SANTA CRUZ BIOTECHNOLOGY, INC.

F4/80 (BM8): sc-52664



BACKGROUND

The epidermal growth factor (EGF)-TM7 family constitutes a group of class B G protein-coupled receptors, which includes CD97, EMR1(EGF-like molecule containing mucin-like hormone receptor 1, designated F4/80 in mouse), EMR2, EMR3, FIRE and ETL. These family members are characterized by an extended extracellular region with several N-terminal EGF domains, and are predominantly expressed on cells of the immune system. The EGF-TM7 protein family are encoded by a gene cluster on human chromosome 19p13.3. The F4/80 molecule is solely expressed on the surface of macrophages and serves as a marker for mature macrophage tissues, including Kupffer cells in liver, splenic red pulp macrophages, brain microglia, gut lamina propria and Langerhans cells in the skin. F4/80/EMR1 undergoes extensive N-linked glycosylation as well as some 0-linked glycosylation. The function of F4/80/EMR1 is unclear, but it is speculated to be involved in macrophage adhesion events, cell migration, or as a G protein-coupled signaling component of macrophages.

CHROMOSOMAL LOCATION

Genetic locus: ADGRE1 (human) mapping to 19p13.3; Adgre1 (mouse) mapping to 17 D.

SOURCE

F4/80 (BM8) is a rat monoclonal antibody raised against cultured bone marrow-derived macrophages of mouse origin.

PRODUCT

Each vial contains 100 $\mu g~lg G_{2a}$ in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

F4/80 (BM8) is recommended for detection of EMR1 of human origin, F4/80 of mouse origin and the corresponding rat homolog by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and flow cytometry (1 μ g per 1 x 10⁶ cells).

Suitable for use as control antibody for EMR1 siRNA (h): sc-72157, F4/80 siRNA (m): sc-42865, EMR1 shRNA Plasmid (h): sc-72157-SH, F4/80 shRNA Plasmid (m): sc-42865-SH, EMR1 shRNA (h) Lentiviral Particles: sc-72157-V and F4/80 shRNA (m) Lentiviral Particles: sc-42865-V.

Molecular Weight of F4/80: 160 kDa.

Positive Controls: WEHI-3 cell lysate: sc-3815 or M1 whole cell lysate: sc-364782.

STORAGE

Store at 4° C, **D0 NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

SELECT PRODUCT CITATIONS

- Blumer, M.J., et al. 2008. Localization of tartrate-resistant acid phosphatase (TRAP), membrane type-1 matrix metalloproteinases (MT1-MMP) and macrophages during early endochondral bone formation. J. Anat. 213: 431-441.
- Shi, X. 2010. Resident macrophages in the cochlear blood-labyrinth barrier and their renewal via migration of bone-marrow-derived cells. Cell Tissue Res. 342: 21-30.
- 3. Fan, Y.Y., et al. 2011. Nicotinic acetylcholine receptor α 7 subunit is timedependently expressed in distinct cell types during skin wound healing in mice. Histochem. Cell Biol. 135: 375-387.
- Mouline, C.C., et al. 2012. Monocytes differentiation upon treatment with a peptide corresponding to the C-terminus of activated T cell-expressed Tirc7 protein. J. Cell. Physiol. 227: 3088-3098.
- Osawa, Y., et al. 2013. Tumor necrosis factor-α promotes cholestasisinduced liver fibrosis in the mouse through tissue inhibitor of metalloproteinase-1 production in hepatic stellate cells. PLoS ONE 8: e65251.
- 6. Yan, L., et al. 2014. Role of OGR1 in myeloid-derived cells in prostate cancer. Oncogene 33: 157-164.
- Wen, G., et al. 2015. A novel role of matrix metalloproteinase-8 in macrophage differentiation and polarization. J. Biol. Chem. 290: 19158-19172.
- Vance, M., et al. 2016. AAV gene therapy for MPS1-associated corneal blindness. Sci. Rep. 6: 22131.
- Kim, D.S., et al. 2017. Smac mimetics and oncolytic viruses synergize in driving anticancer T-cell responses through complementary mechanisms. Nat. Commun. 8: 344.
- Guo, Y., et al. 2018. Tim-3 exacerbates kidney ischaemia/reperfusion injury through the TLR-4/NFκB signalling pathway and an NLR-C4 inflammasome activation. Clin. Exp. Immunol. 193: 113-129.
- Huang, Y.W., et al. 2019. Wound healing can be improved by (-)-epigallocatechin gallate through targeting Notch in streptozotocin-induced diabetic mice. FASEB J. 33: 953-964.
- Bumdelger, B., et al. 2020. Disruption of Osteoprotegerin has complex effects on medial destruction and adventitial fibrosis during mouse abdominal aortic aneurysm formation. PLoS ONE 15: e0235553.
- Su, L., et al. 2021. Potential role of senescent macrophages in radiationinduced pulmonary fibrosis. Cell Death Dis. 12: 527.



See **F4/80 (C-7): sc-377009** for F4/80 antibody conjugates, including AC, HRP, FITC, PE, and Alexa Fluor[®] 488, 546, 594, 647, 680 and 790.