

## Anti-ARD1 Antibody

Catalog #	Source	Reactivity	Applications
CQA3253	Rabbit	H, M, R	WB, IH, IF/IC
<b>Description</b>	Rabbit polyclonal antibody to ARD1		
<b>Immunogen</b>	Recombinant full length protein of human ARD1		
<b>Purification</b>	The antibody was purified by immunogen affinity chromatography.		
<b>Specificity</b>	Recognizes endogenous levels of ARD1 protein.		
<b>Clonality</b>	Polyclonal		
<b>Conjugation</b>			
<b>Form</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
<b>Dilution</b>	WB (1/500 - 1/2000), IH (1/50 - 1/200), IF/IC (1/50 - 1/200)		
<b>Gene Symbol</b>	NAA10		
<b>Alternative Names</b>	ARD1; ARD1A; TE2; N-alpha-acetyltransferase 10; N-terminal acetyltransferase complex ARD1 subunit homolog A; NatA catalytic subunit Naa10		
<b>Entrez Gene</b>	8260 (Human); 56292 (Mouse)		
<b>SwissProt</b>	P41227 (Human); Q9QY36 (Mouse)		
<b>Storage/Stability</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.		

**Application key:** E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, CHIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

**Species reactivity key:** H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb- Rabbit, S- Sheep, Z- Zebrafish

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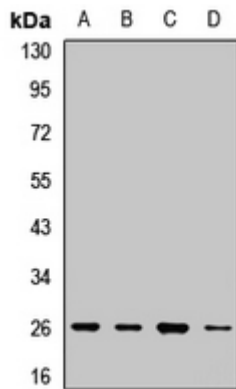
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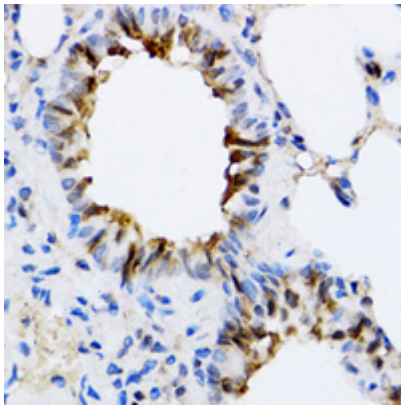
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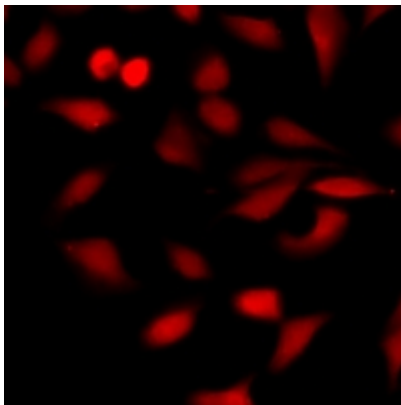
## Product Data Sheet



Western blot analysis of ARD1 expression in Hela (A), HepG2 (B), mouse thymus (C), rat brain (D) whole cell lysates. (Predicted band size: 24; 26 kD; Observed band size: 26 kD)



Immunohistochemical analysis of ARD1 staining in rat lung formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of ARD1 staining in U2OS cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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