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### **Extracellular Loop 3 (EL3) and EL3-Proximal Transmembrane Helix 7 of the Mammalian Type I and Type II GnRH Receptors Determine Differential Ligand Selectivity to GnRH-I and GnRH-II**

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**ABBREVIATIONS:** GnRH, gonadotropin-releasing hormone; gmGnRHR-2, green monkey type II GnRH receptor; GPCR, G protein-coupled receptor

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### ABSTRACT

Mammalian type I and type II GnRH receptors (GnRHRs) show differential ligand preference for GnRH-I and GnRH-II, respectively. Using a variety of chimeric receptors based on green monkey GnRHR-2 (gmGnRHR-2), a representative type II GnRHR, and rat GnRHR, a representative type I GnRHR, the present study elucidated specific domains responsible for this ligand selectivity. A chimeric gmGnRHR-2 with the extracellular loop 3 (EL3) and EL3-proximal transmembrane helix 7 (TMH7) of rat GnRHR showed a great increase in ligand sensitivity to GnRH-I but not to GnRH-II. Point mutation studies indicate that four amino acids, Leu/Phe<sup>7.38</sup>, Leu/Phe<sup>7.43</sup>, Ala/Pro<sup>7.46</sup>, and Pro/Cys<sup>7.47</sup> in TMH7 are critical for ligand selectivity as well as receptor conformation. Further, a combinatory mutation (Pro<sup>7.31</sup>-Pro<sup>7.32</sup>-Ser<sup>7.33</sup> motif to Ser-Glu-Pro in EL3 and Leu<sup>7.38</sup>, Leu<sup>7.43</sup>, Ala<sup>7.46</sup>, and Pro<sup>7.47</sup> to those of rat GnRHR) in gmGnRH-2 exhibited an approximately 500-fold increased sensitivity to GnRH-I, indicating that these residues are critical for discriminating GnRH-II from GnRH-I. [Trp<sup>7</sup>]GnRH-I and [Trp<sup>8</sup>]GnRH-I but not [His<sup>5</sup>]GnRH-I exhibit a higher potency in activating wild type gmGnRHR-2 than native GnRH-I, indicating that amino acids at positions 7 and 8 of GnRHs are more important than position 5 for differential recognition by type I and type II GnRHRs. Collectively, this study suggests a molecular coevolution of ligands and their receptors and facilitate the understanding of the molecular interaction between GnRHs and their cognate receptors.

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Gonadotropin-releasing hormone receptor (GnRHR), a rhodopsin-like G protein coupled receptor (GPCR) is one of the most extensively studied receptors due to its dual significance both for understanding reproductive biology and for the development of medical therapies (Sealfon et al., 1997). It is now well-established that most vertebrates including human have at least two forms of GnRH (White et al., 1998; Fernald and White, 1999). One form (GnRH-I, also called mammalian GnRH) is primarily synthesized in the hypothalamus, while the other form (GnRH-II, also called chicken GnRH-II) is widely expressed in the brain and peripheral tissues. While GnRH-I is known to regulate the secretion and synthesis of gonadotropins in the pituitary, the exact function of GnRH-II is largely unknown. The receptor for GnRH-I was first isolated from mammalian pituitary cells (Reinhart et al., 1992; Tsutsumi et al., 1992; Kaiser et al., 1992), and called mammalian type I GnRHR. Recently, receptors having a high affinity for GnRH-II were identified in nonmammalian and mammalian species (Tensen et al., 1997; Illing et al., 1999; Wang et al., 2001; Millar et al., 2001; Neill et al., 2001; Bogerd et al., 2002; Seong et al., 2003). Mammalian type II GnRHR is closer in structure to nonmammalian GnRHRs than mammalian type I GnRHR. Mammalian type II GnRHR, like nonmammalian GnRHRs, contains the intracellular C-terminal tail which is functionally important for desensitization and internalization (Heding et al., 1998; Willars et al., 1999), while mammalian type I GnRHR does not have a C-terminal tail. Mammalian type II and nonmammalian GnRHRs have Asp<sup>2.50</sup> and Asp<sup>7.49</sup> in the transmembrane helices (TMHs) 2 and 7, respectively, while mammalian type I GnRHRs contain Asp<sup>2.50</sup> and Asn<sup>7.49</sup> which are known to be important for receptor conformation and signal transduction (Blomenröhr et al., 1997; Mitchell et al., 1998).

Mammalian type II GnRHR has a higher affinity for GnRH-II than GnRH-I, while

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the opposite is true for mammalian type I GnRHR. However, the factors that determine such differential ligand selectivity are poorly understood. Mutagenesis studies combined with computational modeling have identified a number of residues which are involved in ligand binding (Davidson et al., 1996; Flanagan et al., 2000; Hoffmann et al., 2000; Hövelmann et al., 2002). GnRH-II differs from GnRH-I by 3 amino acids at positions 5, 7, and 8, thus, searching for residues that may interact with them would help us to understand the mechanism underlying differential ligand selectivity. Recently, it was proposed that Tyr<sup>5</sup> and Leu<sup>7</sup> of GnRH-I interact with Tyr<sup>6,58</sup> and Trp<sup>2,64</sup> of mammalian type I GnRHR (Hövelmann et al., 2002). However, as Tyr<sup>6,58</sup> and Trp<sup>2,64</sup> are also conserved in mammalian type II GnRHR, these residues alone cannot account for differential ligand selectivity. An acidic amino acid, Glu/Asp<sup>7,32</sup> in EL3 of mammalian type I GnRHR is known to confer ligand specificity for GnRH-I by an electrostatic interaction with Arg<sup>8</sup> of GnRH-I (Flanagan et al., 1994; Fromme et al., 2001). However, this is not fully explanatory as some nonmammalian GnRHRs have an acidic amino acid (e.g. Glu<sup>7,32</sup> for bfGnRHR-2 and Asp<sup>7,32</sup> for catfish GnRHR) at this homologous position, yet these receptors respond better to GnRH-II than GnRH-I (Wang et al., 2001). Recently, we demonstrated that the positions of Ser and Pro flanking Glu/Asp<sup>7,32</sup> are critical determinants for ligand selectivity (Wang et al., 2004). Replacement of the Ser-Glu-Pro (SEP) motif by Pro-Glu-Ser (PES) in mammalian type I GnRHR induced an increased sensitivity to GnRH-II but the opposite to GnRH-I. Moreover, mutation of a Ser-Gln-Ser (SQS) motif to SEP in bullfrog type I GnRHR (bfGnRHR-1) showed an increased sensitivity to GnRH-II but a decreased sensitivity to GnRH-I (Wang et al., 2004). However, this study found no reversed ligand selectivity when the Pro-Glu-Tyr (PEY) motif in bfGnRHR-2 was replaced by SEP, suggesting the involvement of other

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residues in ligand selectivity.

Sequence alignments showed that the EL3-proximal TMH7 of mammalian type II GnRHR has a high degree of sequence identity with that of nonmammalian GnRHRs but not with that of mammalian type I GnRHR. In the present study, using rat GnRHR and green monkey (gm) GnRHR-2 as models for representative mammalian type I and type II GnRHRs, respectively, we addressed whether EL3 and/or EL3-proximal TMH7 determine differential ligand selectivity. Domain swapping and site-directed mutagenesis studies suggest that the Pro-Pro-Ser (PPS) motif in EL3 and Leu<sup>7.38</sup>, Leu<sup>7.43</sup>, Ala<sup>7.46</sup>, and Pro<sup>7.47</sup> in TMH7 of gmGnRHR-2 are critical for discriminating GnRH-II from GnRH-I.

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### Materials and Methods

**Materials.** GnRH-I (pyroGlu-His-Trp-Ser-Tyr-Gly-Leu-Arg-Pro-GlyNH<sub>2</sub>), GnRH-II ([His<sup>5</sup>, Trp<sup>7</sup>, Tyr<sup>8</sup>]GnRH-I), [His<sup>5</sup>]GnRH-I, [Trp<sup>7</sup>]GnRH-I, [Trp<sup>8</sup>]GnRH-I, and [Trp<sup>7</sup>, Leu<sup>8</sup>]GnRH-I were synthesized by AnyGen (Gwangju, Korea). The c-fos-luc vector containing -711 ~ +45 sequence of the human c-fos promoter constructed in the pFLASH vector, was a kind gift from Dr. R. Prywes, Columbia University, NY. Vent DNA polymerase was purchased from New England Biolab (Beverly, MA). All oligonucleotides were synthesized from GenoTech (Daejeon, Korea). GH<sub>3</sub> cell lines stably expressing gmGnRHR-2 or rat GnRHR were established as described previously (Acharjee et al., 2002; Wang et al., 2003).

**Amino Acid Residue Numbering Scheme.** Amino acid residues are numbered according to their positions in gmGnRHR-2. To facilitate the comparison among different GnRHRs, the standard numbering system proposed by Ballesteros and Weinstein (Ballesteros and Weinstein, 1995) was also used.

**Construction of Wild Type and Mutant GnRHRs.** The cDNA of gmGnRHR-2 subcloned into pcDNA3 (Invitrogen, San Diego, CA) at the *Kpn*I and *Xba*I sites (Wang et al., 2003) was used as a template for creating domain-swapped or site-directed mutants. Domain swapping and site-directed mutagenesis were performed by the PCR overlapping extension method (Wang et al., 2003; Wang et al., 2004). To facilitate the construction of domain-swapped mutants, an exogenously introduced *Eco*RV site at the Asn<sup>7.34</sup> residue and an intrinsic *Bst*XI site or two intrinsic *Bam*HI sites were used. EL3 or EL3-proximal TMH7 of rat GnRHR was amplified using a specific set of primers flanked by the overlapping sequence of gmGnRHR-2 and the appropriate restriction

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endonuclease recognition site, producing rEL3S and rEL3L, respectively. The fragment from the N-terminal to the *EcoRV* site at the Asn<sup>7.34</sup> residue of rEL3S was replaced by the corresponding fragment of rat GnRHR, generating the r6TM chimera. Similarly, the fragment from the N-terminal to the *BamHI* site at the Pro<sup>7.47</sup> residue of gmGnRHR-2 was replaced by the corresponding part of rat GnRHR, producing the r6.5TM chimera. Mutated sequences were confirmed using the Sequenase Version 2.0 DNA Sequencing Kit (USB Corporation, Cleveland, OH) according to the manufacturer's instructions.

**Inositol Phosphate Production Assay.** The inositol phosphate (IP) production assay was performed as previously described (Wang et al., 2003). GH<sub>3</sub> cells ( $1 \times 10^5$  per well) expressing gmGnRHR-2 or rat GnRHR were seeded in 12-well plates and the following day cells were incubated in inositol-free DMEM (Life Technologies, Inc., Rockville, MD) containing 2% dialyzed fetal bovine serum (FBS) and labeled with 1  $\mu$ Ci *myo*-[<sup>3</sup>H]inositol/well (Amersham Pharmacia Biotech, Buckinghamshire, UK) for 18 h. Medium was then removed and cells were washed with 0.5 ml buffer A (140 mM NaCl, 20 mM HEPES, 4 mM KCl, 8 mM D-Glucose, 1 mM MgCl<sub>2</sub>, 1 mM CaCl<sub>2</sub>, 1 mg/ml fatty acid-free BSA). Cells were then preincubated with buffer A containing 10 mM LiCl for 15 min, followed by treatment with graded concentrations (0.01 nM - 10  $\mu$ M) of GnRHs at 37°C for 45 min. The reaction was terminated by removing the incubation medium and adding 0.5 ml of ice-cold 10 mM formic acid. After 30 min at 4°C, the formic acid extracts were transferred into columns containing Dowex anion exchange resin. Total IPs were then eluted with 1 ml of 1 M ammonium formate/0.1 M formic acid, and their radioactivity was determined.

**Luciferase Assay.** Wild type and mutant GnRHRs were transiently transfected into CV-1 cells, which were maintained at 37°C in DMEM with 10% heat-inactivated FBS,



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1 mM glutamate, 100 U of penicillin, and 100 µg/ml streptomycin. Cells were seeded in 24-well plates ( $1 \times 10^5$  per well) and transfection was performed using the SuperFect transfection kit (QIAGEN, Chatsworth, CA) according to the manufacturer's instructions with a minor modification. For each transfection, 100 ng of each receptor cDNA, 200 ng of c-fos-luc vector along with 200 ng of internal control plasmid pCMVβ-Gal were used. One day after transfection, cells were serum-starved for 24 h, and then challenged with GnRH for 6 h (Oh et al., 2003). Cells were harvested and luciferase activity in the cell extract was determined according to standard methods in a Lumat LB9501 (EG & G Berthold, Bad Wildbad, Germany). The luciferase activities were normalized using β-Gal values. Transfection experiments were performed in duplicate and repeated three to five times.

**Binding Assay.** GnRH-II was radioiodinated using the chloramine-T method and purified by chromatography on a Sephadex G-25 (Sigma, St. Louis, MO) column in 0.01 M acetic acid, 0.1% BSA. HeLa cells were transfected with wild type, individual mutant construct, or pcDNA3 (300 ng of DNA/well in 12-well plates) with Effectene (QIAGEN) according to the manufacturer's instructions. Thirty-six hours after transfection, intact cells were washed and incubated with binding buffer (DMEM supplemented with 0.1% BSA, pH 7.4) containing 250,000 cpm  $^{125}$ I-GnRH-II (0.5 ml final volume) at 20°C for 1 h to achieve equilibrium. Specific binding was calculated by subtracting nonspecific binding (presence of 10 µM unlabelled GnRH-II) from total binding. For the displacement binding assay,  $^{125}$ I-GnRH-II was incubated in the presence of graded concentrations of cold GnRH-I or GnRH-II.

**Molecular Modeling.** gmGnRHR-2 was built by MODELLER 6v2 (Sali and Blundell, 1993) based on the crystal structure of bovine rhodopsin (Okada et al., 2002)

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as a template. GnRH-I or GnRH-II was docked onto putative binding sites of gmGnRHR-2 manually using Visual Molecular Dynamics (Humphrey et al., 1996): PyroGlu<sup>1</sup> with Asn<sup>5,39</sup>, His<sup>2</sup> with Asp<sup>2,61</sup>, Trp<sup>3</sup> with Asn<sup>6,48</sup>, Tyr<sup>5</sup> with Tyr<sup>6,58</sup>, Gly<sup>10</sup> with Asp<sup>2,61</sup> and Asn<sup>2,65</sup>. The models for gmGnRHR-2/GnRH-II, mutant gmGnRHR-2/GnRH-I, and mutant gmGnRHR-2/GnRH-II were built by mutating corresponding residues in the gmGnRHR-2/GnRH-I model and underwent energy minimization and molecular dynamics annealing simulations in the MODELLER. The final models showing good geometry were confirmed by PROCHECK (Laskowski et al., 1993). The contacts between ligands and receptors were analyzed using Ligplot (Wallace et al., 1995). Figures of the models were drawn using Visual Molecular Dynamics (Humphrey et al., 1996).

