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# Datasheet for ABIN7150227 anti-POLD1 antibody (Catalytic Subunit)





## Overview

Quantity:	100 µg
Target:	POLD1
Binding Specificity:	AA 1-123, Catalytic Subunit
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This POLD1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunoprecipitation (IP)

## Product Details

Immunogen:	Recombinant Human DNA polymerase delta catalytic subunit protein (1-123AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

## Target Details

Target:	POLD1
Alternative Name:	POLD1 (POLD1 Products)
Background:	Background: As the catalytic component of the trimeric (Pol-delta3 complex) and tetrameric
	DNA polymerase delta complexes (Pol-delta4 complex), plays a crucial role in high fidelity

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genome replication, including in lagging strand synthesis, and repair. Exh	nibits both DNA
polymerase and $3$ \\\'- to $5$ \\\'-exonuclease activities (PubMed:16510448)	, PubMed:19074196,
PubMed:20334433, PubMed:24035200, PubMed:24022480). Requires th	e presence of
accessory proteins POLD2, POLD3 and POLD4 for full activity. Depending	g upon the absence
(Pol-delta3) or the presence of POLD4 (Pol-delta4), displays differences i	n catalytic activity.
Most notably, expresses higher proofreading activity in the context of Po	l-delta3 compared with
that of Pol-delta4 (PubMed:19074196, PubMed:20334433). Although bot	h Pol-delta3 and Pol-
delta4 process Okazaki fragments in vitro, Pol-delta3 may be better suite	ed to fulfill this task,
exhibiting near-absence of strand displacement activity compared to Po	l-delta4 and stalling on
encounter with the 5\\\'-blocking oligonucleotides. Pol-delta3 idling proc	ess may avoid the
formation of a gap, while maintaining a nick that can be readily ligated (F	oubMed:24035200).
Along with DNA polymerase kappa, DNA polymerase delta carries out ap	proximately half of
nucleotide excision repair (NER) synthesis following UV irradiation (PubN	/led:20227374). Under
conditions of DNA replication stress, in the presence of POLD3 and POLI	04, may catalyze the
repair of broken replication forks through break-induced replication (BIR)	(PubMed:24310611).
Involved in the translesion synthesis (TLS) of templates carrying 06-met	hylguanine or abasic
sites (PubMed:19074196).	

Aliases: Polymerase (DNA directed) delta 1 catalytic subunit antibody, CDC2 antibody, CDC2 homolog antibody, CRCS10 antibody, DNA directed DNA polymerase delta 1 antibody, DNA directed polymerase delta 1 antibody, DNA pol delta 1 antibody, DNA polymerase delta catalytic subunit antibody, DNA polymerase subunit delta p125 antibody, DPOD1\_HUMAN antibody, MDPL antibody, POLD antibody, POLD 1 antibody, POLD1 antibody, Polymerase (DNA directed) delta 1 catalytic subunit 125 kDa antibody, Polymerase (DNA) delta 1 catalytic subunit antibody, Polymerase DNA directed delta 1 catalytic subunit 125kD antibody, polymerase, DNA, delta antibody

UniProt:	P28340
Pathways:	Telomere Maintenance, DNA Damage Repair, DNA Replication, Chromatin Binding, Synthesis of
	DNA

## **Application Details**

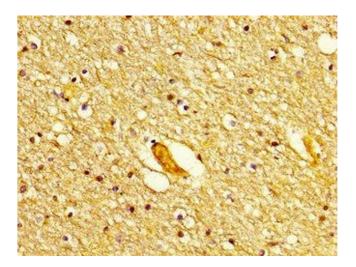
Application Notes:	Recommended dilution: WB:1:500-1:5000, IHC:1:200-1:500, IP:1:200-1:2000,
Restrictions:	For Research Use only

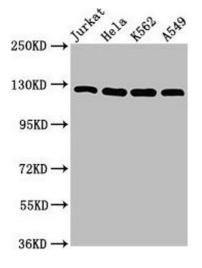
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## Handling

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.

### Images





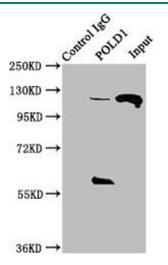
#### Immunohistochemistry

**Image 1.** IHC image of ABIN7150227 diluted at 1:300 and staining in paraffin-embedded human brain 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 30min 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.

#### Western Blotting

**Image 2.** Western Blot Positive WB detected in: Jurkat whole cell lysate, Hela whole cell lysate, K562 whole cell lysate, A549 whole cell lysate All lanes: POLD1 antibody at 5 µg/mL Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 124 kDa Observed band size: 124 kDa

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#### Western Blotting

**Image 3.** Immunoprecipitating POLD1 in Hela whole cell Iysate Lane 1: Rabbit control IgG instead of ABIN7150227 in Hela whole cell Iysate. For western blotting, a HRPconjugated Protein G antibody was used as the secondary antibody (1/2000) Lane 2: ABIN7150227 (8 µg) + Hela whole cell Iysate (500 µg) Lane 3: Hela whole cell Iysate (10 µg)

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