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Title Page

Thermodynamics and Docking of Agonists to the β2-Adrenoceptor Determined Using [^{3}H]-(R,R')-4-Methoxyfenoterol as the Marker Ligand

Lawrence Toll, Karolina Pajak, Anita Plazinska, Krzysztof Jozwiak, Lucita Jimenez, Joseph A. Kozocas, Mary J. Tanga, James E. Bupp, Irving W. Wainer,

Department of Chemistry, Medical University of Lublin, Lublin, Poland (KP, AP, KJ), SRI International, Menlo Park, CA, USA (LJ, JAK, MJT, JEP); Laboratory of Clinical Investigation, National Institute on Aging Intramural Research Program, Baltimore, MD, USA (IWW)

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Running Title Page

Running Title: Thermodynamics docking of β2-AR agonists

Corresponding author:

Lawrence Toll

Torrey Pines Institute for Molecular Studies

11350 SW Village Parkway

Port St. Lucie, FL 34987, USA

Phone: 772-345-4714

FAX: 772-345-3649

ltoll@tpims.org

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Abbreviations: G Protein Coupled Receptor, GPCR; Comparative Molecular Field Analysis, CoMFA; fenoterol, FEN; β-AR, β-adrenergic receptor; polyethyleneimine, PEI;

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Abstract

G protein coupled receptors (GPCRs) are integral membrane proteins that change conformation subsequent to ligand binding so that they can transduce signals from an extracellular ligand to a variety of intracellular components. The detailed interaction of a molecule with a GPCR is a complicated process that is influenced by the receptor conformation, thermodynamics, and ligand conformation and stereoisomeric configuration. To better understand the molecular interactions of fenoterol analogs with the β2-adrenergic receptor, we developed a new agonist radioligand for binding assays. [^{3}H]-(R,R')-methoxyfenoterol was used to probe the binding affinity for a series of fenoterol stereoisomers and derivatives. The results suggest that the radioligand binds with high affinity to an agonist conformation of the receptor, which represents about 25% of the total β_2-AR receptor population as determined with the antagonist [^{3}H]-CGP-12177. The β_2-AR agonists tested in this study have considerably higher affinity for the agonist conformation of the receptor and Ki values determined for fenoterol analogs model much better the cAMP activity of the β_2-AR elicited by these ligands. The thermodynamics of binding are also different when interacting with an agonist conformation, being purely entropy driven for each fenoterol isomer, rather than a mixture of entropy and enthalpy when the fenoterol isomers binding was determined using [^{3}H]CGP-12177. Finally, computational modeling identified the molecular interactions involved in agonist binding and allow for the prediction of additional novel β_2-AR agonists. The study underlines the possibility of using defined radioligand structure to probe a specific conformation of such shape-shifting system as the β_2-adrenoceptor.

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Introduction

The binding affinity of a compound to the β_2-adrenoceptor (β_2-AR) is routinely determined using competitive binding assays based upon the concentration dependent displacement of a marker radioligand. While this technique is used to characterize the binding of β_2-AR agonists and antagonists, the most often employed marker ligands are non-selective β-AR antagonists, which often have significant binding affinities to the β_1-AR, β_2-AR and β_3-AR (Baker, 2005; Brodde et al., 1983; Hoffmann et al., 2004; Joseph et al., 2004; Nikulin et al., 2006; Perrone et al., 2008; Staehelin et al., 1983; Toews et al., 1983). These radioligands include (-)-3-[^{125}I]-iodocyanopindolol ([^{125}I]-CYP) (Brodde et al., 1983), [^{3}H]-CGP-12177 (Staehelin et al., 1983), [^{125}I]-iodopindolol (Toews et al., 1983) and [^{3}H]-dihydroalprenolol (Perrone et al., 2008; Staehelin et al., 1983). One of the most widely used markers, [^{3}H]-CGP-12177, is a non-conventional antagonist of the β_1-AR that binds at two sites on the receptor (Joseph et al., 2004). Thus, it is not clear which site or sites on the β_2-AR interact with [^{3}H]-CGP-12177, nor how these interactions affect the identification and characterization of β_2-AR agonists.

The concerns associated with the use of [^{3}H]-CGP-12177 as a marker ligand was evident in our recent work on the characterization of the individual stereoisomers of the β_2-AR agonist fenoterol (Fen) and a series of Fen analogues, c.f. Fig. 1 (Jozwiak et al., 2007; Jozwiak et al., 2010a; Jozwiak et al., 2010b; Toll et al., 2011). During the course of these studies, we determined the binding thermodynamics of the four stereoisomers of Fen, (R,R')-, (R,S')-, (S,R')- and (S,S')-Fen to the β_2-AR (Jozwiak et al., 2010a). All of these compounds are full agonists of the β_2-AR with respect to the stimulation of cAMP

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accumulation in HEK293 cells stably transfected with β_2-AR (HEK-β_2-AR), and the binding studies were performed using membranes obtained from these cells. In these studies, [^{3}H]-CGP-12177 was used as the marker ligand and the binding affinities were expressed as K_{iCGP} values. The results indicated that there were significant stereochemistry-based differences in the binding mechanisms as this process was entropy-driven when (R,R')- and (R,S')-Fen were the ligands, while the binding of the (S,R')- and (S,S')- isomers was an enthalpy-driven process. In addition the calculated Hill's coefficients (n values) also differed as an n = ~1 was calculated for the (R,R')- and (R,S')-isomers and an n = ~2 was determined for the (S,R')- and (S,S')-isomers. These binding studies have also been used to develop Comparative Molecular Field Analysis (CoMFA) models for the interaction of the Fen analogs, and other agonists and antagonists for the β_2-AR, and we have been able to model the differential interactions of the stereoisomers. However, binding studies using the antagonist [^{3}H]-CGP-12177 may only explore a portion of the binding interactions.

It is well known that GPCRs, such as the β_2-AR, bind ligands in multiple conformations. In particular, antagonists seem to have similar affinities to most or all receptor conformations, while an agonist will stabilize the receptor in a conformation for which it has high affinity, but it will bind with low affinity to other "antagonist" conformations (Kent et al., 1980). Therefore, our (and most other) previous studies have primarily examined the interaction of Fen analogs and other agonists with the antagonist conformation. This is evident if one compares the binding affinity of (R,R')-Fen when competing with [^{3}H]-CGP-12177, K_i = 345 nM, versus its potency for stimulation of cAMP accumulation in the same cells, EC_{50} = 0.30 nM (Jozwiak et al., 2010b).

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In order to examine the binding of Fen analogs to a different conformation of the β_2-AR, we synthesized [^{3}H]-(R,R')-4-methoxyfenoterol, [^{3}H]-MFen, Fig.1, to use as the marker ligand in receptor binding studies (Kozocas et al., 2010). We have previously characterized MFen as a potent and selective β_2-AR agonist which stimulates cAMP accumulation in HEK-β_2-AR cells, (Toll et al., 2011), induces cardiomyocyte contractility in a mouse cardiomyocyte model, (Jozwiak et al., 2010b), and inhibits 1321N1 mitogenesis (Toll et al., 2011). Here we report the initial study of the use of this compound as a marker in the determination of β_2-AR agonist binding affinities, examine the thermodynamics of Fen binding to an "agonist" conformation, and employ molecular dynamics calculations to perform in silico docking of Fen analogs to this agonist conformation.

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Materials and Methods

Materials

The Fen analogues used in this study, Fig. 1, were synthesized as previously described (Jozwiak et al., 2007) and the preparation of [^{3}H]-MFen (25 Ci/mmol), Fig. 1, has been recently reported (Kozocas et al., 2010). Dulbecco's Modified Eagle Medium (DMEM) was purchased from Lonza Walkersville, Inc. (Walkersville, MD), fetal bovine serum (FBS) was purchased from Mediatech, Inc. (Manassas, VA), penicillin-streptomycin and geneticin (G418) were purchased from Invitrogen (Carlsbad, CA), sodium chloride and calcium chloride were purchased from Mallinckrodt (Phillipsburg NJ) and (*rac*)-propranolol, (R)-isoproterenol, ICI-118-551, Tris-HCl, Trizma Base, potassium chloride, magnesium chloride and D-(+)-glucose were purchased from Sigma-Aldrich (St. Louis, MO).

Membrane binding studies

HEK cells stably transfected with cDNA encoding human β_2-AR (HEK-β_2-AR, provided by Dr. Brian Kobilka, Stanford University Medical Center, Palo Alto, CA) were grown in DMEM containing 10% FBS and 0.05% penicillin-streptomycin with 400 μg/ml G418. The cells were scraped from the 150 x 25 mm plates and centrifuged at 500 x g for 5 min. The pellet was washed twice by homogenization in Tris-HCl [50 mM, pH 7.7] and centrifugation at 27,000 x g for 10 min. The pellet was resuspended in Tris-HCl [15 mM, pH 7.4], containing 120 mM sodium chloride, 5.4 mM potassium chloride, 1.8 mM calcium chloride, 0.8 mM magnesium chloride, and 5 mM glucose. The binding assays contained 3.9 nM [^{3}H]-MFen and 60 μg cell membranes, in a volume of 1.0 ml. The mixture was incubated at 25°C for 2 h and filtered over glass fiber filters soaked in 0.05%

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polyethyleneimine (PEI). Nonspecific binding was determined using 10 μM (*rac*)-propranolol. The reaction was terminated by filtration using a Tomtec 96 harvester (Orange, CT) through glass fiber filters. Bound radioactivity was counted on a Pharmacia Biotech beta-plate liquid scintillation counter (Piscataway, NJ) and expressed in counts per minute. Saturation experiments were conducted using concentrations ranging from 0.2 to 20 nM [^{3}H]-MFen.

Docking simulations

The 2RH1.pdb and 3POG.pdb molecular models of the β_2-AR were used in the simulated docking studies. Ligand molecules were prepared using HyperChem 6.03 (HyperCube Inc., Gainesville, FL) software using Model Build procedure. Molegro Virtual Docker (MVD v. 2010.4.0.0) software was employed for docking simulations. The MolDock SE search algorithm was used, and the number of searching runs was set to 100. The following parameters were set during docking simulation: population size = 50, maximum iteration = 1500, energy threshold = 100.00, max steps = 300. The estimation of ligand-protein interactions was described by the MVD implemented scoring functions: MolDock Score, Rerank Score, Hbond Score, Similarity Score, Docking Score.

Statistical analysis

Results were analyzed by non-linear regression analysis using the program Graphpad/Prism (ISI, San Diego, CA). For competition experiments, IC_{50} values and Hill's coefficients (n) were determined using at least six concentrations of each Fen analog. The Ki values were calculated by the method of Cheng and Prusoff (Cheng and Prusoff, 1973).

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Results

Characterization of [^{3}H]-MFen binding to membranes from HEK-β_2-AR cells

Saturation analysis of [^{3}H]-MFen binding to membranes from HEK-β_2-AR cells indicated, a single binding component with a K_{dMFen} of 4.88 ± 0.41 nM and a B_{max} of 2136 ± 114 fmol/mg protein. Non-specific binding represented less than 20% of total binding (Fig. 2). The binding was inhibited by the non-selective β-AR antagonist (*rac*)-propranolol and the selective β_2-AR antagonist ICI 118,551 indicating that [^{3}H]-MFen specifically bound to the β_2-AR. The calculated K_{dMFen} value was ~100-fold lower than the previously reported β_2-AR affinity of MFen determined using the same cellular membranes and [^{3}H]-CGP-12177 as the marker ligand, K_{iCGP} = 473 nM, Table 1 (Jozwiak et al., 2007). The calculated B_{max} value was also lower than the previously reported value, 8,901 ± 1,161 fmol/mg protein, also determined using [^{3}H]-CGP-12177 as the marker ligand (Jozwiak et al., 2007). These results suggest that [^{3}H]-MFen binds with high affinity to an agonist conformation of the receptor, and that, under these conditions, the agonist conformation represents only about 25% of the total β_2-AR receptor population.

Unlike saturation analysis, the kinetic analysis of [^{3}H]-MFen binding suggested that there was more than one binding conformation or component. The analysis of the relationship between specific binding and time revealed that the data fit better to a two-component model of binding than a single component, Fig. 3, making it impossible to calculate definitive kinetic binding constants. Non-linear regression analysis of the data indicated that the association rate appears to be biphasic, with k_{on} of 1.21 min^{-1} for about 35% of the sites and 0.016 min^{-1} for the remaining 65%. The same results were obtained

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from the analysis of the k_{off} data as approximately 35% of the receptors have a k_{off} of 0.018 min^{-1} and 65% have a k_{off} of 0.20 min^{-1}. The data supports the supposition that [^{3}H]-MFen binds to at least two sites on or conformations of the β_2-AR, although this cannot be readily detected by saturation analysis.

Determination of β_2-AR binding affinities using [^{3}H]-MFen as the marker ligand

[^{3}H]-MFen was used as the marker ligand in the determination of the β_2-AR binding affinities (K_{iMFen}) of 22 fenoterol analogs, isoproterenol, propranolol and ICI-118-551 and the data is presented in Table 1. In these experiments a low concentration (3.9 nM) of [^{3}H]-MFen was used consequently, greater than 75% of the binding represents the high affinity binding conformation. The K_{iMFen} value observed for the non-selective β-AR antagonist (*rac*)-propranolol, 3.69 nM, was nearly 10-fold higher than the K_{iCGP} value, 0.46 nM, determined using [^{3}H]-CGP-12177 as was the calculated affinity of the selective β_2-AR antagonist ICI 118,551, K_{iMFen}, 2.52 nM and K_{iCGP} = 0.60 nM, Table 1. The opposite result was obtained with the β_2-AR agonist isoproterenol as the K_{iMFen} was 79-fold lower than the K_{iCGP}, 2.44 nM and 192 nM, respectively, and more in line with the EC_{50} value that induced cAMP accumulation in HEK-β_2-AR cells, 0.20 nM, Table 1. The calculated Hill's coefficients for the agonist compounds were slightly less than 1.0, suggesting there is some heterogeneity of binding, consistent with the saturation and kinetic experiments. Interestingly, the Hill coefficient for the antagonists *rac*-propranolol and ICI 118,551 were 2.10 and 1.84, respectively, suggesting that some degree of positive cooperativity of binding may exist.

The data obtained with the 22 fenoterol analogs tested in this study are consistent with the supposition that K_{iMFen} values better reflect the β_2-AR agonist properties than

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K_{iCGP} values. The apparent affinities uniformly increased when [^{3}H]-MFen was the marker ligand relative to the K_{iCGP} values for each of the compounds tested, Table 1 (Competition curves are shown in Supplemental Figure S1). The changes ranged from a 116-fold decrease in K_i (increase in affinity) for (R,R')-MFen, to a 4-fold decrease when (S,S')-1-naphthylfenoterol and (R,S')-2-naphthylfenoterol were studied. Nevertheless, there was a good correlation between the Ki values using the two radioligands, with a correlation coefficient of $R^2 = 0.7899$ (see Supplemental Figure S2). In general, the magnitude of the change was dependent upon the configuration at the β-OH carbon with an R configuration producing a greater enhancement in the binding affinity when [^{3}H]-MFen was the marker ligand.

When the K_i values were compared to the corresponding EC_{50} values determined for the stimulation of cAMP accumulation in HEK-β_2-AR cells, the magnitude of the K_{iMFen} values were more reflective of this activity than the K_{iCGP} values, Table 1. For each compound, the binding affinity using [^{3}H]-MFen was closer to the EC50 value for cAMP accumulation that was the Ki for [^{3}H]-CGP-12177 binding. In addition, the pKi [^{3}H]-MFen is significantly better correlated with pEC_{50} cAMP than the pKi [^{3}H]-CGP-12177 , 0.5532 vs. 0.4143, respectively (Supplemental Figure S2). These results suggest that the high affinity site probed by [^{3}H]-MFen is associated with the cAMP activity of the β_2-AR and that all of the agonists used in this study, bind to this site to a greater extent than to a lower affinity site probed by [^{3}H]-CGP-12177.

