Stereospecific requirement of cholesterol in the function of the serotonin_{1A} receptor

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The serotonin_{1A} receptor is an important member of the G protein-coupled receptor (GPCR) family. It is involved in the generation and modulation of a variety of cognitive and behavioral functions and serves as a drug target. Previous work from our laboratory has established the sensitivity of the function of the serotonin_{1A} receptor to membrane cholesterol. Solubilization of the hippocampal serotonin_{1A} receptor utilizing the zwitterionic detergent CHAPS is accompanied by loss of cholesterol and results in reduction in specific ligand binding. Replenishment of cholesterol to solubilized membranes restores specific ligand binding to the receptor. We utilized this strategy of sterol replenishment of solubilized membranes to explore the stereospecific stringency of cholesterol for receptor function. We used two stereoisomers of cholesterol, ent-cholesterol (enantioomer of cholesterol) and epi-cholesterol (a diastereomer of cholesterol), for this purpose. Importantly, we show here that while ent-cholesterol could replace cholesterol in supporting receptor function, epi-cholesterol could not. These results imply that the requirement of membrane cholesterol for the serotonin_{1A} receptor function is diastereospecific, yet not enantiospecific. Our results extend and help define specificity of the interaction of membrane cholesterol with the serotonin_{1A} receptor, and represent the first report utilizing ent-cholesterol to examine stereospecificity of GPCR-cholesterol interaction.

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1. Introduction

The G protein-coupled receptor (GPCR) superfamily comprises the largest and most diverse group of proteins in mammals and is involved in information transfer (signal transduction) from outside the cell to the cellular interior [1–3]. GPCRs are typically seven transmembrane domain proteins, regulate physiological responses to a diverse array of stimuli, and mediate multiple physiological processes. Due to this reason, GPCRs have emerged as major drug targets in all clinical areas [4]. It is estimated that ~50% of clinically prescribed drugs target GPCRs [5].

The serotonin_{1A} (5-HT_{1A}) receptor is a representative member of the GPCR family and is implicated in the generation and modulation of various cognitive, behavioral, and developmental functions [6–8]. Ligands that bind to the serotonin_{1A} receptor are reported to possess potential therapeutic effects in anxiety or stress-related disorders [6]. As a consequence, the serotonin_{1A} receptor serves as an important target in the development of therapeutic agents for neuropsychiatric disorders such as anxiety and depression [9]. Since GPCRs are integral membrane proteins with multiple transmembrane passes, the interaction of GPCRs with membrane lipids is an important determinant in their structure and function [10–14]. In fact, an important feature observed in recently solved high resolution crystal structures of GPCRs (such as rhodopsin [15], β_{1}-adrenergic receptor [16], β_{2}-adrenergic receptor [17,18] and A2A adenosine receptor [19]) is the close association of cholesterol molecules to the receptor. Previous work from our laboratory comprehensively demonstrated the requirement of membrane cholesterol in the organization, dynamics, and function of the serotonin_{1A} receptor ([20–22]; reviewed in Refs. [11,12,14]).

Cholesterol is an essential and representative membrane lipid in higher eukaryotes and is crucial in membrane organization, dynamics, function, and sorting [23,24]. A hallmark of membrane cholesterol is its nonrandom distribution in domains (or pools) in biological and model membranes [25–28]. These domains are believed to be crucial since various cellular processes such as membrane sorting and trafficking [29], signal transduction [30], and the entry of pathogens [31,32] have been attributed to these types of domains. The role of cholesterol in
the function and organization of membrane proteins and receptors constitutes an emerging and exciting area of research [10–14]. The detailed mechanism underlying the effect of membrane cholesterol on the structure and function of membrane proteins and receptors is not clear and appears to be complex [12,33,34]. A possible mechanism by which membrane cholesterol has been proposed to modulate the function of membrane receptors is by a direct (specific) interaction, which could induce a conformational change in the receptor. An alternative mechanism envisages an indirect way by altering the membrane physical properties in which the receptor is embedded. Yet another possibility could be a combination of both. A particular kind of proposed specific interaction is based on the concept of “nonannular” binding sites of membrane lipids in membrane proteins [34,35]. Nonannular sites are characterized by lack of accessibility to the annular lipids, i.e., these sites cannot be displaced by competition with annular lipids [36,37].

As mentioned above, earlier work from our laboratory has comprehensively demonstrated the requirement of membrane cholesterol in the function of the serotonin1A receptor [11,12,14]. An important aspect of our results is that the interaction between cholesterol and the serotonin1A receptor was shown to be considerably stringent since immediate biosynthetic precursors of cholesterol (differing with cholesterol merely in a double bond) were not able to maintain receptor function [21,38,39]. In order to further explore the degree of structural (stereospecific) stringency necessary for the ligand binding function of the serotonin1A receptor, we examined whether stereoisomers of cholesterol [enantiomer of cholesterol (ent-cholesterol), or diastereomer of cholesterol (epi-cholesterol); see Fig. 1] could support the ligand binding function of the receptor. We show that while ent-cholesterol could replace cholesterol in supporting receptor function, epi-cholesterol could not.

