

Product Data Sheet

Anti-Histone H2A.X (Phospho-S139) Antibody

Catalog #	Source	Reactivity	Applications		
CQA8445	Mouse	Н, М	WB, IF/IC		
Description	М	Mouse monoclonal antibody to Histone H2A.X (Phospho-S139)			
Immunogen	KL	LH-conjugated synthetic p	hosphopeptide corresponding to residues surrounding		
	S1	S139 of human Histone H2A.X protein. The exact sequence is proprietary.			
Purification	Tł	he antibody was purified	by immunogen affinity chromatography.		
Specificity	Re	Recognizes endogenous levels of Histone H2A.X protein only when phosphorylated			
	at	t S139.			
Clonality	М	Ionoclonal			
Conjugation					
Form	Lie	quid in PBS containing 50	% glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3.		
Dilution	W	/B (1/500 - 1/1000), IF/IC (1/50 - 1/100)		
Gene Symbol	H	2AX			
Alternative Names		H2AX; Histone H2AX; H2a/x; Histone H2A.X			
Entrez Gene		3014 (Human); 15270 (Mouse)			
SwissProt	P1	16104 (Human); P27661 (Mouse)		
Storage/Stabi	lity Sh	nipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	fre	eeze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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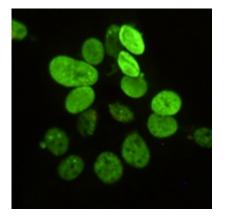
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Western blot analysis of Histone H2A.X (Phospho-S139) expression in NIH3T3 treated with Hydroxyurea (A) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 15 kD)



Immunofluorescent analysis of Histone H2A.X (Phospho-S139) staining in A549 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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