Effect of GTP on β2-AR binding

Experiments were conducted to determine how GTP and GTPγS affect binding of [^{3}H]-MFen and compare that to their effect on the binding of the antagonist [^{3}H]-CGP-12177.

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GTP and its non-hydrolyzable analog GTPγS both dose dependently reduced binding of [^{3}H]-MFen, although the potency of GTP and particularly GTPγS were lower than what might be expected from literature values (Fig. 4). Presumably this is not competitive antagonism but reflect a GTP-induced decrease in agonist affinity. Interestingly, GTP and GTPγS did not have much effect on [^{3}H]-CGP-12177 binding. As expected, [^{3}H]-CGP-12177 binding remained high in the presence of GTP. However, surprisingly, GTP did not induce a decrease in affinity of R,R'-fenoterol, and slightly increased the affinity of R,R'-methoxyfenoterol (Table 2).

Thermodynamic studies

In a previous study, the effect of temperature on the binding of (R,R')-Fen, (R,S')-Fen, (S,R')-Fen, (S,S')-Fen, propranolol and isoproterenol was determined using [^{3}H]-CGP-12177 as the radioligand and the data were subjected to van't Hoff analysis (Jozwiak et al., 2010a). In the current study, the temperature dependence of the K_{iMFen} values of the same test compounds was determined at 4°C, 25°C and 37°C, Table 3. The three temperature points were used to construct van't Hoff plots ($\ln(1/Ki)$ vs. $1/T$), which were further employed to calculate enthalpic and entropic contribution to the free energy change of binding by linear regression of the equation $\ln(1/K_i) = \frac{\Delta S^\circ}{R} - \frac{\Delta H^\circ}{R}\frac{1}{T}$. ΔH°.

ΔS° and ΔG° values calculated using [^{3}H]-MFen affinity data are presented in Table 4. The calculations indicate that the binding of all of the test compounds was purely entropy driven, $\Delta H^\circ > 0$ and $-T\Delta S^\circ < 0$. These results differ from the results of the previous study using [^{3}H]-CGP-12177 as the marker ligand in which the binding of (S,S')-Fen was an enthalpy driven process (ΔH° = -35.8 kJ/mol; $-T\Delta S^\circ$ =+6.3 kJ/mol) and the binding of

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(S,R')-Fen, isoproterenol and propranolol were combined enthalpy/entropy driven processes, Table 3 (Jozwiak et al., 2010a). This observation was confirmed using the approach developed by Borea, et al., c.f. (Merighi et al., 2010), in which the data obtained in this study and the data previously obtained using [^{3}H]-CGP-12177 as the radioligand (Jozwiak et al., 2010a) were placed in a scatter plot of $-T\Delta S°$ versus $\Delta H°$. As seen in Fig. 5, all of the data from the current study was located within the quadrant associated with an entropy driven process, which is in contrast to the data obtained using [^{3}H]-CGP-12177, in which the data span quadrants.

Simulated docking studies

The hypothesis that [^{3}H]-CGP-12177 and [^{3}H]-MFen can be used to probe different conformations of the β_2-AR was tested using simulated receptor-ligand docking studies. Two molecular models of the β_2-AR binding site have been recently reported and were used in the studies: 1) the 2rh1.pdb model {β_2-AR-In} derived using β_2-AR co-crystalized with (*S*)-carazolol, which is regarded as an inactive form of the receptor (Rasmussen et al., 2007); and 2) the 3pog.pdb model {β_2-AR-Ac} obtained from β_2-AR co-crystalized with the agonist, BI-167,107 and the NB90 nanobody, which is regarded as an active form of the receptor (Rasmussen et al., 2011).

The lowest energy poses obtained in docking simulations of (S)-CGP-12177 were obtained with the β_2-AR-In model, Fig. 6a. In these simulations, the position of (S)-CGP-12177 shares a number of similarities with the position (S)-carazolol co-crystalized in the binding site of the β_2-AR-In model. As depicted in Fig 6a, there is a network of four hydrogen bonds formed between the amino and the beta-hydroxy moieties of (S)-CGP-12177 and two protein residues, Asp^{113} of TM3 and Asn^{312} of TM7. In addition (S)-

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CGP-12177 forms a hydrogen bond with Ser^{203} of TM5. A similar interaction was originally observed between (S)-carazolol and the β_2-AR in the crystal model, 2RH1.pdb. Thus, docking of (S)-CGP-12177 reveals the mode of binding conserved for other antagonists or inverse agonists like (S)-carazolol or ICI-118,551 interacting with the receptor.

The inward shift of the TM5 in the ligand binding domain of the β_2-AR-Ac model results in a condition such that the (R,R')-MFen molecule can achieve a similar network of four hydrogen bonds between amino and beta-hydroxy moieties of the ligand and Asp^{113} and Asn^{312} residues, as depicted in Fig. 6b. In addition, Ser^{203} and Ser^{207}, of TM5, interact with two *meta*-hydroxyl moieties of MFen, and Lys^{305} of TM7 interacts with the 4'-methoxy moiety of the ligand. The latter interaction for compounds like formoterol and fenoterol (having para-hydroxy moiety at the aminoalkyl tail) was recently proposed as an important factor in disrupting the ionic lock switch between Lys^{305} and Asp^{192} of extracellular loop 2 occurring in the inactive state and postulated to break during the activation of the receptor (Bokoch et al., 2010). Thus the docking simulations support the assumption that binding of (R,R')-MFen should stabilize the active form of the receptor.

The binding of the marker ligands to the β_2-AR-Ac and β_2-AR-In models was also examined using the scoring functions, MolDockScore values, generated by MVD software, which energetically characterizes the simulated ligand – receptor complexes. The MolDockScore value calculated for the (R,R')-MFen- β_2-AR-Ac complex, shown in Fig.6b, was significantly lower than the analogous value calculated for (R,R')-MFen- β_2-AR-In complex, -136.98 kJ/mol vs. -132.53 kJ/mol, respectively. Interestingly, the difference in MolDockScore values was very small in docking simulations between

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antagonist (S)-CGP-12177 and β_2-AR-In (Fig.5a), compared to β_2-AR-Ac models, -116.60 kJ/mol and -115.82 kJ/mol, respectively. This result is consistent with the observation that antagonists bind with roughly equal affinity to agonist and antagonist conformations of GPCRs and suggests that the binding of (S)-CGP-12177 to the inactive conformation is slightly more favorable than its interaction with active conformation.

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Discussion

The thermodynamics of the binding of agonists and antagonists to β-ARs have been described as fundamentally different processes in which the binding of an agonist is enthalpy-driven while the binding of an antagonist is entropy-driven (Contreras et al., 1986; Miklavc et al., 1990; Weiland et al., 1979). This observation was generalized as the principle of "thermodynamic agonist-antagonist discrimination" (Borea et al., 2000). We have recently reported the results of a study of the binding thermodynamics of (R,R')-Fen, (S,S')-Fen , (R,S')-Fen, and (S,R')-Fen to the β_2-AR (Jozwiak et al., 2010a). In this study, the binding affinities were determined at five different temperatures using [^{3}H]-CGP-12177 as the marker ligand and cellular membranes obtained from HEK-β_2-AR cells. The data indicated that the binding of (S,S')- and (S,R')-Fen were predominately enthalpy-driven processes while the binding of (R,R')- and (R,S')-Fen were entropy-driven. Since all of the Fen stereoisomers were full β_2-AR agonists in the HEK-β_2-AR cells, the results were inconsistent with the principle of "thermodynamic agonist-antagonist discrimination." In the discussion of this inconsistency we suggested that the results of our study might reflect the fact that the β_2-AR exists in an inactive (R) conformation and one or more ligand-specific active conformations (R*^{n}) (Seifert and Dove, 2009) and that displacement binding studies using [^{3}H]-CGP-12117, a high-affinity neutral antagonist (Baker et al., 2008), may reflect the relative affinity of the Fen stereoisomers for the inactive receptor state. We also suggested that a potential approach to clarifying these interactions was to conduct the displacement binding studies with the β_2-AR agonist [^{3}H]-(R,R')-MFen and this manuscript reports the results of this study.

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The data from the current study indicate that the binding properties of [^{3}H]-MFen are what one would expect for a high affinity β_2-AR agonist, and similar to those previously described for [^{3}H]-formoterol and earlier studies with [^{3}H]hydroxybenzylisoproterenol (Lefkowitz and Williams, 1977; Mak et al., 1994) . [^{3}H]-MFen has high affinity for β_2-AR with a K_d = 4.88 nM, and binding is decreased by the presence of GTP analogs. In addition, saturation and kinetic analyses indicate that the binding occurs at a single high affinity binding site, or perhaps two independent but high affinity sites, which appears to differ from the site probed by [^{3}H]-CGP-12177.

While the data from the saturation analysis indicated that [^{3}H]-MFen binds to a single high affinity conformation of the receptor, the results from association and dissociation experiments suggested binding to two conformations of the receptor, as both curves fit 2-binding site models better than single site models. In addition, low Hill coefficients for the agonist inhibition of [^{3}H]-MFen binding also are consistent with two potential binding sites or conformations. These results suggest the potential of two independent high affinity receptor conformations, since kinetic experiments, but not equilibrium experiments, such as saturation, can identify different binding components with roughly equal affinity but different kinetics. These results are consistent with biophysical experiments that demonstrated two kinetically distinguishable conformational states after agonist binding (Swaminath et al., 2004).

In the analysis of the [^{3}H]-MFen saturation binding studies with the membranes from the HEK-β_2-AR, the maximum binding capacity, B_{max} value, was 2136 fmol/mg protein. This was significantly lower than the B_{max} value calculated for [^{3}H]-CGP-12177 binding to the same membranes, 8,901 fmol/mg protein (Jozwiak et al., 2007). The most

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reasonable explanation for this observation is that in the HEK-β_2-AR cell line, the majority of the β_2-AR receptors reside in a conformation that has low affinity for [^{3}H]-MFen. Therefore, the data suggest that, in these saturation binding studies, [^{3}H]-MFen probes a subset of the conformations probed by [^{3}H]-CGP-12177. In the binding experiments, the low affinity binding is absent. This is consistent with the results from the docking studies in which the difference in MolDockScore values was very small in docking simulations between (*S*)-CGP-12177 and either β_2-AR-In (Fig.6a), and β_2-AR-Ac models, -116.60 kJ/mol and -115.82 kJ/mol, respectively, while the difference for MFen was relatively much larger (-136.98 kJ/mol vs. -132.53 kJ/mol). Due to the high dissociation rate of agonists from low affinity sites, presumably, bound radioactivity dissociates during the washing in a filtration assay, and accordingly, the [^{3}H]-agonist does not appear to easily recognize a low affinity binding site. This is consistent with the Hill coefficients close to but slightly less than 1.0 for agonists inhibiting [^{3}H]-MFen binding. Interestingly, agonists also have high Hill coefficients when inhibiting [^{3}H]-CGP-12177 (see Supplemental Table T1). Furthermore, GTP and GTPγS, which stabilize a low affinity agonist conformation, have very little effect on [^{3}H]-CGP-12177 binding, or on the ability of the fenoterol analogs to compete with [^{3}H]-CGP-12177 binding. Together, these data indicate that [^{3}H]-CGP-12177 binds preferably to and stabilizes the low affinity agonist conformation, without much overlap with the high affinity agonist conformation, the one to which [^{3}H]-MFen presumably binds.

The assumption that [^{3}H]-MFen binds to a high affinity conformation of the β_2-AR is supported by the comparative K_i values determined using [^{3}H]-MFen and [^{3}H]-CGP-12177 as the marker ligands. For the Fen analogs and (*R*)-isoproterenol, which are

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all full β_2-AR agonists in the HEK-β_2-AR cell line, each of the K_{iMFen} values were considerably lower (i.e. higher affinity) than the corresponding K_{iCGP} values, Table 1. The magnitude of the increase in binding affinity when using [^{3}H]-MFen was, to a great extent, dependent upon the configuration at the β-OH carbon, with an R configuration producing a greater enhancement in the binding affinity when [^{3}H]-MFen was the marker ligand. (R,R')-Fen and (R,S')-Fen far prefer binding to the high affinity state probed by [^{3}H]-MFen as the K_{iCGP}/K_{iMFen} ratios are 86-fold and 20-fold, respectively. (S,R')-Fen and (S,S')-Fen have little preference between the two states as the K_{iCGP}/K_{iMFen} ratios are 6-fold and 8-fold, respectively. The opposite effect was observed with the two β-AR antagonists used in this study, propranolol and ICI-118-551, as the K_{iMFen} values were 8-fold and 4-fold higher than the corresponding K_{iCGP} values, Table 1. The apparent decrease in binding affinity when using [^{3}H]-MFen suggests that the antagonists have a lower, but still significant affinity for the high affinity agonist conformation of the receptor. For each of the agonists, the K_{iMFen} values were consistent with ligand potency and there was a considerably better correlation between binding affinity and functional activity measured as EC_{50} values for stimulation of cAMP accumulation.

The data from the thermodynamic studies also support the hypothesis that the binding studies using [^{3}H]-MFen reflect ligand binding to a subset of the conformations probed by [^{3}H]-CGP-12177. The results indicate that the binding to the conformation probed by [^{3}H]-MFen was entropy-driven for all of the competing ligands used in the thermodynamic section of the study, including the antagonists, Table 3. In our previous studies using [^{3}H]-CGP-12177, the binding of (S,S')-fenoterol to the β_2-AR was found to be a purely enthalpy-driven process, the binding of (S,R')-fenoterol, (R)-isoproterenol

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and *rac*-propranolol were enthalpy-entropy driven and the binding of (R,S')-fenoterol and (R,R')-fenoterol were purely entropy-driven, Table 3, Fig. 5. (Jozwiak et al., 2010a). In all, these results suggest that the thermodynamic properties obtained using [^{3}H]-CGP-12177 represent the sum total of multiple factors including unequal distributions of high and low affinity receptor conformations leading to a mixture of enthalpy and entropy driven processes.

The hypothesis that [^{3}H]-MFen can be used to explore an active, high affinity conformation of the β_2-AR was tested using simulated receptor-ligand docking studies employing the 2rh1.pdb model {β_2-AR-In} regarded as an inactive form of the receptor (Rasmussen et al., 2007) and the 3pog.pdb model {β_2-AR-Ac} which is regarded as an active form of the receptor (Rasmussen et al., 2011). The docking of [^{3}H]-CGP-12177 and [^{3}H]-MFen in the β_2-AR-In and β_2-AR-Ac models confirmed a conserved binding mode proposed earlier for this group of molecules (Weis and Kobilka, 2008), with amino and β-hydroxy groups of both ligands trapped in the network of cross-interactions with Asp^{113} and Asn^{312} residues. Small differences in topological organization of ligand binding sites in the β_2-AR-In and β_2-AR-Ac models allows both MFen and CGP-12177 to adopt positions in which their aromatic ring systems may exercise optimized interactions with Ser^{203} ((S)-CGP-12177 and (R,R')-MFen) and Ser^{207} ((R,R')-MFen) of TM5, Fig. 6. The comparison of the MolDockScore functions generated during docking simulations suggest that (R,R')-MFen should preferentially bind to the β_2-AR-Ac model while (*S*)-CGP-12177 would bind to both the β_2-AR-In and β_2-AR-Ac models, with a slight preference for the inactive conformation of the receptor. This is consistent with the relative K_i values obtained in the saturation binding studies utilizing the two probes.

