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Mouse OLFR12 cDNA Clone in Bacterial Expression Vector (His-GST)

Overview				
Quantity:	500 ng			
Gene:	Olfr12 (OLFR12)			
Species:	Mouse			
Fusion tag:	His-GST			
Insert:	cDNA			
Vector:	Bacterial Expression Vector			
Application:	Cloning (Clon)			
Product Details				
Purpose:	Bacterial expression of Mouse Olfr12 with His-GST			
Insert Length:	1121 bp			
Vector Backbone:	pPB-His-GST			
Promoter:	T7 Promoter			
Bacterial Resistance:	Kanamycin			
Expression Type:	Transient			
Specificity:	5-Nhel and 3-Xhol			
	Fusion tag: Dual N-terminal tag, 6X Histidine followed by Glutathione-S-Transferase Protein			
	which is cleavable with TEV (Size 27.9 kDa)			
Sequencing Primer:	GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-			
	GCTAGTTATTGCTCAGCGG-3'			
Target Details				
Gene:	Olfr12 (OLFR12)			

Target Details Olfr12 Alternative Name: NCBI Accession: NM_206896 **Application Details Application Notes:** The pPB vectors are low-medium copy number vectors in which the gene expression is driven by the strong T7 promoter. Below are some basic guidelines for using the pPB vectors for protein production: 1. The pPB vectors are designed to be used with E. coli strains that are DE3 lysogens i.e. the host E. coli cell has a source of T7 RNA polymerase. 2. Recombinant protein induction is usually done at OD600 of 0.6-1.2 using Isopropyl β-D-1thiogalactopyranoside (IPTG) at a final concentration of 0.05 -1mM. 3. The ideal concentration of IPTG must be determined empirically for each recombinant protein/cell-line. Similarly, the length of time and temperature for induction provide other variables that need to be optimized on a case-to-case basis. 4. For toxic proteins, it is recommended to go for shorter induction time and also to try and suppress basal recombinant gene expression through (a) addition of glucose or use of pLysS plasmid. Please note that special cell-lines are also available in the market that cater to expression of toxic proteins. 5. Once grown for the desired length of time, harvest cells by centrifugation and either freeze the cells at -80°C (as such or after re-suspending in the desired buffer) or proceed with the purification. Restrictions: For Research Use only Handling Format: Liquid Buffer: 10 mM Tris-HCI, 1 mM EDTA, pH 8.0 Storage: -20 °C

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Expiry Date:

Storage Comment:

Product cited in:

Johnson, Drugan, Miller, Evans: "38" in:, Vol. 1363, Issue Nucleic acids research, pp. 28-39, (

1 year when stored at -20° C or lower in a non-frost free freezer.

12 months

1991)