

## **Product Data Sheet**

### **Anti-Histone H3 Antibody**

Catalog #	Source	Reactivity	Applications
CPA3483	Rabbit	H, M, R, B, C, P, Z	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody to His	tone H3
Immunogen	KLH-	conjugated synthetic peptide	e encompassing a sequence within the N-term
	regio	n of human Histone H3. The	exact sequence is proprietary.
Purification	The a	antibody was purified by imr	nunogen affinity chromatography.
Specificity	Reco	gnizes endogenous levels of	Histone H3 protein.
Clonality	Polyc	lonal	
Conjugation			
Form	Liqui	d in 0.42% Potassium phosp	hate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB (	1/500 - 1/1000), IH (1/100 - 1,	200), IF/IC (1/100 - 1/500)
Gene Symbol	H3F3	A	
Alternative Na	ames H3.3	A; H3F3; H3.3B; Histone H3.	3
Entrez Gene	3020	, 3021 (Human); 15078, 150	81 (Mouse); 100361558, 117056 (Rat)
SwissProt	P842	43 (Human); P84244 (Mous	e); P84245 (Rat)
Storage/Stabi	lity Ship	oed at 4°C. Upon delivery ali	quot and store at -20°C for one year. Avoid
	freez	e/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

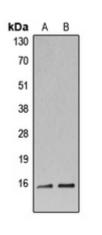
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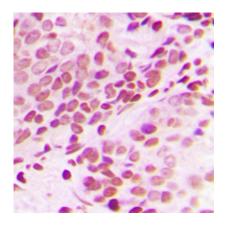
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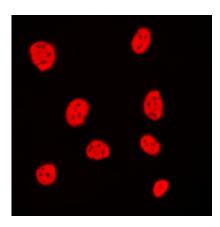
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Western blot analysis of Histone H3 expression in HeLa (A), A431 (B) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 15 kD)



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 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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