abcam

Product datasheet

Anti-Epoxide hydrolase antibody [15B2AD9] ab110307

3 Images

Overview

Product name Anti-Epoxide hydrolase antibody [15B2AD9]

Description Mouse monoclonal [15B2AD9] to Epoxide hydrolase

Host species Mouse

Tested applications Suitable for: IHC-P, Flow Cyt, ICC/IF

Species reactivity Reacts with: Human

Immunogen Tissue, cells or virus. This information is proprietary to Abcam and/or its suppliers.

Positive control Human HDFn cells Human normal colon FFPE tissue

General notes

This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Product was previously marketed under the MitoSciences sub-brand.

Properties

Form Liquid

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

Storage buffer pH: 7.5

Preservative: 0.02% Sodium azide Constituent: HEPES buffered saline

Purity Proprietary Purification

Purification notes The purity of ab110307 is near homogeneity as judged by SDS-PAGE (purity >95%). The

antibody was produced in vitro using hybridomas grown in serum-free medium, and then purified

by biochemical fractionation.

Clonality Monoclonal

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Clone number 15B2AD9

lsotype lgG1 **Light chain type** kappa

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab110307 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
IHC-P		Use a concentration of 5 µg/ml.
Flow Cyt		Use $1\mu g$ for 10^6 cells. $\underline{ab170190}$ - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.
ICC/IF		Use a concentration of 10 µg/ml.

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FunctionBiotransformation enzyme that catalyzes the hydrolysis of arene and aliphatic epoxides to less

reactive and more water soluble dihydrodiols by the trans addition of water.

Tissue specificity Found in liver.

Involvement in diseaseNote=In some populations, the high activity haplotype tyr113/his139 is overrepresented among

women suffering from pregnancy-induced hypertension (pre-eclampsia) when compared with

healthy controls.

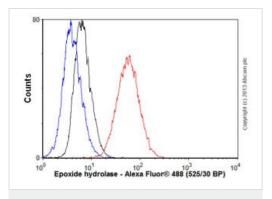
Defects in EPHX1 are a cause of familial hypercholanemia (FHCA) [MIM:607748]. FHCA is a

disorder characterized by elevated serum bile acid concentrations, itching, and fat malabsorption.

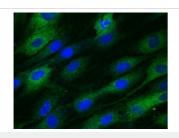
Sequence similarities Belongs to the peptidase S33 family.

Cellular localization Microsome membrane. Endoplasmic reticulum membrane.

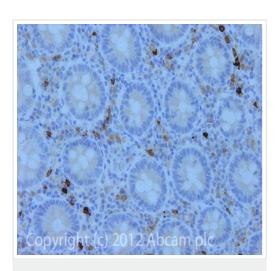
Images



Flow Cytometry - Anti-Epoxide hydrolase antibody [15B2AD9] (ab110307)



Immunocytochemistry/ Immunofluorescence - Anti-Epoxide hydrolase antibody [15B2AD9] (ab110307)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Epoxide hydrolase antibody [15B2AD9] (ab110307)

Overlay histogram showing HepG2 cells stained with ab110307 (red line). The cells were fixed with 4% paraformaldehyde (10 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 (ab110307, 1 μ g/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-mouse lgG (H&L) (ab150113) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG1 [ICIGG1] (ab91353, 1 μ 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.

Immunocytochemistry image of ab110307-stained Human HDFn cells.

The cells were paraformaldehyde fixed (4%, 20 min) and Triton X-100 permeabilized (0.1%, 15 min). The cells were incubated with ab110307 at 4 μ g/ml for 2 hours at room temperature or over night at 4°C. The secondary antibody was (green) Alexa Fluor® 488 goat anti-mouse lgG (H+L) used at a 1/1000 dilution for 1 hour. 10% Goat serum was used as the blocking agent for all blocking steps. DAPI was used to stain the cell nuclei (blue).

Target protein locates mainly in microsomal/ER.

IHC image of Epoxide hydrolase staining in Human normal colon formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab110307, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times

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