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## Contribution of the transmembrane domain 6 of melanocortin-4 receptor to peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity

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### Abstract

The melanocortin receptor (MCR) subtype family is a member of the GPCR superfamily and each of them has a different pharmacological profile regarding the relative potency of the endogenous and synthetic melanocortin peptides. Substitution of Trp with DNaI (2') in  $\gamma$ -MSH resulted in the loss of binding affinity and potency at hMC4R. However, the molecular mechanism of this ligand selectivity is unclear. In this study, we utilized chimeric receptors and site-directed mutagenesis approaches to investigate the molecular basis of MC4R responsible for peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity. Cassette substitutions of the second, third, fourth, fifth, and sixth TM of the human MC4R (hMC4R) with the homologous regions of hMC1R were constructed and the binding affinity of peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH at these chimeric receptors was evaluated. Our results indicate that the cassette substitutions of TM2, TM3, TM4 and TM5 of hMC4R with homologous regions of the hMC1R did not significantly increase peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity and potency but substitution of the TM6 of the hMC4R with the same region of the hMC1R significantly enhances [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity and potency. Further site-directed mutagenesis study indicates that four amino acid residues, Phe267, Tyr268, Ile269 and Ser270, in TM6 of the hMC4R may play an important role in [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selective activity at MC4R.

### Keywords

MC4R;  $\gamma$ -MSH; MC1R; Obesity; GPCR; Agonist

### 1. Introduction

The melanocortin system plays an important role in the regulation of animal physiology and pathophysiology [1–3]. This system consists of melanocortin peptides, melanocortin receptors (MCRs) and endogenous antagonists (agouti and agouti-related protein). The initial transcript of the pro-opiomelanocortin (POMC) gene is a large hormone precursor post-translationally processed into a number of biologically active peptides. These peptides include  $\alpha$ -,  $\beta$ -, and  $\gamma$ -melanocyte stimulating hormone (MSH), adrenocorticotrophic hormone (ACTH), the lipotropins, and  $\beta$ -endorphin. Melanocortin receptor subtype family is a member of the GPCR

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superfamily and each subtype has different physiological functions [4–6]. Each of the melanocortin receptor subtypes has a different pattern of tissue expression in the central nervous system and peripheral tissue and each receptor also has its own signature profile regarding the relative potency of the endogenous and synthetic melanocortin peptides. The MC1R is mainly expressed in melanocytes and monocytes and plays an important role in pigmentation and anti-inflammatory reactions [6]. The MC2R is mainly expressed in the adrenal gland and involved in adrenal steroidogenesis and adipocyte lipogenesis [7]. The MC3R and MC4R are expressed in the hypothalamus and play an important role in the regulation of food intake and body weight control [8,9]. The MC5R is expressed in the exocrine gland and is involved in the regulation of the exocrine gland function [4], Agouti and AGRP are endogenous antagonists of melanocortin receptors [10,11].

Energy homeostasis is a highly regulated process involving interacting signals between a variety of anorexigenic and orexigenic peptides, proteins and signaling molecules [12–15]. Genetic studies have highlighted the importance of the central melanocortin system in the control of mammalian energy homeostasis in mice and humans [16,17]. Mutations of the pro-opiomelanocortin gene, MC3R and MC4R gene result in an obese phenotype [18–20]. MC3R and MC4R are expressed in hypothalamic nuclei known to play different role in control of body weight [21–24]. MC3R knockout mice had normal food intake with increased body weight and body fat [23,24]. MC4R knockout mice were hyperphagia with increased body weight and fat mass [23]. Mutations of MC3R and MC4R have also been identified in human obese patients [25–29]. Mutations of hMC4R gene have been identified the most common cause of monogenic obesity because they are detected in almost 5–6% children with early onset severe obesity [30]. Over-expression of the melanocortin antagonists, agouti and AGRP, in transgenic animals also possesses an obese phenotype [31–34].

The melanocortin receptors consist of a single polypeptide featuring seven  $\alpha$ -helical transmembrane domains (TMs), an extracellular N-terminus, three extracellular loops, three intracellular loops and an intracellular C-terminus. Many structural features conserved in other G-protein-coupled receptors are found in the melanocortin receptors; the consensus N-linked glycosylation sites near the amino terminus, a palmitoylation site in the COOH-terminal tail, and sites for phosphorylation in the first and third intracellular domains and in the COOH-terminal tail [5,7,21]. All melanocortin receptors contain the conserved DRY motif at the C-terminal end of TM3 and contain a C-terminal cysteine that may function as a fatty acid acylation site. However, the melanocortin receptors lack several features found in most G-protein-coupled receptors; one, or two, cysteine residues in the first and second extracellular loops and prolines found in the fourth and fifth TMs.

All melanocortin peptide agonists contain a His-(L/D)Phe-Arg-Trp consensus sequence that is considered the “message” sequence, responsible for selectivity and activation of the melanocortin receptors [35,36]. The bioactive conformation of agonists peptides involves a  $\beta$ -turn on the His-(L/D)Phe-Arg-Trp motif [35,36] which is likely to serve as an organizing scaffold to orient the side chains of the message sequence in the best orientation for interaction with the receptor. Extensive structure-function studies on  $\alpha$ -MSH have created many new, more potent, and enzyme-resistant analogs of the melanocortin peptides which include synthetic agonists NDP-MSH [Nle<sup>4</sup>, DPhe<sup>7</sup>]- $\alpha$ -MSH and Melanotan II (MTII) and antagonists such as SHU9119 and HS204 [37–39]. Recently, it is reported that substitution of a bulky hydrophobic amino acid in Trp<sup>8</sup> of the  $\gamma$ -MSH with  $\beta$ -(2-naphthyl)-D-alanine (DNaI) ([Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]- $\gamma$ -MSH) significantly decreased agonist selectivity at MC4R [40]. However, the molecular basis of MC4R responsible for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity is unknown. In order to gain insights into the molecular determinants of the hMC4R responsible for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity, we have utilized chimeric receptor and site-directed mutagenesis approaches to investigate the role of the TMs of MC4R in [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -

MSH selectivity. The chimeric receptors and single amino acid mutated receptor of the MC4R were constructed and tested. Our results suggest that four amino acid residues of TM6 of MC4R are crucial for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity at MC4R.

## 2. Methods and materials

### 2.1. Reagents

[Nle<sup>4</sup>-D-Phe<sup>7</sup>]-MSH (NDP-MSH) was purchased from Peninsula Laboratories (Belmont, CA). 3-Isobutyl-methylxanthine (IBMX) from Sigma, and [<sup>125</sup>I] NDP-MSH from PerkinElmer Life Sciences (Boston, MA). The HEK-293 cell line was purchased from ATCC (Manassas, VA) and DMEM, lipofectamine from Life Technologies (Rockville MD). [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH was synthesized using method previous reported [40].

