# abcam

# Product datasheet

# Anti-GUCY1B3 antibody [EPR8822] - BSA and Azide free ab232491

אלבעבע RabMAb

画像数7

#### 製品の概要

製品名 Anti-GUCY1B3 antibody [EPR8822] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR8822] to GUCY1B3 - BSA and Azide free

由来種 Rabbit

特異性 The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for

mouse and rat.

アプリケーション 適用あり: WB, IHC-P

種交差性 交差種: Mouse, Rat, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール IHC-P: Human kidney tissue.

特記事項 ab232491 is the carrier-free version of ab154841.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for

increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.

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

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

**バッファー** pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

**ポリ/モノ** モノクローナル

**クローン名** EPR8822

アイソタイプ lgG

# アプリケーション

**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ab232491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		Use at an assay dependent concentration. Predicted molecular weight: 71 kDa.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.  We do not guarantee IHC-P for mouse and rat.

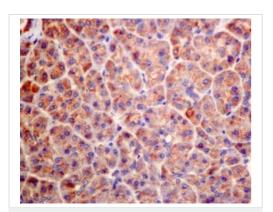
#### ターゲット情報

**配列類似性** Belongs to the adenylyl cyclase class-4/guanylyl cyclase family.

Contains 1 guanylate cyclase domain.

細胞内局在 Cytoplasm.

### 画像



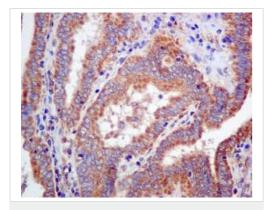
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GUCY1B3 antibody

[EPR8822] - BSA and Azide free (ab232491)

Immunohistochemical analysis of formalin-fixed, paraffin-embedded Human pancreas tissue, labeling GUCY1B3 using unpurified <a href="mailto:ab154841">ab154841</a> at a 1/100 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab154841).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



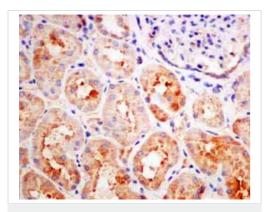
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GUCY1B3 antibody

[EPR8822] - BSA and Azide free (ab232491)

Immunohistochemical analysis of formalin-fixed, paraffin-embedded Human thyroid gland carcinoma tissue, labeling GUCY1B3 using unpurified <u>ab154841</u> at a 1/100 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab154841</u>).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



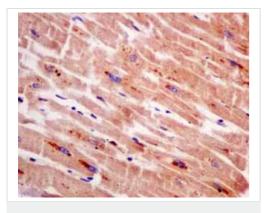
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GUCY1B3 antibody

[EPR8822] - BSA and Azide free (ab232491)

Immunohistochemical analysis of paraffin embedded Human normal kidney tissue using unpurified <u>ab154841</u> showing +ve staining.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab154841</u>).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



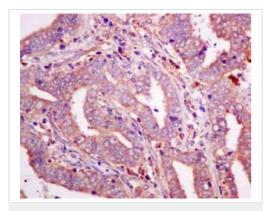
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GUCY1B3 antibody

[EPR8822] - BSA and Azide free (ab232491)

Immunohistochemical analysis of paraffin embedded Human heart tissue using unpurified <u>ab154841</u> showing +ve staining.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab154841).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



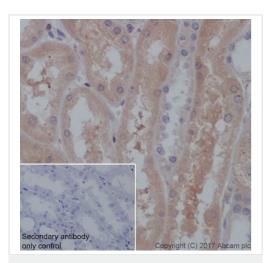
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GUCY1B3 antibody

[EPR8822] - BSA and Azide free (ab232491)

Immunohistochemical analysis of paraffin embedded Human endometrial carcinoma tissue using unpurified <u>ab154841</u> showing +ve staining.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab154841</u>).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.

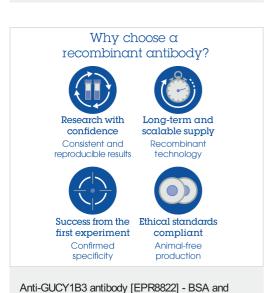


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GUCY1B3 antibody

[EPR8822] - BSA and Azide free (ab232491)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue sections labeling GUCY1B3 with purified <u>ab154841</u> at 1:400 dilution (2.54 µg/ml). Heat mediated antigen retrieval was performed using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0). Tissue was counterstained with Hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab154841).



Azide free (ab232491)

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- Extensive multi-media technical resources to help you

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