# abcam

### Product datasheet

## Alexa Fluor® 488 Anti-UCH37 antibody [EPR4896] ab223411

יעלאעבע RabMAb

#### 画像数 2

製品名 Alexa Fluor® 488 Anti-UCH37 antibody [EPR4896]

製品の詳細 Alexa Fluor® 488 Rabbit monoclonal [EPR4896] to UCH37

由来種 Rabbit

Alexa Fluor® 488. Ex: 495nm, Em: 519nm

適用あり: IHC-Fr

種交差性 交差種: Rat

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

ポジティブ・コントロール IHC-Fr: normal rat kidney tissue sections

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity

交差が予測される動物種: Human 🔷

- 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**.

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製品の概要

標識

アプリケーション

免疫原

特記事項

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Store In the Dark.

**バッファー** pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 1% BSA, 30% Glycerol (glycerin, glycerine), PBS

精製度 Protein A purified

**ポリモ**ノクローナル **ウローン名** EPR4896

アイソタイプ lgG

#### 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ab223411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
IHC-Fr		1/1000.

#### ターゲット情報

機能 Protease that specifically cleaves 'Lys-48'-linked polyubiquitin chains. Deubiquitinating enzyme

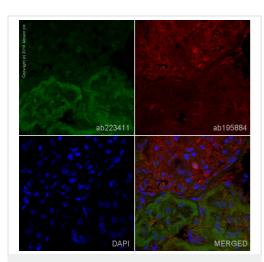
associated with the 19S regulatory subunit of the 26S proteasome.

**配列類似性** Belongs to the peptidase C12 family.

細胞内局在 Cytoplasm. Nucleus. Associates with the proteasome 19S subunit in the cytoplasm. Associates

with the INO80 complex in the nucleus.

#### 画像



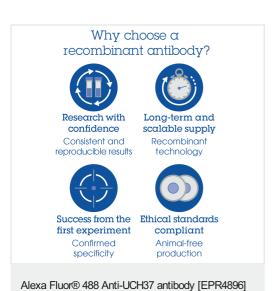
Immunohistochemistry (Frozen sections) - Alexa Fluor® 488 Anti-UCH37 antibody [EPR4896] (ab223411)

IHC image of UCH37 staining in a section of frozen normal rat kidney.

The section was fixed using 10% formaldehyde in 1XPBS for 10 minutes. No antigen retrieval step was performed prior to staining. Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab223411 at 1/1000 dilution (shown in green) and counterstained using **ab195884**, Rat monoclonal to Tubulin (Alexa Fluor® 647), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue). The section was then mounted using Fluoromount®.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

For other IHC staining systems (automated and non-automated), customers should optimize variable parameters such as antigen retrieval conditions, antibody concentrations and incubation times.



(ab223411)

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