I. Introduction and Historical Perspective

The classification of histamine receptors has to date been based on rigorous classical pharmacological analysis, and as yet, the classification of the three histamine receptors that have been defined by this process, (i.e., the H1-, H2-, and H3-receptors) have not been added to because of more recent molecular biological approaches (Schwartz et al., 1991, 1995; Hill, 1990; Leurs et al., 1995b). The scant number of known histamine receptors, compared with the plethora of receptors for some other endogenous substances, probably reflects the relative neglect of histamine rather than a paucity of its receptors. There is some preliminary evidence of heterogeneity of the known histamine receptors (which will be reviewed later in this article), but the acceptance of additional subtypes still awaits the identification of “sequence differences” within a single species and the development of selective agonists and antagonists providing the structural, recognition, and transductional information necessary for reliable classification.

The first histamine receptor antagonists (popularly referred to as the classical antihistamines but now called H1-receptor antagonists) were synthesized (Bovet and Staub, 1936; Bovet, 1950) over 20 years after the discovery (Barger and Dale, 1910) and descriptions of some of the physiological effects (Dale and Laidlaw, 1910) of histamine. These accomplishments had been preceded, as for some other endogenous biogenic amines, by its synthesis as a chemical curiosity (Windaus and Vogt, 1907). Early studies of the antihistamines were qualitative, for example, the demonstration of their ef-
fectiveness in protecting against bronchospasm produced in guinea pigs by anaphylaxis or administration of histamine (Bovet and Staub, 1936). Though qualitative, these studies yielded compounds, e.g., mepyramine (pyrilamine), that remain major ligands to define histamine receptors.

These antagonists were shown to reduce the effects of histamine on many tissues, notably vascular and extravascular smooth muscle (e.g., guinea pig ileum), but it became apparent that some of the effects of histamine were refractory to these classical antihistamines (Loew, 1947). For example, histamine-stimulated gastric secretion was shown to be unresponsive to three different antihistamines (Ashford et al., 1949). The vasodilator response to histamine in the cat was shown to be only partly sensitive to an antihistamine, leading to the suggestion that histamine causes vasodilatation by combining with more than one receptor (Folkw et al., 1948).

The application of the method of Schild (Arunlakshana and Schild, 1959) to the classification of receptors revealed that the pA$_2$ (−log $K_B$) value of mepyramine for antagonism of the positive chronotropic effect of histamine on the right atrium of the guinea pig differed from mepyramine’s pA$_2$ value for antagonism of the contractile response to histamine in guinea pig ileum, implying that the receptors involved were distinct (Arunlakshana and Schild, 1959; Trendelenburg, 1960). The histamine receptor in guinea pig ileum and in other tissues that showed the same or similar pA$_2$ value for these early antihistamines was then named the H$_1$-receptor (Ash and Schild, 1966). As the relative potencies of these histamine antagonists and histamine agonists on gastric acid secretion, relaxation of rat uterus, and chronotropy of the guinea pig right atrium differed from those on the H$_1$-receptor, it was concluded that a separate histamine receptor was involved in these responses.

The development of specific antagonists (H$_2$-antagonists) for this novel receptor represents a classic example of rational drug design (Black et al., 1972; Black, 1989) and showed the “practical value” (Green and Maayani, 1987; Jenkinson, 1987) of a quantitative approach to the analysis of receptor antagonism (Arunlakshana and Schild, 1959). Burimamide was the first compound to be described (Black et al., 1972) that had a higher pA$_2$ for antagonism of the histamine-mediated responses on guinea pig atrium and rat uterus than the pA$_2$ determined for antagonism of the contractile response to histamine in guinea pig ileum. Burimamide was also able to reduce gastric acid secretion in dogs and humans and to reduce the blood pressure response of the cat to histamine (Black et al., 1972). A large number of more potent and selective H$_2$-receptor antagonists have since been developed (Cooper et al., 1990), although further quantitative investigations of the antagonist potency of burimamide on other histamine-mediated responses contributed to the definition and classification of the histamine H$_3$-receptor (Arrang et al., 1983).

The third histamine receptor was also defined by a functional assay. Histamine was found to inhibit its own synthesis and release in rat cerebral cortical slices, and the effects of H$_1$- and H$_2$-receptor agonists and antagonists indicated a distinct receptor (Arrang et al., 1983, 1987b). A highly selective agonist, R-(α)-methylhistamine, and antagonist, thioperamide, clearly defined the H$_3$-receptor (Arrang et al., 1987). Since that time, considerable efforts have been made to develop other H$_3$-receptor–selective agonists and antagonists (Garbarg et al., 1992; Jansen et al., 1992; Van der Goot et al., 1992; Vollinga et al., 1994; Ganellin et al., 1995; Ligneau et al., 1995; Stark et al., 1996b,c).

Table 1 summarizes some of the operational characteristics used to define the nature of the histamine receptor involved in different tissue responses. Histamine derivatives are numbered according to the system given in figure 1 (Black and Ganellin, 1974).

## II. Histamine H$_1$-Receptor

### A. Distribution and Function

The study of the distribution of histamine H$_1$-receptors in different mammalian tissues has been greatly aided by the development of selective radioligands for this particular histamine receptor subtype. [$^3$H]mepyramine was originally developed in 1977 (Hill et al., 1977) and since that time has been used successfully to detect H$_1$-receptors in a wide variety of tissues including: mammalian brain; smooth muscle from airways, gastrointestinal tract, genitourinary system, and the cardiovascular system; adrenal medulla; and endothelial cells and lymphocytes (Hill, 1990). In some tissues and cells, however, it is notable that [$^3$H]mepyramine additionally binds to secondary non-H$_1$–receptor sites (Chang et al., 1979a; Hill and Young, 1980; Hadfield et al., 1983; Mitsuhashi and Payan, 1988; Arias-Montano and Young, 1993; Dickenson and Hill, 1994; Leurs et al., 1995b). In rat liver, in which [$^3$H]mepyramine predominantly binds to a protein homologous with debrisoquine 4-hydroxylase cytochrome P450 (Fukui et al., 1990), quinine can be used to inhibit this nonspecific binding. This observation has led Liu et al. (1992) to suggest that quinine may be used to inhibit binding to other lower affinity sites. However, it is clear that not all secondary binding sites for [$^3$H]mepyramine are sensitive to inhibition by quinine (Dickenson and Hill, 1994). Thus, in DDT$_1$MF-2 cells, a 38 to 40 kDa protein has been isolated, which binds H$_1$-receptor antagonists with $K_D$ values in the micromolar range (Mitsuhashi and Payan, 1988; Mitsuhashi et al., 1989) but which is not sensitive to inhibition by quinine (Dickenson and Hill, 1994). Nevertheless, DDT$_1$MF-2 cells can be shown to additionally possess [$^3$H]mepyramine binding sites that have the characteristics of H$_1$-receptors (i.e., $K_D$ values in the nanomolar range) and to mediate functional responses, which are clearly produced by histamine H$_1$-receptor.
activation (Dickenson and Hill, 1992; White et al., 1993; Dickenson and Hill, 1994).

Other radioligands that have been used to study histamine H₁-receptors are [3H]mianserin (Peroutka and Snyder, 1981), [3H]dopamine (Tran et al., 1981; Kamba and Richelson, 1984; Taylor and Richelson, 1982), [125I]iodobolpyramine (Bouthenet et al., 1988), [125I]iodoazidocephedrine (Ruat et al., 1988), and [3H][(+)-N-methyl-4-methylidiphenhydramine (Treherne and Young, 1988b). [125I]Iodobolpyramine has been used for autoradiographic localization of H₁-receptors in guinea pig brain, although less success has been achieved in rat brain (Körner et al., 1986; Bouthenet et al., 1988). The very slow dissociation of [3H]mepyramine from H₁-receptors at low temperatures (e.g., 4°C) does, however, mean that this ligand can also be used for autoradiography (Palacios et al., 1981a,b; Rotter and Frostholm, 1986). [125I]Iodoazidocephedrine is a very potent H₁-receptor antagonist that can bind irreversibly to H₁-receptors following irradiation with ultraviolet light (Ruat et al., 1988). [11C]Mepyramine and [11C]dopamine have also proved useful for imaging histamine H₁-receptors in the living human brain (Villemagne et al., 1991; Yanai et al., 1992, 1995).