**Data Analysis.** Analyses were performed using nonlinear regression, and the data were expressed as sigmoid dose-response curves. GnRH concentrations inducing half-maximal stimulation ( $EC_{50}$ ), half maximal inhibition ( $IC_{50}$ ), and maximal fold increases ( $E_{max}$ ) were calculated using GraphPad PRISM3 software (GraphPad, San Diego, CA). All data are presented as mean  $\pm$  SEM. The data were analyzed by one-way ANOVA followed by the Bonferroni Test.  $p < 0.05$  was considered statistically significant.

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### Results

**Differential Ligand Selectivity of Mammalian and Nonmammalian GnRHRs.** The ligand selectivities of rat GnRHR and gmGnRHR-2 were examined using two different methods: IP production and c-fos promoter-driven luciferase (c-fos-luc) assays. For the IP assay, GH<sub>3</sub> cells stably expressing rat GnRHR or gmGnRHR-2 were used (Wang et al., 2003; Maiti et al., 2003), and for c-fos-luc assay, CV-1 cells transiently expressing rat GnRHR or gmGnRHR-2 were used. As for rat GnRHR, GnRH-I showed a lower EC<sub>50</sub> than GnRH-II, indicating that rat GnRHR has a higher sensitivity to GnRH-I than GnRH-II. However, gmGnRHR-2 responded better to GnRH-II than GnRH-I in both assay systems (Fig. 1 and Table 1). Regarding GnRHR, GnRH-I had a 7.4- and 5.9-fold higher potency than GnRH-II in IP and c-fos-luc assay systems, respectively. For gmGnRHR-2, GnRH-II was 204-fold (IP assay) and 239-fold (c-fos-luc assay) more potent than GnRH-I (Table 1). Since c-fos-luc was more sensitive than the IP assay system, we used the c-fos-luc system in ensuing experiments.

**EL3 and EL3-proximal TMH7 Are Involved in Differential Ligand Selectivity.** Sequence alignment showed that EL3-proximal TMH7 of mammalian type II GnRHR has a high degree of sequence identity with that of nonmammalian GnRHRs but not mammalian type I GnRHR. Further, it was suggested that the proximal region of TMH7 of GnRHR affects the conformation of EL3 (Petry et al., 2002). We therefore presumed that both EL3 and the EL3-proximal TMH7 may be involved in differential ligand selectivity. To address this possibility, EL3 alone or together with EL3-proximal TMH7 of gmGnRHR-2 was swapped with that of rat GnRHR, designated rEL3S or rEL3L, respectively. Swapping EL3 alone did not induce a significant change in sensitivity for

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either GnRH-I or GnRH-II such that chimeric rEL3S, like wild type gmGnRHR-2, showed a higher sensitivity to GnRH-II than GnRH-I (Fig. 2A). Interestingly, rEL3L showed a great increase in sensitivity to GnRH-I but not to GnRH-II (Fig. 2B and Table 2), indicating that EL3-proximal TMH7 in gmGnRHR-2 is likely important for the discrimination between them. The functional importance of EL3-proximal TMH7 was further confirmed by additional chimeric receptors that have N-terminals to EL3 or to EL3-proximal TMH7 of rat GnRHR, denoted r6TM or r6.5TM, respectively. The chimeric receptor r6TM, which has the EL3-proximal TMH7 sequence of gmGnRHR-2, has a high sensitivity to GnRH-II and a low sensitivity to GnRH-I, characteristics of a type II receptor (Fig. 2C). In contrast, r6.5TM containing the EL3-proximal TMH7 sequence of rat GnRHR has the ligand sensitivity, characteristics of a type I GnRHR (Fig. 2D), again confirming that EL3-proximal TMH7 in gmGnRHR-2 is critical for ligand selectivity.

**Identification of rEL3L Amino Acids Involved in Ligand Selectivity.** As sequence alignment showed a six-amino acid difference in EL3-proximal TMH7 between gmGnRHR-2 and rat GnRHR (Fig. 2E), we postulated that one of them may be responsible for ligand selectivity. Thus, six individual amino acids were reciprocally changed in the rEL3L chimeric receptor. Point mutation of Phe<sup>7.37</sup> to Ile, Phe<sup>7.38</sup> to Leu, or Ala<sup>7.42</sup> to Gly did not induce significant changes in ligand selectivity compared to rEL3L (Fig. 3A-B and Table 2). Point mutation of Pro<sup>7.46</sup> to Ala or Cys<sup>7.47</sup> to Pro completely suppressed receptor function in response to either GnRH-I or GnRH-II (Fig. 3C and D). Finally, the mutation of Phe<sup>7.43</sup> to Leu in rEL3L showed a significant decrease in sensitivity to GnRH-I (Fig. 3A).

It was previously shown that Glu<sup>7.32</sup> of mouse GnRHR is a critical residue conferring

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ligand specificity for Arg<sup>8</sup> of GnRH-I (Flanagan et al., 1994). Further, we recently demonstrated that the positions of Ser and Pro flanking Glu<sup>7.32</sup> are crucial for the ligand selectivity between mammalian and nonmammalian GnRHRs (Wang et al., 2004). Therefore, we sought to determine whether these amino acids are critical for the ligand selectivity in the chimeric receptor rEL3L. The Glu<sup>7.32</sup> or Ser<sup>7.31</sup>-Glu<sup>7.32</sup>-Pro<sup>7.33</sup> (SEP) motif in rEL3L was changed to Gln or Pro-Pro-Ser (PPS), respectively. A mutation of Glu<sup>7.32</sup> to Gln significantly decreased sensitivity to GnRH-I but not to GnRH-II (Fig. 3C-D and Table 2). Moreover, replacement of the SEP motif by PPS greatly increased sensitivity to GnRH-II, while it slightly decreased sensitivity to GnRH-I (Fig. 3C-D and Table 2). These results suggest that the SEP/PPS motif together with amino acids in TMH7 is important in ligand sensitivity.

**Identification of Amino Acids in Wild Type gmGnRHR-2 That Confer Differential Ligand Selectivity.** As we observed that the amino acid residues Phe<sup>7.43</sup>, Pro<sup>7.46</sup>, and Cys<sup>7.47</sup> in rEL3L critically affected receptor activation and ligand selectivity, we further examined the function of these residues in wild type gmGnRHR-2. Two mutants, L<sup>7.43</sup>F and A<sup>7.46</sup>P had no receptor activity (Fig. 4). The mutant P<sup>7.47</sup>C had essentially the same ligand selectivity as the wild type gmGnRHR-2 (Fig. 4). As we failed to observe reversed ligand selectivity by a single mutation, we postulated that multiple amino acids are involved in ligand selectivity. To address this, double or triple mutants with different combinations of Leu<sup>7.43</sup>, Ala<sup>7.46</sup>, and Pro<sup>7.47</sup> were constructed. Interestingly, a double mutant A<sup>7.46</sup>P/P<sup>7.47</sup>C exhibited an improvement in ligand selectivity for both GnRH-I and GnRH-II (Fig. 5A-B). The double mutant L<sup>7.43</sup>F/P<sup>7.47</sup>C showed a decrease in receptor efficacy in both GnRH-I and GnRH-II but showed a slight increase in sensitivity for GnRH-I (Fig. 5A-B). The double mutant L<sup>7.43</sup>F/A<sup>7.46</sup>P

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did not respond to GnRH stimulation (Fig. 5A-B). A triple mutant L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C exhibited a large increase in ligand sensitivity for both GnRH-I and GnRH-II. As compared to wild type gmGnRHR-2, the L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant showed a ~ 200-fold increased sensitivity to GnRH-I and a ~ 20-fold increased sensitivity to GnRH-II (Fig. 5A-B).

As the PPS/SEP motif in EL3 and Leu<sup>7.43</sup>, Ala<sup>7.46</sup>, and Pro<sup>7.47</sup> residues in TMH7 affect ligand selectivity, we examined a combinatory effect of these two motifs. The gmGnRHR-2 with SEP/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutation revealed a slight decrease in sensitivity for both GnRH-I and GnRH-II (Fig. 5C-D). This mutant also showed a decrease in E<sub>max</sub> values for both GnRHs compared with the L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant, which may be due to low receptor expression (Table 2). Interestingly, additional mutations rEL3S/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C (Fig. 7B), SEP/I<sup>7.37</sup>F/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C (Fig. 5C-D) or SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C (Fig. 5C-D) increased sensitivity to GnRH-I and decreased sensitivity to GnRH-II (Table 2).

**Ligand Binding Affinities.** Ligand affinities of wild type, rEL3L, and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutants were determined using a competition binding assay. For the binding assay, HeLa cells were used as they, when transfected with the receptors, have a much higher binding capacity than CV-1 cells. It should be noted that HeLa cells, in the c-fos-luc assay system, give the EC<sub>50</sub> values similar to those of CV-1 cells when we applied the same receptor and ligand (data not shown). HeLa cells, however, have a high basal c-fos-luc activity, therefore their fold-increases are usually much lower than those in CV-1 cells (Oh et al., 2003). <sup>125</sup>I-GnRH-II (250,000 cpm) was applied to HeLa cells expressing wild type and mutant receptors in the presence of graded concentrations of cold GnRH-I or GnRH-II. Log IC<sub>50</sub> for GnRH-I in cells

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expressing rEL3L ( $-7.87 \pm 0.22$ ) and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C ( $-8.30 \pm 0.11$ ) were significantly lowered compared to that in cells expressing the wild type receptor ( $-6.69 \pm 0.21$ ) (Fig. 6A), indicating an increased affinity for GnRH-I in the mutant receptors. Log IC<sub>50</sub> for GnRH-II in cells expressing the wild type, rEL3L, and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutants were  $-9.58 \pm 0.27$ ,  $-10.19 \pm 0.23$ , and  $-9.96 \pm 0.27$ , respectively (Fig. 6B), showing that ligand affinities for GnRH-II in mutant receptors do not change as drastically as those for GnRH-I.

Relative ligand binding of mutant constructs was determined using <sup>125</sup>I-GnRH-II in the absence or presence of cold GnRH-II (10 μM). For wild type gmGnRHR-2, total and nonspecific binding were  $2.3 \pm 0.2\%$  ( $5826 \pm 50$  cpm) and  $0.60 \pm 0.2\%$  ( $1567 \pm 30$  cpm), respectively. Total binding for other mutant receptors ranged from 1.16 to 6.83%, while nonspecific bindings for other receptors were the same as that for the wild type receptor. Mutants that did not respond to GnRHs (rEL3L/P<sup>313</sup>A, rEL3L/C<sup>314</sup>P, L<sup>7.43</sup>F, and A<sup>7.46</sup>P) were unable to bind radioiodinated GnRH-II. rEL3S, L<sup>7.43</sup>F/P<sup>7.47</sup>C, and SEP/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C showed relatively low binding; rEL3L/PPS and A<sup>7.46</sup>P/P<sup>7.47</sup>C had higher binding than gmGnRHR-2. Other mutants exhibited 49.7% to 178.6% binding compared with wild type gmGnRHR-2 (Table 2).

**Ligand Sensitivity for Chimeric GnRHs.** Natural and chimeric GnRHs, in which amino acids at positions 5, 7, and 8 were substituted, were used to examine ligand sensitivity of gmGnRHR-2, rEL3L, and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutants. For wild type gmGnRHR-2, all chimeric GnRHs ([His<sup>5</sup>]GnRH-I, [Trp<sup>7</sup>]GnRH-I, [Trp<sup>8</sup>]GnRH-I, and [Trp<sup>7</sup>, Leu<sup>8</sup>]GnRH-I) exhibited a higher potency than GnRH-I (Table 3). In particular, substitution of the amino acid residues at positions 7 and 8 of

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GnRH-I greatly increased potency to activate gmGnRHR-2. It should be noted that chimeric ligands [His<sup>5</sup>]GnRH-I and [Trp<sup>7</sup>]GnRH-I which, like GnRH-I, retain Arg<sup>8</sup> showed a 100 - 200-fold increased potency for either rEL3L or SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C that have enhanced sensitivity to GnRH-I (Fig. 7 and Table 3). [Trp<sup>8</sup>]GnRH-I revealed a 20 - 50-fold increased sensitivity for rEL3L and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C compared to that for wild type gmGnRHR-2. Interestingly, rEL3L and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C had similar sensitivity to chimeric GnRHs. (Fig. 7 and Table 3). This result supports the idea that positions 7 and 8 in GnRH are important for conferring its specificity.

**Molecular Modeling.** To support our biochemical data we constructed models to simulate the interaction of GnRHs with wild-type gmGnRHR-2 and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant (Fig. 8). Overall, the models agree well with previous reports (Hövelmann et al., 2002; Wang et al., 2004): two cystein residues (Cys<sup>113</sup> and Cys<sup>188</sup>) of the receptors are close as they are involved in a disulfide bond; pGlu<sup>1</sup> of the ligands formed hydrogen bonds with Asn<sup>5.39</sup> of the receptors; Trp<sup>3</sup> of the ligands was located in the aromatic cage formed by Trp<sup>6.48</sup>, Phe<sup>5.43</sup>, and Tyr<sup>6.52</sup> of the receptors; Arg<sup>8</sup> of GnRH-I formed an ionic interaction with Glu<sup>7.32</sup> of the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant.

GnRH-II and GnRH-I differ by three residues: His/Tyr<sup>5</sup>, Trp/Leu<sup>7</sup>, and Tyr/Arg<sup>8</sup>. Trp<sup>7</sup> of GnRH-II made a hydrophobic contact with Pro<sup>7.32</sup> of gmGnRHR-2. Tyr<sup>8</sup> had an interaction with Pro<sup>7.32</sup> and His<sup>7.36</sup> (Fig. 8A). In addition, Trp<sup>7</sup> formed a hydrogen bond with the carbonyl oxygen of Val<sup>7.30</sup>, and Tyr<sup>8</sup> with the backbone of Ser<sup>7.33</sup>. However, in the GnRH-I/gmGnRHR-2 complex, Leu<sup>7</sup> formed a hydrophobic contact with Tyr<sup>6.58</sup>. Arg<sup>8</sup> moved to EL2 and interacted with Val<sup>4.67</sup> via a hydrophobic interaction (Fig. 8B).



