RbAp48 antibody [13D10]

Cat. No. GTX70234

Host	Mouse
Clonality	Monoclonal
lsotype	lgG2b
Applications	WB, ICC/IF, IP, IHC
Reactivity	Human

References (5) Package 100 µl

Applications

Application Note

*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	1:500-1:3000
ICC/IF	1:100-1:1000
IP	1:500
IHC	Assay dependent
Not tested in other applications.	

d in other applications.

Calculated MW

48 kDa. (<u>Note</u>)

Properties	
Form	Liquid
Buffer	PBS
Preservative	No preservative
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	1 mg/ml (Please refer to the vial label for the specific concentration.)
Immunogen	The complete coding region (amino acids 1-425) of RbAp48 expressed in E. coli.
Purification	Protein G purified From tissue culture supernatant
Conjugation	Unconjugated



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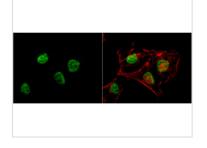


Note

For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

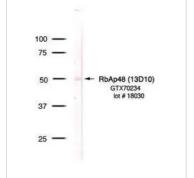
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DATA IMAGES



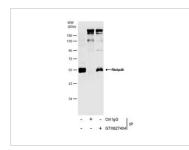
GTX70234 ICC/IF Image

RbAp48 antibody [13D10] detects RbAp48 protein at nucleus by immunofluorescent analysis. Sample: HeLa cells were fixed in 4% paraformaldehyde at RT for 15 min. Green: RbAp48 protein stained by RbAp48 antibody [13D10] (GTX70234) diluted at 1:200. Red: phalloidin, a cytoskeleton marker, stained by phalloidin (invitrogen, A12380) diluted at 1:200. Blue: Hoechst 33342 staining.



GTX70234 WB Image

Detection of RbAp48 in HeLa nuclear extract (GTX24105)

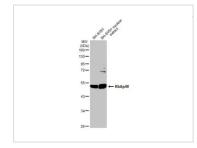


GTX70234 IP Image

Immunoprecipitation of RbAp48 protein from HeLa whole cell extract using 5 μ g of RbAp48 antibody [13D10] (GTX70234).

Western blot analysis was performed using RbAp48 antibody [13D10] (GTX70234).

EasyBlot HRP-conjugated anti mouse IgG antibody (GTX221667-01) was used to detect the primary antibody, and the signal was developed with Trident ECL plus-Enhanced.



GTX70234 WB Image

SH-SY5Y whole cell and nuclear extracts (30 µg) were separated by 10% SDS-PAGE, and the membrane was blotted with RbAp48 antibody [13D10] (GTX70234) diluted at 1:2000. The HRP-conjugated anti-mouse IgG antibody (GTX213111-01) was used to detect the primary antibody.



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