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Datasheet for ABIN4711624 Human SPAG7 cDNA Clone in Bacterial Expression Vector (His-MBP)

Overview

Quantity:	500 ng	
Gene:	SPAG7	
Species:	Human	
Fusion tag:	His-MBP	
Insert:	cDNA	
Vector:	Bacterial Expression Vector	
Application:	Cloning (Clon)	

Product Details

Vector Backbone: Promoter: Bacterial Resistance:	Bacterial expression of Human SPAG7 with His-MBP	
Vector Backbone: Promoter: Bacterial Resistance:	601 hn	
Promoter: Bacterial Resistance:	684 bp	
Bacterial Resistance:	pPB-His-MBP	
	T7 Promoter	
Expression Type:	Kanamycin	
	Transient	
	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)	
	MBP Forward primer: 5'-CGCAGATGTCCGCTTTCTGG-3', T7 terminator primer: 5'- GCTAGTTATTGCTCAGCGG-3'	

Target Details

G	eı	ne	2
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SPAG7

Alternative Name:

SPAG7 (SPAG7 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

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	
Dublications		
Publications		
Product cited in:	Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (
	1991)	