

AP31973PU-N**Polyclonal Antibody to APOBEC3A - Aff - Purified****Alternate names:**

Apolipoprotein B mRNA editing enzyme, Phorbol-1, catalytic polypeptide-like 3A

Quantity:

0.1 mg

Concentration:

0.5 mg/ml

Background:

PHO1 a member of the cytidine deaminase gene family. The PHO1 gene is one of seven related genes or pseudogenes found in a cluster, thought to result from gene duplication, on chromosome 22. Members of the cluster encode proteins that are structurally and functionally related to the C to U RNA-editing cytidine deaminase APOBEC1. It is thought that the proteins may be RNA editing enzymes and have roles in growth or cell cycle control. This gene encodes a protein that lacks the zinc binding activity and may be an expressed pseudogene.

Uniprot ID:[P31941](#)**NCBI:**[NP_001180218.1](#)**GeneID:**[100913187](#)**Host:**

Goat

Immunogen:

Synthetic peptide with sequence from the internal region of Human APOBEC3A (NP_663745.1).

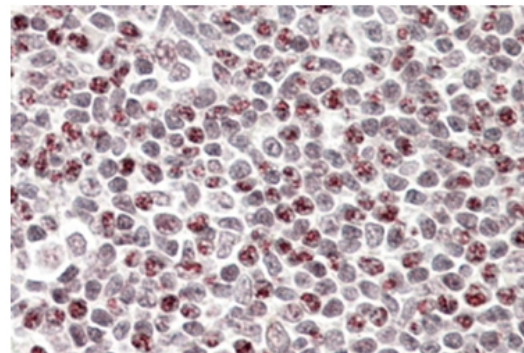
Genename: APOBEC3A**AA Sequence:**

C-TSNFNNGIGRHKTY

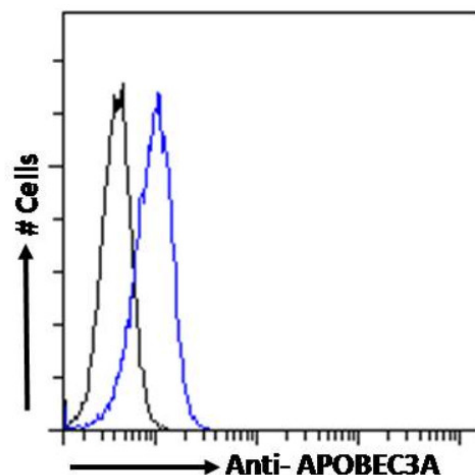
Format:**State:** Liquid purified Ig fraction**Purification:** Immunoaffinity Chromatography**Buffer System:** Tris saline, pH~7.3**Preservatives:** 0.02% Sodium Azide**Stabilizers:** 0.5% BSA**Applications:****Peptide ELISA:** 1/8000 (Detection limit).**Western blot:** 3-6 µg/ml. Approx. 26-28kDa band observed in Human Spleen, Tonsil and Peripheral Blood Monocytes (PBM) lysates (calculated MW of 23.0kDa according to Human NP_663745.1). The observed molecular weight corresponds to earlier findings with different antibodies from other commercial sources. Recommended concentration: 0.1-1µg/ml. Primary incubation 1 hour at room temperature.**Immunohistochemistry on Paraffin Sections:** 3-6 µg/ml. In paraffin embedded Human Tonsil shows staining of distinct parts of the nucleus in lymphoid cells.**Immunofluorescence:** 10 µg/ml. Strong expression of the protein seen in the nucleus of A431 and A549 cells. **Flow Cytometry:** 10 µg/ml. Flow cytometric analysis of Jurkat cells.

Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.

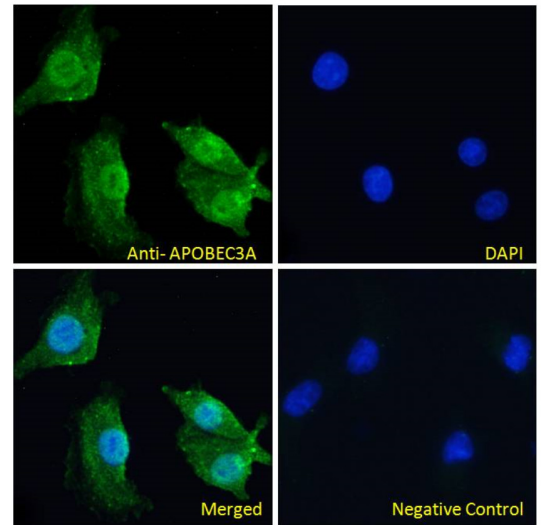
Specificity:	This antibody recognizes Human Phorbolin 1 / APOBEC3A. Other species not tested.
Storage:	Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing. Shelf life: one year from despatch.
Pictures:	Paraffin Embedded Human Tonsil stained with APOBEC3A antibody Cat.-No AP31973PU-N at 3.8 µg/ml. Steamed antigen retrieval with citrate buffer pH 6, AP-staining.



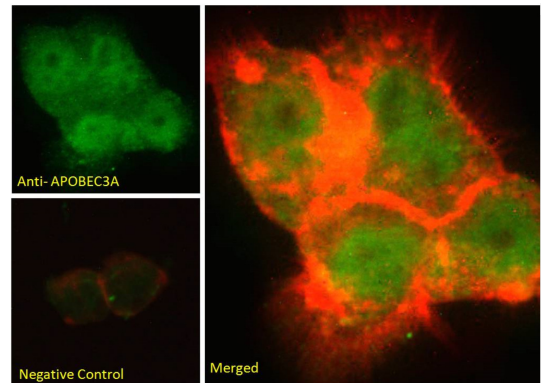
AP31973PU-N Flow cytometric analysis of paraformaldehyde fixed Jurkat cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody



AP31973PU-N Immunofluorescence analysis of paraformaldehyde fixed A549 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing nuclear and some cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



AP31973PU-N Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing nuclear staining. Actin filaments were stained with phalloidin (red). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



AP31973PU-N (1µg/ml) staining of Human Spleen (A), Tonsil (B) and (0.1ug/ml) PBM (C) lysate (35µg protein in RIPA buffer). Detected by chemiluminescence.

