abcam

Product datasheet

Anti-YY1 antibody [EPR4652] - Nuclear Loading Control ab109237

יעלאעבע RabMAb

★★★★ <u>10 Abreviews</u> 45 References

画像数 14

製品の概要

製品名 Anti-YY1 antibody [EPR4652] - Nuclear Loading Control

製品の詳細 Rabbit monoclonal [EPR4652] to YY1 - Nuclear Loading Control

由来種 Rabbit

アプリケーション 適用あり: WB, IHC-P, ICC/IF, Flow Cyt (Intra), ChIC/CUT&RUN-seq

適用なし: ChIP or IP

種交差性 交差種: Mouse, Rat, Human

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

ポジティブ・コントロール WB: HeLa, Daudi, Y79, and HuT-78 cell lysates, mouse and rat heart tissue. IHC-P: Human

kidney, tonsil and cervix carcinoma tissues. ICC/IF: HeLa and HUT-78 cells.

特記事項 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**.

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.

Stable for 12 months at -20°C.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol, 59% PBS, 0.05% BSA

精製度 Protein A purified

ポリモノ モノクローナル

アイソタイプ

ΙgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab109237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
WB	★★★★ (7)	1/2000 - 1/10000. Predicted molecular weight: 45 kDa.
IHC-P	★★★ ★ (1)	1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. See IHC antigen retrieval protocols. For unpurified use at 1/250 - 1/500.
ICC/IF		1/50. For unpurified use at 1/100 - 1/250.
Flow Cyt (Intra)		Use at an assay dependent concentration.
ChIC/CUT&RUN-seq		Use at an assay dependent concentration.

追加情報

Is unsuitable for ChIP or IP.

ターゲット情報

LAIA	Falls
森	配

Multifunctional transcription factor that exhibits positive and negative control on a large number of cellular and viral genes by binding to sites overlapping the transcription start site. May play an important role in development and differentiation. The function of YY1 as an activator or a repressor is specified by the presence of other proteins. For example it acts as a repressor in absence of adenovirus E1A protein but as an activator in its presence.

配列類似性

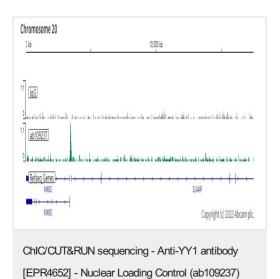
Belongs to the YY transcription factor family.

Contains 4 C2H2-type zinc fingers.

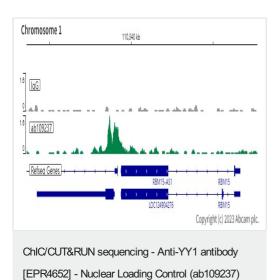
細胞内局在

Nucleus matrix. Associated with the nuclear matrix.

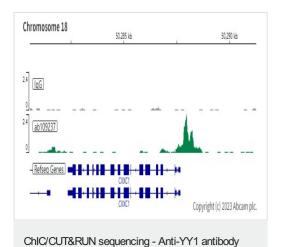
画像



ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ μ L, 2.5 x 10^5 K-562 (Human chronic myelogenous leukemia lymphoblast) cells and 5 μ g of ab109237 [EPR4652]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative lgG control **ab172730** is also shown. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.

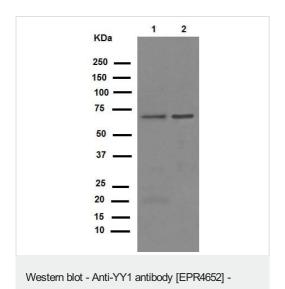


ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ μ L, 2.5 x 10^5 K-562 (Human chronic myelogenous leukemia lymphoblast) cells and 5 μ g of ab109237 [EPR4652]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative lgG control <u>ab172730</u> is also shown. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.



[EPR4652] - Nuclear Loading Control (ab109237)

ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ μ L, 2.5 x 10^5 K-562 (Human chronic myelogenous leukemia lymphoblast) cells and 5 μ g of ab109237 [EPR4652]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative lgG control <u>ab172730</u> is also shown. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.



Nuclear Loading Control (ab109237)

All lanes : Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237) at 1/10000 dilution (purified)

Lane 1 : HeLa (Human epithelial cell line from cervix adenocarcinoma) cell lysate

Lane 2: Daudi (Human Burkitt's lymphoma cell line) cell lysate

Lysates/proteins at 20 µg per lane.

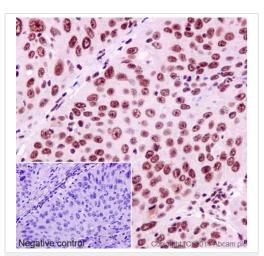
Secondary

All lanes : Peroxidase-conjugated goat anti-rabbit lgG (H+L) at 1/1000 dilution

Predicted band size: 45 kDa **Observed band size:** 68 kDa

Blocking buffer and concentration: 5% NFDM/TBST.