2. Materials and methods

2.1. Materials

CHAPS, cholesterol, M(3)CD, DMPC, DPH, EDTA, EGTA, MgCl2, MnCl2, iodoacetamide, PEG, PMSF, serotonin, sucrose, polyethylenimine, sodium azide, and Tris were obtained from Sigma Chemical Co. (St. Louis, MO). 3-Epicholesterol (5-cholesten-3-ol), to be denoted as epi-cholesterol, was obtained from Steraloids (Newport, RI). The enantiomer of cholesterol (ent-cholesterol) was synthesized as previously described [40,41]. BCA reagent for protein estimation was from Pierce (Rockford, IL). [3H]-8-OH-DPAT (sp. activity 106 Ci/mmol) was purchased from DuPont New England Nuclear (Boston, MA). GF/B glass microfiber filters were from Whatman International (Kent, U.K.). All solvents used were of analytical grade. All other chemicals used were of the highest purity available. Water was purified through a Millipore (Bedford, MA) Milli-Q system and used throughout. Fresh bovine brains were obtained from a local slaughterhouse within 10 min of death and the hippocampal region was carefully dissected out. The hippocampi were ash frozen in liquid nitrogen and stored at −70 °C till further use.

2.2. Methods

2.2.1. Preparation of native hippocampal membranes

Native hippocampal membranes were prepared as described previously [42]. Bovine hippocampal tissue (~50 g) was homogenized as 10% (w/v) in a polytron homogenizer in 2.5 mM Tris, 0.32 M sucrose, 5 mM EDTA, 5 mM EGTA, 0.02% sodium azide, 0.24 mM PMSF, 10 mM iodoacetamide, pH 7.4 buffer. The homogenate was centrifuged at 900 × g for 10 min at 4 °C. The resultant supernatant was filtered through four layers of cheesecloth and centrifuged at 50,000 × g for 20 min at 4 °C. The pellet obtained was suspended in 10 vol. of 50 mM Tris, 1 mM EDTA, 0.24 mM PMSF, 10 mM iodoacetamide, pH 7.4 buffer using a hand-held Dounce homogenizer and centrifuged at 50,000 × g for 20 min at 4 °C. This procedure was repeated until the supernatant was clear. The final pellet (native hippocampal membranes) was suspended in a minimum volume of 50 mM Tris, pH 7.4 (buffer A), homogenized using a hand-held Dounce homogenizer, flash frozen in liquid nitrogen and stored at −70 °C. Protein concentration was assayed using the BCA reagent [43].

2.2.2. Solubilization of native membranes

Hippocampal membranes (HM) were solubilized as described previously using the zwitterionic detergent CHAPS [44–46]. CHAPS-solubilized membrane was precipitated using PEG in order to remove NaCl from the solubilized extract, since agonist binding of the serotonin1A receptor is inhibited by NaCl [42]. This procedure also removes detergent. The PEG-precipitated CHAPS-solubilized membrane (referred to as solubilized membrane (SM)) was suspended in buffer A and used immediately for sterol replenishment and radioligand binding assays.

2.2.3. Sterol replenishment of solubilized membranes

Solubilized membranes were replenished with ent-cholesterol, epi-cholesterol or cholesterol using water soluble complexes of M(3)CD and the respective sterol. The complex was prepared by dissolving required amounts of the sterol (ent-cholesterol, epi-cholesterol or cholesterol)
and M|CD in a ratio of 1:10 (mol/mol) in buffer A by constant vortexing at room temperature (~23 °C). Stock solutions (typically 2 mM of ent-cholesterol, epi-cholesterol or cholesterol:20 mM M|CD) of this complex were freshly prepared prior to each experiment. Sterol replenishments were carried out at a protein concentration of ~2 mg/ml by incubating solubilized membranes with 1 mM sterol:10 mM M|CD complex for 30 min in buffer A at 25 °C under constant shaking. Membranes were then spun down at 100,000 × g for 1 h at 4 °C, suspended in the same buffer, and immediately used for radioligand binding assays.