### 2.2. Characterization of the pharmacological profile of the peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSR

**2.2.1. Construction of the melanocortin chimeric receptors**—The amino acid sequences of the human (h) MC1R and human (h) MC4R were examined by hydrophobicity plot (Genetics Computer Group, Inc., Madison, WI) and manually by comparing their sequences to a previously published alignment of seven transmembrane G-protein coupled receptor  $\alpha$ -helices [41]. The chimeric receptors utilized in these studies are schematically diagramed in Fig. 1. The chimeric receptors were constructed by polymerase chain reaction (PCR) using Pfu polymerase (Stratagene, La Jolla, CA) [42]. The human MC4R served as template. During an initial round of PCR, partial-length receptor fragments were generated. The sequence of one of the PCR primer oligonucleotides consisted of the transmembrane domain of interest coupled to a portion of the extracellular domain required to form the chimeric receptor. The second oligonucleotide primer consisted of either the 5' or 3' end of the MC4R. Receptor fragments were separated by agarose gel electrophoresis and used in a second round of PCR in which full-length chimeric receptor constructs were assembled by cycling the appropriate fragments together for 10 cycles prior to adding both 5' and 3' receptor primers. The chimeric receptors were subcloned into the eukaryotic expression vector pcDNA 3.1 (Invitrogen, Carlsbad, CA). The entire coding region of the chimeric receptors was sequenced to confirm that the desired sequences will be present and that no sequence errors had been introduced by University of Alabama at Birmingham Sequence Core. The hMC4R/hMC1R (TM2) chimera encoded amino acids Met 1 to Ile83 and Ile103 to Tyr332 of hMC4R and amino acids Cys78 to Val97 of hMC1R. The hMC4R/hMC1R (TM3) chimera encoded amino acids Met 1 to Ile121 and Ile143 to Tyr332 of hMC4R and amino acids Asp117 to Ala137 of hMC1R. The hMC4R/hMC1R (TM4) chimera encoded amino acids Met 1 to Cys172 and Ser188 to Tyr332 of hMC4R and amino acids Ile168 to Tyr182 of hMC1R. The hMC4R/hMC1R (TM5) chimera encoded amino acids Met 1 to Val193 and Leu231 to Tyr332 of hMC4R and amino acids Leu189 to Val205 of hMC1R. The hMC4R/hMC1R (TM6) chimera encoded amino acids Met 1 to Ile251 and Cys271 to Tyr332 of hMC4R and amino acids Gly248 to Leu266 of hMC1R. The chimeric receptors were then subcloned into the eukaryotic expression vector pCDNA 3.1 (Invitrogen; Carlsbad, CA).

**2.2.2. Cell culture and transfection**—The HEK-293 cell line was utilized in this study. The cells were cultured in DMEM medium containing 10% bovine fetal serum and HEPES. Cells at 80% confluence were washed twice, and the receptor constructs were transfected into cells using lipofectamine (Life Technologies, Rockville MD). The permanently transfected clonal cell lines were selected by resistance to the neomycin analogue G418 [43].

**2.2.3. Binding assays**—Binding experiments were performed using the conditions previously described [44]. Briefly, after removal of the media, cells were incubated with non-radioligand [DPhe<sup>6</sup>]- $\gamma$ -MSH or [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH from 10<sup>-10</sup> to 10<sup>-6</sup> M in 0.5 mL

MEM containing 0.2% BSA and  $2 \times 10^5$  cpm of  $^{125}\text{I}$ -NDP-MSH for 1 h. The binding reactions were terminated by removing the media and washing the cells twice with MEM containing 0.2% BSA. The cells were then lysed with 0.2N NaOH, and the radioactivity in the lysate was quantified in an analytical gamma counter (PerkinElmer, Shelton, CT). Nonspecific binding was determined by measuring the amount of  $^{125}\text{I}$ -label bound on the cells in the presence of excess  $10^{-6}$  M unlabeled ligand. Specific binding was calculated by subtracting nonspecifically bound radioactivity from total bound radioactivity. Binding data are reported as  $\text{IC}_{50}$ .

**2.2.4. cAMP assay**—Cellular cAMP generation was measured using a competitive binding assay kit (TRK 432, Amersham, Arlington Heights, IL). Briefly, cell culture media was removed, and cells were incubated with 0.5 mL Earle's Balanced Salt Solution (EBSS), containing the melanocortin agonist  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  or  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  ( $10^{-10}$ – $10^{-6}$  M), for 1 h at  $37^\circ\text{C}$  in the presence of  $10^{-3}$  M isobutylmethylxanthine. The reaction was stopped by adding ice-cold 100% ethanol (500  $\mu\text{L}$ /well). The cells in each well were scraped, transferred to a 1.5-mL tube, and centrifuged for 10 min at  $1900 \times g$ , and the supernatant was evaporated in a  $55^\circ\text{C}$  water bath with pre-purified nitrogen gas. cAMP content was measured as previously described, according to instructions accompanying the assay kit [45].

**2.2.5. Receptor expression by using FACs**—To determine whether the receptor proteins are expressed at the cell surface, we generated a chimeric hMC4R with the N terminal fusion to the FLAG (Flag-hMC4R). The FLAG protein is an eight amino acid peptide (Asp-Tyr-Lys-Asp-Asp-Asp-Lys) and FLAG-tagged receptor expression vectors have been widely used to purify receptor protein and to determine the protein expression [47]. Cells transfected with the chimeric receptors were harvested using 0.2% EDTA and washed twice with phosphate buffer saline (PBS). Aliquots of  $3 \times 10^6$  cells were centrifuged and fixed with 3% paraformaldehyde in PBS (pH 7.4). The cells were incubated with 50  $\mu\text{L}$  of 10  $\mu\text{g}/\text{mL}$  murine anti-FLAG M1 monoclonal antibody (Sigma, Catalog No. 316) in incubation buffer for 45 min. Under this condition the primary antibody binds only to receptors located at the cell surface. The cells were collected by centrifugation and washed three times with incubation buffer. The cell pellets were suspended in 100  $\mu\text{L}$  of incubation buffer containing CY<sup>TM</sup>3-conjugated Affinity Pure Donkey Anti-Mouse Ig G (ImmunoResearch Lab, Inc., West Grove, PA) and incubated at room temperature for 30 min. Flow cytometry was performed on a fluorescence-activated cell sorter (Becton Dickinson FACStar plus six parameter cytometer/sorter with a dual Argon ion laser, San Jose, CA) [46]. The results were analyzed using the software CellQuest (Beckton-Dickinson Immunocytometry Systems, San Jose, CA).

