

Anti-HuD + HuC antibody [EPR19098] ab184267

リコンビナント **RabMAb**

★★★★★ **2 Abreviews** **9 References** 画像数 23

製品の概要

製品名	Anti-HuD + HuC antibody [EPR19098]
製品の詳細	Rabbit monoclonal [EPR19098] to HuD + HuC
由来種	Rabbit
特異性	This antibody shows very weak cross reactivity to HuB in WB test. Please contact our Scientific support team for more information.
アプリケーション	適用あり: Flow Cyt (Intra), Indirect ELISA, WB, IHC-P, IHC-Fr, ICC/IF, IP
種交差性	交差種: Mouse, Rat, Human
免疫原	Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.
ポジティブ・コントロール	WB: Mouse His-tagged HuC and His-GST-tagged HuD recombinant proteins; Human fetal brain; Mouse and rat brain; Neuro-2a and SH-SY5Y cell lysates. IHC-P: Human cerebral cortex, colon and glioma tissues; Mouse cerebral cortex and stomach tissues; Rat cerebral cortex and colon tissues. IHC-Fr: Mouse cerebral cortex and stomach tissues; Rat cerebral cortex and colon tissues. ICC/IF: SH-SY5Y and Neuro-2a cells. Flow Cyt: SH-SY5Y and Neuro-2a cells. IP: Mouse brain lysate. I-ELISA: Mouse Elavl3 and 4.
特記事項	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb[®] patents.</p>

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 long term. Avoid freeze / thaw cycle.
バッファー	pH: 7.2 Preservative: 0.01% Sodium azide

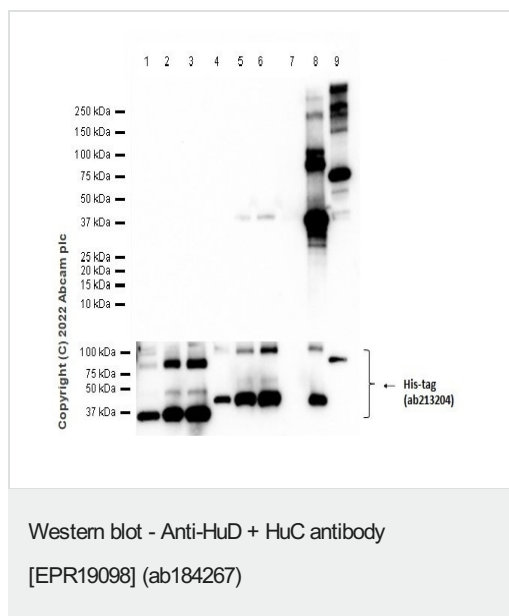
	Constituents: PBS, 40% Glycerol, 0.05% BSA
精製度	Protein A purified
ポリ/モノ	モノクローナル
クローン名	EPR19098
アイソタイプ	IgG

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ab184267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)		1/500.
Indirect ELISA		Use a concentration of 1 µg/ml.
WB		1/1000. Detects a band of approximately 39 kDa (predicted molecular weight: 40 kDa).
IHC-P		1/4000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
IHC-Fr		1/500. Antigen retrieval: Heated citrate solution (10mM citrate pH 6.0 + 0.05% Tween-20).
ICC/IF		1/500.
IP		1/30.

画像



All lanes : Anti-HuD + HuC antibody [EPR19098] (ab184267) at 1/1000 dilution

Lane 1 : His-tagged Mouse HUR recombinant protein (aa1-326) at 0.01 µg

Lane 2 : His-tagged Mouse HUR recombinant protein (aa1-326) at 0.05 µg

Lane 3 : His-tagged Mouse HUR recombinant protein (aa1-326) at 0.1 µg

Lane 4 : His-tagged mouse HuB recombinant protein (aa1-360) at 0.01 µg

Lane 5 : His-tagged mouse HuB recombinant protein (aa1-360) at 0.05 µg

Lane 6 : His-tagged mouse HuB recombinant protein (aa1-360) at 0.1 µg

Lane 7 : Blank

Lane 8 : His-tagged mouse HuC recombinant protein (aa1-367) at 0.01 µg

Lane 9 : His-GST-tagged mouse HuD recombinant protein (aa1-385) at 0.01 µg

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/20000 dilution

Predicted band size: 40 kDa

Observed band size: 37,75 kDa

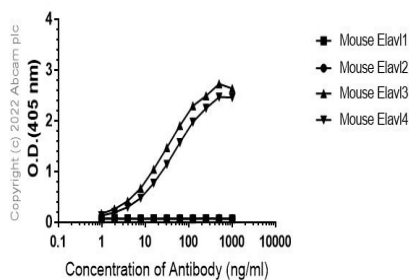
Exposure time: 10 seconds

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

ab184267 shows very weak cross reactivity with HuB.

