

Product datasheet for CF500915

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

RALBP1 Mouse Monoclonal Antibody [Clone ID: OTI6G10]

Product data:

Product Type: Primary Antibodies

Clone Name: OTI6G10

Applications: FC, IF, IHC, IP, WB

Recommended Dilution: WB 1:2000, IHC 1:50, IF 1:100, Flow 1:100, IP: 4ug/mL

Reactivity: Human, Mouse, Rat

Host: Mouse Isotype: IgG2a

Clonality: Monoclonal

Immunogen: Full length human recombinant protein of human RALBP1 (NP_006779) produced in HEK293T

cell.

Formulation: Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose)

Reconstitution Method: For reconstitution, we recommend adding 100uL distilled water to a final antibody

concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific)

Purification: Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: Unconjugated

Storage: Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Predicted Protein Size: 76.0 kDa

Gene Name: ralA binding protein 1

Database Link: NP 006779

Entrez Gene 19765 MouseEntrez Gene 84014 RatEntrez Gene 10928 Human

Q15311





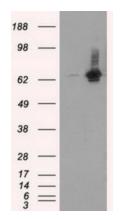
Background:

Can activate specifically hydrolysis of GTP bound to RAC1 and CDC42, but not RALA. Mediates ATP-dependent transport of S-(2,4-dinitrophenyl)-glutathione (DNP-SG) and doxorubicin (DOX) and is the major ATP-dependent transporter of glutathione conjugates of electrophiles (GS-E) and DOX in erythrocytes. Can catalyze transport of glutathione conjugates and xenobiotics, and may contribute to the multidrug resistance phenomenon. Serves as a scaffold protein that brings together proteins forming an endocytotic complex during interphase and also with CDC2 to switch off endocytosis, One of its substrates would be EPN1/Epsin

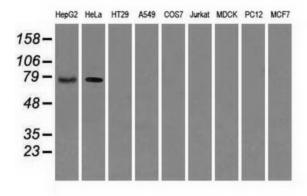
Synonyms: RIP1; RLIP1; RLIP76

Protein Pathways: Pancreatic cancer, Pathways in cancer

Product images:

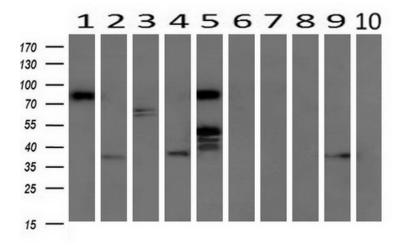


HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY RALBP1 ([RC201524], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-RALBP1. Positive lysates [LY402031] (100ug) and [LC402031] (20ug) can be purchased separately from OriGene.

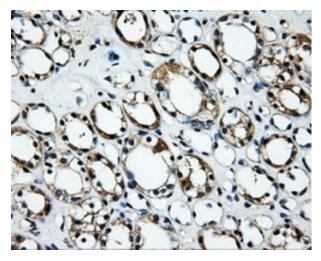


Western blot analysis of extracts (35ug) from 9 different cell lines by using anti-RALBP1 monoclonal antibody.

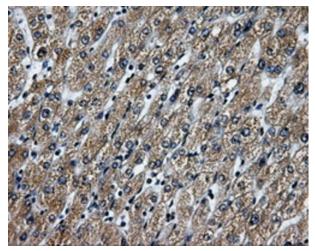




Western blot analysis of extracts (10ug) from 10 Human tissue by using anti-RALBP1 monoclonal antibody at 1:200 (1: Testis; 2: Omentum; 3: Uterus; 4: Breast; 5: Brain; 6: Liver; 7: Ovary; 8: Thyroid gland; 9: colon;10: spleen).

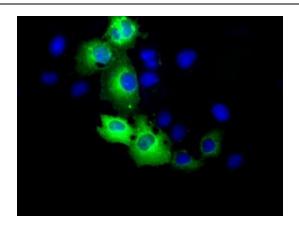


Immunohistochemical staining of paraffinembedded Kidney tissue within the normal limits using anti-RALBP1mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

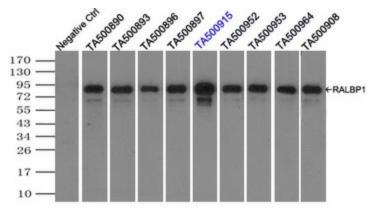


Immunohistochemical staining of paraffinembedded liver tissue within the normal limits using anti-RALBP1mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

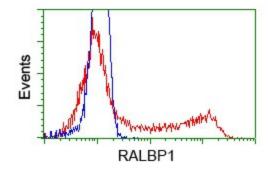




Anti-RALBP1 mouse monoclonal antibody ([TA500915]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY RALBP1 ([RC201524]).



Immunoprecipitation (IP) of RALBP1 by using TrueMab monoclonal anti-RALBP1 antibodies (Negative control: IP without adding anti-RALBP1 antibody.). For each experiment, 500ul of DDK tagged RALBP1 overexpression lysates (at 1:5 dilution with HEK293T lysate), 2ug of anti-RALBP1 antibody and 20ul (0.1mg) of goat anti-mouse conjugated magnetic beads were mixed and incubated overnight. After extensive wash to remove any non-specific binding, the immunoprecipitated products were analyzed with rabbit anti-DDK polyclonal antibody.



HEK293T cells transfected with either pCMV6-ENTRY RALBP1 ([RC201524]) (Red) or empty vector control plasmid (Blue) were immunostained with anti-RALBP1 mouse monoclonal ([TA500915]), and then analyzed by flow cytometry.