### 2.3. Statistical analysis

Each experiment was performed in duplicate three separate times. The mean value of the dose-response data of binding and cAMP production was fit to a sigmoid curve with a variable slope factor using non-linear squares regression analysis (Graphpad Prism, Graphpad Software, San Diego, CA). Data are expressed as mean  $\pm$  S.E. Significant difference was assessed by one-way ANOVA with  $P < 0.05$  considered to be statistically significant.

## 3. Results

### 3.1. Peptide $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$ and $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$ binding and activity at MC4R and MC1R

Peptide  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  and  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  are  $\gamma\text{-MSH}$  analogues (Table 1). However, the binding affinities and potencies of these two analogues at MC1R and MC4R are different [40]. To determine  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  and  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  binding affinity at wild type hMC4R and hMC1R (hMC4R-WT and hMC1R-WT), the ability of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  and  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  to displace  $^{125}\text{I}$  NDP-MSH binding at hMC4R-WT and

hMC1R-WT were first examined as shown in Fig. 2. [DPhe<sup>6</sup>]- $\gamma$ -MSH dose-dependently displaces <sup>125</sup>I NDP-MSH binding at the hMC1R WT and hMC4R WT (Fig. 2A). To examine the ability of [DPhe<sup>6</sup>]- $\gamma$ -MSH to activate hMC1R or MC4R, [DPhe<sup>6</sup>]- $\gamma$ -MSH stimulated cAMP production was determined. Consistent with binding data, peptide [DPhe<sup>6</sup>]- $\gamma$ -MSH dose-dependently increased cAMP generation at the hMC1R-WT and hMC4R WT (Fig. 2B). [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH dose-dependently displaces <sup>125</sup>I NDP-MSH binding at the hMC1R WT and hMC4R-WT but binding affinity is significantly decreased at hMC4R (Fig. 2C). To examine the ability of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH to activate hMC1R or hMC4R, [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH stimulated cAMP production was determined. Consistent with binding data, peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH dose-dependently increased cAMP generation at the hMC1R-WT but its potency was significantly reduced at hMC4R. [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH becomes a partial agonist at hMC4R (Fig. 2D). Our results indicate that [DPhe<sup>6</sup>]- $\gamma$ -MSH is a non selective agonist for the MC1R and MC4R but [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH becomes a partial agonist at the MC4R which is consistent with the previous report [40]. Their K<sub>i</sub> and EC<sub>50</sub> are shown in Table 2.

### 3.2. Effects of substitutions of the transmembrane domain of the hMC4R with the corresponding regions of the hMC1R on [DPhe<sup>6</sup>]- $\gamma$ -MSH and (Pro<sup>5</sup>, DNaI (2')<sup>8</sup>)- $\gamma$ -MSH specific bindings and signaling

To investigate the molecular determinant of hMC4R responsible for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity, a domain-exchange strategy was used to localize regions of the hMC4R responsible for peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH activity. [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH is a full agonist at hMC1R but partial agonist at MC4R. The advantage of using the hMC4R as a template is that replacement of hMC4R TMs with the corresponding region of hMC1R may increase [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity and potency if the specific TM of the MC1R is involved in [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity. Cassette substitutions of the second, third, fourth, fifth, and sixth TMs of the hMC4R with homologous regions of the hMC1R were constructed. The first, and seventh TMs were not chosen for investigation because our previous data suggested that 1TM and 7TM of melanocortin receptors were not important in ligand binding [43,44,48]. In order to reduce the possibility of the receptor tertiary structure alteration by the entire TM domain exchange, upper half TM region of the hMC4R was replaced with the corresponding region of the hMC1R (Fig. 1).

To determine whether chimeric receptor proteins are expressed at the cell surface and to quantify receptor expression level, the antigenic epitope FLAG sequence was inserted into the NH<sub>2</sub> terminus of hMC4R-WT or chimeric receptors using polymerase chain reaction [47]. Our results indicate that the FLAG signal was detected by FACs at hMC4R-WT and chimeric receptors. The expression levels of all chimeric receptors were not significantly different from that of wild type receptor (Table 2).

To determine whether the receptor domain exchange alters receptor function [DPhe<sup>6</sup>]- $\gamma$ -MSH binding affinity and potency were evaluated at these chimeric receptors. Our results indicate that unlabelled [DPhe<sup>6</sup>]- $\gamma$ -MSH dose-dependently displaces <sup>125</sup>I-NDP-MSH binding at these chimeric receptors and all chimeric receptors possess high [DPhe<sup>6</sup>]- $\gamma$ -MSH binding affinity (Figs. 3 and 4Figs. 3A and 4A). Consistent with the binding data, our results also indicate that [DPhe<sup>6</sup>]- $\gamma$ -MSH dose dependently increased cAMP generation at these chimeric receptors (Figs.3B and 4B). The domain exchange did not significantly alter [DPhe<sup>6</sup>]- $\gamma$ -MSH potency and suggest that the tertiary structures of the chimeric receptors were not grossly disrupted and that the normal function of the receptor was retained. Their K<sub>i</sub> and EC<sub>50</sub> values are shown in Table 2. NDP-MSH has similar pharmacological profile at these chimeric receptors compared to that of [DPhe<sup>6</sup>]- $\gamma$ -MSH (data are not shown).

After confirming that substitution of the receptor domains did not alter receptor tertiary structure and function, we then investigated which region of the hMC4R is responsible for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selective binding and potency, HEK cells expressing these chimeric receptors were incubated with labeled <sup>125</sup>I NDP-MSH and various concentrations of unlabelled [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH and its binding affinity was assessed. Our results indicate that substitutions of TM2, TM3, TM4 and TM5 of the hMC4R with the corresponding regions of the hMC1R (hMC4R/TM2hMC1R, hMC4R/TM3hMC1R, hMC4R/TM4hMC1R and hMC4R/TM5hMC1R) did not significantly increase peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity (Figs. 3C and 4C). However, substitution of the hMC4R TM6 with the corresponding region of the hMC1R (hMC4R/TM6hMC1R) significantly increased peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity (Fig. 4C). The IC<sub>50</sub> values of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH are summarized in Table 2. To determine whether the substitutions of the hMC4R with the corresponding regions of the hMC1R will also increase [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH potency [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH stimulated cAMP production were determined. Our results indicate that hMC4R/TM2hMC1R, hMC4R/TM3hMC1R, hMC4R/TM4hMC1R and hMC4R/TM5hMC1R did not significantly increase [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH potency (Fig. 3D and 4D). However, substitutions of the hMC4R TM6 with the corresponding regions of the hMC1R (hMC4R/TM6hMC1R) did significantly increase peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH potency which is consistent with the binding results (Fig. 4D). Their K<sub>i</sub> and EC<sub>50</sub> are shown in Table 2.