Band around 80kda detected by anti-His tag antibody should be dimer.

Indirect ELISA antibody dose-response curve
antigen at 1000 ng/ml



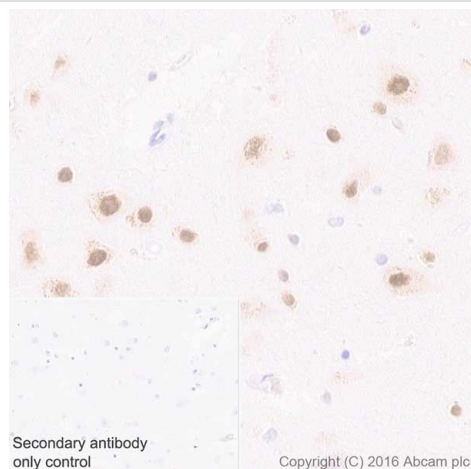
Indirect ELISA - Anti-HuD + HuC antibody
[EPR19098] (ab184267)

Indirect ELISA showing primary antibody ab184267 binding to Mouse Elavl1 (HuR); Mouse Elavl2 (HuB); Mouse Elavl3 (HuC); Mouse Elavl4 (HuD). Antigen concentration is 1000 ng/ml.

Substrate solution is p-nitrophenyl phosphate(PNPP).

Binding of ab184267 was assessed in a serial dilution range 1000-0 ng/ml.

Binding was detected using the secondary antibody, Alkaline Phosphatase-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L) at 1/2500 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody
[EPR19098] (ab184267)

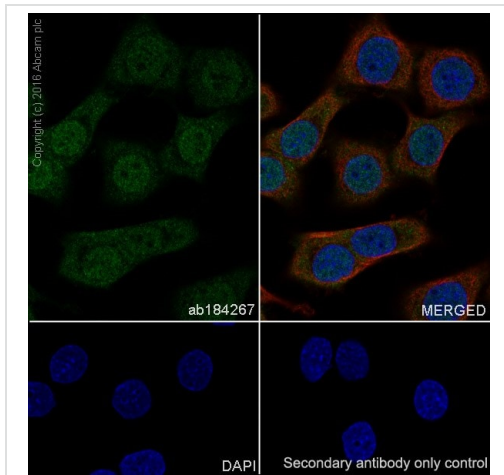
Immunohistochemical analysis of paraffin-embedded human cerebral cortex tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Nuclear and weakly cytoplasmic staining on neurons of human cerebral cortex is observed [PMID: 22007133].

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunocytochemistry/ Immunofluorescence - Anti-HuD + HuC antibody [EPR19098] (ab184267)

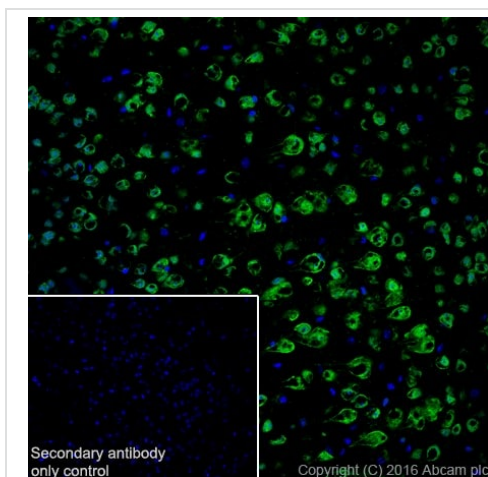
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized SH-SY5Y (Human neuroblastoma cell line from bone marrow) cells labeling HuC + HuD with ab184267 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor[®] 488) (**ab150077**) secondary antibody at 1/1000 dilution (green).

Confocal image showing cytoplasmic and nuclear staining on SH-SY5Y cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with **ab195889** (Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594)) at 1/200 dilution (red).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor[®] 488) (**ab150077**) at 1/1000 dilution.



