



## Cubilin Polyclonal Antibody

| Catalog No         | BYab-13186   |
|--------------------|--|
| Isotype            | lgG  |
| Reactivity         | Human;Rat;Mouse;   |
| Applications       | IHC;IF;WB;ELISA  |
| Gene Name          | CUBN   |
| Protein Name       | Cubilin  |
| Immunogen          | Synthesized peptide derived from the N-terminal region of human Cubilin.   |
| Specificity        | Cubilin Polyclonal Antibody detects endogenous levels of Cubilin protein.  |
| Formulation        | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.  |
| Source             | Polyclonal, Rabbit,IgG   |
| Purification       | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.  |
| Dilution           | WB 1:500-2000 IHC: 1/100 - 1/300. ELISA: 1/40000 IF 1:50-200   |
| Concentration      | 1 mg/ml  |
| Purity             | ≥90%   |
| Storage Stability  | -20°C/1 year   |
| Synonyms           | CUBN; IFCR; Cubilin; 460 kDa receptor; Intestinal intrinsic factor receptor;<br>Intrinsic factor-cobalamin receptor; Intrinsic factor-vitamin B12 receptor   |
| Observed Band      | 400kD  |
| Cell Pathway       | Apical cell membrane ; Peripheral membrane protein . Cell membrane ;<br>Peripheral membrane protein . Membrane, coated pit . Endosome . Lysosome<br>membrane ; Peripheral membrane protein . Lacks a transmembrane domain and<br>depends on interaction with AMN for location at the plasma membrane<br>(PubMed:29402915, PubMed:30523278). Colocalizes with AMN and LRP2 in the<br>endocytotic apparatus of epithelial cells (By similarity)  |
| Tissue Specificity | Detected in kidney cortex (at protein level) (PubMed:9572993). Expressed in kidney proximal tubule cells, placenta, visceral yolk-sac cells and in absorptive intestinal cells. Expressed in the epithelium of intestine and kidney.   |
| Function           | disease:Defects in CUBN are a cause of recessive hereditary megaloblastic<br>anemia 1 (MGA1) [MIM:261100]; also referred to as MGA1 Norwegian type or<br>Imerslund-Grasbeck syndrome (I-GS). MGA1 is due to selective malabsorption of<br>vitamin B12. Defects in vitamin B12 absorption lead to impaired function of<br>thymidine synthase. As a consequence DNA synthesis is interrupted. Rapidly<br>dividing cells involved in erythropoiesis are particularly affected.,domain:The CUB |
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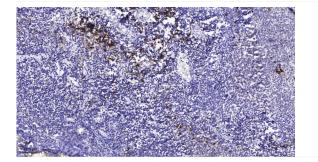
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|                           | domains 5 to 8 mediate binding to GIF and ALB. CUB domains 1 and 2 mediate<br>interaction with LRP2.,function:Cotransporter which plays a role in lipoprotein,<br>vitamin and iron metabolism, by facilitating their uptake. Binds to ALB, MB, Kappa<br>and lambda-light chains, TF, hemoglobin, GC, SCGB1A1, APOA1, high density<br>lipoprotein, and the GIF-cobalamin complex. The binding of all ligands required<br>calcium. Serves as important transporter in sev |
|---------------------------|---|
| Background                | Cubilin (CUBN) acts as a receptor for intrinsic factor-vitamin B12 complexes. The role of receptor is supported by the presence of 27 CUB domains. Cubulin is located within the epithelium of intestine and kidney. Mutations in CUBN may play a role in autosomal recessive megaloblastic anemia. [provided by RefSeq, Jul 2008],   |
| matters needing attention | Avoid repeated freezing and thawing!  |
| Usage suggestions         | This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.   |

## **Products Images**



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).

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