

Datasheet for ABIN5530641  
**anti-FAM96A antibody (C-Term)**



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3 Images

## Overview

Quantity:	400 µL
Target:	FAM96A
Binding Specificity:	AA 109-138, C-Term
Reactivity:	Human, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This FAM96A antibody is un-conjugated
Application:	Western Blotting (WB), Immunofluorescence (IF), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))

## Product Details

Immunogen:	This FA96A antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 109-138 amino acids from the C-terminal region of human FA96A.
Isotype:	Ig Fraction
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.

## Target Details

Target:	FAM96A
Alternative Name:	FA96A ( <a href="#">FAM96A Products</a> )
Molecular Weight:	18 kDa

## Target Details

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Gene ID: 84191

UniProt: [Q9H5X1](#)

## Application Details

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Application Notes: For WB starting dilution is: 1:1000

For IHC-P starting dilution is: 1:50~100

For IF starting dilution is: 1:10~50

Restrictions: For Research Use only

## Handling

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Format: Liquid

Concentration: 0.5 mg/mL

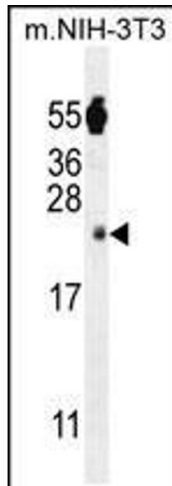
Buffer: Supplied in PBS with 0.09 % (W/V) sodium azide.

Preservative: Sodium azide

Precaution of Use: This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

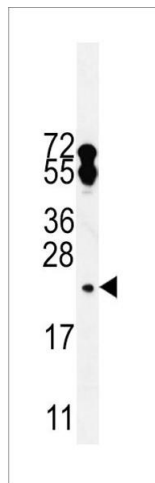
Storage: 4 °C,-20 °C

Storage Comment: Store at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high temperatures.



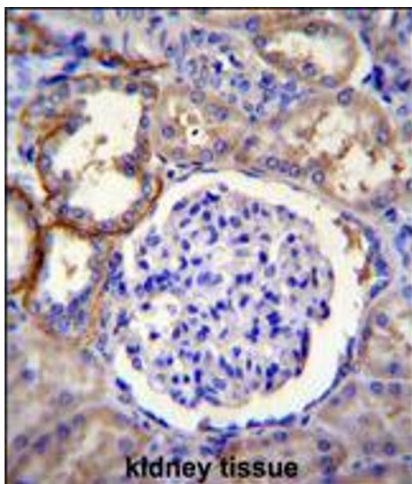
#### Western Blotting

**Image 1.** Western blot analysis in mouse NIH-3T3 cell line lysates (35ug/lane).



#### Western Blotting

**Image 2.** Western blot analysis in WiDr cell line lysates (35ug/lane).



#### Immunohistochemistry

**Image 3.** FA96A Antibody immunohistochemistry analysis in formalin fixed and paraffin embedded human kidney tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.