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Datasheet for ABIN4718166 Mouse BC030867 cDNA Clone in Bacterial Expression Vector (His-MBP)

Overview

Quantity:	500 ng
Gene:	BC030867
Species:	Mouse
Fusion tag:	His-MBP
Insert:	cDNA
Vector:	Bacterial Expression Vector
Application:	Cloning (Clon)

Product Details

Purpose:Bacterial expression of Mouse BC030867 with His-MBPInsert Length:2543 bpVector Backbone:pPB-His-MBPPromoter:T7 PromoterBacterial Resistance:KanamycinExpression Type:TransientSpecificity:5-Nhel and 3-Xhol Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which cleavable with Thrombin (Size 43 kDa)		
Vector Backbone:pPB-His-MBPPromoter:T7 PromoterBacterial Resistance:KanamycinExpression Type:TransientSpecificity:5-Nhel and 3-Xhol Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which	Purpose:	Bacterial expression of Mouse BC030867 with His-MBP
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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	Vector Backbone:	pPB-His-MBP
Expression Type: Transient Specificity: 5-Nhel and 3-Xhol Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which	Promoter:	T7 Promoter
Specificity: 5-Nhel and 3-Xhol Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which	Bacterial Resistance:	Kanamycin
Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which	Expression Type:	Transient
	Specificity:	Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which is
Sequencing Primer: MBP Forward primer: 5'-CGCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'- GCTAGTTATTGCTCAGCGG-3'	Sequencing Primer:	

Target Details

Gene:

BC030867

Target Details	
Alternative Name:	BC030867
NCBI Accession:	NM_153544
Application Details	
Application Notes:	The pPB vectors are low-medium copy number vectors in which the gene expression is driver
	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-1-
	thiogalactopyranoside (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

Expiry Date:

Storage Comment:

Publications

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

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Publications
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1991)