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# anti-MSRA antibody (AA 24-235)





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#### Overview

Quantity:	100 μL
Target:	MSRA
Binding Specificity:	AA 24-235
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This MSRA antibody is un-conjugated
Application:	Immunohistochemistry (IHC), ELISA

#### **Product Details**

Immunogen:	Recombinant Human Mitochondrial peptide methionine sulfoxide reductase protein (24-235AA)
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

## Target Details

Target:	MSRA
Alternative Name:	MSRA (MSRA Products)
Background:	Background: Has an important function as a repair enzyme for proteins that have been
	inactivated by oxidation. Catalyzes the reversible oxidation-reduction of methionine sulfoxide in

proteins to methionine.

Aliases: Cytosolic methionine S sulfoxide reductase antibody, Methionine sulphoxide reductase A antibody, Mitochondrial peptide methionine sulfoxide reductase antibody, MSR A antibody, msrA antibody, MSRA\_HUMAN antibody, peptide met (0) reductase antibody, Peptide Met(0) reductase antibody, Peptide methionine (S) S oxide reductase antibody, Peptide-methionine (S)-S-oxide reductase antibody, PMSR antibody, Protein methionine S oxide reductase antibody, Protein-methionine-S-oxide reductase antibody

UniProt:

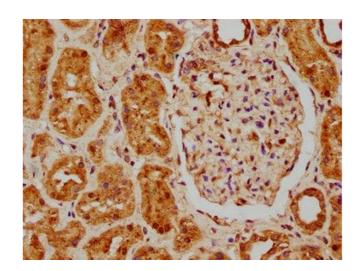
Q9UJ68

### **Application Details**

Application Notes:	Recommended dilution: IHC:1:200-1:500,
Restrictions:	For Research Use only
Handling	
Format:	Liquid

Buffer:	Preservative: 0.03 % Proclin 300
	Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4

	Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.



#### **Immunohistochemistry**

Image 1. IHC image of ABIN7159929 diluted at 1:200 and staining in paraffin-embedded human kidney tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.