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On the other hand, Trp<sup>7</sup> of GnRH-II formed hydrophobic contacts with Glu<sup>7.32</sup> and His<sup>7.36</sup> of the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant, and Tyr<sup>8</sup> made a hydrogen bond with the backbonal nitrogen of Glu<sup>7.32</sup> (Fig. 8C). In the complex of GnRH-I with the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant, Leu<sup>7</sup> formed hydrophobic contact with Tyr<sup>6.58</sup>, and Arg<sup>8</sup> had an ionic interaction with Glu<sup>7.32</sup> (Fig. 8D). Mutation of the four amino acids at TMH7 altered intramolecular interactions. For instance, the hydrogen bond between Leu<sup>7.43</sup> and Ala<sup>7.46</sup> of gmGnRHR-2 was suppressed in the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant. In addition, novel hydrophobic contacts of Phe<sup>7.38</sup> with Leu<sup>6.54</sup>, Phe<sup>7.43</sup> with Asp<sup>2.61</sup>, and Pro<sup>7.46</sup> with Leu<sup>7.44</sup> were formed in the mutant receptor. It is also notable that the various intramolecular interactions were highly dependent upon ligand type. Hydrophobic contacts of Phe<sup>7.43</sup> with Leu<sup>1.42</sup> and Val<sup>2.57</sup>, Cys<sup>7.47</sup> with Gly<sup>1.49</sup>, and hydrogen bonds between Cys<sup>7.47</sup> and Ser<sup>1.45</sup> and Asn<sup>1.50</sup> were present in the mutant receptor/GnRH-I complex but these interactions were absent in the mutant receptor/GnRH-II complex. In contrast, the hydrophobic contacts of Phe<sup>7.38</sup> with Leu<sup>6.53</sup> and Pro<sup>7.46</sup> with Val<sup>2.53</sup> were present in the mutant receptor/GnRH-II complex.

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### Discussion

The present study demonstrates that replacement of EL3 and EL3-proximal TMH7 of gmGnRHR-2 with those of rat GnRHR greatly improves ligand sensitivity to GnRH-I but not to GnRH-II. Site-directed mutations on gmGnRHR-2 and back mutations on the domain-swapped receptor show that the PPS motif in EL3 and Leu<sup>7.38</sup>, Leu<sup>7.43</sup>, Ala<sup>7.46</sup>, and Pro<sup>7.47</sup> in TMH7 of gmGnRHR-2, and the corresponding residues of rat GnRHR are responsible for differential ligand sensitivity to GnRH-I and GnRH-II.

It was suggested that not only Glu/Asp<sup>7.32</sup> but also the positions of Ser and Pro flanking Glu/Asp<sup>7.32</sup> in EL3 of mammalian type I GnRHR determine high selectivity for GnRH-I (Wang et al., 2004; Fromme et al., 2004). These findings indicate that a local conformation of EL3 is critical for differential ligand selectivity among nonmammalian, mammalian type I, and type II GnRHRs. However, replacement of EL3 from gmGnRHR-2 alone does not affect ligand selectivity to GnRH-I and GnRH-II. Similarly, substitution of SEP for the PEY motif of bfGnRHR-2 does not alter ligand sensitivity to GnRH-I and GnRH-II (Wang et al., 2004). These observations suggest that other amino acid residues/motifs are involved in the selectivity of GnRH. Our study strongly suggests that in mammalian type II GnRHR, EL3-proximal TMH7 in addition to EL3 participates in differential ligand selectivity. The importance of EL3-proximal TMH7 in ligand sensitivity is supported by the observation that rat GnRHR with the entire TMH7 of gmGnRHR-2 (r6TM) exhibits a significant decrease in sensitivity for GnRH-I. Mutations of Pro<sup>7.47</sup> to Cys combined with the mutation of Leu<sup>7.43</sup> to Phe and/or Ala<sup>7.46</sup> to Pro significantly increases sensitivity for GnRH-I, while mutations of a single amino acid residue at these positions does not affect ligand selectivity to GnRH-I, suggesting

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that the combination of each amino acid in TMH7 is critical for differential ligand selectivity.

Based on Millar's classification (Millar et al., 2004) we aligned the sequences of EL3 and proximal TMH7 of various GnRHR subtypes: human-1 and rat-1 for mammalian type I receptors, green monkey-2 and marmoset-2 for type II mammalian receptors, bullfrog-3 and *Xenopus*-2 for nonmammalian type II receptors, Japanese medaka-1 and bullfrog-2 for type III receptors, and finally Japanese medaka-2, bullfrog-2, and catfish-1 for nonmammalian type I receptors. Leu<sup>7.40</sup>, Leu<sup>7.44</sup>, and Asn<sup>7.45</sup> residues are conserved between gmGnRHR-2 and the mammalian type I receptors, but these residues are also largely conserved in many other nonmammalian GnRHRs. Thus, these amino acid residues are not specific to mammalian GnRHR subtypes. After excluding the amino acids, which are conserved throughout the GnRHR subtypes, we found that at least 4 residues, Iso<sup>7.37</sup>, Leu<sup>7.38</sup>, Gly<sup>7.42</sup>, and Leu<sup>7.43</sup> in gmGnRHR-2 are different from those in mammalian type I receptors but are highly conserved in nonmammalian GnRHRs (Fig. 9). Among these, Iso<sup>7.37</sup> and Gly<sup>7.42</sup> are not likely to contribute to GnRH-I selectivity, as back mutations of these residues in the rEL3L receptor did not significantly affect sensitivity to GnRH-I. Two amino acids, Ala<sup>7.47</sup> and Pro<sup>7.48</sup> in gmGnRHR-2 are different from either those in type I mammalian GnRHR or in nonmammalian GnRHRs. Thus, they are unique to the type II mammalian receptors.

Double or triple mutations of Leu<sup>7.43</sup>, Ala<sup>7.46</sup>, and Pro<sup>7.47</sup> in TMH7 significantly increases ligand sensitivity to GnRH-I. Currently, we cannot clearly explain how the combined mutation L<sup>7.38</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C increases GnRH-I sensitivity. It is unlikely that these residues have direct interactions with GnRH-I as they are deeply buried in the three-dimensional structure. Rather, the mutation on these residues may play a role in

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modulating conformation of the binding pocket in EL3. Consistently, our molecular modeling data show no direct interactions of these residues with the ligand. It is of interest to note that inter- and intra-molecular interactions of the mutant receptor could be modified by the ligand type applied, indicating that conformational changes in these residues may be closely related to the alteration in the ligand binding pocket of EL3.

Amino acids at positions 7.46 and 7.47 in TMH7 appear critical for receptor conformation and stability. The mutation of Ala<sup>7.46</sup> to Pro in wild type gmGnRHR-2 and the mutation of Pro<sup>7.46</sup> to Ala or Cys<sup>7.47</sup> to Pro in rEL3L impair receptor responsiveness. Extremely low binding of these mutant receptors to GnRH suggests that this impairment can be ascribed to the loss of binding activity or receptor stability. It is known that a Pro residue leads to a local constraint on the polypeptide chain conformation due to its pyrrolidine ring structure. Thus, Pro at a proper position in TMH7 appears to be important for receptor conformation/stability in wild type and mutant gmGnRHR-2. The occurrence of two successive Pro residues found in the rEL3L/C<sup>7.47</sup>P, and A<sup>7.46</sup>P mutants might disrupt receptor conformation/stability as the loss of responsiveness of A<sup>7.46</sup>P mutant can be rescued by a double mutation (A<sup>7.46</sup>P/P<sup>7.47</sup>C). Further, Pro<sup>7.46</sup> in the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant has an intramolecular contact with Leu<sup>7.44</sup>, which is different from that of the wild type receptor in which Ala<sup>7.46</sup> at the same position has a hydrophobic interaction with Leu<sup>7.43</sup>. Pro<sup>7.47</sup> alone in gmGnRHR-2 could not critically alter receptor conformation/stability as the mutation of Pro<sup>7.47</sup> to Cys did not affect ligand potency. The Cys residue at position 7.47 is highly conserved in nonmammalian and mammalian type I GnRHRs as well as in many other GPCRs, except for gmGnRHR-2 (Fig. 9). Thus, it may be possible that mutation of Pro<sup>7.47</sup> to Cys helps to form a more stable conformation. Mutation of Leu<sup>7.43</sup> to Phe in gmGnRHR-2 abolishes

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the receptor responsiveness to ligand, which can be rescued by a double mutation (L<sup>7.43</sup>F/P<sup>7.47</sup>C). In a three-dimensional structure, Leu<sup>7.43</sup> and Pro<sup>7.47</sup> are spatially very close. Therefore, it is postulated that a mutation of Leu<sup>7.43</sup> to Phe might cause a steric hindrance, which can be reversed by a further replacement of Pro<sup>7.47</sup> to Cys. Moreover, double mutations (L<sup>7.43</sup>F/P<sup>7.47</sup>C or A<sup>7.46</sup>P/P<sup>7.47</sup>C) not only rescue the activity of the L<sup>7.43</sup>F or A<sup>7.46</sup>P mutant but also significantly increase the ligand sensitivity to both GnRH-I and GnRH-II. Further, an approximately 100-fold increase in sensitivity toward GnRH-I was observed in a triple mutant L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C compared to that of wild type gmGnRHR-2. Such an increase in sensitivity to ligands suggests that this motif is crucially involved in receptor activation. Thus, it is likely that Pro<sup>7.47</sup> in the wild type gmGnRHR-2 and Pro<sup>7.46</sup> in mutant receptors are involved in TMH movements, contributing to GPCR activation/inactivation by forming molecular hinges or swivels (Sansom and Weinstein, 2000; Stitham et al., 2002).

It is noteworthy that replacement of EL3 and EL3-proximal TMH7 or mutations of amino acids in these regions did not decrease sensitivity to GnRH-II, instead there was a slightly increased sensitivity to GnRH-II. It is well known that an acidic amino acid at position 7.32 in EL3 is required for high affinity binding with Arg<sup>8</sup> of GnRH-I. It appears that such an acidic residue also plays a certain role in interaction with Tyr<sup>8</sup> of GnRH-II. Using a molecular model, Blumenröhr et al. (2002) suggested that Tyr<sup>8</sup> of GnRH-II interacts with Glu<sup>7.32</sup> in EL3 of the catfish GnRHR. Consistently, our molecular model also showed that Tyr<sup>8</sup> of GnRH-II has contact with Glu<sup>7.32</sup> of the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant. It should be noted that Arg<sup>8</sup> of GnRH-I did not interact with the PPS motif of wild type gmGnRHR-2 while Tyr<sup>8</sup> of GnRH-II had contact with Pro<sup>7.32</sup> of wild type gmGnRHR-2. Thus, at least in the gmGnRHR-2

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structure, GnRH-II may not discriminate the receptor with the PPS motif from the receptor with the SEP motif.

Substitution of His for Tyr<sup>5</sup> of GnRH-I did not alter its potency to activate wild type gmGnRHR-2, rEL3L, or SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C, suggesting that position 5 of GnRH does not largely contribute to receptor-ligand interaction. This result is consistent with previous reports (Blomenröhr et al., 2002; Wang et al., 2003). Substitution of Trp for Leu<sup>7</sup> or Arg<sup>8</sup> in GnRH-I significantly increased the ability to activate gmGnRHR-2, indicating the importance of positions 7 and 8 in recognition of mammalian type II GnRHR. The SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant showed similar ligand sensitivity with that of rEL3L, implying that multiple residues are required for distinguishing GnRH-II from GnRH-I. It is worthy of note that gmGnRHR-2 shows a high sequence identity in EL3-proximal TMH7 with nonmammalian GnRHRs but a relatively low sequence identity with that of mammalian type I GnRHR (Fig. 9). The evolutionary divergence of EL3 and TMH7 between mammalian type I and type II GnRHR, therefore, may confer the differential selectivity toward GnRH-I and GnRH-II.

In summary, our studies demonstrate that EL3 and EL3-proximal TMH7 are responsible for differential ligand selectivity between mammalian type I and type II GnRHRs. The elucidation of specific domains responsible for ligand selectivity may facilitate the understanding of ligand and receptor molecular coevolution, the mechanism of ligand-mediated GnRHR activation, and the development of novel drugs.

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### LEGENDS for FIGURES

**Fig. 1.** Ligand selectivity of rat GnRHR or gmGnRHR-2. A, IP assays were performed using GH<sub>3</sub> cells which stably express rat GnRHR (rat) or gmGnRHR-2 (gm2). Cells were treated with increasing concentrations of GnRH-I or GnRH-II for 30 min. B, c-fos promoter-driven luciferase (c-fos-luc) activity was examined in CV-1 cells that were cotransfected with 200 ng c-fos-luc reporter vector plus rat GnRHR or gmGnRHR-2. Twenty-four hours after transfection, cells were serum-starved for 18 h then treated for 6 h with GnRH-I or GnRH-II. Cell lysates were used for luciferase assays. Dashed lines are dose-responses of rat (○) and gm2 (●) to GnRH-I and solid lines are dose-responses of rat (□) and gm2 (■) to GnRH-II.

**Fig. 2.** Ligand selectivity of chimeric GnRHRs. A-D, Chimeric receptors rEL3S (A), rEL3L (B), r6TM (C), r6.5TM (D) were transfected into CV-1 cells and c-fos-luc activity was measured. Dashed lines are dose-response c-fos-luc activity of wild type gmGnRHR-2 (gm2) or rat GnRHR (rat) and solid lines are that of chimeric receptors. Thin gray lines in the receptor diagram represent the portion of gmGnRHR-2, while thick blue lines are the region from rat GnRHR. E, The amino acid sequence alignment of EL3 and TMH7 among wild type and chimeric receptors. The amino acid numbers are seen beside the sequences. Insertions in rEL3S and rEL3L are underlined. Amino acids that are further characterized in ensuing experiments are shown in bold and indicated by arrows with the position numbers.

**Fig. 3.** Point mutation of the rEL3L mutant. CV-1 cells were transfected with 200 ng c-

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fos-luc reporter plasmid plus rEL3L or rEL3L with point mutations. A-B, c-fos-luc activity in cells expressing rEL3L/F<sup>7.37</sup>I, rEL3L/F<sup>7.38</sup>L, rEL3L/A<sup>7.42</sup>G, rEL3L/F<sup>7.43</sup>L in response to GnRH-I (A) and GnRH-II (B). C-D, c-fos-luc activity in cells expressing rEL3L/P<sup>7.46</sup>A, rEL3L/C<sup>7.47</sup>P, rEL3L/E<sup>7.32</sup>Q, and rEL3L/PPS in response to GnRH-I (C) and GnRH-II (D). Dashed and solid lines are data obtained from rEL3L and the mutant receptors, respectively.

**Fig. 4.** Point mutations of wild type gmGnRHR-2. CV-1 cells were transiently transfected with gmGnRHR-2 (gm2) and point-mutated receptors L<sup>7.43</sup>F, A<sup>7.46</sup>P, and P<sup>7.47</sup>C. Cells were treated with different concentrations of GnRH-I (A) or GnRH-II (B) for 6 h.

