propionyl]-NH$_2$ (Agonist) [28] | – | – |
| Comments | TFLLR-NH$_2$ is selective relative to the PAR$_2$ receptor [159, 958]. | 2-Furoyl-LIGRL-O-NH$_2$ activity was measured via calcium mobilisation in HEK 293 cells which constitutively coexpress human PAR$_1$ and PAR$_2$. | – | – |

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Comments: 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 (TMRSS15, P98073) on pancreatic-derived trypsinogen (PRSS1, P00777); trypsin, a family of enzymes (α/β TPSAB1, Q15661; γ1 TPSG1, Q8NR82; δ1 TPXD1, Q9BZ73) 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).

Further reading on Proteinase-activated receptors

Adams MN et al. (2011) Structure, function and pathophysiology of protease activated receptors. Pharmacol. Ther. 130: 248-82 [PMID:21277892]
Canto I et al. (2012) Allosteric modulation of protease-activated receptor signaling. Mini Rev Med Chem 12: 804-11 [PMID:22681248]
García PS et al. (2010) The role of thrombin and protease-activated receptors in pain mechanisms. Thromb. Haemost. 103: 1145-51 [PMID:20431855]
Hollenberg MD et al. (2002) International Union of Pharmacology. XXVIII. Proteinase-activated receptors. Pharmacol. Rev. 54: 203-17 [PMID:12037136]
Ramachandran R et al. (2012) Targeting proteinase-activated receptors: therapeutic potential and challenges. Nat Rev Drug Discov 11: 69-86 [PMID:22212680]
Soh UJ et al. (2010) Signal transduction by protease-activated receptors. Br. J. Pharmacol. 160: 191-203 [PMID:20423334]

QRFP receptor

Overview: The human gene encoding the QRFP receptor (QRFR, 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 [1131]. 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, QRFR (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 Gal1 (35%) receptors.

Nomenclature
HGNC, UniProt
QRFP, Q96P65
Endogenous agonists
QRFP43 (QRFP, P83859) [311, 587, 1923] – Rat, QRFP26 (QRFF) [311, 910]
Agonists
LV-2172 [1448]
Selective antagonists
compound 25e (pIC50 7.3) [628, 629]
Labelled ligands
[^125I]QRFP43 (human) (Agonist) [587, 1063, 1923]

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

Further reading on QRFP receptor

Fukusumi S et al. (2006) Recent advances in mammalian RFamide peptides: the discovery and functional analyses of PrRP, RFRPs and QRFP. Peptides 27: 1073-86 [PMID:16500002]
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 [112, 713]) may be divided into two pairs, RXFP1/2 and RXFP3/4. Endogenous agonists at these receptors are a number of heterodimeric peptide hormones structurally related to insulin: relaxin-1 (RLN1, P04808), relaxin (RLN2, P04090), relaxin-3 (RLN3, Q8WXF3) (also known as INSL7), insulin-like peptide 3 (INSL3 (INSL3, P51460)) and INSL5 (INSL5, Q9YSQ6).

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 [1755] and porcine relaxin may have a higher efficacy than human relaxin (RLN2, P04090) [714]. Relaxin-3 (RLN3, Q8WXF3) has differential affinity for RXFP2 between species; mouse and rat RXFP2 have a higher affinity for relaxin-3 (RLN3, Q8WXF3) [1754]. At least two binding sites have been identified on RXFP1 and RXFP2: 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 [714, 1885]. The unique N-terminal LDLa module of RXFP1 and RXFP2 is essential for receptor signalling [1756].

Nomenclature RXFP1 RXFP2 RXFP3 RXFP4 HGNC, UniProt RXFP1, Q9HBX9 RXFP2, Q8WXD0 RXFP3, Q9NSD7 RXFP4, Q8TDU9

Potency order of endogenous ligands relaxin (RLN2, P04090) = relaxin-1 (RLN1, P04808) > relaxin-3 (RLN3, Q8WXF3) [1885]

Endogenous antagonists – –

Antagonists B-R13/17K H2 relaxin (pEC\textsubscript{50} 5.7–6.7) [827, 1446]

Selective antagonists –

Allosteric modulators ML290 (Agonist) (pEC\textsubscript{50} 7) [2146, 2149]