H₁-receptors have been extensively studied in blood vessels (Barger and Dale, 1910; Dale and Laidlaw, 1910; Folkow et al., 1948; Black et al., 1972) and other smooth muscle preparations (Ash and Schild, 1966; Black et al., 1972; Marshall, 1955; Hill, 1990). In smooth muscles, such as the guinea pig ileum, which freely generate muscle action potentials, modulation of action-potential discharge by low concentrations of histamine is an important mechanism by which tension is increased (Bolton, 1979; Bolton et al., 1981; Bülbbring and Burnstock, 1960). In guinea pig ileum, there is also evidence

![Fig. 1. Numbering for histamine derivatives.](image)

**TABLE 1**
Operational characteristics of histamine receptors

<table>
<thead>
<tr>
<th>Receptor</th>
<th>Location</th>
<th>Response</th>
<th>Agonists</th>
<th>Antagonists</th>
</tr>
</thead>
<tbody>
<tr>
<td>Histamine H₁</td>
<td>Most smooth muscle, endothelial cells, adrenal medulla, heart, CNS</td>
<td>Smooth muscle contraction, stimulation of NO formation, endothelial cell contraction, increased vascular permeability, stimulation of hormone release, negative inotropism, depolarization (block of leak potassium current) and increased neuronal firing, inositol phospholipid hydrolysis and calcium mobilization, hyperpolarization by Ca²⁺-dependent potassium current</td>
<td>Histamineᵃ</td>
<td>Mepyramine (+) and (−)</td>
</tr>
<tr>
<td></td>
<td>Gastric parietal cells, vascular smooth muscle, suppressor T cells, neutrophils, CNS, heart, uterus (rat)</td>
<td>Stimulation of gastric acid secretion, smooth muscle relaxation, stimulation of adenyl cyclase, positive chronotropic and inotropic effects on cardiac muscle, decreased firing rate, hyperpolarization or facilitation of signal transduction in CNS, block of Ca²⁺-dependent potassium conductance (1 AHP, accommodation of firing, after-hyperpolarization), increase of hyperpolarization-activated current, inhibition of lymphocyte function</td>
<td>Histamineᵃ</td>
<td>Cimetidine</td>
</tr>
<tr>
<td>Histamine H₂</td>
<td>CNS, peripheral nerves (heart, lung, gastrointestinal tract), endothelium, enterochromaffin cells</td>
<td>Inhibition of neurotransmitter release, endothelium-dependent relaxation of rabbit middle cerebral artery, inhibition of gastric acid secretion (dog), increase in smooth muscle voltage-dependent Ca²⁺ current, inhibition of firing of tuberomammillary (histaminergic) neurons</td>
<td>Histamineᵃ</td>
<td>R-α-methylhistamine</td>
</tr>
<tr>
<td>Histamine H₃</td>
<td>CNS, central nervous system.</td>
<td>Inhibition of neurotransmitter release, endothelium-dependent relaxation of rabbit middle cerebral artery, inhibition of gastric acid secretion (dog), increase in smooth muscle voltage-dependent Ca²⁺ current, inhibition of firing of tuberomammillary (histaminergic) neurons</td>
<td>Histamineᵃ</td>
<td>Nα-methylhistamine</td>
</tr>
</tbody>
</table>

ᵃ Nonselective.
ᵇ H₂-antagonist.
that a component of the contractile response to histamine is mediated by inositol 1,4,5-trisphosphate–induced mobilization of intracellular calcium (Morel et al., 1987; Bolton and Lim, 1989; Donaldson and Hill, 1986b). In nonexcitable smooth muscles, such as airway and vascular smooth muscle, contractile responses to H₁-receptor stimulation primarily involve mobilization of calcium from intracellular stores as a consequence of inositol phospholipid hydrolysis (Matsumoto et al., 1986; Kotlikoff et al., 1987; Takuwa et al., 1987; Hall and Hill, 1988; Paniettiere et al., 1989; Van Amsterdam et al., 1989).

In vascular endothelial cells, H₁-receptor stimulation leads to several cellular responses including: (a) changes in vascular permeability (particularly in postcapillary venules) as a result of endothelial cell contraction (Majno and Palade, 1961; Majno et al., 1968; Meyrick and Brigham, 1983; Grega, 1986; Killackey et al., 1986; Svensjo and Grega, 1986); (b) prostacyclin synthesis (McIntyre et al., 1985; Brotherton, 1986; Carter et al., 1988; Resink et al., 1987); (c) synthesis of platelet-activating factor (McIntyre et al., 1985); (d) release of prostacyclin from human T lymphocytes using [125I]iodobolpyramine (Villemain et al., 1990) and shown to increase [Ca^{2+}] (Kitamura et al., 1996).

Histamine H₁-receptors have long been established to be present in the adrenal medulla and to elicit the release of catecholamines (Emmelin and Muren, 1949; Staszewska-Barczak and Vane, 1965; Robinson, 1982; Livett and Marley, 1986; Noble et al., 1988). Thus, histamine can induce the release of both adrenaline and noradrenaline from cultured bovine adrenal chromaffin cells (Livett and Marley, 1986). In these cells, histamine can also stimulate phosphorylation of the catecholamine biosynthesis enzyme tyrosine hydroxylase via a mechanism that involves release of intracellular calcium (Bunn et al., 1995). In addition to its effects on catecholamine synthesis and release from adrenal chromaffin cells, histamine can also elicit the release of leucine- and noradrenaline from cultured bovine adrenal chromaffin cells, histamine can also elicit the release of catecholamines (Emmelin and Muren, 1949; Staszewska-Barczak and Vane, 1965; Robinson, 1982; Livett and Marley, 1986; Noble et al., 1988). Thus, histamine can induce the release of both adrenaline and noradrenaline from cultured bovine adrenal chromaffin cells (Livett and Marley, 1986). In these cells, histamine can also stimulate phosphorylation of the catecholamine biosynthesis enzyme tyrosine hydroxylase via a mechanism that involves release of intracellular calcium (Bunn et al., 1995). In addition to its effects on catecholamine synthesis and release from adrenal chromaffin cells, histamine can also elicit the release of leucine- and noradrenaline from cultured bovine adrenal chromaffin cells, histamine can also elicit the release of leucine- and methionine-enkephalin (Bommer et al., 1987). Furthermore, after prolonged exposure to histamine, there is a marked increase in messenger ribonucleic acid-encoding proenkephalin A (Bommer et al., 1987; Kley, 1988; Wan et al., 1989).

In human atrial myocardium and guinea pig ventricle, histamine produces negative inotropic effects (Guo et al., 1984; Genovese et al., 1988; Zavecz and Levi, 1978). In human myocardium, this response is associated with inhibitory effects on heart rate and can be unmasked when the positive effects of histamine on the rate and force of contraction (mediated via H₂-receptors) are attenuated by conjoint administration of adenosine or adenosine A₁-receptor agonists (Genovese et al., 1988). However, in guinea pig left atria (Reinhardt et al., 1974, 1977; Steinberg and Holland, 1975; Hattori et al., 1983, 1988a) and rabbit papillary muscle (Hattori et al., 1988b), histamine produces a positive inotropic response via a mechanism that is not associated with a rise in adenosine 3c,5c-cyclic monophosphate (cAMPb) levels (see Hill, 1990).

Histamine H₁-receptors are widely distributed in mammalian brain (Hill, 1990; Schwartz et al., 1991). In human brain, higher densities of H₁-receptors are found in neocortex, hippocampus, nucleus accumbens, thalamus, and posterior hypothalamus, whereas cerebellum and basal ganglia show lower densities (Chang et al., 1979b; Kamba and Richelson, 1984; Martinez-Mir et al., 1990; Villemagne et al., 1991; Yanai et al., 1992). The distributions in rat (Palacios et al., 1981a) and guinea pig (Palacios et al., 1981b; Bouthenet et al., 1988) are similar to each other and to humans with the exception that the guinea pig cerebellum shows high density (Ruat and Schwartz, 1989; Chang et al., 1979b; Hill and Young, 1980; Palacios et al., 1981b; Bouthenet et al., 1988). In most brain areas, there was overlap of H₁-receptor binding sites and messenger ribonucleic acid levels except in hippocampus and cerebellum in which the discrepancy is likely to reflect the presence of abundant H₁-receptors in dendrites of pyramidal and Purkinje cells, respectively (Traiffort et al., 1994). Histamine H₁-receptor activation causes inhibition of firing and hyperpolarization in hippocampal neurons (Haas, 1981) and an amphetamine-sensitive outward current in olfactory bulb interneurons (Jahn et al., 1995), effects most likely produced by intracellular Ca²⁺ release. However, many other notably vegetative ganglia (Christian et al., 1989), hypothalamic supraoptic (Haas et al., 1975), brainstem (Gerber et al., 1990; Khatib et al., 1990), thalamic (McCormick and Williamson, 1991), and human cortical neurons (Reiner and Kamondi, 1994) are excited by histamine H₁-receptor activation through a block of a potassium conductance.

**B. H₁-Selective Ligands**

Although a large number of compounds have been synthesized as selective and competitive antagonists of the histamine H₁-receptor (see for example Casy, 1977; Ganellin, 1982), chemical effort directed at the generation of highly potent and selective H₁-receptor agonists has not achieved the same success. Modification of the ethylamine side chain of histamine is not favorable for H₁-receptor agonism (Leurs et al., 1995b). Furthermore, resolution of the enantiomers of the chiral compounds generated by methylation of the α- or β-positions did not reveal any stereoselectivity of the side chain for the H₁-receptor (Arrang et al., 1987; Leurs et al., 1995). Alkylation of the side chain amine group does not dras...
tically reduce H1-receptor activity, but N\textsuperscript{N}- and N\textsuperscript{O},N\textsuperscript{O}-
dimethylhistamine are also potent agonists for the H\textsubscript{3}
receptor (table 2; fig. 2; Arrang et al., 1983). Modification
of the imidazole moiety of histamine has been the most
successful approach for obtaining agonists with selectivity
for the H\textsubscript{1}-receptor. Replacement of the imidazole
moiety of histamine by other aromatic heterocyclic ring
structures in 2-pyridylethylamine and 2-thiazolylethy-
lanine yields two compounds with selectivity for the
H\textsubscript{1}-receptor (table 2; fig. 2). Both compounds act as full
agonists in producing contraction of guinea pig ileum
(Donaldson and Hill, 1986c), but in other tissues (e.g.,
guinea pig cerebral cortical slices or DDT\textsubscript{1}MF-2 cells),
2-pyridylethylamine behaves as a low-efficacy agonist
(Donaldson and Hill, 1986a; White et al., 1993). Substi-
tutions in the 2-position of the imidazole ring of hista-
mine have produced compounds that are the most selec-
tive H\textsubscript{1}-agonists available (Zingel et al., 1995). Thus,
2(3-bromophenyl)histamine and 2\{3-(trifluoromethyl)-
phenyl\}histamine are both relatively potent and highly
selective H\textsubscript{1}-agonists available (Zingel et al., 1995). Thus,
both compounds appear to be potent H\textsubscript{1}-agonists
in guinea pig ileum (Leschke et al., 1995), although some
of the halogenated 2-phenylhistamines are low-efficacy
agonists in DDT\textsubscript{1}MF-2 cells (Zingel et al., 1990; White et
al., 1993) and in guinea pig aorta (Leschke et al., 1995)
and can exhibit partial agonist properties.