### 3.3. Mutations of non-conserved amino acid residues in TM6 of the hMC4R for peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding and signaling

Substitution of TM6 of the hMC4R with the corresponding region of the hMC1R significantly increases peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity and potency suggest that non-conserved amino acid residues of the TM6 of the hMC4R may be involved in peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH specific binding. Five amino acid residues of the TM6 are identified to be non-conserved between MC4R and MC1R. To determine whether these amino acid residues are involved in peptide [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH specific binding and activity, these residues were individually mutated with the corresponding amino acid residues of hMC1R and evaluated. Our results indicate that these mutant receptors were expressed at cell surface and the expression levels showed no significant variation compared with that of wild type receptor (Table 3), The binding affinities and potencies of DPhe- $\gamma$ -MSH at these mutated receptors are similar to that of hMC4R (Fig. 5A and B). The binding affinity of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH at mutations, Ile266Ala, Phe267Leu, and Ser270Leu was also similar to that of the hMC4R-WT but the binding affinity of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH at mutations Tyr268Ile, Ile269Val was increased (Fig. 5C). Consistent with the binding results, the mutations, Ile266Ala, Phe267Leu, and Ser270Leu did not significantly increase [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH-mediated cAMP production but the mutations of Tyr268Ile, Ile269Val increased [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH potency (Fig. 5D). Their K<sub>i</sub> and EC<sub>50</sub> are shown in Table 3.

Single mutation of hMC4R did not dramatically increase [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity and potency, we therefore examined the combination of four amino acid mutations in TM6 on [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity. The multiple amino acid mutations, including Phe267Leu, Tyr268Ile, Ile269Val and Ser270Leu, were created and tested. Our results indicate that this multiple amino acid mutation of TM6 did result in an increase of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity at MC4R. The binding affinity of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH at this multiple mutations was significantly increased (Fig. 6A). Consistent with the binding results, this multiple mutations did significantly increase [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH-mediated cAMP production (Fig. 6B), suggesting that these four amino acid residues maybe involved in [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity.

## 4. Discussion

We have identified that several key amino acid residues of TM6 of MC4R are crucial for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity at the hMC4R by utilizing chimeric receptors and site-directed mutagenesis.

Melanocortin agonists bind in a  $\beta$ -turn conformation that organizes the message sequence (His-L/DPhe-Arg-Trp) in an optimal arrangement for binding and activation of the receptors. Extensive pharmacological studies have demonstrated that the tripeptide D-Phe-Arg-Trp in NDP-MSH is important not only for ligand binding but also for receptor activation at MC1R, MC3R and MC4R [44,49,50]. When Phe, Arg and Trp in MSH were substituted with alanine, the peptide binding affinities and potencies were significantly decreased at these receptors [51–53]. However, substitution of DPhe in position 7 of MTII with a D-(2'-naphthyl)-D-alanine (D-NaI (2')) (SHU9119) and Phe in position 6 of  $\gamma$ -MSH with DNaI(2') switches peptide from agonist to antagonist at MC3R and MC4R but retain agonist activity at MC1R and MC5R [39,40]. This information provides an initial estimate of the main contacts between peptides and receptor.

In the absence of a crystal structure of the MC4R, free or in complex with a ligand, little is known about the exact orientation of MC4 agonists in their binding pocket. Homology models of GPCRs based on bovine rhodopsin in combination with site-directed mutagenesis have been successfully used in the past to provide a structural framework for both ligand binding and functional studies. Accumulated evidence also indicates that TM3 of hMC3R and hMC4R was crucial for DNaI (2')7-SHU9119 ligand antagonist selectivity [50,54].

The Trp9 (super) residue of MSH was identified to be crucial for peptide binding affinity at hMC4R [52]. The Ala substitution resulted in a significant decrease of ligand binding affinity at hMC4R [44,55]. The bulky lipophilic amino acid derivative *c*-Hex Ala9 resulted in a large decrease in affinity but still retained efficacy [55], implying that the aromatic nature of Trp9- $\alpha$ -MSH is crucial for ligand binding affinity at hMC4R [44,50]. Pharmacological studies indicate that the core sequence His-Phe-Arg-Trp of  $\gamma$ -MSH is crucial for  $\gamma$ -MSH binding affinity and potency although  $\gamma$ -MSH has seven amino acid residues different from  $\alpha$ -MSH. The last four amino acids in the C-terminal region of  $\gamma$ -MSH are not important for its biological activity and selectivity at melanocortin receptors [56]. Substitution of the residue Trp8 of the  $\gamma$ -MSH with D-(2'-naphthyl)-D-alanine (D-NaI (2')<sup>8</sup>), [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH, significantly decreases its binding affinity and potency at MC4R [40]. Pogozheva et al. have modeled both the active and inactive states of hMC4R and propose that TMs movement accompany MC4R activation [57], TM6 shifts outward and undergoes counterclockwise movement, changing the position of TM6 into the binding pocket. Several residues of the hMC4R have been identified to be involved in NDP-MSH-induced receptor activation [55, 57,58]. However, substitution of Trp with DNaI (2) in  $\gamma$ -MSH [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH resulted in a significantly reduced MC4R activation but not for hMC1R. The aim of this study was to explore the molecular mechanism of MC4R responsible for this ligand selectivity using chimeric receptor and site-directed mutagenesis. The hMC4R and the hMC1R share a number of structural, functional and pharmacologic similarities. In the putative TM regions, the hMC4R and hMC1R show 67% identity. [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]- $\gamma$ -MSH is a potent agonist at MC1R but significantly decreased binding affinity and potency at MC4R. We speculate that the unique residues of the hMC4R might play an important role in this ligand selectivity. To determine which unique residue of the hMC4R is involved in this ligand selectivity, a chimeric receptor approach between hMC4R and hMC1R was utilized. We anticipate that the chimeric receptor using MC4R as a template will gain [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH binding affinity instead of using the chimeric receptor of the MC1R as a template losing peptide binding affinity. Our results demonstrate that the replacement of MC4R TM6 by MC1R TM6 results in increased

binding and potency of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH. This led us to the hypothesis that the D-(2'-naphthyl)-D-alanine (D-NaI (2')<sup>8</sup>) in [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH may interfere in the TM6 shift outward and counterclockwise movement, physically hindering the conformational changes necessary to induce receptor activation. Comparison of the amino acid sequences of TM6 between the hMC4R and the hMC1R revealed that five amino acid residues are different. They are residues, Ile266, Phe267, Tyr268, Ile269 and Ser270, which could potentially confer affinity characteristics of [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH. We examined the roles of these residues on ligand selectivity using a site directed mutagenesis study. However, our results indicate that single amino acid substitution of these residues did not significantly increase ligand efficacy, suggesting that multiple amino acid residues of TM6 may be important for this ligand selectivity. We then make a multiple amino acid mutation, including Phe267, Tyr268, Ile269 and Ser270 in TM6. These receptor residues at hMC4R were replaced with the corresponding residues of hMC1R and tested. Our results support our hypothesis that these residues are crucial for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity at MC4R. The replacements of these residues increased [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity at MC4R. The possible mechanism is that these mutations remove the steric hindrances of TM6 movement and induce receptor activation. However, substitution of TM6 of hMC4R with the corresponding region of hMC1R neither bring binding nor potency to the same level of hMC1R, suggesting that other areas close to this site may play a role and this is important for future potential investigation.

