

KD-Validated Anti-DYNLL1 Recombinant Rabbit Monoclonal Antibody



Catalog #: 61946

Aliases

DYNLL1; Dynein Light Chain LC8-Type 1; DLC1; DLC8; PIN; DNCL1; LC8; Protein Inhibitor Of Neuronal Nitric Oxide Synthase; Dynein, Cytoplasmic, Light Polypeptide 1; Dynein Light Chain 1, Cytoplasmic; 8 KDa Dynein Light Chain; DNCLC1; Hd1c1; HDLC1; Cytoplasmic Dynein Light Polypeptide; LC8a

Background

Gene Name: DYNLL1

NCBI Gene Entry: [8655](#)

UniProt Entry: [P63167](#)

Application Information

Molecular Weight: Predicted, 10 kDa, observed, 10 kDa

Clonality: Rabbit monoclonal antibody

Clone ID: 23GB5245

Species Reactivity: Human, mouse, rat

Applications Tested: Western blotting (WB), flow cytometry (FCM), immunocytochemistry (IC), immunohistochemistry (IHC)

Immunogen

A synthesized peptide derived from human DYNLL1

Isotype

Rabbit IgG

Storage Buffer

Supplied in PBS (pH 7.4) containing 50% glycerol, and 0.02% sodium azide.

Storage

Store at -20 °C for one year.

