abcam

Product datasheet

Anti-CRMP1 + CRMP2 antibody [EPR7792] ab129082

Recombinant RabMAb

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Overview



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Learn more about K_D

Storage buffer	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: PBS, 40% Glycerol, 0.05% BSA	
Purity	Protein A purified	
Clonality	Monoclonal	
Clone number	EPR7792	
lsotype	lgG	

Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab129082 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

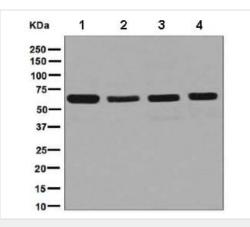
Application	Abreviews	Notes
Flow Cyt (Intra)		1/10 - 1/1000. <u>ab172730</u> - Rabbit monoclonal IgG, is suitable for use as an isotype control with this antibody.
WB		1/10000 - 1/50000. Detects a band of approximately 64 kDa (predicted molecular weight: 62 kDa).
IP		1/10 - 1/100.
IHC-P		 1/1000. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. For unpurified use at 1/100 - 1/250. See <u>IHC antigen retrieval protocols</u>. The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat.
ICC/IF		1/50 - 1/100.

Target

Cellular localization

CRMP1: Cytoplasm. Cytoplasm > cytoskeleton > centrosome. Cytoplasm > cytoskeleton > spindle. Associated with centrosomes and the mitotic spindle during metaphase. CRMP2: Cytoplasm.

Images



Western blot - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) All lanes : Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) at 1/10000 dilution

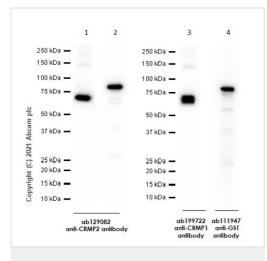
Lane 1 : PC12 cell lysate Lane 2 : Jurkat cell lysate Lane 3 : U87 MG cell lysate Lane 4 : NIH 3T3 cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : HRP labelled goat anti-rabbit at 1/2000 dilution

Predicted band size: 62 kDa Observed band size: 64 kDa



Western blot - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) Lanes 1-2 : Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) at 1/1000 dilution Lane 3 : Anti-CRMP1 antibody [EP14521] (<u>ab199722</u>) at 1/1000 dilution

Lane 4 : Anti-GST antibody [EPR4236] (ab111947) at 1/1000 dilution

Lanes 1 & 3 : Recombinant Human CRMP1 protein (<u>ab202599</u>) Lanes 2 & 4 : Recombinant Human CRMP2 protein (<u>ab114269</u>)

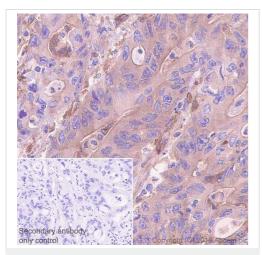
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 62 kDa

Exposure time: 3 seconds

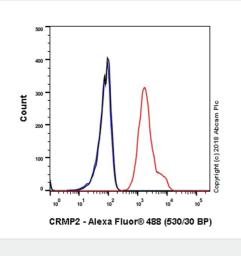
Blocking and dilution buffer: 5% NFDM /TBST



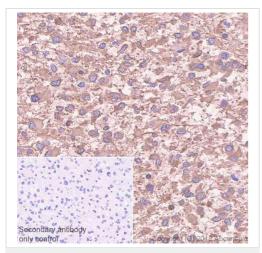
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human colon carcinoma tissue sections labeling CRMP2 with Purified ab129082 at 1:1000 dilution (0.12 µg/ml). Heat mediated antigen retrieval was performed using **ab93684** (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use)was used as the secondary antibody. Negative control:PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



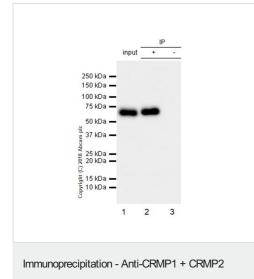
Immunohistochemistry (Formalin/PFA-fixed paraffin-



