# abcam

# Product datasheet

# Anti-YY1 antibody [EPR4652] - BSA and Azide free ab232573

ועלשעבע RabMAb

1 References 画像数 10

#### 製品の概要

製品名 Anti-YY1 antibody [EPR4652] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR4652] to YY1 - BSA and Azide free

由来種 Rabbit

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

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 IHC-P: Human cervix carcinoma tissue.

特記事項 ab232573 is the carrier-free version of ab109237.

> 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® 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® 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. Do Not Freeze.

**バッファー** pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

**ポリ/モノ** モノクローナル **ウローン名** EPR4652

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ab232573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
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.  See IHC antigen retrieval protocols.
WB		Use at an assay dependent concentration. Predicted molecular weight: 45 kDa.
ICC/IF		Use at an assay dependent concentration.
ChIC/CUT&RUN-seq		Use at an assay dependent concentration.
Flow Cyt (Intra)		Use at an assay dependent concentration.

追加情報 Is unsuitable for ChIP or IP.

## ターゲット情報

機能

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

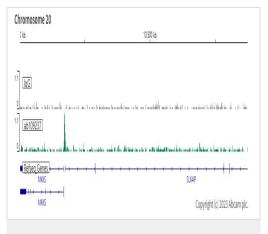
absence of adenovirus E1A protein but as an activator in its presence.

**配列類似性** Belongs to the YY transcription factor family.

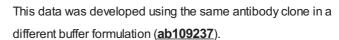
Contains 4 C2H2-type zinc fingers.

**細胞内局在** Nucleus matrix. Associated with the nuclear matrix.

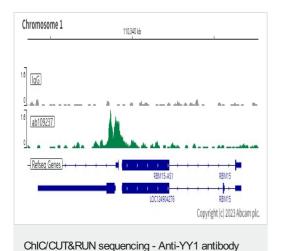
. . . .



ChIC/CUT&RUN sequencing - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)



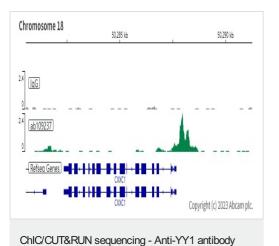
ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ $\mu$ L, 2.5 x 10^5 K-562 (Human chronic myelogenous leukemia lymphoblast) cells and 5  $\mu$ g of <u>ab109237</u> [EPR4652]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The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



[EPR4652] - BSA and Azide free (ab232573)

This data was developed using the same antibody clone in a different buffer formulation (<u>ab109237</u>).

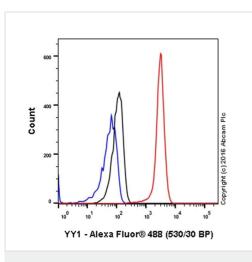
ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ $\mu$ L, 2.5 x 10^5 K-562 (Human chronic myelogenous leukemia lymphoblast) cells and 5  $\mu$ g of <u>ab109237</u> [EPR4652]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The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



[EPR4652] - BSA and Azide free (ab232573)

This data was developed using the same antibody clone in a different buffer formulation (ab109237).

ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ $\mu$ L, 2.5 x 10^5 K-562 (Human chronic myelogenous leukemia lymphoblast) cells and 5  $\mu$ g of <u>ab109237</u> [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.



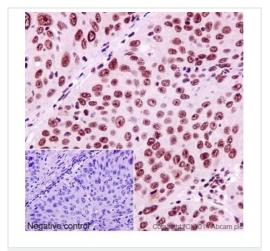
Flow Cytometry (Intracellular) - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

ab109237 staining YY1 in the human cell line HeLa (human 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<sup>®</sup> 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)

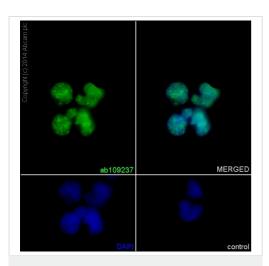
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab109237).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab109237</u>).

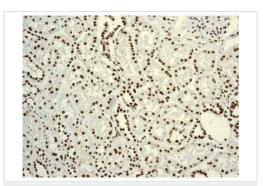


Immunocytochemistry/ Immunofluorescence - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

Immunocytochemistry/Immunofluorescence analysis of HUT-78 cells labelling YY1 with purified <u>ab109237</u> at 1/50. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. <u>ab150077</u>, an Alexa Fluor<sup>®</sup> 488-conjugated goat anti-rabbit IgG (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<sup>®</sup> 594-conjugated goat anti-mouse IgG (1/500).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab109237).

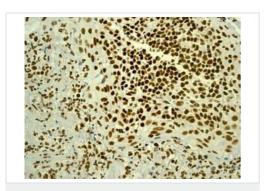


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab109237</u>).

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

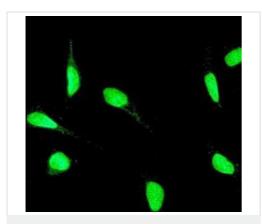


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab109237).

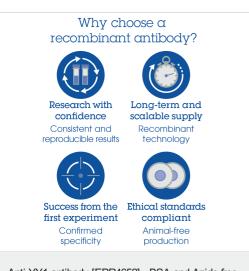
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunocytochemistry/ Immunofluorescence - Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab109237).



Anti-YY1 antibody [EPR4652] - BSA and Azide free (ab232573)

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