# Mouse CSF1R / MCSF Receptor / CD115 Protein (His Tag), Biotinylated

Catalog Number: 50059-M08H-B



# **General Information**

## Gene Name Synonym:

Al323359; CD115; CSF-1R; Csfmr; Fim-2; Fms; M-CSF-R; M-CSFR

#### **Protein Construction:**

A DNA sequence encoding the mouse CSF1R (NP\_001032948.2) extracellular domain (Met1-Ser511) was fused with a polyhistidine tag at the C-terminus. The expressed protein was biotinylated in vitro.

Source: Mouse

Expression Host: HEK293 Cells

**QC** Testing

**Purity:** > 97% as determined by SDS-PAGE.

#### **Bio Activity:**

1.Measured by its binding ability in a functional ELISA. 2.Immobilized mouse IL34-His (Cat:50055-M08H) at  $10\mu g/mL$  (100 $\mu L/well$ ) can bind?biotinylated mouse CSF1R (Cat:50059-M08H-B), The EC $_{50}$  of biotinylated mouse CSF1R is 0.1-0.4 $\mu g/mL$ .

#### **Endotoxin:**

< 1.0 EU per µg protein as determined by the LAL method.

#### Stability:

Samples are stable for up to twelve months from date of receipt at -70  $^{\circ}\mathrm{C}$ 

Predicted N terminal: Ala 20

## **Molecular Mass:**

The recombinant mouse CSF1R consists of 503 amino acids and has a predicted molecular mass of 56.8 kDa.

#### Formulation:

Lyophilized from sterile Sterile PBS

Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as protectants before lyophilization. Specific concentrations are included in the hardcopy of COA. Please contact us for any concerns or special requirements.

# **Usage Guide**

## Storage:

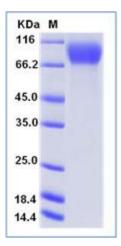
Store it under sterile conditions at  $-20^{\circ}$ C to  $-80^{\circ}$ C upon receiving. Recommend to aliquot the protein into smaller quantities for optimal storage.

Avoid repeated freeze-thaw cycles.

#### Reconstitution:

Detailed reconstitution instructions are sent along with the products.

#### SDS-PAGE:



# **Protein Description**

M-CSFR encoded by the proto-oncogene c-fms is the receptor for colony stimulating factor 1 (CSF1R), a cytokine involved in the proliferation, differentiation, and activation of macrophages. This cell surface glycoprotein is consisted by an extracellular ligand-binding domain, a single membrane-spanning segment, and an intracellular tyrosine kinase domain. Binding of CSF1 activates the receptor kinase, leading to "autophosphorylation" of receptor subunits and the concomitant phosphorylation of a series of cellular proteins on tyrosine residues. CSF1R is a tyrosine kinase receptor that is absolutely required for macrophage differentiation and thus occupies a central role in hematopoiesis. CSF1 and its receptor (CSF1R, product of c-fms protooncogene) were initially implicated as essential for normal monocyte development as well as for trophoblastic implantation. This apparent role for CSF1/CSF1R in normal mammary gland development is very intriguing because this receptor/ligand pair has also been found to be important in the biology of breast cancer in which abnormal expression of CSF1 and its receptor correlates with tumor cell invasiveness and adverse clinical prognosis. Tumor cell expression of CSF1R is under the control of several steroid hormones (glucocorticoids and progestins) and the binding of several bHLH transcription factors, while tumor cell expression of CSF-1 appears to be regulated by other hormones, some of which are involved in normal lactogenic differentiation. However, studies have demonstrated that CSF1 and CSF1R have additional roles in mammary gland development during pregnancy and lactation. The role of CSF1 and CSF1R in normal and neoplastic mammary development that may elucidate potential relationships of growth factor-induced biological changes in the breast during pregnancy and tumor progression.

#### References

1.Sherr CJ. (1990) The colony-stimulating factor 1 receptor: pleiotropy of signal-response coupling. Lymphokine Res. 9(4): 543-8. 2.Kacinski BM. (1997) CSF-1 and its receptor in breast carcinomas and neoplasms of the female reproductive tract. Mol Reprod Dev. 46(1): 71-4. 3.Sapi E, *et al.* (1999) The role of CSF-1 in normal and neoplastic breast physiology. Proc Soc Exp Biol Med. 220(1): 1-8.

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