Labelled ligands \[\textsuperscript{33}P\text{relaxin (human)} (Agonist) [714, 1885], europium-labelled relaxin (Agonist) [1778], \[\textsuperscript{125}I\text{relaxin (human)} (Agonist) [1395], \[\textsuperscript{33}P\text{relaxin (human)} (Agonist) [714, 1885]

\[\textsuperscript{125}I\text{relaxin-3 (human)} (Agonist) [1186], \[\textsuperscript{125}I\text{relaxin-3-B/INSL5 A chimera (Agonist) [1184], europium-labelled relaxin-3-B/INSL5 A chimera (Agonist) [749], europium-labelled INSLS (pK\textsubscript{d} 8.3) [749], europium-labelled INSL5 (pK\textsubscript{d} 8.3) [749], europium-labelled relaxin-3-B/INSL5 A chimera (Agonist) [749], europium-labelled insulin-like peptide 3 (INSL3 (INSL3, P51460)) and INSL5 (INSL5, Q9YSQ6).

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 [1755] and porcine relaxin may have a higher efficacy than human relaxin (RLN2, P04090) [714]. Relaxin-3 (RLN3, Q8WXF3) has differential affinity for RXFP2 between species; mouse and rat RXFP2 have a higher affinity for relaxin-3 (RLN3, Q8WXF3) [1754]. At least two binding sites have been identified on RXFP1 and RXFP2: 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 [714, 1885]. The unique N-terminal LDLa module of RXFP1 and RXFP2 is essential for receptor signalling [1756].

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Comments: Relaxin is the cognate peptide ligand for RXFP1 and is in extended Phase III clinical trials for the treatment of acute heart failure [1322]. Relaxin has vasodilatory, anti-fibrotic, angiogenic, anti-apoptotic and anti-inflammatory effects. Small molecule allosteric agonists such as ML290 have been developed [1787, 2149]. The antifibrotic actions of relaxin are dependent on the angiotensin receptor AT2 [344]. RXFP2 and its cognate ligand INSL3 have a more specialized role with mutations reported in patients with cryptorchidism [538]. cAMP elevation is the major signalling pathway for RXFP1 and RXFP2 [834, 835], but RXFP1 also activates MAP kinases, nitric oxide signalling, tyrosine kinase phosphorylation and relaxin can interact with glucocorticoid receptors [716]. RXFP1 signalling involves lipid rafts, residues in the C-terminus of the receptor and activation of phosphatidylinositol-3-kinase [717] and pre-assembled protein complexes [715]. Receptor expression profiles suggest that RXFP3 is a brain neuropeptide receptor and RXFP4 a gut hormone receptor with the relaxin-3/RXFP3 system modulating feeding [596, 597, 749, 1777, 1830], anxiety [1694, 2204], and reward and motivated goal-directed behaviours [821, 1694, 2055]. Relaxin-3 (RLN3, Q8WXF3) is an agonist at RXFP3 and RXFP4 whereas INSL5 (INSL5, Q9Y5Q6) 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. INSL5 is secreted from enteroendocrine L cells and the INSL5/RXFP4 system controls food intake and glucose homeostasis [685]. RXFP3 and RXFP4 couple to G_{i,o} and inhibit adenyl cyclase [1186, 2014], and also cause Erk1/2 phosphorylation [2014]. RXFP4 also causes phosphorylation of p38MAPK, Akt and S6RP [51]. There is evidence that at RXFP3, relaxin (RLN2, P04090) is a biased ligand compared to the cognate ligand relaxin-3.

Further reading on Relaxin family peptide receptors

Bathgate RA et al. (2013) Relaxin family peptides and their receptors. Physiol. Rev. 93: 405-80 [PMID:23303914]
Du XJ et al. (2010) Cardiovascular effects of relaxin: from basic science to clinical therapy. Nat Rev Cardiol 7: 48-58 [PMID:19935741]
Halls ML et al. (2015) International Union of Basic and Clinical Pharmacology. XCV. Recent advances in the understanding of the pharmacology and biological roles of relaxin family peptide receptors 1-4, the receptors for relaxin family peptides. Pharmacol. Rev. 67: 389-440 [PMID:25761609]
Ivell R et al. (2011) Relaxin family peptides in the male reproductive system—a critical appraisal. Mol. Hum. Reprod. 17: 71-84 [PMID:20952422]
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 (SST1-SST5; nomenclature as agreed by the NC-IUPHAR Subcommittee on Somatostatin Receptors [829]). Activation of these receptors produces a wide range of physiological effects throughout the body including the inhibition of secretion of many hormones. 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 [427].