Mepyramine (also known as pyrilamine) is the refer-
ence selective and high-affinity H\textsubscript{1}-receptor antagonist
(table 3; Hill, 1990). Other classical H\textsubscript{1}-antagonists that
have been used for characterization purposes include
chlorpheniramine, tripelennamine, promethazine, and
diphenhydramine (fig. 3). Some of these, however, poss-
several compounds between the three different histamine
receptors (table 3) does not guarantee an unambiguous
characterization. This can only be achieved by appro-
imate quantitative assessment of receptor antagonism,
preferably with a range of compounds of very different
chemical structure. The stereoisomers of chlorphenira-
mine are particularly useful in this regard (table 3). The
enantiomers of 4-methyl-diphenhydramine and brom-
pheniramine also differ by two orders of magnitude in
their affinity for the H\textsubscript{1}-receptor (Chang et al., 1979b;
Treherne and Young, 1988b). The geometric isomer
trans-triprodine is three orders of magnitude more pot-
tent than its cis counterpart and is one of the most
potent H\textsubscript{1}-antagonists available for the guinea pig H\textsubscript{1} re-
ceptor (tables 3 and 4; Ison et al., 1973). The tricyclic
antidepressants amitriptyline and doxepin are also very
potent H\textsubscript{1}-receptor antagonists (K\textsubscript{D} 0.6 and 0.1 nM re-
respectively; Figge et al., 1979; Aceves et al., 1985).

At therapeutic dosages, many of the classical H\textsubscript{1}-ant-
ithamines give rise to sedative side effects that have
been attributed to occupancy of H\textsubscript{1}-receptors in the cen-
tral nervous system (CNS) (Schwartz et al., 1981; Ni-
colson et al., 1991; Leurs et al., 1995b). Most of the
classical H\textsubscript{1}-antihistamines, including promethazine and ( + )-chlorpheniramine, readily cross the blood-brain
barrier. However, several compounds that penetrate
poorly into the CNS and appear to be devoid of central depressant effects are now available (fig. 4). These in-
clude terfenadine (Rose et al., 1982; Wiech and Martin,
1982), astemizole (Laduron et al., 1982; Niemegeers et
al., 1982), mequitazine (Uzan and Le Fer, 1979), lorata-
dine (Ahn and Barnett, 1986), acrivastine (Leighton et
al., 1983; Cohen et al., 1985), cetirizine (Timmerman,
1982), astemizole (Laduron et al., 1982; Niemegeers et
al., 1982), mequitazine (Uzan and Le Fer, 1979), lorata-
dine (Ahn and Barnett, 1986), acrivastine (Leighton et
al., 1983; Cohen et al., 1985), cetirizine (Timmerman,
1992b), and temelastine (Brown et al., 1986; Calcutt et
al., 1987). The pK\textsubscript{A} values for these agents are given in
table 5 (Ter Laak et al., 1994).

C. Receptor Structure

Photoaffinity binding studies using \(^{252}\text{I}j\)iodoazido-
phenylpyrpyramine and subsequent sodium dodecyl sulfate-
polyacrylamide gel electrophoresis analysis have indi-
cated that the H\textsubscript{1}-receptor protein has a molecular
weight of 56 kDa under reducing conditions in rat,
guinea pig, and mouse brain (Ruat et al., 1988, 1990b;
Ruat and Schwartz, 1989). Similarly, studies in bovine
adrenal medullar membranes with another photoaffini-
ity ligand \(^{3}\text{H}j\)azidobenzamide (Yamashita et al., 1991b)
found labeled peptides in the size range 53 to 58 kDa. Interestingly, the specifically labeled H₁-receptor (with [¹²⁵]Iodoazidophenpyramine) in guinea pig heart was found to have a substantially higher molecular weight, although there is no obvious difference in the pharmacological characteristics of the H₁-receptor in this tissue (Ruat et al., 1990a).

The bovine adrenal medulla H₁-receptor was cloned in 1991 by expression cloning in the *Xenopus* oocyte system (Yamashita et al., 1991a). The deduced amino acid se-
sequence represents a 491 amino acid protein with a calculated molecular weight of 56 kDa (table 6). The protein has the seven putative transmembrane (TM) domains expected of a G-protein-coupled receptor and possesses N-terminal glycosylation sites. A striking feature of the proposed structure is the very large third intracellular loop (212 amino acids) and relatively short (17 amino acids) intracellular C terminal tail. The availability of the bovine sequence and lack of introns has enabled the H1-receptor to be cloned from several species (table 6) including rat (Fujimoto et al., 1993), guinea pig (Horio et al., 1993; Traiffert et al., 1994), mouse (Inove et al., 1996), and human (De Backer et al., 1993; Fukui et al., 1994; Moguilevsky et al., 1994; Smit et al., 1996c). The human histamine H1-receptor gene has now been localized to chromosome 3 bands 3p14-p21 (Le Coniat et al., 1994).

At the present time, these different clones should be regarded as true species homologues of the histamine H1-receptor, even though there are notable differences between them in some antagonist potencies (table 4). Unfortunately, the number of H1-receptor antagonists evaluated in binding studies in cells transfected with the different recombinant receptors is rather limited. Nevertheless, it is clear that the stereoisomers of chlorpheniramine show marked differences between species. For example, the guinea pig H1-receptor has a KD of 0.9 nM for (+)-chlorpheniramine, whereas for the rat H1-receptor, the value is nearer 8 nM (table 4). Similar differences for this compound and others (notably mepyramine and triprolidine) have been reported for the native H1-receptors in guinea pig and rat brain, respectively (table 4; Chang et al., 1979b; Hill and Young, 1980; Hill, 1990). Such species differences may also explain why [125I]iodobolpyramine can label guinea pig CNS H1-receptors but is unable to detect H1-receptors in rat brain (Körner et al., 1986; Boucheniet et al., 1988). The native H1-receptor protein has been solubilized from both guinea pig and rat brain membranes (Toll and Snyder, 1982; Treherne and Young, 1988a), and the solubilized receptor retains the same differences in H1-antagonist potency for (+)-chlorpheniramine as that observed in membranes (Toll and Snyder, 1982). What is not clear, however, is why mepyramine appears to be more potent as an antagonist of the recombinant rat H1-receptor (expressed in C6 cells) than it is of the native H1-receptor in rat brain membranes (table 4; Chang et al., 1979b; Hill and Young, 1980; Fujimoto et al., 1993). The recombinant study performed in rat C6 cells (Fujimoto et al., 1993) is complicated by the presence of a low level of endogenous H1-receptors (Peakman and Hill, 1994), but a high affinity for mepyramine (KD = 1 nM) has been deduced from functional studies in untransfected C6 cells (table 4; Peakman and Hill, 1994).

Site-directed mutagenesis has begun to shed some light on the binding domains for H1-agonists and -antagonists. Amino acid sequence alignment of the cloned histamine H1- and H2-receptors (see fig. 5) has led to the suggestion that the third (TM3) and fifth (TM5) transmembrane domains of the receptor proteins are responsible for binding histamine (Birdsall, 1991; Timmerman, 1992a). Aspartate (107) in TM3 of the human H1-receptor, which is conserved in all aminergic receptors, has

### TABLE 3

**Antagonist dissociation constants at histamine receptors**

<table>
<thead>
<tr>
<th>Antagonist</th>
<th>H1 (nM)</th>
<th>H2 (nM)</th>
<th>H3 (nM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Doxepin</td>
<td>0.06</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Triprolidine</td>
<td>0.1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Temelastine</td>
<td>0.3</td>
<td>&gt;10</td>
<td></td>
</tr>
<tr>
<td>Mepyramine</td>
<td>0.4</td>
<td>5.2</td>
<td>&gt;1</td>
</tr>
<tr>
<td>(+)-Chlorpheniramine</td>
<td>0.4</td>
<td>1.2</td>
<td>&gt;58</td>
</tr>
<tr>
<td>(-)-Chlorpheniramine</td>
<td>204</td>
<td>1.2</td>
<td>&gt;58</td>
</tr>
<tr>
<td>Diphenhydrame</td>
<td>1.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Promethazine</td>
<td>1.2</td>
<td>3.0</td>
<td></td>
</tr>
<tr>
<td>Chlorpromazine</td>
<td>1.2</td>
<td>5.9</td>
<td></td>
</tr>
<tr>
<td>Tripelenamine</td>
<td>3.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Arpmidone</td>
<td>20</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cimetidine</td>
<td>450</td>
<td>800</td>
<td>33</td>
</tr>
<tr>
<td>Metiamide</td>
<td>n.d.</td>
<td>920</td>
<td>2.5</td>
</tr>
<tr>
<td>Ranitidine</td>
<td>&gt;100</td>
<td>200</td>
<td>&gt;1.2</td>
</tr>
<tr>
<td>Famotidine</td>
<td>n.d.</td>
<td>17</td>
<td></td>
</tr>
<tr>
<td>Zolantidine</td>
<td>6.2</td>
<td>25</td>
<td>&gt;10</td>
</tr>
<tr>
<td>Mefitidine</td>
<td>&gt;24</td>
<td>24</td>
<td>100</td>
</tr>
<tr>
<td>Tiotidine</td>
<td>&gt;30</td>
<td>15</td>
<td>&gt;12</td>
</tr>
<tr>
<td>Iodoaminpotentin</td>
<td>1.1</td>
<td>2.5</td>
<td></td>
</tr>
<tr>
<td>Impromidine</td>
<td>3.4</td>
<td></td>
<td>65</td>
</tr>
<tr>
<td>Burimamide</td>
<td>320</td>
<td>7.8</td>
<td>70</td>
</tr>
<tr>
<td>Thioperamide</td>
<td>&gt;100</td>
<td>&gt;10</td>
<td>4</td>
</tr>
<tr>
<td>Isohendrenioprop</td>
<td>n.d.</td>
<td>n.d.</td>
<td>0.25</td>
</tr>
<tr>
<td>Cloopenidone</td>
<td>&gt;10</td>
<td>&gt;10</td>
<td>0.13</td>
</tr>
<tr>
<td>Imperoxypant</td>
<td>1.4</td>
<td>5.3</td>
<td>5</td>
</tr>
<tr>
<td>GR174737</td>
<td>126</td>
<td>250</td>
<td>4</td>
</tr>
</tbody>
</table>

