# abcam

# Product datasheet

# Anti-GCLM antibody [EPR6667] ab126704



★★★★★ 2 Abreviews 33 References 画像数7

#### 製品の概要

製品名 Anti-GCLM antibody [EPR6667]

製品の詳細 Rabbit monoclonal [EPR6667] to GCLM

由来種 Rabbit

アプリケーション 適用あり: WB, IP, IHC-P

適用なし: Flow Cyt

種交差性 交差種: Mouse, Rat, Human

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

ポジティブ・コントロール 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

特記事項 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**.

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

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

精製度 Protein A purified

ポリモノ モノクローナル

クローン名 **EPR6667** 

アイソタイプ lgG

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab126704の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB	*** <u>*</u>	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.

追加情報

Is unsuitable for Flow Cyt.

#### ターゲット情報

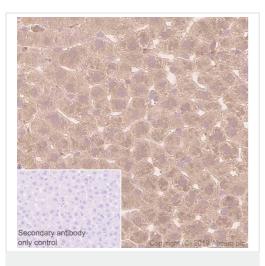
組織特異性 In all tissues examined. Highest levels in skeletal muscle.

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

1/2.

**配列類似性** Belongs to the aldo/keto reductase family. Glutamate--cysteine ligase light chain subfamily.

#### 画像



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 <a href="mailto:ab93684">ab93684</a> (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.



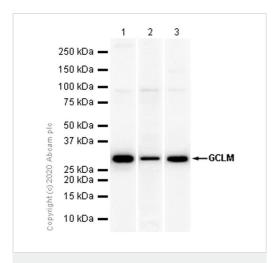
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GCLM antibody
[EPR6667] (ab126704)

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 <a href="mailto:ab93684">ab93684</a> (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.

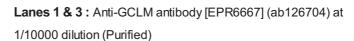


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GCLM antibody
[EPR6667] (ab126704)

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  $\mu$ 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)



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 lysate

Lane 3: PC-12 (Rat adrenal gland pheochromocytoma) whole cell
lysate

## Secondary

Lanes 1 & 3: Goat Anti-Rabbit IgG 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

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

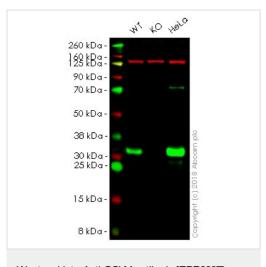
Lane 1: Wild-type HAP1 whole cell lysate

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



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

**Lanes 1 - 3:** Merged signal (red and green). Green - ab126704 observed at 31 kDa. Red - loading control, <u>ab130007</u>, 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 <a href="mailto:ab130007">ab130007</a> (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 <a href="mailto:ab216773">ab216773</a> and Goat anti-Mouse lgG H&L (IRDye® 680RD) preabsorbed <a href="mailto:ab216776">ab216776</a> secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.

Immunoprecipitation - Anti-GCLM antibody

[EPR6667] (ab126704)

Purified ab126704 at 1/20 dilution (0.6 $\mu$ g) 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.

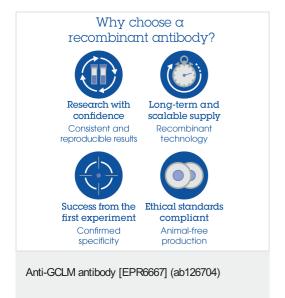
Lane 3 (-): Rabbit monoclonal lgG (<u>ab172730</u>) instead of ab126704 in HeLa whole cell lysate.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) (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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