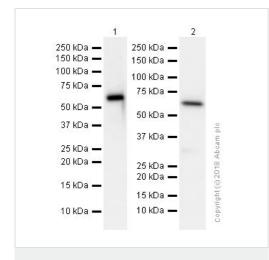
Flow Cytometry (Intracellular) - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) Intracellular Flow Cytometry analysis of NIH/3T3 (Mouse embryonic fibroblast) cells labeling CRMP2 with Purified ab129082 at 1/20 dilution (10µg/ml) (red). Cells were fixed with 4% Paraformaldehyde. A Goat anti rabbit IgG (Alexa Fluorr[®] 488, **ab150077**) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal IgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082)



antibody [EPR7792] (ab129082)



Western blot - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human glioma tissue sections labeling CRMP2 with Purified ab129082 at 1:1000 dilution (0.12 µg/ml). Heat mediated antigen retrieval was performed using **ab93684** (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use)was used as the secondary antibody. Negative control:PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

ab129082 (purified) at 1:20 dilution (0.5µg) immunoprecipitating CRMP2 in U-87 MG whole cell lysate. Lane 1 (input): U-87 MG (Human glioblastoma-astrocytoma epithelial cell) whole cell lysate 10µg Lane 2 (+): ab129082 & U-87 MG whole cell lysate Lane 3 (-): Rabbit monoclonal lgG (<u>ab172730</u>) instead of ab129082 in U-87 MG whole cell lysate For western blotting, VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) was used for detection at 1:1000 dilution. Blocking and diluting buffer: 5% NFDM/TBST.

All lanes : Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) at 1/50000 dilution

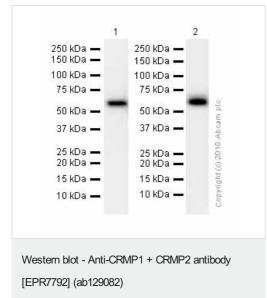
Lane 1 : U-87 MG (Human glioblastoma-astrocytoma epithelial cell) whole cell lysates

Lane 2 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates

Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 62 kDa Observed band size: 64 kDa U-87 MG (Human glioblastoma-astrocytoma epithelial cell) whole cell lysates



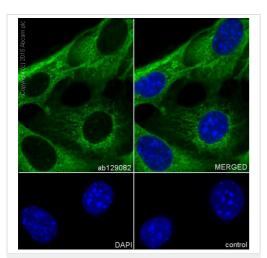
All lanes : Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) at 1/100000 dilution

Lane 1 : NIH/3T3 (Mouse embryonic fibroblast) whole cell lysates Lane 2 : PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysates

Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

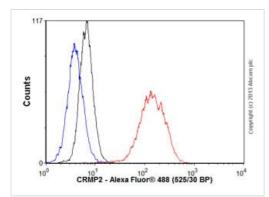
Predicted band size: 62 kDa Observed band size: 64 kDa



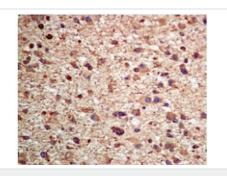
Immunocytochemistry/ Immunofluorescence - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) ab129082 staining CRMP2 in NIH/3T3 (mouse embryonic fibroblast) cells by ICC/IF

(Immunocytochemistry/immunofluorescence). Cells were fixed with 100% methanol. Samples were incubated with primary antibody at a dilution of 1/500. A goat anti rabbit IgG (Alexa Fluor[®] 488) (**ab150077**) was used as the secondary antibody at a dilution of 1/1000. DAPI was used as a nuclear counterstain.

Negative control 1: PBS only.

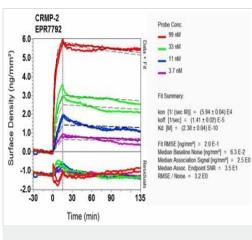


Flow Cytometry (Intracellular) - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) Overlay histogram showing SH-SY5Y cells stained with ab129082 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab129082, 1/1000 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluorr[®] 488 goat anti-rabbit lgG (H&L) (<u>ab150077</u>) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (0.1μ g/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



ab129082, at a dilution of 1/100, staining CRMP2 in paraffin embedded Human astrocytoma tissue by Immunohistochemistry. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082)



OI-RD Scanning - Anti-CRMP1 + CRMP2 antibody [EPR7792] (ab129082) Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about KD



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