**Fig. 5.** Multiple mutations in EL3 and TMH7 of gmGnRHR-2. CV-1 cells were transfected with wild type gmGnRHR-2 or each mutant. A-B, Cells expressing A<sup>7.46</sup>P/P<sup>7.47</sup>C, L<sup>7.43</sup>F/P<sup>7.47</sup>C, L<sup>7.43</sup>F/A<sup>7.46</sup>P, and L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C were treated with different concentrations of GnRH-I (A) or GnRH-II (B) for 6 h. C-D, Cells expressing combinatory mutants SEP/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C, rEL3S/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C, SEP/I<sup>7.37</sup>F/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C were treated with GnRH-I (C) or GnRH-II (D). Dashed lines are data from wild type receptor (gm2) and solid lines are from mutant receptors, respectively.

**Fig. 6.** Competition binding assays for the wild type and mutant receptors. <sup>125</sup>I-GnRH-II was applied to HeLa cells expressing the wild type (gm2), rEL3L, and SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant receptors in the presence of graded

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concentrations of cold GnRH-I (A) or GnRH-II (B).

**Fig. 7.** Ligand selectivity of the chimeric receptors to various chimeric GnRHs. CV-1 cells were transfected with wild type (gm2) (A), rEL3L (B), or SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C (C). Cells were treated with different concentrations of wild type and chimeric GnRHs for 6 h.

**Fig. 8.** Molecular models for the interaction of GnRH-II with wild-type gmGnRHR-2 (A), GnRH-I with wild-type gmGnRHR-2 (B), GnRH-II with the SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant (C), and GnRH-I with SEP/L<sup>7.38</sup>F/L<sup>7.43</sup>F/A<sup>7.46</sup>P/P<sup>7.47</sup>C mutant (D). TMHs 1 to 6 of GnRHRs were drawn as cylinders in gray. TMH 7 and ligands were drawn as tubes in yellow and green, respectively. The following residues were drawn as a ball and stick-model with carbon atoms in cyan, oxygen atoms in red, nitrogen atoms in blue, and sulfur atoms in yellow. Pro/Ser<sup>7.31</sup>, Pro/Glu<sup>7.32</sup>, Ser/Pro<sup>7.33</sup>, Leu/Phe<sup>7.38</sup>, Ile/Phe<sup>7.43</sup>, Ala/Pro<sup>7.46</sup>, Pro/Cys<sup>7.47</sup>, Cys<sup>3.25</sup> and Cys<sup>4.78</sup> of receptors, and His/Tyr<sup>5</sup>, Trp/Leu<sup>7</sup>, and Tyr/Arg<sup>8</sup> of ligands are identified with numbers.

**Fig. 9.** Amino acid sequence alignment of EL3 and TMH7 in GnRHRs and other GPCRs. The amino acids in EL3 and TMH7 responsible for ligand selectivity are shown in bold and indicated by arrows. The amino acids that are highly conserved among mammalian type II and nonmammalian GnRHRs, but differ from the mammalian type I receptor are shaded. Note that the Cys residue at position 7.48 is highly conserved in other GPCRs. Sequences used are human-1 GnRHR (Accession No. NM\_000406), rat-1



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GnRHR (NM\_031038), green (G) monkey-2 GnRHR (AF353988), marmoset-1 GnRHR (AF368286), Xenopus-2 GnRHR (AF257320), bullfrog-1-3 GnRHR (AF144063, AF153913, and AF144062), Japanese (J) medaka-1-2 GnRHR (AB057677 and AB057676), catfish-1 GnRHR (X97497), human angiotensin II 1 receptor (AF245699), human bradykinin B2 receptor (NM\_000623), human neuropeptide Y1 receptor (NM\_000909), human opioid mu1 receptor (NM\_001008505), human somatostatin-1 receptor (NM\_001049), and human oxytocin receptor (AY389507).

TABLE 1.

TABLE 1. The differential ligand selectivity between rat GnRHR (rat) and gmGnRHR-2 (gm2)

	IP (EC <sub>50</sub> , Log M)		c-fos-luc (EC <sub>50</sub> , Log M)	
	GnRH-I	GnRH-II	GnRH-I	GnRH-II
<b>gm2</b>	-6.45 ± 0.11	-8.76 ± 0.14 (204)	-5.83 ± 0.10	-8.21 ± 0.09 (239)
<b>rat</b>	-8.55 ± 0.14	-7.68 ± 0.15 (7.4)	-8.31 ± 0.06	-7.54 ± 0.07 (5.9)

Data represent the mean ± SEM from three independent experiments.  
Numbers between parentheses represent fold difference in sensitivity  
between GnRH-I and GnRH-II

TABLE 2.

TABLE 2. Relative binding,  $E_{\max}$  (fold-induction) and  $EC_{50}$  (log M) of various GnRH receptors

Receptors	Relative binding (% gm2)	GnRH-I		GnRH-II	
		$E_{\max}$	$EC_{50}$	$E_{\max}$	$EC_{50}$
gm2	100 ± 1.2	9.33 ± 0.35	-6.00 ± 0.10	13.82 ± 0.35	-8.54 ± 0.13
rat	98.8 ± 1.8	12.49 ± 0.26	-8.31 ± 0.06*	8.78 ± 0.29	-7.54 ± 0.07*
r6TM	89.5 ± 1.1	9.59 ± 0.33	-7.16 ± 0.12*	5.45 ± 0.20	-7.95 ± 0.12
r6.5TM	168.1 ± 4.5	17.14 ± 0.38	-9.18 ± 0.08*	17.53 ± 0.35	-8.48 ± 0.07
rEL3S	21.8 ± 0.1	5.79 ± 0.18	-6.64 ± 0.10	7.72 ± 0.21	-8.21 ± 0.09
rEL3L	110.1 ± 5.4	6.24 ± 0.14	-8.38 ± 0.09*	6.37 ± 0.08	-8.76 ± 0.05
rEL3L/F7.37I	110.3 ± 9.4	6.83 ± 0.31	-8.25 ± 0.16*	10.77 ± 0.53	-8.24 ± 0.18
rEL3L/F7.38L	138.7 ± 1.1	7.63 ± 0.35	-7.98 ± 0.18*	10.56 ± 0.29	-8.34 ± 0.10
rEL3L/A7.42G	49.7 ± 3.4	6.31 ± 0.22	-8.25 ± 0.09*	5.43 ± 0.12	-8.79 ± 0.10
rEL3L/F7.43L	52.7 ± 0.1	3.89 ± 0.27	-7.81 ± 0.20*	4.61 ± 0.19	-8.63 ± 0.18
rEL3L/P7.46A	3.8 ± 0.5	N.d			
rEL3L/C7.47P	1.5 ± 1.0	N.d			
rEL3L/E7.32Q	70.7 ± 0.2	3.53 ± 0.14	-7.28 ± 0.15*	4.61 ± 0.14	-8.47 ± 0.14
rEL3L/PPS	332.2 ± 7.3	12.50 ± 0.27	-7.61 ± 0.07*	16.78 ± 0.35	-9.36 ± 0.09*
L7.43F	1.7 ± 1.2	N.d			
A7.46P	3.8 ± 0.9	N.d			
P7.47C	72.1 ± 2.1	6.46 ± 0.26	-5.93 ± 0.08	12.29 ± 0.57	-8.29 ± 0.17
L7.43F/A7.46P	3.2 ± 2.8	N.d			
L7.43F/P7.47C	16.8 ± 1.2	3.57 ± 0.07	-7.03 ± 0.07*	4.94 ± 0.15	-8.79 ± 0.17
A7.46P/P7.47C	303.3 ± 1.4	10.53 ± 0.61	-7.17 ± 0.17*	11.00 ± 0.29	-9.65 ± 0.12*
L7.43F/A7.46P/P7.47C	178.6 ± 8.6	5.93 ± 0.20	-8.21 ± 0.13*	7.06 ± 0.20	-9.82 ± 0.13*
SEP/L7.43F/A7.46P/P7.47C	11.1 ± 0.5	2.92 ± 0.08	-7.87 ± 0.12*	2.53 ± 0.10	-9.69 ± 0.26*
rEL3S/L7.43F/A7.46P/P7.47C	54.6 ± 2.1	4.68 ± 0.13	-8.76 ± 0.15*	6.14 ± 0.11	-9.59 ± 0.10*
SEP/I7.37F/L7.38F/ L7.43F/A7.46P/P7.47C	51.2 ± 1.3	3.99 ± 0.14	-8.59 ± 0.17*	4.76 ± 0.17	-9.32 ± 0.17*
SEP/L7.38F/L7.43F/A7.46P/P7.47C	62.1 ± 1.7	5.27 ± 0.13	-8.79 ± 0.11*	5.79 ± 0.14	-9.43 ± 0.11*

Values represent the mean ± SEM of three independent experiments performed in duplicate. Binding was expressed as percentage of specific binding of gmGnRHR-2 (gm2). N.d., not determined. \*,  $p < 0.05$  vs. wild type gm2

TABLE 3.

TABLE 3.  $E_{\max}$  (fold-induction) and  $EC_{50}$  (log M) for various chimeric GnRHs

Chimeric GnRH	gm2		rEL3L		SEP/L <sup>7.38</sup> F/L <sup>7.43</sup> F/A <sup>7.46</sup> P/P <sup>7.47</sup> C	
	$E_{\max}$	$EC_{50}$	$E_{\max}$	$EC_{50}$	$E_{\max}$	$EC_{50}$
GnRH-I	9.33 ± 0.35	-6.00 ± 0.10	6.24 ± 0.14	-8.38 ± 0.09	5.27 ± 0.13	-8.79 ± 0.11
[His <sup>5</sup> ]GnRH-I	8.66 ± 1.11	-6.38 ± 0.34	6.82 ± 0.39	-8.51 ± 0.22	4.78 ± 0.21	-8.53 ± 0.18
[Trp <sup>7</sup> ]GnRH-I	10.73 ± 0.05	-7.15 ± 0.05*	3.38 ± 0.14	-9.42 ± 0.26*	3.00 ± 0.19	-8.90 ± 0.34
[Trp <sup>8</sup> ]GnRH-I	9.48 ± 0.11	-6.74 ± 0.03*	4.80 ± 0.17	-8.16 ± 0.15	3.42 ± 0.20	-8.30 ± 0.24
[Trp <sup>7</sup> , Leu <sup>8</sup> ]GnRH-I	12.11 ± 0.42	-7.47 ± 0.11*	5.68 ± 0.38	-8.43 ± 0.27	5.94 ± 0.43	-7.59 ± 0.23*

Data represent the mean ± SEM of three independent experiments performed in duplicate.