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In conclusion, the results of this study indicate that [^{3}H]-MFen can be used as a probe in the determination of binding affinities to the β_2-AR. The data also indicate that this compound binds to a high affinity active conformation of the receptor and allows for the characterization of selective β_2-AR agonists. [^{3}H]-MFen may also be useful in the experimental verification of predictions made using the β_2-AR-Ac model and for QSAR studies aimed at the development of highly selective and active β_2-AR agonists.

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Authorship Contributions.

Participated in research design: Toll, Wainer, Jozwiak

Conducted experiments: Jimenez, Pajak, Plazinska

Contributed new reagents of analytic tools: Kozocas, Tanga, Bupp

Performed data analysis: Jimenez, Toll, Jozwiak

Wrote or contributed to the writing of the manuscript: Toll, Wainer, Jozwiak

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Footnotes:

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Legends for Figures

Figure 1. The structures of the compounds used in this study, where only the (R,R') configurations of the stereoisomers are presented.

Figure 2. Saturation binding of [^{3}H]-MFen to membranes obtained from HEK-β_2-AR cells, showing non-specific, specific and total binding. Non-specific binding was determined in the presence of 10 μM propranolol. Data shown is from a single experiment conducted in triplicate. This experiment was repeated two additional times with similar results.

Figure 3. Dissociation (A) and association (B) kinetics of [^{3}H]-MFen binding to membranes obtained from HEK-β_2-AR cells, where: solid lines represent a 2-site model and dashed lines represent best fit to a one-site binding model.

Figure 4. Effect of GTP and GTPγS on [^{3}H]-MFen binding. Binding was conducted to HEK-β2-AR cell membranes as described above in the presence of various concentrations of GTP, GTPγS, and propranolol as a standard. Values shown are average ± SD of three experiments conducted in triplicate.

Figure 5. Scatter plot of $-T\Delta S^\circ$ versus ΔH° values for the compounds used in the thermodynamic studies where the values denoted by **O** were previously determined using [^{3}H]-CGP-12177 as the marker ligand (Jozwiak et al., 2010b) and the values denoted by ■ were determined in this study using [^{3}H]-MFen as the marker ligand.

Figure 6. Molecular models of a) (*S*)-CGP-12177 interacting with an inactive model of β_2-AR (2RH1.pdb, β_2-AR-In) and b) (R,R')-MFen interacting with an active model of β_2-AR (3POG.pdb, β_2-AR-Ac) obtained during docking simulations. For clarity of both figures, TM1, TM2 and extracellular loop2 were hidden and remaining transmembrane

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segments are color coded, TM3 –red; TM4 – green; TM5 – magenta; TM6 – yellow and TM7 – blue. Only the residues forming hydrogen bonds (shown as green arrows) with a ligand molecule are shown explicitly. All aliphatic hydrogen atoms are hidden.

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	Receptor Binding					cAMP Stimulation
Compound	[^{3}H]-CGP-12177[a]		[^{3}H]-MFen		K_{iCGP}/ K_{iMFen}	
	K_i (nM)	**HS**	**K_i (nM)**	**HS**		**EC_{50}(nM)**
(R)-Isoproterenol	192 ± 24	0.85 ± 0.1	2.44 ± 0.28	0.78 ± 0.1	79	0.2
Propranolol	0.46 ± 0.06	1.24 ± 0.1	3.69 ± 1.36	1.88 ± 0.3	0.1	NA
ICI-118,551	0.60 ± 0.3	1.34 ± 0.4	2.52 ± 0.29	2.01 ± 0.2	0.2	NA
(R,R')-Fen	345 ± 33.8	0.92 ± 0.1	4.00 ± 0.75	0.76 ± 0.1	86	0.3
(R,S')-Fen	3,695 ± 245	0.81 ± 0.1	183 ± 30.0	0.97 ± 0.1	20	4.7
(S,R')-Fen	10,330 ± 1,405	1.02 ± 0.1	1,827 ± 117	0.92 ± 0.1	6	8.5
(S,S')-Fen	27,749 ± 6,816	ND	3,370 ± 210	1.34 ± 0.2	8	580.2
(R,R')-MFen	473 ± 35	0.98 ± 0.1	4.09 ± 0.55	0.80 ± 0.1	116	0.3
(R,S')-MFen	1,929 ± 135	1.01 ± 0.1	26.1 ± 2.44	1.00 ± 0.1	74	2
(S,R')-MFen	,268 ± 508	1.28 ± 0.1	91.3 ± 32.01	0.86 ± 0.1	58	7.2
(S,S')-MFen	5,880 ± 2,722	2.30 ± 0.3	2,870 ± 234	1.68 ± 0.5	6	33.2
(R,R')-PhFen	,864 ± 248	0.97 ± 0.02	26.6 ± 1.49	0.79 ± 0.1	70	ND
(R,R')-1-NapFen	41 ± 38	1.06 ± 0.2	3.66 ± 0.42	0.88 ± 0.2	66	12.5
(R,S')-1-NapFen	341 ± 32	0.93 ± 0.01	3.67 ± 1.32	0.86 ± 0.1	93	2.7
(S,R')-1-NapFen	1,783 ± 208	1.06 ± 0.1	57.6 ± 4.73	0.91 ± 0.04	31	66.7
(S,S')-1-NapFen	2,535 ± 295	1.12 ± 0.1	615 ± 85.2	0.99 ± 0.04	4	29.7
(R,R')-2-NapFen	404 ± 97	0.97 ± 0.04	4.52 ± 1.14	0.83 ± 0.1	89	0.4
(R,S')-2-NapFen	509 ± 5	1.06 ± 0.1	134 ± 10.7	1.04 ± 0.2	4	7.6
(R,R')-NH2Fen	2,933 ± 238	1.01 ± 0.1	42.8 ± 11.3	0.83 ± 0.05	69	2.42
(R,S')-NH2Fen	7,937 ± 561	1.07 ± 0.03	187 ± 42.6	0.91 ± 0.03	42	ND
(S,R')-NH2Fen	23,125 ± 2,093	ND	463 ± 103	1.00 ± 0.1	50	ND
(R,R')-EtFen	1,273 ± 81	1.01 ± 0.01	39.1 ± 5.38	0.93 ± 0.1	33	2.8
(R,S')-EtFen	5,758 ± 833	2.07 ± 0.4	294 ± 45.1	0.87 ± 0.2	20	16.6
(R,R')-MNF	277 ± 11	1.07 ± 0.09	13.3 ± 2.72	0.86 ± 0.1	21	3.9
(R,S')-MNF	317 ± 6	1.06 ± 0.02	12.7 ± 1.83	0.90 ± 0.2	25	4

Table 1. The β_2-adrenoreceptor (β_2-AR) binding affinities (K_i) determined using either [^{3}H]-4-methoxyfenoterol (K_{iMFen}) or [^{3}H]-CGP-12177 (K_{iCGP}) as the marker ligand and

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membranes obtained from HEK-β_2-AR cells and the induced cAMP accumulation in HEK-β_2-AR cells presented as EC_{50} values. The K_i values were determined at 25°C and are presented as ± SEM with $n \geq 3$, see text for experimental details. ND = Not Determined. For binding experiments this was because binding affinities were too low to obtain Hill coefficients. [b] data obtained from Jozwiak et al., 2010a; Jozwiak et al., 2010b; Toll et al., 2011)

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Table 2. Effect of GTP on agonist inhibition of [^{3}H]-CGP-12177 binding. Binding was conducted as described above in the presence and absence of 10 μM GTP.

Compound	-GTP	+GTP
propranolol	2.27 ± 0.78	2.03 ± 1.26
R,R'-fenoterol	560 ± 116	553 ± 151
R,R'-methoxyfenoterol	411 ± 229	220 ± 111

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Table 3. The influence of temperature on the binding to the β_2-AR of the antagonist propranolol and the agonists isoproterenol, (R,R')-Fen, (R,S')-Fen, (S,R')-Fen and (S,S')-Fen using [^{3}H](R,R')-MFen as the maker ligand where n = 3. All values are Ki except for (R,R')-MFen which are Kd values derived from saturation analysis. These Kd values were used to calculate Ki for the remaining compounds, as described in Materials and Methods. See text for experimental procedures.

	4°C	**25°C**	**37°C**
Compound	K_i (nM)	K_i (nM)	K_i (nM)
(R,R')-MFen	5.68 ± 1.35 nM	4.88 ± 0.41 nM	3.66 ± 0.82 nM
Propranolol	4.74 ± 1.94	3.69 ± 1.36	3.66 ± 0.66
Isoproterenol	4.40 ± 0.59	2.44 ± 0.28	1.89 ± 0.53
(R,R')-Fen	7.97 ± 3.79	4.00 ± 0.75	2.59 ± 0.20
(R,S')-Fen	187.40 ± 35	183 ± 30	83.00 ± 3.22
(S,R')-Fen	3,338 ± 764	1,827 ± 117	1,798.50 ± 198
(S,S')-Fen	3,800 ± 482	3,370 ± 210	1,639 ± 340

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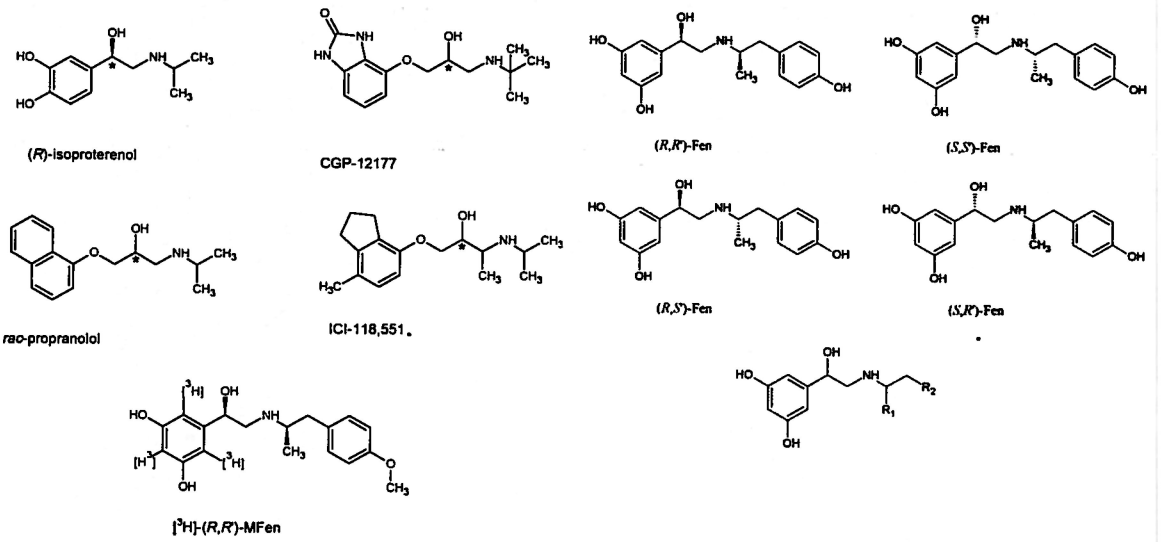
Table 4. Thermodynamic parameters of binding to the β_2-AR, using [^{3}H]-MFen as the marker ligand as compared to the previously reported parameters obtained using [^{3}H]-CGP-12177 as the marker ligand (Jozwiak et al., 2010a). *Distance* parameter is the Euclidean distance describing a shift of the (ΔH°; $-T\Delta S^\circ$) point, determined in the [^{3}H]-MFen experiment with respect to the [^{3}H]-CGP-12177 derived point, as illustrated in Fig. 4.

	$\Delta H^\circ_{\text{MFen}}$ **[kJ/mol]**	$\Delta H^\circ_{\text{CGP}}$ **[kJ/mol]**	$-T\Delta S^\circ_{\text{MFen}}$ **[kJ/mol]**	$-T\Delta S^\circ_{\text{CGP}}$ **[kJ/mol]**	$\Delta G^\circ_{\text{MFen}}$ **[kJ/mol]**	$\Delta G^\circ_{\text{CGP}}$ **[kJ/mol]**	***distance*** **[kJ/mol]**
Propranolol	+5.98 (±0.04)	–4.5	-54.32 (±0.04)	–48.95	-48.3 (±0.06)	–53.5	11.8
(*R*)-Isoproterenol	+18.4 (±0.13)	–18.03	-67.9 (±0.13)	–19.5	-49.5 (±0.2)	–37.6	60.6
(R,R')-Fen	+24.1 (±0.52)	0	-72.6 (±1.4)	–38.8	-48.5 (±1.5)	–38.8	41.5
(R,S')-Fen	+15.3 (±3.0)	+7.1	-54.9 (±1.2)	–40.0	-39.6 (±3.2)	–31.1	17.0
(S,R')-Fen	+18.4 (±0.13)	–23.0	-47.0 (±1.1)	–8.1	-32.7 (±1.6)	–31.1	53.9
(S,S')-Fen	+16.2 (±2.2)	–35.8	-48.5 (±2.2)	+6.3	-32.2 (±3.1)	–29.5	75.6

Figure 1



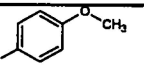
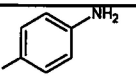
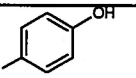
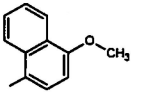
Compounds	R1	R2
MethoxyFen (MFen)	$-CH_3$	
PhenylFen (PhFen)	$-CH_3$	
1-naphtyl-Fen (1-NapFen)	$-CH_3$	
2-naphtyl-Fen (2-NapFen)	$-CH_3$	
Amino-Fen (NH2Fen)	$-CH_3$	
Ethyl-Fen (EtFen)	$-CH_2CH_3$	
4'-methoxy-1-naphtyl-Fen (MNFen)	$-CH_3$	

Figure 2.

