

Research use only. Not for use in diagnostic procedures.

# human ERG K<sup>+</sup> channel

Product No.: RBHERGM400UA
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Lot No.: 2055543

### Material Provided

Membranes:	1 x 400 units / 400 μL frozen aliquot
Product Information	
Cellular Background:	HEK293
GenBank Accession Number:	NM_000238 (KCNH2, transcript variant 1)
Unit Size:	3 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**

 $B_{max}$  and  $K_d$  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 ( $B_{max}$  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):	1.2
Expression Level (B <sub>max</sub> ):	3.0 pmol/mg membrane protein.
$K_d$ for [ <sup>125</sup> I]-BeKm-1 :	0.03 nM
Protein Concentration:	3 μg/μL

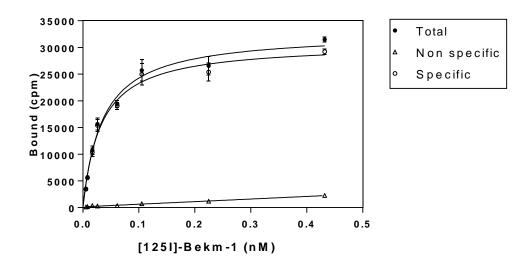
(1) Smith, P.K., et al. (1985). Anal. Biochem. 150, 76-85.

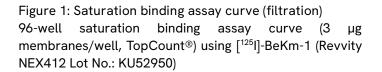


### **Recommended Assay Conditions**

Assay Buffer:	20 mM Hepes-Tris pH 7.2, 0.1 mM KCl, 0.1% BSA
Wash Buffer:	20 mM Tris-HCl pH 7.3, 150 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 BeKm-1 (Calbiochem 198800) 0.125 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 27 °C
4 - Filtration:	aspirate and wash 9 x 500 $\mu L$ with ice cold wash buffer over GF/C filter (presoaked in 0.3% PEI).

## Lot Specific Data







#### **Typical Product Data**

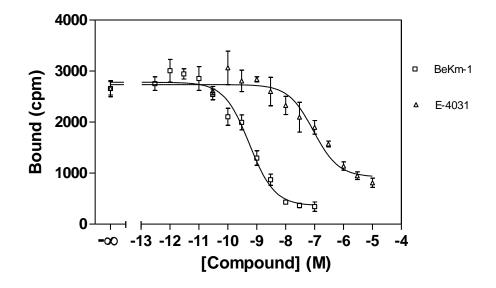


Figure 2: Competition binding assay curve (filtration) 96-well competition binding assay curve (3  $\mu$ g membranes/well, TopCount<sup>®</sup>). Recommended radioligand concentration = 0.1 nM.

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

Reference Compounds	Ki
	(nM)
BeKm-1	0.41
E-4031	68

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**Revvity** 940 Winter Street Waltham, MA 02451 USA

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