

## **Product Data Sheet**

## **Anti-CUBN Antibody**

Catalog #	Source	e Reactivity	Applications		
CPA4380	Rabbit	н	WB, IH		
Description		Rabbit polyclonal antibody	to CUBN		
Immunogen		KLH-conjugated synthetic p	eptide encompassing a sequence within the N-term		
		region of human CUBN. Th	e exact sequence is proprietary.		
Purification		The antibody was purified	by immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	els of CUBN protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/5	0 - 1/100)		
Gene Symbol		CUBN			
Alternative Na	ames	IFCR; Cubilin; 460 kDa rece	ptor; Intestinal intrinsic factor receptor; Intrinsic		
		factor-cobalamin receptor;	Intrinsic factor-vitamin B12 receptor		
Entrez Gene		8029 (Human)			
SwissProt		O60494 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon deliv	ery 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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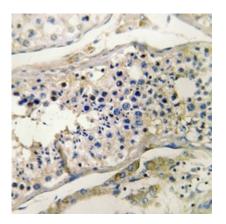
# Cohesion

kDa A

For research purposes only, not for human use

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Western blot analysis of CUBN expression in human kidney (A) whole cell lysates. (Predicted band size: 398 kD; Observed band size: 398 kD)



Immunohistochemical analysis of CUBN staining in human testis 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.

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