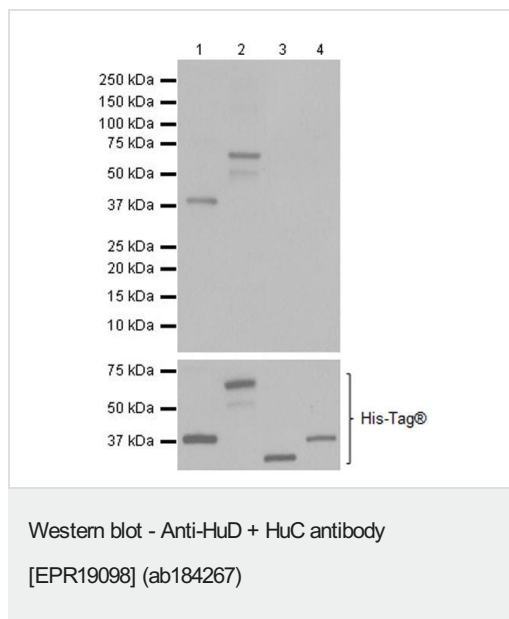
Immunohistochemistry (Frozen sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen mouse cerebral cortex tissue labeling HuC + HuD with ab184267 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor[®] 488) (**ab150077**) secondary antibody at 1/1000 dilution (green).

Cytoplasmic and nuclear staining on neurons of mouse cerebral cortex is observed.

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor[®] 488) (**ab150077**) at 1/1000 dilution.



All lanes : Anti-HuD + HuC antibody [EPR19098] (ab184267) at 1/2000 dilution

Lane 1 : His-tagged mouse HuC recombinant protein (aa1-367)

Lane 2 : His-GST-tagged mouse HuD recombinant protein (aa1-385)

Lane 3 : His-tagged mouse HuR recombinant protein (aa1-360)

Lane 4 : His-tagged mouse HuB recombinant protein (aa1-360)

Lysates/proteins at 0.01 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/100000 dilution

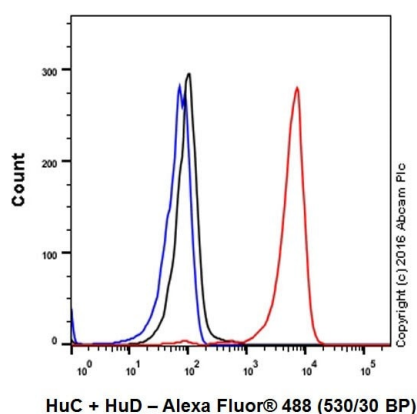
Predicted band size: 40 kDa

Observed band size: 39,68 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

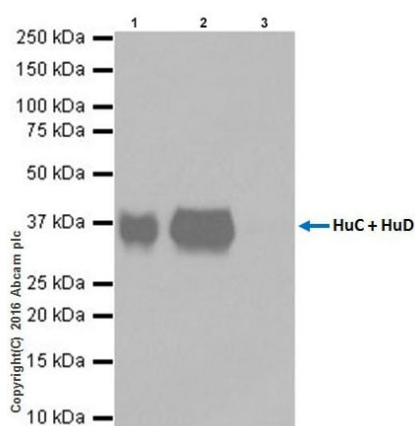
Exposure time: 0.5 second.

Please note that the band around 55kDa in lane 2 is most likely the HuD protein with tags.



Flow Cytometry (Intracellular) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed SH-SY5Y (Human neuroblastoma cell line from bone marrow) cells labeling HuC + HuD with ab184267 at 1/500 dilution (red) compared with a rabbit monoclonal IgG isotype control (**ab172730**; black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody; blue). Goat anti rabbit IgG (Alexa Fluor® 488) at 1/2000 dilution was used as the secondary antibody.



Immunoprecipitation - Anti-HuD + HuC antibody [EPR19098] (ab184267)

HuC + HuD was immunoprecipitated from 0.35 mg of mouse brain lysate with ab184267 at 1/30 dilution.

Western blot was performed from the immunoprecipitate using ab184267 at 1/1000 dilution.

VeriBlot for IP Detection Reagent (HRP) (**ab131366**), was used for detection at 1/10000 dilution.