In conclusion, we have utilized chimeric receptor and site-directed mutagenesis approaches to identify critical amino acid residues of MC4R responsible for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity and potency. We believe that identification of the molecular basis of hMC4R responsible for different ligand-mediated receptor binding and signaling will provide valuable information of the molecular mechanism of MC4R in obesity development and provide important information for rational design of the MC4R potent and selective agonist which can be used for obesity treatment in the future. Our results provide the first evidence that the TM6 of the hMC4R is critical for [Pro<sup>5</sup>, DNaI (2')<sup>8</sup>]- $\gamma$ -MSH selectivity and potency.

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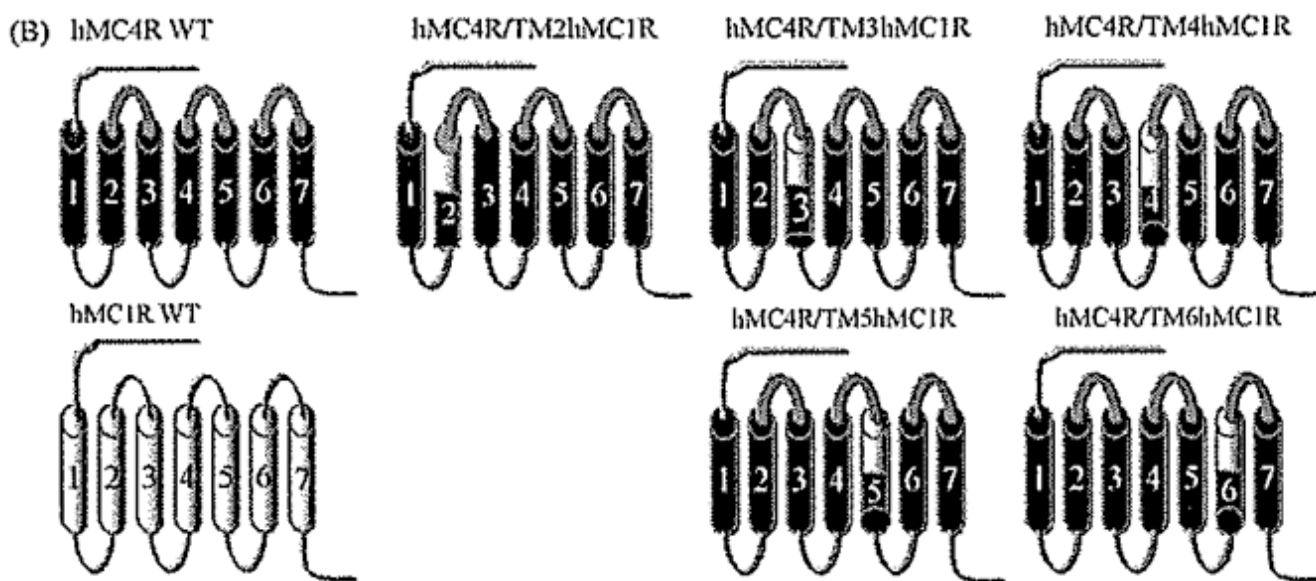


