



Code No. KAL-KT118-S

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  $\mu$  1 / vial)

Format Mouse monoclonal antibody 0.25mg/ml

Buffer PBS [not containing the additive agent and is filtered through a 0.22 µm filter]

Storage Store below -20°C

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

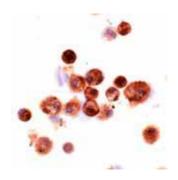
Clone No. SRA-C6
Subclass IgG1

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

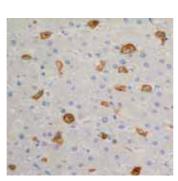
recombinant protein corresponding to amino acid 131-451 of human type I MSR-A, were fused with mouse NS-1 myeloma cells. The hybridoma cell line with positive reaction was grown in ascitic fluid of BALB/c mouse, from which

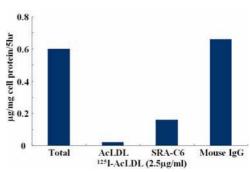
the antibody was purified by Protein G affinity chromatography. immunohistochemistry:  $5.0 \mu$  g/ml, western blotting :  $1.0 \mu$  g/ml,

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



Working dilution for





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  $\mu$  g/ml): Inhibitory effect of anti-human SR-A antibody on the degradation of <sup>125</sup>I-AcLDL by human monocyte-derived macrophages(day7)

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## [Specificity]

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

M  $\phi$ : 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

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