**Values determined in functional assays from guinea pig ileum contraction (H1), biochemical determinations in guinea pig cerebral cortical slices (H1), chronotropic responses in guinea pig atria (H2), cyclic AMP accumulation in guinea pig hippocampal slices (H3), inhibition of histamine release in rat cerebral cortical slices (H3), and inhibition of transmurally stimulated guinea pig ileum (H3). n.d., not determined.**
been shown to be essential for the binding of histamine and H1-receptor antagonists to the H1-receptor (Ohta et al., 1994). In the α2- and β2-adrenoceptors, two serine residues in TM5 accept the phenolic hydroxyl groups of the catechol ring of noradrenaline. In the H1-receptor, the residues corresponding to asparagine (198) and threonine (194) are in corresponding positions in TM5 of the human H1-receptor. However, substitution of an alanine for threonine (194) did not influence either agonist or antagonist binding (Ohta et al., 1994; Moguilevsky et al., 1995). Substitution of alanine (198) for asparagine (198) substantially decreased agonist, but not antagonist affinity (Ohta et al., 1994; Moguilevsky et al., 1995). Similar mutations to the corresponding residues (threonine (203) and asparagine (207) in the guinea pig H1-receptor sequence produce very similar results (Leurs et al., 1994a). It is interesting to note, however, that whereas 2-methylhistamine is similarly affected by the asparagine207 alanine mutation, the H1-selective agonists 2-thiazolylethylamine, 2-pyridylethylamine, and 2-(3-bromophenyl)histamine are much less affected by this mutation (Leurs et al., 1994a). These data suggest

**TABLE 4**

Species variation in H1-receptor antagonist potency (K_i, nM)

<table>
<thead>
<tr>
<th>Antagonist</th>
<th>Guinea pig h1 (CHO)</th>
<th>Human H1 (CHO)</th>
<th>Human H1 (brain)</th>
<th>Rat h1 (C6)</th>
<th>Rat h1 (brain)</th>
<th>Bovine H1 (Adrenal Medulla)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mepyramine</td>
<td>0.7</td>
<td>0.8</td>
<td>1.1, 4.0</td>
<td>1.0</td>
<td>9.1</td>
<td>2.6</td>
</tr>
<tr>
<td>(+)-Chlorpheniramine</td>
<td>0.9</td>
<td>0.8</td>
<td>3.5, 2.5</td>
<td>4.2</td>
<td>7.5 (4.4)%</td>
<td>9.1</td>
</tr>
<tr>
<td>(−)-Chlorpheniramine</td>
<td>103</td>
<td>200</td>
<td>316</td>
<td>350</td>
<td>540 (&gt;620)%</td>
<td>760</td>
</tr>
<tr>
<td>Triprolidine</td>
<td>0.7</td>
<td>0.2</td>
<td>1.0</td>
<td>3.7</td>
<td>2.0</td>
<td>n.d.</td>
</tr>
</tbody>
</table>

Unless otherwise stated, values show K_i determinations from inhibition of [3H]mepyramine binding. n.d., not determined.

_a_ h1 = transfected H1-receptor cDNA.

_b_ H1 = native/endogenous H1-receptor.

_c_ Values in parentheses show the values obtained from functional studies of the endogenous H1-receptor present in rat C6 cells (Peakman & Hill, 1994).
that asparagine (207) interacts with the N\textsuperscript{ε}-nitrogen of the imidazole ring of histamine. Furthermore, Leurs et al. (1995a) have recently shown that lysine (200) interacts with the N\textsuperscript{ε}-nitrogen of histamine and is important for the activation of the H\textsubscript{1}-receptor by histamine and the nonimidazole agonist, 2-pyridylethylamine. Interestingly, however, the lysine (200) alanine mutation did not alter the binding affinity of 2-pyridylethylamine to the guinea pig H\textsubscript{1}-receptor (Leurs et al., 1995).

\[ \text{CH}_3 \text{CH}_2 \text{CH}_2 \text{NH} \]
\[ \text{CH}_3 \text{CH} = \text{C} \text{H} \]
\[ \text{C}_6 \text{H}_4 \text{N} \text{C} \text{H} \]
\[ \text{CH}_2 \text{CH}_2 \text{OH} \]
\[ \text{CH}_2 \text{C}_6 \text{H}_4 \text{N} \text{CH}_2 \text{CH}_2 \text{CH}_2 \text{CH}_2 \text{NH}_2 \]
\[ \text{CH}_3 \text{CH}_2 \text{CO}_2 \text{H} \]

**D. Signal Transduction Mechanisms**

The primary mechanism by which histamine H\textsubscript{1}-receptors produce functional responses in cells is the activation of phospholipase C via a pertussis toxin-insensitive G-protein that is probably related to the G\textsubscript{q/11} family of G-proteins (Hill, 1990; Leurs et al., 1995b). The number of tissues and cell types in which a histamine H\textsubscript{1}-receptor–mediated increase in either inositol phosphate accumulation or intracellular calcium mobilization has been described is extensive, and further details are provided in several comprehensive reviews (Hill, 1990; Hill and Donaldson, 1992; Leurs et al., 1995b). Stimulation by histamine of \[^{3}H\]inositol phosphate accumulation and calcium mobilization has also been observed in Chinese hamster ovary (CHO) cells transfected with the human, bovine, and guinea pig H\textsubscript{1}-receptor complementary deoxyribonucleic acid (cDNA) (Leurs et al., 1994c; Smit et al., 1996c; Iredale et al., 1993; Megson et al., 1995). It is worth noting, however, that in some tissues, histamine can stimulate inositol phospholipid hydrolysis independently of H\textsubscript{1}-receptors. Thus, in the longitudinal smooth muscle of guinea pig ileum and neonatal

\[ \text{CH}_3 \text{CH}_2 \text{CH}_2 \text{NH} \]
\[ \text{CH}_3 \text{CH} = \text{C} \text{H} \]
\[ \text{C}_6 \text{H}_4 \text{N} \text{C} \text{H} \]
\[ \text{CH}_2 \text{CH}_2 \text{OH} \]
\[ \text{CH}_2 \text{C}_6 \text{H}_4 \text{N} \text{CH}_2 \text{CH}_2 \text{CH}_2 \text{CH}_2 \text{NH}_2 \]
\[ \text{CH}_3 \text{CH}_2 \text{CO}_2 \text{H} \]
rat brain (Donaldson and Hill, 1985, 1986b; Claro et al., 1987), a component can be identified in the response to histamine that is resistant to inhibition by H1-receptor antagonists. It remains to be established, however, whether these effects are due to “tyramine-like” effects of histamine on neurotransmitter release (Bailey et al., 1987; Young et al., 1988a) or direct effects of histamine on the associated G-proteins (Seifert et al., 1994).

In addition to effects on the inositol phospholipid signaling systems, histamine H1-receptor activation can lead to activation of several other signaling pathways, many of which appear to be secondary to changes in intracellular calcium concentration or the activation of protein kinase C. Thus, histamine can stimulate nitric oxide synthase activity (via a Ca2+/calmodulin-dependent pathway) and subsequent activation of soluble guanylyl cyclase in a variety of different cell types (Schmidt et al., 1990; Leurs et al., 1991a; Yuan et al., 1993; Casale et al., 1985; Duncan et al., 1980; Hattori et al., 1990; Sertl et al., 1987). Arachidonic acid release and the synthesis of arachidonic acid metabolites such as prostacyclin and thromboxane A₂ can also be enhanced by H1-receptor stimulation (Carter et al., 1988; Resink et al., 1987; Leurs et al., 1994c; Muriyama et al., 1990).

Interestingly, in CHO-K1 cells transfected with the guinea pig H1-receptor, the histamine-stimulated release of arachidonic acid is partially inhibited (approximately 40%) by pertussis toxin, whereas the same response in HeLa cells possessing a native H1-receptor was resistant to pertussis toxin treatment (Leurs et al., 1994c). The reason for this difference remains to be established, but it does caution against the use of signal transduction pathways in highly expressed recombinant cell systems as a primary receptor classification tool.

This point is best illustrated by the fact that in intact cellular systems, H1-receptor activation can produce substantial changes in the intracellular levels of cAMP. In most tissues, histamine H1-receptor activation does not activate adenyl cyclase directly but acts to amplify direct cAMP responses to histamine H2-, adenosine A2-, and vasoactive intestinal polypeptide receptors (Palacios et al., 1978; Al-Gadi and Hill, 1987, 1985; Donaldson et al., 1989; Garbarg and Schwartz, 1988; Magistretti and Schorderet, 1985; Marley et al., 1991). In many of these cases, a role for both intracellular Ca2⁺ ions and protein kinase C has been implicated in this augmentation response (Al-Gadi and Hill, 1987; Schwabe et al., 1978; Garbarg and Schwartz, 1988). In CHO cells transfected with the bovine or guinea pig H1-receptor, H1-
receptor activation can also lead to both direct cAMP responses and to an enhancement of forskolin-stimulated cAMP formation (Leurs et al., 1994c; Sanderson et al., 1996).

