abcam

Product datasheet

Anti-DCTN1/p150-glued antibody [EPR26464-28] ab302629



重组 RabMAb

15 图像

概述

产品名称 Anti-DCTN1/p150-glued抗体[EPR26464-28]

描述 兔单克隆抗体[EPR26464-28] to DCTN1/p150-glued

宿主 Rabbit

特异性 IHC application not suitable for human and rat species.

经测试应用 适用于: WB, IHC-P, IHC-Fr, ICC/IF, Flow Cyt (Intra), IP

种属反应性 与反应: Mouse, Human

不与反应: Rat

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

阳性对照 WB: HeLa transfected with siRNA specifically targeting DCTN1 and a scrambled siRNA control,

> Neuro-2a, and NIH/3T3 whole cell lysates, mouse brain and cerecbellum tissue lysates. IHC-P: Mouse cerebrum, spinal cord, and dorsal root ganglion FFPE tissue sections. IHC-Fr: Mouse spinal cord fresh frozen tissue section. ICC/IF: HeLa and Neuro-2a cell lines. Flow Cyt (Intra):

HeLa and Neuro-2a cell lines. IP: HeLa and Neuro-2a whole cell lysates.

常规说明 This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

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**.

性能

形式 Liquid

存放说明 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.

存储溶液 pH: 7.20

Preservative: 0.01% Sodium azide

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

纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR26464-28

同种型 IgG

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab302629于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度:实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		1/1000. Detects a band of approximately 150 kDa (predicted molecular weight: 141 kDa).
IHC-P		1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
IHC-Fr		1/100.
ICC/IF		1/50.
Flow Cyt (Intra)		1/50.
IP		1/30.

靶标

功能

Required for the cytoplasmic dynein-driven retrograde movement of vesicles and organelles along microtubules. Dynein-dynactin interaction is a key component of the mechanism of axonal transport of vesicles and organelles.

组织特异性

疾病相关

Brain.

Defects in DCTN1 are the cause of distal hereditary motor neuronopathy type 7B (HMN7B) [MIM:607641]; also known as progressive lower motor neuron disease (PLMND). HMN7B is a neuromuscular disorder. Distal hereditary motor neuronopathies constitute a heterogeneous group of neuromuscular disorders caused by selective degeneration of motor neurons in the anterior horn of the spinal cord, without sensory deficit in the posterior horn. The overall clinical picture consists of a classical distal muscular atrophy syndrome in the legs without clinical sensory loss. The disease starts with weakness and wasting of distal muscles of the anterior tibial and peroneal compartments of the legs. Later on, weakness and atrophy may expand to the proximal muscles of the lower limbs and/or to the distal upper limbs.

Defects in DCTN1 are a cause of susceptibility to amyotrophic lateral sclerosis (ALS) [MIM:105400]. ALS is a neurodegenerative disorder affecting upper and lower motor neurons, and resulting in fatal paralysis. Sensory abnormalities are absent. Death usually occurs within 2 to 5 years. The etiology is likely to be multifactorial, involving both genetic and environmental factors. Defects in DCTN1 are the cause of Perry syndrome (PERRYS) [MIM:168605]; also called

parkinsonism with alveolar hypoventilation and mental depression. Perry syndrome is a neuropsychiatric disorder characterized by mental depression not responsive to antidepressant drugs or electroconvulsive therapy, sleep disturbances, exhaustion and marked weight loss.

Parkinsonism develops later and respiratory failure occurred terminally.

序列相似性 Belongs to the dynactin 150 kDa subunit family.

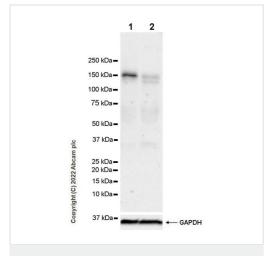
Contains 1 CAP-Gly domain.

翻译后修饰 Ubiquitinated by a SCF complex containing FBXL5, leading to its degradation by the

proteasome.

细胞定位 Cytoplasm. Cytoplasm > cytoskeleton.

图片



Western blot - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

All lanes : Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629) at 1/1000 dilution

Lane 1: HeLa (human cervix adenocarcinoma epithelial cell), transfected with scrambled siRNA control, whole cell lysate

Lane 2: Hela transfected with siRNA specifically targeting DCTN1, whole cell lysate

Lysates/proteins at 20 µg per lane.

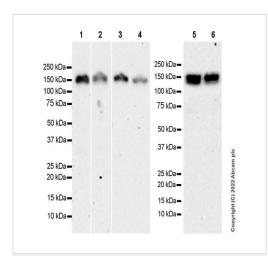
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

Predicted band size: 141 kDa Observed band size: 150 kDa

Exposure time: 114 seconds

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



Western blot - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

All lanes : Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629) at 1/1000 dilution

Lane 1 : HeLa (human cervix adenocarcinoma epithelial cell), whole cell lysate

Lane 2: SH-SY5Y (human neuroblastoma epithelial cell), whole cell lysate

Lane 3 : Neuro-2a (mouse neuroblastoma neuroblast), whole cell lysate

Lane 4: NIH/3T3 (mouse embryonic fibroblast), whole cell lysate

Lane 5: Mouse brain tissue lysate

Lane 6: Mouse cerecbellum tissue lysate

Lysates/proteins at 20 µg per lane.

