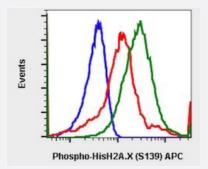
H2AX (phospho S139) monoclonal antibody, clone 1E4 (APC)

Catalog # MAB23467 Size 100 Reactions

Applications



Flow Cytometry

Flow cytometric analysis of HEK 293T cells with H2AX (phospho S139) monoclonal antibody, clone 1E4 (APC) (Cat # MAB23467). Untreated and unstained as negative control (blue) or untreated (red) or treated with UV and TPA (green).

Specification	
Product Description	Rabbit monoclonal antibody raised against synthetic phosphopeptide of human H2AX.
Immunogen	A synthetic phosphopeptide corresponding to residues surrounding S139 of human H2AX.
Host	Rabbit
Reactivity	Human, Mouse
Form	Liquid
Conjugation	APC
Purification	Protein A/G purification
lsotype	lgG1, kappa
Recommend Usage	Flow Cytometry (5 uL/10 ⁶ cells) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.4 (0.2% BSA, 0.09% sodium azide).
Storage Instruction	Store at 4°C.

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Product Information

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

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Gene Info — H2AFX	
Entrez GenelD	<u>3014</u>
Gene Name	H2AFX
Gene Alias	H2A.X, H2A/X, H2AX
Gene Description	H2A histone family, member X
Omim ID	<u>601772</u>
Gene Ontology	Hyperlink
Gene Summary	Histones are basic nuclear proteins that are responsible for the nucleosome structure of the chro mosomal fiber in eukaryotes. Two molecules of each of the four core histones (H2A, H2B, H3, an d H4) form an octamer, around which approximately 146 bp of DNA is wrapped in repeating units, called nucleosomes. The linker histone, H1, interacts with linker DNA between nucleosomes and f unctions in the compaction of chromatin into higher order structures. This gene encodes a member of the histone H2A family, and generates two transcripts through the use of the conserved stem-l oop termination motif, and the polyA addition motif. [provided by RefSeq
Other Designations	H2AX histone

Pathway

• Systemic lupus erythematosus

Disease

Azoospermia



- Breast cancer
- Breast Neoplasms
- DNA Damage
- Genetic Predisposition to Disease
- Lymphoma
- Oligospermia
- Ovarian cancer
- Prostate cancer
- Prostatic Neoplasms
- Urinary Bladder Neoplasms