| 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   | pasireotide   | pasireotide   | pasireotide   | pasireotide   |
|              | [1740]        | [1740]        | [1740]        | [1740]        | [1740]        |
| Selective agonists | L-797,591 [1669], Des-Ala1,2,5-(D-Trp8,IAMP9)SRIF [512] | L-054,522 [2172], BIM 23027 [283], octreotide [238, 1540, 1805, 1806, 1807, 2172] | L-796,778 [1669] | L-803,087 [1669] | BIM 23052 [1325, 1805, 1806, 1807], L-817,818 [1669] |
| Selective antagonists | SRA880 (pKd 8–8.1) [831] | [D-Tyr8]CYN 154806 (pKd 8.1–8.9) [1478] | NVP ACQ090 (pKi 7.9) [832] | – | – |
| Labelled ligands | – | [125]I[Tyr3 SMS 201-995 (Agonist)] [1805, 1806] | [125]I[BIM23027 (Agonist)] [811] – | – | [125]I[Tyr3 SMS 201-995 (Agonist)] [1805, 1806] |

Comments: [125]I[Tyr11-SRIF-14], [125]I[LTT-SRIF-28], [125]I[CGP 23996] and [125]I[Tyr10-CST14] may be used to label somatostatin receptors nonselectively. A number of nonpeptide subtype-selective agonists have been synthesised [1669]. Octreotide and lanreotide are being used in the treatment of SST2-expressing neuroendocrine tumors and pasireotide for SST5-expressing neuroendocrine tumors. A novel peptide somatostatin analogue, veldoreotide (somatoprim), has affinity for SST2, SST4 and SST5 receptors and is a potent inhibitor of GH secretion [1586, 1793].

Further reading on Somatostatin receptors

Colao A et al. (2011) Resistance to somatostatin analogs in acromegaly. Endocr. Rev. 32: 247-71 [PMID:21123741]

Hoyer D et al. (2000) Somatostatin receptors. In The IUPHAR Compendium of Receptor Characterization and Classification, 2nd edn. Edited by Watson SF, Girdlestone D: IUPHAR Media: 354-364

Schulz S et al. (2014) Fine-tuning somatostatin receptor signalling by agonist-selective phosphorylation and dephosphorylation: IUPHAR Review S. Br. J. Pharmacol. 171: 1591-9 [PMID:24328848]

Weckbecker G et al. (2003) Opportunities in somatostatin research: biological, chemical and therapeautic aspects. Nat Rev Drug Discov 2: 999-1017 [PMID:14654798]

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

Full Contents of Concise Guide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Succinate receptor

G protein-coupled receptors → Succinate receptor

**Overview:** Nomenclature as recommended by NC-IUPHAR [414]. The Succinate receptor has been identified as being activated by physiological levels of the Krebs’ cycle intermediate succinate and other dicarboxylic acids such as maleate in 2004. Since its pairing with its endogenous ligand, the receptor has been the focus of intensive research and its role has been evidenced in various (patho)physiological processes such as regulation of renin production, retinal angiogenesis or immune response.

| Nomenclature | succinate receptor |
|--------------|--------------------|
| HGNC, UniProt | SUCNR1, Q9BXA5     |
| Endogenous agonists | succinic acid [762, 1854] |

**Comments:** In humans, there is the possibility of two open-reading frames (ORFs) for SUCNR1, allowing the generation of 330 or 334 amino acid proteins Wittenberger et al.[2127] noted that the 330-AA protein was more likely to be expressed given the Kozak sequence surrounding the second ATG. Some databases report SUCNR1 as being 334-AA long.

**Further reading on Succinate receptor**

Ariza AC et al. (2012) The succinate receptor as a novel therapeutic target for oxidative and metabolic stress-related conditions. *Front Endocrinol (Lausanne)* 3: 22. [PMID:22649411]

de Castro Fonseca M et al. (2016) GPR91: expanding the frontiers of Krebs cycle intermediates. *Cell Commun. Signal* 14:3 [PMID:26739054]

Tachykinin receptors

G protein-coupled receptors → Tachykinin receptors

**Overview:** Tachykinin receptors (provisional nomenclature as recommended by NC-IUPHAR [587]) are activated by the endogenous peptides substance P (*TAC1, P20366*) (SP), neurokinin A (*TAC1, P20366*) (NKA; previously known as substance K), neurokinin α (*neuromedin L1, 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 sequence: Phe-x-Gly-Leu-Met. Marked species differences in in vitro pharmacology exist for all three receptors, in the context of nonpeptide ligands. Antagonists such as aprepitant and fosaprepitant were approved by FDA and EMA, in combination with other antiemetic agents, for the prevention of nausea and vomiting associated with emetogenic cancer chemotherapy.

