For Research Use Only

OCRL Polyclonal antibody

Catalog Number: 17695-1-AP

Featured Product

5 Publications

BC094726

GenBank Accession Number:



Basic Information

Catalog Number: 17695-1-AP

Size: GenelD (NCBI): 150ul, Concentration: 500 µg/ml by 4952

Nanodrop and 267 µg/ml by Bradford Full Name:

method using BSA as the standard;

 Source:
 Calculated MW:

 Rabbit
 893 aa, 103 kDa

 Isotype:
 Observed MW:

 IgG
 105 kDa

Immunogen Catalog Number:

AG11900

Purification Method:
Antigen affinity purification

Recommended Dilutions: WB 1:500-1:2000

IP 0.5-4.0 ug for IP and 1:200-1:1000

for WB

oculocerebrorenal syndrome of Lowe HHC 1:100-1:400

Applications

Tested Applications:

IHC, IP, WB, ELISA

Cited Applications:

IF, IHC, WB

Species Specificity: human, mouse, rat Cited Species:

human, mouse

Note-IHC: suggested antigen retrieval with

TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate

buffer pH 6.0

Positive Controls:

WB: HeLa cells, HEK-293 cells, mouse brain tissue, rat

brain tissue, SH-SY5Y cells

IP : HeLa cells,

IHC: mouse kidney tissue,

Background Information

OCRL is also named as INPP5F, OCRL1 and belongs to the 5-phosphatase gene family and that Lowe syndrome represents an inborn error of inositol phosphate metabolism(PMID: 9430698). The protein product of the gene that when mutated is responsible for Lowe syndrome, or oculocerebrorenal syndrome (OCRL), is an inositol polyphosphate 5-phosphatase. It may function in lysosomal membrane trafficking by regulating the specific pool of phosphatidylinositol 4,5-bisphosphate that is associated with lysosomes. It has 2 isoforms produced by alternative splicing. Defects in OCRL are the cause of Lowe oculocerebrorenal syndrome (OCRL) and Dent disease type 2 (DD2). This antibody is specific to OCRL

Notable Publications

| Author | Pubmed ID | Journal | Application |
|-----------------|-----------|-------------------------|-------------|
| Nana Sakakibara | 34586410 | Nephrol Dial Transplant | WB |
| Yu Zhang | 34488756 | BMC Med Genomics | IHC |
| Hequn Liu | 32393163 | J Neurodev Disord | WB |

Storage

Storage:

Store at -20°C. Stable for one year after shipment.

Storage Buffer

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

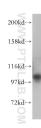
*** 20ul sizes contain 0.1% BSA

For technical support and original validation data for this product please contact:

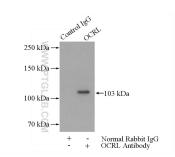
T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA)

E: proteintech@ptglab.com W: ptglab.com This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

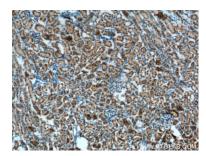
Selected Validation Data



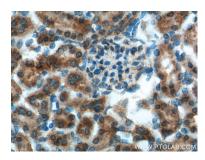
HeLa cells were subjected to SDS PAGE followed by western blot with 17695-1-AP (OCRL antibody) at dilution of 1:300 incubated at room temperature for



IP Result of anti-OCRL (IP:17695-1-AP, 4ug; Detection:17695-1-AP 1:300) with HeLa cells lysate 1080ug.



Immunohistochemical analysis of paraffinembedded mouse kidney tissue slide using 17695-1-AP (OCRL Antibody) at dilution of 1:200 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunohistochemical analysis of paraffinembedded mouse kidney tissue slide using 17695-1-AP (OCRL Antibody) at dilution of 1:200 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).