



Product Datasheet

Anti-YB1/YBX1 Antibody (orb76271)

Description Anti-YB1/YBX1 Antibody. Tested in Flow Cytometry, IHC, WB applications. This

antibody reacts with Human, Mouse, Rat.

Species/Host Rabbit

Reactivity Human, Mouse, Rat

Conjugation Unconjugated

Tested Applications FC, IHC, WB

Immunogen A synthetic peptide corresponding to a sequence in the middle region of human

YB1, identical to the related rat and mouse sequences.

Form/Appearance Lyophilized

Concentration Adding 0.2 ml of distilled water will yield a concentration of 500 μg/ml.

Storage Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -

20°C in small aliquots to prevent freeze-thaw cycles.

Note For research use only

Application notes Western blot, 0.1-0.5µg/ml, Human, Mouse, Rat Immunohistochemistry(Paraffin-

embedded Section), 2-5 μ g/ml, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μ g/1x106 cells, Human. Add 0.2ml of distilled water will yield a concentration of

500ug/ml

Isotype Rabbit IgG

Clonality Polyclonal

Antibody Type Primary Antibody

MW 50 kDa

Uniprot ID P67809

Biorbyt Ltd.

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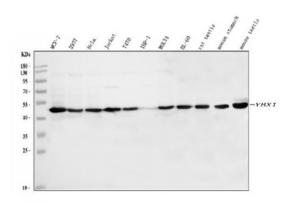
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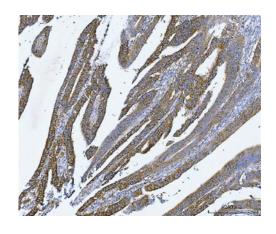


Expiration Date

12 months from date of receipt.



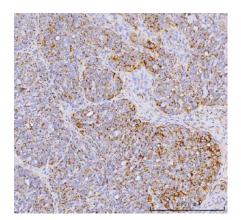
Western blot analysis of YBX1 using anti-YBX1 antibody (orb76271). Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel)/90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: human MCF-7 whole cell lysates, Lane 2: human 293T whole cell lysates, Lane 3: human HeLa whole cell lysates, Lane 4: human Jurkat whole cell lysates, Lane 5: human T47D whole cell lysates, Lane 6: human THP-1 whole cell lysates, Lane 7: human MOLT4 whole cell lysates, Lane 8: human HL-60 whole cell lysates, Lane 9: rat testis tissue lysates, Lane 10: mouse stomach tissue lysates, Lane 11: mouse testis tissue lysates. After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-YBX1 antigen affinity purified polyclonal antibody (Catalog # orb76271) at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # orb90503) with Tanon 5200 system. A specific band was detected for YBX1 at approximately 50 kDa. The expected band size for YBX1 is at 36 kDa.



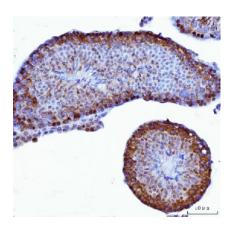
IHC analysis of YBX1 using anti-YBX1 antibody (orb76271). YBX1 was detected in a paraffin-embedded section of human rectal cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-YBX1 Antibody (orb76271) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



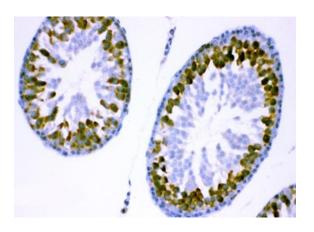




IHC analysis of YBX1 using anti-YBX1 antibody (orb76271). YBX1 was detected in a paraffin-embedded section of human ovarian cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-YBX1 Antibody (orb76271) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



IHC analysis of YBX1 using anti-YBX1 antibody (orb76271). YBX1 was detected in a paraffin-embedded section of mouse testis tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-YBX1 Antibody (orb76271) overnight at 4°C. Peroxidase Conjugated Goat Antirabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.

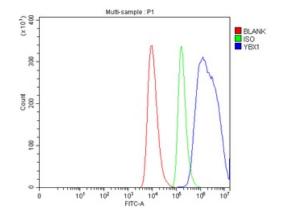


IHC analysis of YBX1 using anti-YBX1 antibody (orb76271). YBX1 was detected in a paraffin-embedded section of rat testis tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-YBX1 Antibody (orb76271) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.

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Flow Cytometry analysis of HEL cells using anti-YBX1 antibody (orb76271). Overlay histogram showing HEL cells stained with orb76271 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-YBX1 Antibody (orb76271, 1 μ g/1x10^6 cells) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (5-10 μ g/1x10^6 cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1 μ g/1x10^6) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

 $\begin{aligned} & \text{Email: } \underline{\text{info@biorbyt.com}}, \, \underline{\text{support@biorbyt.com}} \\ & \text{Phone: } \underline{+1 \ (415) \ 906-5211} \ \big| \ \text{Fax: } \underline{+1 \ (415) \ 651-8558} \end{aligned}$