# abcam

# Product datasheet

# Anti-DYNLL1/PIN antibody [EP1660Y] - BSA and Azide free ab232343





重组 RabMAb

## 8 图像

#### 概述

产品名称 Anti-DYNLL1/PIN抗体[EP1660Y] - BSA and Azide free

描述 兔单克隆抗体[EP1660Y] to DYNLL1/PIN - BSA and Azide free

宿主 Rabbit

特异性 ab51603 recognizes DLC8. The mouse and rat recommendation is based on the WB results. We

do not guarantee IHC-P for mouse and rat.

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

种属反应性 与反应: Mouse, Rat, Human

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

表位 The epitope for this antibody is on the N-terminus, AA2-14.

阳性对照 IHC-P: Human breast carcinoma tissue.

常规说明 ab232343 is the carrier-free version of ab51603.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.

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<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **patents**.

性能

形式 Liquid

**存放说明** Shipped at 4°C. Store at +4°C. Do Not Freeze.

**存储溶液** pH: 7.2

Constituent: PBS

**无载体** 是

纯**度** Protein A purified

**克隆** 单克隆

**克隆编号** EP1660Y

**同种型** IgG

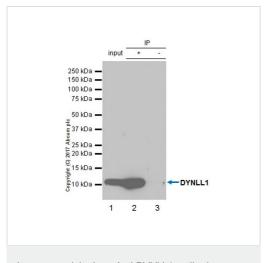
应用

The Abpromise guarantee Abpromise™承诺保证使用ab232343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		Use at an assay dependent concentration. Detects a band of approximately 10 kDa (predicted molecular weight: 10 kDa).
IP		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. 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.
ICC/IF		Use at an assay dependent concentration.
Flow Cyt (Intra)		Use at an assay dependent concentration.

## 图片



Immunoprecipitation - Anti-DYNLL1 antibody [EP1660Y] - BSA and Azide free (ab232343)

**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 (+): <u>ab51603</u> & 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

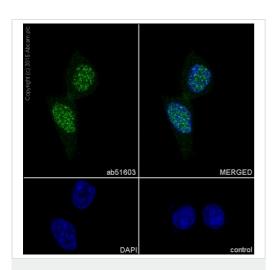
For western blotting, VeriBlot for IP Detection Reagent (HRP) (ab131366) was used for detection at 1:1000 dilution.

Blocking and diluting buffer: 5% NFDM/TBST.

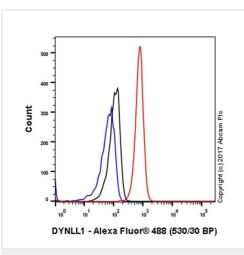
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).

Immunocytochemistry/ Immunofluorescence analysis of MCF7 (Human breast adenocarcinoma epithelial cell) cells labeling DYNLL1 with Purified <a href="mailto:ab51603">ab51603</a> at 1:100 dilution (6.7µg/ml). Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. <a href="mailto:ab150077">ab150077</a> Goat anti rabbit lgG(Alexa Fluor<sup>®</sup> 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.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).



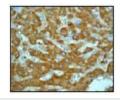
Immunocytochemistry/ Immunofluorescence - Anti-DYNLL1 antibody [EP1660Y] - BSA and Azide free (ab232343)



Flow Cytometry (Intracellular) - Anti-DYNLL1/PIN antibody [EP1660Y] - BSA and Azide free (ab232343)

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 lgG (Alexa Fluor<sup>®</sup> 488) (**ab150077**) was used as the secondary antibody at a 1/2000 dilution. Black - Rabbit monoclonal lgG (**ab172730**). Blue (unlabeled control) - Cells without incubation with the primary and secondary antibodies.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).

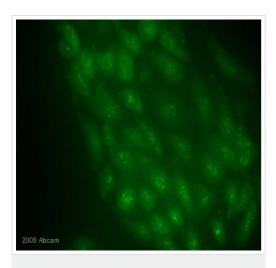


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DYNLL1 antibody

[EP1660Y] - BSA and Azide free (ab232343)

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).

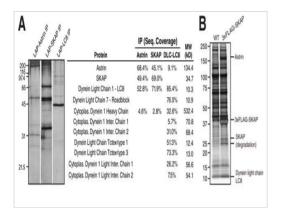


Immunocytochemistry/ Immunofluorescence - Anti-DYNLL1/PIN antibody [EP1660Y] - BSA and Azide free (ab232343)

This image is courtesy of an anonymous Abreview.

Unpurified <u>ab51603</u> staining DLC8 in mouse kidney cells 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<sup>®</sup>488-conjugated Goat polyclonal to rabbit IgG (1/1000) was used as secondary antibody.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).

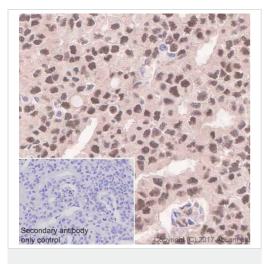


Immunoprecipitation - Anti-DYNLL1/PIN antibody [EP1660Y] - BSA and Azide free (ab232343)

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 <u>ab51603</u> 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.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).

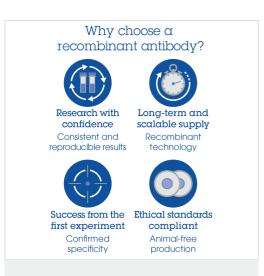


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DYNLL1 antibody

[EP1660Y] - BSA and Azide free (ab232343)

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab51603).



Anti-DYNLL1/PIN antibody [EP1660Y] - BSA and Azide free (ab232343)

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