Recommended Dilutions

Western Blotting (WB): 1:1,000-1:5,000

Flow Cytometry (FCM): 1:2,000

Immunocytochemistry (IC): 1:100-1:1,000

Immunohistochemistry (IHC): 1:100-1:200

SUPPORT

SUPPORT@GENUINBIOTECH.COM
TEL: +1-540-855-7041

ORDERS

SALES@GENUINBIOTECH.COM
FAX: +1-540-855-7041

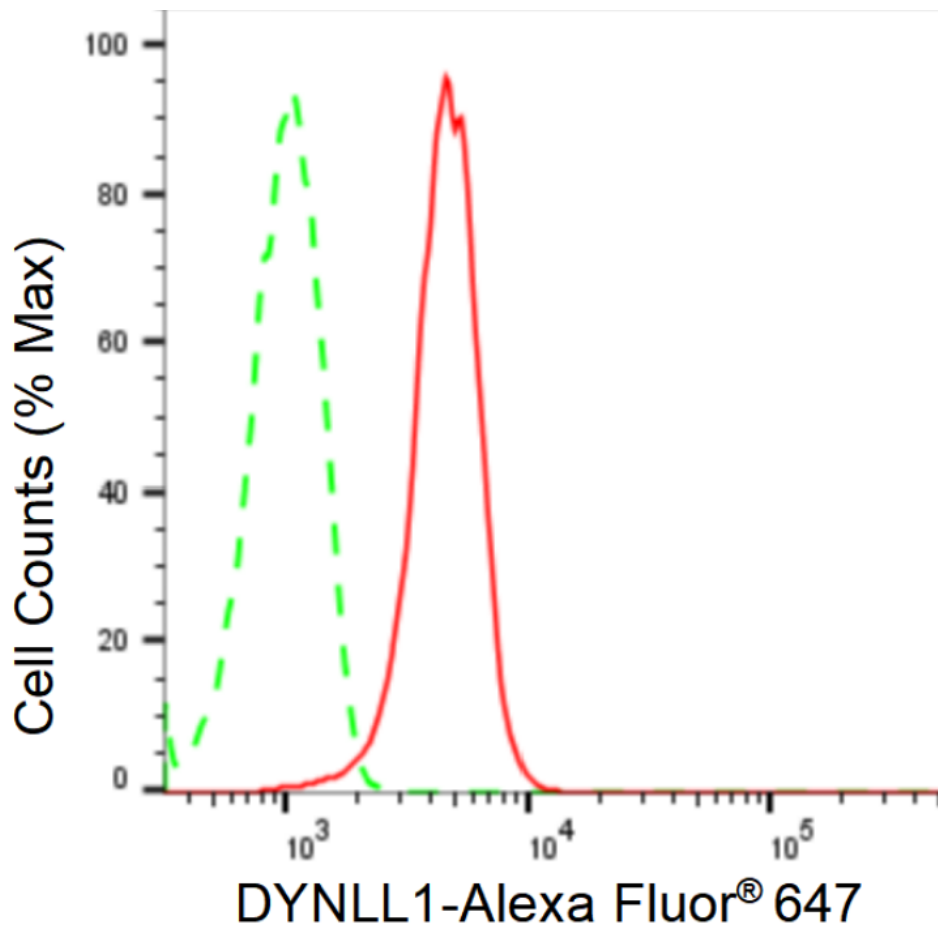
WWW.GENUINBIOTECH.COM

KD-Validated Anti-DYNLL1 Recombinant Rabbit Monoclonal Antibody

PAGE 2

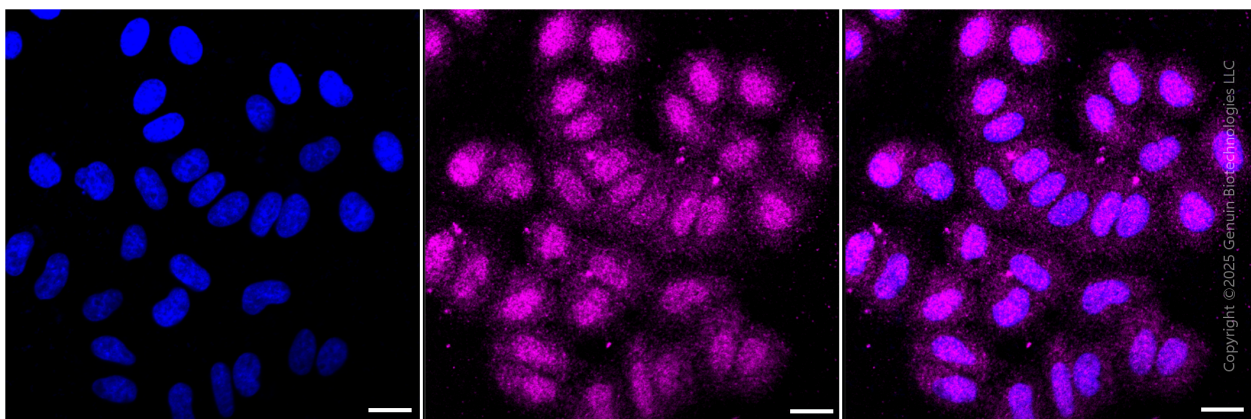
Note: This product is for research use only.

Validation Data



Copyright ©2024 Genuin Biotechnologies LLC

Flow cytometric analysis of DYNLL1 expression in HepG2 cells using DYNLL1 antibody (Cat#61946, 1:2,000). Green, isotype control; red, DYNLL1.



