

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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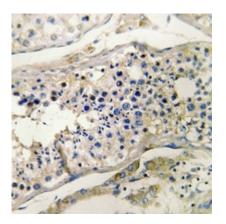
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kDa A

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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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