Human IL21 ORF mammalian expression plasmid (Codon Optimized), N-HA tag

Catalog Number: HG10584-NY

General Information
Gene: interleukin 21
Official Symbol: IL21
Synonym: Za11, IL-21
Source: Human
cDNA Size: 405bp
Plasmid: pCMV3-HA-IL21

Description
Lot: Please refer to the label on the tube
Sequence Description:
Identical with the Gene Bank Ref. ID sequence.
Restriction site: KpnI + XbaI (6kb + 0.43kb)
Vector: pCMV3-SP-N-HA
Shipping carrier:
Each tube contains approximately 10 μg of lyophilized plasmid.

Storage:
The lyophilized plasmid can be stored at ambient temperature for three months.

Quality control:
The plasmid is confirmed by full-length sequencing with primers in the sequencing primer list.

Sequencing primer list:

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>pCMV3-F</td>
<td>5’ CAGGTGTCCACTCCAGGTCCAAG 3’</td>
</tr>
<tr>
<td>pcDNA3-R</td>
<td>5’ GGCACTAGAAGGCACAGTCGAGG 3’</td>
</tr>
</tbody>
</table>

Or

<table>
<thead>
<tr>
<th>Primer</th>
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</tr>
</thead>
<tbody>
<tr>
<td>Forward T7</td>
<td>5’ TAATACGACTCTATAGGG 3’</td>
</tr>
<tr>
<td>ReverseBGH</td>
<td>5’ TAGAAGGCACAGTCGAGG 3’</td>
</tr>
</tbody>
</table>

E.coli strains for transformation (recommended but not limited)
Most commercially available competent cells are appropriate for the plasmid, e.g. TOP10, DH5α and TOP10F’.

Plasmid Resuspension protocol
1. Centrifuge at 5,000 × g for 5 min.
2. Carefully open the tube and add 100 μl of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin to concentrate the liquid at the bottom. Speed is less than 5000 × g.
5. Store the plasmid at -20 °C.

The plasmid is ready for:
- Restriction enzyme digestion
- PCR amplification
- E. coli transformation
- DNA sequencing

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Fax: +86-10-51029969  ●  Tel:+86- 400-890-9989  ●  http://www.sinobiological.com
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Vector Information

All of the pCMV vectors are designed for high-level stable and transient expression in mammalian hosts. High-level stable and non-replicative transient expression can be carried out in most mammalian cells. The vectors contain the following elements:

- Human enhanced cytomegalovirus immediate-early (CMV) promoter for high-level expression in a wide range of mammalian cells.
- Hygromycin resistance gene for selection of mammalian cell lines.
- A Kozak consensus sequence to enhance mammalian expression.

Physical Map of Plasmid :

- pCMV3-SP-N-HA
- Vector Name
- Vector Size: 6146bp
- Vector Type: Mammalian Expression Vector
- Expression Method: Constitutive, Stable / Transient
- Promoter: CMV
- Antibiotic Resistance: Kanamycin
- Selection In Mammalian Cells: Hygromycin
- Protein Tag: HA
**pCMV3-SP-N-HA** (suitable for secretary and membrane protein expression)

**Physical Map**

- **CMV promoter**
- **enhancer**
- **TT primer**
- **Kozak**
- **Signal Peptide**
- **N-HA**
- **linker**
- **MCS**
- **BGH reverse primer**

**pCMV3-SP-N-HA** (suitable for secretary and membrane protein expression)

**Comments for pCMV3-SP-N-HA:**

- CMV promoter: bases 250-837
- enhancer: bases 838-1445
- SV40 early promoter: bases 2387-2756
- Hygromycin ORF: bases 2774-3799
- pUC origin: bases 4442-5115
- Kanamycin ORF: bases 5189-6004

**Description**

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<tr>
<td>Protein Tag</td>
<td>HA</td>
</tr>
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<td>Sequencing Primer</td>
<td>Forward: T7(TAATACGACTCACTATAGGG) Reverse: BGH(TAGAAGGCACAGTCGAGG)</td>
</tr>
</tbody>
</table>

**Schematic of pCMV3-SP-N-HA Multiple Cloning Sites**

```
1145  GGTGTCACCTCCGACCATGTTAACTCTTAAATAGGG GCCGCCACC
       Kpn I  Hind III

1175  AAG CTT GGT ACC ATGCCCATGCTGTCTGCTGCTGCCCTGCTTGGCTGGAAGCTG

1535  TAT CCT TAG GAC GTG CCT GAC TAC GCC GTG GGA GGC GGT AGC GCT
       Hind III

1588  GCT AGC GGA TCC GTT AAC CTT AAG ACC GGT GAT ATC ATC GAT TAA A
      Nho I Bam HI Hpa II Age I Eco R V Stop Codon

1629  CTC GAG TCT AGA GGC GCC GCC GAATTC GGG CCC GTTAAAC
       Xba I Not I Apa I

1667  CCGCTGAGACGGCCTGCTGCTGCTGCTTCTA GTTGGCACGATCTGTTGTG
       BGH Primer binding site
```

**pCMV3-SP-N-HA** is recommended for constructing the N-HA tag secretary and membrane proteins expression vector which containing a naïve signal peptide. An universal signal peptide is used to instead the naïve signal peptide.