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HIST1H2BC (Ab-116) Antibody

 $17KD \rightarrow$

 $10 \text{KD} \rightarrow$

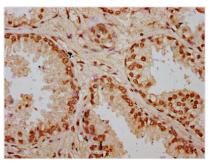
Product Code	CSB-PA010403OA116nacHU	
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.	
Uniprot No.	P62807	
Immunogen	Peptide sequence around site of Lys (116) derived from Human Histone H2B type 1-C/E/F/G/I	
Raised In	Rabbit	
Species Reactivity	Human	
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:100-1:1000, IHC:1:10-1:100, IF:1:1-1:10	
Form	Liquid	
Conjugate	Non-conjugated	
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4	
Purification Method	Antigen Affinity Purified	
Isotype	IgG	
Clonality	Polyclonal	
Alias	Histone H2B type 1-C/E/F/G/I (Histone H2B.1 A) (Histone H2B.a) (H2B/a) (Histone H2B.g) (H2B/g) (Histone H2B.h) (H2B/h) (Histone H2B.k) (H2B/k) (Histone H2B.I) (H2B/I), HIST1H2BC; HIST1H2BE; HIST1H2BF; HIST1H2BG; HIST1H2BI, H2BFL; H2BFH; H2BFG; H2BFA; H2BFK	
Immunogen Species	Homo sapiens (Human)	
Research Area	Epigenetics and Nuclear Signaling	
Target Names	HIST1H2BC	
Image	$72KD \rightarrow$ Western Blot $72KD \rightarrow$ Positive WB detected in: K562 whole cell lysate $55KD \rightarrow$ All lanes: HIST1H2BC antibody at 0.72µg/ml Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 14 kDa Observed band size: 14 kDa	

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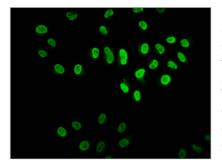


CUSABIO TECHNOLOGY LLC

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IHC image of CSB-PA010403OA116nacHU diluted at 1:20 and staining in paraffin-embedded human prostate cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence staining of Hela cells with CSB-PA010403OA116nacHU at 1:2.5, counterstained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).