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Datasheet for ABIN4833747

# **Human IWS1 cDNA Clone in Bacterial Expression Vector (His-GST)**

Application: Cloning  Product Details  Purpose: Bacter  Insert Length: 159 bp	
Species: Human Fusion tag: His-GS Insert: cDNA Vector: Bacter Application: Cloning Product Details Purpose: Bacter Insert Length: 159 bp	
Fusion tag: His-GS Insert: cDNA  Vector: Bacter  Application: Clonin  Product Details  Purpose: Bacter  Insert Length: 159 bp	
Insert: cDNA  Vector: Bacter  Application: Cloning  Product Details  Purpose: Bacter  Insert Length: 159 bp	n
Vector: Bacter Application: Clonin  Product Details  Purpose: Bacter Insert Length: 159 bp	TST TS
Application: Cloning  Product Details  Purpose: Bacter  Insert Length: 159 bp	
Product Details  Purpose: Bacter  Insert Length: 159 bp	rial Expression Vector
Purpose: Bacter Insert Length: 159 bp	g (Clon)
Insert Length: 159 bp	
	rial expression of Human IWS1 with His-GST
Vector Backbone: pPB-H	
•	is-GST
Promoter: T7 Pro	omoter
Bacterial Resistance: Kanan	nycin
Expression Type: Transi	ent
Specificity: 5-Nhel	l and 3-Xhol
Fusion	tag: Dual N-terminal tag, 6X Histidine followed by Glutathione-S-Transferase Protein
which	is cleavable with TEV (Size 27.9 kDa)
Sequencing Primer: GST Fo	orward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-
GCTAC	GTTATTGCTCAGCGG-3'
Target Details	
Gene: IWS1	

Alternative Name:

IWS1 (IWS1 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)