\*,  $p < 0.05$  vs. GnRH-I

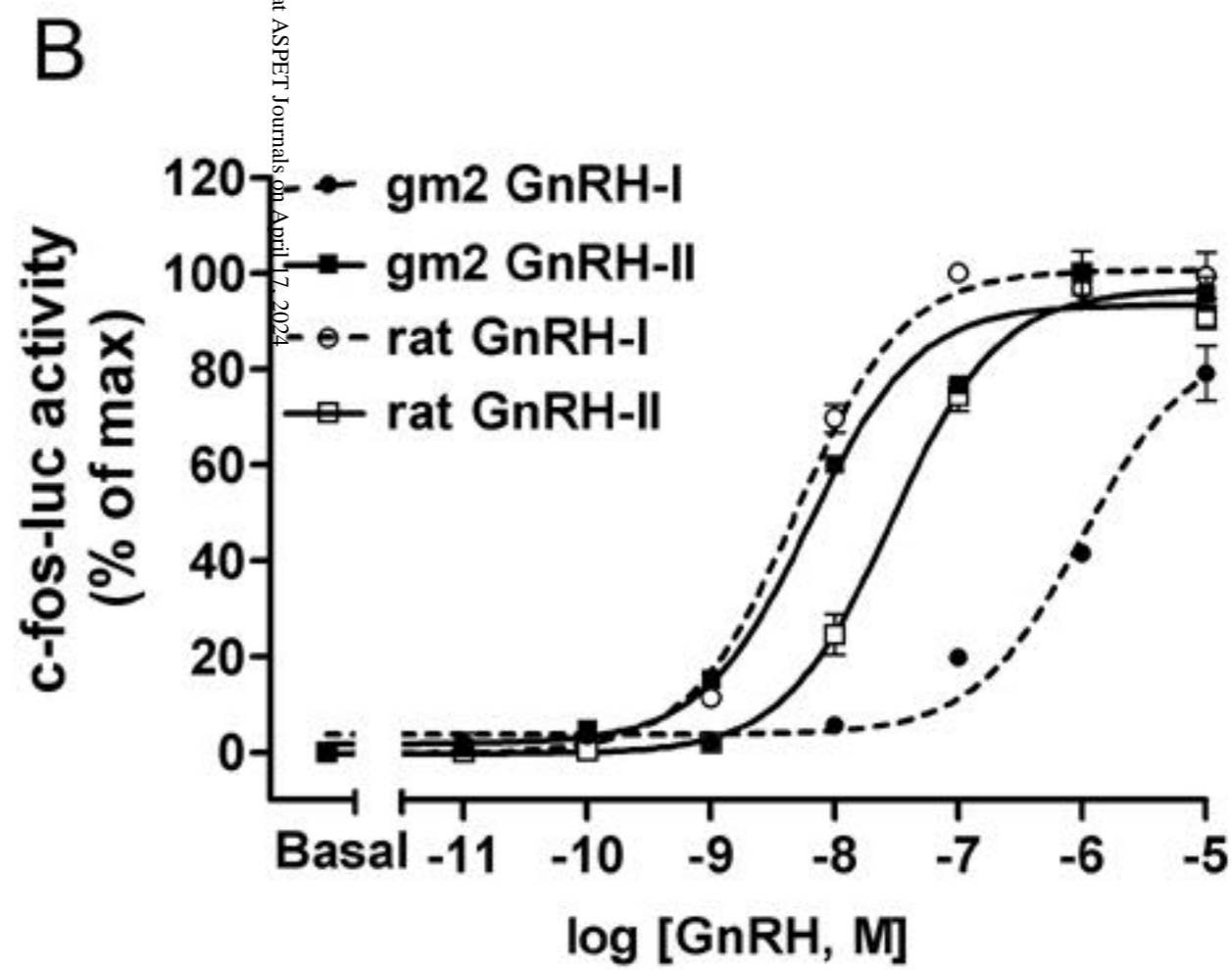
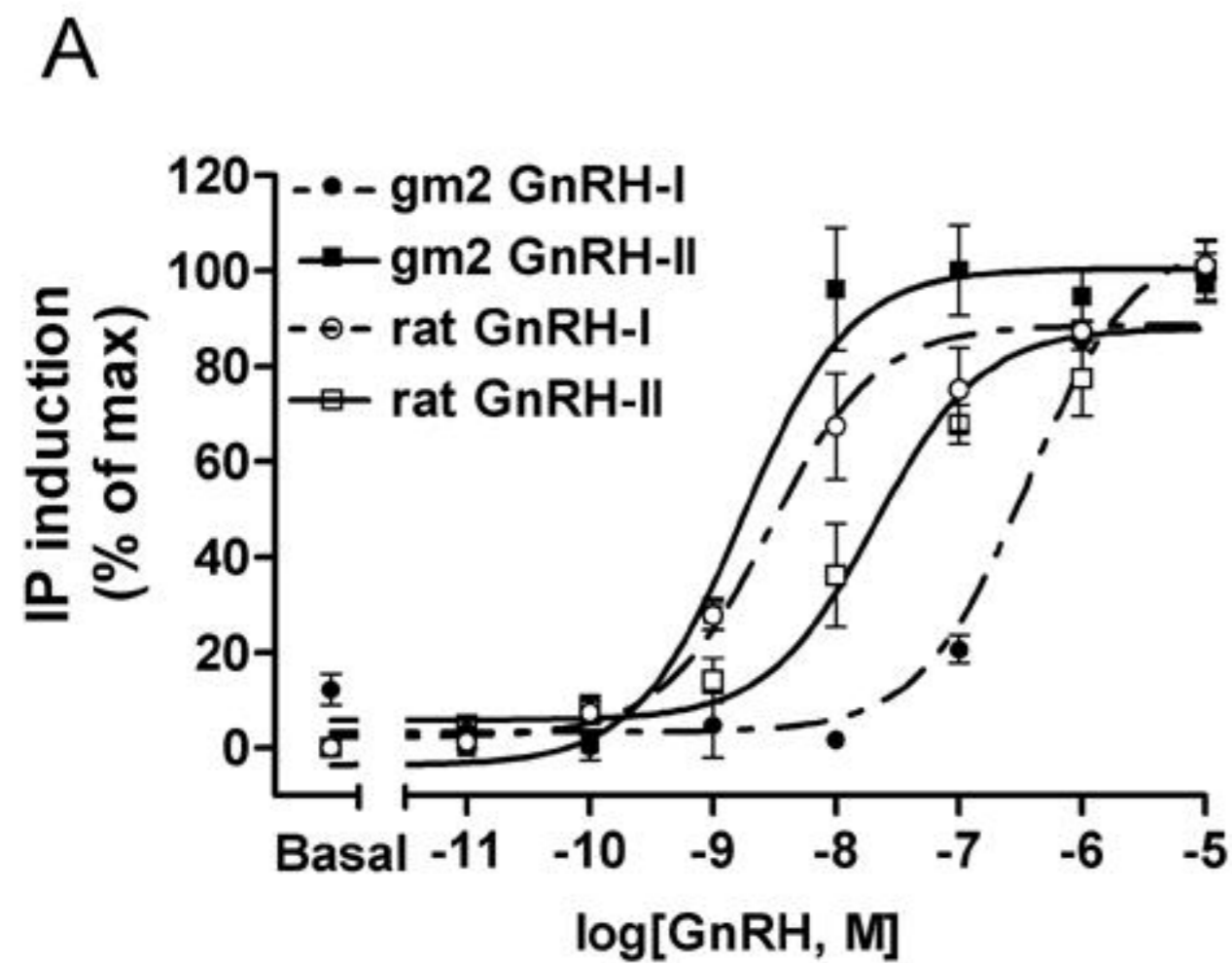
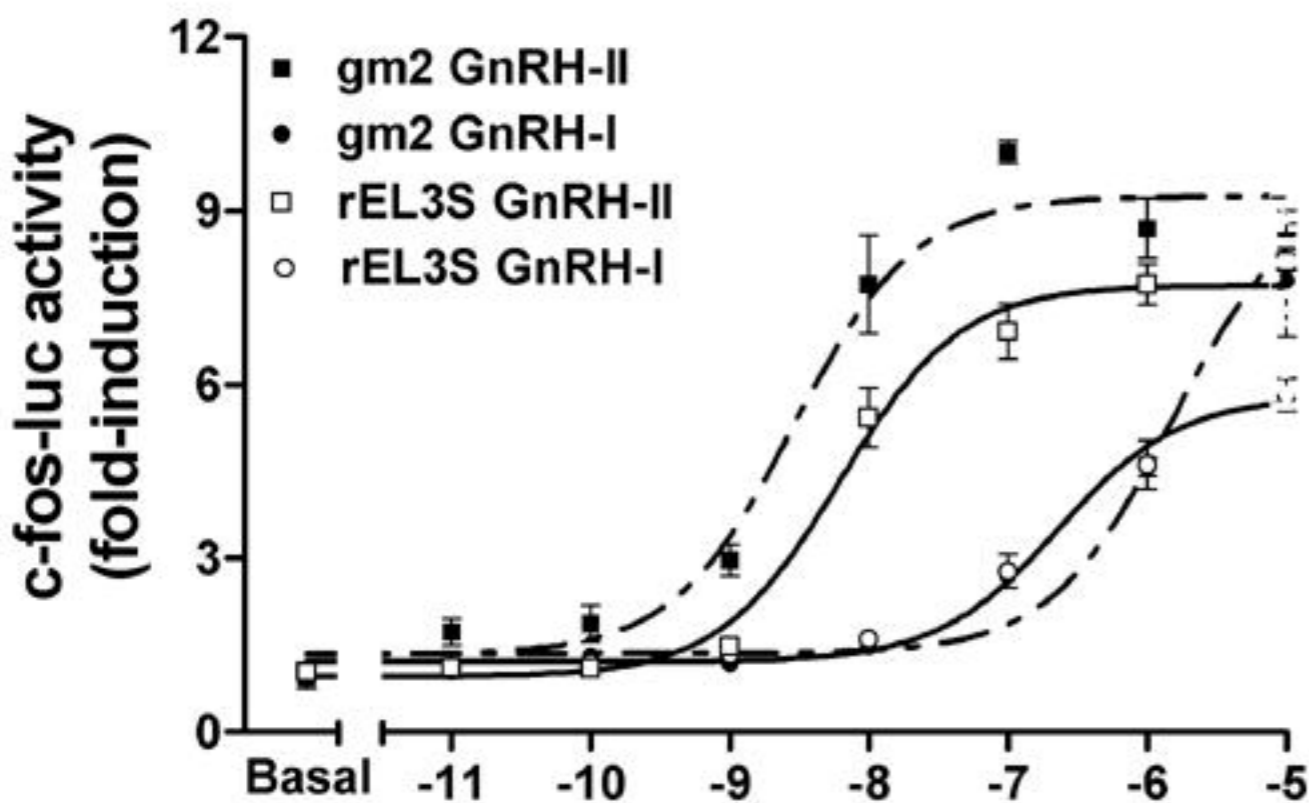
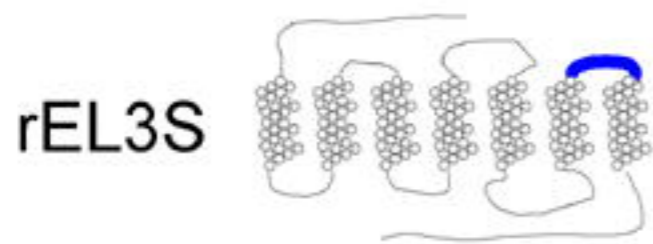
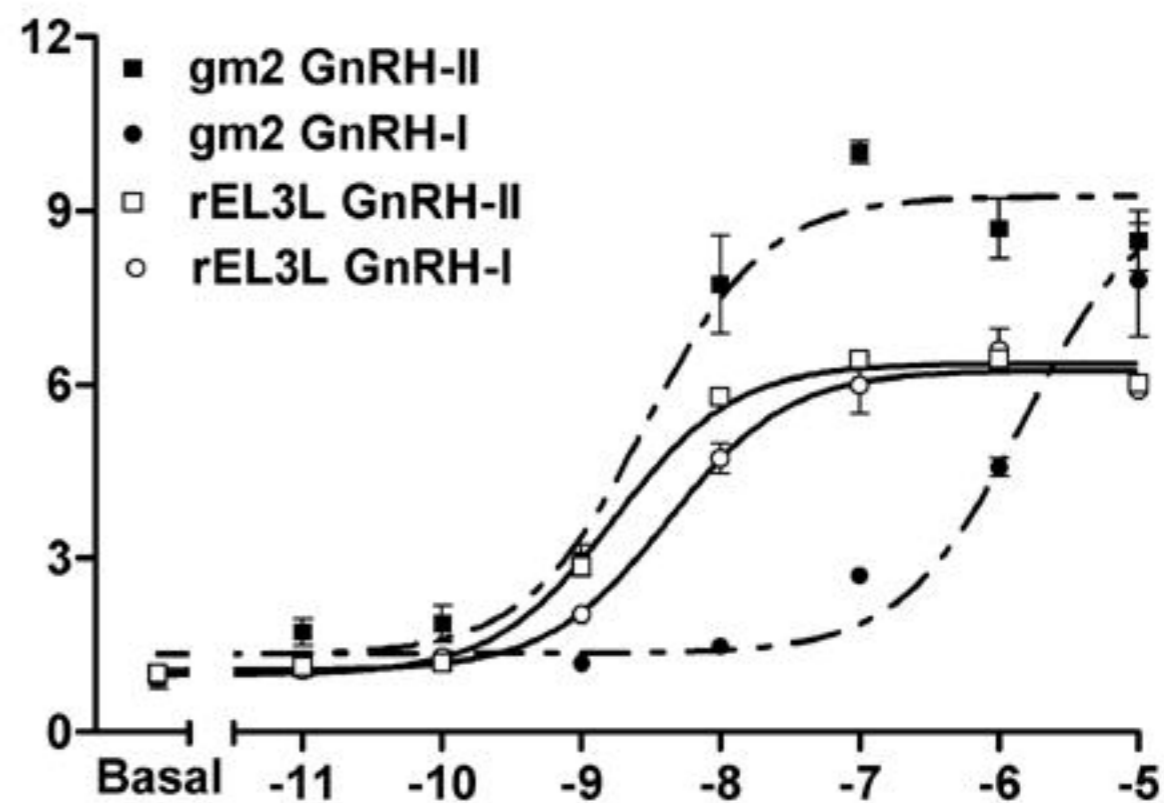
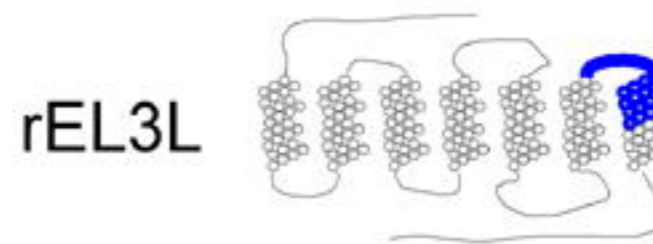


Fig. 2.

