

**Mouse Monoclonal Antibody to YWHAB**  
**Purified Mouse Monoclonal Antibody**  
**Catalog # AO2325a****Specification**

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**Mouse Monoclonal Antibody to YWHAB - Product Information**

Application	E, WB, FC, ICC
Primary Accession	<a href="#">P31946</a>
Reactivity	Human, Mouse, Rat
Host	Mouse
Clonality	Monoclonal
Isotype	Mouse IgG2b
Calculated MW	28kDa KDa

**Description**

This gene encodes a protein belonging to the 14-3-3 family of proteins, members of which mediate signal transduction by binding to phosphoserine-containing proteins. This highly conserved protein family is found in both plants and mammals. The encoded protein has been shown to interact with RAF1 and CDC25 phosphatases, suggesting that it may play a role in linking mitogenic signaling and the cell cycle machinery. Two transcript variants, which encode the same protein, have been identified for this gene.;

**Immunogen**

Purified recombinant fragment of human YWHAB (AA: 1-246) expressed in E. Coli.

**Formulation**

Purified antibody in PBS with 0.05% sodium azide

**Application Note**

ELISA: 1/10000; WB: 1/500 - 1/2000; IHC: 1/200 - 1/1000; ICC: 1/200 - 1/1000; FCM: 1/200 - 1/400

**Mouse Monoclonal Antibody to YWHAB - Additional Information**

**Gene ID** 7529

**Other Names**

HS1; GW128; YWHAA; KCIP-1; HEL-S-1

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

Mouse Monoclonal Antibody to YWHAB is for research use only and not for use in diagnostic or therapeutic procedures.

**Mouse Monoclonal Antibody to YWHAB - Protein Information**

**Name** YWHAB

**Function**

Adapter protein implicated in the regulation of a large spectrum of both general and specialized signaling pathways. Binds to a large number of partners, usually by recognition of a phosphoserine or phosphothreonine motif. Binding generally results in the modulation of the activity of the binding partner. Negative regulator of osteogenesis. Blocks the nuclear translocation of the phosphorylated form (by AKT1) of SRPK2 and antagonizes its stimulatory effect on cyclin D1 expression resulting in blockage of neuronal apoptosis elicited by SRPK2. Negative regulator of signaling cascades that mediate activation of MAP kinases via AKAP13.

**Cellular Location**

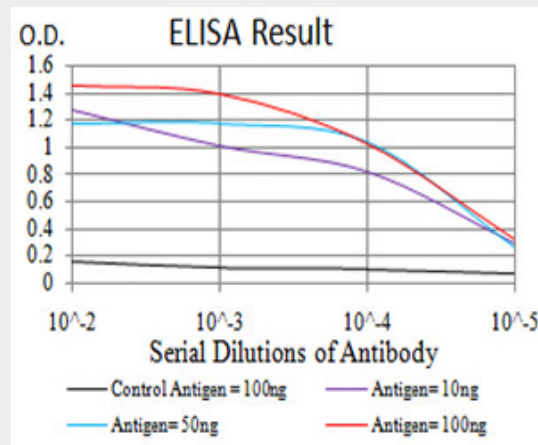
Cytoplasm. Melanosome. Note=Identified by mass spectrometry in melanosome fractions from stage I to stage IV

**Mouse Monoclonal Antibody to YWHAB - Protocols**

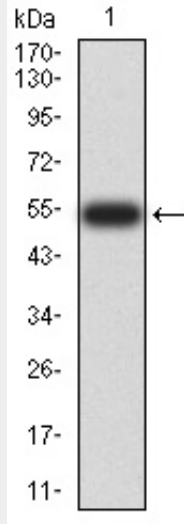
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

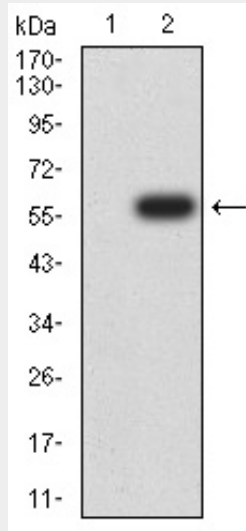
**Mouse Monoclonal Antibody to YWHAB - Images**



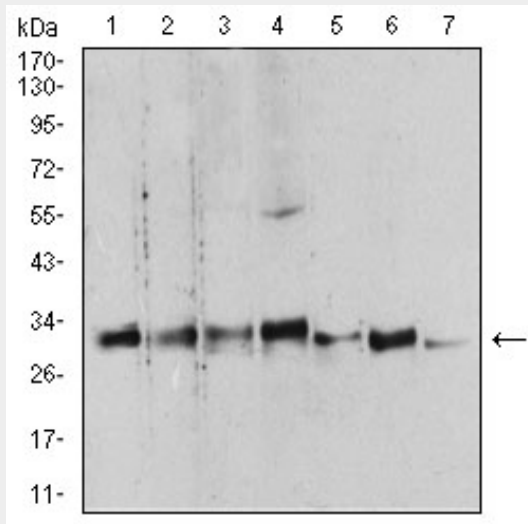
Black line: Control Antigen (100 ng);Purple line: Antigen (10ng); Blue line: Antigen (50 ng); Red line:Antigen (100 ng)



