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Datasheet for ABIN4697970

Human BAALC cDNA Clone in Bacterial Expression Vector (His-MBP)

Quantity:500 ngGene:BAALCSpecies:HumanFusion tag:His-MBPInsert:cDNAVector:Bacterial ExpressionApplication:Cloning (Clon)Product DetailsPurpose:Insert Length:438 bpVector Backbone:pPB-His-MBP	
Species: Human Fusion tag: His-MBP Insert: cDNA Vector: Bacterial Expressi Application: Cloning (Clon) Product Details Purpose: Bacterial expressi Insert Length: 438 bp	
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Product Details Purpose: Bacterial expressi Insert Length: 438 bp	ion Vector
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	ion of Human BAALC with His-MBP
Vector Backbone: pPB-His-MBP	
Promoter: T7 Promoter	
Bacterial Resistance: Kanamycin	
Expression Type: Transient	
Specificity: 5-Nhel and 3-Xho	
Fusion tag: Dual N	N-terminal tag, 6X Histidine followed by Maltose Binding Protein which is
cleavable with Th	rombin (Size 43 kDa)
Sequencing Primer: MBP Forward prin	mer: 5'-CGCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'-
GCTAGTTATTGC	TCAGCGG-3'
Target Details	
Gene: BAALC	

Alternative Name:

BAALC (BAALC Products)

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-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
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, (1991)