# -online.com genomics

## Datasheet for ABIN4728522 Mouse MEX3A cDNA Clone in Bacterial Expression Vector (His-MBP)

#### Overview

Quantity:	500 ng
Quantity.	500 Hg
Gene:	MEX3A
Species:	Mouse
Fusion tag:	His-MBP
Insert:	cDNA
Vector:	Bacterial Expression Vector
Application:	Cloning (Clon)

#### Product Details

Purpose:	Bacterial expression of Mouse Mex3a with His-MBP
Insert Length:	5776 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:

Alternative Name: Mex3a (MEX3A Products)   NCBI Accession: NM_001029890   Application Details Image: Comparison of the product of the pr	
Application Details    Application Notes: The pPB vectors are low-medium copy number vectors in which the gene expression   by the strong T7 promoter. Below are some basic guidelines for using the pPB vectors for protein production:	
Application Notes: The pPB vectors are low-medium copy number vectors in which the gene expression   by the strong T7 promoter. Below are some basic guidelines for using the pPB vectors for protein production:	
by the strong T7 promoter. Below are some basic guidelines for using the pPB vectors for protein production:	
Below are some basic guidelines for using the pPB vectors for protein production:	is driven
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 recombin	ant
protein/cell-line. Similarly, the length of time and temperature for induction provide ot	her
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 tr	y and
suppress basal recombinant gene expression through (a) addition of glucose or use c	of pLysS
plasmid. Please note that special cell-lines are also available in the market that cater t expression of toxic proteins.	to
5. Once grown for the desired length of time, harvest cells by centrifugation and either	r freeze
the cells at -80°C (as such or after re-suspending in the desired buffer) or proceed wit	h 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, (

```
Publications
```

1991)