



**ABCLONAL BIOTECHNOLOGY, INC.**

## M-CSF Receptor Rabbit pab Antibody

### Anti M-CSF Receptor antibody

<b>Catalog Number:</b>	A0736	<b>Quantity:</b>	100ul
<b>Lot Number:</b>	A00009	<b>Species:</b>	Rabbit
<b>Gene ID:</b>	1436	<b>Swiss Prot:</b>	P07333

### DESCRIPTION

<b>Description</b>	Rabbit polyclonal to Human M-CSF Receptor
<b>Species</b>	Rabbit
<b>Applications</b>	WB FC
<b>Reactivity</b>	H
<b>Immunogen</b>	A recombinant protein of human M-CSF Receptor
<b>Other Name</b>	CD_antigen=CD115; Flags: Precursor; Proto-oncogene c-Fms; Macrophage colony-stimulating factor 1 receptor;CSF-1-R; FMS;CSF1R;

### PROPERTIES

<b>Form</b>	Liquid
<b>Storage instructions</b>	Upon delivery aliquot and store at -20°C or -80°C.
<b>Storage buffer</b>	PBS with 0.1% Sodium Azide, 50% Glycerol,
<b>Purity</b>	Affinity purification
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	IgG

### APPLICATION

<b>WB</b>	WB :1/200-500
<b>FC</b>	FC:1/10-50



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### BACKGROUND

Macrophage-colony stimulating factor (M-CSF, CSF-1) receptor is an integral membrane tyrosine kinase encoded by the *c-fms* proto-oncogene. M-CSF receptor is expressed in monocytes (macrophages and their progenitors) and drives growth and development of this blood cell lineage. (1-3). Binding of M-CSF to its receptor induces receptor dimerization, activation and autophosphorylation of cytoplasmic tyrosine residues used as docking sites for SH2-containing signaling proteins (4). There are at least five major tyrosine autophosphorylation sites. Tyr723 (Tyr721 in mouse) is located in the kinase insert (KI) region. Phosphorylated Tyr723 binds the p85 subunit of PI3 kinase as well as PLC- $\gamma$  2 (5). Phosphorylation of Tyr809 provides a docking site for Shc (5). Overactivation of this receptor can lead to a malignant phenotype in various cell systems (6). The activated M-CSF receptor has been shown to be a predictor of poor outcome in advanced epithelial ovarian carcinoma (7) and breast cancer (8).

After initial dimerization and autophosphorylation, the CSF-1 receptor undergoes regulated intramembrane proteolysis (RIP) which involves proteolytic processing of this membrane protein and results in release of extracellular domain, intramembrane cleavage and release of the cytoplasmic domain into the cytosol (9). The activated intracellular domain then moves to the nucleus and regulates transcription of specific genes (10). It has been shown that the processing and down modulation of CSF-1 receptor is a continuous process and its rate increases substantially in response to a variety of stimuli including PMA, LPS, tumor necrosis factor, IL-2, IL-4 and its physiological ligand CSF-1 (9).

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6. Morley, G. M. et al. (1999) *Oncogene* 18, 3076-3084.
7. Toy, E. P. et al. (2001) *Gynecol. Oncol.* 80, 194-200.
8. Maher, M. G. et al. (1998) *Clin. Cancer Res.* 4, 1851-1856.
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