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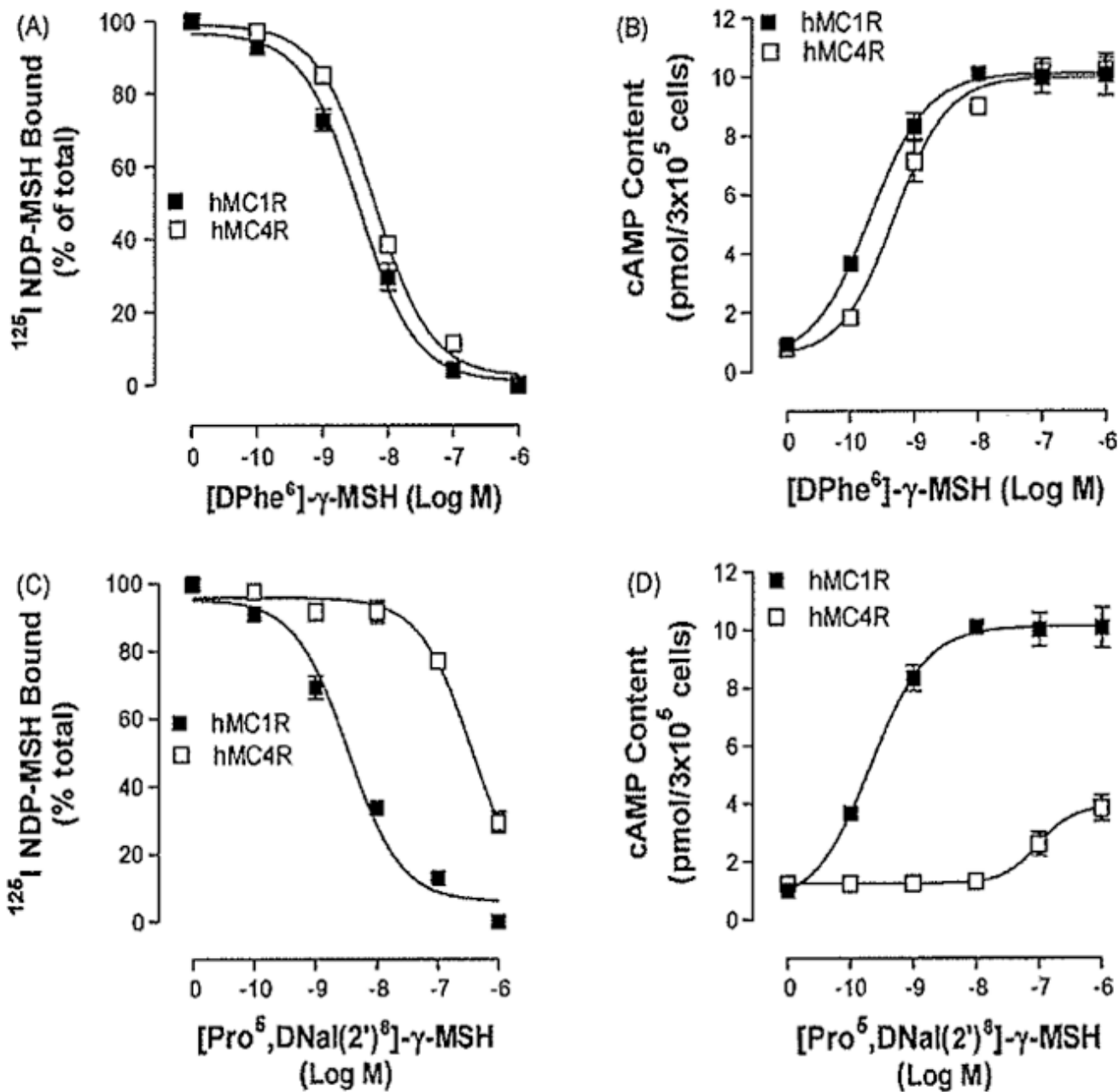
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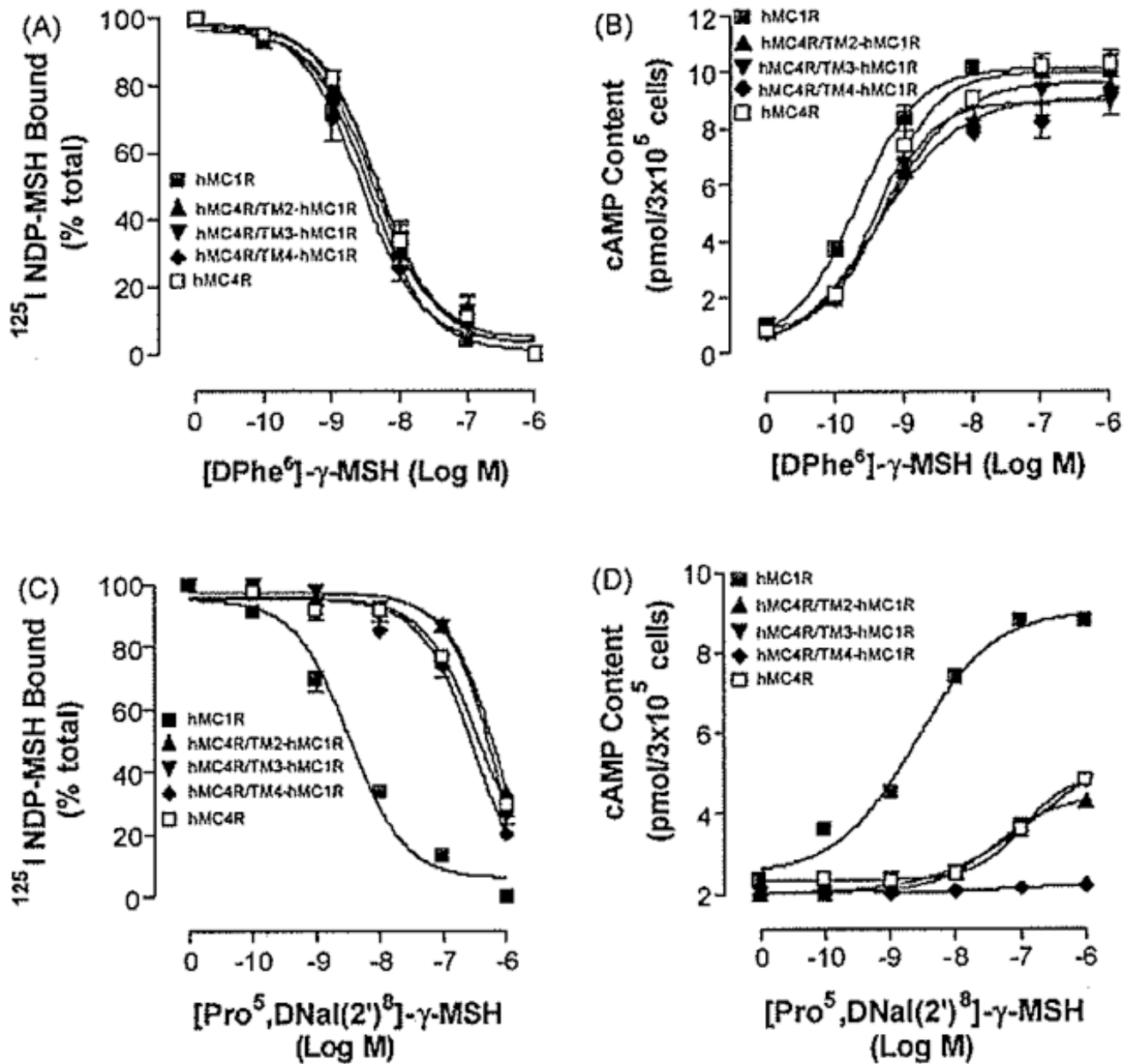
		<b>TM2</b>			
(A)	hMC1R	71	SPMYCFICCL	ALSDLLVSGT	NVLETA
	hMC4R	77	SPMYFFICSL	AVADMLVSVS	NGSETI
		<b>TM3</b>			
	hMC1R	117	DNVIDVITCS	SMLSSLCLFLG	AIAVDRY
	hMC4R	122	DNVIDSVICS	SLLASICSL	SIAVDRY
		<b>TM4</b>			
	hMC1R	159	PRARQAVAAI	WVASVVFSTL	FIAY
	hMC4R	164	KRVGIIISCI	WAACTVSGIL	FITY
		<b>TM5</b>			
	hMC1R	189	LLCLVVFLLA	MLVLMVLYV	HML
	hMC4R	194	IICLITMFFT	MLALMASLYV	HMF
		<b>TM6</b>			
	hMC1R	240	AVTLTILLGI	FFLCWGPFLL	HLTLIVL
	hMC4R	244	AITLTILIGV	FVVCWAPFLL	HLIFYIS