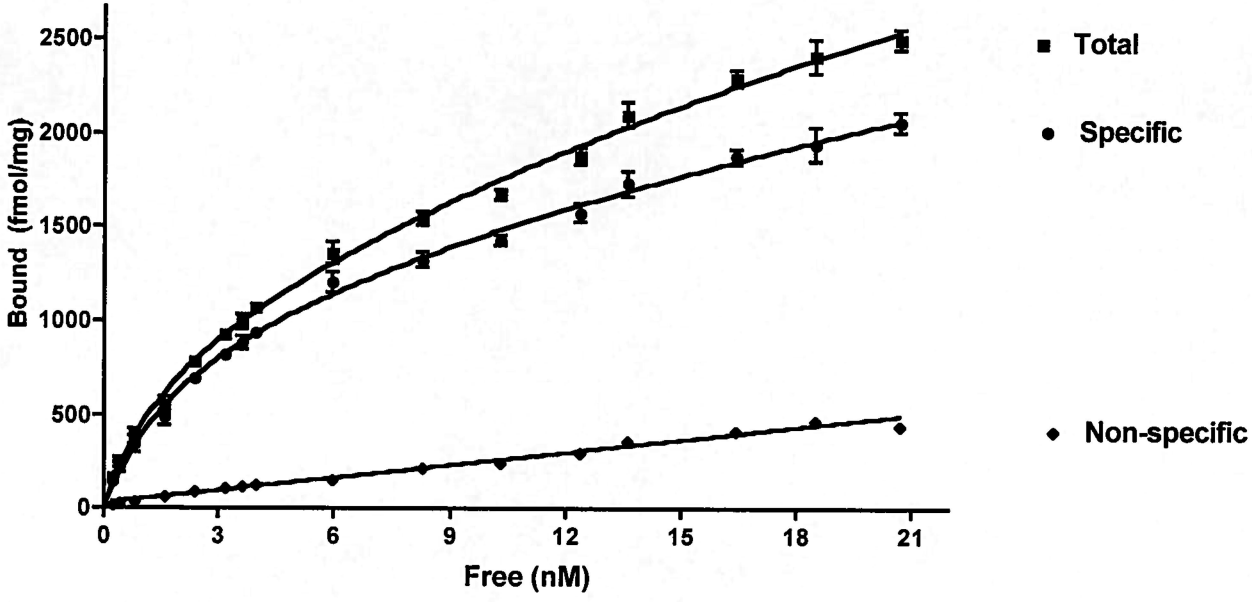
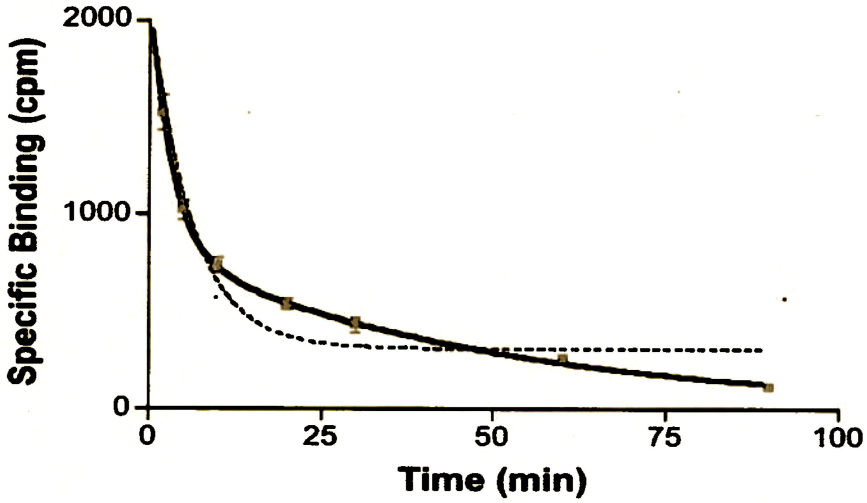
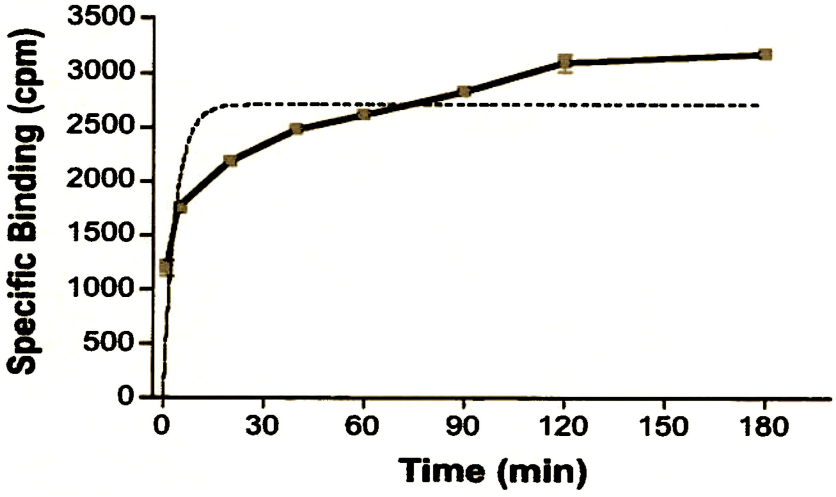

Bound (fmol/mg)
2500
2000
1500
1000
500
0
0
3
6
9
12
15
18
21
Free (nM)
Total
Specific
Non-specific

Figure 3


Specific Binding (cpm)
2000
1000
0
0
25
50
75
100
Time (min)

A


Specific Binding (cpm)
3500
3000
2500
2000
1500
1000
500
0
0
30
60
90
120
150
180
Time (min)

B

Figure 4

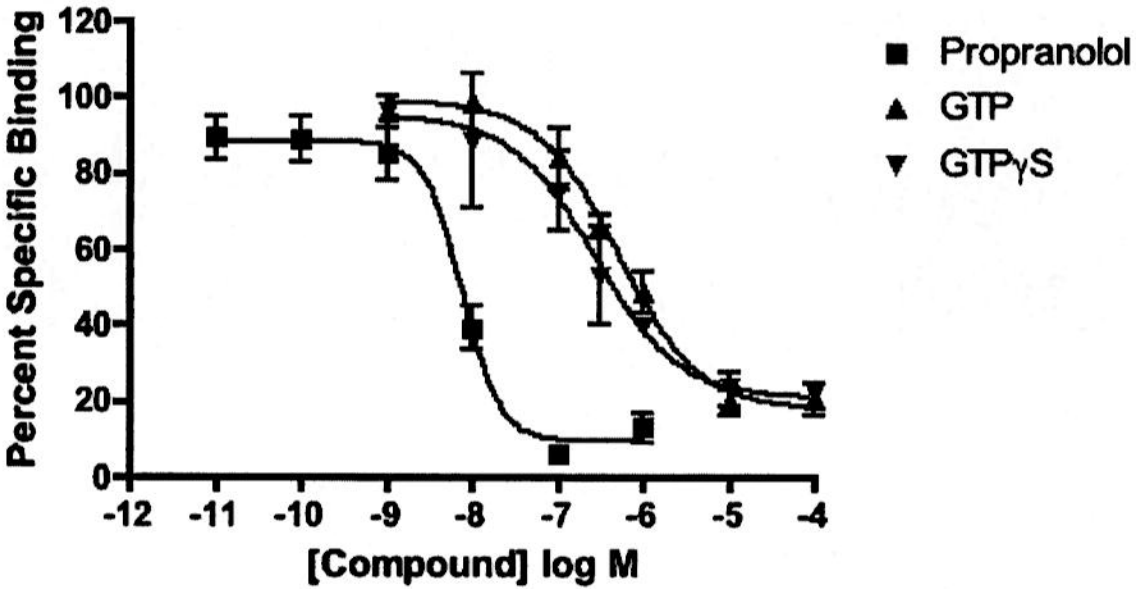

Percent Specific Binding
120
100
80
60
40
20
0
-12
-11
-10
-9
-8
-7
-6
-5
-4
[Compound] log M
Propranolol
GTP
GTPγS

Figure 5

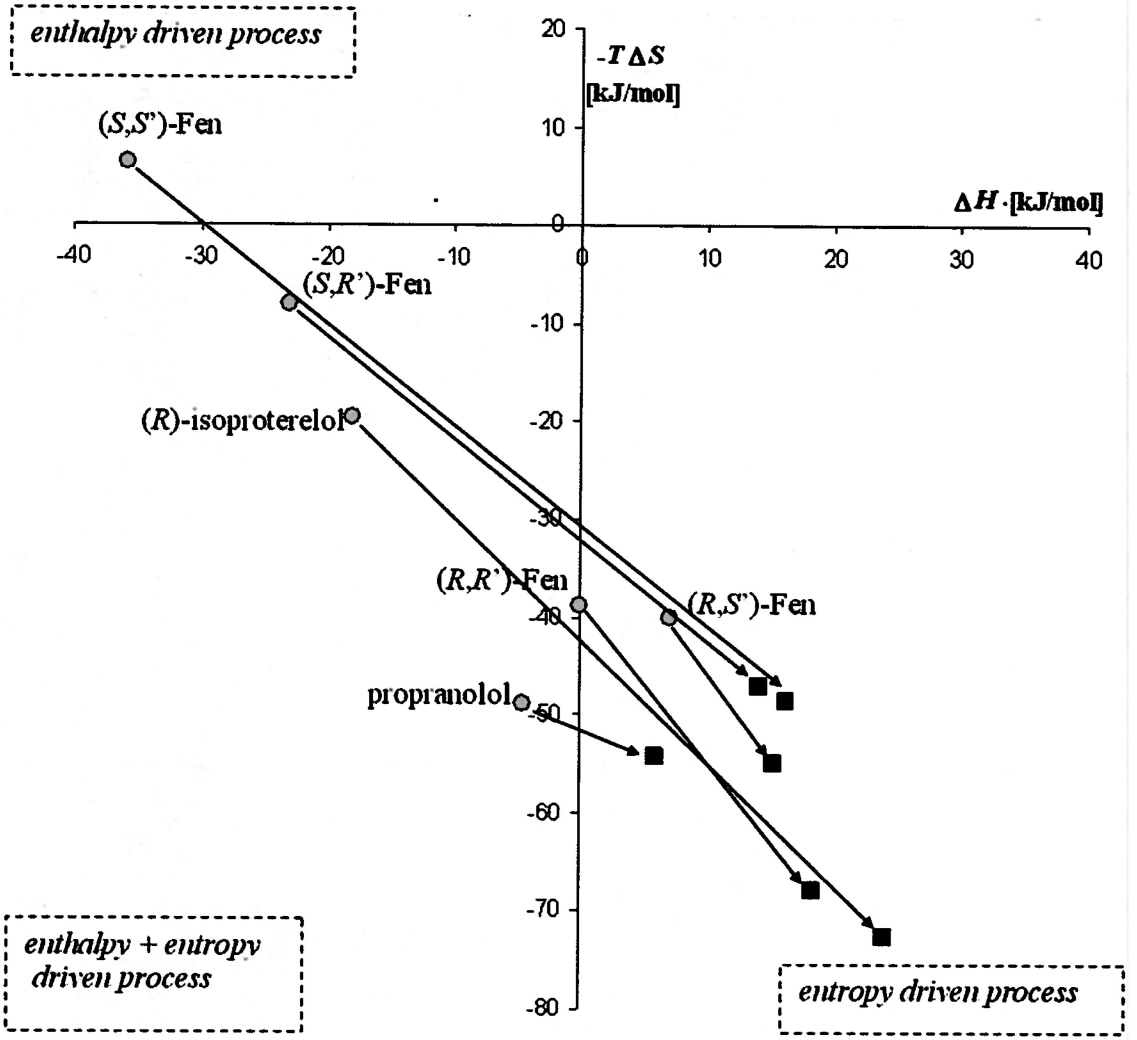
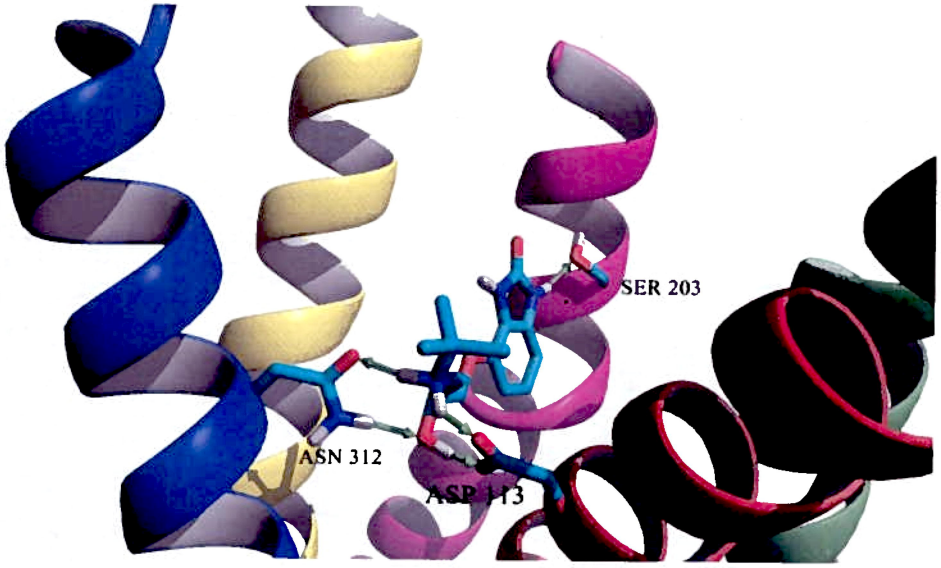
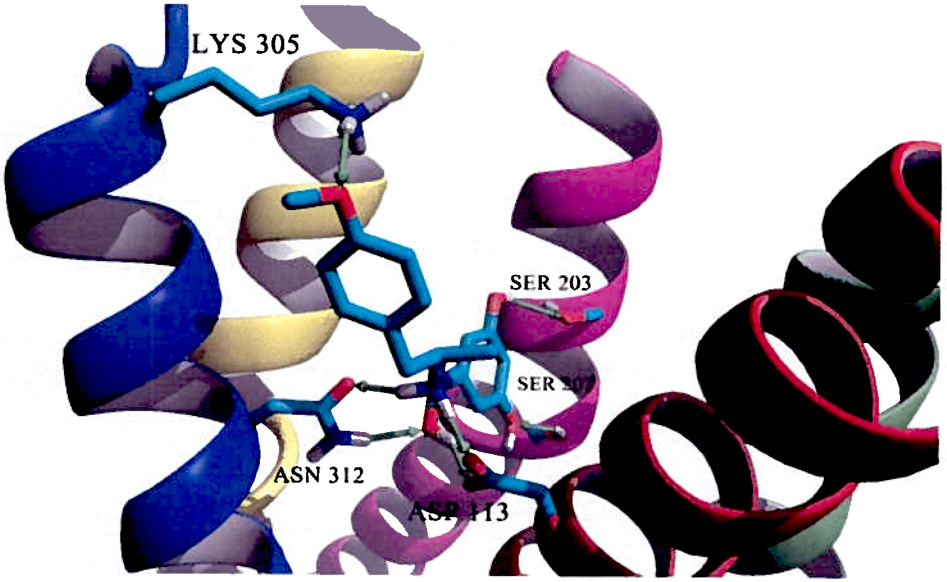


Figure 6



A



B

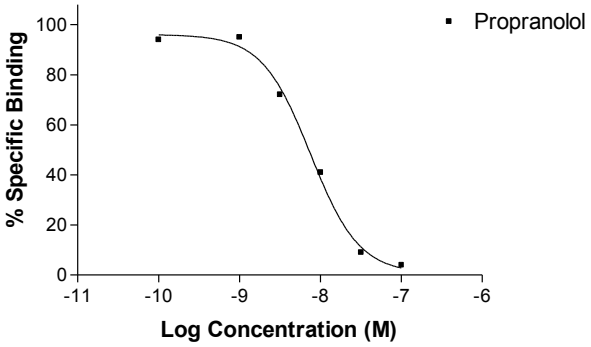
Molecular Pharmacology

Thermodynamics and Docking of Agonists to the β2-Adrenoceptor Determined Using [^{3}H]-(R,R')-4-Methoxyfenoterol as the Marker Ligand

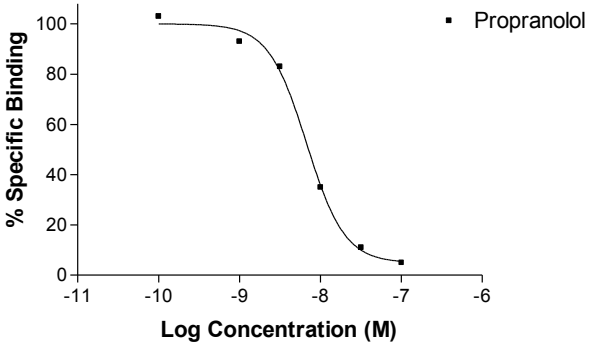
Lawrence Toll, Karolina Pajak, Anita Plazinska, Krzysztof Jozwiak, Lucita Jimenez, Joseph A. Kozocas, Mary J. Tanga, James E. Bupp, Irving W. Wainer,

Supplemental Figure S1. Inhibition curves generated using [^{3}H]M-Fen. Each experiment was conducted in triplicate. Data was analyzed using GraphPad Prism.

