For Research Use Only

CEP89, CCDC123 Monoclonal antibody

Catalog Number: 68112-1-Ig



Purification Method:

Basic Information

Catalog Number: GenBank Accession Number:

68112-1-lg BC136328 Protein G purification

Size:GeneID (NCBI):CloneNo.:150ul , Concentration: 1000 ug/ml by849021F12C5

 Nanodrop;
 UNIPROT ID:
 Recommended Dilutions:

 Source:
 Q96ST8
 WB 1:1000-1:4000

 Mouse
 Full Name:
 IF-P 1:200-1:800

Isotype: coiled-coil domain containing 123

IgG1 Calculated MW:
Immunogen Catalog Number: 783 aa, 90 kDa
AG28339 Observed MW:
90 kDa

Applications

Tested Applications:

WB, IF-P, ELISA WB: HeLa cells, A549 cells, Neuro-2a cells, A431 cells,

Positive Controls:

 Species Specificity:
 SH-5Y5Y cells, U-251 cells, SH-SY5Y cells

 human, mouse
 IF-P: mouse eye tissue, hTERT-RPE1 cells

Background Information

CCDC 123 (as known as CEP123), also named as CEP89, encodes for a protein required for ciliogenesis. It plays a role in mitochondrial metabolism by modulating complex IV activity. It has been shown that CEP123 is localized to the distal appendages of the mother centriolecep and the localization of CEP123 is cell cycle-dependent with its levels decreasing during mitosis. CEP123 depletion can cause defects in ciliary vesicle formationcep and prevent the formation of a ciliary vesicle at the distal end of the mother centriole. It is possible that CEP123 is involved in regulating the recruitment of membranes to the centrosome through its interaction with CEP290 (PMID:23575228, 23789104, 23348840).

Storage

Storage

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

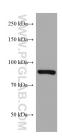
Storage Buffer

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

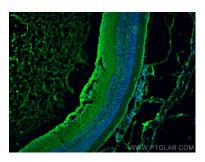
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

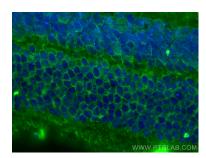
Selected Validation Data



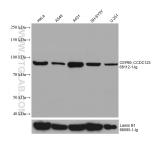
Neuro-2a cells were subjected to SDS PAGE followed by western blot with 68112-1-lg (CCDC123 antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours.



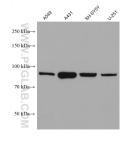
Immunofluorescent analysis of (4% PFA) fixed mouse eye tissue using CEP89, CCDC123 antibody (68112-1-1g, Clone: 1F12C5) at dilution of 1:400 and Coralite®488-Conjugated Goat Anti-Mouse IgG(H+L).



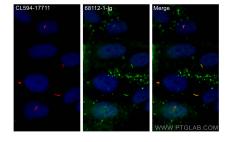
Immunofluorescent analysis of (4% PFA) fixed mouse eye tissue using CEP89, CCDC123 antibody (68112-1-1g, Clone: 1F12C5) at dilution of 1:400 and Coralite® 488-Conjugated Goat Anti-Mouse IgG(H+L).



Various lysates were subjected to SDS PAGE followed by western blot with 681.12-1-lg (CEP89, CCDC 123 antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours. The membrane was stripped and reblotted with Lamin B1 Monoclonal antibody (66095-1-lg) as loading control.



Various lysates were subjected to SDS PAGE followed by western blot with 68112-1-lg (CCDC 123 antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours.



Immunofluorescent analysis of (4% PFA) fixed hTERT-RPE1 cells using CEP89, CCDC 123 antibody (68112-1-lg, Clone: 1F12C5) at dilution of 1:400 and Multi-rAb Coralite ® Plus 488-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (RGAM002), Coralite®594 ARL13B antibody (CL594-17711, red).