**Further reading on Tachykinin receptors**

Gilissen J et al. (2016) Insight into SUCNR1 (GPR91) structure and function. *Pharmacol. Ther.* 159: 56-65 [PMID:26808164]

Peti-Peterdi J., (2010) High glucose and renin release: the role of succinate and GPR91. *Kidney Int.* 78 (12): 1214-7. [PMID:20861827]

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
### Nomenclature

| Gene Symbol | Name          |
|-------------|---------------|
| TACR1       | NK₁ receptor  |
| TACR2       | NK₂ receptor  |
| TACR3       | NK₃ receptor  |

### HGNC, UniProt

| Gene Symbol | Gene Name          |
|-------------|--------------------|
| TACR1       | TACR1              |
| TACR2       | TACR2              |
| TACR3       | TACR3              |

### Potency order of endogenous ligands

| Ligand          | Potency Order     |
|-----------------|-------------------|
| substance P     | > neurokinin A    |
| neurokinin A    | > neurokinin B    |
| neurokinin B    | > substance P     |

### Agonists

- Substance P-OMe
- Selective agonists:
  - [Sar⁹, Met(O₂)¹¹]SP
  - Septide
  - [Pro⁹]SP
  - [Lys⁵, Me-Leu⁹, Nle¹⁰]NKA-(4-10)
  - GR64349
  - [β-Ala⁸]neurokinin A-(4-10)
  - [Phe(Me)⁷]neurokinin B
  - Senktide

### Selective antagonists

| Antagonist       | pKi/pIC₅₀ |
|------------------|-----------|
| Aprepitant       | 10.1      |
| CP 99994         | 9.8       |
| CP67580          | 7.7       |
| GR94800          | 9.5       |
| saredutant       | 9.4–9.7   |
| NEN10627         | 7.7       |
| [3H]saredutant   | 9.7       |
| [125I]L703,606   | 9.5       |
| [125I]BH-[Sar⁹, Met(O₂)¹¹]SP | 9.7 |
| [3H]GR100679     | 9.2       |
| [125I]NKA        | 9.4–9.7   |
| [3H]osanetant    | 9.3–9.7   |
| [3H]senktide     | 8.4–9.7   |
| [125I]MePhe⁷NKB  | 8.4–9.7   |

### Labelled ligands

- [125I]BH-[Sar⁹, Met(O₂)¹¹]SP (Agonist)
- [125I]NKA (human, mouse, rat)(Agonist)
- [3H]NKA (human, mouse, rat)(Agonist)
- [3H]GR100679 (Antagonist)
- [125I]L703,606 (Antagonist)
- [3H]GR100679 (Antagonist)
- [125I]MePhe⁷NKB

### Comments

The NK₁ receptor has also been described to couple to G proteins other than Gq/11 [1680]. 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 (SwissProtP30098) with structural similarities to the NK₃ receptor was found to respond to NKB when expressed in Xenopus oocytes or Chinese hamster ovary cells [459, 1049].

### Further reading on Tachykinin receptors

- Douglas SD et al. (2011) Neurokinin-1 receptor: functional significance in the immune system in reference to selected infections and inflammation. *Ann. N. Y. Acad. Sci.* 1217: 83-95
- Foord SM et al. (2005) International Union of Pharmacology. XLVI. G protein-coupled receptor list. *Pharmacol Rev* 57: 279-288
- Jones S et al. (2008) The neurokinin 1 receptor: a potential new target for anti-platelet therapy? *Curr Opin Pharmacol* 8: 114-9
- Rance NE et al. (2010) Neurokinin B and the hypothalamic regulation of reproduction. *Brain Res.* 1364: 116-28
- Rojas C et al. (2012) Pharmacological mechanisms of 5-HT₃ and tachykinin NK₁ receptor antagonism to prevent chemotherapy-induced nausea and vomiting. *Eur. J. Pharmacol.* 684: 1-7
- Steinhoff MS et al. (2014) Tachykinins and their receptors: contributions to physiological control and the mechanisms of disease. *Physiol. Rev.* 94: 265-301

