

# Histone H2A.X (Acetyl Lys5) Polyclonal Antibody

Catalog # AP63459

#### **Product Information**

Application WB Primary Accession P16104

Reactivity Human, Mouse, Rat

HostRabbitClonalityPolyclonalCalculated MW15145

#### **Additional Information**

**Gene ID** 3014

Other Names H2AFX; H2AX; Histone H2A.x; H2a/x

**Dilution** WB~~WB: 1:1000-2000

Format PBS, pH 7.4, containing 0.09% (W/V) sodium azide as Preservative and 50%

Glycerol.

Storage Conditions -20°C

#### **Protein Information**

**Name** H2AX ( <u>HGNC:4739</u>)

**Function** Variant histone H2A which replaces conventional H2A in a subset of

nucleosomes. 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. Required for

checkpoint-mediated arrest of cell cycle progression in response to low doses of ionizing radiation and for efficient repair of DNA double strand breaks

(DSBs) specifically when modified by C-terminal phosphorylation.

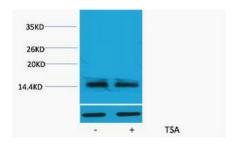
Cellular Location Nucleus. Chromosome

## **Background**

Variant histone H2A which replaces conventional H2A in a subset of nucleosomes. 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. Required for checkpoint-mediated arrest of cell cycle progression in response to low doses of ionizing radiation and for efficient repair of DNA double strand breaks (DSBs) specifically when modified by C- terminal phosphorylation.

### **Images**



Western blot analysis of extracts from Hela cells, untreated (-) or treated, 1:5000.. Secondary antibody was diluted at 1:20000 cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003,Inventbiotech,MN,USA).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.