Histone H3.1 pAb

Orders =

400-6123-828

Abmost

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□200 µl

BACKGROUND

Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling.

REFERENCES

1. Mehedint MG et al. Choline deficiency alters global histone methylation and epigenetic marking at the Re1 site of the calbindin 1 gene. FASEB J 24:184-95.

SOURCE

Histone H3.1 polyclonal antibody is produced by immunizing rabbit with a synthetic peptide corresponding to residues of human H3.1(NP 003520.1).

SPECIFICITY

Histone H3.1 polyclonal antibody detects endogenous levels of total histone H3.1.

STORAGE

Store at -20°C. Stable for one year from the date of shipment.

REACTIVITY

H, Mk, M, ChHm, soybean, Arabidopsis thaliana

ISOTYPE

Rabbit IgG

PREDICTED MOLECULAR WEIGHT

17 kDa

APPLICATION TESTED

WB 1: 5000-1: 20000

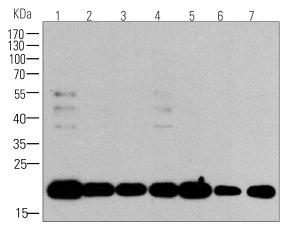
WB 1:500 (soybean, Arabidopsis thaliana)

For application specific protocols please see the web page at www. abmart. com. cn/ document, aspx

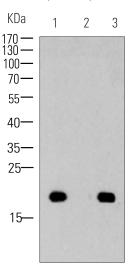
SECONDARY ANTIBODY

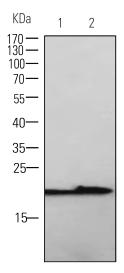
Use an anti-RABBIT secondary antibody.

APPLICATION DATA



Western blot analysis of lysates from various cell lines: Lane1-7: Hela, HEK293, Cos-7, NIH/3T3, CHO-K1, PC12, C6 Primary antibody: 1: 5000





Western blot analysis and peptide inhibition:

293T nuclear extract (Lane1); Histone H3.1 pAb pre-adsorbed with 2µM antigen petptide (lane2); Histone H3.1 pAb pre-absortbed with 2µM negative control peptide (lane3).

Primary antibody: 1: 20000

Western blot analysis of lysates from various cell lines:

Lane1-2: soybean, Arabidopsis thaliana

Primary antibody: 1: 500

Applications Key: WB —Western blot, IP—Immunoprecipitation, IHC—Immunohistochemistry, ChIP—Chromatin Immunoprecipitation, IF—Immunofluorescence