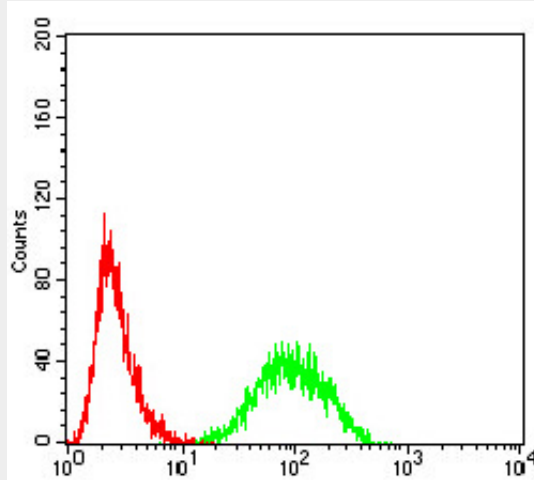
Western blot analysis using YWHAB mAb against human YWHAB (AA: 1-246) recombinant protein. (Expected MW is 54 kDa)



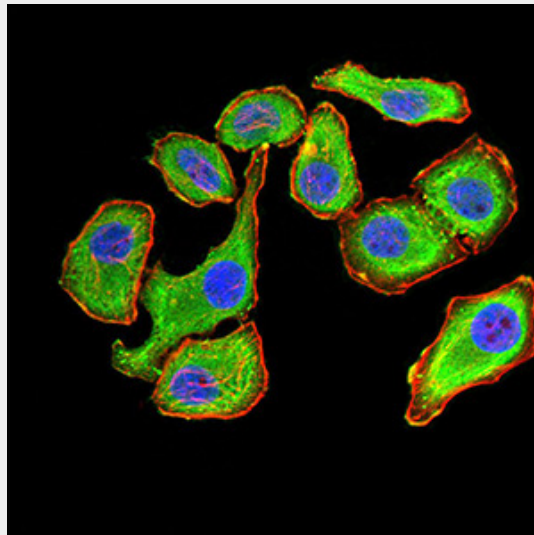
Western blot analysis using YWHAB mAb against HEK293 (1) and YWHAB (AA: 1-246)-hIgGFc transfected HEK293 (2) cell lysate.



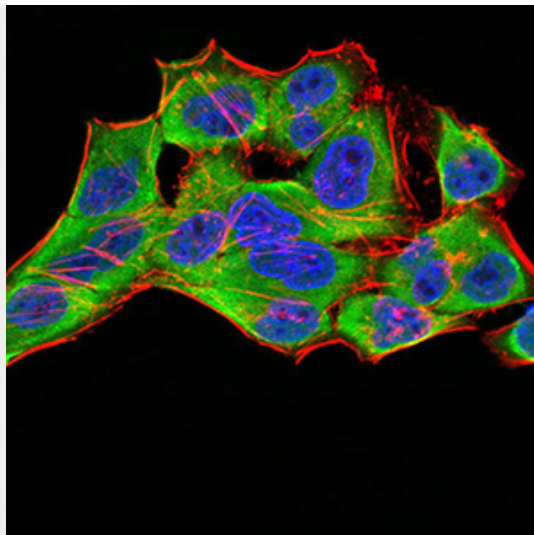
Western blot analysis using YWHAB mouse mAb against Hela (1), NIH/3T3 (2), C6 (3), A431 (4), K562 (5), PC-12 (6), and U937 (7) cell lysate.



Flow cytometric analysis of Hela cells using YWHAB mouse mAb (green) and negative control (red).



Immunofluorescence analysis of GC-7901 cells using YWHAB mouse mAb (green). Blue: DRAQ5 fluorescent DNA dye. Red: Actin filaments have been labeled with Alexa Fluor- 555 phalloidin. Secondary antibody from Fisher



Immunofluorescence analysis of HeLa cells using YWHAB mouse mAb (green). Blue: DRAQ5 fluorescent DNA dye. Red: Actin filaments have been labeled with Alexa Fluor- 555 phalloidin. Secondary antibody from Fisher

#### **Mouse Monoclonal Antibody to YWHAB - References**

1.BMC Res Notes. 2014 Feb 20;7:97. ; 2.Mol Biol Rep. 2012 Dec;39(12):10647-53.;