


### Anti-DYNLL1/PIN antibody [EP1660Y] ab51603

KO 評価済 リコンビナント RabMAb

★★★★★ 7 Abreviews 35 References 画像数 12

#### 製品の概要

製品名	Anti-DYNLL1/PIN antibody [EP1660Y]
製品の詳細	Rabbit monoclonal [EP1660Y] to DYNLL1/PIN
由来種	Rabbit
特異性	ab51603 recognizes DLC8. 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	<b>適用あり:</b> Flow Cyt (Intra), WB, IHC-P, IP, ICC/IF
種交差性	<b>交差種:</b> Mouse, Rat, Human <b>交差が予測される動物種:</b> Drosophila melanogaster 
免疫原	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.
エピトープ	The epitope for this antibody is on the N-terminus, AA2-14.
ポジティブ・コントロール	WB: HeLa cell lysate; Mouse testis tissue lysate. IHC-P: Human liver tissue. Flow Cyt (intra): HeLa cells.
特記事項	This product is a recombinant monoclonal antibody, which offers several advantages including: <ul style="list-style-type: none"> <li>- High batch-to-batch consistency and reproducibility</li> <li>- Improved sensitivity and specificity</li> <li>- Long-term security of supply</li> <li>- Animal-free production</li> </ul> For more information <a href="#">see here</a> . Our RabMAb <sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a> .

#### 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. Stable for 12 months at -20°C.
バッファー	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.21% BSA

精製度	Protein A purified
ポリ/モノ	モノクローナル
クローン名	EP1660Y
アイソタイプ	IgG

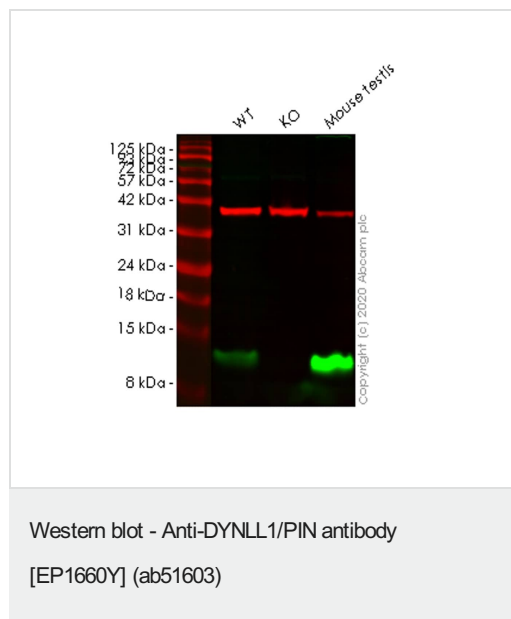
## アプリケーション

**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ab51603の使用に適用されず  
 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)		1/2300.
WB	★★★★★ (6)	1/1000 - 1/10000. Detects a band of approximately 10 kDa (predicted molecular weight: 10 kDa).
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat.
IP		1/30. <b>For unpurified use at 1/100.</b>
ICC/IF	★★★★★ (1)	1/100 - 1/250.

## ターゲット情報

<b>機能</b>	<p>Acts as one of several non-catalytic accessory components of the cytoplasmic dynein 1 complex that are thought to be involved in linking dynein to cargos and to adapter proteins that regulate dynein function. Cytoplasmic dynein 1 acts as a motor for the intracellular retrograde motility of vesicles and organelles along microtubules. May play a role in changing or maintaining the spatial distribution of cytoskeletal structures.</p> <p>Binds and inhibits the catalytic activity of neuronal nitric oxide synthase.</p> <p>Promotes transactivation functions of ESR1 and plays a role in the nuclear localization of ESR1.</p> <p>Regulates apoptotic activities of BCL2L11 by sequestering it to microtubules. Upon apoptotic stimuli the BCL2L11-DYNLL1 complex dissociates from cytoplasmic dynein and translocates to mitochondria and sequesters BCL2 thus neutralizing its antiapoptotic activity.</p>
<b>組織特異性</b>	Ubiquitous.
<b>配列類似性</b>	Belongs to the dynein light chain family.
<b>翻訳後修飾</b>	Phosphorylation at Ser-88 appears to control the dimer-monomer transition. According to PubMed:15193260, it is phosphorylated at Ser-88 by PAK1, however, according to PubMed:18650427, the DYNLL1 dimer is not accessible for PAK1 and the phosphorylation could not be demonstrated in vitro.
<b>細胞内局在</b>	Cytoplasm, cytoskeleton. Nucleus. Mitochondrion. Upon induction of apoptosis translocates together with BCL2L11 to mitochondria.



