abcam

Product datasheet

Anti-Catalase antibody [12C2DB9] - Peroxisome Marker ab110292



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Overview

Product name Anti-Catalase antibody [12C2DB9] - Peroxisome Marker

Description Mouse monoclonal [12C2DB9] to Catalase - Peroxisome Marker

Host species Mouse

Tested applications

Suitable for: ICC, IP, Flow Cyt

Species reactivity

Reacts with: Rat, Cow, Human

Immunogen Tissue, cells or virus. This information is considered to be commercially sensitive.

Positive control HeLa cells, HepG2 cells, HL60 cells, Cow heart, Rat liver Flow cyt: HAP1-WT cells.

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 Ammonium Sulphate Precipitation

Purification notes Purity near homogeneity as judge by SDS-PAGE. The antibody was produced in-vitro using

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hybridomas grown in serum-free medium and then purified by biochemical fractionation.

Clonality Monoclonal
Clone number 12C2DB9

IsotypeIgG1Light chain typekappa

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab110292 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	*****(1)	Use a concentration of 5 µg/ml.
IP		Use at an assay dependent concentration.
Flow Cyt		Use a concentration of 1 µg/ml. <u>ab170190</u> - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.

Target

Function Occurs in almost all aerobically respiring organisms and serves to protect cells from the toxic

effects of hydrogen peroxide. Promotes growth of cells including T-cells, B-cells, myeloid

leukemia cells, melanoma cells, mastocytoma cells and normal and transformed fibroblast cells.

Involvement in disease Defects in CAT are the cause of acatalasia (ACATLAS) [MIM:115500]; also known as

acatalasemia. This disease is characterized by absence of catalase activity in red cells and is

often associated with ulcerating oral lesions.

Sequence similarities Belongs to the catalase family.

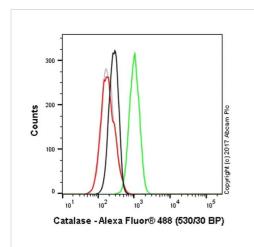
Post-translational

modifications

The N-terminus is blocked.

Cellular localization Peroxisome.

Images

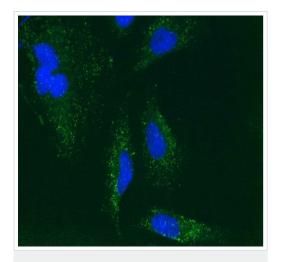


Flow Cytometry - Anti-Catalase antibody [12C2DB9] - Peroxisome Marker (ab110292)

Overlay histogram showing HAP1 wildtype (green line) and HAP1-CAT knockout cells (red line) stained with ab110292. The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab110292, 1µg/ml) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-mouse IgG (H&L) presorbed (ab150117) at 1/2000 dilution for 30 min at 22°C.

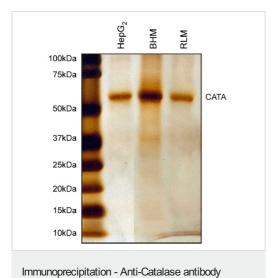
A mouse IgG1 isotype control antibody (ab170190) was used at the same concentration and conditions as the primary antibody (HAP1 wildtype - black line, HAP1-CAT knockout - grey line). Unlabelled sample was also used as a control (this line is not shown for the purpose of simplicity).

Acquisition of >5,000 events were collected using a 50 mW Blue laser (488nm) and 530/30 bandpass filter.



Immunocytochemistry - Anti-Catalase antibody [12C2DB9] - Peroxisome Marker (ab110292)

Immunocytochemical staining of Human HeLa cells. The cells were paraformaldehyde fixed (4%, 20 minutes) and Triton X-100 permeabilized (0.1%, 15 minutes). The cells were incubated with the ab110292 (5 μ g/ml) for 2 hours at room temperature or over night at 4°C. The secondary antibody, Alexa Fluor® 488 goat antimouse lgG (green), was 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). The target protein locates mainly in peroxisomes.



[12C2DB9] - Peroxisome Marker (ab110292)

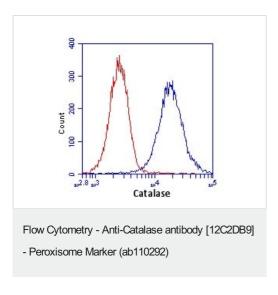
Immunoprecipitation of Catalase using ab110292 from:

Lane 1: HepG2 cells

Lane 2: Cow heart sample

Lane 3: Rat liver sample

Predicted band size is 60 kDa.



Flow cytometric analysis of HL-60 cells using ab110292 at 1 μ g/mL, (blue) or an isotype control (red).

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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