

Anti-H1-4 antibody



Catalog Number: 176509

Product name

Anti-H1-4 antibody

Specificity

Human, Mouse, Rat

Antibody description

Rabbit monoclonal antibody to H1-4

Preparation

This antigen of this antibody was synthetic phospho-peptide corresponding to residues surrounding thr17 of human h1.4.

Formulation

Liquid, 1*TBS (pH7.4), 0.05% BSA, 40% Glycerol.
Preservative: 0.05% Sodium Azide.

Storage

Store at +4°C after thawing. Aliquot store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.

Clonality

Monoclonal

Ig Type

IgG

Applications

WB, ICC/IF, IHC-P

Dilutions

WB: 1:500-1:1,000

ICC/IF: 1:50-1:200

IHC-P: 1:50-1:200

Validations

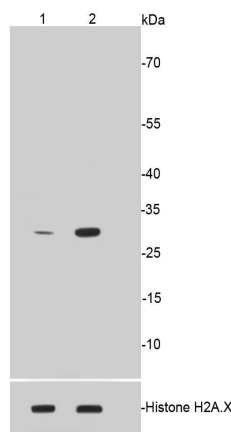


Fig1:: Western blot analysis of Phospho-Histone H1.3(T17)+Histone H1.4(T17) on different lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:40,000 dilution was used for 1 hour at room temperature.; Positive control::; Lane 1: Untreated CRC whole cell lysates; Lane 2: CRC cells treated with 1.5ug/ml Colcemid for 12 hours whole cell lysates

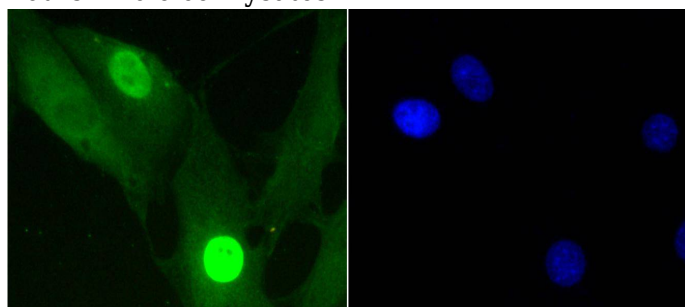


Fig2:: ICC staining of Phospho-Histone H1.3(T17)+Histone H1.4(T17) in NIH/3T3 cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with the primary antibody (1/50) for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Rabbit IgG was used as the secondary antibody at 1/1,000 dilution. The nuclear counter stain is DAPI (blue).

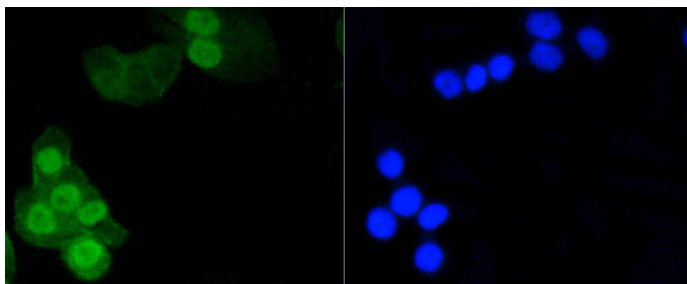


Fig3:; ICC staining of Phospho-Histone H1.3(T17)+Histone H1.4(T17) in CRC cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with the primary antibody (1/50) for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Rabbit IgG was used as the secondary antibody at 1/1,000 dilution. The nuclear counter stain is DAPI (blue).

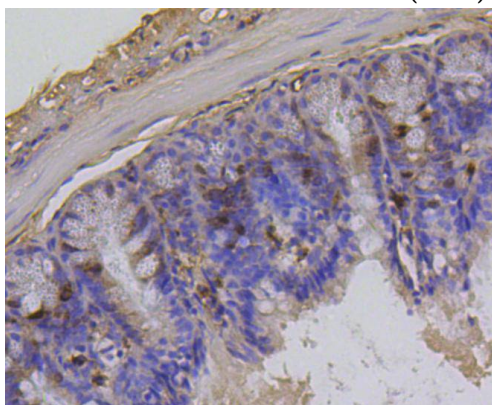


Fig4:; Immunohistochemical analysis of paraffin-embedded mouse colon cancer tissue using anti-Phospho-Histone H1.3(T17)+Histone H1.4(T17) antibody. Counter stained with hematoxylin.

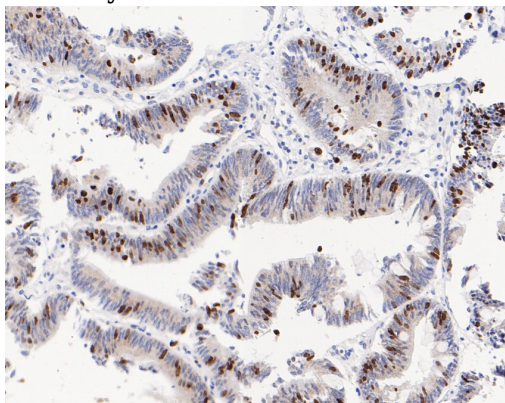


Fig5:; Immunohistochemical analysis of paraffin-

embedded human colon carcinoma tissue using anti-Phospho-Histone H1.3(T17)+Histone H1.4(T17) antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (1/200) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

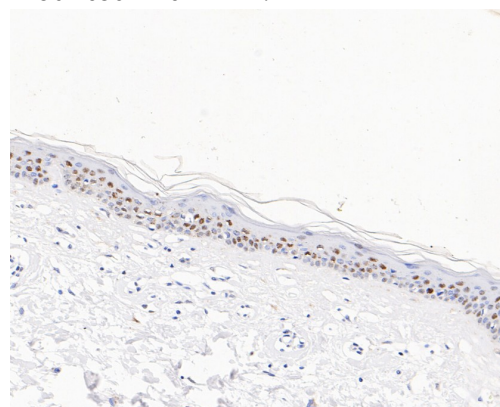
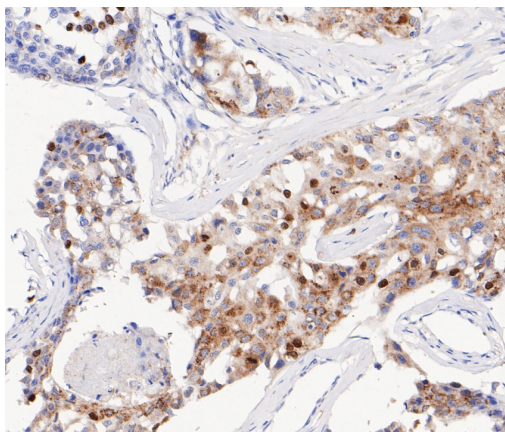


Fig6:; Immunohistochemical analysis of paraffin-embedded human skin tissue using anti-Phospho-Histone H1.3(T17)+Histone H1.4(T17) antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (1/200) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.



embedded human breast carcinoma tissue using anti-Phospho-Histone H1.3(T17)+Histone H1.4(T17) antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (1/200) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

Fig7.; Immunohistochemical analysis of paraffin-