

# Recombinant Phospho-Histone H3 (Ser10) Monoclonal Antibody

catalog number: AN302105L

Note: Centrifuge before opening to ensure complete recovery of vial contents.

## **Description**

Reactivity Human; Mouse

**Immunogen** Peptide. This information is proprietary to PTMab

Host Rabbit Isotype lgG, κ Clone A829

**Purification** Protein Apurified

Buffer PBS, 50% glycerol, 0.05% Proclin 300, 0.05% protein protectant.

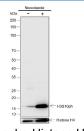
#### **Applications Recommended Dilution**

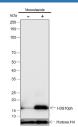
1:1000-1:5000 **WB** 

6μg antibody/100μg chromatin **ChIP** 

1:50-1:100 ΙP

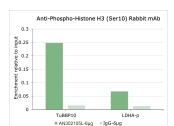
#### Data





Antibody at dilution of 1:2000. (-): HeLa, (+):HeLa + Nocodazole (100ng/mL, 18h)

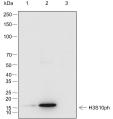
> Observed-MW:17 kDa Calculated-MW:17 kDa



Chromatin immunoprecipitation analysis of HeLa+Nocodazole(100ng/mL, 18h) immunoprecipitated DNA by real-time PCR using primers specific for the human LDHA-P and TuBBP10. The data are presented as enrichment of each sample relative to the total amount of input chromatin at each amplicon.

Western Blot with Phospho-Histone H3 (Ser10) Monoclonal Western Blot with Phospho-Histone H3 (Ser10) Monoclonal Antibody at dilution of 1:2000. (-): NIH-3T3, (+): NIH-3T3+Serum(0%, overnight) add Nocodazole(100ng/mL, 18h) add Calyculin A (100nM, 1h)

> Observed-MW:17 kDa Calculated-MW:17 kDa



Immunoprecipitation analysis using anti-Phospho-Histone H3 (Ser10) Monoclonal Antibody. Western blot was performed from the immunoprecipitate using Phospho-Histone H3 (Ser10) Monoclonal Antibody at a dilution of 1:100. Lane 1: 10% Input, Lane 2: Phospho-Histone H3 Monoclonal Antibody, Lane 3: Rabbit monoclonal IgG

> Observed-MW:17 kDa Calculated-MW:17 kDa

Isotype

# **Preparation & Storage**

# For Research Use Only

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Shipping Ice bag

# **Background**

Histones are subject to a variety of enzyme catalyzed modifications, including acetylation, methylation, phosphorylation, ubiquitylation, etc. Crotonylation of lysine is a newly identified reversible modification controlling chromosome structure and gene transcription. The reversible lysine crotonylation has been well demonstrated in eukaryotic histones from worm to human. The unique structure and genomic localization of histone lysine crotonylation suggest that it is mechanistically and functionally different from histone lysine acetylation. Specifically, in both human somatic and mouse male germ cell genomes, histone crotonylation marks either active promoters or potential enhancers. Crotonylation of histone H4 at Lys5 may play a vital role in the epigenetic modulation, including chromatin remodeling and DNA transcriptional regulation.

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