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## anti-SAA2 antibody (AA 19-94)





#### Overview

Quantity:	100 μg
Target:	SAA2
Binding Specificity:	AA 19-94
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SAA2 antibody is un-conjugated
Application:	Immunohistochemistry (IHC), ELISA

## **Product Details**

Immunogen:	Recombinant Human Serum amyloid A-2 protein (19-94AA)
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

## Target Details

Target:	SAA2
Alternative Name:	SAA2 (SAA2 Products)
Background:	Background: Major acute phase reactant. Apolipoprotein of the HDL complex.  Aliases: SAA2 antibody, Serum amyloid A-2 protein antibody, SAA2 antibody

#### **Target Details**

UniProt:	PODJI9
Pathways:	Toll-Like Receptors Cascades

### **Application Details**

Application Details	
Application Notes:	Recommended dilution: IHC:1:500-1:1000,
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin

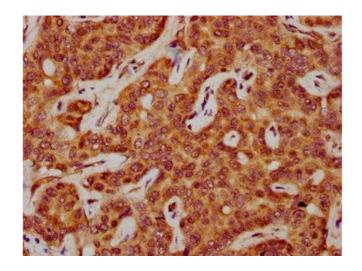
Storage: -20 °C,-80 °C

Storage Comment: Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

handled by trained staff only.

#### **Images**

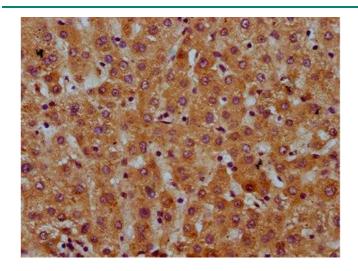
Precaution of Use:



#### **Immunohistochemistry**

This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be

Image 1. IHC image of ABIN7169368 diluted at 1:600 and staining in paraffin-embedded human liver cancer 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.



#### **Immunohistochemistry**

Image 2. IHC image of ABIN7169368 diluted at 1:600 and staining in paraffin-embedded human liver 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.