III. Histamine H$_2$-Receptor

A. Distribution and Function

Unlike the situation with H$_1$-selective radioligands, attempts to map the distribution of H$_2$-receptors by using radiolabeled H$_2$-receptor antagonists have met with variable success (Hill, 1990). Thus, [³H]cimetidine and [³H]ranitidine have proved unsuitable as H$_2$-radioligands, and in the case of cimetidine, the binding to sites specifically labeled with the radioligand is potently inhibited by imidazoles that have very low H$_2$-receptor binding affinities (Burkard, 1978; Kendall et al., 1980; Smith et al., 1980; Bristow et al., 1981; Warrender et al., 1983). More success has been achieved with [³H]tiotidine, which has a higher affinity for the H$_2$-receptor (table 7) in guinea pig brain, lung parenchyma, and CHO-K1 cells transfected with the human H$_2$-receptor cDNA (Gajtkowski et al., 1983; Norris et al., 1984; Sterk et al., 1986; Foreman et al., 1985a; Gantz et al., 1991a), although studies in rat brain were not successful (Maayani et al., 1982). At the present time, [¹²⁵I]iodoazidopotentidine has successfully been used for irreversible labeling (Ruat et al., 1990b; Hirschfeld et al., 1992).

Most information to date on the distribution of histamine H$_2$-receptor, however, has been provided by functional studies in different tissues (Hill, 1990). Histamine H$_2$-receptor–stimulated cAMP accumulation or adenylyl cyclase activity has been demonstrated in a variety of tissues including brain (Hegstrand et al., 1976; Green et al., 1977; Kanof et al., 1977; Palacios et al., 1978; Gajtkowski et al., 1983; Al-Gadi and Hill, 1985, 1987), gastric cells (Soll and Wollin, 1979; Gespach et al., 1982), and cardiac tissue (Johnson et al., 1979a,b; Kanof and Greengard, 1979a; Johnson, 1982). Histamine H$_2$-receptors have a potent effect on gastric acid secretion, and the inhibition of this secretory process by H$_2$-receptor antagonists has provided evidence for an important physiological role of histamine in the regulation of gastric secretion (Black et al., 1972; Black and Shankley, 1985; Soll and Berglindh, 1987). High concentrations of histamine are also present in cardiac tissues of most animal species and can mediate positive chronotropic and inotropic effects on atrial or ventricular tissues via H$_2$-receptor stimulation (Black et al., 1972; Inui and Imamura, 1976; Levi et al., 1982; Hattori and Levi, 1984; Hescheler et al., 1987; Levi and Alloatti, 1988). H$_2$-receptor–mediated smooth muscle relaxation has also been documented in airway, uterine, and vascular smooth muscle (Black et al., 1972; Reinhardt and Ritter, 1979; Gross et al., 1981; Eyre and Chand, 1982; Edvinsson et al., 1983; Foreman et al., 1985b; Ottosson et al., 1989). Finally, histamine H$_2$-receptors can inhibit a variety of functions within the immune system (Hill, 1990). H$_2$-receptors on basophils

<table>
<thead>
<tr>
<th>Receptor</th>
<th>Ligand</th>
<th>$K_D$ (nM)</th>
<th>Tissue</th>
</tr>
</thead>
<tbody>
<tr>
<td>H$_1$-receptor</td>
<td>[³H]Mepyramine</td>
<td>0.8</td>
<td>Guinea pig brain$^a$</td>
</tr>
<tr>
<td></td>
<td>[¹²⁵I]Iodobolpyramine</td>
<td>0.1</td>
<td>Guinea pig brain$^b$</td>
</tr>
<tr>
<td></td>
<td>[¹²⁵I]Iodoazidophenpyramine</td>
<td>0.01</td>
<td>Guinea pig cerebellum$^c$</td>
</tr>
<tr>
<td></td>
<td>[¹¹C]Mepyramine</td>
<td>1.0</td>
<td>Human brain (in vivo)$^d$</td>
</tr>
<tr>
<td></td>
<td>[¹¹C]Doxepin</td>
<td>0.1</td>
<td>Human brain (in vivo)$^e$</td>
</tr>
<tr>
<td>H$_2$-receptor</td>
<td>[³H]Tiotidine</td>
<td>25</td>
<td>Guinea pig brain$^f$</td>
</tr>
<tr>
<td></td>
<td>[¹²⁵I]Iodoaminopotentidine</td>
<td>0.3</td>
<td>Guinea pig brain$^g$</td>
</tr>
<tr>
<td></td>
<td>[¹²⁵I]Iodoazidopotentidine</td>
<td>10</td>
<td>Guinea pig brain$^g$</td>
</tr>
<tr>
<td>H$_3$-receptor</td>
<td>[¹¹I]R-(α)-methylhistamine</td>
<td>0.5</td>
<td>Rat brain$^h$</td>
</tr>
<tr>
<td></td>
<td>[¹¹I]N'-methylhistamine</td>
<td>2.0</td>
<td>Rat cerebral cortex$^1$</td>
</tr>
<tr>
<td></td>
<td>[¹²⁵I]Iodoproxyfan</td>
<td>0.3</td>
<td>Rat cerebral cortex$^1$</td>
</tr>
<tr>
<td></td>
<td>[³H]GR168320</td>
<td>0.065</td>
<td>Rat striatum$^i$</td>
</tr>
<tr>
<td></td>
<td>[³H]</td>
<td>0.1</td>
<td>Rat cerebral cortex$^1$</td>
</tr>
</tbody>
</table>

$^a$ Hill et al. (1981)  
$^b$ Körner et al. (1986)  
$^c$ Ruat et al. (1988)  
$^d$ Villemagne et al. (1991)  
$^e$ Yanai et al. (1995)  
$^f$ Gajtkowski et al. (1983)  
$^g$ Jansen et al. (1992)  
$^h$ Ligneau et al. (1994)  
$^i$ Brown et al. (1994)  
$^j$ Clark and Hill (1995)  
$^k$ Jansen et al. (1992)  
$^l$ Brown et al. (1994)
and mast cells have been shown to negatively regulate the release of histamine (Bourne et al., 1971; Lichtenstein and Gillespie, 1975; Lett-Brown and Leonard, 1977; Ting et al., 1980; Plaut and Lichtenstein, 1982). Furthermore, there is increasing evidence that H2-receptors on lymphocytes can inhibit antibody synthesis, T-cell proliferation, cell-mediated cytolysis, and cytokine production (Bourne et al., 1971; Melmon et al., 1974, 1981; Griswold et al., 1984; Khan et al., 1985, 1986; Sansoni et al., 1985; Melmon and Khan, 1987). In the CNS, histamine H2-receptor activation can inhibit nerve cells (Haas and Bucher, 1975; Haas and Wolf, 1977), but the most intriguing action is a block of the long-lasting after-hyperpolarization and the accommodation of firing, an effect with a remarkably long duration leading to potentiation of excitation in rodents (Haas and Konnerth, 1983; Haas and Greene, 1986) and human brain (Haas et al., 1988). A slow excitation is also common (Greene and Haas, 1989; Phelan et al., 1990). Synaptic transmission in the hippocampus is profoundly enhanced (Kostopoulos et al., 1988), and synaptic plasticity is induced or enhanced (Brown et al., 1995). An increase of the hyperpolarization-activated current has also been described in thalamic relay neurons (McCormick and Williamson, 1991). Indications for non-cAMP mediated actions of H2-receptor activation are given by Haas et al. (1978) and Jahn et al. (1995).

**B. H2-Selective Ligands**

The initial definition of the H1- and H2-subclasses of histamine receptor by Ash and Schild (1966) and Black and colleagues (1972) led to a successful search for H2-receptor selective antagonists with clinical relevance for the treatment of peptic ulcer. Burimamide was the first compound developed that showed selectivity for the H2-receptor (Black et al., 1972), but more recent work has shown that this compound is a more potent H3-receptor antagonist (Arrang et al., 1983). Cimetidine and metiamide were developed directly from burimamide (Black et al., 1974; Brimblecombe et al., 1975; Ganellin, 1978). Since then, a large number of compounds have been developed with H2-receptor antagonist properties (see Ganellin (1992) for review). These include ranitidine (Bradshaw et al., 1979), tiotidine (Yellin et al., 1979), nizatidine (Lin et al., 1986), famotidine (Takeda et al., 1982), and mifentidine (Donetti et al., 1984), which have been extensively used for characterization purposes (table 3; fig. 6). Iodoaminopotentidine (Ki = 2.5 nM) is one of the most potent H2-receptor antagonists available, and, as mentioned above, this compound has been used as a successful radioligand (Hirschfeld et al., 1992). Most H2-receptor antagonists are polar compounds and penetrate poorly into the CNS. Although this property is of great use for selective actions on peripheral tissues (e.g., gastric mucosa), it does limit the use of the com-

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![Chemical Structures](image-url)

**FIG. 6.** Histamine H2-receptor antagonists.
pounds for the in vivo evaluation of H2-receptor function within the CNS. However, one compound (zolantidine) is a potent and selective brain-penetrating histamine H2-receptor antagonist (table 3; Calcutt et al., 1988; Young et al., 1988b). Both cimetidine and ranitidine have been shown to demonstrate inverse agonism on histamine H2-receptors transfected into CHO cells (Smit et al., 1996a). Thus, in CHO cells expressing high levels of H2-receptors, in which a considerable constitutive activation of H2-receptors was demonstrated, cimetidine and ranitidine inhibited basal adenyl cyclase activity (Smit et al., 1996a). In contrast, burimamide behaved as a neutral antagonist (Smit et al., 1996a).