**All lanes** : Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603) at 1/1000 dilution

**Lane 1** : Wild-type HeLa cell lysate

**Lane 2** : DYNLL1 knockout HeLa cell lysate

**Lane 3** : Mouse testis tissue lysate

Lysates/proteins at 20 µg per lane.

### Secondary

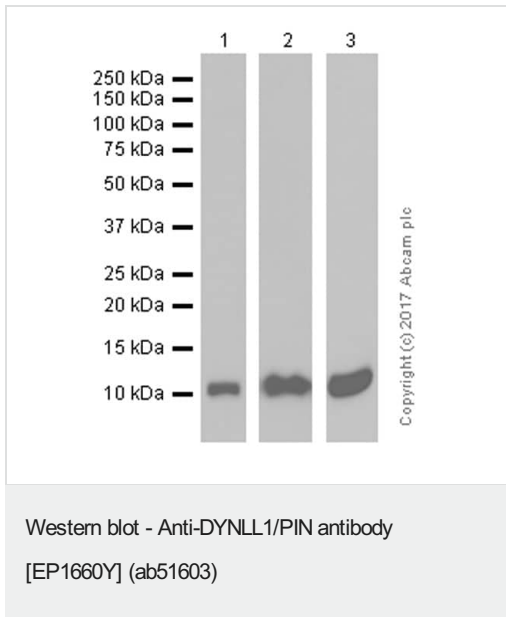
**All lanes** : Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ([ab216773](#)) at 1/10000 dilution

**Predicted band size:** 10 kDa

**Observed band size:** 10 kDa

**Lanes 1-3:** Merged signal (red and green). Green - ab51603 observed at 10 kDa. Red - loading control [ab8245](#) observed at 36 kDa.

ab51603 Anti-DYNLL1/PIN antibody [EP1660Y] was shown to specifically react with DYNLL1/PIN in wild-type HeLa cells. Loss of signal was observed when knockout cell line [ab265265](#) (knockout cell lysate [ab257414](#)) was used. Wild-type and DYNLL1/PIN knockout samples were subjected to SDS-PAGE. ab51603 and Anti-GAPDH antibody [6C5] - Loading Control ([ab8245](#)) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ([ab216773](#)) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed ([ab216776](#)) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



**All lanes :** Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603) at 1/10000 dilution (purified)

**Lane 1 :** HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates

**Lane 2 :** Mouse testis lysates

**Lane 3 :** Rat testis lysates

Lysates/proteins at 20 µg per lane.

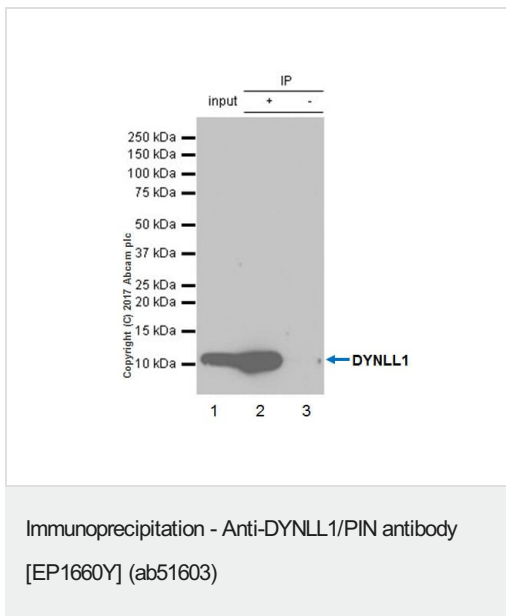
### Secondary

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

**Predicted band size:** 10 kDa

**Observed band size:** 10 kDa

Blocking and diluting buffer: 5% NFDm/TBST



ab51603 (purified) at 1:30 dilution (2ug) immunoprecipitating DYNLL1/PIN in HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate.

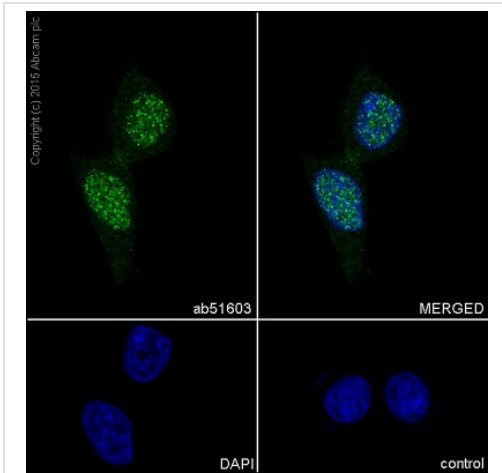
**Lane 1 (input):** HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate 10ug

