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# Datasheet for ABIN5647000 anti-ACP1 antibody (AA 33-61)

3 Images



### Overview

Quantity:	0.08 mL
Target:	ACP1
Binding Specificity:	AA 33-61
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Application:	Western Blotting (WB), Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF), Flow Cytometry (FACS)

## Product Details

Immunogen:	A portion of amino acids 33-61 from the human protein was used as the immunogen for this ACP1 antibody.
Isotype:	Ig Fraction
Purification:	Antigen affinity purified

# Target Details

Target:	ACP1
Alternative Name:	Acid Phosphatase / ACP1 (ACP1 Products)
Background:	ACP1 belongs to the phosphotyrosine protein phosphatase family of proteins. It functions as an acid phosphatase and a protein tyrosine phosphatase by hydrolyzing protein tyrosine
	phosphate to protein tyrosine and orthophosphate. This enzyme also hydrolyzes

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Target Details	
	orthophosphoric monoesters to alcohol and orthophosphate. This gene is genetically polymorphic, and three common alleles segregating at the corresponding locus give rise to six phenotypes. Each allele appears to encode at least two electrophoretically different isozymes, Bf and Bs, which are produced in allele-specific ratios.
UniProt:	P24666
Application Details	
Application Notes:	Western blot: 1:1000,IHC (Paraffin): 1:50-1:100,Flow Cytometry: 1:10-1:50,Immunofluorescence: 1:10-1:50
Restrictions:	For Research Use only
Handling	
Buffer:	In 1X PBS, pH 7.4, with 0.09 % sodium azide
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C

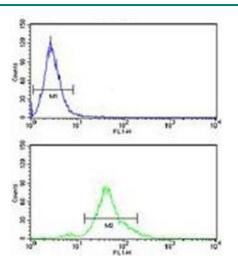
Storage Comment: Aliquot the ACP1 antibody and store frozen at -20°C or colder. Avoid repeated freeze-thaw cycles.

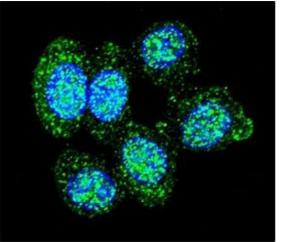
#### Images

#### Western Blotting

**Image 1.** Western blot analysis of ACP1 antibody and human HeLa lysate. Predicted molecular weight 14-18 kDa (isoforms 1-3).

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#### **Flow Cytometry**

**Image 2.** ACP1 antibody flow cytometry analysis of human HeLa cells (bottom histogram) compared to a negative control (top histogram). FITC-conjugated goat-anti-rabbit secondary Ab was used for the analysis.

#### Immunofluorescence

**Image 3.** Confocal immunofluorescent analysis of ACP1 antibody with human HeLa cells followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green). DAPI was used as a nuclear counterstain (blue).

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