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PKD2 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP14827b

Specification

PKD2 Antibody (C-term) - Product Information

Application	**WB,E**
Primary Accession	Q13563
Other Accession	NP_000288.1
Reactivity	**Mouse**
Host	**Rabbit**
Clonality	**Polyclonal**
Isotype	**Rabbit IgG**
Calculated MW	**109691**
Antigen Region	**874-903**

PKD2 Antibody (C-term) - Additional Information

Gene ID 5311

Other Names
Polycystin-2, Autosomal dominant polycystic kidney disease type II protein, Polycystic kidney disease 2 protein, Polycystwin, R48321, PKD2

Target/Specificity
This PKD2 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 874-903 amino acids from the C-terminal region of human PKD2.

Dilution
WB~~1:1000
E~~Use at an assay dependent concentration.

Format
Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage
Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions
PKD2 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

PKD2 Antibody (C-term) - Protein Information

Name PKD2 (HGNC:9009)

Function Forms a nonselective cation channel (PubMed:11854751, PubMed:11991947, PubMed:15692563, PubMed:26269590, PubMed:27071085, PubMed:31441214, PubMed:39009345). Can function as a homotetrameric ion channel or can form heteromer with PKD1 (PubMed:31441214, PubMed:33164752). Displays distinct function depending on its subcellular localization and regulation by its binding partners (PubMed:11854751, PubMed:11991947, PubMed:27214281, PubMed:29899465). In primary cilium functions as a cation channel, with a preference for monovalent cations over divalent cations that allows K(+), Na(+) and Ca(2+) influx, with low selectivity for Ca(2+) (PubMed:27071085). Involved in fluid-flow mechanosensation by the primary cilium in renal epithelium (By similarity). In the endoplasmic reticulum, likely functions as a K(+) channel to facilitate Ca(2+) release (By similarity). The heterotetrameric PKD1/PKD2 channel has higher Ca(2+) permeability than homomeric PKD2 channel and acts as a primarily Ca(2+)-permeable channel (PubMed:31441214). Interacts with and acts as a regulator of a number of other channels, such as TRPV4, TRPC1, IP3R, RYR2, ultimately further affecting intracellular signaling, to modulate intracellular Ca(2+) signaling (PubMed:11854751, PubMed:11991947, PubMed:27214281, PubMed:29899465). Together with TRPV4, forms mechano- and thermosensitive channels in cilium (PubMed:18695040). In cardiomyocytes, PKD2 modulates Ca(2+) release from stimulated RYR2 receptors through direct association (By similarity). Also involved in left-right axis specification via its role in sensing nodal flow; forms a complex with PKD1L1 in cilia to facilitate flow detection in left- right patterning (By similarity). Acts as a regulator of cilium length together with PKD1 (By similarity). Mediates systemic blood pressure and contributes to the myogenic response in cerebral arteries though vasoconstriction (By similarity).

Cellular Location
Cell projection, cilium membrane; Multi-pass membrane protein. Endoplasmic reticulum membrane; Multi-pass membrane protein. Cell membrane; Multi-pass membrane protein. Basolateral cell membrane. Cytoplasmic vesicle membrane. Golgi apparatus {ECO:0000250|UniProtKB:O35245}. Vesicle Secreted, extracellular exosome Note=PKD2 localization to the plasma and ciliary membranes requires PKD1. PKD1:PKD2 interaction is required to reach the Golgi apparatus form endoplasmic reticulum and then traffic to the cilia (By similarity). Retained in the endoplasmic reticulum by interaction with PACS1 and PACS2 (PubMed:15692563). Detected on kidney tubule basolateral membranes and basal cytoplasmic vesicles (PubMed:10770959) Cell surface and cilium localization requires GANAB (PubMed:27259053) Detected on migrasomes and on extracellular exosomes in urine (PubMed:21406692). Preferentially localized to the dorsal side of immotile cilia (By similarity). {ECO:0000250|UniProtKB:O35245, ECO:0000269|PubMed:15692563, ECO:0000269|PubMed:21406692, ECO:0000269|PubMed:27259053}

Tissue Location
Detected in fetal and adult kidney (PubMed:10770959). Detected at the thick ascending limb of the loop of Henle, at distal tubules, including the distal convoluted tubule and cortical collecting tubules, with weak staining of the collecting duct (PubMed:10770959). Detected on placenta syncytiotrophoblasts (at protein level) (PubMed:26269590). Strongly expressed in ovary, fetal and adult kidney, testis, and small intestine. Not detected in peripheral leukocytes.

PKD2 Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation

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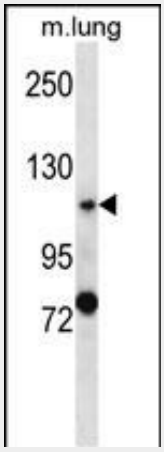
- Flow Cytomety
- Cell Culture

PKD2 Antibody (C-term) - Images



PKD2 Antibody (C-term) (Cat. #AP14827b) western blot analysis in mouse lung tissue lysates (35ug/lane).This demonstrates the PKD2 antibody detected the PKD2 protein (arrow).

PKD2 Antibody (C-term) - Background

This gene encodes a member of the polycystin protein family. The encoded protein contains multiple transmembrane domains, and cytoplasmic N- and C-termini. The protein may be an integral membrane protein involved in cell-cell/matrix interactions. The encoded protein may function in renal tubular development, morphology, and function, and may modulate intracellular calcium homoeostasis and other signal transduction pathways. This protein interacts with polycystin 1 to produce cation-permeable currents. Mutations in this gene have been associated with autosomal dominant polycystic kidney disease.

PKD2 Antibody (C-term) - References

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Bailey, S.D., et al. Diabetes Care 33(10):2250-2253(2010)
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