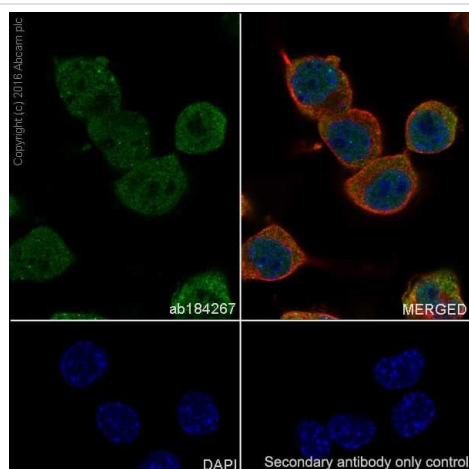
Lane 1: Mouse brain lysate, 10 µg (Input).

Lane 2: ab184267 IP in mouse brain lysate.

Lane 3: Rabbit monoclonal IgG (**ab172730**) instead of ab184267 in mouse brain lysate.

Blocking and dilution buffer and concentration: 5% NFDm/TBST.

Exposure time: 30 seconds.



Immunocytochemistry/ Immunofluorescence - Anti-HuD + HuC antibody [EPR19098] (ab184267)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized Neuro-2a (Mouse neuroblastoma cell line) cells labeling HuC + HuD with ab184267 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (**ab150077**) secondary antibody at 1/1000 dilution (green).

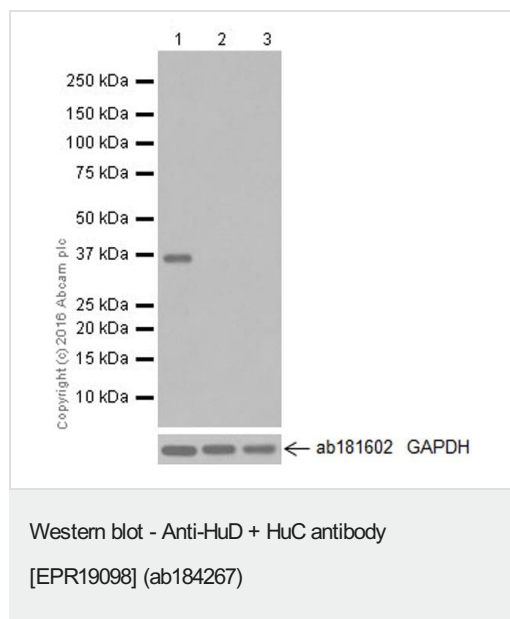
Confocal image showing cytoplasmic and nuclear staining on Neuro-2a cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with **ab195889** (Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594)) at 1/200 dilution (red).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor®

488) ([ab150077](#)) at 1/1000 dilution.



All lanes : Anti-HuD + HuC antibody [EPR19098] (ab184267) at 1/1000 dilution

Lane 1 : Human fetal brain lysate

Lane 2 : Human fetal heart lysate

Lane 3 : Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : VeriBlot for IP Detection Reagent (HRP) ([ab131366](#)) at 1/4000 dilution

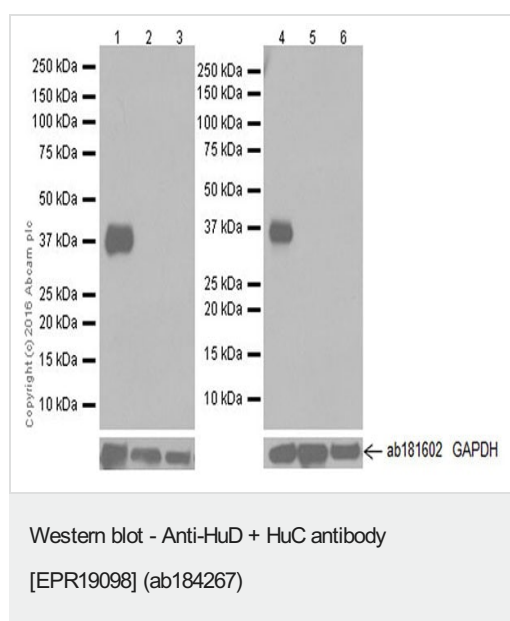
Predicted band size: 40 kDa

Observed band size: 39 kDa

Exposure time: 8 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

The expression profile is consistent with what has been described in the literature (PMID: 9016658; PMID:8535975).