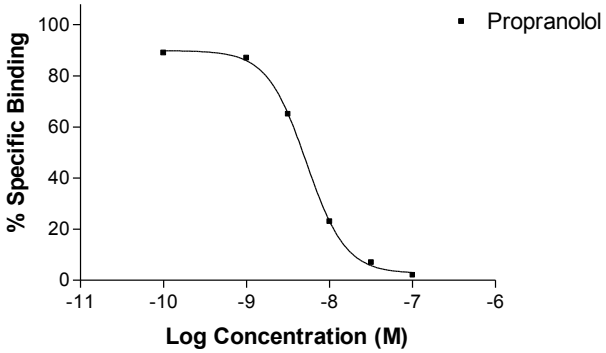
Supplemental Figure S1



HILLSLOPE	-1.450
EC50	7.5480e-009



HILLSLOPE	-1.850
EC50	6.7610e-009



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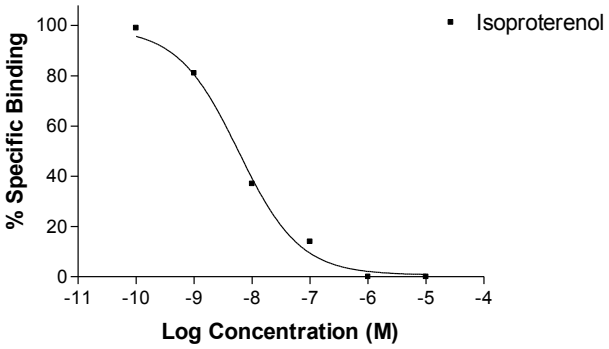
EC50	5.2590e-009



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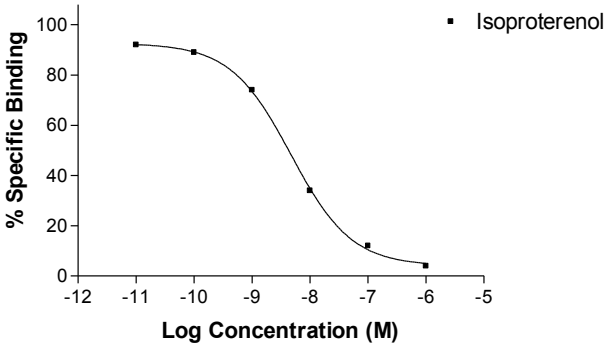
EC50	5.8640e-009



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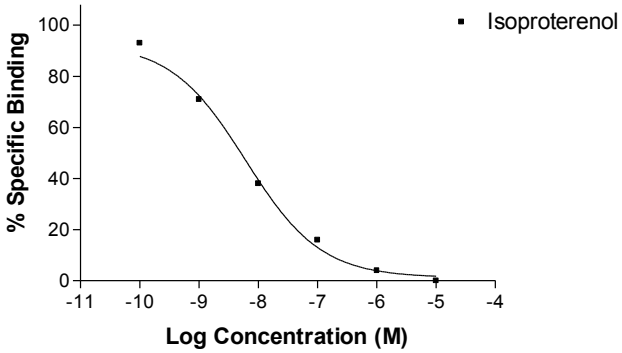
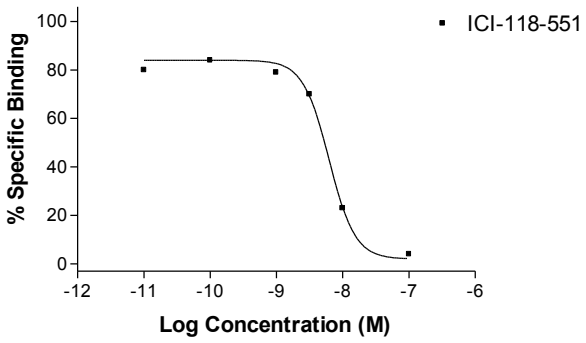
EC50	4.6850e-009



HILLSLOPE	-0.6851
EC50	6.1160e-009



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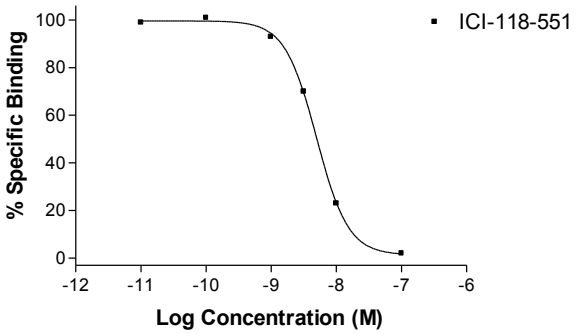
EC50	6.2460e-009



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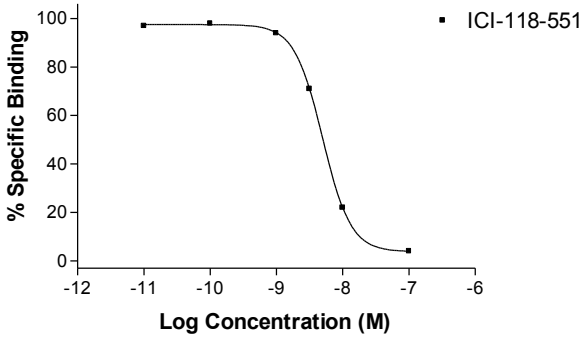
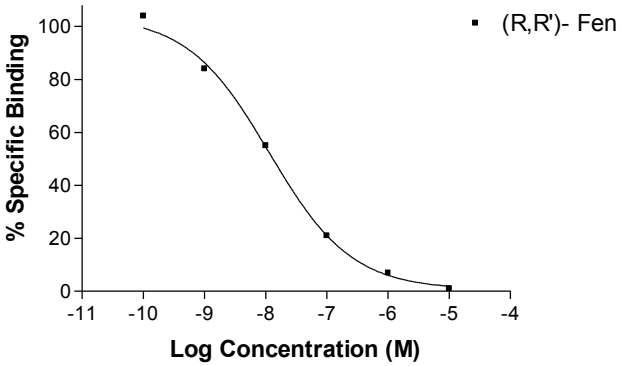
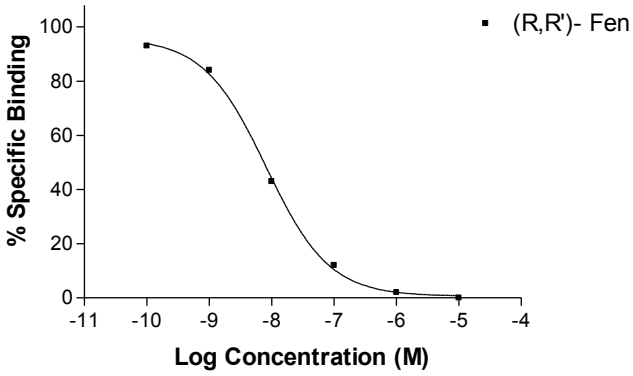
EC50	4.9850e-009



HILLSLOPE	-2.042
EC50	4.9870e-009



HILLSLOPE	-0.6476
EC50	1.1460e-008



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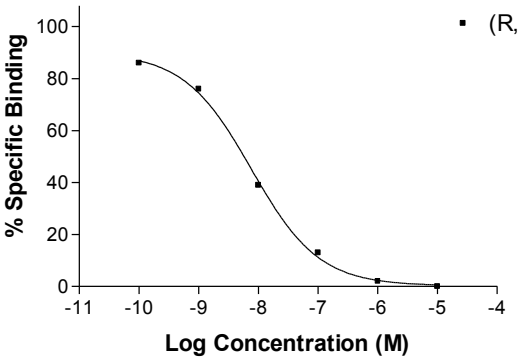
EC50	8.1880e-009



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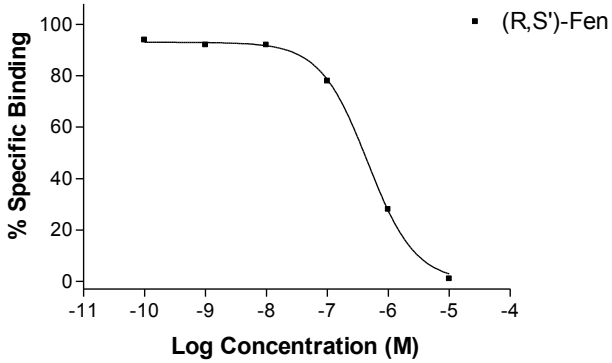
EC50	7.6590e-009



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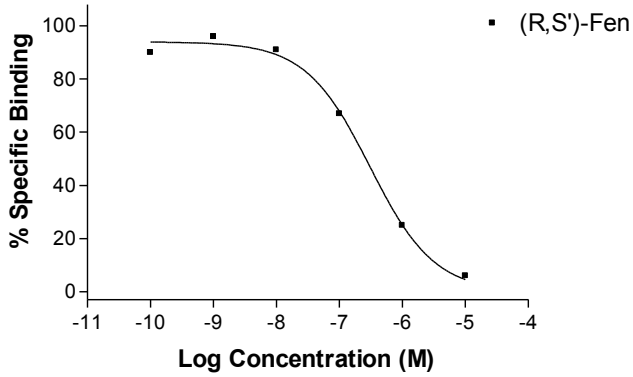
EC50	4.5580e-007



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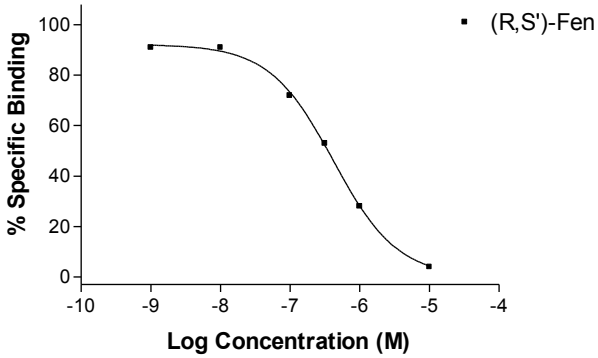
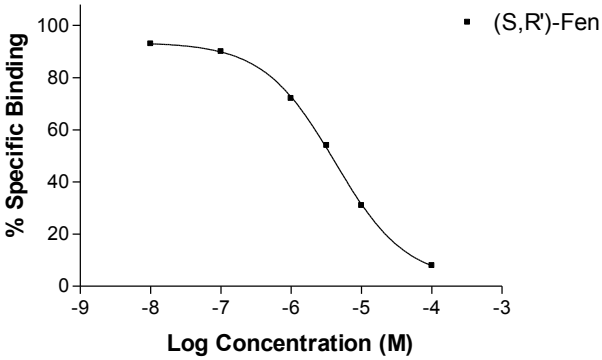
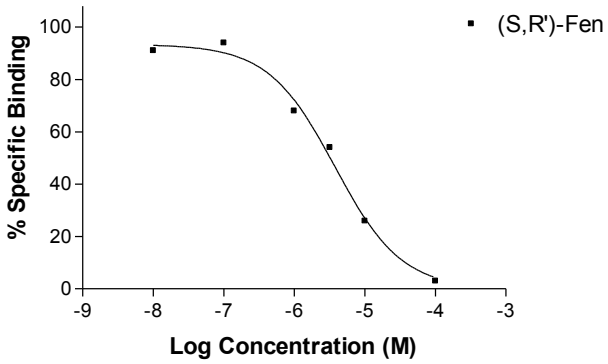
EC50	3.0490e-007



HILLSLOPE	-0.9427
EC50	4.1830e-007



HILLSLOPE	-0.8530
EC50	4.1570e-006



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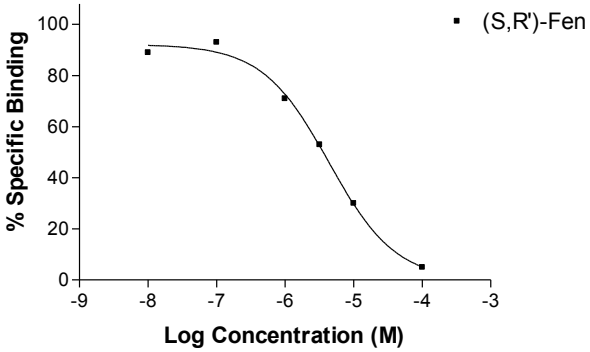
EC50	3.7650e-006



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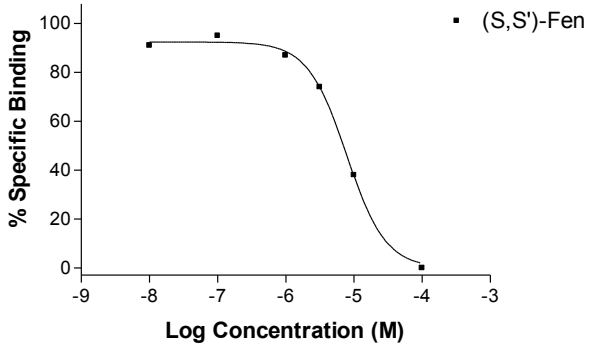
EC50	4.3620e-006



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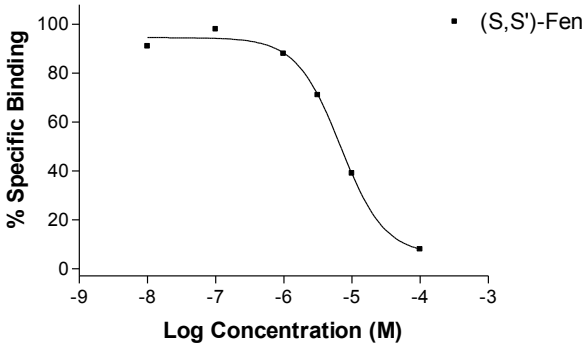
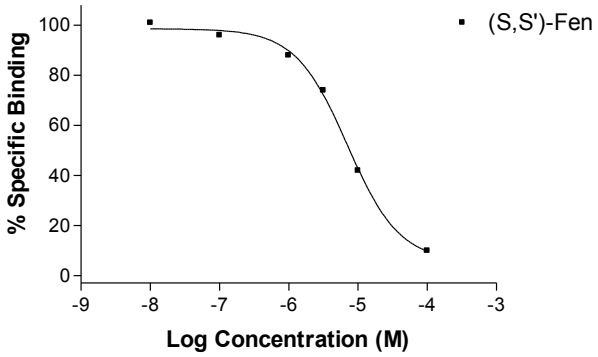
EC50	7.8290e-006



HILLSLOPE	-1.340
EC50	6.8060e-006



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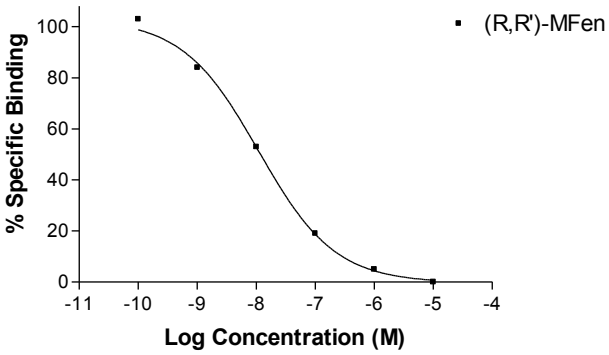
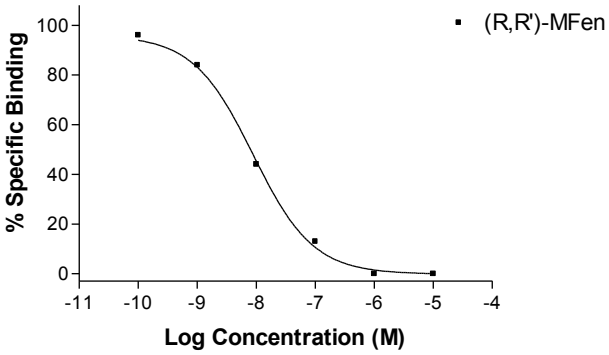
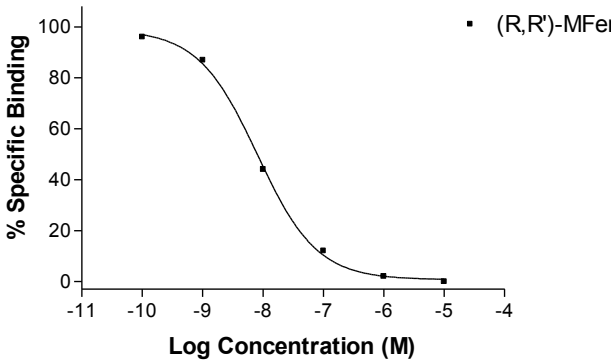
EC50	7.0590e-006



