

Datasheet for ABIN7168107
anti-RNASEH2A antibody (AA 201-299)[Go to Product page](#)

1 Image

Overview

Quantity:	100 µg
Target:	RNASEH2A
Binding Specificity:	AA 201-299
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This RNASEH2A antibody is un-conjugated
Application:	Immunohistochemistry (IHC), ELISA

Product Details

Immunogen:	Recombinant Human Ribonuclease H2 subunit A protein (201-299AA)
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

Target:	RNASEH2A
Alternative Name:	RNASEH2A (RNASEH2A Products)
Background:	Background: Catalytic subunit of RNase HII, an endonuclease that specifically degrades the RNA of RNA:DNA hybrids. Participates in DNA replication, possibly by mediating the removal of

Target Details

lagging-strand Okazaki fragment RNA primers during DNA replication. Mediates the excision of single ribonucleotides from DNA:RNA duplexes.

Aliases: RNASEH2A antibody, RNASEH1 antibody, RNH1A antibody, Ribonuclease H2 subunit A antibody, RNase H2 subunit A antibody, EC 3.1.26.4 antibody, Aicardi-Goutieres syndrome 4 protein antibody, AGS4 antibody, RNase H(35) antibody, Ribonuclease H1 large subunit antibody, RNase H1 large subunit antibody, Ribonuclease H1 subunit A antibody

UniProt: [O75792](#)

Application Details

Application Notes: Recommended dilution: IHC:1:200-1:500,

Restrictions: For Research Use only

Handling

Format: Liquid

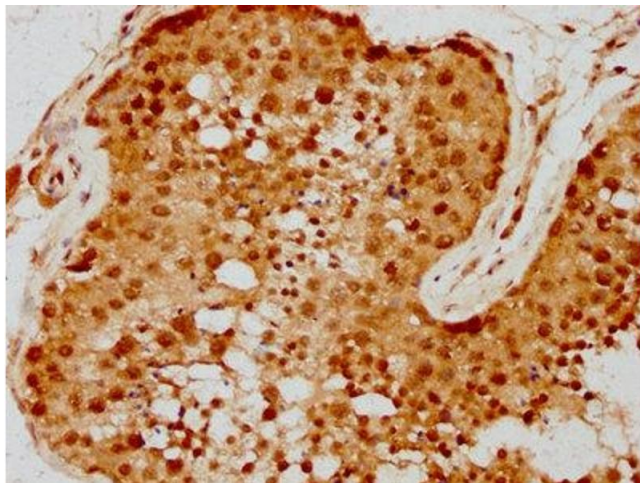
Buffer: Preservative: 0.03 % Proclin 300
Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4

Preservative: 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 ABIN7168107 diluted at 1:300 and staining in paraffin-embedded human testis tissue performed on a Leica Bond™ 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.