All lanes : Anti-HuD + HuC antibody [EPR19098] (ab184267) at 1/1000 dilution

Lane 1 : Mouse brain lysate

Lane 2 : Mouse heart lysate

Lane 3 : Mouse kidney lysate

Lane 4 : Rat brain lysate

Lane 5 : Rat heart lysate

Lane 6 : Rat kidney lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/100000 dilution

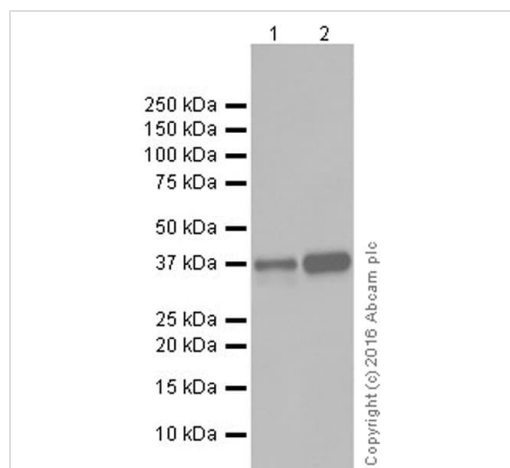
Predicted band size: 40 kDa

Observed band size: 39 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: Lane 1-3: 1 minute; Lane 4-6: 5 seconds.

The expression profile is consistent with what has been described in the literature (PMID: 9016658; PMID: 8535975).



Western blot - Anti-HuD + HuC antibody
[EPR19098] (ab184267)

All lanes : Anti-HuD + HuC antibody [EPR19098] (ab184267) at 1/5000 dilution

Lane 1 : Neuro-2a (Mouse neuroblastoma cell line) whole cell lysate

Lane 2 : SH-SY5Y (Human neuroblastoma cell line from bone marrow) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

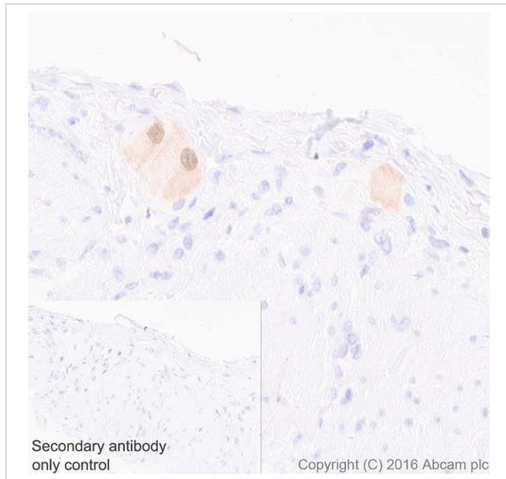
All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/100000 dilution

Predicted band size: 40 kDa

Observed band size: 39 kDa

Exposure time: 5 seconds

Blocking/Dilution Buffer: 5% NFDM/TBST



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

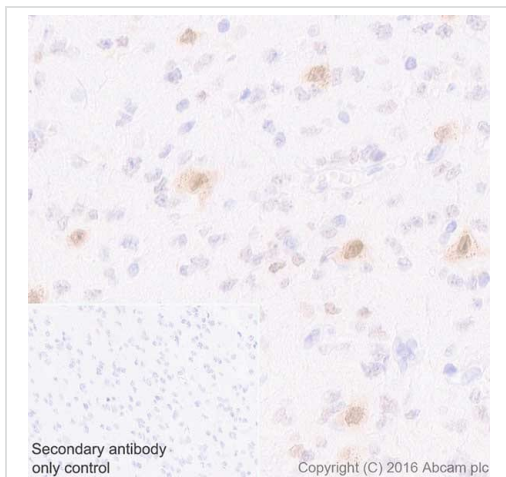
Immunohistochemical analysis of paraffin-embedded human colon tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Nuclear and weakly cytoplasmic staining on myenteric ganglia of human colon is observed [PMID: 16918730].

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

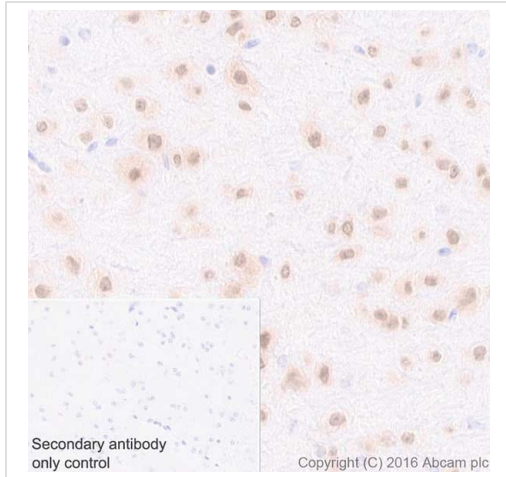
Immunohistochemical analysis of paraffin-embedded human glioma tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Nuclear and weakly cytoplasmic staining on human glioma is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

