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Datasheet for ABIN4750328

Rat OLR1338 cDNA Clone in Bacterial Expression Vector (His-MBP)

Overview				
Quantity:	500 ng			
Gene:	OLR1338			
Species:	Rat			
Fusion tag:	His-MBP			
Insert:	cDNA			
Vector:	Bacterial Expression Vector			
Application:	Cloning (Clon)			
Product Details				
Purpose:	Bacterial expression of Rat Olr1338 with His-MBP			
Insert Length:	948 bp			
Vector Backbone:	pPB-His-MBP			
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 Maltose Binding Protein which is			
	cleavable with Thrombin (Size 43 kDa)			
Sequencing Primer:	MBP Forward primer: 5'-CGCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'-			
	GCTAGTTATTGCTCAGCGG-3'			
Target Details				
Gene:	OLR1338			

Target Details Olr1338 Alternative Name: NCBI Accession: NM_001000789 **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

Publications

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)