

DESCRIPTION

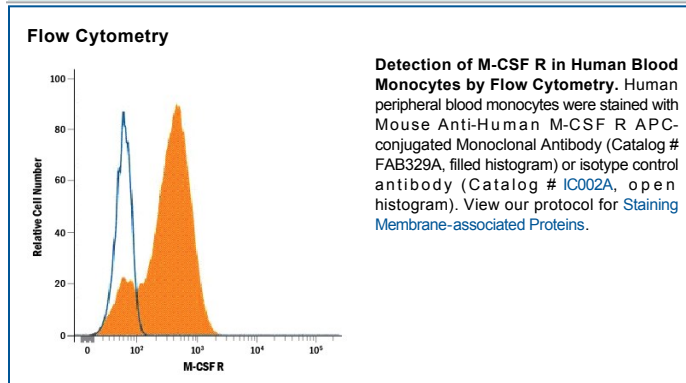
| | |
|---------------------------|--|
| Species Reactivity | Human |
| Specificity | Detects human M-CSF R in direct ELISAs and Western blots. Does not cross-react with recombinant human (rh) GM-CSF R α or rhGM-CSF R β . |
| Source | Monoclonal Mouse IgG ₁ Clone # 61708 |
| Purification | Protein A or G purified from hybridoma culture supernatant |
| Immunogen | Mouse myeloma cell line NS0-derived recombinant human M-CSF R Ile20-Glu512 (Pro54Ala) Accession # P07333.2 |
| Conjugate | Allophycocyanin Excitation Wavelength: 620-650 nm Emission Wavelength: 660-670 nm |
| Formulation | Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details. *Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions. |

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

| | Recommended Concentration | Sample |
|-----------------------|----------------------------------|---------------|
| Flow Cytometry | 10 μ L/10 ⁶ cells | See Below |

DATA



PREPARATION AND STORAGE

Shipping The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage **Protect from light. Do not freeze.**

- 12 months from date of receipt, 2 to 8 °C as supplied.

BACKGROUND

M-CSF Receptor, the product of the *c-fms* proto-oncogene, is a member of the type III subfamily of receptor tyrosine kinases that also includes receptors for SCF and PDGF. These receptors each contain five immunoglobulin-like domains in their extracellular domain (ECD) and a split kinase domain in their intracellular region (1-4). M-CSF Receptor is expressed primarily on cells of the monocyte/macrophage lineage, dendritic cells, stem cells and in the developing placenta (1). Human M-CSF receptor cDNA encodes a 972 amino acid (aa) type I membrane protein with a 19 aa signal peptide, a 493 aa extracellular region containing the ligand-binding domain, a 25 aa transmembrane domain, and a 435 aa cytoplasmic domain. The human M-CSF R ECD shares 60%, 64%, 72%, 75%, 75%, and 76% aa identity with mouse, rat, bovine, canine, feline, and equine M-CSF R, respectively. Activators of protein kinase C induce TACE/ADAM17 cleavage of the M-CSF receptor, releasing the functional ligand-binding extracellular domain (5). M-CSF binding induces receptor homodimerization, resulting in transphosphorylation of specific cytoplasmic tyrosine residues and signal transduction (6). The intracellular domain of activated M-CSF R binds more than 150 proteins that affect cell proliferation, survival, differentiation and cytoskeletal reorganization. Among these, PI3 Kinase, P42/44 ERK, and c-Cbl are key transducers of M-CSF R signals (3, 4). M-CSF R engagement is continuously required for macrophage survival and regulates lineage decisions and maturation of monocytes, macrophages, osteoclasts and dendritic cells (3, 4). M-CSF R and integrin $\alpha\beta_3$ share signaling pathways during osteoclastogenesis, and deletion of either causes osteopetrosis (7, 8). In the brain, microglia expressing increased M-CSF R are concentrated with Alzheimer's A β peptide, but their role in pathogenesis is unclear (9, 10).

References:

1. deParseval, N. *et al.* (1993) *Nucleic Acids Res.* **21**:750.
2. Rothwell, V.M. and L.R. Rohrschneider (1987) *Oncogene Res.* **1**:311.
3. Chitu, V. and E.R. Stanley (2006) *Curr. Opin. Immunol.* **18**:39.
4. Ross, F.P. and S.L. Teitelbaum (2005) *Immunol. Rev.* **208**:88.
5. Roida, E. *et al.* (2001) *J. Immunol.* **166**:1583.
6. Yeung, Y. *et al.* (1998) *J. Biol. Chem.* **273**:17128.
7. Dai, X. *et al.* (2002) *Blood* **99**:111.
8. Faccio, R. *et al.* (2003) *J. Clin. Invest.* **111**:749.
9. Li, M. *et al.* (2004) *J. Neurochem.* **91**:623.
10. Mitrasinovic, O.M. *et al.* (2005) *J. Neurosci.* **25**:4442.