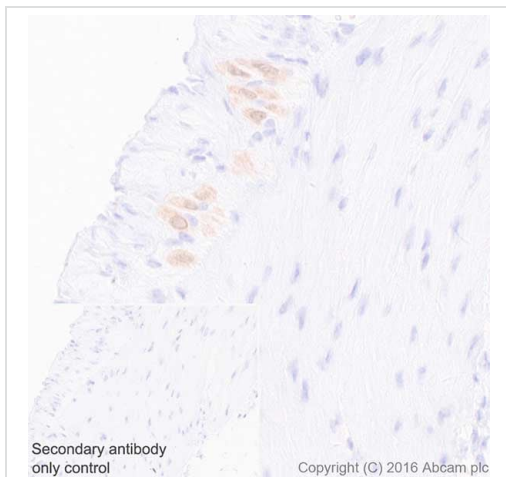
Immunohistochemical analysis of paraffin-embedded mouse cerebral cortex tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Nuclear and weakly cytoplasmic staining on neurons of mouse cerebral cortex is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

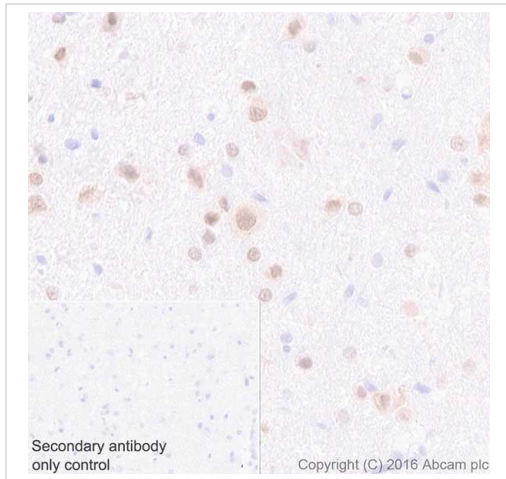
Immunohistochemical analysis of paraffin-embedded mouse stomach tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Cytoplasmic and nuclear staining on myenteric ganglia of mouse stomach is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

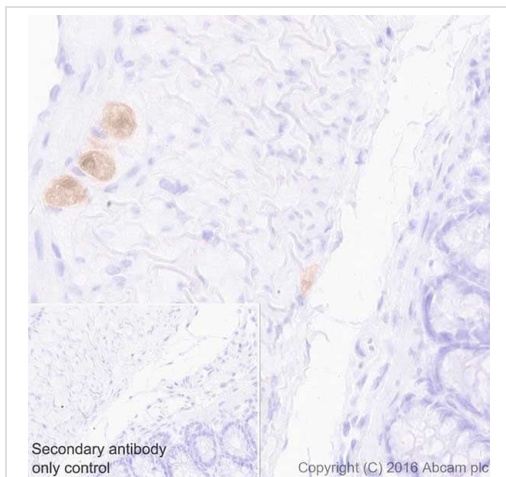
Immunohistochemical analysis of paraffin-embedded rat cerebral cortex tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Nuclear and weakly cytoplasmic staining on neurons of rat cerebral cortex is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

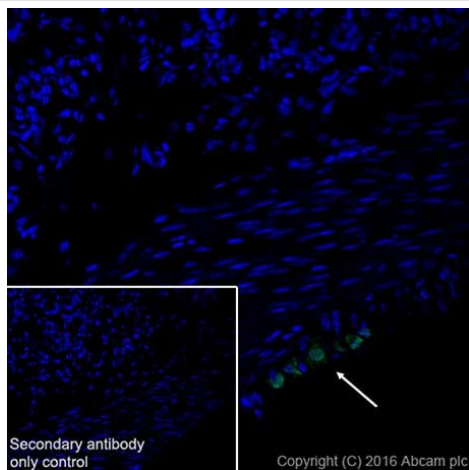
Immunohistochemical analysis of paraffin-embedded rat colon tissue labeling HuC + HuD with ab184267 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Cytoplasmic and nuclear staining on myenteric ganglia of rat colon is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