Copyright ©2025 Genuin Biotechnologies LLC

SUPPORT

SUPPORT@GENUINBIOTECH.COM
TEL: +1-540-855-7041

ORDERS

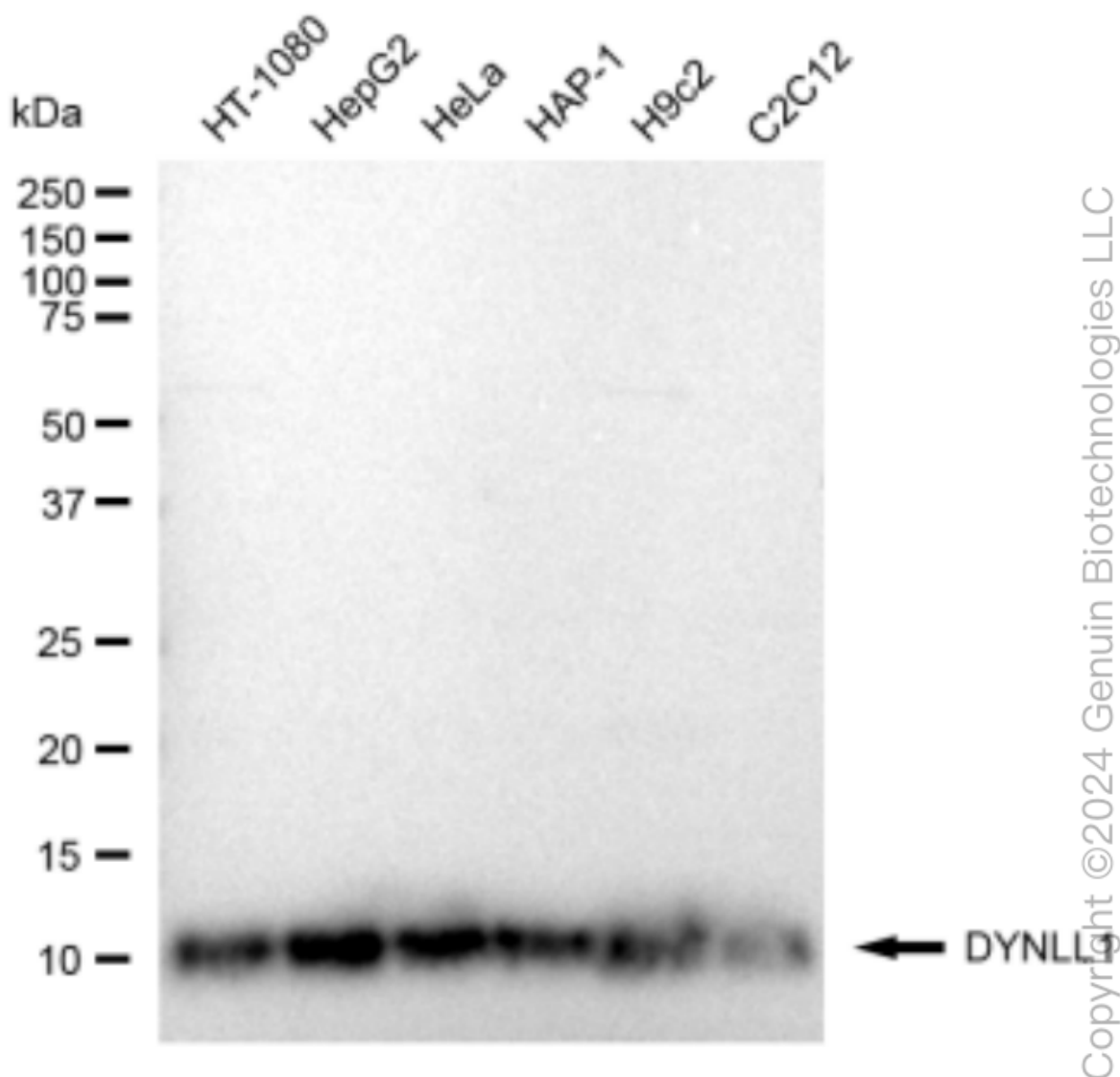
SALES@GENUINBIOTECH.COM
FAX: +1-540-855-7041

WWW.GENUINBIOTECH.COM

KD-Validated Anti-DYNLL1 Recombinant Rabbit Monoclonal Antibody

PAGE 3

Nuclei were stained blue with DAPI; DYNLL1 was stained magenta with Alexa Fluor® 647. Images were taken using Leica stellaris 5. Protein abundance based on laser Intensity and smart gain: Medium. Scale bar, 20 μ m.



Western blotting analysis using anti-DYNLL1 antibody (Cat#61946). Total cell lysates (30 μ g) from various cell lines were loaded and separated by SDS-PAGE. The blot was incubated with anti-DYNLL1 antibody (Cat#61946, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody (Cat#201, 1:20,000) respectively. Image was developed using FeQ™ ECL Substrate Kit (Cat#226).