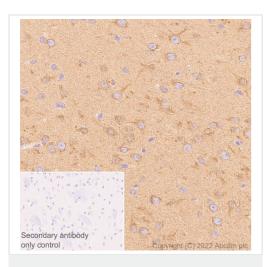
Secondary

All lanes : Goat Anti-Rabbit $\lg G \ H\&L \ (HRP) \ (\underline{ab97051})$ at 1/100000 dilution

Predicted band size: 141 kDa Observed band size: 150 kDa

Exposure time: 3 minutes

Blocking and diluting buffer and concentration: 5% NFDM/TBST This blot was developed using a high sensitivity ECL substrate.

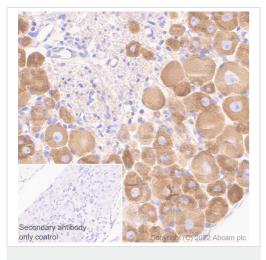


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunohistochemical analysis of paraffin-embedded mouse cerebrum tissue labeling DCTN1/p150-glued with ab302629 at 1/1000 (0.539 μ g/ml) followed by a ready to use Leica DS9800 (Bond Polymer Refine Detection kit). Positive staining on mouse cerebrum (PMID:17122035). The section was incubated with ab302629 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

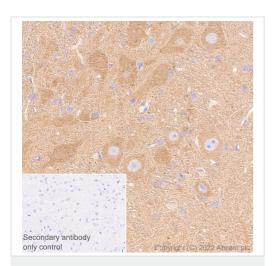


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunohistochemical analysis of paraffin-embedded mouse dorsal root ga tissue labeling DCTN1/p150-glued with ab302629 at 1/1000 (0.539 μg/ml) followed by a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit). Positive staining on mouse dorsal root ganglion (PMID:23874158). The section was incubated with ab302629 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

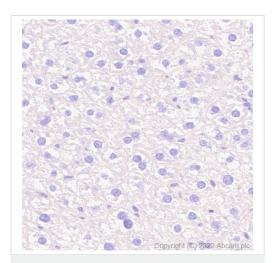


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunohistochemical analysis of paraffin-embedded mouse spinal cord tissue labeling DCTN1/p150-glued with ab302629 at 1/1000 (0.539 µg/ml) followed by a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit). Positive staining on mouse spinal cord (PMID:23408943, 17122035). The section was incubated with ab302629 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

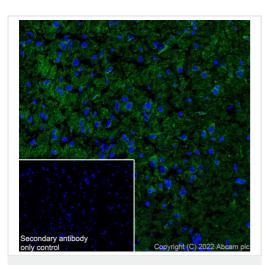


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunohistochemical analysis of paraffin-embedded mouse liver tissue labeling DCTN1/p150-glued with ab302629 at 1/1000 (0.539 µg/ml) followed by a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit). Negative control: no staining on mouse liver (PMID:1836789). The section was incubated with ab302629 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond™ Polymer Refine Detection kit).

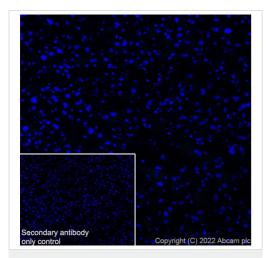
Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins



Immunohistochemistry (Frozen sections) - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen mouse spinal cord (fresh) tissue labeling DCTN1/p150-glued with ab302629 at 1/100 (1.078 µg/ml) dilution followed by **ab150081** Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed at 1/1000 (2 µg/ml) dilution (Green). Positive staining on mouse spinal cord is observed. The nuclear counterstain was DAPI (Blue).

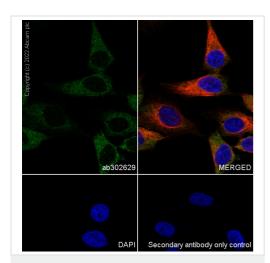
Secondary antibody control: Secondary antibody is <u>ab150081</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed at 1/1000 (2 µg/ml) dilution.



Immunohistochemistry (Frozen sections) - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen mouse liver (fresh) tissue labeling DCTN1/p150-glued with ab302629 at 1/100 (1.078 μg/ml) dilution followed by <u>ab150081</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed at 1/1000 (2 μg/ml) dilution (Green). Negative control: no staining on mouse liver (PMID:1836789) is observed. The nuclear counterstain was DAPI (Blue).

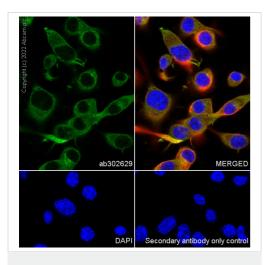
Secondary antibody control: Secondary antibody is <u>ab150081</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed at 1/1000 (2 µg/ml) dilution.