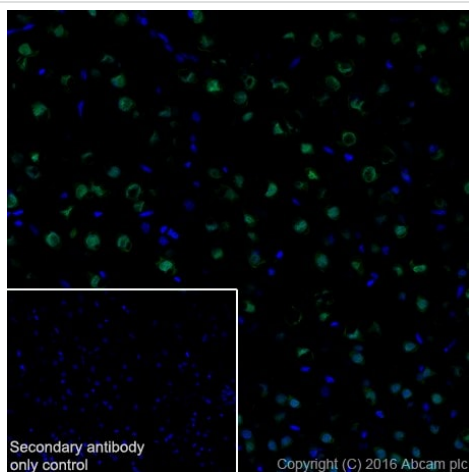
Immunohistochemistry (Frozen sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen mouse stomach tissue labeling HuC + HuD with ab184267 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) ([ab150077](#)) secondary antibody at 1/1000 dilution (green).

Cytoplasmic and nuclear staining on myenteric ganglia of mouse stomach is observed (white arrow).

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor® 488) ([ab150077](#)) at 1/1000 dilution.



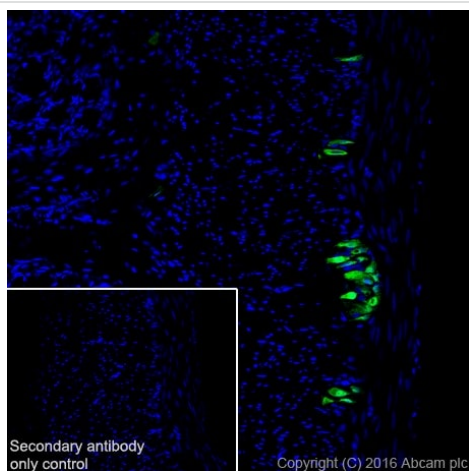
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Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen rat cerebral cortex tissue labeling HuC + HuD with ab184267 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) ([ab150077](#)) secondary antibody at 1/1000 dilution (green).

Cytoplasmic and nuclear staining on neurons of rat cerebral cortex is observed.

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor® 488) ([ab150077](#)) at 1/1000 dilution.



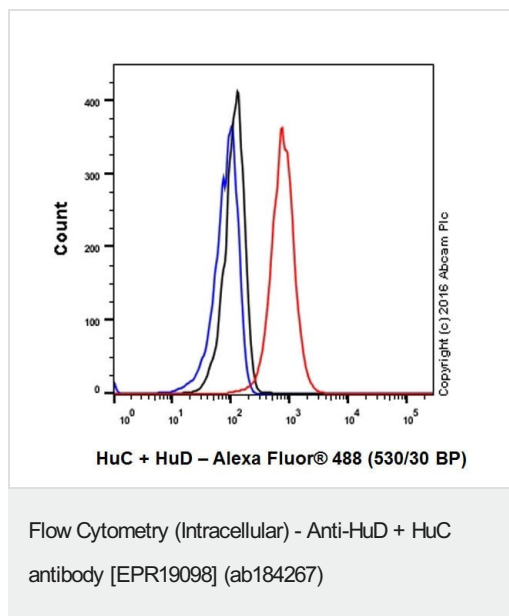
Immunohistochemistry (Frozen sections) - Anti-HuD + HuC antibody [EPR19098] (ab184267)

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen rat colon tissue labeling HuC + HuD with ab184267 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) ([ab150077](#)) secondary antibody at 1/1000 dilution (green).

Cytoplasmic and nuclear staining on myenteric ganglia of rat colon is observed.





The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor® 488) ([ab150077](#)) at 1/1000 dilution.



Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed Neuro-2a (Mouse neuroblastoma cell line) cells labeling HuC + HuD with ab184267 at 1/500 dilution (red) compared with a rabbit monoclonal IgG isotype control (**ab172730**; black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody; blue). Goat anti rabbit IgG (Alexa Fluor® 488) at 1/2000 dilution was used as the secondary antibody.

Why choose a recombinant antibody?

 <p>Research with confidence Consistent and reproducible results</p>	 <p>Long-term and scalable supply Recombinant technology</p>
 <p>Success from the first experiment Confirmed specificity</p>	 <p>Ethical standards compliant Animal-free production</p>

Anti-HuD + HuC antibody [EPR19098] (ab184267)

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