

# Product datasheet for AP11346PU-N

# APOBEC3A\_B (N-term) Rabbit Polyclonal Antibody

### **Product data:**

#### OriGene Technologies, Inc.

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Product Type:	Primary Antibodies	
Applications:	IHC, WB	
Recommended Dilution:	Peptide ELISA: 1/1,000. Western blot: 1/100-1/500. Immunohistochemistry: 1/50-1/100.	
Reactivity:	Human	
Host:	Rabbit	
lsotype:	lg	
Clonality:	Polyclonal	
Immunogen:	KLH conjugated synthetic peptide between 1-30 amino acids from the N-terminal region of human PHO1.	
Specificity:	This antibody is specific to APOBEC3A/PHO1 (N-term).	
Formulation:	PBS containing 0.09% (W/V) Sodium Azide as preservative. State: Purified State: Liquid purified Ig fraction.	
Concentration:	lot specific	
Purification:	This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.	
Conjugation:	Unconjugated	
Storage:	Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing.	
Stability:	Shelf life: one year from despatch.	
Gene Name:	APOBEC3A and APOBEC3B deletion hybrid	
Database Link:	<u>Entrez Gene 100913187 Human</u> <u>P31941</u>	

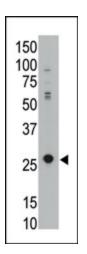


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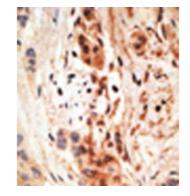
CRIGENE APOBEC3A_B (N-term) Rabbit Polyclonal Antibody – AP11346PU-N	
Background:	PHO1 a member of the cytidine deaminase gene family. The PHO1 gene is one of seven related genes or pseudogenes found in a cluster, thought to result from gene duplication, on chromosome 22. Members of the cluster encode 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.
Synonyms:	Phorbolin-1

Synonyms.	
Note:	Calculated MW: 23012 Da

# **Product images:**



Western blot analysis of anti-PHO1 (N-term) Pab to detect PHO1 in jurkat cell lysate.



Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by AEC staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.

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