4-Methylhistamine was the first agonist described that had any selectivity for the H2-receptor (Black et al., 1972), although more potent and selective H2-agonists are now available (table 2). It is noteworthy that many of the selective H2-agonists exhibit H1-or H3-antagonist properties (see table 2); consequently the demonstration of H2-agonism in a given tissue or cell type needs confirming with H2-agonists. Impromidine is approximately 48 times more potent than histamine in mediating atrial chronotropic responses, but in several other H2-receptor–containing tissues, its relative potency and efficacy are lower (Durant et al., 1978; Leurs et al., 1995b). A large number of impromidine analogues have been synthesized and evaluated for H2-agonism. These studies have led to the development of the potent H2-agonists, impromidine and arpromidine (table 2; Timmerman, 1992c). Arpromidine and analogues are potential candidates for treatment of congestive heart failure (Buschauer, 1989; Buschauer and Baumann, 1991; Mörsdorf et al., 1990). Another potent H2-agonist has been derived as an analogue of dimaprit by considering cyclic adenylyl cyclase (Fukushima et al., 1995).

C. Receptor Structure

Photoaffinity binding studies using [125I]iodoazidotentidine and sodium dodecyl sulfate-polyacrylamide gel electrophoresis have suggested that the H2-receptor in guinea pig hippocampus and striatum has a molecular weight of 59 kDa (Ruát et al., 1990b). However, comparison with the calculated molecular weights (40.2 to 40.5 kDa) for the recently cloned H2-receptors (table 6) suggests that the native H2-receptor in guinea pig brain is glycosylated. Consistent with this proposal, it is noteworthy that all of the cloned H2-receptor proteins possess N-glycosylation sites in the N-terminus region (Gantz et al., 1991a,b; Ruát et al., 1991; Traiffort et al., 1995). Removal of these glycosylation sites by site-directed mutagenesis of the H2-receptor has been shown to decrease the sensitivity of the H2-receptor to adrenaline and to reduce the efficacy of the H2-receptor in response to histamine (Gantz et al., 1999). Similarly, changing the aspirate (186) of TM5 to an alanine resulted in complete loss of tiotidine binding without affecting the EC50 for histamine-stimulated cAMP formation (Gantz et al., 1992). Changing the threonine (190) to an alanine, however, resulted in a lower Kd for tiotidine and a reduction in both the maximal cAMP response and histamine EC50 value (Gantz et al., 1992). Mutation of Asp (186) and Gly (187) in the canine H2-receptor (to Ala (186) and Ser (187), however, produces a bifunctional receptor that can be stimulated by adrenaline and inhibited by both propranolol and cimetidine (Delvalle et al., 1995). Thus, these data suggest that the pharmacological specificity of the H2-receptor resides in only a few key amino acid residues.

Other site-directed mutagenesis studies on the H2-receptor have been very limited. However, Smit et al. (1996) have identified a residue in the second intracellular loop [leucine (124)] of the rat H2-receptor, which appears necessary for efficient coupling to Gs.

D. Signal Transduction Mechanisms

It is generally accepted that histamine H2-receptors couple to adenylyl cyclase via the GTP-binding protein Gs (Johnson, 1982; Hill, 1990; Leurs et al., 1995b). Histamine is a potent stimulant of cAMP accumulation in many cell types (Johnson, 1982), particularly those of CNS origin (Daly, 1977). Thus, H2-receptor–mediated effects on cAMP accumulation have been observed in...
brain slices (Al-Gadi and Hill, 1985; Palacios et al., 1978), gastric mucosa (Soll and Wollin, 1979; Chew et al., 1980; Batzri et al., 1982; Gespach et al., 1982), fat cells (Grund et al., 1975; Keller et al., 1981), cardiac myocytes (Warbanow and Wollenberger, 1979), vascular smooth muscle (Reinhardt and Ritter, 1979), basophils (Lichtenstein and Gillespie, 1975), and neutrophils (Busse and Sosman, 1977). Furthermore, H2-receptor–mediated cAMP accumulation has been demonstrated in CHO cells transfected with the rat, canine, or human H2-receptor cDNA (Gantz et al., 1991a,b; Leurs et al., 1994b; Fukushima et al., 1995).

Direct stimulation of adenylyl cyclase activity in cell-free preparations has been detected in both brain and cardiac muscle membranes (Hegstrand et al., 1976; Green et al., 1977; Green and Maayani, 1977; Kanof et al., 1977; Johnson et al., 1979a,b; Kanof and Greengard, 1979a,b; Newton et al., 1982; Olianas et al., 1984). However, caution is required regarding the interpretation of receptor characterization studies using histamine-stimulated adenylyl cyclase activity alone (Hill, 1990). A striking feature of studies of histamine H2-receptor–stimulated adenylyl cyclase activity in membrane preparations is the potent antagonism observed with certain neuroleptics and antidepressants (table 8; Spiker et al., 1976; Green et al., 1977; Green and Maayani, 1977; Kanof and Greengard, 1978; Green, 1983). It is notable, however, that most of the neuroleptics and antidepressants are approximately 2 orders of magnitude weaker as antagonists of histamine-stimulated cAMP accumulation in intact cellular systems (table 8; Tuong et al., 1980; Kamba and Richelson, 1983; Hill, 1990). One potential explanation of these differences resides within the buffer systems used for the cell-free adenylyl cyclase assays. Some differences in potency of some antidepressants and neuroleptics have been observed when membrane binding of H2-receptors has been evaluated using [125I]iodoaminopotentidine (table 8; Traiffort et al., 1991). However, invariably the differences observed in the K_i values deduced from ligand binding studies in different buffers are not as large as the differences in K_B values obtained from functional studies (table 8). For example, in the case of amitriptyline, no difference was observed in binding affinity in Krebs and Tris buffers (Traiffort et al., 1991).

In addition to G_s-coupling to adenylyl cyclase, there are reports of H2-receptors coupling to other signaling systems. For example, in gastric parietal cells, H2-receptor stimulation has been shown to increase the intracellular free concentration of calcium ions (Chew, 1985, 1986; Chew and Petropoulos, 1991; Malinowska et al., 1988; Delvalle et al., 1992a). A similar calcium response to histamine H2-receptor stimulation has also been observed in HL-60 cells (Mitsuhashi et al., 1989; Seifert et al., 1992) and in hepatoma-derived cells transfected with the canine H2-receptor cDNA (Delvalle et al., 1992b). In these latter cells, the influence on [Ca^{2+}]_i was accompanied by both an increase in inositol trisphosphate accumulation and a stimulation of cAMP accumulation (Delvalle et al., 1992b). Interestingly, the H2-receptor–stimulated calcium and inositol trisphosphate responses in these cells were both inhibited by choleratoxin treatment (but not by pertussis toxin), whereas choleratoxin produced the expected increase in cAMP levels (Delvalle et al., 1992a,b). In single parietal cells, H2-receptors have been shown to release calcium from intracellular calcium stores (Negulescu and Machen, 1988). It should be noted, however, that no effect of H2-agonists was observed on inositol phosphate accumulation or intracellular calcium levels in CHO cells transfected with the human H2-receptor (Leurs et al., 1994a).

### Table 8
Comparison of antagonist K_B values for inhibition of H2-receptor-stimulated adenylyl cyclase activity in membranes and cyclic AMP accumulation in intact cellular systems

<table>
<thead>
<tr>
<th>Antagonist</th>
<th>Antagonist K_B value (μM)</th>
<th>Binding studies (K_B, μM)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Slices^a</td>
<td>Dissociated cells^b</td>
</tr>
<tr>
<td>Cimetidine</td>
<td>0.6</td>
<td>0.5</td>
</tr>
<tr>
<td>Metiamide</td>
<td>0.8</td>
<td>n.d.</td>
</tr>
<tr>
<td>Tiotidine</td>
<td>n.d.</td>
<td>0.03</td>
</tr>
<tr>
<td>Cyproheptadine</td>
<td>5.7</td>
<td>n.d.</td>
</tr>
<tr>
<td>Mianserin</td>
<td>10.0</td>
<td>2.8</td>
</tr>
<tr>
<td>Imipramine</td>
<td>&gt; 10</td>
<td>3.3</td>
</tr>
<tr>
<td>Amitriptyline</td>
<td>3.5</td>
<td>1.9</td>
</tr>
<tr>
<td>Chlorpromazine</td>
<td>3.9</td>
<td>3.0</td>
</tr>
<tr>
<td>Haloperidol</td>
<td>&gt; 10</td>
<td>29</td>
</tr>
</tbody>
</table>

Measurements were made of H2-mediated adenylyl cyclase activity in homogenates of guinea pig hippocampus, impromidine-stimulated cyclic AMP accumulation in guinea pig hippocampal slices, and of H2-mediated cyclic AMP accumulation in dissociated hippocampal tissue. n.d., not determined.

^a Tuong et al. (1980)  
^b Kamba et al. (1983)  
^c Kanof and Greengard (1978)  
^d Kanof and Greengard (1979a,b)  
^e Green et al. (1977)  
^f Traiffort et al. (1991)  
^g Leurs et al. (1994a)
Thus, the effect of H3-receptor stimulation on intracellular calcium signaling may be very cell-specific.

In CHO cells transfected with the rat H3-receptor, H3-receptor stimulation produces both an increase in cAMP accumulation and an inhibition of P2a-receptor–mediated arachidonic acid release (Traiffort et al., 1992b). Interestingly, however, the effect on phospholipase A2 activity (i.e., arachidonic acid release) was not mimicked by forskolin, PGE1, or 8-bromo-cAMP, suggesting a mechanism of activation that is independent of cAMP-mediated protein kinase A activity (Arrang et al., 1992b). However, in CHO cells transfected with the human H3-receptor, no inhibitory effects of H3-receptor stimulation were observed on phospholipase A2 activity (Leurs et al., 1994b). This observation suggests that these cAMP-independent effects might depend on the level of receptor expression or subtle differences between clonal cell lines.

