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Research Use Only. Not for diagnostic or therapeutic use.

EB06675 - Goat Anti-Wiskott-Aldrich Syndrome / WASP Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: WASP, Wiskott-Aldrich syndrome (eczema-thrombocytopenia), WAS,

THC, IMD2, thrombocytopenia 1 (X-linked), Wiskott-Aldrich syndrome

(eczema-thrombocytopenia) protein, THC1, Wiskott-Aldrich syndrome protein

Official Symbol: WAS

Accession Number(s): NP_000368.1

Human GeneID(s): 7454

Important Comments: No cross-reactivity expected with N WASP (WASL).

Immunogen

Peptide with sequence C-SPADKKRSGKKKI, from the internal region of the protein sequence according to NP_000368.1.

Please note the peptide is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:128000.

IHC: Paraffin embedded Human Spleen. Recommended concentration: 5µg/ml.

Immunofluorescence: Strong expression of the protein seen in the nuclei and cytoplasm

of HepG2 and U2OS cells. Recommended concentration: 10µg/ml.

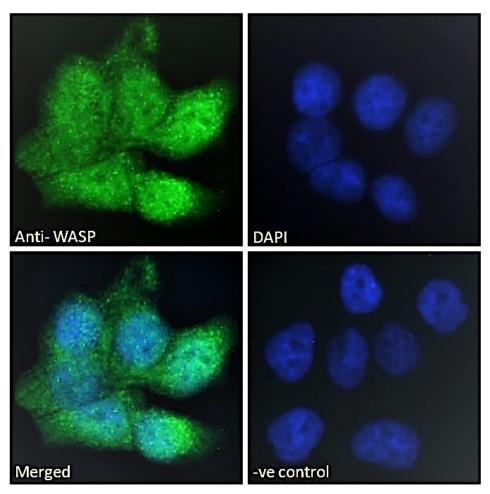
 $\textbf{Flow Cytometry:} \ \ \textbf{Flow cytometric analysis of HepG2 cells.} \ \ \textbf{Recommended concentration:}$

10ug/ml.

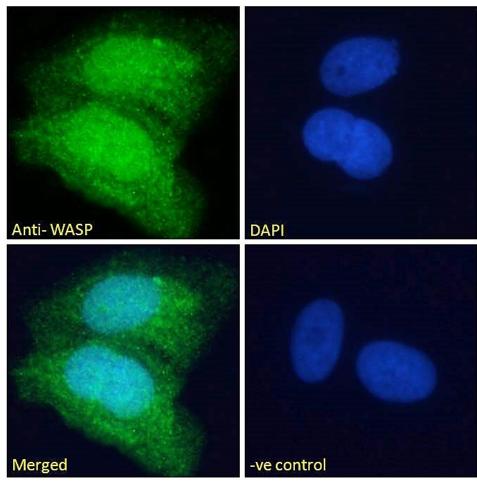
Species Reactivity

Tested: Human

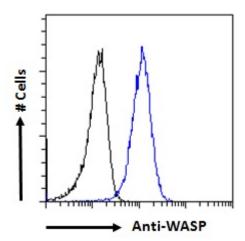
Expected from sequence similarity: Human, Mouse, Rat, Dog



EB06675 Immunofluorescence analysis of paraformaldehyde fixed HepG2 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

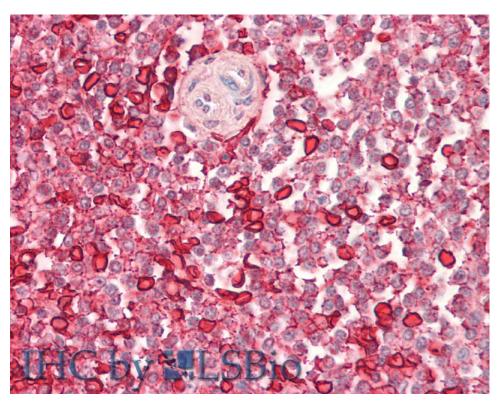


EB06675 Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



EB06675 Flow cytometric analysis of paraformaldehyde fixed HepG2 cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control:

Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.



EB06675 (5μg/ml) staining of paraffin embedded Human Spleen. Steamed antigen retrieval with citrate buffer pH 6, AP-staining.