**Searchable database:** [http://www.guidetopharmacology.org/index.jsp](http://www.guidetopharmacology.org/index.jsp)

**Full Contents of ConciseGuide:** [http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full](http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full)
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 [557]) 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 [1896]. [3H]TRH (human, mouse, rat) is able to label both TRH1 and TRH2 receptors with Kd values of 13 and 9 nM respectively. Synthesis and biology of ring-modified L-Histidine containing TRH analogues has been reported [1316].

| Nomenclature | TRH1 receptor | TRH2 receptor |
|--------------|---------------|---------------|
| HGNC, UniProt | TRHR, P34981  | -             |
| Antagonists   | diazepam (pKᵢ 5.2) [471] – Rat | -             |
| Selective antagonists | midazolam (pKᵢ 5.5) [471] – Rat, chlordiazepoxide (pKᵢ 4.8) [471] – Rat, chlordiazepoxide (pKᵢ 4.7) [1878] – Mouse | -             |
| Comments      | -             | A class A G protein-coupled receptor: not present in man |

Further reading on Thyrotropin-releasing hormone receptors
Bílek R et al. (2011) TRH-like peptides. Physiol Res 60: 207-15 [PMID:21114375]
Foord SM et al. (2005) International Union of Pharmacology. XLVI. G protein-coupled receptor list. Pharmacol Rev 57: 279-288 [PMID:15914470]
Nillni EA. (2010) Regulation of the hypothalamic thyrotropin releasing hormone (TRH) neuron by neuronal and peripheral inputs. Front Neuroendocrinol 31: 134-56 [PMID:20074684]

Trace amine receptor
G protein-coupled receptors → Trace amine receptor

Overview: Trace amine-associated receptors were discovered from a search for novel S-HT receptors [189], 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 receptor [1244]) has affinity for the endogenous trace amines tyramine, β-phenylethylamine and octopamine in addition to the classical amine dopamine [189]. Emerging evidence suggests that TA1 is a modulator of monoaminergic activity in the brain [2151] with TA1 and dopamine D₂ receptors shown to form constitutive heterodimers when co-expressed [519]. In addition to trace amines, receptors can be activated by amphetamine-like psychostimulants, and endogenous thyronamines.

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature
HGNC, UniProt
TA1 receptor
TAAR1, Q96RJ0

Potency order of endogenous ligands
Agonists
tyramine > β-phenylethylamine > octopamine = dopamine [189]
Antagonists
ROS166017 [1648]
EPPTB (Inverse agonist) (pIC50 5.1) [205]

Labelled ligands
[3H]tyramine (Agonist) [189]

Comments: In addition to TA1, in man there are up to 5 functional TAAR genes (TAAR2,5,6,8,9). See [189] for detailed discussion. The product of the gene TAAR2 (also known as GPR58) appears to respond to β-phenylethylamine > tyramine and to couple through Gs [189]. TAAR5, in some individuals, and TAAR4 are pseudogenes in man, although functional in rodents. The signalling characteristics and pharmacology of TAAR5 (PNR, Putative Neurotransmitter Receptor: TAAR5, O14804), TAAR6 (Trace amine receptor 4, TaR-4: TAAR6, 96RI8), TAAR8 (Trace amine receptor 5, GPR102: TAAR8, O14804), and TAAR9 (trace amine associated receptor 9: TAAR9, 96RI9) are lacking. The thyronamines, endogenous derivatives of thyroid hormone, have affinity for rodent cloned trace amine receptors, including TA1 [1728]. An antagonist EPPTB has recently been described with a pKd of 9.1 at the mouse TA1 but >5.3 for human TA1 [1863].

Further reading on Trace amine receptor
Maguire JJ et al. (2009) International Union of Pharmacology. LXXII. Recommendations for trace amine receptor nomenclature. Pharmacol. Rev. 61:1 - 8 [PMID:19325074]
Pei Y et al. (2016) Trace Amines and the Trace Amine-Associated Receptor 1: Pharmacology, Neurochemistry, and Clinical Implications. Front Neurosci 10: 148 [PMID:27092049]

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 [466, 557, 2032]) 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 [138]. 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 [389, 1195, 1379, 1476]. Human urotensin-II (UTS2, O95399), an 11-amino-acid peptide [389], retains the cyclohexapeptide sequence of goby U-II that is thought to be important in ligand binding [224, 1003]. 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 [388]. A second endogenous ligand for the UT has been discovered in rat [1890]. This is the urotensin II-related peptide (UTS2B, Q765I0), 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, Q765I0) are predicted for the mature mouse and human peptides [472]. UT exhibits relatively high sequence identity with somatostatin, opioid and galanin receptors [2032].