IV. Histamine H3-Receptor

A. Distribution and Function

The high apparent affinity of R-(α)-methylhistamine for the histamine H3-receptor has enabled the use of this compound as a radiolabeled probe (Arrang et al., 1987). This compound has been successfully used to identify a single binding site in rat cerebral cortical membranes, which in phosphate buffer has the pharmacological characteristics of the H3-receptor (Arrang et al., 1987, 1990). [3H]R-(α)-methylhistamine binds with high affinity (K_D = 0.3 nM) to rat brain membranes, although the binding capacity is generally low (approximately 30 fmol/mg protein; Arrang et al., 1987). Autoradiographic studies with [3H]R-(α)-methylhistamine have demonstrated the presence of specific thioperamide-inhibitable binding in several rat brain regions, particularly cerebral cortex, striatum, hippocampus, olfactory nucleus, and the bed nuclei of the stria terminalis, which receive ascending histaminergic projections from the magnocellular nuclei of the posterior hypothalamus (Arrang et al., 1987; Pollard et al., 1993). H3-receptors have also been visualized in human brain and the brain of nonhuman primates (Martinez-Mir et al., 1990). H3-receptor binding has been additionally characterized using [3H]R-(α)-methylhistamine in guinea pig cerebral corticosubcortical membranes (Kilpatrick and Michel, 1991), guinea pig lung (Arrang et al., 1987), guinea pig intestine, and guinea pig pancreas (Korte et al., 1990). Nα-methylhistamine has also proved successful as a radiolabeled probe for the H3-receptor. Although the relative agonist activity of Nα-methylhistamine (with respect to histamine) is fairly similar for all three histamine receptor subtypes (table 2), the binding affinity of histamine and Nα-methylhistamine for the H3-receptor is several orders of magnitude higher than for either the H1- or H2-receptors (Hill et al., 1977; Ruat et al., 1990b). This ligand can identify high-affinity H3-receptor sites in both guinea pig (Korte et al., 1990) and rat (West et al., 1990; Kathman et al., 1993; Clark and Hill, 1995) brain.

The binding of 3H-agonists to H3-receptors in brain tissues has been shown to be regulated by guanine nucleotides, implying a linkage to heterotrimeric G-proteins (Arrang et al., 1987, 1990; Zweig et al., 1992; Clark and Hill, 1995). The binding of H3-receptor agonists also seems to be sensitive to several cations. Magnesium and sodium ions have been shown to inhibit [3H]R-(α)-methylhistamine binding in rat and guinea pig brain (Kilpatrick and Michel, 1991), and the presence of calcium ions has been reported to reveal heterogeneity of agonist binding (Arrang et al., 1990). The inhibitory effect of sodium ions on agonist binding means that higher B_max values are usually obtained in sodium-free Tris buffers compared with that in Na/K phosphate buffers (Clark and Hill, 1995). West et al. (1990) have suggested that multiple histamine H3-receptor subtypes exist in rat brain (termed H3A and H3B) on the basis of [3H]Nα-methylhistamine binding in rat cerebral cortical membranes in 50 mM Tris buffer. Under these conditions, the selective H3-antagonist thioperamide can discriminate two affinity binding states (West et al., 1990). However, Clark and Hill (1995) have noted that the observed heterogeneity of thioperamide binding is dependent on the concentration of sodium ions or guanine nucleotides within the incubation medium. Thus, in the presence of 100 mM sodium chloride, thioperamide binding conforms to a single binding isotherm (Clark and Hill, 1995). The simplest interpretation of these data is that the H3-receptor can exist in different conformations for which thioperamide, but not agonists or other H3-antagonists (e.g., clobenpropit), can discriminate. Clark and Hill (1995) have suggested that the equilibrium between these conformations is altered by guanine nucleotides or sodium ions. If this hypothesis is correct, it is likely that the different binding sites represented resting, active, or G-protein–coupled conformations of the H3-receptor. Furthermore, if thioperamide preferentially binds to uncoupled receptors, then this compound should exhibit negative efficacy in functional assays.

More recently, radiolabeled H3-receptor antagonists have become available. The first compound to be developed was [125I]iodophenpropit, which has been used to successfully label H3-receptors in rat brain membranes (Jansen et al., 1992). Inhibition curves for thioperamide and iodophenpropit were consistent with interaction with a single binding site, but H3-receptor agonists were able to discriminate high- [4 nM for R-(α)-methylhistamine] and low- [0.2 μM for R-(α)-methylhistamine] affinity binding sites (Jansen et al., 1992). More recently, [3H]GR16820 (Brown et al., 1994) and [125I]iodoproxyfan (Ligneau et al., 1994) have also proved useful as high-affinity radiolabeled H3-antagonists. [125I]iodoproxyfan (Stark et al., 1996a) is the most potent and selective ligand available at the present time with a K_D of 65 pm (Ligneau et al., 1994). In rat striatum, in the
presence of guanine nucleotides such as guanosine 5'-O-(3-thiotriphosphate) (GTPγS), 40% of the binding sites exhibited a 40-fold lower affinity for \( H_3 \)-agonists, providing further evidence for a potential linkage of \( H_3 \)-receptors to G-proteins (Ligneau et al., 1994). \(^{[3H]}\)thioperamide and \(^{[3H]}\)5-methylthioperamide have also been used to label \( H_3 \)-receptors in rat brain membranes (Alves-Rodrigues et al., 1996; Yanai et al., 1994). However, \(^{[3H]}\)thioperamide was shown to bind additionally to low-affinity, high-capacity, non \( H_3 \)-receptor sites in this tissue (Alves-Rodrigues et al., 1996).

In addition to data obtained from ligand binding studies, evidence for the localization of histamine \( H_3 \)-receptors has also come from functional studies, primarily involving inhibition of neurotransmitter release. The \( H_3 \)-receptor was first characterized as an autoreceptor-regulating histamine synthesis and release from rat cerebral cortex, striatum, and hippocampus (Arrang et al., 1983, 1985b, c 1987a, 1988a,b). \( H_3 \)-receptor–mediated inhibition of histamine release has also been observed in human cerebral cortex (Arrang et al., 1988a). Differences in the distribution of \( H_3 \)-receptor binding sites and the levels of histidine decarboxylase (an index of histaminergic nerve terminals) suggested at an early stage that \( H_3 \)-receptors were not confined to histamine-containing neurons within the mammalian CNS (Arrang et al., 1987; Van der Werf and Timmerman, 1989). This has been confirmed by the observations that \( H_3 \)-receptors can regulate serotonergic (Schlicker et al., 1988), noradrenergic (Schlicker et al., 1989, 1992), cholinergic (Clapham and Kilpatrick, 1992), and dopaminergic (Schlicker et al., 1993) neurotransmitter release in mammalian brain. Histamine \( H_3 \)-receptor activation inhibits the firing of the histamine-neurons in the posterior hypothalamus through a mechanism different from autoreceptor functions found on other aminergic nuclei, presumably a block of Ca\(^{2+}\)-current (Haas, 1992). Electrophysiological evidence for reduction of excitatory neurotransmitter release (glutamate) has been presented by Brown and Reymann (unpublished data, 1996).

Inhibitory effects of \( H_3 \)-receptor activation on neurotransmission have also been documented in the periphery. Thus, \( H_3 \)-receptors have been identified regulating the release of sympathetic neurotransmitters in guinea pig mesenteric artery (Ishikawa and Sperelakis, 1987), human saphenous vein (Molderings et al., 1992), guinea pig atria (Endou et al., 1994; Imamura et al., 1994), and human heart (Imamura et al., 1995). Inhibition of parasympathetic nerve activity has also been observed in guinea pig ileum and human bronchi and tracheal (Trzeciakowski, 1987; Tamura et al., 1988; Ichinose et al., 1989; Ichinose and Barnes, 1989; Hew et al., 1990; Menkveld and Timmerman, 1990; Leurs et al., 1991a,b; Poli et al., 1991). An inhibitory effect of \( H_3 \)-receptor stimulation on release of neuuropeptides (tachykinins or calcitonin gene-related peptide) from sensory C fibers has been reported from airways (Ichinose et al., 1990), meninges (Matsubara et al., 1992), skin (Ohkubo and Shibata, 1995), and heart (Imamura et al., 1996). A modulation of acetylcholine, capsaicin, and substance P effects by histamine \( H_3 \)-receptors in isolated perfused rabbit lungs has also been reported (Delaunois et al., 1995).

There is evidence that \( H_3 \)-receptor stimulation can inhibit the release of neurotransmitters from nonadrenergic-noncholinergic nerves in guinea pig bronchiolus (Burgaud and Oudart, 1994) and ileum (Taylor and Kilpatrick, 1992). Interestingly, in guinea pig ileum, the \( H_3 \)-antagonists beta-histidine and phenylbutanoylhistamine were much less potent as inhibitors of \( H_3 \)-mediated effects on nonadrenergic-noncholinergic transmission than they were as antagonists of histamine release in rat cerebral cortex (Taylor and Kilpatrick, 1992). A similar low potency has been observed for these two antagonists for antagonism of \( H_3 \)-receptor–mediated \(^{[3H]}\)acetylcholine release from rat entorhinal cortex (Clapham and Kilpatrick, 1992) and antagonism of \( H_3 \)-receptor–mediated 5-hydroxytryptamine (5-HT) release from porcine enterochromaffin cells (Schworer et al., 1994). These observations provide support for the possible existence of distinct \( H_3 \)-receptor subtypes, but these responses need to be investigated further to exclude alternative explanations. For example, Arrang et al. (1995) have recently shown that phenylbutanoylhistamine can inhibit \(^{[3H]}\)acetylcholine release from rat entorhinal cortex slices and synaptosomes via a nonhistamine receptor mechanism. Thus, the potency of phenylbutanoylhistamine as an \( H_3 \)-receptor antagonist in these preparations may be greatly underestimated because of the additional nonspecific properties of the drug (Arrang et al., 1995).

