

# Datasheet for ABIN7181545 anti-HIST1H3A antibody (AA 37-48)

# 2 Images



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Quantity:	100 μL		
Target:	HIST1H3A		
Binding Specificity:	AA 37-48		
Reactivity:	Human		
Host:	Rabbit		
Clonality:	Polyclonal		
Conjugate:	This HIST1H3A antibody is un-conjugated		
Application:	ELISA, Immunohistochemistry (IHC)		
Product Details			
Immunogen:	Synthesized peptide derived from Human Histone H3.1 protein (37-48aa)		
Isotype:	IgG		
Cross-Reactivity:	Human		
Purification:	Antigen Affinity Purified		
Target Details			
Target:	HIST1H3A		
Alternative Name:	HIST1H3A (HIST1H3A Products)		
Background:	Background: Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a		

## **Target Details**

template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling. Aliases: Histone H3.1 (Histone H3/a) (Histone H3/b) (Histone H3/c) (Histone H3/d) (Histone H3/f) (Histone H3/h) (Histone H3/j) (Histone H3/k) (Histone H3/l), HIST1H3A, HIST1H3B, HIST1H3C, HIST1H3D, HIST1H3E, HIST1H3F, HIST1H3G, HIST1H3H, HIST1H3I, HIST1H3J, H3FA, H3FL, H3FC, H3FB, H3FD, H3FI, H3FH, H3FK, H3FF, H3FJ

UniProt:

Preservative:

P68431

# **Application Details**

Application Notes:	Recommended dilution: IHC:1:10-1:100,
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300

Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4

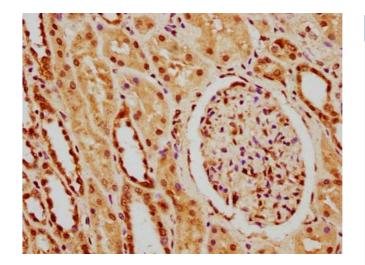
ProClin

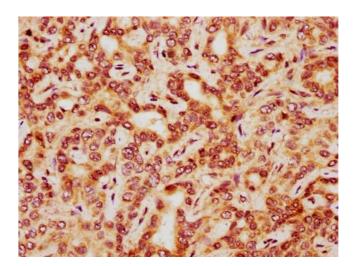
Precaution of Use:

This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Storage: -20 °C,-80 °C

Storage Comment: Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.





## **Immunohistochemistry**

**Image 1.** IHC image of nphHU diluted at 1:20 and staining in paraffin-embedded human kidney tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10 % normal goat serum 30 min at RT. Then primary antibody (1 % BSA) was incubated at 4 °C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

#### **Immunohistochemistry**

**Image 2.** IHC image of nphHU diluted at 1:20 and staining in paraffin-embedded human liver cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10 % normal goat serum 30 min at RT. Then primary antibody (1 % BSA) was incubated at 4 °C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.