# abcam

## Product datasheet

# Anti-p130 antibody [EP2141Y] - BSA and Azide free ab247453



#### 11 Images

#### Overview

**Product name** Anti-p130 antibody [EP2141Y] - BSA and Azide free

**Description** Rabbit monoclonal [EP2141Y] to p130 - BSA and Azide free

**Host species** Rabbit

**Tested applications** Suitable for: Flow Cyt (Intra), IHC-P, WB

Unsuitable for: ICC/IF or IP

Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

**General notes** ab247453 is the carrier-free version of ab76234.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.

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 see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

#### **Properties**

Form Liquid

**Storage instructions** Shipped at 4°C. Store at +4°C. Do Not Freeze.

EP2141Y

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

**Clonality** Monoclonal

Isotype IqG

#### **Applications**

Clone number

The Abpromise guarantee Our Abpromise guarantee covers the use of ab247453 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)		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols.
WB		Use at an assay dependent concentration. Detects a band of approximately 128 kDa (predicted molecular weight: 128 kDa).

**Application notes** Is unsuitable for ICC/IF or IP.

Nucleus.

## **Target**

Post-translational

**Cellular localization** 

modifications

Function

Key regulator of entry into cell division. Directly involved in heterochromatin formation by maintaining overall chromatin structure and, in particular, that of constitutive heterochromatin by stabilizing histone methylation. Recruits and targets histone methyltransferases SUV420H1 and SUV420H2, leading to epigenetic transcriptional repression. Controls histone H4 'Lys-20' trimethylation. Probably acts as a transcription repressor by recruiting chromatin-modifying enzymes to promoters. Potent inhibitor of E2F-mediated trans-activation, associates preferentially with E2F5. Binds to cyclins A and E. Binds to and may be involved in the transforming capacity of the adenovirus E1A protein. May act as a tumor suppressor.

Sequence similarities

Belongs to the retinoblastoma protein (RB) family.

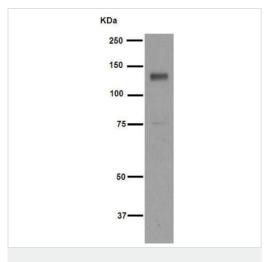
Developmental stage

G0-restricted expression.

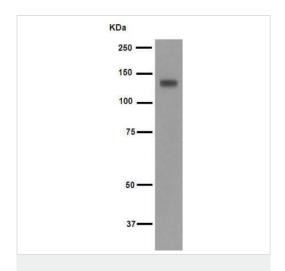
During G0 and early G1 phase of the cell cycle, phosphorylated on Ser-639 and on 5 sites within

the domain B. Phosphorylation on Ser-672 in G1 leads to its ubiquitin-dependent proteolysis.

2



Western blot - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)



Western blot - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)

Anti-p130 antibody [EP2141Y] (ab76234) at 1/1000 dilution (unpurified) + Daudi cell lysate at 10  $\mu g$ 

#### Secondary

Peroxidase-conjugated goat anti-rabbit IgG at 1/1000 dilution

**Predicted band size:** 128 kDa **Observed band size:** 130 kDa

This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.

Anti-p130 antibody [EP2141Y] ( $\underline{ab76234}$ ) at 1/1000 dilution (purified) + Daudi cell lysate at 10  $\mu g$ 

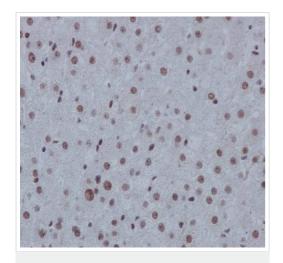
# Secondary

Peroxidase-conjugated goat anti-rabbit lgG (H+L) at 1/1000 dilution

**Predicted band size:** 128 kDa **Observed band size:** 130 kDa

This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.

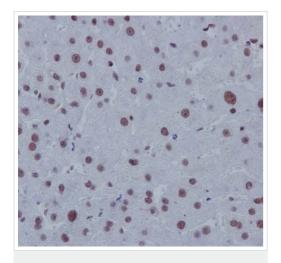
Blocking and dilution buffer: 5% NFDM/TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)

This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.

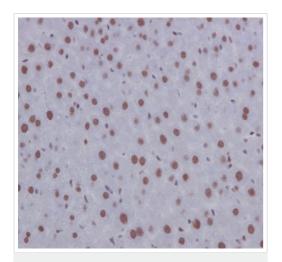
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human liver tissue sections labelling p130 with unpurified <a href="mailto:ab76234">ab76234</a> at 1/250. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)

This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.

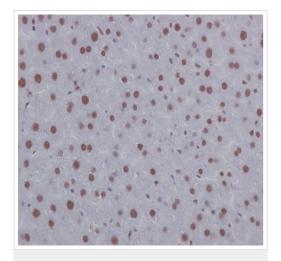
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human liver tissue sections labelling p130 with purified <a href="mailto:ab76234">ab76234</a> at 1/300. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)

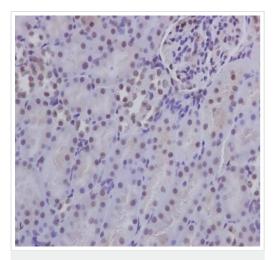
This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse liver tissue sections labelling p130 with unpurified <a href="mailto:ab76234">ab76234</a> at 1/250. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.



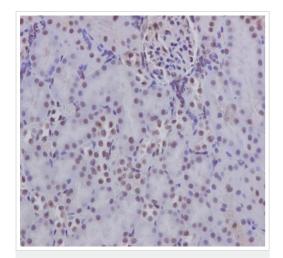
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This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.lmmunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse liver tissue sections labelling p130 with purified <u>ab76234</u> at 1/300. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated antirabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.



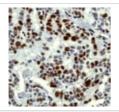
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)

This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.lmmunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat kidney tissue sections labelling p130 with unpurified <u>ab76234</u> at 1/250. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated antirabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.



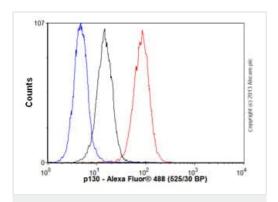
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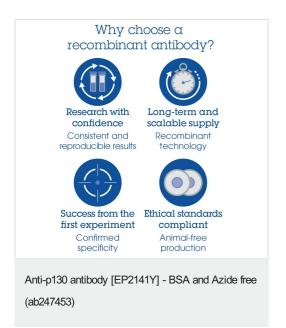
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This data was developed using <u>ab76234</u>, the same antibody clone in a different buffer formulation.lmmunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human breast carcinoma tissue labelling p130 with unpurified <u>ab76234</u> at 1/50. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-p130 antibody [EP2141Y] - BSA and Azide free (ab247453)

This data was developed using ab76234, the same antibody clone in a different buffer formulation. Overlay histogram showing Jurkat cells stained with unpurified ab76234 (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 proteinprotein interactions followed by the antibody (ab76234, 1/100 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-rabbit lgG (H&L) (ab150077) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (1µg/1x10<sup>6</sup> 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. This antibody gave a positive signal in Jurkat cells fixed with 4% paraformaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



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