A

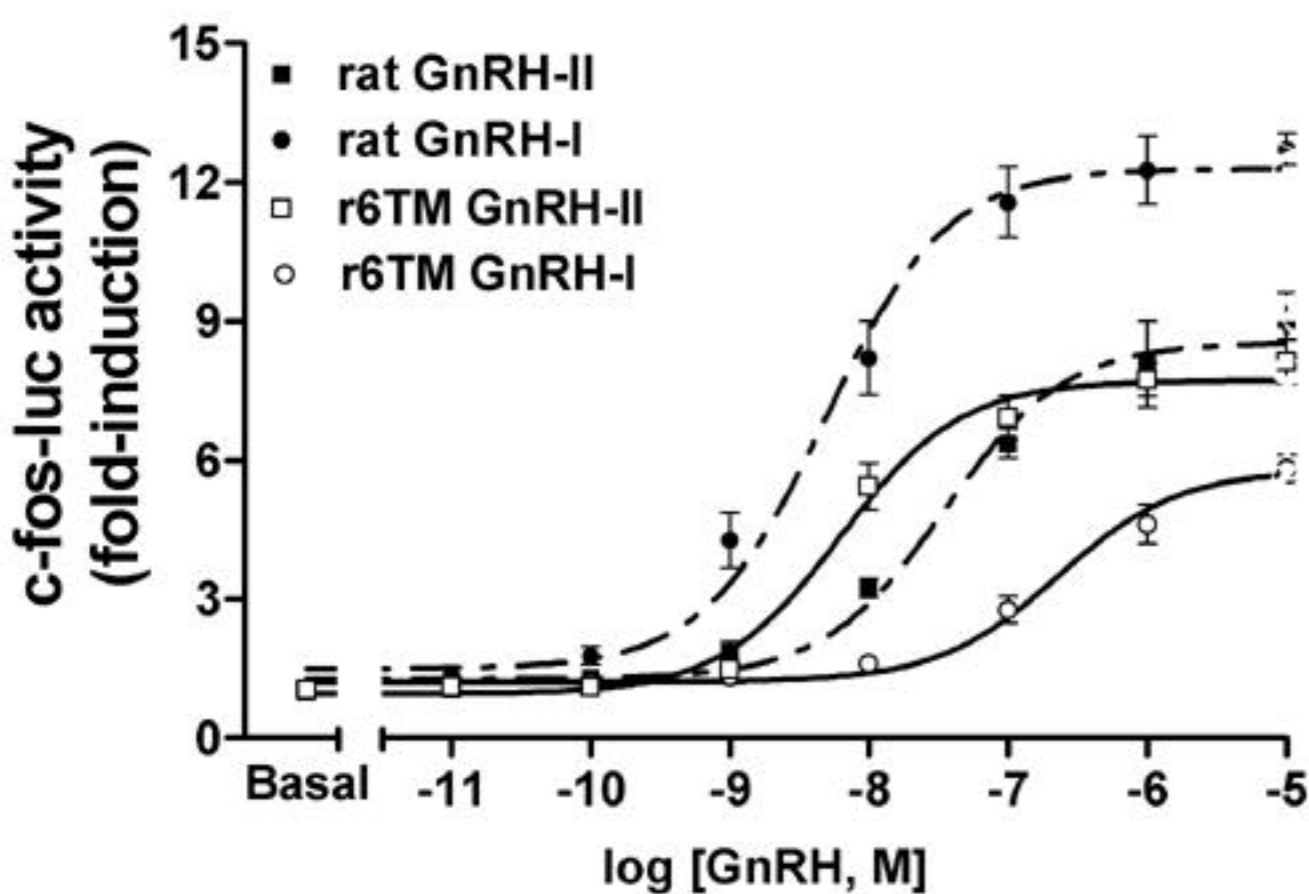


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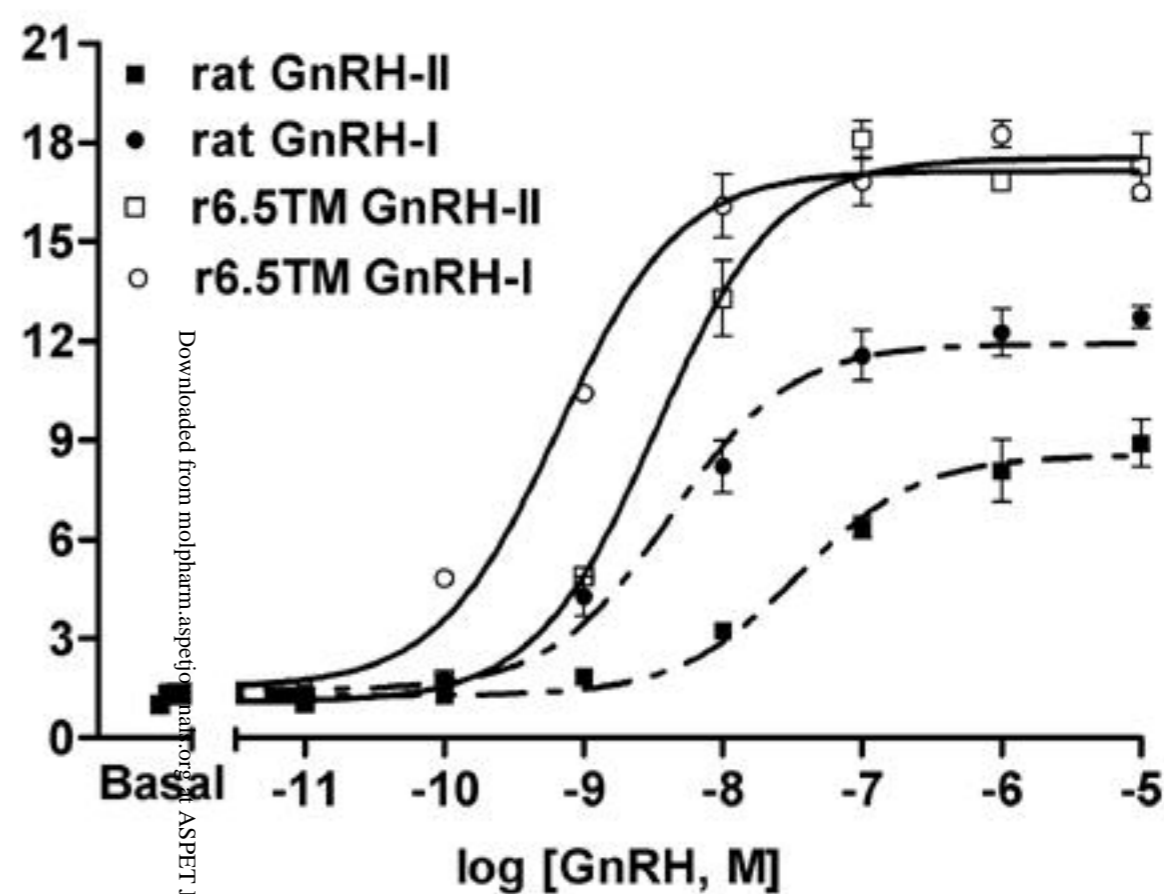


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D



E

			EL3		TMH7			
rGnRHR	288	...WYWF	DPEMLNRV	<b>SE</b> PVN	HF <b>FF</b> LF <b>AFL</b> NP <b>C</b>	FDPL...		320
gmGnRHR-2	286	...WYWF	SPTMLTEV	<b>PP</b> SL <b>S</b>	HI <b>L</b> FL <b>FG</b> LL <b>N</b> A <b>P</b>	LDPL...		318
rEL3L	286	...WYWF	<u>DPEMLNRV</u>	<b>SE</b> PVN	HF <b>FF</b> LF <b>AFL</b> NP <b>C</b>	LDPL...		318
rEL3S	286	...WYWF	<u>DPEMLNRV</u>	<b>SE</b> PVN	HI <b>L</b> FL <b>FG</b> LL <b>N</b> A <b>P</b>	LDPL...		318
r6TM	288	...WYWF	DPEMLNRV	<b>SE</b> PVN	HI <b>L</b> FL <b>FG</b> LL <b>N</b> A <b>P</b>	LDPL...		320
r6.5TM	288	...WYWF	DPEMLNRV	<b>SE</b> PVN	HF <b>FF</b> LF <b>AFL</b> NP <b>C</b>	LDPL...		320
				↑	↑	↑	↑↑	
				7.32	7.38	7.43	7.46	7.47

Fig. 3.

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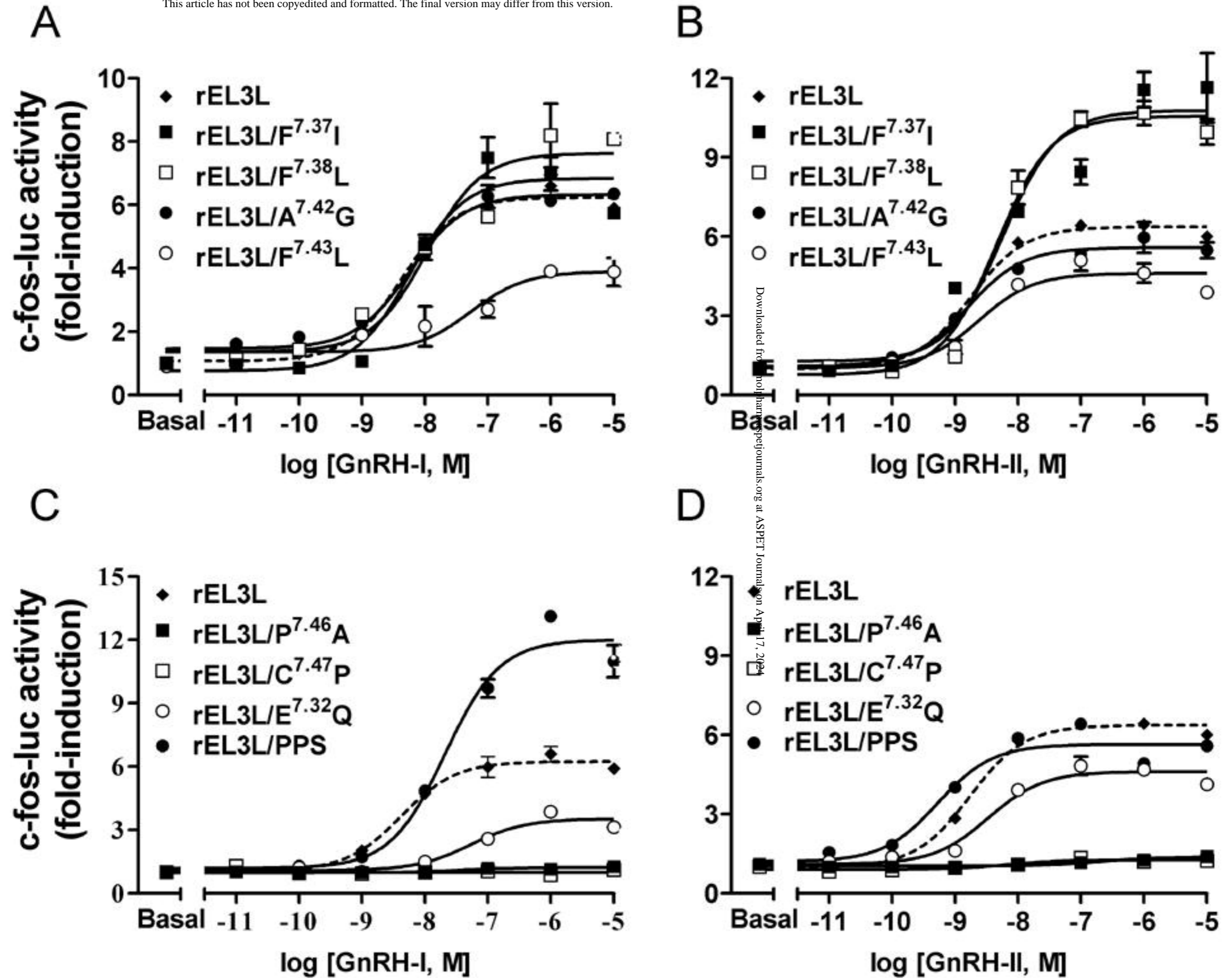


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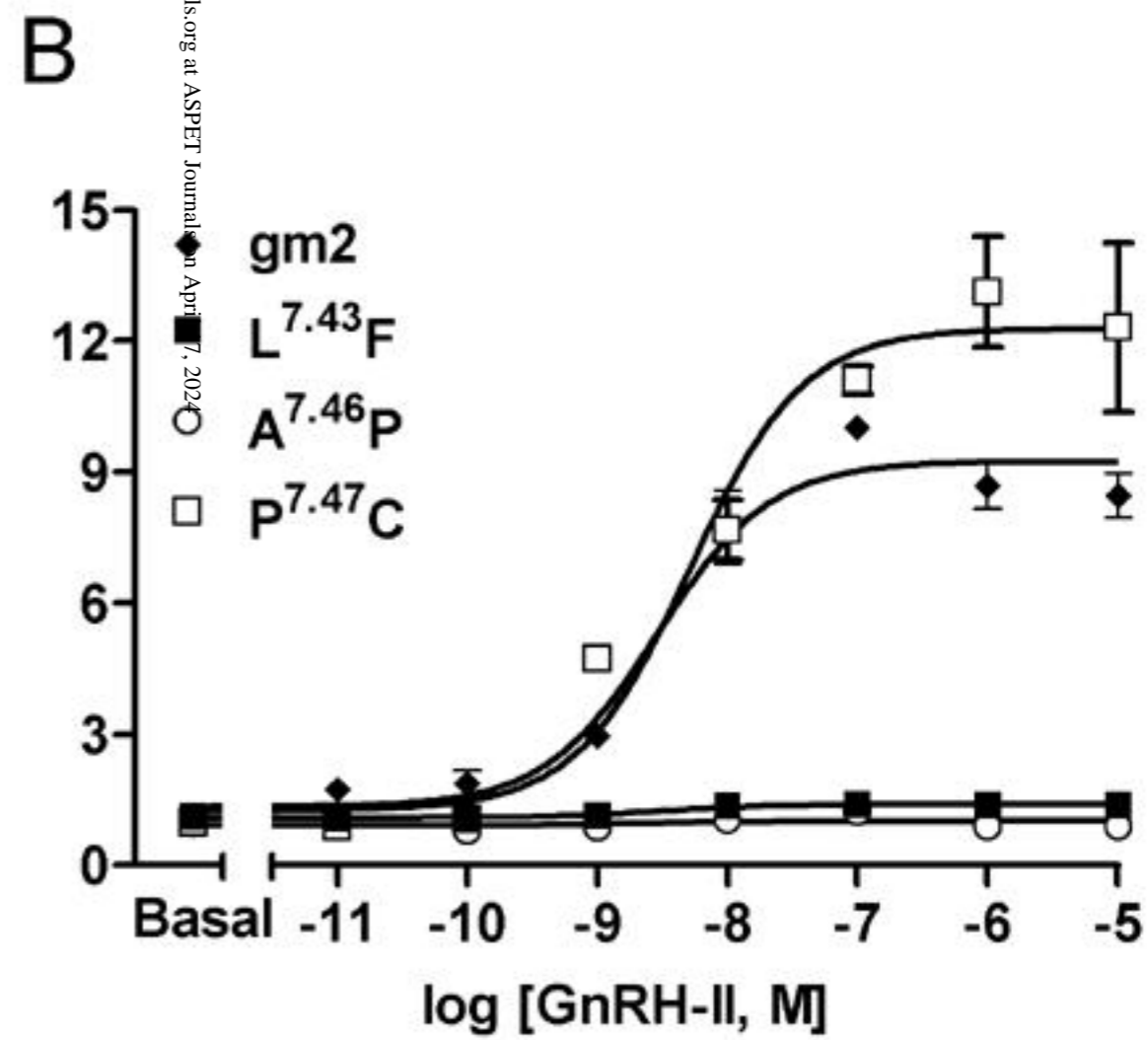
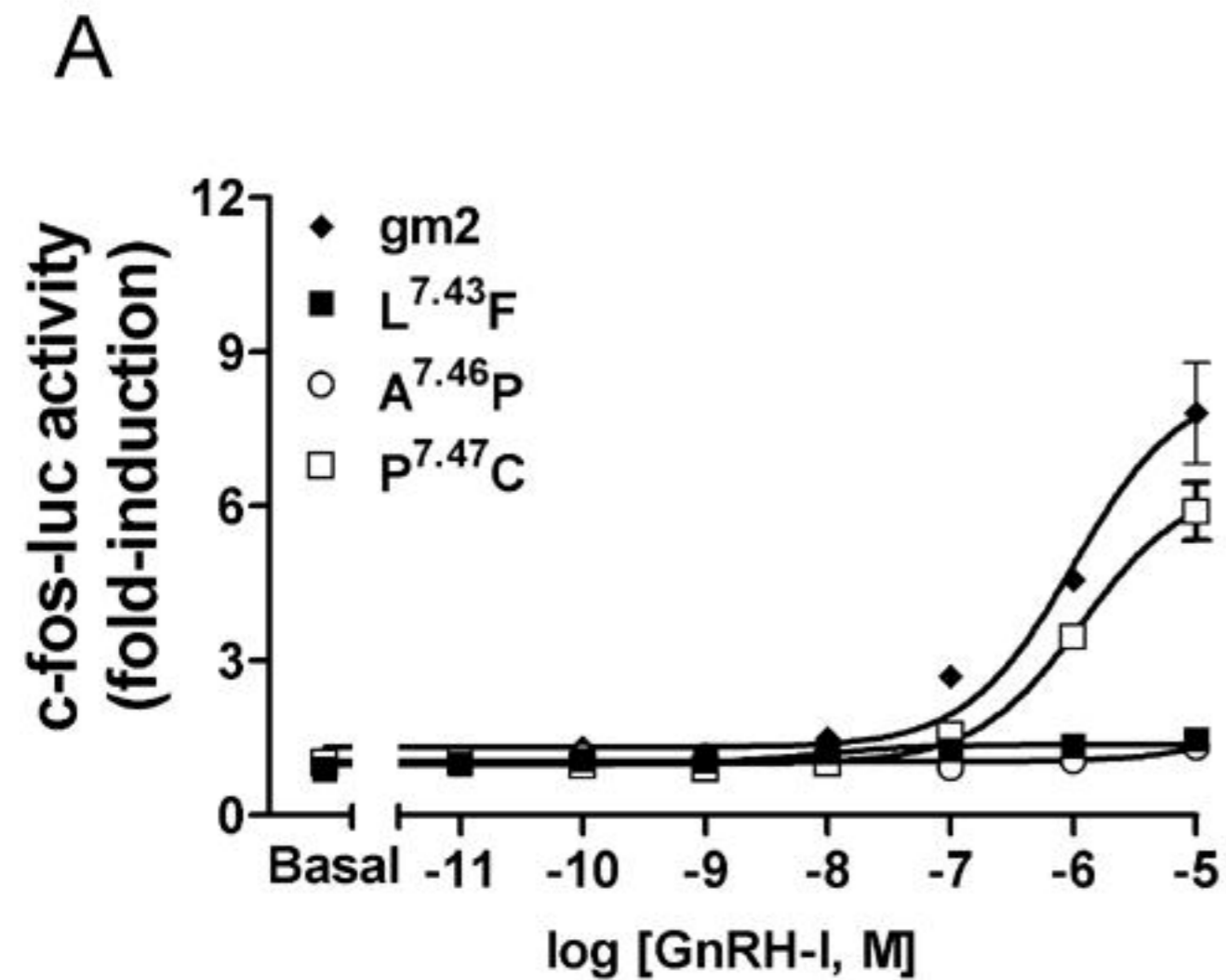




Fig. 5.

