abcam

Product datasheet

Anti-GCLM antibody [EPR6667] ab126704





★★★★★ 2 Abreviews 13 References 7 Images

Overview

Product name Anti-GCLM antibody [EPR6667]

Description Rabbit monoclonal [EPR6667] to GCLM

Host species Rabbit

Suitable for: WB, IP, IHC-P **Tested applications**

Unsuitable for: Flow Cyt

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide within Human GCLM aa 50-150. The exact sequence is proprietary.

Positive control IHC: Rat liver tissue; Mouse cardiac muscle tissue; Human bladder cancer tissue WB: HeLa,

NIH/3T3, PC-12; Wild-type HAP1 whole cell lysate, HeLa cell lysate IP: HeLa cells

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 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.

We are constantly working hard to ensure we provide our customers with best in class antibodies. As a result of this work we are pleased to now offer this antibody in purified format. We are in the process of updating our datasheets. The purified format is designated 'PUR' on our product labels. If you have any questions regarding this update, please contact our Scientific Support team.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

Purity Protein A purified

ClonalityMonoclonalClone numberEPR6667

Isotype IgG

Applications

The Abpromise guarantee Our A

Our Abpromise guarantee covers the use of ab126704 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	★★★★★ (2)	1/1000 - 1/10000. Predicted molecular weight: 31 kDa.
IP		1/10 - 1/100.
IHC-P		1/50 - 1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Application notes Is unsuitable for Flow Cyt.

Target

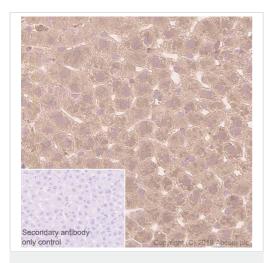
Tissue specificity In all tissues examined. Highest levels in skeletal muscle.

Pathway Sulfur metabolism; glutathione biosynthesis; glutathione from L-cysteine and L-glutamate: step

1/2

Sequence similaritiesBelongs to the aldo/keto reductase family. Glutamate--cysteine ligase light chain subfamily.

Images



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

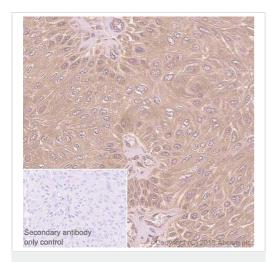
[EPR6667] (ab126704)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Rat liver tissue sections labeling GCLM with purified ab126704 at 1/50 dilution (2.4 µg/mL). Perform heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



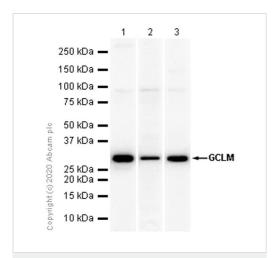
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GCLM antibody
[EPR6667] (ab126704)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Mouse cardiac muscle tissue sections labeling GCLM with purified ab126704 at 1/50 dilution (2.4 µg/mL). Perform heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GCLM antibody
[EPR6667] (ab126704)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human bladder cancer tissue sections labeling GCLM with purified ab126704 at 1/50 dilution (2.4 μ g/mL). Perform heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



Western blot - Anti-GCLM antibody [EPR6667] (ab126704)

Lanes 1 & 3: Anti-GCLM antibody [EPR6667] (ab126704) at 1/10000 dilution (Purified)

Lane 2: Anti-GCLM antibody [EPR6667] (ab126704) at 1/10000 dilution

Lane 1 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

Lane 2 : NIH/3T3 (Mouse embryonic fibroblast) whole cell lysateLane 3 : PC-12 (Rat adrenal gland pheochromocytoma) whole cell

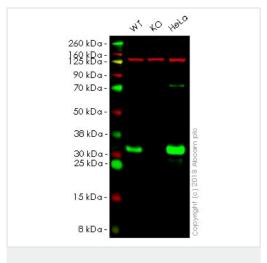
lysate

Secondary

Lanes 1 & 3: Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/20000 dilution

Lane 2: Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/20000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 31 kDa



Western blot - Anti-GCLM antibody [EPR6667] (ab126704)

All lanes: Anti-GCLM antibody [EPR6667] (ab126704) at 1 µg/ml

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: GCLM knockout HAP1 whole cell lysate

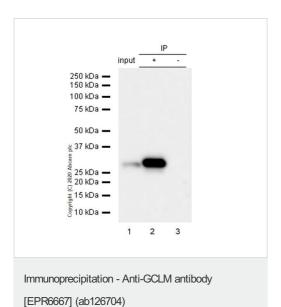
Lane 3: HeLa whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 31 kDa

Lanes 1 - 3: Merged signal (red and green). Green - ab126704 observed at 31 kDa. Red - loading control, ab130007, observed at 130 kDa.

ab126704 was shown to recognize GCLM in wild-type HAP1 cells as signal was lost at the expected MW in GCLM knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and GCLM knockout samples were subjected to SDS-PAGE. Ab126704 and ab130007 (Mouse anti-Vinculin loading control) were incubated overnight at 4°C at 1 µg/ml and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse lgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



Purified ab126704 at 1/20 dilution (0.6µg) immunoprecipitating GCLM in HeLa whole cell lysate.

Lane 1 (input): HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate 10µg

Lane 2 (+): ab126704 + HeLa whole cell lysate.

Lane 3 (-): Rabbit monoclonal IgG (ab172730) instead of ab126704 in HeLa whole cell lysate.

VeriBlot for IP Detection Reagent (HRP) (ab131366) (1/5000 dilution) was used for Western blotting.

Blocking Buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM/TBST.

Observed band size: 31 kDa



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