-online.com genomics

Datasheet for ABIN4832933 Human HJURP cDNA Clone in Bacterial Expression Vector (His-GST)

Overview

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

Product Details

| Purpose: | Bacterial expression of Human HJURP with His-GST | |
|-----------------------|---|--|
| Insert Length: | 2247 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

| G | eı | า | 2 |
|---------|----|---|---|
| \circ | | | - |

| Alternative Name: | HJURP (HJURP 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) | |