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Datasheet for ABIN4632954 Human OR52K2 cDNA Clone in Bacterial Expression Vector (His tag)

Overview

| Quantity: | 500 ng |
|--------------|-----------------------------|
| Gene: | OR52K2 |
| Species: | Human |
| Fusion tag: | His tag |
| Insert: | cDNA |
| Vector: | Bacterial Expression Vector |
| Application: | Cloning (Clon) |

Product Details

| Purpose: | Bacterial expression of Human OR52K2 with His tag |
|-----------------------|-----------------------------------------------------------------------------------------------------------------------|
| Insert Length: | 945 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: | OR52K2 |
|-------------------|--------------------------|
| Alternative Name: | OR52K2 (OR52K2 Products) |

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| Target [| Details |
|----------|---------|
|----------|---------|

NCBI Accession:

NM_001005172

Application Details

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