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Datasheet for ABIN4737697 Mouse TCL1B4 cDNA Clone in Bacterial Expression Vector (His-MBP)

Overview

Quantity:	500 ng
Gene:	TCL1B4
Species:	Mouse
Fusion tag:	His-MBP
Insert:	cDNA
Vector:	Bacterial Expression Vector
Application:	Cloning (Clon)

Product Details

Purpose:	Bacterial expression of Mouse Tcl1b4 with His-MBP
Insert Length:	1031 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:

TCL1B4

Alternative Name: Tcl1b4 NCBI Accession: NM_013774 Application Details Image: Comparison of the problem of the proble	
Application Details Application Notes: The pPB vectors are low-medium copy number vectors in which the gene expression i 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 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 (
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thiogalactonyranosida (IDTC) at a final concentration of 0.05. 1mM	-D-1-
3. The ideal concentration of IPTG must be determined empirically for each recombina	nt
protein/cell-line. Similarly, the length of time and temperature for induction provide oth	er
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 o	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	
Storage Comment: 1 year when stored at -20° C or lower in a non-frost free freezer.	

Expiry Date:

12 months

Publications

Product cited in:

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

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Publications
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1991)