# abcam

### Product datasheet

## Anti-hnRNP C1/C2 antibody [EP3034Y] ab75822

ועלשעבע RabMAb

2 References 画像数5

#### 製品の概要

製品名 Anti-hnRNP C1/C2 antibody [EP3034Y]

製品の詳細 Rabbit monoclonal [EP3034Y] to hnRNP C1/C2

由来種 Rabbit

アプリケーション 適用あり: WB, IHC-P

適用なし: Flow Cyt or IP

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 WB: Mouse and Rat brain lysate, HeLa, and MCF7 lysate. IHC-P: Human breast cancer tissue,

Mouse and Rat testis.

特記事項 This product has switched from a hybridoma to recombinant production method on 23rd February

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

精製度 Protein A purified

モノクローナル ポリモノ

**クローン名** EP3034Y

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ab75822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		1/10000. Detects a band of approximately 34 kDa (predicted molecular weight: 34 kDa).
IHC-P		1/800. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.

追加情報 Is unsuitable for Flow Cyt or IP.

#### ターゲット情報

機能 Binds pre-mRNA and nucleates the assembly of 40S hnRNP particles (PubMed:8264621).

Specifically recognizes and binds N6-methyladenosine (m6A)-containing RNAs, a modification present at internal sites of mRNAs that affects mRNA splicing, processing and stability. M6A alters the local structure in mRNAs and long non-coding RNAs (lncRNAs) via a mechanism named 'm(6)A-switch' to facilitate binding of HNRNPC, leading to regulation of mRNA splicing (PubMed:25719671). Single HNRNPC tetramers bind 230-240 nucleotides. Trimers of HNRNPC tetramers bind 700 nucleotides. May play a role in the early steps of spliceosome assembly and pre-mRNA splicing. Interacts with poly-U tracts in the 3'-UTR or 5'-UTR of mRNA and modulates

PubMed:16010978, PubMed:7567451, PubMed:8264621).

**配列類似性** Belongs to the RRM HNRPC family. RALY subfamily.

Contains 1 RRM (RNA recognition motif) domain.

翻訳後修飾 Phosphorylated on Ser-260 and Ser-299 in resting cells. Phosphorylated on Ser-253 and on 1

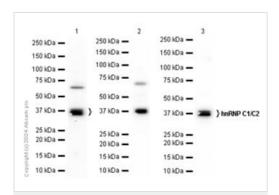
serine residue in the poly-Ser stretch at position 238 in response to hydrogen peroxide.

the stability and the level of translation of bound mRNA molecules (PubMed:12509468,

Sumoylated. Sumoylation reduces affinity for mRNA.

**細胞内局在** Nucleus. Component of ribonucleosomes.

#### 画像



Western blot - Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822)

**All lanes**: Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822) at 1/10000 dilution

Lane 1 : Mouse brain lysate

Lane 2 : Rat brain lysate

Lane 3 : HeLa (Human cervix adenocarcinoma epithelial cell)

whole lysate

Lysates/proteins at 15 µg per lane.

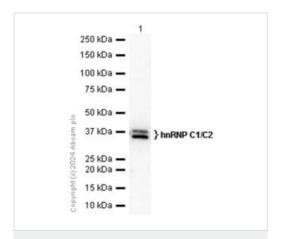
#### Secondary

**All lanes :** Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

**Predicted band size:** 34 kDa **Observed band size:** 34 kDa

Blocking and diluting buffer and concentration: 5% NFDM/TBST.

Exposure time: Lane 1, 60 seconds; Lane 2, 5 seconds; Lane 3, 3 seconds



Western blot - Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822)

Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822) at 1/10000 dilution + MCF7(Human breast adenocarcinoma epithelial cell) whole lysate at 15  $\mu g$ 

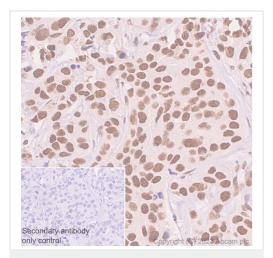
#### Secondary

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 34 kDa Observed band size: 34 kDa

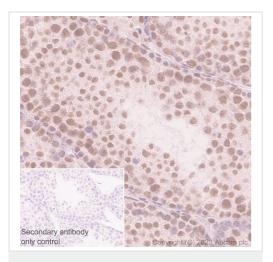
Exposure time: 180 seconds

Blocking and diluting buffer and concentration: 5% NFDM/TBST.



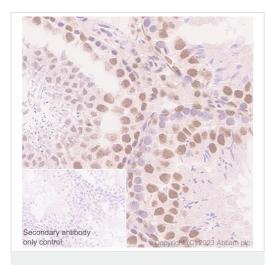
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822)

Immunohistochemical analysis of paraffin-embedded Human breast cancer labeling hnRNP C1/C2 with ab75822 at 1/800 dilution (0.589 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). The section was incubated with ab75822 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822)

Immunohistochemical analysis of paraffin-embedded Mouse testis tissue labeling hnRNP C1/C2 with ab75822 at 1/800 dilution (0.589 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). The section was incubated with ab75822 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-hnRNP C1/C2 antibody [EP3034Y] (ab75822)

Immunohistochemical analysis of paraffin-embedded Rat testis tissue labeling hnRNP C1/C2 with ab75822 at 1/800 dilution (0.589 µg/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). The section was incubated with ab75822 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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