

Catalog: OM161304



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cyclin B1 Antibody (H-433)

| Catalog: OM161304 | |
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| | 100ul |

Product profile

Product name cyclin B1 Antibody (H-433)

Antibody Type Primary Antibodies

Modification Notes In eukaryotic cells, mitosis is initiated following the activation of a protein kinase known variously as matu

ration-promoting factor, M-phase specific histone kinase or M-phase kinase. This protein kinase is composed of a catalytic subunit (Cdc2), a regulatory subunit (cyclin B) and a low molecular weight subunit (p13

-Suc 1). The Cdc/cyclin enzyme is subject to multiple levels of control, of which the regulation of the catal ytic subunit by tyrosine phosphorylation is the best understood. Tyrosine phosphorylation inhibits the Cd

c2/cyclin B enzyme; tyrosine dephosphorylation, occurring at the onset of mitosis, directly activates the p re-MPF complex. Evidence has established that B type cyclins not only act on M-phase regulatory subunit

s of the Cdc2 protein kinase, but also activate the Cdc25A and Cdc25B endogenous tyrosine phosphata

se, of which Cdc2 is the physiological substrate. The specificity of this effect is shown by the inability of e

ither cyclin A or cyclin D1 to display any such stimulation of Cdc25A or Cdc25B.

Key Feature

Clonality Polyclonal

Isotype lgG

Host Species Rabbit

 $\textbf{Tested Applications} \qquad \text{WB ,IP ,IF ,IHC ,FC ,ELISA}$

Species Reactivity Human Mouse Rat

Concentration 1mg/ml

Purification Affinity purified

Target Information

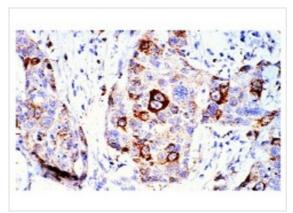
Alternative Names epitope corresponding to amino acids 1-433 representing full length cyclin B1 of human origin

Tissue Specificity epitope corresponding to amino acids 1-433 representing full length cyclin B1 of human origin

Entrez Gene

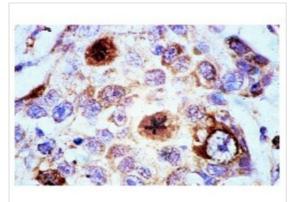
891

Application



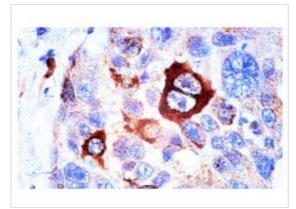
Application

Immunoperoxidase staining of formalin-fixed, paraffin-embedded human breast carcinoma tissue at low magnification.



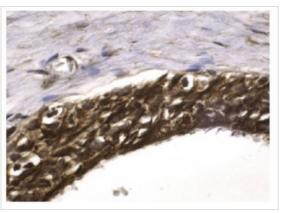
Application

Immunoperoxidase staining of formalin-fixed, paraffin-embedded human breast carcinoma tissue at high magnification. Note staining of selected cells, showing nuclear localization.



Application

Immunoperoxidase staining of formalin-fixed, paraffin-embedded human breast carcinoma tissue at high magnification. Note staining of selected cells, showing cytoplasmic localization.

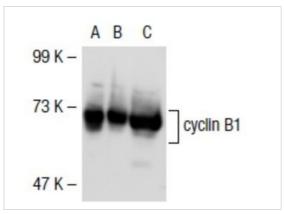


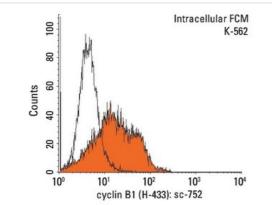
Application

Immunoperoxidase staining of formalin fixed, paraffin-embedded human testis tissue showing nuclear and cytoplasmic staining of cells in seminiferus ducts.

Application

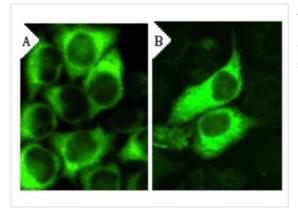
Western blot analysis of cyclin B1 expression in non-transfected: (A) and mouse cyclin B1 transfected: (B) 293T whole cell lysates and K-562 nuclear extract (C).





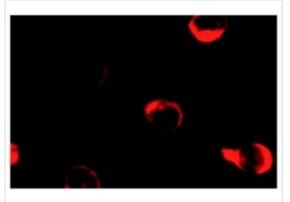
Application

Indirect, intracellular FCM analysis of fixed and permeabilized K-562 cells stained with cyclin B1 (H-433), followed by PE-conjugated goat anti-rabbit $\lg G$: . Black line histogram represents the isotype control, normal rabbit $\lg G$:

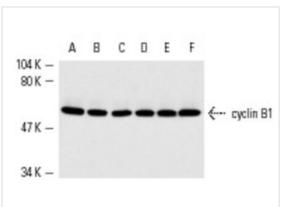


Application

Immunofluorescence staining of methanol-fixed, control HeLa (A) and cyclin B1 siRNA silenced HeLa (B) cells showing diminished cytoplasmic staining in the siRNA silenced cells. Cells probed with cyclin B1 (H-433): .



Application



Application

Western blot analysis of cyclin B1 expression in untreated (A,C,E) and phorbol ester-induced (B,D,F) K-562 (A,B), Jurkat (C,D) and HeLa (E,F) nuclear extracts.

Application Notes

recommended for detection of cyclin B1 of mouse, rat and human origin by WB, IP, IF, IHC(P), ELIS A and indirect FCM using PE-conjugated goat anti-rabbit IgG: sc-3739; also reactive with addition al species, including equine, canine, bovine and porcine:

Additional Information

Form Liquid

Storage Instructions For short-term storage, store at 4° C. For long-term storage, aliquot and store at -20°C or below. Avoid

multiple freeze-thaw cycles.

Storage Buffer phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.

Note The product is for research use only,not for use in diagnostic or therapeutic procedures.

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