



Code No. KT118

For research use only

Anti Human Macrophage Scavenger Receptor A (MSR-A:CD204) Monoclonal Antibody (Clone No. SRA-C6)

Class A macrophage scavenger receptor (MSR-A: CD204) was identified in the search for the receptor molecules that are implicated in the pathological deposition of cholesterol during atherogenesis through receptor-mediated uptake of modified low density lipoprotein (LDL). MSR-A possesses a wide range of ligand-binding specificities and recognize a variety of molecules such as modified LDL including acetylated LDL, oxidized LDL, advanced glycation end products (AGEs), polyribonucleotides such as poly G and poly I and bacterial surface lipids including lipopolysaccharide and lipoteicoic acid.

This antidody was produced from the mouse immunized with recombinant protein of human type I MSR-A and has been proved to be useful for the western blotting and immunohistochemistry. This antibody also inhibits the endocytic degradation of acetylated LDL and oxidized LDL by high glucose-treated human monocyte-derived macrophages and has anti MSR-A neutralizing activity.

This antibody is useful tools for the study of MSR-A in atherogenesis and various other pathological conditions.

Package Size $50 \mu g$ (200 μl / vial)

Format Mouse monoclonal antibody 0.25mg/ml

Buffer PBS [containing 2% Block Ace as a stabilizer, 0.1%Proclin as a bacteriostat]

Storage Store below -20°C

Once thawed, store at 4°C. Repeated freeze-thaw cycles should be avoided.

Clone No. SRA-C6 Subclass IgG1

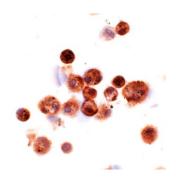
Purification method The spleen cells obtained from MSR-A deficient mouse, immunized with

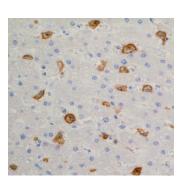
recombinant protein corresponding to amino acid 131-451 of human type I MSR-A, were fused with mouse NS-1 myeloma cells. The hybridoma cell line with positive reaction was grown on non-serum medium, from which the

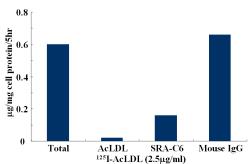
antibody was purified by Protein G affinity chromatography.

Working dilution for immunohistochemistry: 5.0μ g/ml, western blotting : 1.0μ g/ml,

Neutralization: Depends on the experimental design(Application Reference:1)







Left: Human alveolar macrophages(Cytospin preparation): Most macrophages are positive.

Center: Human liver (paraffin section): Kupffer cells are positive

Right: Neutralizing activity of SRA-C6 (20 μ g/ml): Inhibitory effect of anti-human SR-A antibody on the degradation of ¹²⁵I-AcLDL by human monocyte-derived macrophages(day7)

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[Specificity]

| Organ | Reaction | | | Reaction | |
|-----------------|--|---|---------------------------------|--|-------------------------|
| | Positive | Negative | Organ | Positive | Negative |
| Heart | Intramuscular M ϕ (+-) | | Trachea | Mucosal M φ (+-) | |
| Lung | Alveolar M ϕ (+) M ϕ in alveolar septa (+-) | | Esophagus | Interstitial M ϕ (+-) | |
| Liver | Kupffer cells (+) M φ in portal triads(+) | | Stomach | $M \phi$ in lamina propria(+) $M \phi$ in striated muscle(+-) | |
| Kidney | Interstitial M ϕ (+) | | Small and large intestines | $M \phi$ in lamina propria(+) $M \phi$ in striated muscle(+-) | |
| Spleen | Red pulp M φ (+) | Interdigitating cells | Skin | Dermal M φ (+) | Langerhans cells |
| Thymus | Interlobular M ϕ (+) | | Brain (cerebrum and cerebellum) | Perivascular M φ (Mato cells) (+) | |
| Lymph nodes | Sinus M φ (+) | Tingible body M φ Interdigitating cells | Testes | Interstitial M ϕ (+) | |
| Pancreas | Interlobular M φ (+) | | Uterus | Interstitial M φ (+) | |
| Salivary gland | Interlobular M φ (+) | | Ovaries | Interstitial M φ (+) | |
| Thyroid | Interfollicluar M φ (+-) | | Placenta | Hofbauer cells (+) | |
| Parathyroid | Interlobular M φ (+-) | | Bone marrow | M φ (+) | Myeloid precursor cells |
| Adrenals | Interstitial M ϕ (+) | | Blood monocyte | 3 days in culture (+) | Freshly isolated |
| Urinary bladder | Interstitial M φ (+-) | | | | |
| Prostate | Interstitial M ϕ (+-) | | | | |

M ϕ : macrophage \cdot , (+): most cells were positive; (+-): about 10-50% of cells were positive

[Application Reference]

- 1. Fukuhara-Takaki K., Sakai M., Sakamoto Y., Takeya M., Horiuchi S.: Expression of class A scavenger receptor is enhanced by high glucose in vitro and under diabetic conditions in vivo: one mechanism for an increased rate of atherosclerosis in diabetes.: J Biol Chem. 280(5): 3355-3364, 2005
- 2. Tomokiyo R., Jinnouchi K., Honda M., Wada Y., Hanada N., Hiraoka T., Suzuki H., Kodama T., Takahashi K., Takeya M.: Production, characterization, and interspecies reactivities of monoclonal antibodies against human class A macrophage scavenger receptors: Atherosclerosis, 161:123-132, 2002

Manufacturer



Medicinal Chemistry Pharmaceutical Co., Ltd.

Previous manufacturer



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