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Datasheet for ABIN4842951 Human USP35 cDNA Clone in Bacterial Expression Vector (His-GST)

Overview

Quantity:	500 ng	
	USP35	
Gene:	05835	
Species:	Human	
Fusion tag:	His-GST	
Insert:	cDNA	
Vector:	Bacterial Expression Vector	
Application:	Cloning (Clon)	

Product Details

Purpose:	Bacterial expression of Human USP35 with His-GST	
Insert Length:	1815 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

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USP35

Alternative Name:	USP35 (USP35 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-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

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)	