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Datasheet for ABIN4827272

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

Species: Human Fusion tag: His-GST Insert: cDNA Vector: Bacterial Expression Vector Application: Cloning (Clon)  Product Details  Purpose: Bacterial expression of Human BLID with His-GST Insert Length: 327 bp  Vector Backbone: pPB-His-GST  Promoter: 17 Promoter  Bacterial Resistance: Kanamycin  Expression Type: Transient  Specificity: 5-Nhel 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 Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3'  Target Details	Overview	
Fusion tag: His-GST  Insert: cDNA  Vector: Bacterial Expression Vector  Application: Cloning (Clon)  Product Details  Purpose: Bacterial expression of Human BLID with His-GST  Insert Length: 327 bp  Vector Backbone: pPB-His-GST  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 Glutathione-S-Transferase Protein which is cleavable with TEV (Size 27.9 kDa)  Sequencing Primer: GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3'  Target Details	Quantity:	500 ng
Fusion tag: His-GST  Insert: cDNA  Vector: Bacterial Expression Vector  Application: Cloning (Clon)  Product Details  Purpose: Bacterial expression of Human BLID with His-GST  Insert Length: 327 bp  Vector Backbone: pPB-His-GST  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 Glutathione-S-Transferase Protein which is cleavable with TEV (Size 27.9 kDa)  Sequencing Primer: GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3'  Target Details	Gene:	BLID
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Purpose: Bacterial expression of Human BLID with His-GST  Insert Length: 327 bp  Vector Backbone: pPB-His-GST  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 Glutathione-S-Transferase Protein which is cleavable with TEV (Size 27.9 kDa)  Sequencing Primer: GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3'  Target Details	Vector:	Bacterial Expression Vector
Purpose: Bacterial expression of Human BLID with His-GST  Insert Length: 327 bp  Vector Backbone: pPB-His-GST  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 Glutathione-S-Transferase Protein which is cleavable with TEV (Size 27.9 kDa)  Sequencing Primer: GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3'  Target Details	Application:	Cloning (Clon)
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Target Details	Sequencing Primer:	GST Forward primer: 5'-CACGTTTGGTGGTGGCGAC3', T7 terminator primer: 5'-
		GCTAGTTATTGCTCAGCGG-3'
Gene: BLID	Target Details	
	Gene:	BLID

BLID (BLID 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)