

Product Name: LATS1 Rabbit Monoclonal Antibody
Catalog #: AMRe21250



Summary

| | |
|------------------------|----------------------------------|
| Production Name | LATS1 Rabbit Monoclonal Antibody |
| Description | Rabbit Monoclonal Antibody |
| Host | Rabbit |
| Application | WB,IF,IP,ELISA |
| Reactivity | Human,Mouse,Rat |

Performance

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|---------------------|--|
| Conjugation | Unconjugated |
| Modification | Unmodified |
| Isotype | IgG,Kappa |
| Clonality | Monoclonal |
| Form | Liquid |
| Storage | Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles. |
| Buffer | PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA |
| Purification | Protein A |

Immunogen

| | |
|--------------------------|--|
| Gene Name | LATS1 WARTS |
| Alternative Names | Serine/threonine-protein kinase LATS1;Large tumor suppressor homolog 1;WARTS protein kinase;h-warts; |
| Gene ID | 9113.0 |
| SwissProt ID | O95835. |

Application

| | |
|-------------------------|---|
| Dilution Ratio | WB 1:2000-1:10000;IF 1:200-1:1000;ELISA 1:5000-1:20000;IP 1:50-1:200; |
| Molecular Weight | Calculated MW:127kD;Observed MW:140kD |

Background

Cell localization:Cytoplasm.The protein encoded by this gene is a putative serine/threonine kinase that localizes to the

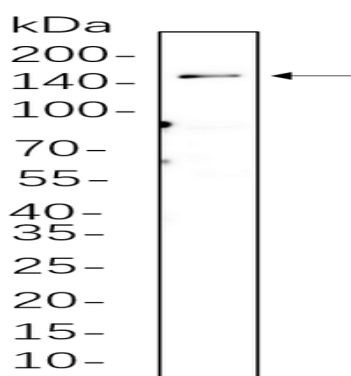
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mitotic apparatus and complexes with cell cycle controller CDC2 kinase in early mitosis. The protein is phosphorylated in a cell-cycle dependent manner, with late prophase phosphorylation remaining through metaphase. The N-terminal region of the protein binds CDC2 to form a complex showing reduced H1 histone kinase activity, indicating a role as a negative regulator of CDC2/cyclin A. In addition, the C-terminal kinase domain binds to its own N-terminal region, suggesting potential negative regulation through interference with complex formation via intramolecular binding. Biochemical and genetic data suggest a role as a tumor suppressor. This is supported by studies in knockout mice showing development of soft-tissue sarcomas, ovarian stromal cell tumors and a high sensitivity to carcinogenic treatment.

Research Area

Image Data



C2C12 cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with primary antibody 1:1000. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody.

Note

For research use only.