SUPPORT

SUPPORT@GENUINBIOTECH.COM
TEL: +1-540-855-7041

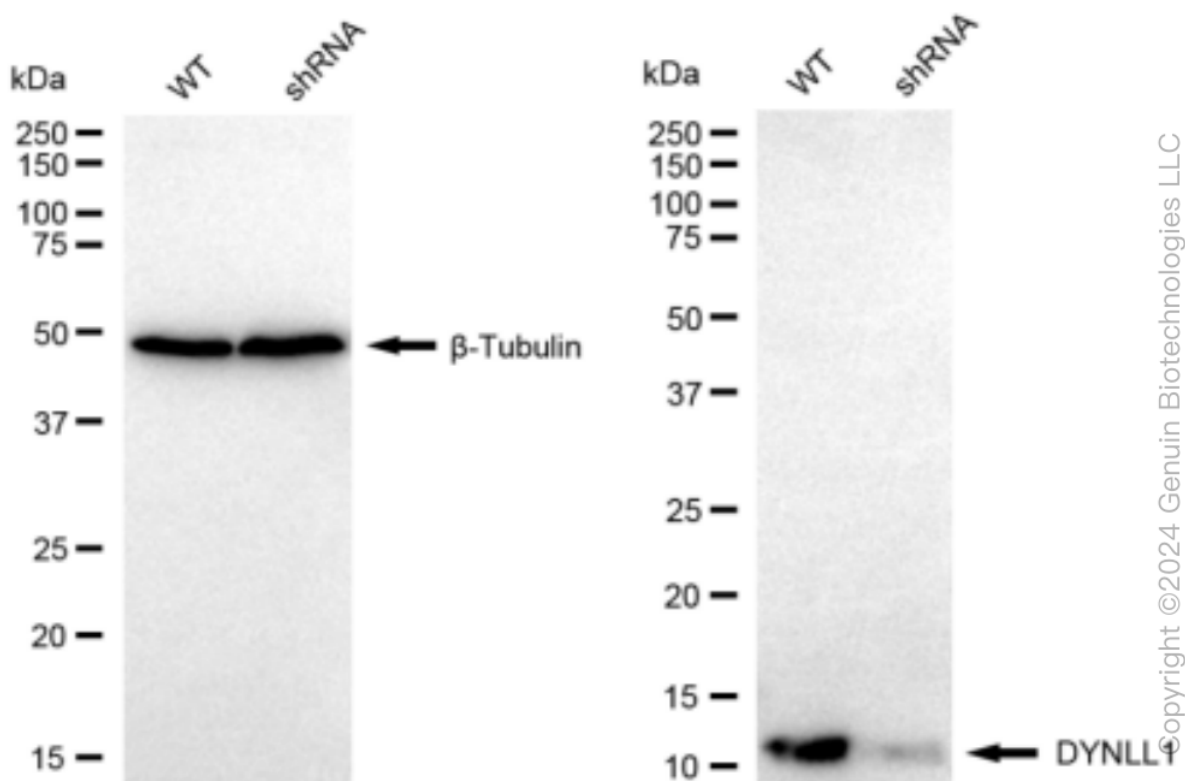
ORDERS

SALES@GENUINBIOTECH.COM
FAX: +1-540-855-7041

WWW.GENUINBIOTECH.COM

KD-Validated Anti-DYNLL1 Recombinant Rabbit Monoclonal Antibody

PAGE 4



Western blotting analysis using anti-DYNLL1 antibody (Cat#61946). DYNLL1 expression in wild type (WT) and DYNLL1 shRNA knockdown (KD) HeLa cells with 30 μ g of total cell lysates. β -Tubulin serves as a loading control. The blot was incubated with anti-DYNLL1 antibody (Cat#61946, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody (Cat#201, 1:20,000) respectively. Image was developed using FeQTM ECL Substrate Kit (Cat#226).

