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Datasheet for ABIN4631537 Human FKBP9 cDNA Clone in Bacterial Expression Vector (His tag)

Overview

Quantity:	500 ng
Gene:	FKBP9
Species:	Human
Fusion tag:	His tag
Insert:	cDNA
Vector:	Bacterial Expression Vector
Application:	Cloning (Clon)

Product Details

Purpose:	Bacterial expression of Human FKBP9 with His tag
Insert Length:	807 bp
Vector Backbone:	pPB-N-His
Promoter:	T7 Promoter
Bacterial Resistance:	Kanamycin
Expression Type:	Transient
Specificity:	5-Nhel and 3-Xhol Fusion tag: A singel N-terminal 6X-Histidine tag which is cleavable with Thrombin (Size 2.3 kDa)
Sequencing Primer:	T7 promoter primer: 5'-TAATACGACTCACTATAGGG-3', T7 terminator primer: 5'- GCTAGTTATTGCTCAGCGG-3'

Target Details

Gene:	FKBP9
Alternative Name:	FKBP9 (FKBP9 Products)

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Application Details Application Notes: The pPB vectors are low-medium copy number vectors in which the gene expression is 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. 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-	
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 host E. coli cell has a source of T7 RNA polymerase.	
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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 recombinar	nt
protein/cell-line. Similarly, the length of time and temperature for induction provide othe	r
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 f	reeze
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	

Target Details

NCBI Accession:

NM_007270

Handling

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)