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## Datasheet for ABIN4834452 Human LDHD cDNA Clone in Bacterial Expression Vector (His-GST)

#### Overview

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

#### Product Details

Purpose:	Bacterial expression of Human LDHD with His-GST
Insert Length:	1524 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

Gene:

LDHD

Application Details Application Notes: Th by Be 1. ho	OHD (LDHD Products)
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Be 1. <sup>-</sup> ho	ne pPB vectors are low-medium copy number vectors in which the gene expression is driven
1. <sup>-</sup> ho	the strong T7 promoter.
ho	elow are some basic guidelines for using the pPB vectors for protein production:
	The pPB vectors are designed to be used with E. coli strains that are DE3 lysogens i.e. the
2.	ost E. coli cell has a source of T7 RNA polymerase.
	Recombinant protein induction is usually done at OD600 of 0.6-1.2 using Isopropyl $\beta$ -D-1-
thi	iogalactopyranoside (IPTG) at a final concentration of 0.05 -1mM.
3.	The ideal concentration of IPTG must be determined empirically for each recombinant
pro	otein/cell-line. Similarly, the length of time and temperature for induction provide other
vai	riables 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
SU	ppress basal recombinant gene expression through (a) addition of glucose or use of pLysS
pla	asmid. Please note that special cell-lines are also available in the market that cater to
ex	pression of toxic proteins.
5. (	Once grown for the desired length of time, harvest cells by centrifugation and either freeze
the	e cells at -80°C (as such or after re-suspending in the desired buffer) or proceed with the
ри	urification.
Restrictions: Fo	or 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
Publications	
Product cited in:	Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (
	1991)