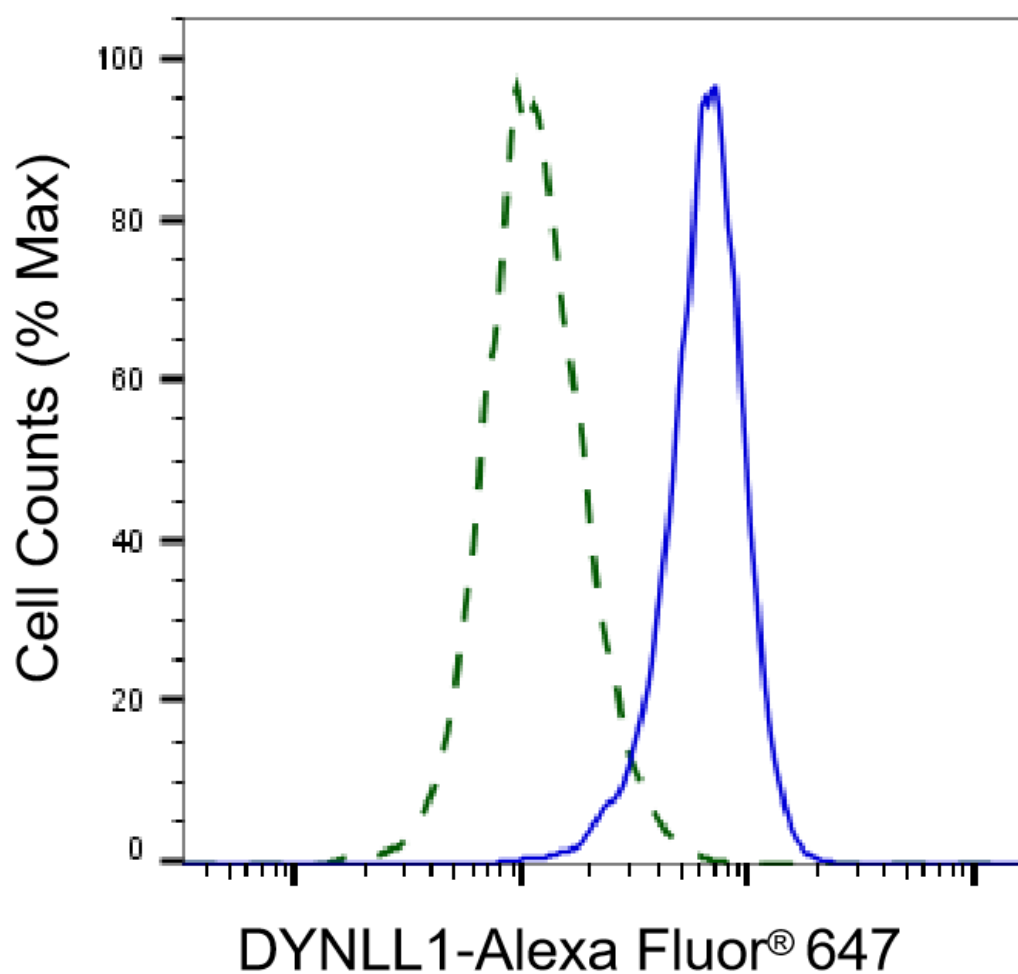
SUPPORT

SUPPORT@GENUINBIOTECH.COM
TEL: +1-540-855-7041

ORDERS

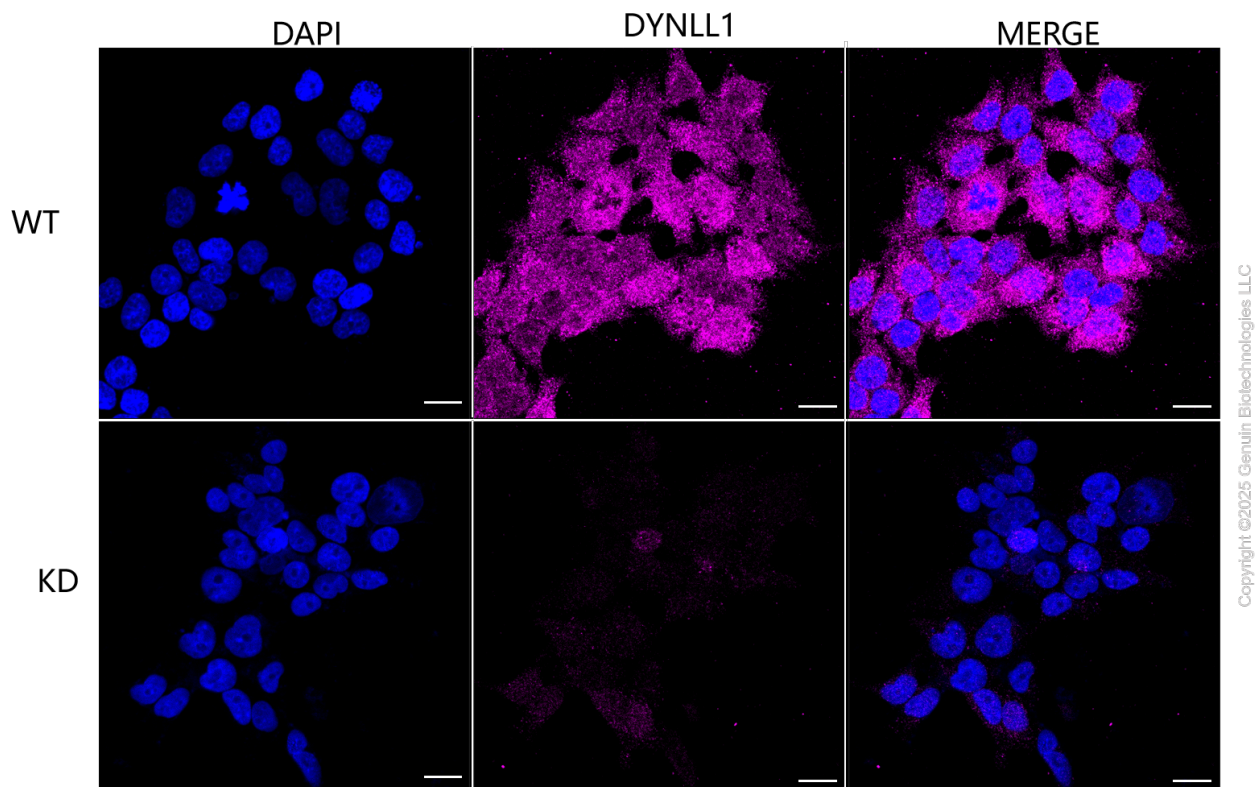
SALES@GENUINBIOTECH.COM
FAX: +1-540-855-7041

