

PRODUCT DATASHEET

ChemiScreen™ NTR1 Neurotensin Receptor Membrane Preparation

CATALOG NUMBER: HTS034M QUANTITY: 200 units

LOT NUMBER: SC20200109 VOLUME/CONCENTRATION: 1 mL, 1 mg/mL

BACKGROUND: Neurotensin is a 13 amino acid peptide that induces hypothermia and mediates analgesia when

administered centrally, and also mediates the behavioral effects of antipsychotic drugs in rodents (Kinkead and Nemeroff, 2004). In addition, several intestinal functions, such as contractility and inflammation, are mediated by neurotensin (Castagliuolo et al., 1999). The actions of neurotensin are mediated by two GPCRs, NTR1 (or NTS1) and NTR2 (or NTS2). NTR1 knockout mice are resistant to the hypothermic, analgesic and contractile effects of neurotensin, indicating that NTR1 is the primary mediator of these effects (Pettibone et al., 2002). Small molecule antagonists, including SR48692 and SR142948A, that are partially selective for NTR1 also interfere with the biological effects of neurotensin (Labbé-Jullié et al., 1998). NTR1 membrane preparations are crude membrane preparations made from our proprietary stable recombinant cell lines to ensure high-level of GPCR surface expression; thus, they are ideal HTS tools for screening of antagonists of neurotensin/NTR1 interactions. The membrane preparations

exhibit a Kd of 0.6 nM for [125I]-neurotensin.

APPLICATIONS: Radioligand binding assay

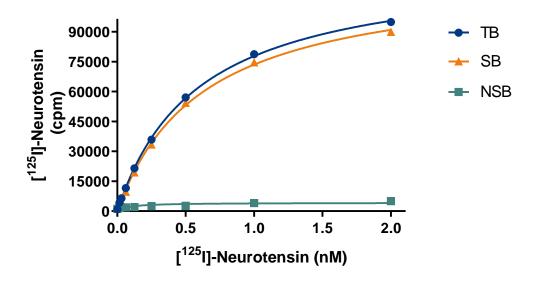


Figure 1. Saturation binding for NTR1. 5μ g/well NTR1 Membrane Preparation was incubated with increasing amount of 125 I-labeled neurotensin in the absence (total binding, TB) or presence (nonspecific binding, NSB) of 200-fold excess unlabeled neurotensin. Specific binding (SB) was determined by subtracting NSB from TB. Sample data from representative.



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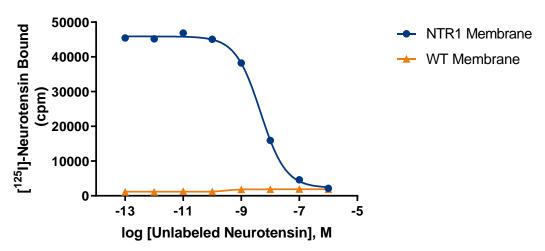


Figure 2. Competition binding for NTR1. 5 μ g/well NTR1 Membrane Preparation (HTS034M) and Wild-Type Chem-1 Membrane Preparation (WT; Cat. # HTS000MC1) were incubated with 0.5 nM 125 I-labeled neurotensin and increasing concentrations of unlabeled neurotensin. More than 5- fold signal:background was obtained with NTR1 Membrane Preparation, whereas no signal was observed in wild-type Membrane Preparation. Representative sample data.

SPECIFICATIONS: 1 unit = 5 µg

B_{max}: 9.77 pmol/mg protein

K_d: 0.6 nM

Signal:background: ≥5-fold

TRANSFECTION: Full length NTSR1 cDNA encoding human NTR1 (Accession number

NM_002531)

Species: Human

HOST CELLS: Chem-1, an adherent mammalian cell line without any endogenous NTR1

expression.

RECOMMENDED ASSAY CONDITIONS: Membranes are mixed with radioactive ligand and unlabeled competitor (see Figures 1 and 2 for concentrations tested) in binding buffer in a nonbinding 96-well plate, and incubated for 1-2 h. Prior to filtration, a GF/C 96-well filter plate is coated with 0.33% polyethyleneimine for 30 min, then washed with 50mM HEPES, pH 7.4, 0.5% BSA. Binding reaction is transferred to the filter plate, and washed 3 times (1 mL per well per wash) with Wash Buffer. The plate is dried and counted.

Binding buffer: 50mM HEPES, 5mM MgCl2, 1mM CaCl2, 0.2% BSA, pH to 7.4 and filter.

Radioligand: [125] Neurotensin (PerkinElmer#: NEX198)

Wash Buffer: 50 mM Hepes, pH 7.4, 500mM NaCl, 0.1% BSA, filtered and stored at 4°C.

One package contains enough membranes for at least 200 assays (units), where a unit is the amount of membrane that will yield greater than 5-fold signal:background with ¹²⁵I-labeled neurotensin at 0.5 nM.

PRESENTATION:

Liquid in packaging buffer: 50 mM Tris pH 7.4, 10% glycerol and 1% BSA with no preservatives.

Packaging method: Membrane proteins were adjusted to the indicated concentration in 1 ml



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packaging buffer, rapidly frozen, and stored at -80°C.

STORAGE/HANDLING:

Store at -70°C. Product is stable for at least 6 months from the date of receipt when stored as directed. Do not freeze and thaw.

REFERENCES:

- 1. Castagliuolo I et al. (1999) Neurotensin is a proinflammatory neuropeptide in colonic inflammation. J. Clin. Invest. 103: 843-849.
- 2. Kinkead B. and Nemeroff C. (2004) Neurotensin, schizophrenia, and antipsychotic drug action. Int. Rev. Neurobiol. 59: 327-349.
- Labbé-Jullié C et al. (1998) Mutagenesis and modeling of the neurotensin receptor NTR1: Identification of residues that are critical for binding SR48692, a nonpeptide neurotensin antagonist. J. Biol. Chem. 273: 16351-16357.
- Pettibone D.J et al. (2002) The effects of deleting the mouse neurotensin receptor NTR1 on central and peripheral responses to neurotensin. J. Pharmacol. Exp. Ther. 300: 305-313.

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