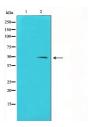


Phospho-Cyclin E1 (Thr395) Ab

Cat.#: AF3235 Size: 100ul,200ul	Concn.: 1mg/ml Source: Rabbit	Mol.Wt.: 48kDa Clonality: Polyclonal
Application:	WB 1:500-1:2000 IHC 1:50-1:200, IF/ICC 1:100-1:500	
Reactivity:	Human	
Purification:	The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.	
Specificity:	Phospho-Cyclin E1 (Thr395) Ab detects endogenous levels of Cyclin E1 only when phosphorylated at Threonine 395	
Immunogen:	A synthesized peptide derived from human Cyclin E1 around the phosphorylation site of Threonine 395	
Uniprot:	P24864	
Description:	a member of the highly conserv members are characterized by a protein abundance through the regulators of CDK kinases.	a dramatic periodicity in
Subcellular Location:	Nucleus.	
Tissue Specificity:	Highly expressed in testis and p bronchial epithelial cells.	olacenta. Low levels in
Similarity:	Belongs to the cyclin family. Cyclin E subfamily.	
Storage Condition and Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Store at -20 °C.Stable for 12 months from date of receipt	



Western blot analysis of Cyclin E1 phosphorylation expression in Paclitaxel treated HeLa whole cell lysates, The lane on the left is treated with the antigen-specific peptide.





AF3235 staining MCF7 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted Ab in 5% w/v milk , 1X TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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