HILLSLOPE	-0.6736
EC50	1.0940e-008



HILLSLOPE	-0.8550
EC50	8.8640e-009



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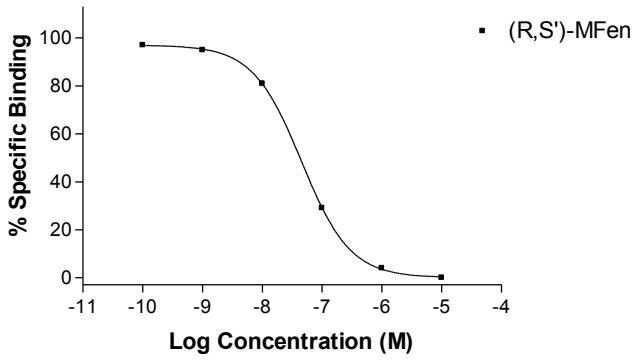
EC50	8.0820e-009



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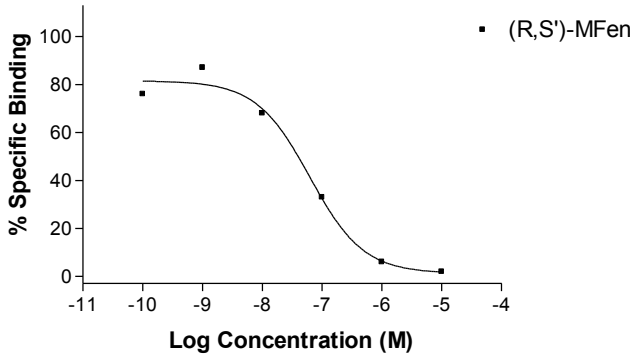
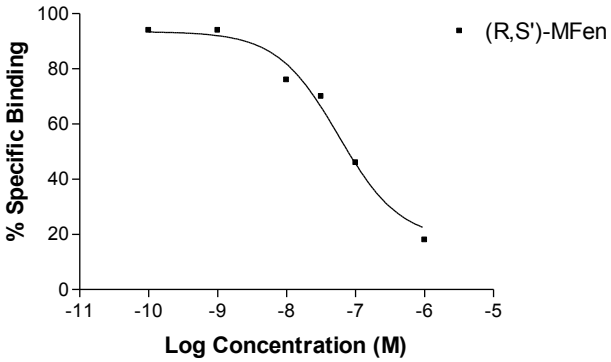
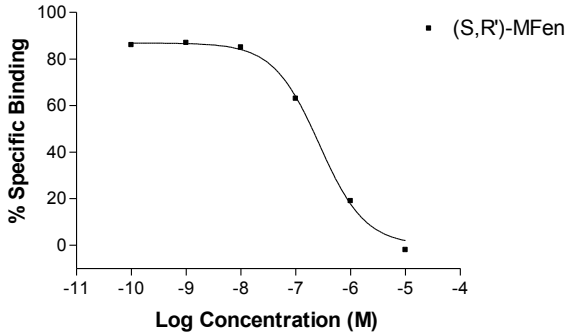
EC50	4.5270e-008



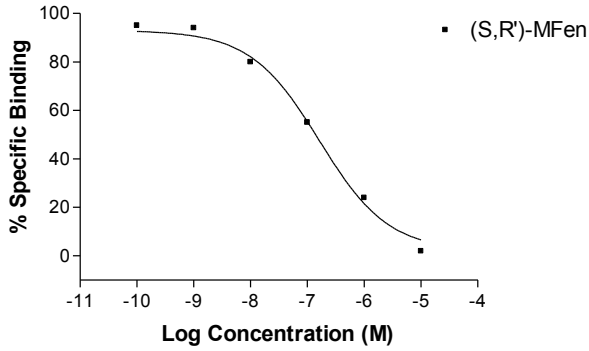
HILLSLOPE	-0.9678
EC50	6.2380e-008



HILLSLOPE	-0.9712
EC50	5.6790e-008



HILLSLOPE	-1.033
EC50	2.6700e-007



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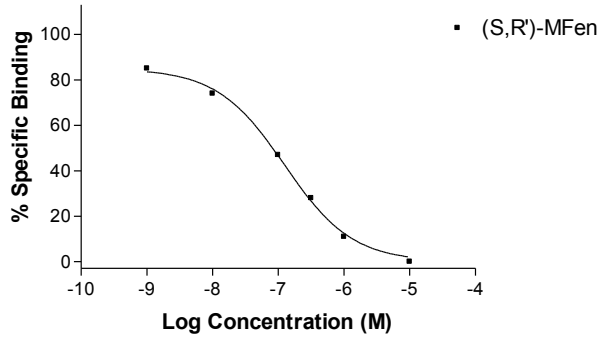
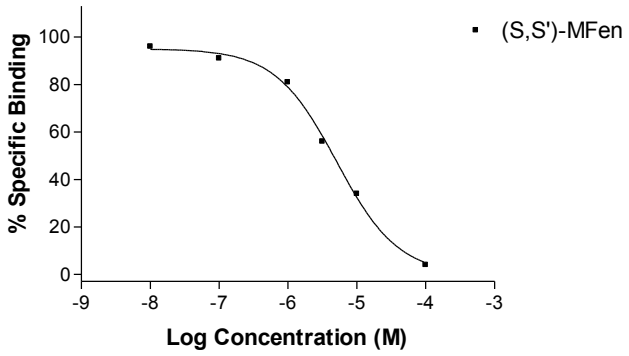
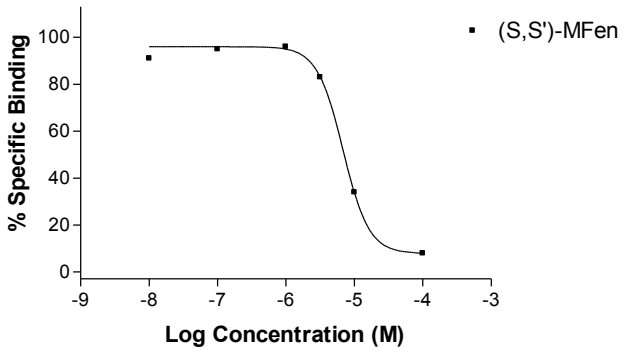
EC50	1.6260e-007



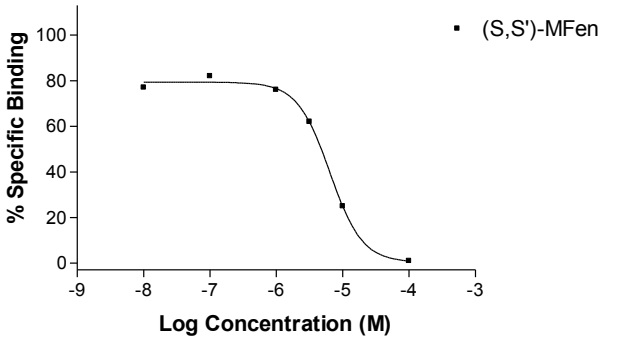
HILLSLOPE	-0.8449
EC50	1.2520e-007



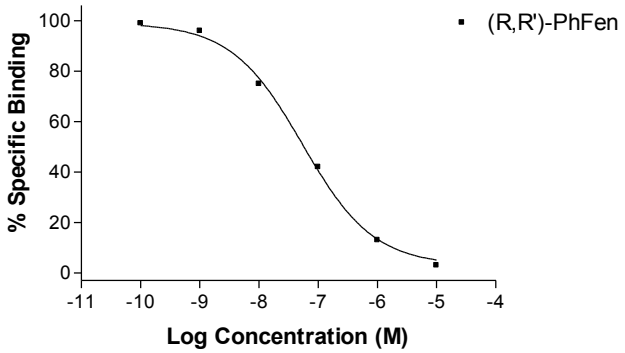
HILLSLOPE	-0.9715
EC50	5.0940e-006



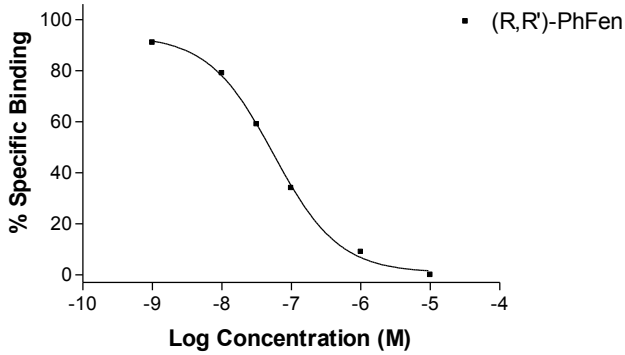
HILLSLOPE	-2.292
EC50	6.8530e-006



HILLSLOPE	-1.775
EC50	6.4160e-006



HILLSLOPE	-0.7225
EC50	5.4170e-008



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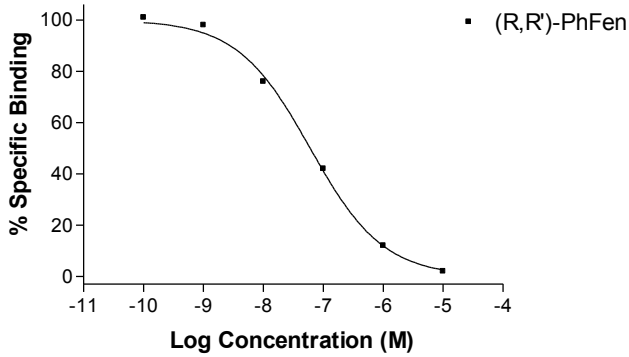
EC50	5.5530e-008



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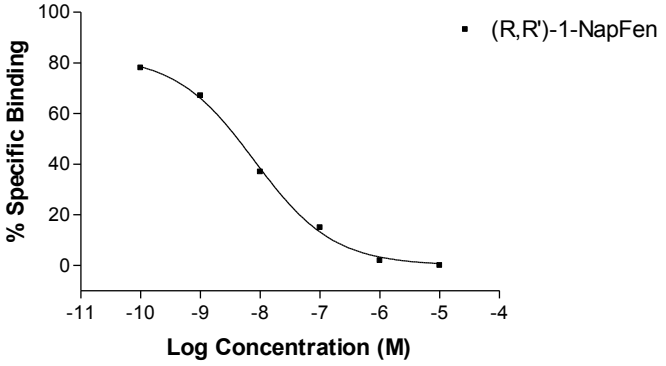
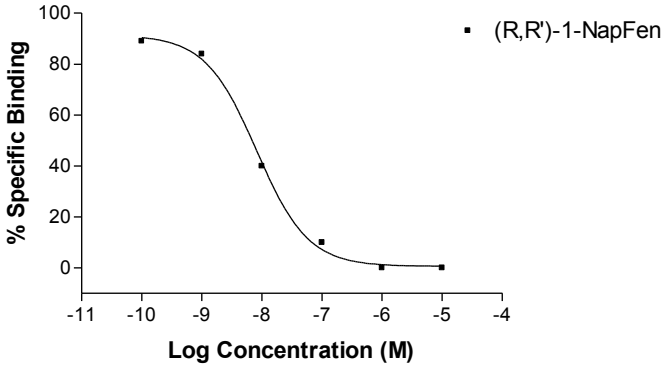
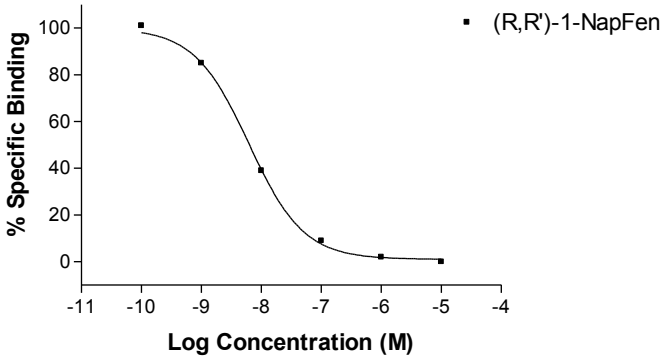
EC50	6.0350e-008



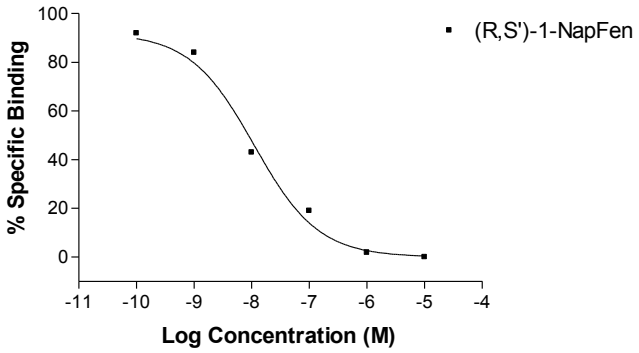
HILLSLOPE	-0.6570
EC50	7.9430e-009



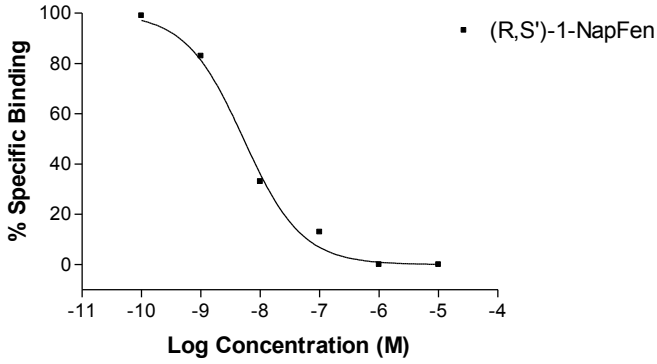
HILLSLOPE	-1.018
EC50	8.1940e-009



HILLSLOPE	-0.9521
EC50	6.2030e-009



HILLSLOPE	-0.7800
EC50	1.1010e-008



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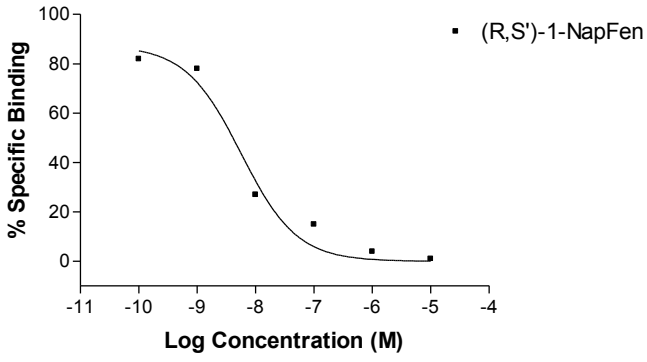
EC50	5.2360e-009