2.2.4. Radioligand binding assays
Receptor binding assays were carried out as described earlier [20] with some modifications. Tubes in duplicate with ~0.8 mg protein in a total volume of 1 ml of 50 mM Tris, 1 mM EDTA, 10 mM MgCl2, 5 mM MnCl2, pH 7.4 buffer were incubated with the radiolabeled agonist [3H]8-OH-DPAT (final concentration in assay tube being 0.29 nM) for 1 h at 25 °C. Nonspecific binding was determined by performing the assay in the presence of 10 μM serotonin. The binding reaction was terminated by rapid filtration under vacuum in a Millipore multipore filtration apparatus through Whatman GF/B 2.5 cm diameter glass microfiber filters (1.0 μm pore size), which were presoaked in 0.15% polyethylenimine for 1 h [47]. Filters were then washed three times with 3 ml of cold water (4 °C) dried, and the retained radioactivity was measured in a Packard Tri-Carb 1500 liquid scintillation counter using 5 ml of scintillation fluid.

2.2.5. Estimation of inorganic phosphate
The concentration of lipid phosphate was determined subsequent to total digestion by perchloric acid [48] using Na2HPO4 as standard. DMPC was used as an internal standard to assess lipid digestion. Samples without perchloric acid digestion produced negligible readings.

2.2.6. Fluorescence anisotropy measurements
Fluorescence anisotropy experiments were carried out using the fluorescent probe 1,6-diphenyl-1,3,5-hexatriene (DPH) as described previously [49]. Steady state fluorescence was measured in a Hitachi F-4010 spectrophotometer using 1 cm path length quartz cuvettes at room temperature (~23 °C). Excitation and emission wavelengths were set at 358 and 430 nm. Excitation and emission slits with bandwidths of 1.5 and 20 nm were used. The optical density of the samples measured at 358 nm was always less than 0.15. Fluorescence anisotropy measurements were performed using a Hitachi polarization accessory. Anisotropy (r) values were calculated from the equation [50]:

\[ r = \frac{I_{NW} - G_{HH}}{I_{NW} + 2G_{HH}} \]

where \( I_{NW} \) and \( I_{HH} \) are the measured fluorescence intensities (after appropriate background subtraction) with the excitation polarizer vertically oriented and the emission polarizer vertically and horizontally oriented, respectively. G is the grating correction factor and is the ratio of the efficiencies of the detection system for vertically and horizontally polarized light and is equal to \( h_{NW}/h_{HH} \). All experiments were done with multiple sets of samples and average values of fluorescence anisotropy are shown in Fig. 3.

2.2.7. Statistical analysis
Significance levels were estimated using Student’s two-tailed unpaired t-test using Graphpad Prism software version 4.0 (San Diego, CA).

3. Results and discussion
The enantiomer of cholesterol (ent-cholesterol) is the non-superimposable mirror image of native (natural) cholesterol (see Fig. 1b). Enantiomers have identical physicochemical properties (except for the direction of rotation of plane-polarized light). As a consequence, the membrane biophysical properties (such as compressibility and phase behavior) remain the same when native cholesterol is replaced with ent-cholesterol [40,51–53]. In addition, both native cholesterol and ent-cholesterol support normal growth of a mutant mammalian cell line [54]. ent-Cholesterol is often utilized to distinguish specific interaction of cholesterol from nonspecific effects [53,55–57]. ent-Cholesterol is a diastereomer of cholesterol in which only the orientation of the hydroxyl group at carbon-3 is inverted relative to native cholesterol (Fig. 1c). Previous studies have shown that the biophysical properties of epi-cholesterol and native cholesterol are different in membranes [40,53 and references therein]. For example, epi-cholesterol and native cholesterol have been reported to differ in their tilt angles, condensing ability, and phase transition properties in membranes [58–61].

Purified membrane proteins are ideally suited for studying lipid–protein interactions. However, purification of membrane proteins poses a considerable challenge. A necessary criterion for purification of an integral membrane protein is that the protein must be carefully removed from the native membrane and dispersed in solution. This process, termed solubilization, is most efficiently accomplished utilizing amphiphilic detergents [62,63]. In this process, proteins and lipids held together in native membranes are dissociated in the presence of a suitable detergent. This results in the formation of small protein and lipid clusters that remain dissolved (solubilized) in the aqueous solution. In our previous work, we partially purified the hippocampal serotonin1A receptor by solubilizing the receptor in a functionally active form using CHAPS, a synthetic zwitterionic detergent, which is mild and non-denaturing [44,64]. The solubilization conditions were highly optimized so as to prevent dissociation and depletion of trimeric G-proteins, which could result from high concentrations of CHAPS [65,66], and therefore helpful in effectively solubilizing GPCRs in a functionally active form. Hipppocampal membranes, solubilized this way, contain the serotonin1A receptor in a relatively pure (enriched) form. Interestingly, it has been previously shown by us [67] and others [68] that solubilization of the serotonin1A receptor by CHAPS leads to a reduction in membrane cholesterol and specific ligand binding to the receptor. More importantly, we previously demonstrated that upon replenishment of solubilized membranes with cholesterol, specific ligand binding of the serotonin1A receptor could be restored [67]. In this paper, we utilized this strategy of sterol replenishment to the solubilized receptor to explore the stereospecific stringency of cholesterol for receptor function utilizing stereoisomers of cholesterol (ent-cholesterol and epi-cholesterol).

