

## Basic Information

<b>Product Name</b>	Anti-ABI2 Antibody		
<b>Gene Name</b>	ABI2		
<b>Source</b>	Rabbit		
<b>Clonality</b>	Polyclonal		
<b>Isotype</b>	IgG		
<b>Species Reactivity</b>	human, mouse, rat		
<b>Tested Application</b>	WB, ICC/IF, FCM		
<b>Contents</b>	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.		
<b>Immunogen</b>	A synthetic peptide corresponding to a sequence at the N-terminus of human ABI2, identical to the related mouse and rat sequences.		
<b>Concentration</b>	500 ug/ml		
<b>Purification</b>	Immunogen affinity purified.		
<b>Observed MW</b>	56-70 kDa		
<b>Dilution Ratios</b>	Western blot (WB):	1:500-2000	
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400	
	Flow Cytometry (Fixed):	1:50-200	

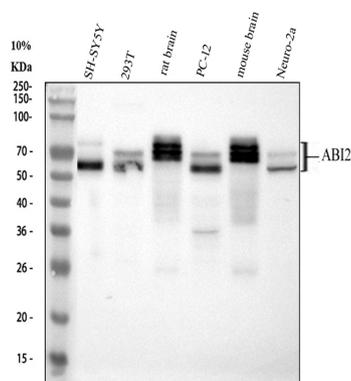
## Storage

12 months from date of receipt, -20°C as supplied.

## Background Information

ABI2 (ABL Interactor 2), is a protein that in humans is encoded by the ABI2 gene. By analysis of a YAC and a BAC, Machado et al. (2000) mapped the ABI2 gene to 2q31-q33. ABI2 possesses a basic N terminus with homology to a homeodomain protein; a central serine-rich region; 3 PEST sequences, which are implicated in susceptibility to protein degradation; several proline-rich stretches; and an acidic C terminus with multiple phosphorylation sites and an SH3 domain. Dai and Pendergast (1995) suggested that the ABI proteins may function to coordinate the cytoplasmic and nuclear functions of the ABL1 tyrosine kinase.

## Selected Validation Data



Western blot analysis of ABI2 using anti-ABI2 antibody (BA3031-2).

The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human SH-SY5Y whole cell lysates,

Lane 2: human 293T whole cell lysates,

Lane 3: rat brain tissue lysates,

Lane 4: rat PC-12 whole cell lysates,

Lane 5: mouse brain tissue lysates,

Lane 6: mouse Neuro-2a whole cell lysates.

After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-ABI2

antigA03957-Aen affinity purified polyclonal antibody (BA3031-2) at

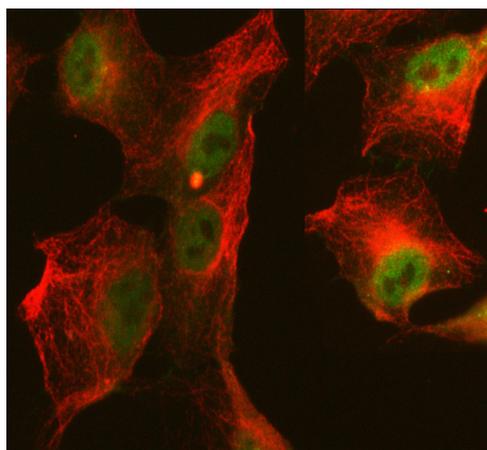
a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP

secondary antibody (Catalog # BA1054). The signal is developed

using ECL Plus Western Blotting Substrate (Catalog # AR1197). A

specific band was detected for ABI2 at approximately 56-70 kDa. The

expected band size for ABI2 is at 56 kDa.



ICC/IF analysis of ABI2 using anti-ABI2 antibody (BA3031-2) and anti-Alpha Tubulin antibody (M03989-3).

ABI2 was detected in an immunocytochemical section of A549 cells.

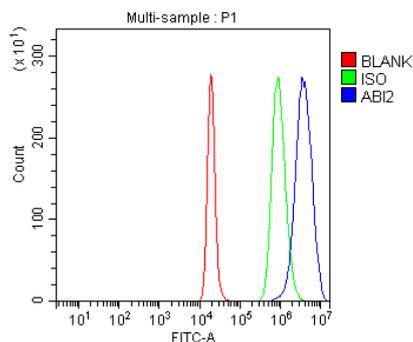
The section was incubated with rabbit anti-ABI2 Antibody (BA3031-2)

at a dilution of 1:100. Fluoro488-conjugated Anti-rabbit IgG

Secondary Antibody (green) (Catalog # BA1127) and Cy3-conjugated

Anti-mouse IgG Secondary Antibody (red) (Catalog # BA1031) were

used as secondary antibody.



Flow Cytometry analysis of SH-SY5Y cells using anti-ABI2 antibody (BA3031-2).

Overlay histogram showing SH-SY5Y cells stained with BA3031-2 (Blue line). To facilitate intrMyelin basic protein/MBPllular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-ABI2 Antibody (BA3031-2) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.