

Product datasheet

Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] ab157455

Recombinant RabMAb

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Overview		
Product name	Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471]	
Description	Rabbit monoclonal [EPR9471] to eEF1A1/EF-Tu+eEF1A1 + eEF1AL3	
Host species	Rabbit	
Specificity	The immunogen used for this product shares 6 continuous identical amino acids with eEF1A2. Cross-reactivity with this protein has not been confirmed experimentally.	
Tested applications	Suitable for: WB, IHC-P, ICC/IF, IP, Flow Cyt (Intra)	
Species reactivity	Reacts with: Mouse, Rat, Human	
Immunogen	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.	
Positive control	WB: HeLa, MCF7, 293T and Neuro-2a cell lysates. GST tagged Recombinant Human EEF1A1 and EEF1AL3 protein	
General notes	 This product is a recombinant monoclonal antibody, which offers several advantages including: High batch-to-batch consistency and reproducibility Improved sensitivity and specificity Long-term security of supply Animal-free production For more information <u>see here</u>. Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb[®] patents</u>. 	

Properties

Form	Liquid	
Storage instructions	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C. Stable for 12 months at -20°C.	
Storage buffer	Preservative: 0.01% Sodium azide Constituents: 40% Glycerol (glycerin, glycerine), 0.05% BSA, 59% PBS	
Purity	Protein A purified	
Clonality	Monoclonal	

Clone number	EPR9471
lsotype	lgG

Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab157455 in the following tested applications.

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

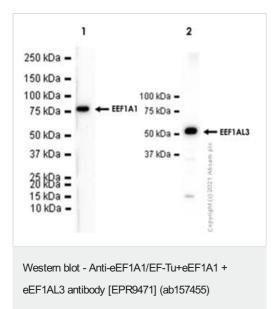
Application	Abreviews	Notes
WB	★★★★ <u></u> (<u>1)</u>	1/40000. Predicted molecular weight: 50 kDa. For unpurified use at 1/1000 - 1/10000.
IHC-P		1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/100.
IP		1/10 - 1/100.
Flow Cyt (Intra)		Use at an assay dependent concentration.

Target

Cellular localization

eEF1A1/EF-Tu: Cytoplasm.

Images



All lanes : Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) at 1/1000 dilution

Lane 1 : GST tagged Recombinant Human EEF1A1 protein (Full length, 76 KDa)

Lane 2 : His tagged Recombinant Human EEF1AL3 protein (Full length, 52 KDa)

Secondary

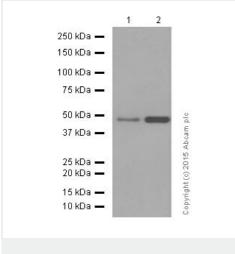
All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 50 kDa

Exposure time:

Lane 1: 180 seconds

Lane 2: 5 seconds



Western blot - Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455)

All lanes : Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) at 1/40000 dilution (purified)

Lane 1 : Rat kidney lysate Lane 2 : Rat spleen lysate

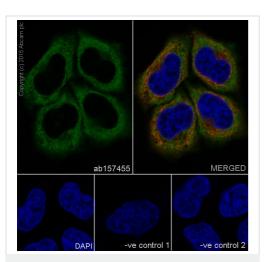
Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/10000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 50 kDa

Blocking buffer and concentration: 5% NFDM/TBST. Diluting buffer and concentration: 5% NFDM /TBST.

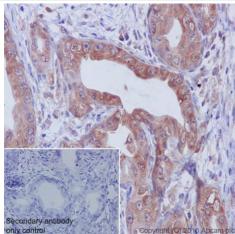


Immunocytochemistry/ Immunofluorescence - AntieEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455)

Immunocytochemistry/Immunofluorescence analysis of HeLa (human cervix adenocarcinoma) cells labelling eEF1A1/EF-Tu with purified ab157455 at 1/100. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. **ab150077**, an Alexa Fluor[®] 488-conjugated goat anti-rabbit IgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain. **ab7291**, a mouse anti-tubulin (1/1000) and **ab150120**, an Alexa Fluor[®] 594-conjugated goat anti-mouse IgG (1/1000) were also used.