**Lane 2 (+):** ab51603 & HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

**Lane 3 (-):** Rabbit monoclonal IgG (**ab172730**) instead of ab51603 in HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

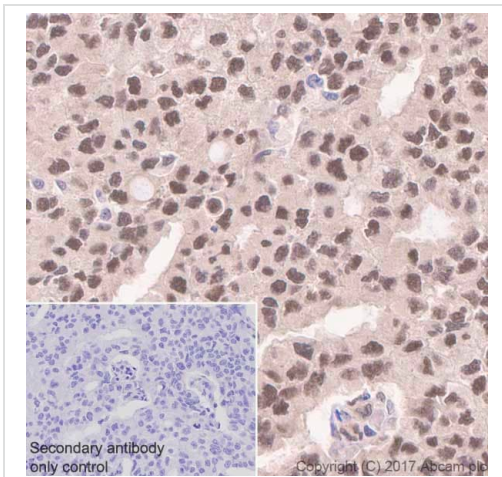
For western blotting, VeriBlot for IP Detection Reagent (HRP) (**ab131366**) was used for detection at 1:1000 dilution.

Blocking and diluting buffer: 5% NFDm/TBST.



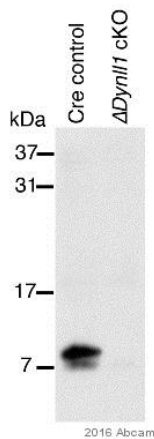
Immunocytochemistry/ Immunofluorescence - Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

Immunocytochemistry/ Immunofluorescence analysis of MCF7 (Human breast adenocarcinoma epithelial cell) cells labeling DYNLL1/PIN with Purified ab51603 at 1:100 dilution (6.7 µg/ml). Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. **ab150077** Goat anti rabbit IgG(Alexa Fluor® 488) was used as the secondary antibody at 1:1000 dilution. DAPI nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human breast carcinoma tissue sections labeling DYNLL1/PIN with Purified ab51603 at 1:500 dilution (1.34 µg/ml). Heat mediated antigen retrieval was performed using **ab93684** (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.



Western blot - Anti-DYNLL1/PIN antibody

[EP1660Y] (ab51603)

This image is courtesy of an abreview submitted by Dr. Jörg Heierhorst

**All lanes :** Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603) at 1/5000 dilution (unpurified)

**Lane 1 :** Primary mouse Mb1-Cre control Eμ-Myc B cell lymphoma (lysate of whole lymphnode)

**Lane 2 :** Primary mouse Mb1-Cre DYNLL1/PIN-conditional knockout Eμ-Myc B cell lymphoma (lysate of whole lymphnode)

**Secondary**

**All lanes :** HRP conjugated polyclonal goat IgG at 1/5000 dilution

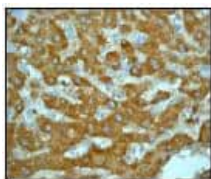
Performed under reducing conditions.

**Predicted band size:** 10 kDa

**Observed band size:** 10 kDa

**Exposure time:** 10 minutes

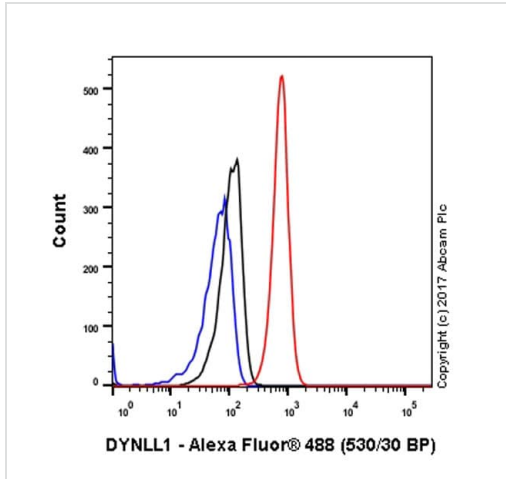
Lymphnodes were dissociated in PBS 2% FBS. Cell suspensions filtered through 70 μm and 40 μm cell strainers, and 300 x g pellets were lysed in modified RIPA buffer (150 mM NaCl, 20 mM Tris pH7.4, 1 mM EDTA, 1 mM EGTA, 10 mM NaF, 1% Triton X-100, 1% sodium deoxycholate, 0.1% SDS, 1 mM PMSF, 1 x protein inhibitor cocktail (Sigma)).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-DYNLL1/PIN antibody

