

Anti-CCL2/MCP1 Antibody

HA500042



Product Type:	Rabbit polyclonal IgG, primary antibodies
Species reactivity:	Human
Applications:	WB, IHC-P, IF-Cell
Molecular Wt:	Predicted band size: 11 kDa.

Description: Acts as a ligand for C-C chemokine receptor CCR2 . Signals through binding and activation of CCR2 and induces a strong chemotactic response and mobilization of intracellular calcium ions Exhibits a chemotactic activity for monocytes and basophils but not neutrophils or eosinophils. May be involved in the recruitment of monocytes into the arterial wall during the disease process of atherosclerosis.Expressed in the seminal plasma, endometrial fluid and follicular fluid (at protein level).

Immunogen: Synthetic peptide within human CCL2/MCP1 aa 50-99.

Positive control: A549 cell lysate, Hela cell lysate, human colon tissue, human colon carcinoma tissue.

Subcellular location: Secreted.

Database links: SwissProt: P13500 Human

Recommended Dilutions:

WB	1:500
IHC-P	1:100-1:500
IF-Cell	1:50-1:200

Storage Buffer: 1*TBS (pH7.4), 0.2% BSA, 50% Glycerol. Preservative: 0.05% Sodium Azide.

Storage Instruction: Store at +4°C after thawing. Aliquot store at -20°C. Avoid repeated freeze / thaw cycles.

Purity: Immunogen affinity purified.

Hangzhou Huaan Biotechnology Co., Ltd.

Orders:0086-571-88062880

Technical:0086-571-89986345

Service mail:support@huabio.cn

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Images

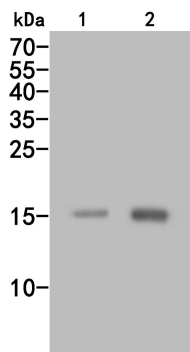


Fig1: Western blot analysis of CCL2/MCP1 on different lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (HA500042, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:5,000 dilution was used for 1 hour at room temperature.

Positive control:

Lane 1: A549 cell lysate

Lane 2: Hela cell lysate

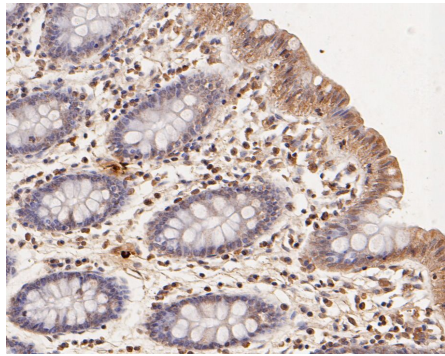


Fig2: Immunohistochemical analysis of paraffin-embedded human colon tissue using anti-CCL2/MCP1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (HA500042, 1/800) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

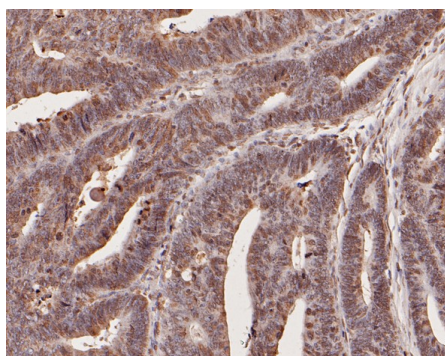


Fig3: Immunohistochemical analysis of paraffin-embedded human colon carcinoma tissue using anti-CCL2/MCP1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (HA500042, 1/200) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

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Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

1. Li Y.S. et. al. The expression of monocyte chemotactic protein (MCP-1) in human vascular endothelium in vitro and in vivo. Mol. Cell. Biochem. 126:61-68(1993).

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