

Datasheet for ABIN7601923

anti-CDT1 antibody (AA 508-546)



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Quantity:	100 μg	
Target:	CDT1	
Binding Specificity:	AA 508-546	
Reactivity:	Human, Mouse, Rat	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This CDT1 antibody is un-conjugated	
Application:	ELISA, Western Blotting (WB), Immunofluorescence (IF), Immunocytochemistry (ICC), Flow Cytometry (FACS)	

Product Details

Purpose:	Anti-CDT1/DUP Antibody Picoband®	
Immunogen:	E.coli-derived human CDT1/DUP recombinant protein (Position: D508-L546).	
Isotype:	IgG	
Cross-Reactivity (Details):	No cross-reactivity with other proteins.	
Characteristics:	Anti-CDT1/DUP Antibody Picoband® (ABIN7601923). Tested in ELISA, Flow Cytometry, IF, ICC, WB applications. This antibody reacts with Human, Mouse, Rat. The brand Picoband indicates this is a premium antibody that guarantees superior quality, high affinity, and strong signals with minimal background in Western blot applications. Only our best-performing antibodies are designated as Picoband, ensuring unmatched performance.	
Purification:	Immunogen affinity purified.	

Target Details

Target:	CDT1
Alternative Name:	CDT1 (CDT1 Products)
Background:	Synonyms: Lumican, Keratan sulfate proteoglycan lumican, KSPG lumican, LUM, LDC, SLRR2D
	Tissue Specificity: Cornea and other tissues.
	Background: CDT1 (Chromatin licensing and DNA replication factor 1) is a protein that in
	humans is encoded by the CDT1 gene. The protein encoded by this gene is involved in the
	formation of the pre-replication complex that is necessary for DNA replication. The encoded
	protein can bind geminin, which prevents replication and may function to prevent this protein
	from initiating replication at inappropriate origins. Phosphorylation of this protein by cyclin A-
	dependent kinases results in degradation of the protein.
Molecular Weight:	60 kDa
Gene ID:	81620
UniProt:	Q9H211
Pathways:	MAPK Signaling, Mitotic G1-G1/S Phases, DNA Replication, Synthesis of DNA
Application Details	
Application Notes:	Western blot, 0.25-0.5 μg/mL, Human, Mouse, Rat
	Immunocytochemistry/Immunofluorescence, 5 μg/mL, Human
	Flow Cytometry (Fixed), 1-3 µg/1x10 ⁶ cells, Human
	ELISA, 0.1-0.5 μg/mL, -
	1. Bicknell, L. S., Bongers, E. M. H. F., Leitch, A., Brown, S., Schoots, J., Harley, M. E., Aftimos, S.,
	Al-Aama, J. Y., Bober, M., Brown, P. A. J., van Bokhoven, H., Dean, J., and 15 others. Mutations in
	the pre-replication complex cause Meier-Gorlin syndrome. Nature Genet. 43: 356-359, 2011. 2.
	Bongers, E. M. H. F., Opitz, J. M., Fryer, A., Sarda, P., Hennekam, R. C. M., Hall, B. D., Superneau,
	D. W., Harbison, M., Poss, A., van Bokhoven, H., Hamel, B. C. J., Knoers, N. V. A. M. Meier-Gorlin
	syndrome: report of eight additional cases and review. Am. J. Med. Genet. 102: 115-124, 2001.
	3. Feingold, M. Meier-Gorlin syndrome. (Letter) Am. J. Med. Genet. 109: 338 only, 2002.
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Adding 0.2 mL of distilled water will yield a concentration of 500 µg/mL.

Handling

Concentration:	500 μg/mL
Buffer:	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na2HPO4.
Storage:	4 °C,-20 °C
Storage Comment:	At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and thawing.