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## Datasheet for ABIN1392366 anti-PKD1L3 antibody (AA 121-220) (Alexa Fluor 647)



Overview

| Quantity:            | 100 µL  |
|----------------------|---|
| Target:              | PKD1L3  |
| Binding Specificity: | AA 121-220  |
| Reactivity:          | Human   |
| Host:                | Rabbit  |
| Clonality:           | Polyclonal  |
| Conjugate:           | This PKD1L3 antibody is conjugated to Alexa Fluor 647   |
| Application:         | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

## Product Details

| Immunogen:            | KLH conjugated synthetic peptide derived from human PKD1L3 |
|-----------------------|--|
| Isotype:              | IgG  |
| Predicted Reactivity: | Human  |
| Purification:         | Purified by Protein A.                                     |

## Target Details

| Target:           | PKD1L3  |
|-------------------|---|
| Alternative Name: | PKD1L3 (PKD1L3 Products)  |
| Background:       | Synonyms: PC1 like 3 protein, Polycystic kidney disease 1 like 3, Polycystic kidney disease |

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 1/2 | Product datasheet for ABIN1392366 | 03/07/2024 | Copyright antibodies-online. All rights reserved. protein 1 like 3, Polycystin 1 like 3, Polycystin 1L3, PK1L3\_HUMAN.

Background: Polycystin-1L3 is a 1,732 amino acid multi-pass membrane protein that contains one PLAT domain, one GPS domain and one C-type lectin domain. Expressed at high levels in placenta and present at lower levels in lung and heart, Polycystin-1L3 is thought to function as an ion-channel regulator that may interact with Polycystin-L and play a role in heteromeric taste channels. The gene encoding Polycystin-1L3 maps to human chromosome 16, which encodes over 900 genes and comprises nearly 3 % of the human genome. The GAN gene is located on chromosome 16 and, with mutation, may lead to giant axonal neuropathy, a nervous system disorder characterized by increasing malfunction with growth. The rare disorder Rubinstein-Taybi syndrome is also associated with chromosome 16, as is Crohn's disease, which is a gastrointestinal inflammatory condition.

## **Application Details**

| Application Notes: | IF(IHC-P) 1:50-200   |  |
|--------------------|--|--|
|                    | IF(IHC-F) 1:50-200   |  |
|                    | IF(ICC) 1:50-200   |  |
| Restrictions:      | For Research Use only  |  |
| Handling           |  |  |
| Format:            | Liquid   |  |
| Concentration:     | 1 µg/µL  |  |
| Buffer:            | Aqueous buffered solution containing 0.01M TBS ( pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol.        |  |
| Preservative:      | ProClin  |  |
| Precaution of Use: | This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only. |  |
| Storage:           | -20 °C   |  |
| Storage Comment:   | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles.                                  |  |
| Expiry Date:       | 12 months  |  |

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