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# Datasheet for ABIN6244194 anti-MKS1 antibody (AA 90-124)

3 Images



### Overview

Quantity:	200 µL
Target:	MKS1
Binding Specificity:	AA 90-124
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This MKS1 antibody is un-conjugated
Application:	Western Blotting (WB), Immunofluorescence (IF), Flow Cytometry (FACS)

## Product Details

Immunogen:	This MKS1 antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 90-124 amino acids from the human region of human MKS1.	
Clone:	RB57948	
Isotype:	Ig Fraction	
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.	

# Target Details

Target:	MKS1
Alternative Name:	MKS1 (MKS1 Products)
Background:	Component of the tectonic-like complex, a complex localized at the transition zone of primary

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	cilia and acting as a barrier that prevents diffusion of transmembrane proteins between the cilia		
	and plasma membranes. Involved in centrosome migration to the apical cell surface during		
	early ciliogenesis. Required for ciliary structure and function, including a role in regulating length		
	and appropriate number through modulating centrosome duplication. Required for cell		
	branching morphology.		
Molecular Weight:	64528		
UniProt:	Q9NXB0		

## Application Details

Application Notes:	IF: 1:25. WB: 1:2000. FC: 1:25
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Purified polyclonal antibody supplied in PBS with 0.09 % (W/V) sodium azide.

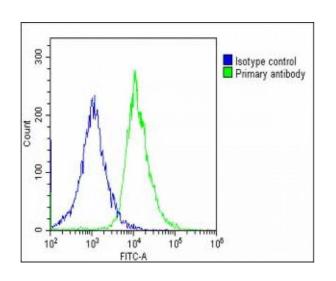
Images	

Expiry Date:

Storage:

Preservative:

Precaution of Use:



Sodium azide

4 °C,-20 °C

6 months

should be handled by trained staff only.

#### Flow Cytometry

This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which

**Image 1.** Overlay histogram showing HepG2 cells stained with (ABIN6244194 and ABIN6578872)(green line). The cells were fixed with 2 % paraformaldehyde (10 min) and then permeabilized with 90 % methanol for 10 min. The cells were then icubated in 2 % bovine serum albumin to block non-specific protein-protein interactions followed by the antibody ((ABIN6244194 and ABIN6578872), 1:25 dilution) for 60 min at 37 °C. The secondary antibody used was Goat-

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Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed(OE188374) at 1/200 dilution for 40 min at 37 °C. Isotype control antibody (blue line) was rabbit IgG1 (1 µ g/1x10^6 cells) used under the same conditions. Acquisition of >10, 000 events was performed.

#### **Western Blotting**

Image 2. All lanes : Anti-MKS1 Antibody (N-Term) at 1:2000 dilution Lane 1: Human kidney lysate Lane 2: Human lung lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 65 kDa Blocking/Dilution buffer: 5 % NFDM/TBST.

#### Immunofluorescence

Image 3. Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1 % Triton X-100 permeabilized HepG2 (human liver hepatocellular carcinoma cell line) cells labeling MKS1 with (ABIN6244194 and ABIN6578872) at 1/25 dilution, followed by Dylight® 488-conjugated goat anti-rabbit IgG (NK179883) secondary antibody at 1/200 dilution (green). Immunofluorescence image showing cytoplasm staining on HepG2 cell line. Cytoplasmic actin is detected with Dylight® 554 Phalloidin (PD18466410) at 1/100 dilution (red). The nuclear counter stain is DI (blue).