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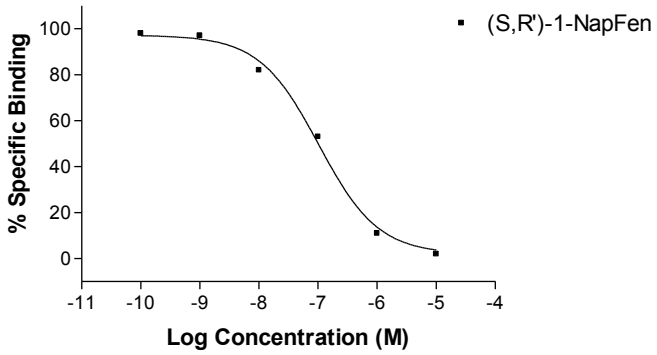
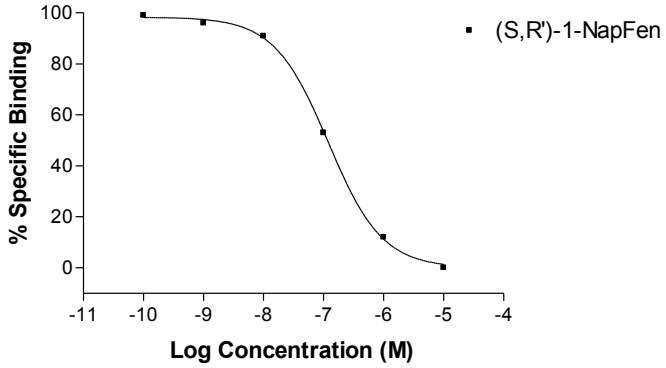
EC50	5.6540e-009



HILLSLOPE	-0.8637
EC50	1.0270e-007



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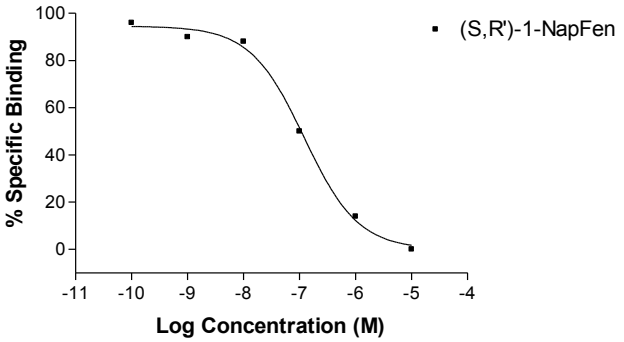
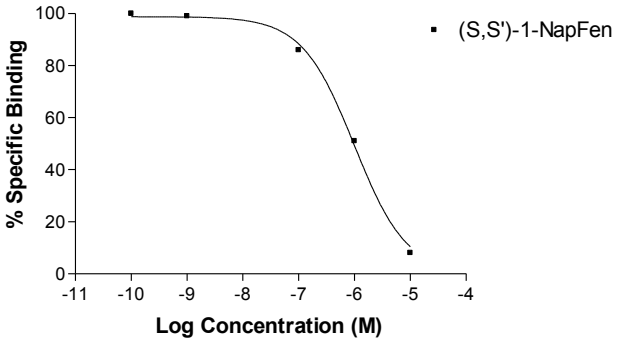
EC50	1.1980e-007



HILLSLOPE	-0.9022
EC50	1.2120e-007



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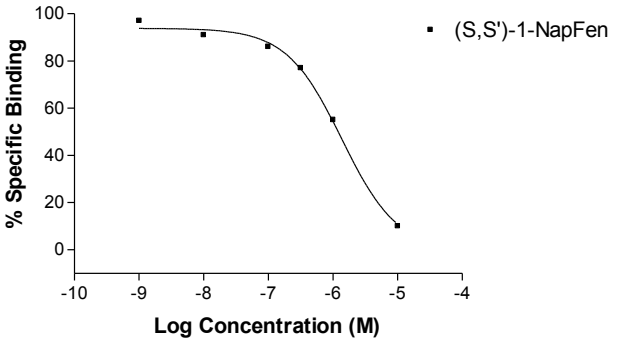
EC50	9.9120e-007



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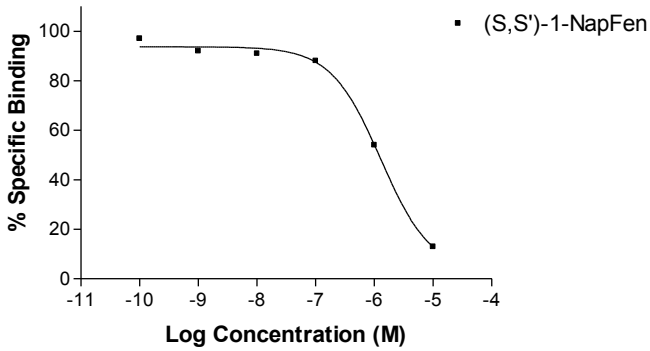
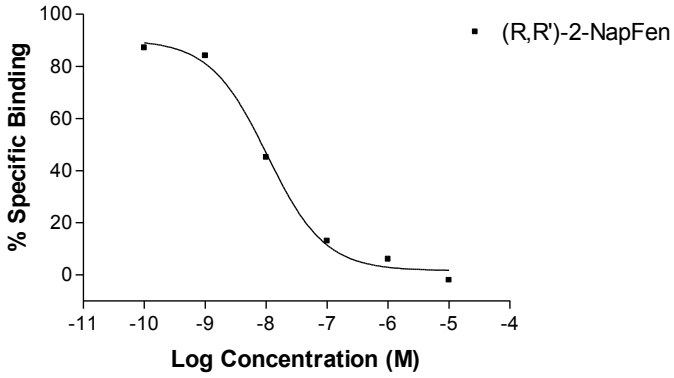
EC50	1.3690e-006



HILLSLOPE	-1.023
EC50	1.2780e-006



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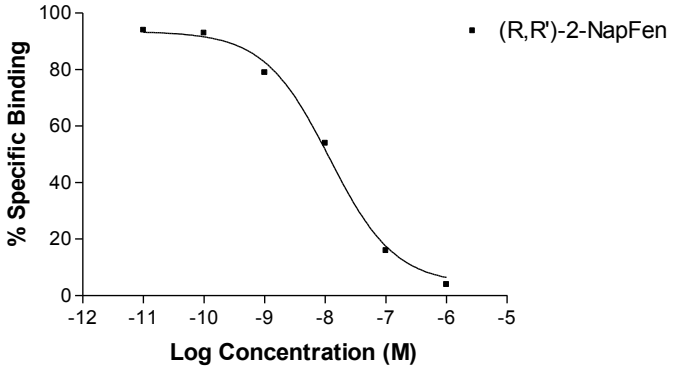
EC50	1.0410e-008



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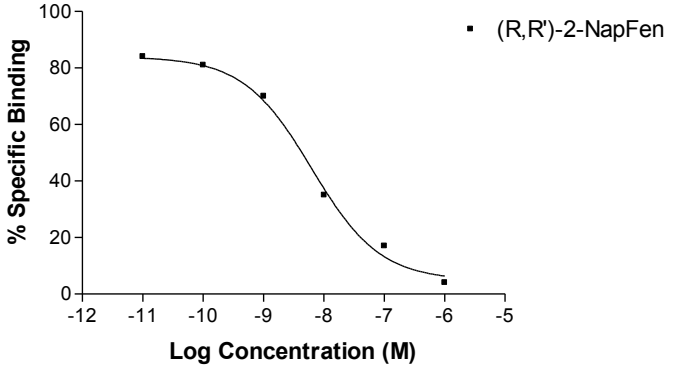
EC50	1.1680e-008



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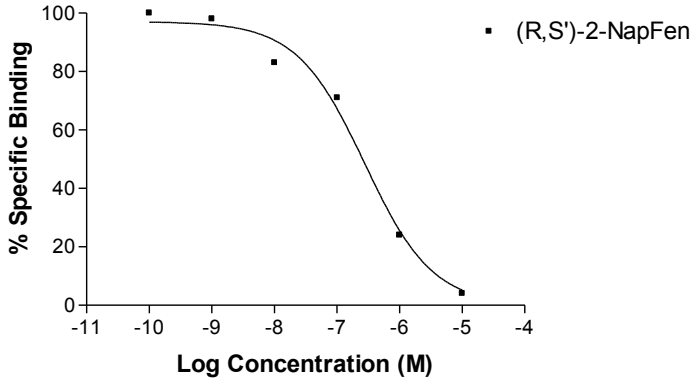
EC50	6.2600e-009



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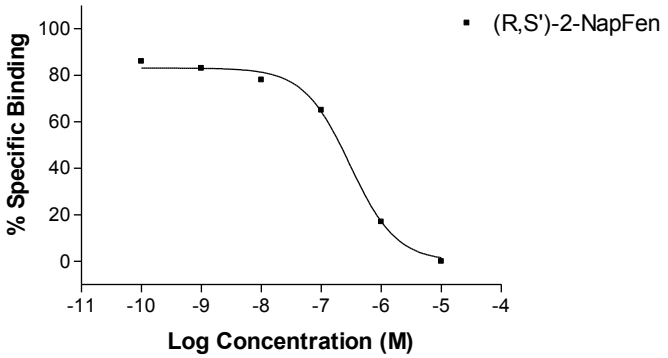
EC50	2.7610e-007



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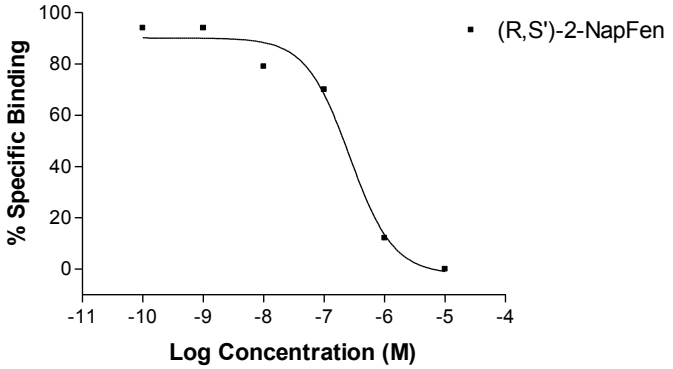
EC50	2.9950e-007



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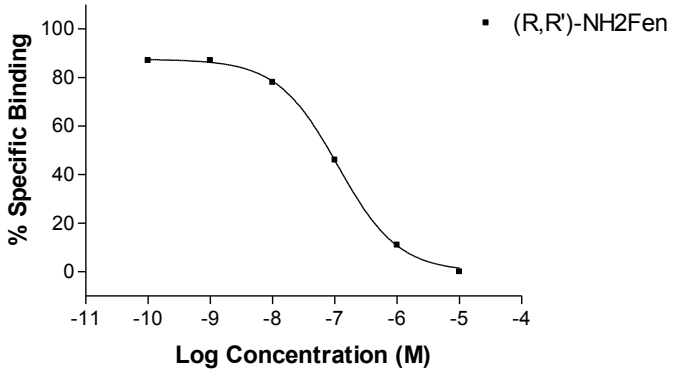
EC50	2.6190e-007



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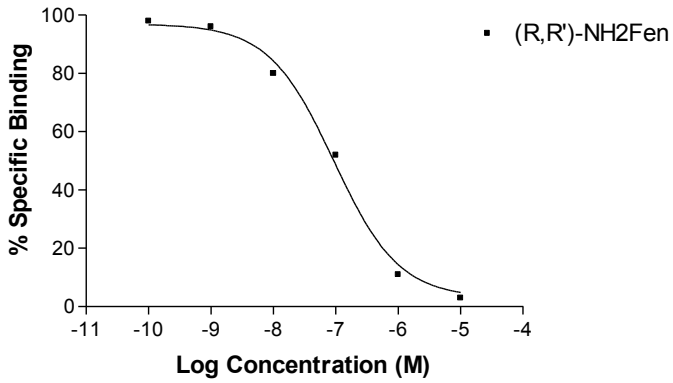
EC50	1.1130e-007



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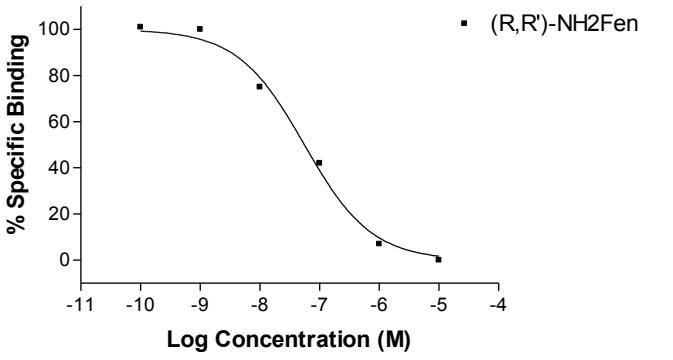
EC50	9.2810e-008



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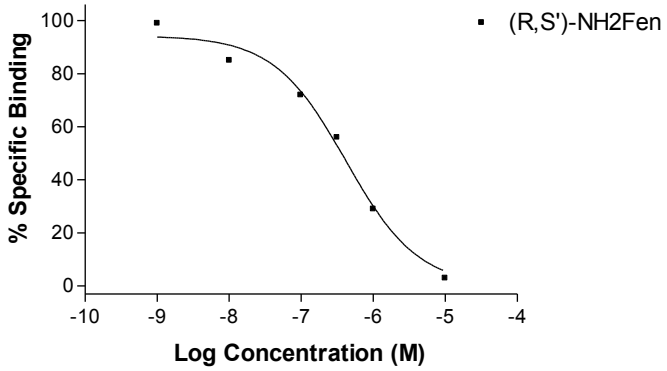
EC50	5.5660e-008



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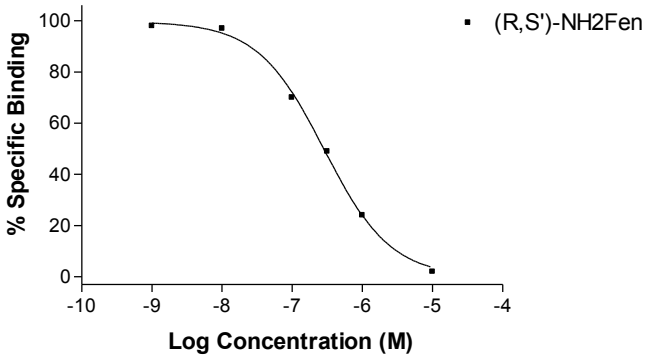
EC50	4.1840e-007



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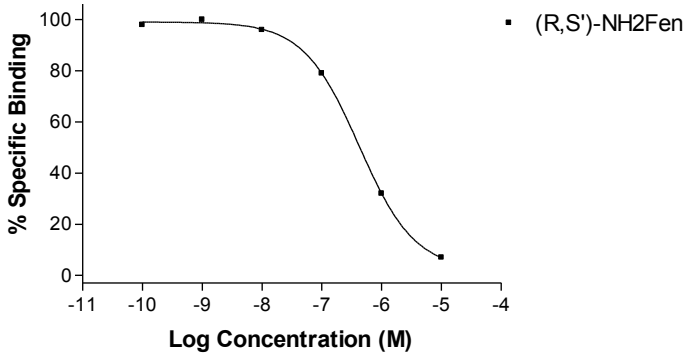
EC50	2.8530e-007



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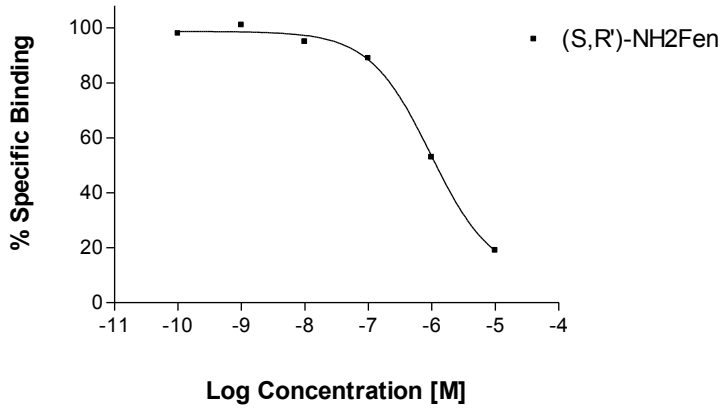
EC50	4.1890e-007



