

Basic Information

Product Name	Anti-APAF1 Antibody	
Gene Name	APAF1	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human	
Tested Application	WB, FCM, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	E.coli-derived human APAF1 recombinant protein (Position: M1-E1248).	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	142 kDa	
Dilution Ratios	Western blot (WB):	1:500-2000
	Flow Cytometry (Fixed):	1:50-200
	Enzyme linked immunosorbent assay (ELISA):	1:100-1000

Storage

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

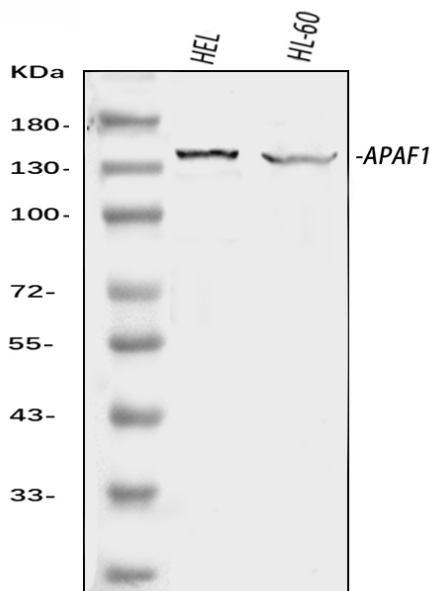
Background Information

Apoptotic peptidase activating factor 1, also known as APAF1, is a protein which in humans is encoded by the APAF1 gene. This gene is mapped to chromosome 12q23. It encodes a cytoplasmic protein that initiates apoptosis. And it is an essential downstream effector of p53-mediated apoptosis. This protein contains several copies of the WD40 repeat domain, a caspase recruitment domain(CARD), and an ATPase domain(NB-ARC). In the presence of cytochrome c and dATP, APAF1 assembles into an oligomeric apoptosome, which is responsible for activation of procaspase-9 and maintenance of the enzymatic activity of processed caspase-9. Furthermore, APAF1 is inactivated in metastatic melanomas, leading to defects in the execution of apoptotic cell death. Additionally, APAF1 has been shown to interact with NLRP1, Caspase-9, APIP, BCL2-like 1 and HSPA4.

Reference

Anti-APAF1 Antibody被引用在1文献中。

Selected Validation Data

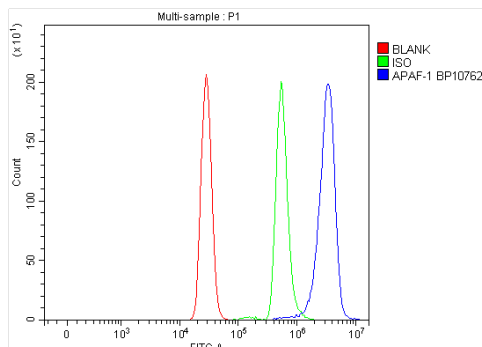


Western blot analysis of APAF1 using anti-APAF1 antibody (A00889-2). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: HEL whole cell lysates,

Lane 2: HL-60 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-APAF1 antigen affinity purified polyclonal antibody (A00889-2) 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 APAF1 at approximately 142 kDa. The expected band size for APAF1 is at 142 kDa.



Flow Cytometry analysis of U937 cells using anti-APAF1 antibody (A00889-2).

Overlay histogram showing U937 cells stained with A00889-2 (Blue line).

To facilitate intracellular 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-APAF1 Antibody (A00889-2) at 1:100 dilution for 30 min at

20°C. DyLight®488 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.