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
### Ut receptor

**Nomenclature**

UT receptor

**HGNC, UniProt**

UTS2R, Q9UKP6

**Endogenous agonists**

urotensin II-related peptide (UTS2R, Q76510) [472, 1243], urotensin-II (UTS2, O95399) [467, 503, 681]

**Selective agonists**

[Pen6]U-(4-11) (human) [681], U-II-(4-11) (human) [681], [3-iodo-Tyr6]U-II-(4-11) (human) [1084], Urolinin [95], FL104 [1139, 1141], AC-7954 [398, 1140]

**Selective antagonists**

JNJ-39319202 (pKi 8.4) [1106], urantide (pKi 8.3) [1536], SB-706375 (pKi 7.2) [445] – Rat, palosuran (pIC50 7.1) [366], SB-436811 (pKi 6.7) [912] – Rat, SB-611812 (pKi 6.6) [1622], S6716 (Inverse agonist) (pIC50 6.4) [554] – Rat, [Cha6]U-II-(4-11) (pKi 6.4) [312] – Rat

**Labelled ligands**

[125I]U-II (human) (Agonist) [42, 198, 312, 1243], [125I]N-biotin-[Ahx0,Bpa3]U-II (human) [454]

**Comments**

In the human vasculature, human urotensin-II (UTS2, O95399) elicits both vasoconstrictor (pD2 9.3-10.1, [1243]) and vasodilator (pIC50 10.3-10.4, [1872]) responses.

### Further reading on Urotensin receptor

Foord SM et al. (2005) International Union of Pharmacology. XLVI. G protein-coupled receptor list. Pharmacol Rev 57: 279-288 [PMID:15914470]

Hunt BD et al. (2010) A rat brain atlas of urotensin-II receptor expression and a review of central urotensin-II effects. Naunyn Schmiedebergs Arch. Pharmacol. 382: 1-31 [PMID:20422157]

Maryanoff BE et al. (2010) Urotensin-II receptor modulators as potential drugs. J. Med. Chem. 53: 2695-708 [PMID:20043680]

Ross B et al. (2010) Role of urotensin II in health and disease. Am. J. Physiol. Regul. Integr. Comp. Physiol. 298: R1156-72 [PMID:20421634]

### 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 [557]) 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).

**Nomenclature**

V1A receptor

AVPR1A, P37288

V1B receptor

AVPR1B, P47901

**Potency order of endogenous ligands**

vasopressin (AVP, P01185) > oxytocin (OXT, P01178)

**Endogenous agonists**

vasopressin (AVP, P01185) [24, 326, 383, 439, 1418, 1571, 1702, 1913, 1914, 1945, 1946, 2162]

**Selective agonists**

F180 [50, 383]

**Antagonists**

conivaptan (pKi 8.2–8.4) [1913, 1914]

**Selective antagonists**

reloclipivaptan (pKi 8.1–9.3) [24, 383, 682, 1571, 1771, 1913, 1945, 1946, 1986]

d(CH2)2[Tyr(Me)2,Arg8]VP (pKi 9)

**Searchable database**: http://www.guidetopharmacology.org/index.jsp

**Full Contents of ConciseGuide**: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Nomenclature

### V₁ₐ receptor

| Labelled ligands | [¹²⁵]OH-LVA (Antagonist) (pKᵩ 10.3–10.4) [334, 383, 1571], [³[H]AVP (human, mouse, rat) (Agonist) [214, 334, 383, 384, 1418, 1571, 1702, 1913, 1914, 1945, 1946, 1986, 2162], [³[H]d(CH₂)₂[Tyr(Me)²]AVP (Antagonist) (pKᵩ 9) |

### V₁₈ receptor

| Labelled ligands | [³[H]AVP (human, mouse, rat) (Agonist) [214, 334, 383, 384, 1418, 1571, 1702, 1913, 1914, 1945, 1946, 1986, 2162] |