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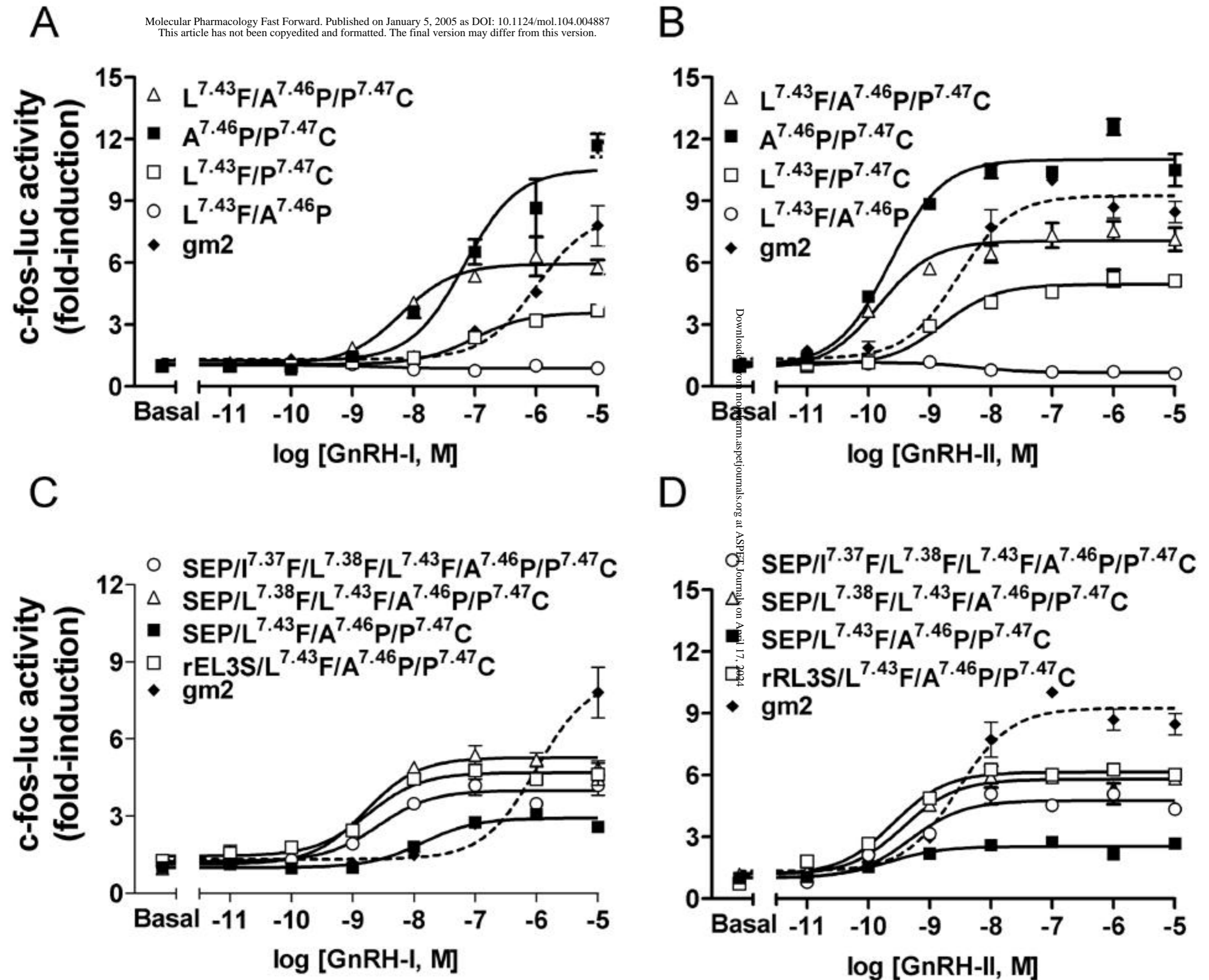


Fig. 6.

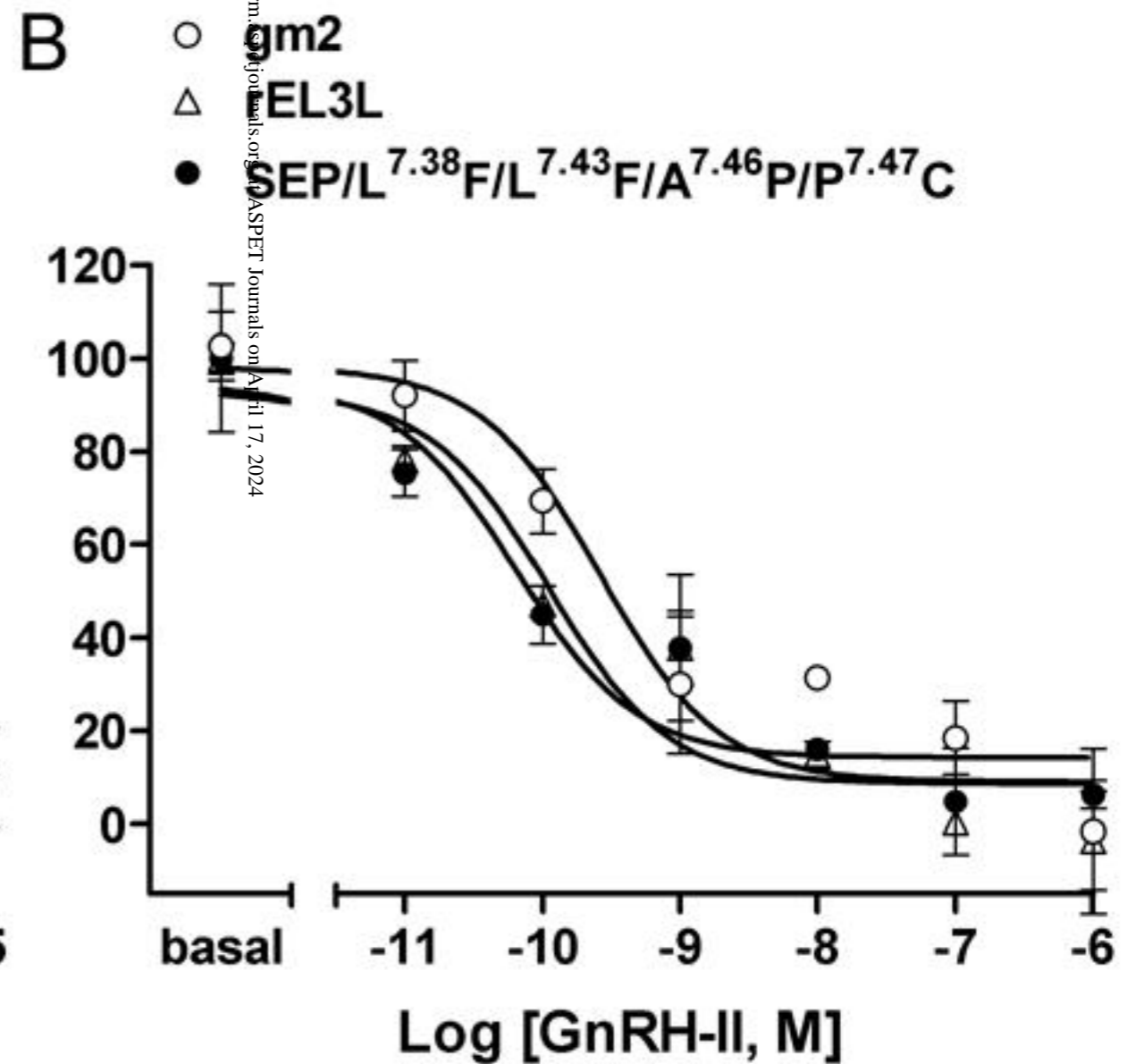
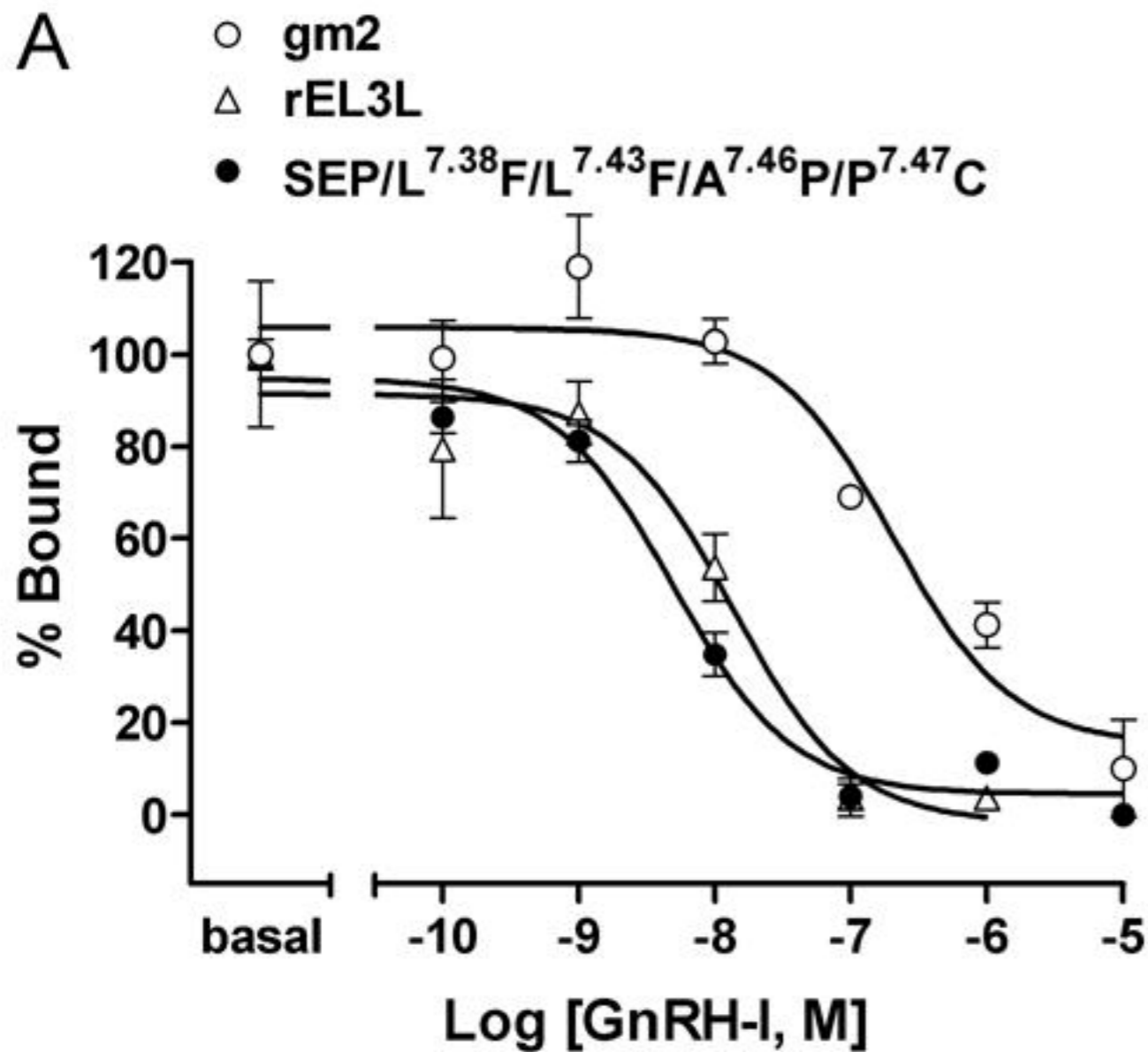


Fig. 7.