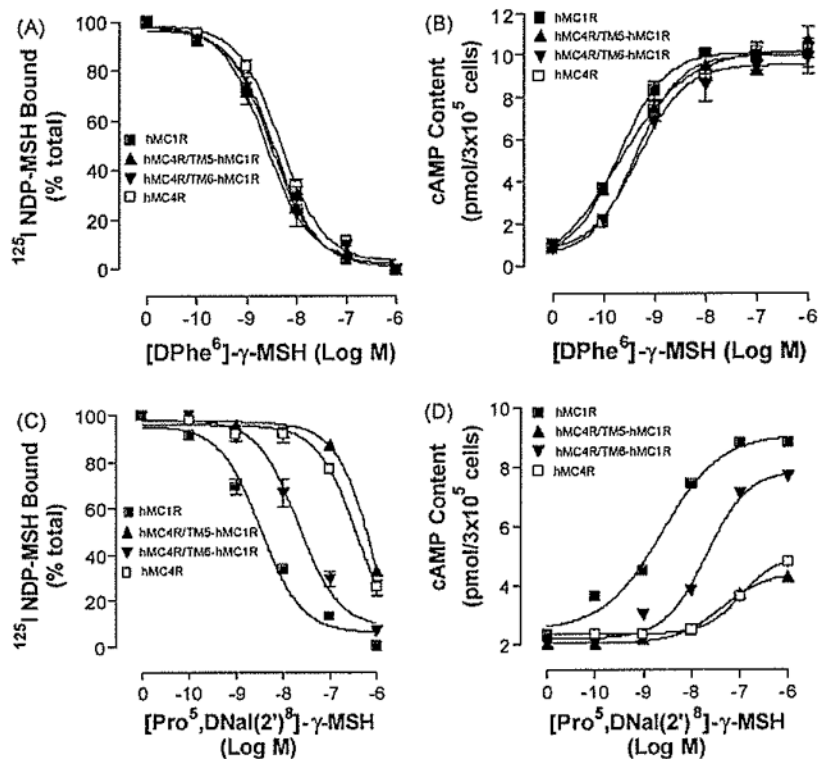
**Fig. 1.** The strategy of making chimeric receptors. Panel A represents the receptor sequences of the hMC4R and hMC1R. Panel B schematically depicts the seven transmembrane structures of the “wild type” (WT) MC4R (drawn with heavy lines) and MC1R (drawn with thin lines) and the structure of the chimeric MC4R with the substituted TMs of the MC1R. Bold font represents the dissimilar residues.



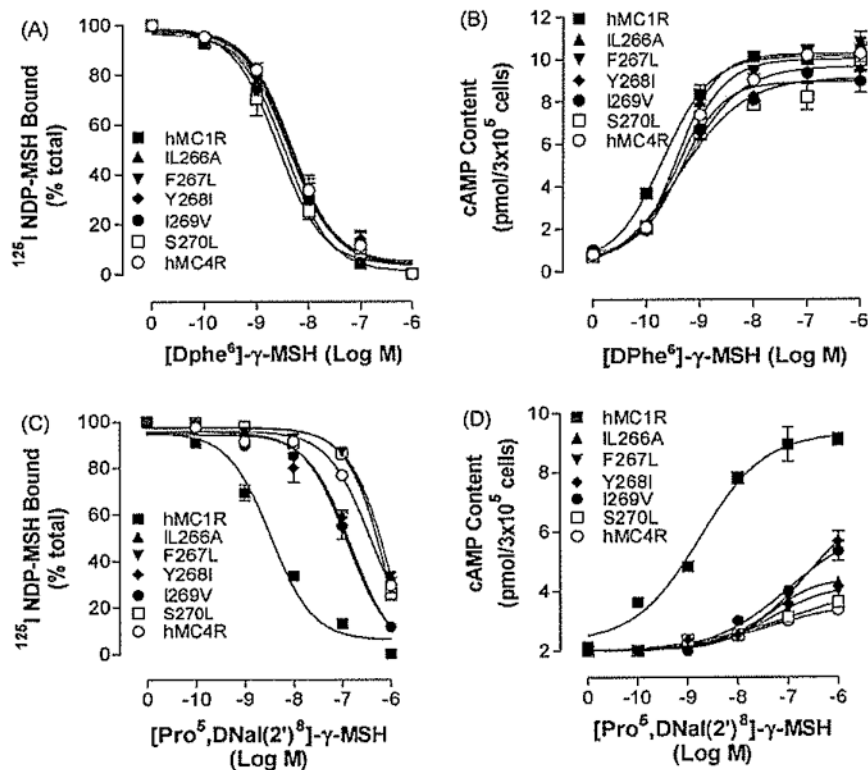
**Fig. 2.** Binding affinities and potencies of peptide [DPhe<sup>6</sup>]-γ-MSH and [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]-γ-MSH and NDP-MSH at the wild type hMC4R and hMC1R. Panel A depicts the binding affinity of [DPhe<sup>6</sup>]-γ-MSH as determined by inhibition of <sup>125</sup>I NDP-MSH binding. Panel B represents the ability of [DPhe<sup>6</sup>]-γ-MSH to stimulate the production of intracellular cAMP. Panel C depicts the binding affinity of [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]-γ-MSH as determined by inhibition of <sup>125</sup>I NDP-MSH binding. Panel D represents the ability of [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]-γ-MSH to stimulate the production of intracellular cAMP. Data points represent the mean ± S.E.M. of at least three independent experiments.



**Fig. 3.** Binding affinities and potencies of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  and  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  at hMC4R/TM2hMC1R, hMC4R/TM3hMC1R and hMC4R/TM3hMC1R. Panel A depicts the binding affinity of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  as determined by inhibition of  $^{125}\text{I}$ NDP-MSH binding at the chimeric receptors. Panel B represents the ability of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  to stimulate the production of intracellular cAMP at the chimeric receptors. Panel G depicts the binding affinity of  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  as determined by inhibition of  $^{125}\text{I}$ NDP-MSH binding at the chimeric receptors. Panel D represents the ability of  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  to stimulate the production of intracellular cAMP at the chimeric receptors. Data points represent the mean  $\pm$  S.E.M. of at least three independent experiments.

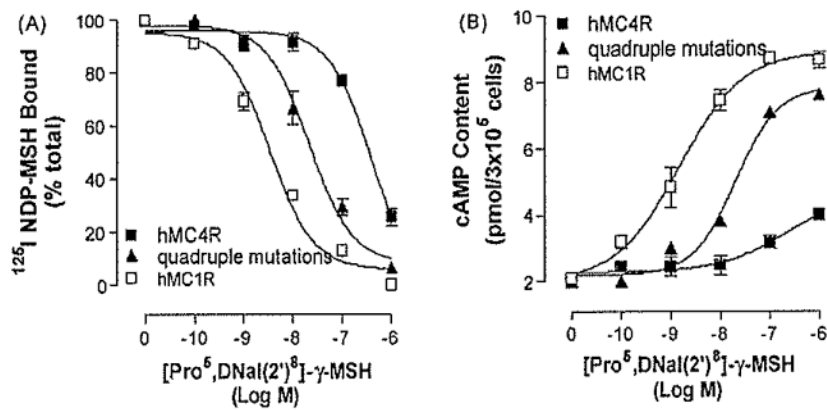


**Fig. 4.** Binding affinities and potencies of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  and  $[\text{Pro}^5, \text{DNAl}(2')^8]\text{-}\gamma\text{-MSH}$  at hMC4R/TM5hMC1R and hMC4R/TM6hMC1R. Panel A depicts the binding affinity of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  as determined by inhibition of  $^{125}\text{I}$ NDP-MSH binding at the chimeric receptors. Panel B represents the ability of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  to stimulate the production of intracellular cAMP at the chimeric receptors. Panel C depicts the binding affinity of  $[\text{Pro}^5, \text{DNAl}(2')^8]\text{-}\gamma\text{-MSH}$  as determined by inhibition of  $^{125}\text{I}$  NDP-MSH binding at the chimeric receptors. Panel D represents the ability of  $[\text{Pro}^5, \text{DNAl}(2')^8]\text{-}\gamma\text{-MSH}$  to stimulate the production of intracellular cAMP at the chimeric receptors. Data points represent the mean  $\pm$  S.E.M. of at least three independent experiments.



