

Anti-Acetyl-Histone H4-K91 antibody (STJ11100937)

STJ11100937

GENERAL INFORMATION

Product Type Primary antibodies

Short Description

Applications WB/IF/ICC/ELISA

Host/Source Rabbit

Reactivity Human/Mouse/Rat/Other

PRODUCT PROPERTIES

Clonality Polyclonal

Clone ID

Concentration
Conjugation
Purification
Dilution Range
Unconjugated
Affinity purification
WB:1:500-1:1000

IF/ICC:1:50-1:200

ELISA:Recommended starting concentration is 1 Mu g/mL. Please optimize the concentration based on your specific assay

requirements.

Formulation PBS with 0.02% Sodium Azide, 50% Glycerol, pH 7.3.

Isotype IgG

Storage Instruction Store at-20°C for up to 1 year from the date of receipt, and avoid repeat freeze-thaw cycles.

TARGET INFORMATION

Gene ID 121504/554313/8294/8359/8360/8361/8362/8363/8364/8365/8366/8367/8368/8370

Gene Symbol H4C1.H4C2.H4C3.H4C4.H4C5.H4C6.H4C8.H4C9.H4C11.H4C12.H4C13.H4C14.H4C15.H4C16

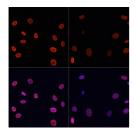
Uniprot ID H4_HUMAN Immunogen ALKRQ

Immunogen Region

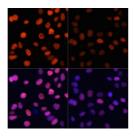
Specificity A synthetic acetylated peptide around K91 of human Histone H4 (NP_003529.1).

Immunogen ALKRQ

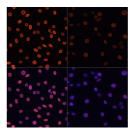
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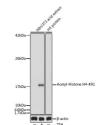
Immunofluorescence analysis of NIH/3T3 cells using Acetyl-Histone H4-K91 Rabbit polyclonal antibody (STJ11100937) at dilution of 1:100. NIH/3T3 cells were treated by TSA (1 uM) at 37 ŰC for 18 hours. Blue: DAPI for nuclear staining.



Immunofluorescence analysis of HeLa cells using Acetyl-Histone HA-K91 Rabbit polyclonal antibody (STJ11100937) at dilution of 1:100. HeLa cells wert treated by TSA (1 uM) at 37 ŰC for 18 hours. Blue DAPI for pucker staining.



Immunofluorescence analysis of C6 cells using Acety Histone H4-K91 Rabbit polyclonal antibod (STJ11109397) at dilution of 1:100. C6 cells were treated by TSA (1 uM) at 37 ŰC for 18 hours. Blue DAPI for nuclear staining.



Western blot analysis of hysates from NIH/3T3 cells using Acetyl-Histone H4-K91 Rabbit polyclona antibody (STJ1110983) at 1:1000 dilution, NIH/3T3 cells were treated by TSA (1 1000 dilution, NIH/3T3 cells were treated by TSA (1 1000 dilution, NIH/3T3 cells were treated by TSA (1 1000 dilution, NIH/3T3 cells were treated by TSA (1 1000 dilution, NIH/3T3 cells were treated by TSA (1 1000 dilution) to the constraint of the cells were treated by TSA (1 1000 dilution) to the cells were treated by the cells were