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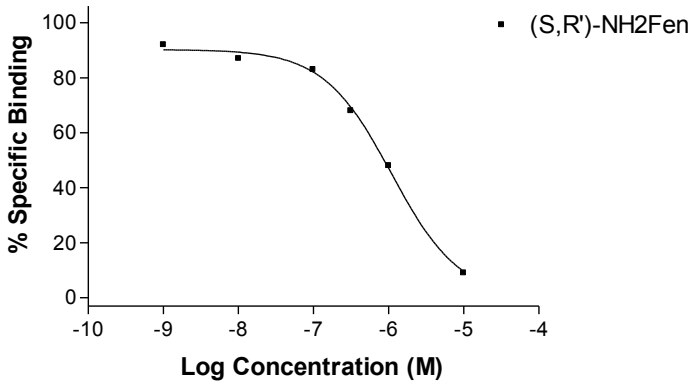
EC50	9.5560e-007



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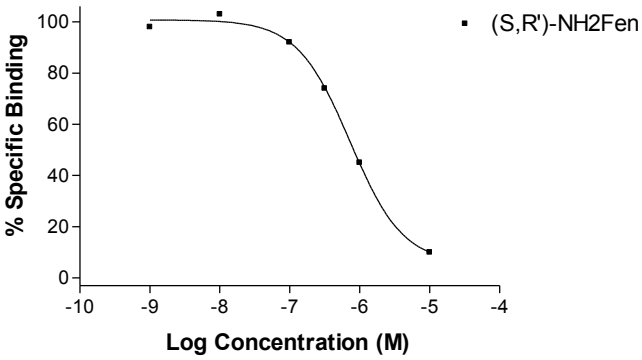
EC50	1.0960e-006



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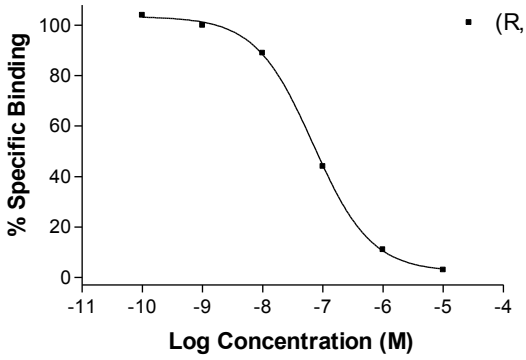
EC50	7.3140e-007



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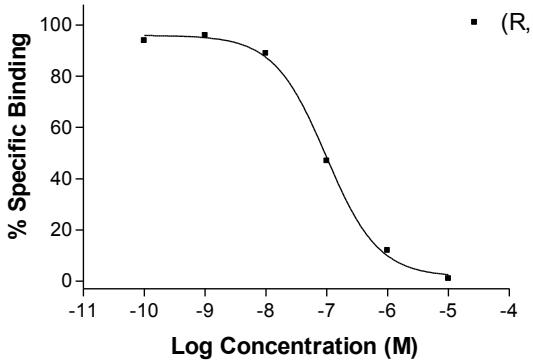
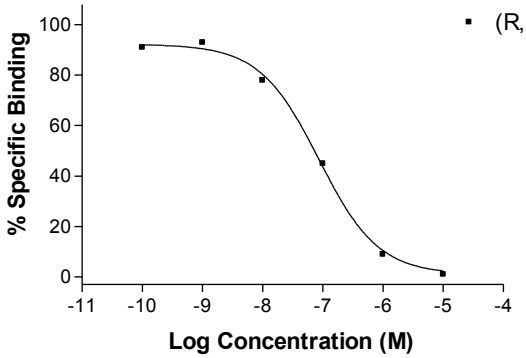
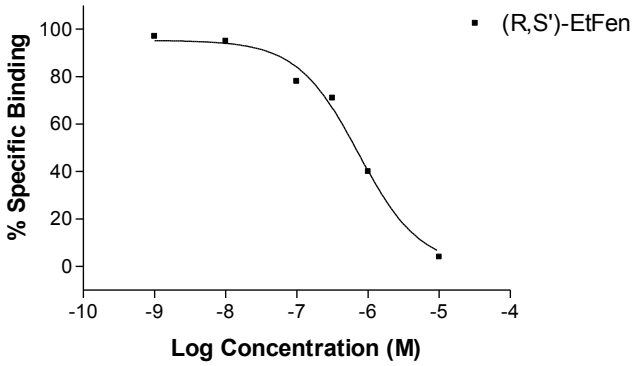
EC50	6.8850e-008



HILLSLOPE	-1.011
EC50	9.7250e-008



HILLSLOPE	-0.8800
EC50	8.5690e-008



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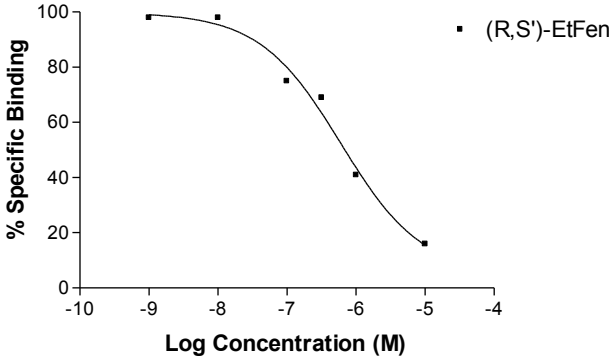

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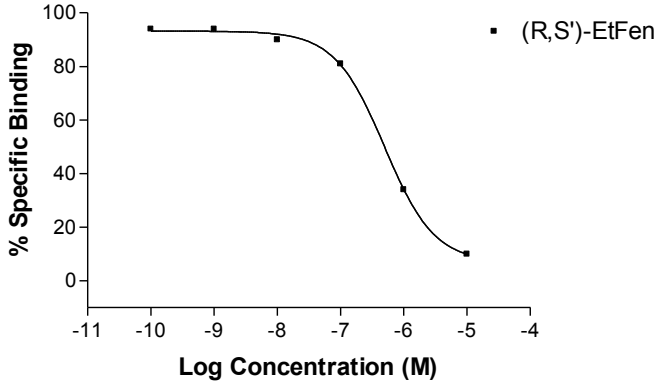

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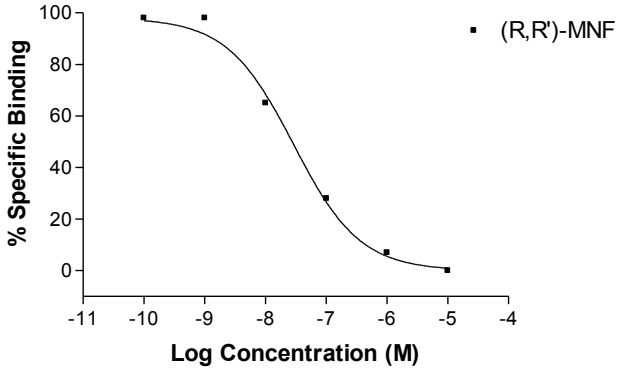
EC50	4.9530e-007



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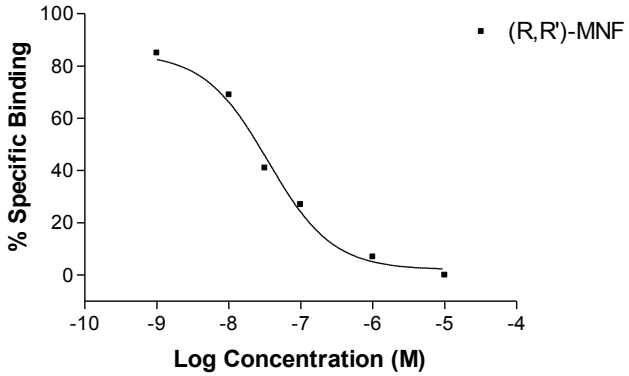
EC50	2.8890e-008



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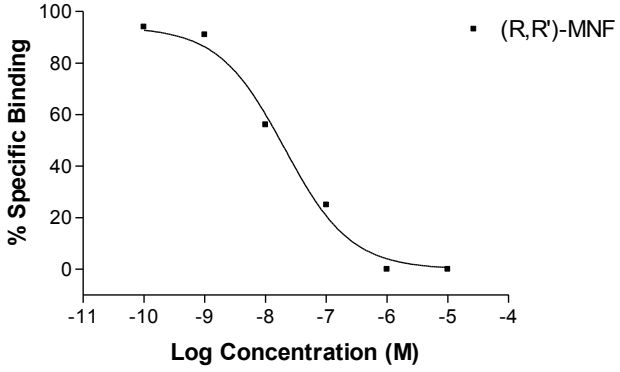
EC50	3.5300e-008



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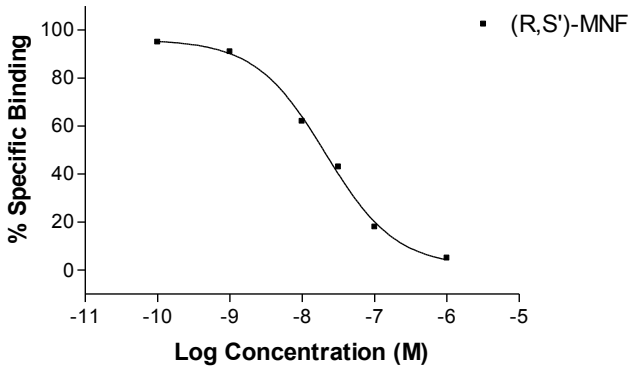

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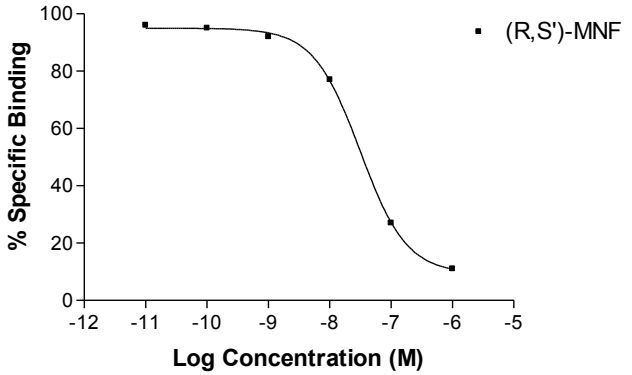
EC50	2.0910e-008



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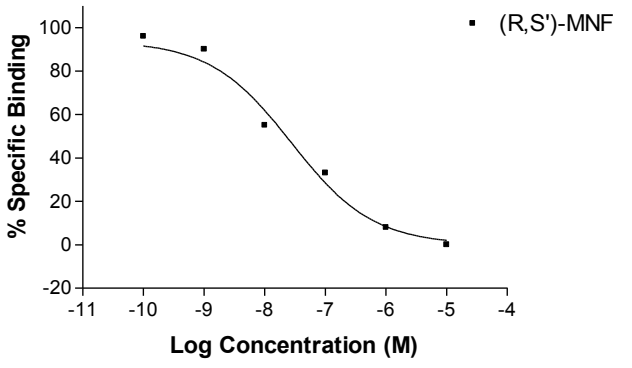
EC50	3.1190e-008



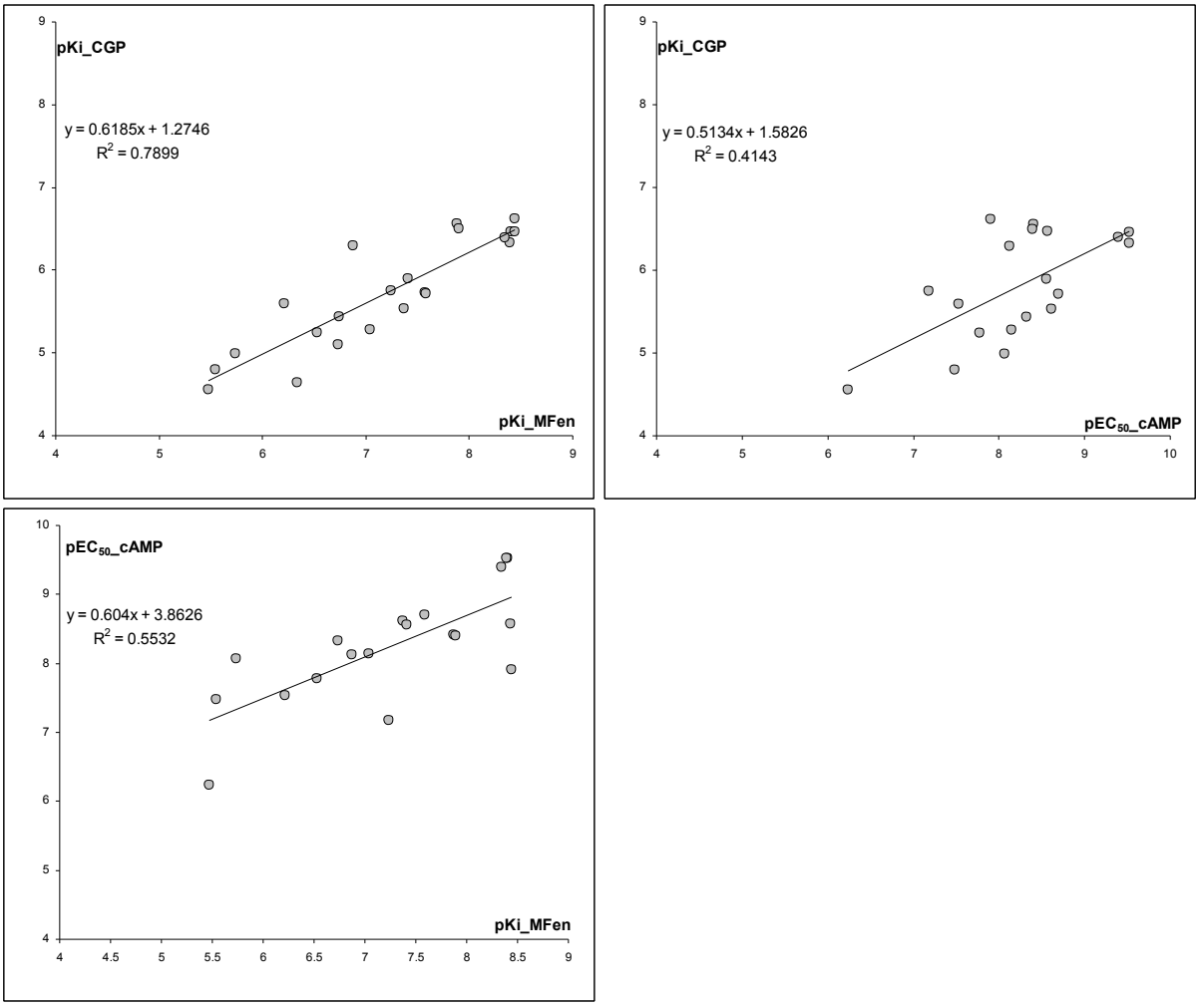
HILLSLOPE	-0.6471
EC50	2.7530e-008

Molecular Pharmacology

Thermodynamics and Docking of Agonists to the β2-Adrenoceptor Determined Using [^{3}H]-(R,R')-4-Methoxyfenoterol as the Marker Ligand

Lawrence Toll, Karolina Pajak, Anita Plazinska, Krzysztof Jozwiak, Lucita Jimenez, Joseph A. Kozocas, Mary J. Tanga, James E. Bupp, Irving W. Wainer,

Supplemental Figure S2. Figure of correlation matrix for pKi values determined with [^{3}H]-CGP-12177 (pKi CGP), [^{3}H]-MFen (pKi MFen) and pEC_{50} values in cAMP accumulation experiments (pEC_{50} cAMP).



Matrix of correlation coefficients.

R^2 values	pKi_MFen	pKi_CGP	pEC_{50}_cAMP
pKi_MFen	1	0.7899	0.5532
pKi_CGP		1	0.4143
pEC_{50}_cAMP			1