**Fig. 5.** Binding affinities and potencies of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  and  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  at mutation of the non-conserved TM6 amino acid residues of hMC4R, Panel A depicts the binding affinity of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  as determined by inhibition of  $^{125}\text{I}$  NDP-MSH binding at the mutated receptors. Panel B represents the ability of  $[\text{DPhe}^6]\text{-}\gamma\text{-MSH}$  to stimulate the production of intracellular cAMP at the mutated receptors. Panel C depicts the binding affinity of  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  as determined by inhibition of  $^{125}\text{I}$  NDP-MSH binding at the mutated receptors. Panel D represents the ability of  $[\text{Pro}^5, \text{DNaI}(2')^8]\text{-}\gamma\text{-MSH}$  to stimulate the production of intracellular cAMP at the mutated receptors. Data points represent the mean  $\pm$  S.E.M. of at least three independent experiments.





**Fig. 6.** Binding affinity and potency of [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]-γ-MSH at multiple mutations of hMC4R. Panel A depicts the binding affinity of [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]-γ-MSH as determined by inhibition of <sup>125</sup>I NDP-MSH binding at this mutated receptor. Panel B represents the ability of [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]-γ-MSH to stimulate the production of intracellular cAMP at the mutated receptor. Data points represent the mean ± S.E.M. of at least three independent experiments.

**Table 1**Sequences of  $\gamma$ -MSH, [DPhe<sup>6</sup>]- $\gamma$ -MSH, and [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]- $\gamma$ -MSH

Sequences of $\gamma$ -MSH analogues	
$\gamma$ -MSH	H-Tyr-Val-Nle-Gly-Pro-Phe-Arg-Trp-Asp-Arg-Phe-Gly-NH <sub>2</sub>
[DPhe <sup>6</sup> ]- $\gamma$ -MSH	H-Tyr-Val-Nle-Gly-Pro-DPhe-Arg-Trp-Asp-Arg-Phe-Gly-NH <sub>2</sub>
[Pro <sup>5</sup> , DNaI(2') <sup>8</sup> ]- $\gamma$ -MSH	H-Tyr-Val-Nle-Gly-Pro-Phe-Arg-DNaI(2')-Asp-Arg-Phe-Gly-NH <sub>2</sub>

The bold letters represent key residues for receptor binding and signaling

Effect of unlabeled ligands on  $^{125}\text{I}$ -NDP-MSH binding, at the HEK cells transfected with the chimeric hMC4 receptors. All determinations were on cells transfected with wild type or chimeric receptors. The receptor expression was determined by FLAG-tagged receptor expression and expressed as % of wild type receptor protein.  $K_i$  values for [DPhe<sup>6</sup>]- $\gamma$ -MSH and [Pro<sup>5</sup>, DNal(2')<sup>8</sup>]- $\gamma$ -were determined from displacement of  $^{125}\text{I}$ -NDP-MSH binding, as described under "Section 2". The data shown are mean  $\pm$  S.E.M of at least three independent experiments.

Table 2

	Receptor expression(%WT)	$^{125}\text{I}$ -NDP-MSH $K_i$ (nM)			cAMP production $EC_{50}$ (nM)	
		[DPhe <sup>6</sup> ]- $\gamma$ -MSH	[Pro <sup>5</sup> , DNal(2') <sup>8</sup> ]- $\gamma$ -MSH	[DPhe <sup>6</sup> ]- $\gamma$ -MSH	[Pro <sup>5</sup> , DNal(2') <sup>8</sup> ]- $\gamma$ -MSH	
hMC4RWT	100	4.1 $\pm$ 0.2	477 $\pm$ 19.4	35 $\pm$	partial agonist	
hMC4R/TM2 hMC1R	98 $\pm$ 12.2	3.7 $\pm$ 0.4	459 $\pm$ 11.5	2.9 $\pm$ 0.3	partial agonist	
hMC4R/TM3 hMC1R	92 $\pm$ 15.7	3.9 $\pm$ 0.5	506 $\pm$ 8.3	3.3 $\pm$ 0.5	partial agonist	
hMC4R/TM4 hMC1R	97.11.2 $\pm$	4.8 $\pm$ 0.6	421 $\pm$ 12.4	3.1 $\pm$ 0.3	partial agonist	
hMC4R/TM5 hMC1R	94 $\pm$ 16.4	2.9 $\pm$ 0.2	569 $\pm$ 13.2	1.7 $\pm$ 0.4	partial agonist	
hMC4R/TM6 hMC1R	96 $\pm$ 12.2	3.9 $\pm$ 0.4	79 $\pm$ 9.7*	3.2 $\pm$ 0.3	116 $\pm$ 65*	
hMC1RWT	100	2.5 $\pm$ 0.2	15.9 $\pm$ 3.8	2.3 $\pm$ 0.5	231 $\pm$ 8.2	

\*  $P < 0.05$

**Table 3**

Mutations of the non-conserved residues in TM6 of the hMC4R on DNaI(2')<sup>8</sup>- $\gamma$ -MSH binding and CAMP production. All determinations were on cells transfected with wild type or mutated receptors.  $K_i$  values for [Pro<sup>5</sup>, DNaI(2')<sup>8</sup>]- $\gamma$ -MSH were determined from displacement of <sup>125</sup>I-NDP-MSH binding, as described under "Section 2". The data shown are mean  $\pm$  S.E.M of at least three independent experiments.

	Receptor expression	[Pro <sup>5</sup> , DNaI(2') <sup>8</sup> ]- $\gamma$ -MSH		
		$K_i$ (nM)	EC <sub>50</sub> (nM)	E <sub>max</sub> (%)
hMC1RWT	100	2.1 $\pm$ 0.1	0.3 $\pm$ 0.02	100
I266T	95 $\pm$ 15.4	133.3 $\pm$ 18.2	Partial agonist	33
F267L	96 $\pm$ 12.2	169.3 $\pm$ 21.2	Partial agonist	32
Y268I	94 $\pm$ 3.5	154.2 $\pm$ 11.1	Partial agonist	45
I269V	92 $\pm$ 7.7	212.9 $\pm$ 30.3	Partial agonist	51
S270L	93 $\pm$ 8.1	124.3 $\pm$ 12.5	Partial agonist	30
hMC4-WT	100	126.6 $\pm$ 13.1	Partial agonist	35