abcam

Product datasheet

Anti-HGD antibody [EPR7874] ab131035

ועלשעבע RabMAb

2 References 画像数4

製品の概要

製品名 Anti-HGD antibody [EPR7874]

製品の詳細 Rabbit monoclonal [EPR7874] to HGD

由来種 Rabbit

アプリケーション 適用あり: WB, IHC-P, ICC/IF

適用なし: Flow Cyt

交差種: Human 種交差性

免疫原 Synthetic peptide within Human HGD aa 400-500 (C terminal). The exact sequence is proprietary.

ポジティブ・コントロール Caco-2, HepG2, and Human fetal liver lysates; Human kidney tissue; HepG2 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**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

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

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

精製度 Protein A purified

ポリモノ モノクローナル

クローン名 EPR7874

アイソタイプ IgG

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ab131035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		1/1000 - 1/10000. Detects a band of approximately 50 kDa (predicted molecular weight: 50 kDa).
IHC-P		1/100 - 1/250. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
ICC/IF		1/100 - 1/250.

追加情報 Is unsuitable for Flow Cyt.

ターゲット情報

組織特異性 Highest expression in the prostate, small intestine, colon, kidney and liver.

אליבוֹל Amino-acid degradation; L-phenylalanine degradation; acetoacetate and fumarate from L-

phenylalanine: step 4/6.

関連疾患 Defects in HGD are the cause of alkaptonuria (AKU) [MIM:203500]. AKU is an autosomal

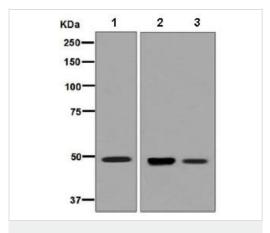
recessive error of metabolism characterized by an increase in the level of homogentisic acid. The

clinical manifestations of AKU are urine that turns dark on standing and alkalinization, black

ochronotic pigmentation of cartilage and collagenous tissues, and spine arthritis.

配列類似性 Belongs to the homogentisate dioxygenase family.

画像



Western blot - Anti-HGD antibody [EPR7874] (ab131035)

All lanes: Anti-HGD antibody [EPR7874] (ab131035) at 1/1000

dilution

Lane 1 : Caco-2 cell lysate

Lane 2 : HepG2 cell lysate

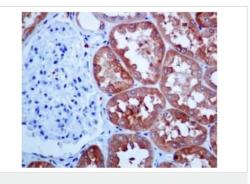
Lane 3: Human fetal liver tissue lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

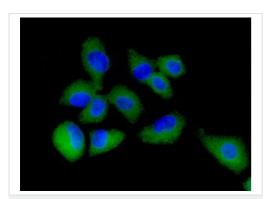
Predicted band size: 50 kDa **Observed band size:** 50 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HGD antibody [EPR7874] (ab131035)

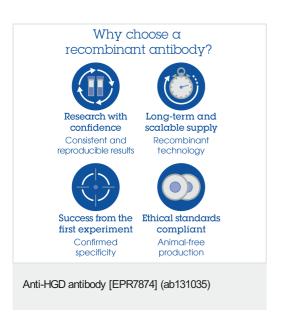
Immunohistochemical analysis of paraffin-embedded Human kidney tissue labelling HGD with ab131035 antibody at 1/100 dilution.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunocytochemistry/ Immunofluorescence - Anti-HGD antibody [EPR7874] (ab131035)

Immunofluorescence analysis of HepG2 cells labelling HGD with ab131035 antibody at 1/100 dilution.



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