Immunocytochemistry/ Immunofluorescence - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized HeLa (human cervix adenocarcinoma epithelial cell) cells labeling DCTN1/p150-glued with ab302629 at 1/50 (10.78 µg/ml) dilution, followed by **ab150081** Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed antibody at 1/1000 (2µg/ml dilution) (Green). Confocal image showing mainly cytoplasmic staining in HeLa cell line is observed. **ab195889** Antialpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor[®] 594) was used to counterstain tubulin at 1/200 (2.5µg/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

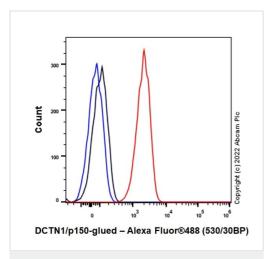
Secondary antibody only control: Secondary antibody is <u>ab150081</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed at 1/1000 ($2\mu g/ml$) dilution.



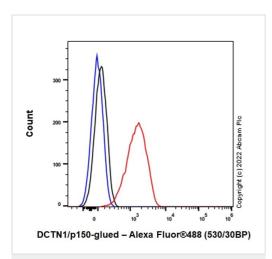
Immunocytochemistry/ Immunofluorescence - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized Neuro-2a (mouse neuroblastoma neuroblast) cells labeling DCTN1/p150-glued with ab302629 at 1/50 (10.78 µg/ml) dilution, followed by ab150081 Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed antibody at 1/1000 (2µg/ml) dilution (Green). Confocal image showing cytoplasmic staining in Neuro-2a cell line is observed. ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor[®] 594) was used to counterstain tubulin at 1/200 (2.5µg/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

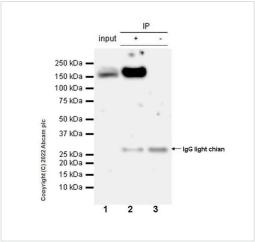
Secondary antibody only control: Secondary antibody is <u>ab150081</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) preadsorbed at 1/1000 ($2\mu g/ml$) dilution.

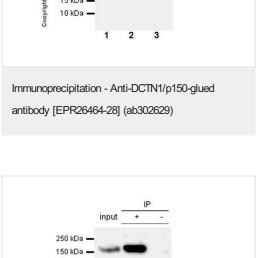


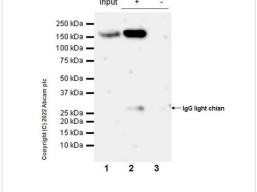
Flow Cytometry (Intracellular) - Anti-DCTN1/p150glued antibody [EPR26464-28] (ab302629) Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized HeLa (human cervix adenocarcinoma epithelial cell) cells labeling DCTN1/p150-glued with ab302629 at 1/50 dilution (1µg) (Red) compared with a Rabbit monoclonal lgG (ab172730) (Black) isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit lgG (Alexa Fluor® 488, ab150081) at 1/2000 dilution was used as the secondary antibody.



Flow Cytometry (Intracellular) - Anti-DCTN1/p150glued antibody [EPR26464-28] (ab302629) Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized Neuro-2a (mouse neuroblastoma neuroblast) cells labeling DCTN1/p150-glued with ab302629 at 1/50 dilution (1µg) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor[®] 488, ab150081) at 1/2000 dilution was used as the secondary antibody.







Immunoprecipitation - Anti-DCTN1/p150-glued antibody [EPR26464-28] (ab302629)

DCTN1/p150-glued was immunoprecipitated from 0.35 mg HeLa (human cervix adenocarcinoma epithelial cell), whole cell lysate 10 µg with ab302629 at 1/30 dilution (2µg in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab302629 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP) (ab131366) was used at 1/5000 dilution.

Lane 1: HeLa (human cervix adenocarcinoma epithelial cell), whole cell lysate 10 μg

Lane 2: ab302629 IP in HeLa whole cell lysate

 $\mbox{\bf Lane 3:} Rabbit\ monoclonal\ lgG\ (\underline{ab172730})\ instead\ of\ ab302629\ in$ HeLa whole cell lysate

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

Exposure time: 3 minutes

DCTN1/p150-glued was immunoprecipitated from 0.35 mg Neuro-2a (mouse neuroblastoma neuroblast), whole cell lysate 10 µg with ab302629 at 1/30 dilution (2µg in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab302629 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP) (ab131366) was used at 1/5000 dilution.

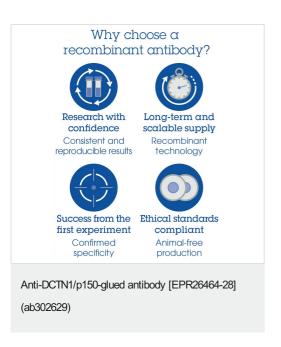
Lane 1: Neuro-2a (mouse neuroblastoma neuroblast), whole cell lysate 10 μg

Lane 2: ab302629 IP in Neuro-2a whole cell lysate

 $\begin{tabular}{ll} \textbf{Lane 3}: Rabbit monoclonal lgG ($\underline{ab172730}$) instead of ab302629 in Neuro-2a whole cell lysate \\ \end{tabular}$

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

Exposure time: 3 minutes



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