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Datasheet for ABIN4837185

# **Human PCIF1 cDNA Clone in Bacterial Expression Vector (His-GST)**

Gene: F Species: H Fusion tag: H Insert: C Vector: E	500 ng PCIF1 Human His-GST cDNA Bacterial Expression Vector Cloning (Clon)
Species: H Fusion tag: H Insert: C Vector: E Application: C	Human  His-GST  cDNA  Bacterial Expression Vector
Fusion tag:  Insert:  Vector:  Application:	His-GST  cDNA  Bacterial Expression Vector
Insert: Converted to the service of	cDNA  Bacterial Expression Vector
Vector: E Application: C	Bacterial Expression Vector
Application:	
	Cloning (Clon)
Product Details	
Purpose: E	Bacterial expression of Human PCIF1 with His-GST
Insert Length: 2	2115 bp
Vector Backbone: p	pPB-His-GST
Promoter:	T7 Promoter
Bacterial Resistance: k	Kanamycin
Expression Type:	Transient
Specificity: 5	5-Nhel and 3-Xhol
F	Fusion tag: Dual N-terminal tag, 6X Histidine followed by Glutathione-S-Transferase Protein
V	which is cleavable with TEV (Size 27.9 kDa)
Sequencing Primer: 0	GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-
(	GCTAGTTATTGCTCAGCGG-3'
Target Details	
Gene: F	

PCIF1 (PCIF1 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-1thiogalactopyranoside (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)