The observed inhibitory effect of \( H_3 \)-receptor stimulation on 5-HT release from porcine enterochromaffin cells in strips of small intestine (Schworer et al., 1994) provides evidence for \( H_3 \)-receptors regulating secretory mechanisms in nonneuronal cells. This observation suggests that \( H_3 \)-receptors may also be present in gastric mast cells or enterochromaffin cells and exert an inhibitory influence on histamine release and gastric acid secretion. Consistent with this suggestion, \( H_3 \)-receptor activation has been shown to inhibit gastric acid secretion in conscious dogs (Soldani et al., 1993). An autoregulation of histamine synthesis by histamine \( H_3 \)-receptors has also been reported in isolated rabbit fundic mucosal cells (Holland et al., 1993).

\( H_3 \)-receptors have been shown to relax rabbit middle cerebral artery via an endothelium-dependent mechanism involving both nitric oxide and prostanoid release (Ea Kim and Oudart, 1988; Ea Kim et al., 1992). Finally, there is a report that \( H_3 \)-receptor activation can stimulate adrenocorticotropic hormone release from the pituitary cell line AtT-20 (Clark et al., 1992).
B. H₃-Receptor Selective Ligands

The initial characterization of the H₃-receptor made use of the relative high affinity of the agonists N⁴-methylhistamine and histamine for the H₃-receptor compared with the H₁- and H₂-receptors together with the H₃-antagonist properties of impropidine (H₂-agonist), burimamide (H₂-antagonist), and betahistine (H₁-agonist) (Arrang et al., 1983, 1985a). Since then, several selective ligands (both agonists and antagonists) have been developed that show little effect on H₁- and H₂-selective ligands (both agonists and antagonists) have been described (table 2). R-α,S-β-dimethylhistamine showed slightly higher potency and even higher selectivity (Lipp et al., 1992). Imetit [S-[2–4(5)]-imidazolylethylisothiourea] is a highly selective, full H₃-agonist that appears to be more potent than R-α-methylhistamine (table 2; Garbarg et al., 1992; Howson et al., 1992; Van der Goot et al., 1992). Both R-α-methylhistamine and imetit have been shown to be active in vivo at low doses (Arrang et al., 1987a; Garbarg et al., 1992). Azomethine derivatives of R-α-methylhistamine were prepared as lipophilic prodrugs to improve the bioavailability of the hydrophilic drug, particularly its entry into the brain (Krause et al., 1995). Immepip is another potent H₃-agonist that has been developed from histamine by extending the alkyl side chain to four methylene groups and incorporating the amino function within a piperidine ring (table 2; Vollinga et al., 1994). Most recently, the H₃-agonist potency of a cyclic, conformationally restricted analogue of histamine (immepyr) has been reported (Shih et al., 1995). This compound has been resolved and the (+)-immepyr shown to have an H₃-binding affinity (Kᵢ = 2.8 nM) one order of magnitude higher than the corresponding (-)-isomer (Shih et al., 1995). In guinea pig ileum, however, (+)-immepyr was one order of magnitude less potent (pD₂ 7.1) than R-α-methylhistamine (pD₂ 8.2) as an H₃-agonist (Shih et al., 1995).

Thioperamide was the first potent and selective H₃-receptor antagonist to be described (Arrang et al., 1987). This compound appears to act as a competitive antagonist in most functional assays of H₃-receptor activity (Arrang et al., 1987; Hew et al., 1990; Menkveld and Timmerman, 1990), although Clark and Hill (1995) have suggested that it may possess inverse agonist properties. More recently, several other potent H₃-antagonists have been described (table 3; fig. 7), including clobenpropit (Kathman et al., 1993), iodophenpropit (Jansen et al., 1992), GR175737 (Clitherow et al., 1996), iodoproxyfan (Ligneau et al., 1994; Schlicker et al., 1996), impentamine (Vollinga et al., 1995; Leurs et al., 1996), ethers (Ganellin et al., 1996; Stark et al., 1996a), and carba-
approximately 70 kDa (Cherifi et al., 1992). However, it remains to be established whether this protein is the histamine H₃-receptor.

D. Signal Transduction Mechanisms

The signal transduction pathways used by the histamine H₃-receptor remain largely subject to speculation, but there is increasing evidence to suggest that this receptor belongs to the superfamily of G-protein–coupled receptors. Evidence for this has largely been obtained from ligand-binding studies involving the modulation by guanine nucleotides of H₃-agonist binding (Arrang et al., 1990; West et al., 1990; Kilpatrick and Michel, 1991; Zweig et al., 1992; Clark and Hill, 1995) and of H₃-agonist inhibition of ³H-antagonist binding (Jansen et al., 1992, 1994; Ligneau et al., 1994). The most direct evidence for a functional H₃-receptor–G-protein linkage has come from studies of [³⁵S]GTPγS binding to rat cerebral cortical membranes (Clark and Hill, 1996). The presence of H₁- and H₂-receptor antagonists (0.1 μM mepyramine and 10 μM tiotidine), both R-α-methylhistamine and N°-methylhistamine produced a concentration-dependent stimulation of [³⁵S]GTPγS binding (EC₅₀ = 0.4 and 0.2 nM, respectively) in rat cerebral cortical membranes (Clark and Hill, 1996). Furthermore, this response was abolished by pretreatment of membranes with pertussis toxin, implying a direct coupling to a Gᵢ or Gₒ protein (Clark and Hill, 1996). Evidence for an involvement of pertussis toxin-sensitive G-proteins in the response to H₃-receptor stimulation has also come from studies of histamine H₃-receptor signaling in human and guinea pig heart (Endou et al., 1994; Imamura et al., 1995). In these tissues, histamine H₃-receptor–stimulation seems to lead to an inhibition of N-type Ca²⁺ channels responsible for voltage-dependent release of noradrenaline (Endou et al., 1994; Imamura et al., 1995).

Very little is known about the intracellular signal transduction pathways initiated by histamine H₃-receptor activation. Several research groups have failed to observe an inhibition of adenylyl cyclase activity in different tissues and cells (Garbarg et al., 1989; Schlicker et al., 1991; Cherifi et al., 1992), which might indicate that H₃-receptors preferentially couple to Gₒ proteins. There is one interesting report of a negative coupling to phospholipase C in the HGT-1 gastric tumor cell line (Cherifi et al., 1992), but this observation needs confirmation by other research.

V. Other Responses to Histamine

A. Potentiation of Responses to N-Methyl-D-Aspartate

Studies in hippocampal cell cultures, acutely dissociated neurons, and Xenopus oocytes expressing the re-
combinant N-methyl-d-aspartate (NMDA) receptor subunits NR2B and NR1 have shown that histamine is able to enhance NMDA-activated currents, independently of the known histamine receptors, via a mechanism that probably involves the polyamine-binding site on the NMDA-receptor complex (Bekkers, 1993; Vorobjev et al., 1993; Williams, 1994; Saysbasili et al., 1995). Histamine and the polyamines spermine and spermidine have also been shown to enhance glutamate toxicity in human NT2-N neurons (Munir et al., 1996). Interestingly, attempts to demonstrate a similar effect of histamine on NMDA-induced currents in rat hippocampal slices, or outside-out patches pulled from the somas of these cells, were without success (Bekkers et al., 1996). However, two studies using conventional and whole cell recording of neurons in the CA1 region of slices of rat hippocampus concluded that the modulation of NMDA-mediated synaptic currents was dependent upon pH (Saysbasili et al., 1995; Janovsky et al., 1995). Thus, at low pH (7.2), histamine enhanced synaptic currents, whereas at pH 7.6 it reduced them. Interestingly, at physiological pH (7.4), no significant action of histamine was seen (Saysbasili et al., 1995).

B. A Role as an Intracellular Messenger?

Although most actions of histamine can be attributed to an extracellular action, there are reports that histamine may have intracellular actions. The activity of the enzyme, histidine decarboxylase, which catalyzes the formation of histamine from histidine, has been observed to be high in several tissues undergoing rapid growth or repair (Ishikawa et al., 1970; Kahlon and Rosengren, 1971; Watanabe et al., 1981; Bartholeyns and Bouclier, 1984; Bartholeyns and Fozard, 1985). These observations have led to the proposal that newly synthesized (nascent) histamine may have a role in cellular proliferation, perhaps via an intracellular site. Some evidence has been accumulated that intracellular histamine levels (or the activity of histidine decarboxylase) can be regulated by tumor-promoting phorbol esters (Saxena et al., 1989). Furthermore, Brandes and colleagues (Saxena et al., 1989; Brandes et al., 1990, 1992) have suggested that N,N-diethyl-2-[4-(phenylmethyl)-phenoxy]ethanamine (DPPE) might be an inhibitor of a specific intracellular histamine receptor (H1C). However, at the present time, the evidence in favor of an intracellular histamine receptor has not been generally accepted, and alternative possibilities need to be explored. For example, the direct effects of histamine, or its analogues, on polyamine sites (Vorobjev et al., 1993; Bekkers, 1993) and heterotrimeric G-proteins (Hagelüken et al., 1995; Seifert et al., 1994) could explain many of the observations to date.

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IUPHAR CLASSIFICATION OF HISTAMINE RECEPTORS