Fig. 2 shows specific binding of the agonist [3H]8-OH-DPAT to serotonin1A receptors in solubilized hippocampal membranes, and upon replenishment of solubilized membranes with either epi-cholesterol or cholesterol. Specific [3H]8-OH-DPAT binding to native hippocampal membranes served as a control for these experiments. The figure shows that the specific [3H]8-OH-DPAT binding to the serotonin1A receptor is reduced upon solubilization to ~55% of the control (native membranes). We attribute this reduction in binding to the loss of membrane cholesterol accompanying solubilization [67]. Subsequent treatment of solubilized membranes with M|CD-cholesterol complex led to considerable recovery (~82%) of specific [3H]8-OH-DPAT binding, due to replenishment of cholesterol. Interestingly, replenishment of solubilized membranes with epi-cholesterol could not restore specific [3H]8-OH-DPAT binding to the receptor and remained at ~41% relative to control (native membranes). These results show that epi-cholesterol is unable to support the ligand binding function of the serotonin1A receptor.

In order to explore the enantioselectivity of cholesterol in its interaction with the serotonin1A receptor, we carried out replenishment of solubilized membranes with ent-cholesterol. As mentioned earlier, ent-cholesterol is often utilized to distinguish specific interaction of cholesterol from nonspecific effects [40,53,55–57]. The effect of replenishment of solubilized membranes with ent-cholesterol is shown in
Fig. 2. Effect of replenishment of cholesterol, epi-cholesterol (epi) and ent-cholesterol (ent) into solubilized membranes (SM) on specific binding of the agonist [3H]8-OH-DPAT to the serotonin1A receptor. Solubilized hippocampal membranes were replenished with cholesterol, epi-cholesterol or ent-cholesterol using 1 mM of respective steroid:10 mM M/CD complex. Values are expressed as percentages of specific binding obtained in native hippocampal membranes (HM). Data shown are means ± S.E. of at least four independent experiments (*, ** and *** correspond to significant (p < 0.05, p < 0.01 and p < 0.0001, respectively) difference in specific ligand binding to cholesterol, epi-cholesterol or ent-cholesterol-replenished membranes relative to solubilized membranes). See Materials and methods for other details.

Fig. 3. Effect of replenishment of cholesterol, epi-cholesterol (epi) and ent-cholesterol (ent) into solubilized membranes (SM) on steady state fluorescence anisotropy of the membrane probe DPH. Solubilized membranes were replenished with cholesterol, epi-cholesterol or ent-cholesterol using 1 mM of the respective steroid:10 mM M/CD complex. Fluorescence anisotropy measurements were performed with membranes containing 50 nmol phospholipid at a probe to phospholipid ratio of 1:180 (mol/mol); at room temperature (−23°C). Values represent means ± S.D. of duplicate points from at least four independent experiments (*** corresponds to p < 0.0001; the change in fluorescence anisotropy was tested against the corresponding value obtained with native hippocampal membranes (HM)). See Materials and methods for other details.
that the requirement of membrane cholesterol for the serotonin1A receptor function is diastereospecific, but not enantiospecific. We have previously shown that immediate biosynthetic precursors of cholesterol, differing with cholesterol in merely a double bond, were not able to support the function of the serotonin1A receptor [21,38,39,45]. In addition, we have shown that the serotonin1A receptor is more compact [76] and stable [49] in the presence of membrane cholesterol. We have very recently shown by coarse-grain molecular dynamics simulation that membrane cholesterol binds preferentially to certain sites on the receptor [77]. A prominent site among these is the cholesterol recognition/interaction amino acid consensus (CRAC) motif, recently identified by us in GPCRs [78].

We show here that a key structural feature of cholesterol for its ability to affect the function of the serotonin1A Receptor is the equatorial configuration of the 3-hydroxyl group. epi-Cholesterol, differing with cholesterol solely in the axial orientation of the 3-hydroxyl group, could not support receptor function, whereas ent-cholesterol which maintains the 3-hydroxy group in the equatorial configuration supports receptor function. Our present results therefore further extend the degree of specificity of the interaction between the serotonin1A receptor and membrane cholesterol. Yet, these results show that this specificity of interaction falls short of achieving enantioselectivity. We conclude that membrane lipid interactions of GPCRs could be of varying specificity and envisage that this type of regulated specificity affects the efficacy of the receptor–ligand interaction and is physiologically important. To the best of our knowledge, our results constitute the first report utilizing ent-cholesterol to explore the stereospecific requirement of cholesterol for GPCR function.

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