WWW.GENUINBIOTECH.COM

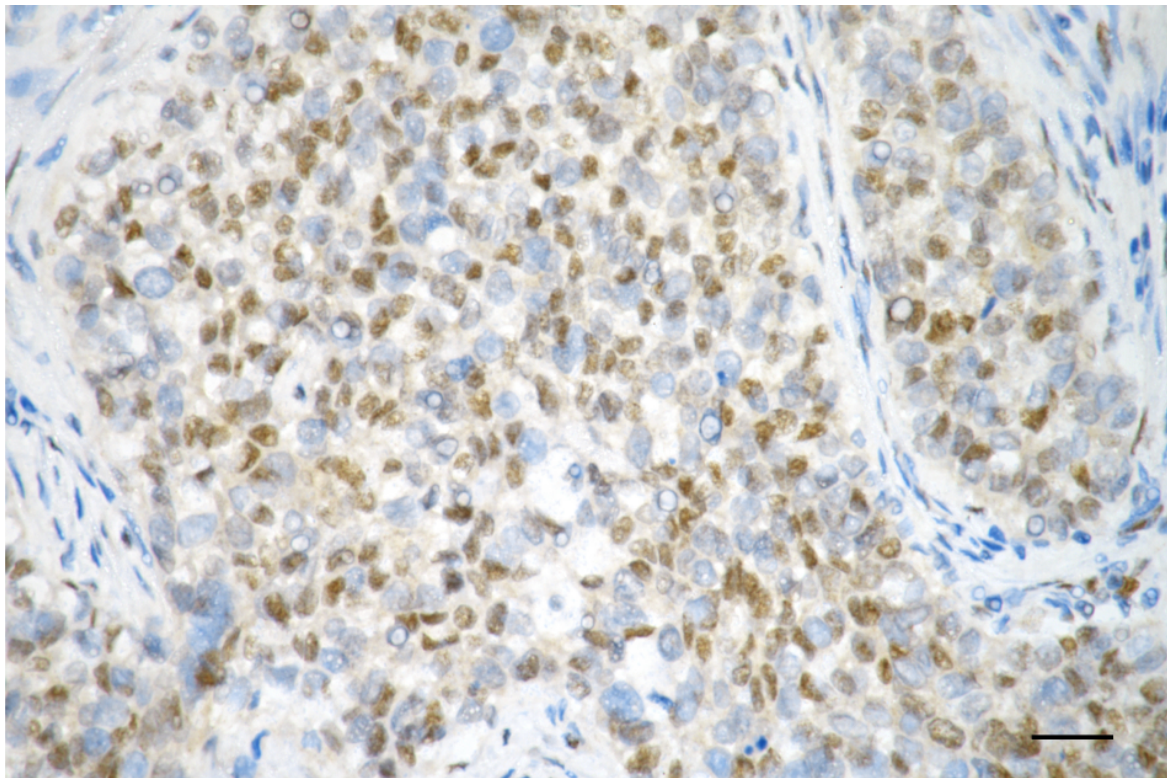


Copyright ©2025 Genuin Biotechnologies LLC

Validation of DYNLL1 knockdown using flow cytometry. Wild-type(WT, Blue) and knockdown(KD, Green) HeLa cells were stained with anti-DYNLL1 antibody (Cat#61946, 1:2,000) and analyzed using BD flow cytometer.



Immunocytochemical staining of HeLa cells using anti-DYNLL1 antibody (Cat#61946, 1:1,000), Top panel: wild-type (WT); Bottom panel: DYNLL1 shRNA knockdown (KD). Nuclei were stained blue with DAPI; DYNLL1 was stained magenta with Alexa Fluor® 647. Scale bar, 20 μ m. Permeabilization: Triton.



Copyright ©2025 Genuin Biotechnologies LLC

Immunohistochemistry was performed on paraffin-embedded human breast carcinoma using anti-DYNLL1 antibody (Cat#61946, 1:200). Antigen retrieval was done in sodium citrate buffer (pH 6.0). DAB was used for detection, with hematoxylin counterstaining. Images were acquired using a Nikon Ci-L Plus microscope (40× objective). Scale bar: 25 μ m.