abcam

Product datasheet

Anti-GLO1 antibody [6F10] ab81461



★★★★★ 3 Abreviews 11 References

画像数4

製品の概要

製品名 Anti-GLO1 antibody [6F10]

製品の詳細 Rat monoclonal [6F10] to GLO1

由来種 Rat

アプリケーション 適用あり: Flow Cyt (Intra), ICC/IF, WB

種交差性 交差種: Mouse, Human, African green monkey

免疫原 Recombinant full length protein. This information is proprietary to Abcam and/or its suppliers.

WB: COS-1, L929 and HeLa whole cell extracts. ICC/IF: Hela cells. HeLa cells: Flow Cyt (Intra). ポジティブ・コントロール

特記事項

ab81461 also reacts with Simian.

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

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid repeated freeze / thaw cycles.

バッファー

Constituents: 50% Glycerol (glycerin, glycerine), PBS

特記事項(精製) The antibody was produced from the hybridoma cultured in serum free medium and purified under

mild conditions by propriety chromatography processes. It is filter sterilized.

一次抗体 備考 ab81461 also reacts with Simian.

ポリ/モノ モノクローナル

クローン名 6F10

アイソタイプIgG2b軽鎖の種類kappa

アプリケーション

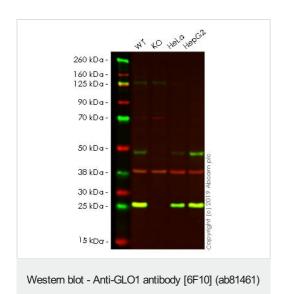
The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab81461の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

| アプリケーション | Abreviews | 特記事項 |
|------------------|-----------|---|
| Flow Cyt (Intra) | | Use 1µg for 10 ⁶ cells. ab18536 - Rat monoclonal lgG2b, is suitable for use as an isotype control with this antibody. |
| ICC/IF | | Use at an assay dependent concentration. |
| WB | | 1/1000. Predicted molecular weight: 21 kDa. |

ターゲット情報

| 機能 | Catalyzes the conversion of hemimercaptal, formed from methylglyoxal and glutathione, to S-lactoylglutathione. |
|-------|--|
| パスウェイ | Secondary metabolite metabolism; methylglyoxal degradation; (R)-lactate from methylglyoxal: step 1/2. |
| 配列類似性 | Belongs to the glyoxalase I family. |

画像



All lanes: Anti-GLO1 antibody [6F10] (ab81461) at 1/1000 dilution

Lane 1 : Wild-type HAP1 whole cell lysate

Lane 2: GLO1 knockout HAP1 whole cell lysate

Lane 3 : HeLa whole cell lysate

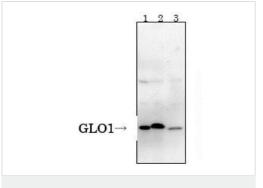
Lane 4 : HepG2 whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 21 kDa

Lanes 1 - 4: Merged signal (red and green). Green - ab81461 observed at 21 kDa. Red - loading control, **ab181602**, observed at 37 kDa.

ab81461 was shown to specifically react with in wild-type HAP1 cells as signal was lost in GLO1 knockout cells. Wild-type and GLO1 knockout samples were subjected to SDS-PAGE. The membrane was blocked with 3% Milk. Ab81461 and ab181602 (Rabbit anti-GAPDH loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/20000 dilution respectively. Blots were developed with Goat anti-Rat lgG H&L (IRDye® 800CW) preabsorbed and Goat anti-Rabbit lgG H&L (IRDye® 680RD) preabsorbed ab216777 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



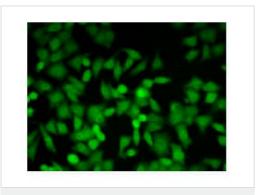
All lanes: Anti-GLO1 antibody [6F10] (ab81461)

Lane 1 : COS-1 (simian) whole cell extract
Lane 2 : L929 (mouse) whole cell extract
Lane 3 : HeLa (human) whole cell extract

Predicted band size: 21 kDa **Observed band size:** 27-29 kDa

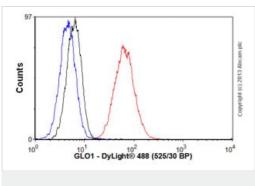
Western blot - Anti-GLO1 antibody [6F10] (ab81461)

Mouse GLO1 shows a single band of 27 kDa while human and simian ones show 29 kDa.



Immunofluorescent staining of HeLa cells with ab81461.

Immunocytochemistry/ Immunofluorescence - Anti-GLO1 antibody [6F10] (ab81461)



Flow Cytometry (Intracellular) - Anti-GLO1 antibody [6F10] (ab81461)

Overlay histogram showing HeLa cells stained with ab81461 (red line). The cells were fixed with 80% methanol (5 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 (ab81461, 1µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-rat lgG (H+L) (ab98386) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was rat lgG2b [RTK4530] (ab18541, 1µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line). Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.

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