abcam

Product datasheet

Anti-Hemopexin antibody [EPR5610] - BSA and Azide free ab226099



6 Images

Overview

Product name Anti-Hemopexin antibody [EPR5610] - BSA and Azide free

Description Rabbit monoclonal [EPR5610] to Hemopexin - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: ICC/IF, WB, IP, IHC-P, Flow Cyt (Intra)

Species reactivity Reacts with: Human

Does not react with: Mouse, Rat

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

Positive control IHC-P: Human liver tissue. Flow Cyt (intra): HepG2 cells. ICC/IF: HepG2 cells

General notes ab226099 is the carrier-free version of ab124935.

> 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[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] 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.

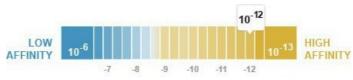
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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Properties

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

Dissociation constant (K_D) $K_D = 8.30 \times 10^{-12} M$



Learn more about K_D

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

ClonalityMonoclonalClone numberEPR5610

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab226099 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
ICC/IF		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 70-75 kDa (predicted molecular weight: 52 kDa).
IP		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
Flow Cyt (Intra)		Use at an assay dependent concentration.

Target

Function Binds heme and transports it to the liver for breakdown and iron recovery, after which the free

hemopexin returns to the circulation.

Tissue specificity Expressed by the liver and secreted in plasma.

Sequence similaritiesBelongs to the hemopexin family.

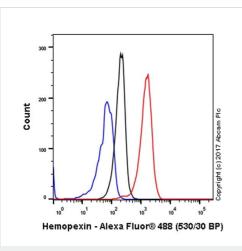
Contains 5 hemopexin-like domains.

Post-translational

modifications

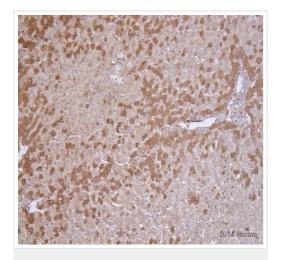
N- and O-glycosylated. O-glycosylated with core 1 or possibly core 8 glycans.

Images



Flow Cytometry (Intracellular) - Anti-Hemopexin antibody [EPR5610] - BSA and Azide free (ab226099) Intracellular Flow Cytometry analysis of HepG2 (Human hepatocellular carcinoma epithelial cell) cells labeling Hemopexin (red) with <u>ab124935</u> at a 1/200 dilution. Cells were fixed with 4% paraformaldehyde and permeabilized with 90% methanol. A goat anti-rabbit lgG (Alexa Fluor[®] 488) (<u>ab150077</u>) was used as the secondary antibody at a 1/2000 dilution. Black - Rabbit monoclonal lgG (Black) (<u>ab172730</u>). Blue (unlabeled control) - Cell without incubation with primary antibody and secondary antibody (Blue).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab124935)



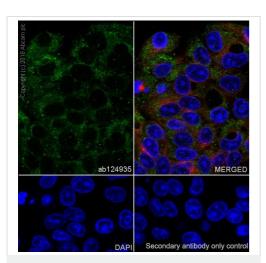
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Hemopexin antibody

[EPR5610] - BSA and Azide free (ab226099)

This image is courtesy of an Abreview submitted by Steffen Rickelt

ab124935 staining Hemopexin in human liver tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffinembedded sections). Tissue was fixed with formaldehyde, permeabilized with 0.2% Triton X-100 in PBS and blocked with 5% milk for 30 minutes at room temperature; antigen retrieval was by heat mediation in Tris pH 9.0. Samples were incubated with primary antibody (1/200 in PBS) for 16 hours at 4°C. An undiluted Biotinconjugated goat anti-rabbit lgG polyclonal was used as the secondary antibody.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab124935).



Immunocytochemistry/ Immunofluorescence - Anti-Hemopexin antibody [EPR5610] - BSA and Azide free (ab226099)

This data was developed using the same antibody clone in a different buffer formulation (<u>ab124935</u>).

Immunocytochemistry analysis of HepG2 (human hepatocellular carcinoma epithelial cell) labeling Hemopexin with purified ${\tt ab124935}$ at 1/200 dilution (10 µg/ml). Cells were fixed with 100% methanol. Goat anti rabbit lgG (Alexa Fluor® 488, ${\tt ab150077}$) at 1/1000 (2 µg/ml) was used as the secondary antibody. ${\tt ab195889}$ Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) 1/200 (2.8 µg/ml) was used as counterstain. Nuclei were stained blue with DAPI.

Negative control: PBS instead of the primary antibody.



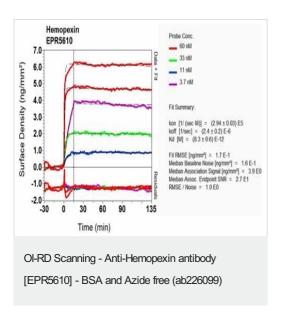
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Hemopexin antibody

[EPR5610] - BSA and Azide free (ab226099)

<u>ab124935</u>, at 1/250 dilution, staining Hemopexin in paraffinembedded Human liver tissue by Immunohistochemistry.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab124935</u>).

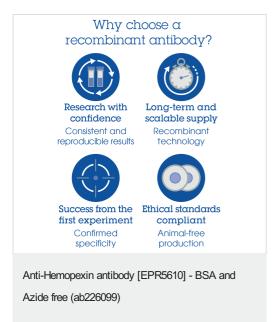
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about K_D

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab124935</u>).



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