Control 1: primary antibody (1/100) and secondary antibody, <u>ab150120</u>, an Alexa Fluor[®] 594-conjugated goat anti-mouse IgG (1/500).

Control 2: <u>**ab7291**</u> (1/1000) and secondary antibody, <u>**ab150077**</u>, an Alexa Fluor[®] 488-conjugated goat anti-rabbit IgG (1/500).

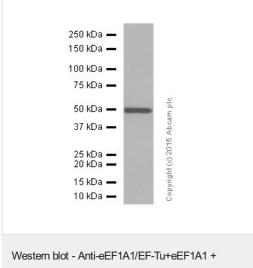


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455)

eEF1A1- Alexa Fluor® 488 (530/30 BP)

Flow Cytometry (Intracellular) - Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human cervix carcinoma tissue labelling eEF1A1/EF-Tu with purified ab157455 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. **ab97051**, a goat anti-rabbit IgG H&L (HRP) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma) cells labeling eEF1A1/EF-Tu with purified ab157455 at 1/50 dilution (10ug/ml) (red). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. A Goat anti rabbit IgG (Alexa Fluor[®] 488) (1/2000 dilution) was used as the secondary antibody. Rabbit monoclonal IgG (Black) was used as the isotype control, cells without incubation with primary antibody and secondary antibody (Blue) was used as the unlabeled control.



Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) at 1/40000 dilution (purified) + Mouse kidney lysate at 20 μg

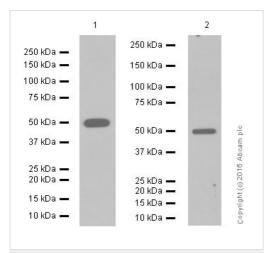
Secondary

Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/10000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 50 kDa

Western blot - Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455)

Blocking buffer and concentration: 5% NFDM/TBST. Diluting buffer and concentration: 5% NFDM/TBST.



Western blot - Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) **All lanes** : Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) at 1/50000 dilution (purified)

Lane 1 : MCF-7 cell lysate Lane 2 : HeLa cell lysate

Lysates/proteins at 20 µg per lane.

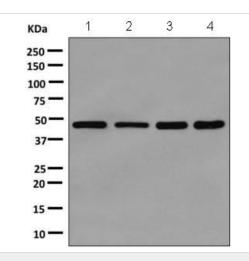
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 20 μg (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 50 kDa

Blocking buffer and concentration: 5% NFDM/TBST. Diluting buffer and concentration: 5% NFDM /TBST.

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Western blot - Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) **All lanes** : Anti-eEF1A1/EF-Tu+eEF1A1 + eEF1AL3 antibody [EPR9471] (ab157455) at 1/1000 dilution (unpurified)

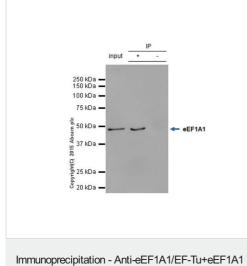
Lane 1 : HeLa cell lysate Lane 2 : MCF7 cell lysate Lane 3 : 293T cell lysate Lane 4 : Neuro-2a cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

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

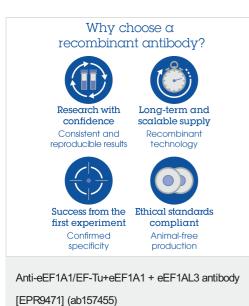
Predicted band size: 50 kDa



+ eEF1AL3 antibody [EPR9471] (ab157455)

ab157455 (purified) at 1/30 immunoprecipitating eEF1A1/EF-Tu in HeLa whole cell lysate. 10 ug of cell lysate was present in the input. For western blotting, a HRP-conjugated Veriblot for IP Detection Reagent (**ab131366**) (1/1,500) was used for detection. A rabbit monoclonal IgG (**ab172730**) was used intead of **ab128913** as a negative control (Lane 3).

Blocking buffer and concentration: 5% NFDM/TBST. Diluting buffer and concentration: 5% NFDM/TBST.



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