

# Histone H3 polyclonal antibody

Catalog: BS67367

Host: Rabbit

oit

Reactivity: Human, Mouse, Rat, Bovine, Chicken, Pig, Zebrafish

# **BackGround:**

The nucleosome, made up of four core histone proteins (H2A, H2B, H3, and H4), is the primary building block of chromatin. Originally thought to function as a static scaffold for DNA packaging, histones have now been shown to be dynamic proteins, undergoing multiple types of post-translational modifications, including acetylation, phosphorylation, methylation, and ubiquitination. Histone methylation is a major determinant for the formation of active and inactive regions of the genome and is crucial for the proper programming of the genome during development . Arginine methylation of histones H3 (Arg2, 17, 26) and H4 (Arg3) promotes transcriptional activation and is mediated by a family of protein arginine methyltransferases (PRMTs), including the co-activators PRMT1 and CARM1 (PRMT4) . In contrast, a more diverse set of histone lysine methyltransferases has been identified, all but one of which contain a conserved cata-SET domain originally identified lytic in the Drosophila Su(var)3-9, Enhancer of zeste, and Trithorax proteins. Lysine methylation occurs primarily on histones H3 (Lys4, 9, 27, 36, 79) and H4 (Lys20) and has been implicated in both transcriptional activation and silencing. Methylation of these lysine residues coordinates the recruitment of chromatin modifying enzymes containing methyl-lysine binding modules such as chromodomains (HP1, PRC1), PHD fingers (BPTF, ING2), tudor domains (53BP1), and WD-40 domains (WDR5). The discovery of histone demethylases, such as PADI4, LSD1, JMJD1, JMJD2, and JHDM1, has shown that methylation is a reversible epigenetic marker.

#### **Product:**

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide. **Molecular Weight:** 

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## ~ 17 kDa

**Swiss-Prot:** 

P68431; Q71DI3; P84243

#### **Purification&Purity:**

The antibody was purified by immunogen affinity chromatography.

# **Applications:**

WB (1/500 - 1/1000), IHC (1/50 - 1/200), IF/ICC (1/50 - 1/200)

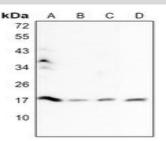
#### **Storage&Stability:**

Store at  $4 \,^{\circ}{\rm C}$  short term. Aliquot and store at -20  $^{\circ}{\rm C}$  long term. Avoid freeze-thaw cycles.

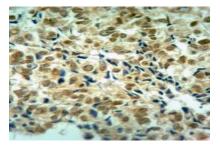
## **Specificity:**

Recognizes endogenous levels of Histone H3 protein.

#### **DATA:**



Western blot analysis of Histone H3 expression in HEK293T (A), DLD (B), U2OS (C), rat muscle (D) whole cell lysates.



Immunohistochemical analysis of Histone H3 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at

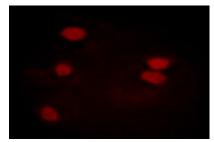
Bioworld technology, co. Ltd.Add:No 9, weidi road Qixia District Nanjing, 210046,<br/>P. R. China.Email:info@biogot.comTel:0086-025-68037686Fax:0086-025-68035151



# **PRODUCT DATA SHEET**

## **Bioworld Technology,Inc.**

room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of Histone H3 staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

# Note:

For research use only, not for use in diagnostic procedure.

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