

# Anti-CDC25C Antibody

Rabbit polyclonal antibody to CDC25C Catalog # AP59983

#### **Product Information**

**Application** WB, IF/IC, IHC

Primary Accession P30307
Other Accession P48967

**Reactivity** Human, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Calculated MW 53365

## **Additional Information**

Gene ID 995

Other Names M-phase inducer phosphatase 3; Dual specificity phosphatase Cdc25C

**Target/Specificity** KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human CDC25C. The exact sequence is proprietary.

**Dilution** WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 - 1/500)

IF/IC~~N/A IHC~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 -

1/500)

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

**Storage** Store at -20 °C.Stable for 12 months from date of receipt

#### **Protein Information**

Name CDC25C

**Function** Functions as a dosage-dependent inducer in mitotic control. Tyrosine

protein phosphatase required for progression of the cell cycle

(PubMed:<u>8119945</u>). When phosphorylated, highly effective in activating G2 cells into prophase (PubMed:<u>8119945</u>). Directly dephosphorylates CDK1 and

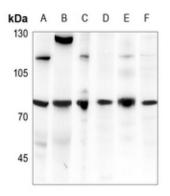
activates its kinase activity (PubMed:8119945).

Cellular Location Nucleus

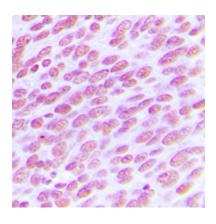
# **Background**

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human CDC25C. The exact sequence is proprietary.

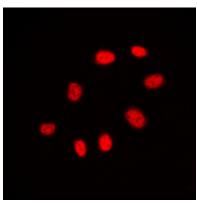
## **Images**



Western blot analysis of CDC25C expression in rat brain (A), mouse lung (B), CT26 (C), PC12 (D), U87MG (E), Hela (F) whole cell lysates.



Immunohistochemical analysis of CDC25C staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of CDC25C staining in A431 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.