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Human DECR1 cDNA Clone in Bacterial Expression Vector (His-MBP)

Gene: DECR1 Species: Human Fusion tag: His-MBP Insert: cDNA Vector: Bacterial Expression Vector Application: Cloning (Clon) Product Details Purpose: Bacterial expression of Human DECR1 with His-MBP Insert Length: 1009 bp Vector Backbone: pPB-His-MBP 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 Maltose Binding Protein which is cleavable with Thrombin (Size 43 kDa) Sequencing Primer: MBP Forward primer: 5'-GCGAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3' Target Details	Overview	
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Fusion tag: His-MBP Insert: cDNA Vector: Bacterial Expression Vector Application: Cloning (Clon) Product Details Purpose: Bacterial expression of Human DECR1 with His-MBP Insert Length: 1008 bp Vector Backbone: pPB-His-MBP Promoter: 77 Promoter Bacterial Resistance: Kanamycin Expression Type: Transient Specificity: 5-Nhel and 3-Xhol Fusion tag: Dual N-terminal tag, 6X Histidine followed by Maltose Binding Protein which is cleavable with Thrombin (Size 43 kDa) Sequencing Primer: MBP Forward primer: 5'-CGCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3' Target Details	Gene:	DECR1
Insert: cDNA Vector: Bacterial Expression Vector Application: Cloning (Clon) Product Details Purpose: Bacterial expression of Human DECR1 with His-MBP Insert Length: 1008 bp Vector Backbone: pPB-His-MBP 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 Maltose Binding Protein which is cleavable with Thrombin (Size 43 kDa) Sequencing Primer: MBP Forward primer: 5'-GCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3' Target Details	Species:	Human
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Gene: DECR1	Target Details	
	Gene:	DECR1

DECR1 (DECR1 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)