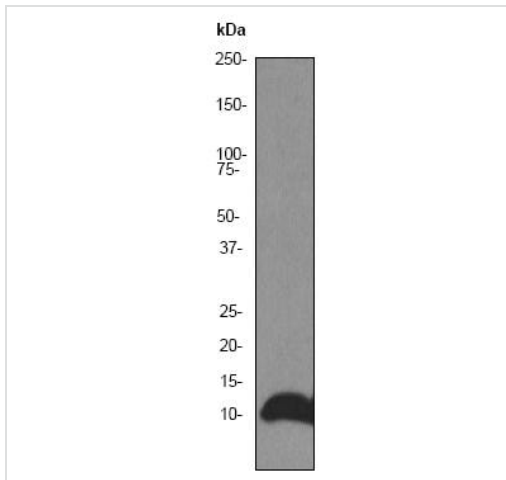
[EP1660Y] (ab51603)

Immunohistochemical staining of paraffin embedded human liver using unpurified ab51603 (1/100).



Flow Cytometry (Intracellular) - Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma) cells labeling DYNLL1/PIN (red) with purified ab51603 at a 1/2300 dilution. Cells were fixed with 4% paraformaldehyde and permeabilized with 90% methanol. A goat anti-rabbit IgG (Alexa Fluor® 488) (**ab150077**) was used as the secondary antibody at a 1/2000 dilution. Black - Rabbit monoclonal IgG (**ab172730**). Blue (unlabeled control) - Cells without incubation with the primary and secondary antibodies.



Western blot - Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

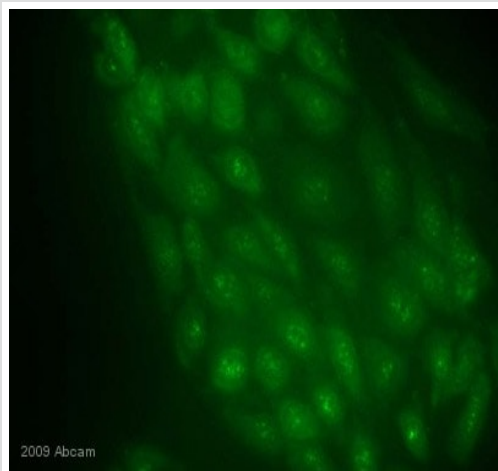
Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603) at 1/10000 dilution (unpurified) + HeLa cell lysate at 10 µg

**Secondary**

Goat anti-rabbit HRP at 1/2000 dilution

**Predicted band size:** 10 kDa

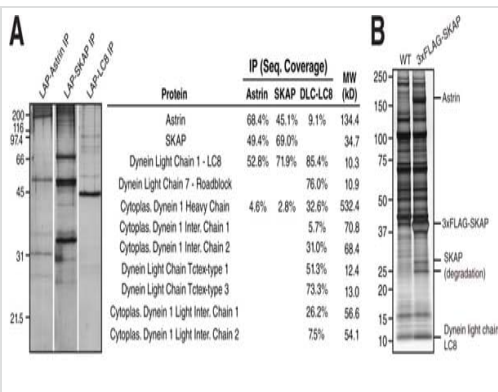
**Observed band size:** 10 kDa



Immunocytochemistry/ Immunofluorescence - Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

This image is courtesy of an anonymous Abreview

Unpurified ab51603 staining DLC8 in mouse kidney cells by ICC/IF (immunocytochemistry/immunofluorescence). Cells were fixed with methanol, permeabilized with 0.1% Triton and blocked with 1% milk for 1 hour at room temperature. The sample was incubated with primary antibody (1/400; 1% milk in PBS) for 16 hours at 4°C. An Alexa Fluor®488-conjugated Goat polyclonal to rabbit IgG (1/1000) was used as secondary antibody.



Immunoprecipitation - Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

Image from Schmidt JC et al, J Cell Biol. 2010 Oct 18;191(2):269-80. Epub 2010 Oct 11, Fig 2. DOI 10.1083/jcb.201006129

Unpurified ab51603 used in IP. SKAP and Astrin form a complex. (A, left) Silver-stained gels showing a one-step IP of GFPLAP-Astrin, GFPLAP-SKAP, or GFPLAP-LC8. (A, right) Data from the mass spectrometric analysis of the purifications indicating the percent sequence coverage from each IP. (B) Silver-stained gel showing the purification of FLAG-SKAP from chicken DT40 cells relative to controls. The indicated proteins were identified by excising them from a gel and analyzing them by mass spectrometry.



### Why choose a recombinant antibody?



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Recombinant technology



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**Ethical standards compliant**  
Animal-free production

Anti-DYNLL1/PIN antibody [EP1660Y] (ab51603)

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