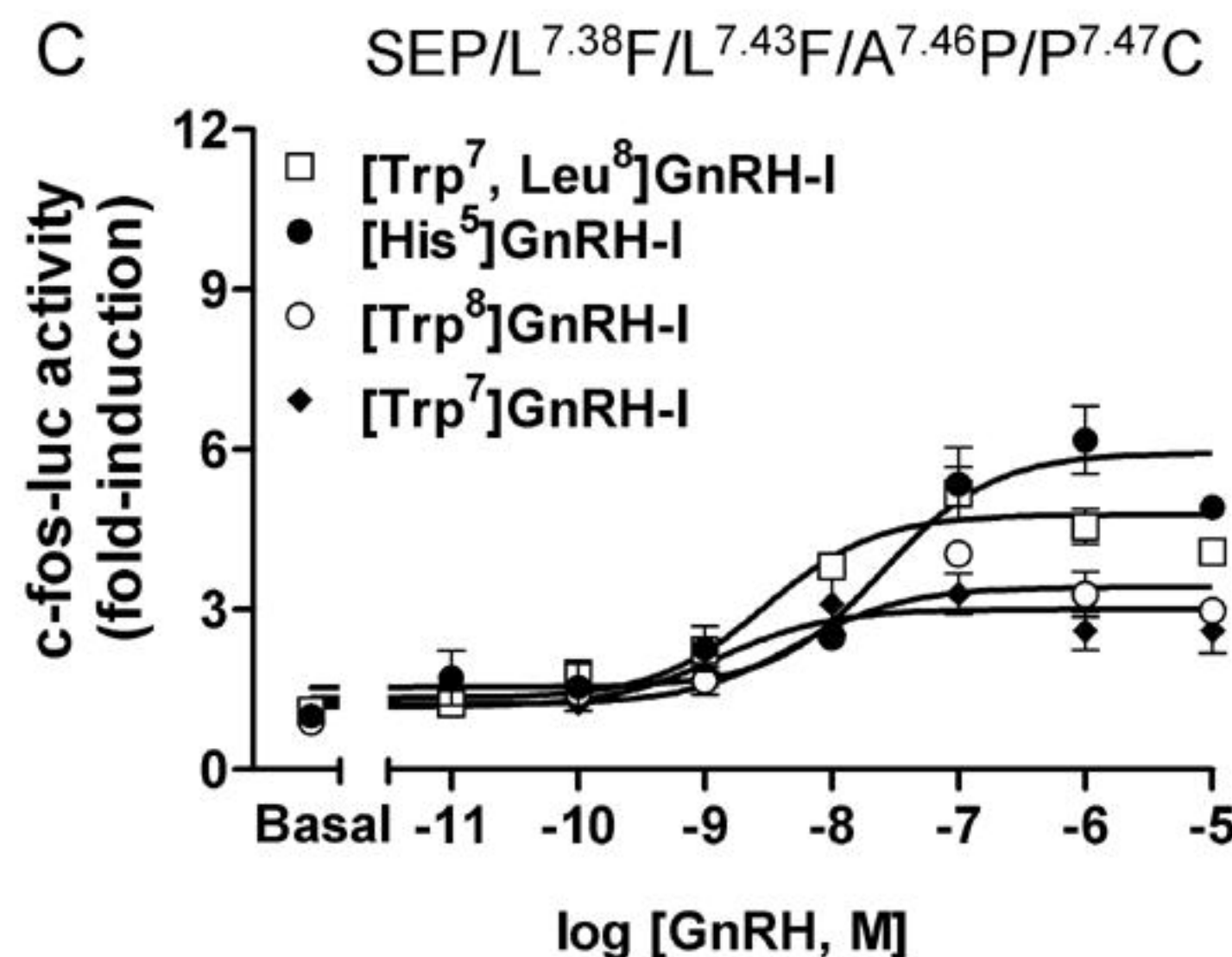
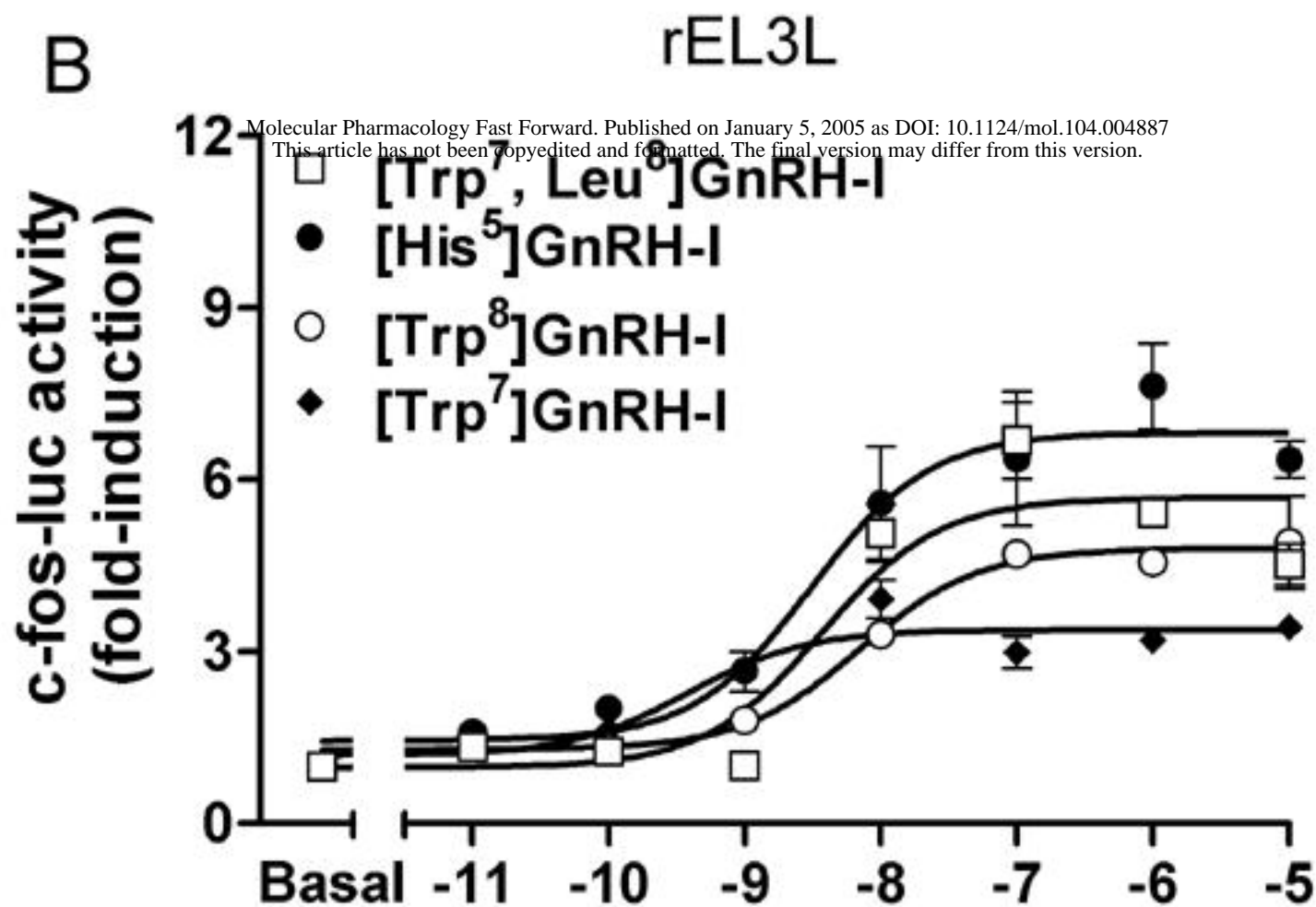
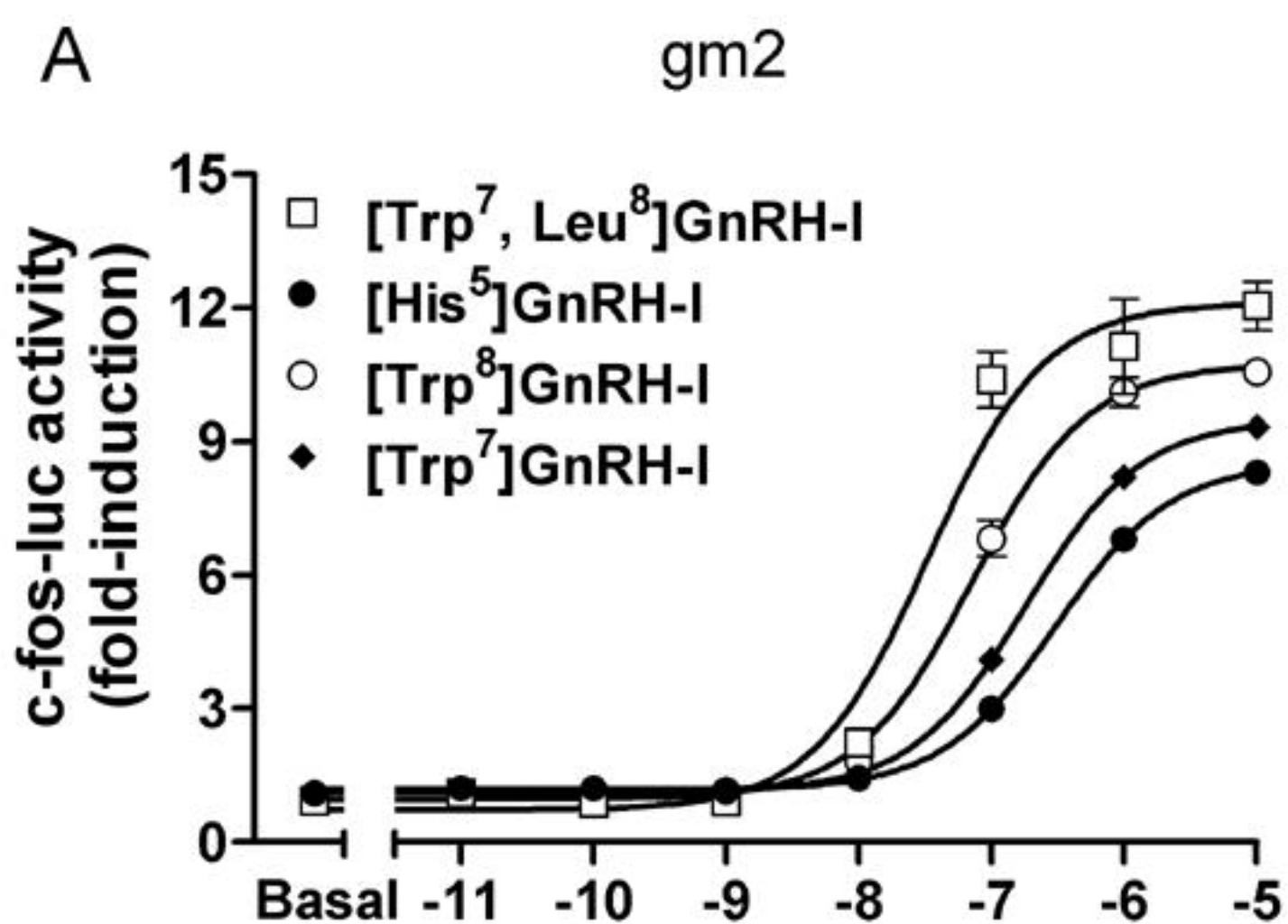


Fig. 8.

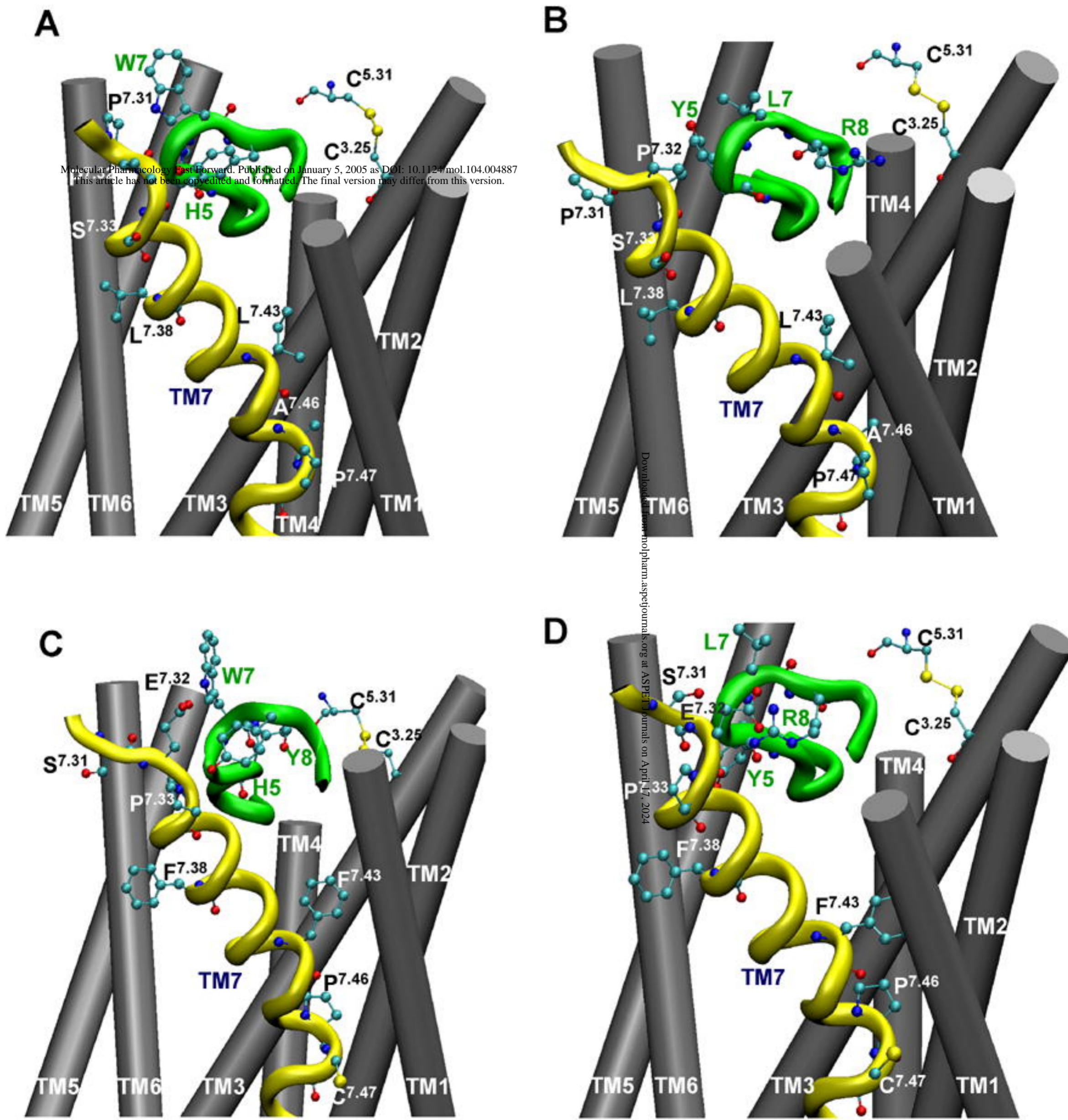


Fig. 9.

	<b>EL3</b>		<b>TMH7</b>	
	↓ ↓ ↓		↓   ↓   ↓ ↓	
Human-1	WYWFDPEMLN--LV <b>SDP</b> VN		H <b>F</b> FLFA <b>F</b> LN <b>PC</b> F	DPLIYGYFS
Rat-1	WYWFDPEMLN--RV <b>SEP</b> VN		H <b>F</b> FLFA <b>F</b> LN <b>PC</b> F	DPLIYGYFS
G-Monkey-2	WYWFSPTMLT--EV <b>PPS</b> IS		H <b>I</b> LFLF <b>G</b> LN <b>A</b> PL	DPLLYGAFT
Marmoset-2	WYWFSPSMLS--EV <b>PPS</b> IS		H <b>I</b> LFLF <b>G</b> LN <b>A</b> PL	DPLLYGAFT
Bullfrog-3	WYWFSPPEMLTSRKV <b>PPS</b> IS		H <b>I</b> LFLF <b>G</b> LN <b>T</b> CL	DPIIYGLFT
Xenopus-2	WYWFSPPEMLTEEKV <b>PPS</b> IS		H <b>I</b> LFLF <b>G</b> LN <b>T</b> CL	DPIIYGLFT
J-Medaka-1	WYWFFPDDLEG-KV <b>SHS</b> IT		H <b>I</b> LFI <b>F</b> LN <b>T</b> CL	DPIIYGLFT
Bullfrog-1	WYWFYPEIMEE-KV <b>SQS</b> T		H <b>I</b> LFI <b>F</b> LN <b>V</b> ACL	DPITYGLFT
J-Medaka-2	WYWFQPDMLR--VT <b>PEY</b> VH		H <b>I</b> L <b>F</b> V <b>F</b> LN <b>T</b> CC	DPVIYGFYT
Bullfrog-2	WYWFQPEMIY--LT <b>PEY</b> VH		H <b>S</b> L <b>F</b> LN <b>L</b> H <b>T</b> CT	DPLVYGLYT
Catfish-1	WYWFQPQMLH--VI <b>PDY</b> VH		H <b>V</b> <b>F</b> <b>F</b> V <b>F</b> LN <b>T</b> CC	DPVIYGFFT
Human angiotensin II 1 receptor			IAYFNN <b>C</b> L	NPLFYGFLG
Human bradykinin B2 receptor			MAYSNS <b>C</b> L	NPLVYVIVG
Human neuropeptide Y1 receptor			TAMIST <b>C</b> V	NPIFYGFLN
Human opioid mu1 receptor			LGYTNS <b>C</b> L	NPVLYAFLD
Human somatostatin-1 receptor			LGYANS <b>C</b> A	NPILYGFSL
Human oxytocin receptor			LASLNS <b>C</b> C	NPWIYMLFT

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