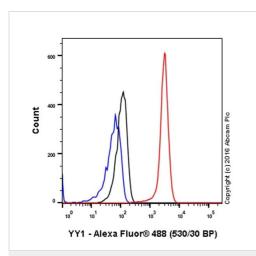
Diluting buffer and concentration: 5% NFDM /TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-YY1 antibody [EPR4652]

- Nuclear Loading Control (ab109237)

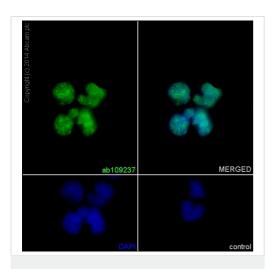
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human cervix carcinoma tissue labelling YY1 with purified ab109237 at 1/500. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. ab97051, a HRP-conjugated goat anti-rabbit lgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.



Flow Cytometry (Intracellular) - Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237) ab109237 staining YY1 in the human cell line HeLa (Human epithelial cell line from cervix adenocarcinoma) by intracellular flow cytometry. Cells were fixed with 4% paraformaldehyde, permiabilised with 90% methanol and the sample was incubated with the primary antibody at a dilution of 1/30. A goat anti rabbit IgG (Alexa Fluor[®] 488) at a dilution of 1/2000 was used as the secondary antibody.

Isoytype control: Rabbit monoclonal IgG (Black).

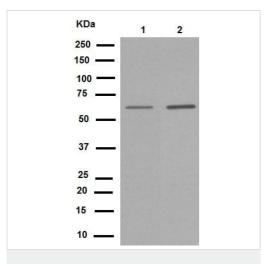
Unlabelled control: Cell without incubation with primary antibody and secondary antibody (Blue).



Immunocytochemistry/ Immunofluorescence - Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237)

Immunocytochemistry/Immunofluorescence analysis of HUT-78 cells labelling YY1 with purified ab109237 at 1/50. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. **ab150077**, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain.

Control: primary antibody (1/50) and secondary antibody, **ab150120**, an Alexa Fluor[®] 594-conjugated goat anti-mouse IgG (1/500).



Western blot - Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237)

All lanes : Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237) at 1/50000 dilution (purified)

Lane 1: Y79 (Human retinoblastoma cell line) cell lysate

Lane 2: HuT-78 cell lysate

Lysates/proteins at 10 µg per lane.

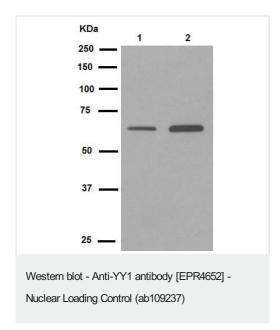
Secondary

All lanes : Peroxidase-conjugated goat anti-rabbit lgG (H+L) at 1/1000 dilution

Predicted band size: 45 kDa Observed band size: 68 kDa

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.



All lanes: Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237) at 1/2000 dilution (purified)

Lane 1: Mouse heart Lane 2: Rat heart

Lysates/proteins at 10 µg per lane.

Secondary

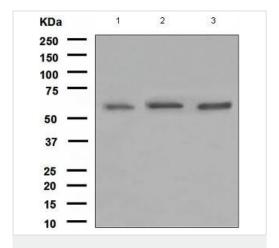
All lanes: Peroxidase-conjugated goat anti-rabbit lgG (H+L) at 1/1000 dilution

Predicted band size: 45 kDa

Observed band size: 68 kDa

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.



Western blot - Anti-YY1 antibody [EPR4652] -Nuclear Loading Control (ab109237)

All lanes: Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237) at 1/1000 dilution (unpurified)

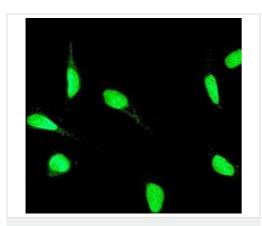
Lane 1: Daudi (Human Burkitt's lymphoma cell line) cell lysate

Lane 2: Y79 (Human retinoblastoma cell line) cell lysate

Lane 3: HuT-78 cell lysate

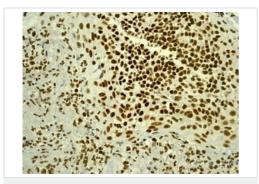
Lysates/proteins at 10 µg per lane.

Predicted band size: 45 kDa



Immunocytochemistry/ Immunofluorescence - Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237)

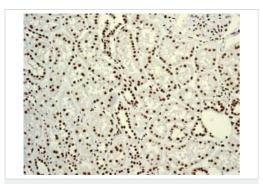
Immunocytochemistry/Immunofluorescence analysis of HeLa cells labelling YY1 with unpurified ab109237 at 1/100.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis human tonsil tissue labelling YY1 with unpurified ab109237 at 1/250.

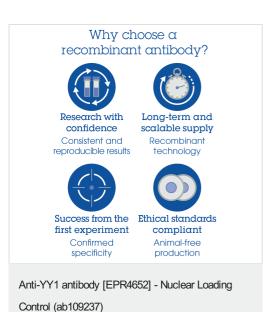
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-YY1 antibody [EPR4652] - Nuclear Loading Control (ab109237)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis human kidney tissue labelling YY1 with unpurified ab109237 at 1/250.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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