Canine CSF1R Gene cDNA clone plasmid

Catalog Number: DG70086-G

General Information

Gene: colony stimulating factor 1 receptor

Official Symbol: CSF1R

Synonym: CSF1R

Source: Canine

cDNA Size: 2904bp

RefSeq: XM_546306.3

Plasmid: pGEM-dCSF1R

Description

Lot: Please refer to the label on the tube

Sequence Description:

Identical with the Gene Bank Ref. ID sequence except for the point mutations: 554 C/T, 1495 C/T resulting in the amino acid Thr substitution by Met, Pro substitution by Ser and 219 T/C, 1044 A/G, 1539 G/T not causing the amino acid variation.

Vector:

pGEM-T

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>
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<tbody>
<tr>
<td>M13-47</td>
<td>5' GCCAGGGTTTTCCCCAGTCACGAC 3'</td>
</tr>
<tr>
<td>RV-M</td>
<td>5' GAGCGGATAACAAATTTCACACAGG 3'</td>
</tr>
</tbody>
</table>

Other M13 primers can also be used as sequencing primers.

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

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⁻.
Vector Information
The pGEM-T vector is a high-efficiency TA cloning vector which contains multiple cloning sites as shown below. The pGEM-T vector is 3.0kb in size and contains the ampicillin resistance gene for selection. The coding sequence was inserted by TA cloning.

Physical Map of pGEM-T: