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## Datasheet for ABIN4705075 Human KLRC4 cDNA Clone in Bacterial Expression Vector (His-MBP)

#### Overview

Quantity:	500 ng	
Gene:	KLRC4	
Species:	Human	
Fusion tag:	His-MBP	
Insert:	cDNA	
Vector:	Bacterial Expression Vector	
Application:	Cloning (Clon)	

#### Product Details

Purpose:	Bacterial expression of Human KLRC4 with His-MBP	
Insert Length:	477 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'-CGCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'-GCTAGTTATTGCTCAGCGG-3'	

### Target Details

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Alternative Name:	KLRC4 (KLRC4 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			
Formati	Liquid		

Target Details

Format:	Liquid		
Buffer:	10 mM Tris-HCl, 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)		