

Research use only. Not for use in diagnostic procedures.

Membrane Target Systems™

human Chemokine CXCR3 Receptor

Product No.: ES-142-M400UA

Lot No.: 2490724

Material Provided

Membranes: $1 \times 400 \text{ units} / 400 \mu \text{L}$ frozen aliquot

Product Information

Cellular Background: CHO-K1

GenBank Accession Number: X95876

Unit Size: 2 µg protein / unit

Storage Buffer: 50 mM Tris-HCL (pH 7.4), 0.5mM EDTA, 10mM MgCl₂, 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_{max} and K_d are determined using radioactive saturation binding assays (Figure 1). Protein concentration is determined using the BCA method ⁽¹⁾. Ratio-to-Reference (RTR) is determined by dividing the maximal signal of the current lot (B_{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): 1.4

Expression Level (B_{max}): 21 pmol/mg membrane protein.

 K_d for $[^{125}I]$ -I-TAC: 0.15 nM

Protein Concentration: 2 µg/µL

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



Recommended Assay Conditions

Assay Buffer: 25 mM Hepes pH 7.4, 10 mM MgCl₂, 1 mM CaCl₂, 0.5% BSA

Wash Buffer: 25 mM Hepes pH 7.4, 5 mM MgCl₂, 1 mM CaCl₂, 500 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 human I-TAC (CXCL11) (Peprotech 300-46) 1 μΜ

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 27 °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

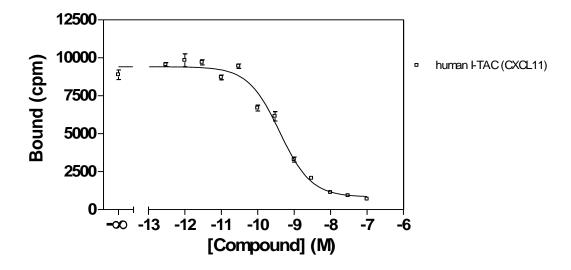


Figure 1: Saturation binding assay curve (filtration) 96-well saturation binding assay curve (2 µg membranes/well, TopCount®) using [125]-I-TAC (Revvity NEX376 Lot No.: JKA2380)



Typical Product Data

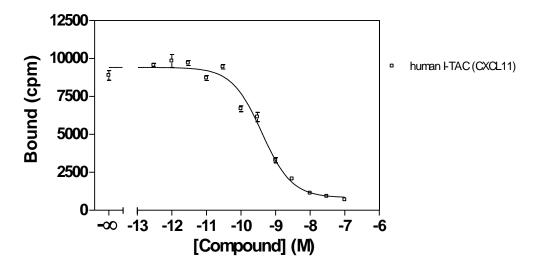


Figure 2: Competition binding assay curve (filtration) 96-well competition binding assay curve (2 μ g membranes/well, TopCount®). Recommended radioligand concentration = 0.04 nM.

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

Reference Compounds	Ki
	(nM)
human I-TAC (CXCL11)	0.34

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