

Research use only. Not for use in diagnostic procedures.

Membrane Target Systems™

# human Melanocortin MC<sub>5</sub> Receptor

Product No.: RBXMC5M400UA

Lot No.: 2382492

#### Material Provided

Membranes: 1 x 400 units / 400 µL frozen aliquot

#### **Product Information**

Cellular Background: HEK293-EBNA

GenBank Accession Number: NM\_005913

Unit Size: 0.8 µg protein / unit

Storage Buffer: 50 mM Tris-HCL (pH 7.4), 0.5mM EDTA, 10mM MgCl<sub>2</sub>, 10% sucrose.

Storage Conditions: Store at -80°C. Freeze-thaw is not recommended as it can affect

product performance and homogeneity. In order to minimize negative impact of freeze-thawing, flash freeze in liquid nitrogen for

30 seconds prior to transferring to -80°C.

Stability: This product is stable for at least 3 years from reception if used and

stored under recommended conditions.

### **Quality Control**

 $B_{\text{max}}$  and  $K_{\text{d}}$  are determined using radioactive saturation binding assays (Figure 1). Protein concentration is determined using the BCA method <sup>(1)</sup>. Ratio-to-Reference (RTR) is determined by dividing the maximal signal of the current lot ( $B_{\text{max}}$  in fmoles) by the maximal signal of a pre-defined reference tested in parallel. RTR is an indicator of lot-to-lot consistency. \*We certify that these results meet our quality release criteria.

Ratio-to-Reference (RTR): 1.14

Expression Level (B<sub>max</sub>): 5.8 pmol/mg membrane protein.

Kd for  $\lceil ^{125}I \rceil$ -(Nle<sup>4</sup>, D-Phe<sup>7</sup>)- $\alpha$ -MSH: 1.09 nM

Protein Concentration: 0.8 μg/μL

(1) Smith, P.K., et al. (1985). Anal. Biochem. 150, 76-85.



## **Recommended Assay Conditions**

Assay Buffer: 25 mM Hepes pH 7.0, 1.5 mM CaCl<sub>2</sub>, 1 mM MgSO<sub>4</sub>, 100 mM NaCl, 0.2% BSA,

1 mM 1,10-phenanthroline, 1 Complete™ protease inhibitor tablet (EDTA

free)/100 mL

Wash Buffer: 25 mM Hepes pH 7.0, 1.5 mM CaCl<sub>2</sub>, 1 mM MgSO<sub>4</sub>, 100 mM NaCl

Binding Protocol: Binding assays are performed in 200 µL total volume according to the

following conditions:

1 - Membrane dilution: 0.05 mL of membranes + 7.45 mL assay buffer (1: 150 dilution)

2 - Incubation: 25 μL of incubation buffer or (Nle<sup>4</sup>,D-Phe<sup>7</sup>)-α-MSH (Bachem H-1100) 3 μM

final for non specific binding (Saturation binding assay)

For competition binding assay: 25 µL of reference compounds at

decreasing concentrations (see figure 2)

25 µL of radioligand at the appropriate concentration (see graph below)

150 µL of diluted membranes

3 - Incubation time: 60 minutes at 37 °C

4 - Filtration: aspirate and wash 9 x 500 μL with ice cold wash buffer over GF/C filter

(presoaked in 0.5 % PEI).

## Lot Specific Data

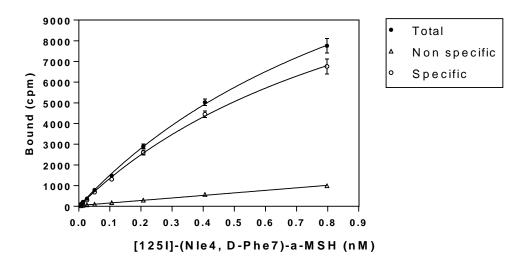


Figure 1: Saturation binding assay curve (filtration) 96-well saturation binding assay curve (0.8 μg membranes/well, TopCount®) using [125]-(Nle4, D-Phe7)-α-MSH (Revvity NEX352 Lot No.: IM22381)



## Typical Product Data

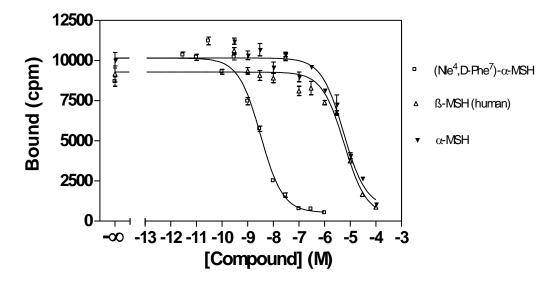


Figure 2: Competition binding assay curve (filtration) 96-well competition binding assay curve (0.8  $\mu$ g membranes/well, TopCount®). Recommended radioligand concentration = 0.8 nM.

\*Even though two sites can be observed occasionally with some ligands, the data presented is derived from single site fitting.

Reference Compounds	Ki
	(nM)
(Nle <sup>4</sup> ,D-Phe <sup>7</sup> )-α-MSH	2.3
ß-MSH (human)	4196
α-MSH	4004

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