

Product Data Sheet

Anti-Histone H2A Antibody

Catalog #	Source	Reactivity	Applications
CPA6462	Rabbit	H, M, R, B, C, Mk	WB, IH
Description	Ra	abbit polyclonal antibody to Hi	istone H2A
Immunogen	KL	H-conjugated synthetic peptid.	le encompassing a sequence within the center
	re	gion of human Histone H2A. T	he exact sequence is proprietary.
Purification	Th	ne antibody was purified by imp	munogen affinity chromatography.
Specificity	Re	ecognizes endogenous levels of	f Histone H2A protein.
Clonality	Ро	blyclonal	
Conjugation			
Form	Lic	quid in 0.42% Potassium phosp	phate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	nd 0.01% sodium azide.	
Dilution	W	'B (1/500 - 1/1000), IH (1/50 - 1/2	200)
Gene Symbol	HI	ST1H2AG; HIST1H2AI; HIST1H2	2AK; HIST1H2AL; HIST1H2AM
Alternative Na	ames H2	2AFP; H2AFC; H2AFD; H2AFI; H	12AFN; Histone H2A type 1; H2A.1; Histone H2A/p
Entrez Gene	83	329, 8330, 8332, 8336, 8969 (H	Human); 319164, 319165, 319166, 319167, 319170,
	31	19171, 319172, 319191, 66543	33 (Mouse)
SwissProt	PC)COS8 (Human); P22752 (Mous	se); P02262 (Rat)
Storage/Stabi	lity Sh	nipped at 4°C. Upon delivery ali	liquot 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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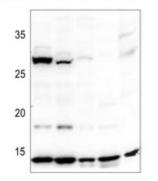
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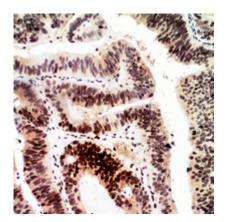
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kDa A B C D E



Western blot analysis of Histone H2A expression in U87MG (A), A549 (B), SGC7901 (C), rat lung (D), mouse brain (E) whole cell lysates. (Predicted band size: 14 kD; Observed band size: 14 kD)



Immunohistochemical analysis of Histone H2A staining in human colon 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 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.

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