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Datasheet for ABIN4711486

# **Human SND1 cDNA Clone in Bacterial Expression Vector (His-MBP)**

Quantity:500 ngGene:SND1Species:HumanFusion tag:His-MBPInsert:cDNAVector:Bacterial Expression VectorApplication:Cloning (Clon)Product DetailsPurpose:Purpose:Bacterial expression of Human SND1 with His-MBInsert Length:2658 bpVector Backbone:pPB-His-MBPPromoter:T7 PromoterBacterial Resistance:KanamycinExpression Type:Transient	P
Species: Human  Fusion tag: His-MBP  Insert: cDNA  Vector: Bacterial Expression Vector  Application: Cloning (Clon)  Product Details  Purpose: Bacterial expression of Human SND1 with His-MB  Insert Length: 2658 bp  Vector Backbone: pPB-His-MBP  Promoter: T7 Promoter  Bacterial Resistance: Kanamycin	P
Fusion tag: His-MBP  Insert: cDNA  Vector: Bacterial Expression Vector  Application: Cloning (Clon)  Product Details  Purpose: Bacterial expression of Human SND1 with His-MB  Insert Length: 2658 bp  Vector Backbone: pPB-His-MBP  Promoter: T7 Promoter  Bacterial Resistance: Kanamycin	P
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Expression Type: Transient	
Specificity: 5-Nhel and 3-Xhol	
Fusion tag: Dual N-terminal tag, 6X Histidine follow	ved by Maltose Binding Protein which is
cleavable with Thrombin (Size 43 kDa)	
Sequencing Primer: MBP Forward primer: 5'-CGCAGATGTCCGCTTTCT	ΓGG-3', T7 terminator primer: 5'-
GCTAGTTATTGCTCAGCGG-3'	
Target Details	
Gene: SND1	

Alternative Name:

SND1 (SND1 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  $\beta$ -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)