### V₂ receptor

| Labelled ligands | [³[H]AVP (human, mouse, rat) (Agonist) [334, 383, 384, 1418, 1702, 1913, 1914, 1946, 2162], [³[H]dDAVP (Agonist) [334, 384, 1946], [³[H]desGly-NH₂[d(CH₂)₂[Tyr(Me)²,Thr4,Orn₈]OT (pKᵩ 8.6) |

| Antagonists | conivaptan (pKᵩ 9.4) [397], tolvaptan (pKᵩ 9.4) [2162], satavaptan (pKᵩ 8.4–9.3) [24, 383, 384, 1770, 1913, 1914, 1986], Lixivaptan (Inverse agonist) (pKᵩ 8.9–9.2) [33, 1771], d(Leu₄)LVP (pKᵩ 6.9–8.4) [1771], mozavaptan (Inverse agonist) (pKᵩ 7.4–8.1) [384, 1771, 1913, 1946, 2162, 2163] |

### OT receptor

| Labelled ligands | [³[H]OT (human, mouse, rat)(Agonist) [334, 583, 895, 998], [¹¹¹In]DOTA-dLVT (pKᵩ 8.3) [333] |

### Comments

The V₂ receptor exhibits marked species differences, such that many ligands (d(Leu₄)VP and [³[H]desGly-NH₂[d(CH₂)₂[Tyr(Me)²,Thr₄,Orn₈]OT) exhibit low affinity at human V₂ receptors [29]. Similarly, [³[H]d(D-Arg⁸)VP is V₂ selective in the rat, not in the human [1702]. The gene encoding the V₂ receptor is polymorphic in man, underlying nephrogenic diabetes insipidus [152]. D([Cha⁴]AVP is selective only for the human and bovine V₁₈ receptors [439], while d([Leu⁴]LVP has high affinity for the rat V₁₈ receptor [1535].

### Further reading on Vasopressin and oxytocin receptors

Bartz JA et al. (2011) Social effects of oxytocin in humans: context and person matter. Trends Cogn. Sci. (Regul. Ed.) 15: 301-9 [PMID:21696997]

Knepper MA. (2012) Systems biology in physiology: the vasopressin signaling network in kidney. Am. J. Physiol., Cell Physiol. 303: C1115-24 [PMID:22932685]

Koshimizu TA et al. (2012) Vasopressin V₁a and V₁b receptors: from molecules to physiological systems. Physiol. Rev. 92: 1813-64 [PMID:23073632]

Manning M et al. (2012) Oxytocin and vasopressin agonists and antagonists as research tools and potential therapeutics. J. Neuroendocrinol. 24: 609-28 [PMID:22375852]

Meyer-Lindenberg A et al. (2011) Oxytocin and vasopressin in the human brain: social neuropeptides for translational medicine. Nat. Rev. Neurosci. 12: 524-38 [PMID:21852800]

Neumann ID et al. (2012) Balance of brain oxytocin and vasopressin: implications for anxiety, depression, and social behaviors. Trends Neurosci. 35: 649-59 [PMID:22974560]

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

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

Vasopressin and oxytocin receptors S116
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 [739, 740]) are activated by the endogenous peptides VIP (P01282), PACAP-38 (ADCYAPI, P18509), PACAP-27 (ADCYAPI, 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 comparable affinity for the PACAP peptides, PACAP-27 (ADCYAPI, P18509) and PACAP-38 (ADCYAPI, P18509), and VIP (VIP, P01282), whereas PACAP-27 (ADCYAPI, P18509) and PACAP-38 (ADCYAPI, 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 (ADCYAPI, P18509), PACAP-27 (ADCYAPI, P18509) and VIP (VIP, P01282) with comparable affinity [411]. PG 99-465 [1374] has been used as a selective VPAC2 receptor antagonist in a number of physiological studies, but has been reported to have significant activity at VPAC1 and PAC1 receptors [446]. 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 [1383]. Two deletion variants of maxadilan, M65 [1994] and Max.d.4 [1384] have been reported to be PAC1 receptor antagonists, but these peptides have not been extensively characterised.

Nomenclature

| HGNC, UniProt | PAC1 receptor | VPAC1 receptor | VPAC2 receptor |
|---------------|---------------|----------------|---------------|
| ADCYAPI1, P41586 | VIPR1, P32241 | VIP (VIP, P01282), PACAP-27 (ADCYAPI, P18509), PACAP-38 (ADCYAPI, P18509) | VIP (VIP, P01282), PACAP-27 (ADCYAPI, P18509), PACAP-38 (ADCYAPI, P18509) |
| Potency order of endogenous ligands | PHI (VIP, P01282) | GHRH (GHRH, P01286), PHI (Pig) | GHRH (GHRH, P01286), PHI (Pig) |
| Selective agonists | PHI (Pig) | PHI (Pig) | PHI (Pig) |
| Selective antagonists | PHI (Pig) | PHI (Pig) | PHI (Pig) |
| Labelled ligands | PHI (Pig) | PHI (Pig) | PHI (Pig) |

Comments: Subtypes of PAC1 receptors have been proposed based on tissue differences in the potencies of PACAP-27 (ADCYAPI, P18509) and PACAP-38 (ADCYAPI, P18509); these might result from differences in G protein coupling and second messenger mechanisms [2018], or from alternative splicing of PAC1 receptor mRNA [1859].

