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anti-POLR1C antibody (Internal Region)



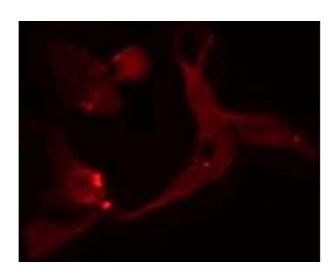
Image



Overview $100 \, \mu L$ Quantity: Target: POLR1C Binding Specificity: Internal Region Reactivity: Human, Mouse Host: Rabbit Clonality: Polyclonal Conjugate: This POLR1C antibody is un-conjugated Application: Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC) **Product Details** Immunogen: A synthesized peptide derived from human POLR1C, corresponding to a region within the internal amino acids. Isotype: IgG POLR1C Antibody detects endogenous levels of total POLR1C. Specificity: Predicted Reactivity: Pig, Bovine, Horse, Sheep, Rabbit, Dog The antiserum was purified by peptide affinity chromatography using SulfoLinkTM Coupling Purification: Resin (Thermo Fisher Scientific). **Target Details** POLR1C Target:

Target Details

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Alternative Name:	POLR1C (POLR1C Products)
Background:	Description: DNA-dependent RNA polymerase catalyzes the transcription of DNA into RNA using the four ribonucleoside triphosphates as substrates. Common component of RNA polymerases I and III which synthesize ribosomal RNA precursors and small RNAs, such as 5S rRNA and tRNAs, respectively. RPAC1 is part of the Pol core element with the central large cleft and probably a clamp element that moves to open and close the cleft (By similarity). Gene: POLR1C
Molecular Weight:	39 kDa
Gene ID:	9533
UniProt:	015160
Application Details	
Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 $\%$ sodium azide and 50 $\%$ glycerol.
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:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



Immunofluorescence (fixed cells)

Image 1. ABIN6275077 staining A549 cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod