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**Product Name: WASP/Wiskott-Aldrich syndrome protein Rabbit Monoclonal Antibody**  
**Catalog #: AMRe87336**

For research use only.

## Summary

<b>Description</b>	Recombinant rabbit monoclonal antibody
<b>Host</b>	Rabbit
<b>Application</b>	WB,ICC/IF,FC
<b>Reactivity</b>	Human
<b>Conjugation</b>	Unconjugated
<b>Modification</b>	Unmodified
<b>Isotype</b>	IgG
<b>Clonality</b>	Monoclonal
<b>Form</b>	Liquid
<b>Concentration</b>	
<b>Storage</b>	Aliquot and store at -20°C (valid for 12 months). Avoid freeze/thaw cycles.
<b>Shipping</b>	Ice bags
<b>Buffer</b>	Supplied in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% sodium azide and 0.05% protective protein. Stable for 12 months from date of receipt.
<b>Purification</b>	Affinity Purification

## Application

<b>Dilution Ratio</b>	WB 1:2000-1:20000,ICC/IF 1:20-1:50,FC 1:20-1:50
<b>Molecular Weight</b>	Calculated MW:53 kDa; Observed MW:60 kDa

## Antigen Information

<b>Gene Name</b>	WASP/Wiskott-Aldrich syndrome protein
<b>Alternative Names</b>	THC; IMD2; SCNX; THC1; WASP; WASPA
<b>Gene ID</b>	7454
<b>SwissProt ID</b>	P42768
<b>Immunogen</b>	A synthetic peptide of human WASP/Wiskott-Aldrich syndrome protein

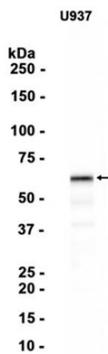
## Background

The Wiskott-Aldrich syndrome (WAS) family of proteins share similar domain structure, and are involved in transduction of signals from receptors on the cell surface to the actin cytoskeleton. The presence of a number of different motifs suggests that

they are regulated by a number of different stimuli, and interact with multiple proteins. Recent studies have demonstrated that these proteins, directly or indirectly, associate with the small GTPase, Cdc42, known to regulate formation of actin filaments, and the cytoskeletal organizing complex, Arp2/3. Wiskott-Aldrich syndrome is a rare, inherited, X-linked, recessive disease characterized by immune dysregulation and microthrombocytopenia, and is caused by mutations in the WAS gene. The WAS gene product is a cytoplasmic protein, expressed exclusively in hematopoietic cells, which show signalling and cytoskeletal abnormalities in WAS patients. A transcript variant arising as a result of alternative promoter usage, and containing a different 5' UTR sequence, has been described, however, its full-length nature is not known. [provided by RefSeq, Jul 2008]

## Research Area

## Image Data



Western blot analysis of extracts from U-937 cells using WASP/Wiskott-Aldrich syndrome protein Rabbit Monoclonal Antibody at 1:1000.