Further reading on VIP and PACAP receptors
Harmar AJ et al. (1998) International Union of Pharmacology. XVIII. Nomenclature of receptors for vasoactive intestinal peptide and pituitary adenylate cyclase-activating polypeptide. Pharmacol Rev 50: 265-270 [PMID:9647867]
Harmar AJ et al. (2012) Pharmacology and functions of receptors for vasoactive intestinal peptide and pituitary adenylate cyclase-activating polypeptide: IUPHAR review 1. Br. J. Pharmacol. 166: 4-17 [PMID:22289055]
Reglodi D et al. (2012) Effects of pituitary adenylate cyclase activating polypeptide in the urinary system, with special emphasis on its protective effects in the kidney. Neuropeptides 46: 61-70 [PMID:21621841]
Smith CB et al. (2012) Is PACAP the major neurotransmitter for stress transduction at the adrenomedullary synapse? J. Mol. Neurosci. 48: 403-12 [PMID:22610912]

Searchable database: http://www.guidetopharmacology.org/index.jsp
Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full
Searchable database: http://www.guidetopharmacology.org/index.jsp

Full Contents of ConciseGuide: http://onlinelibrary.wiley.com/doi/10.1111/bph.13878/full

S.P.H. Alexander et al. The Concise Guide to PHARMACOLOGY 2017/18: G protein-coupled receptors. British Journal of Pharmacology (2017) 174, S17–S129

1155. Levoye A et al. (2006) [1678767]
1156. Lewis TA et al. (2004) [15482930]
1157. Li X et al. (2004) [8969379]
1158. Li AH et al. (1998) [9703464]
1159. Li J et al. (2004) [15027861]
1160. Li L et al. (2002) Neuropharmacology 43: 295
1161. Li R et al. (2013) [23239822]
1162. Li X et al. (2012) [12013525]
1163. Liang BT Urso R Sambraski E et al. (2010) In Adenosine Receptors from Cell Biology to PharmacologyEdited by Borea P: Springer: 257-280 [ISBN: 9789048131440]
1164. Liang M et al. (2000) [10074402]
1165. Liang TS et al. (2001) [11714831]
1166. Liepa A et al. (2004) [15102946]
1167. Liaw CW et al. (2009) [19630353]
1168. Liebscher I et al. (2011) [21097509]
1169. Litgott SB. (2003) [15091997]
1170. Ligneux X et al. (2000) [11090994]
1171. Liljebri C et al. (1995) [7830272]
1172. Lim HD et al. (2012) [19375563]
1173. Limonta P et al. (1999) [10079080]
1174. Lim HD et al. (2003) [14726258]
1175. Lin DC et al. (2002) [11886876]
1176. Lin DC et al. (2012) [22859723]
1177. Lin L et al. (1999) [10458611]
1178. Lin Q et al. (1999) [9890897]
1179. Linden J et al. (1999) [10496592]
1180. Lindsay CW et al. (2004) [15537338]
1181. Lippert K et al. (1999) [10852255]
1182. Linz K et al. (2014) [24713140]
1183. Listschig S et al. (1999) [10051528]
1184. Liu C et al. (2005) [15465295]
1185. Liu C et al. (2003) [14522967]
1186. Liu C et al. (2003) [15422968]
1187. Liu C et al. (2012) [22436474]
1188. Liu C et al. (2012) [15161071]
1189. Liu C et al. (2009) [19047060]
1190. Liu C et al. (2011) [21796211]
1191. Liu C et al. (2012) [22267580]
1192. Liu C et al. (2009) [19369576]
1193. Liu P et al. (2011) [24900283]
1194. Liu Q et al. (2019) [10581185]
1195. Liu Q et al. (2009) [20004959]
1196. Liu S et al. (1998) [8822540]
1197. Liu W et al. (2012) [22798613]
1198. Llinares M et al. (1999) [10231175]
1199. Lobo MK et al. (2007) [17934457]
