

For Research Use Only

Cyclin E1 Polyclonal antibody

Catalog Number: 11554-1-AP

Featured Product

267 Publications



Basic Information

Catalog Number: 11554-1-AP	GenBank Accession Number: BC035498	Purification Method: Antigen affinity purification
Size: 150ul , Concentration: 550 µg/ml by Nanodrop;	GeneID (NCBI): 898	Recommended Dilutions: WB 1:500-1:2000 IHC 1:400-1:1600
Source: Rabbit	Full Name: cyclin E1	
Isotype: IgG	Calculated MW: 410 aa, 47 kDa	
Immunogen Catalog Number: AG2110	Observed MW: 47 kDa	

Applications

Tested Applications: FC, IHC, WB, ELISA	Positive Controls:
Cited Applications: CoIP, IF, IHC, WB	WB : HT-29 cells, NIH/3T3 cells, mouse heart tissue, HeLa cells, Jurkat cells, human lung tissue, MCF-7 cells, HepG2 cells, K-562 cells, human placenta tissue
Species Specificity: human, mouse	IHC : mouse testis tissue, human placenta tissue
Cited Species: human, rat, mouse, zebrafish, pig, Artemia sinica	
Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0	

Background Information

Cyclin E1 (CCNE1) is a member of the highly conserved cyclin family, whose members are characterized by a dramatic periodicity in protein abundance through the cell cycle. CCNE1, an essential cyclin activating Cdk2, regulates the G1-S phase transition of the mammalian cell division cycle. Its timing expression plays a direct role in the initiation of DNA replication, the control of histone biosynthesis, and the centrosome cycle. CCNE1 is associated with disease progression in various malignancies and is associated clinically with poor prognosis. Two bands of Cyclin E1 were expressed in U2OS and MDA-MB-231 cells (PMID:9858585, PMID: 24112607).

Notable Publications

Author	Pubmed ID	Journal	Application
Yong Zhu	34660260	Front Oncol	WB
Yong-Li Zhang	34679694	Antioxidants (Basel)	WB
Hao Yang	27708221	Oncotarget	WB

Storage

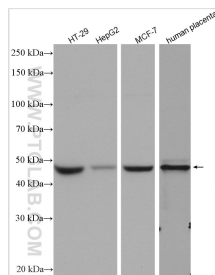
Storage:
Store at -20°C. Stable for one year after shipment.
Storage Buffer:
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

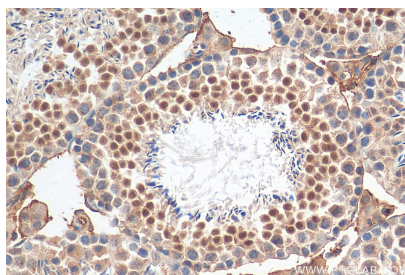
For technical support and original validation data for this product please contact:
T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA)
E: proteintech@ptglab.com
W: ptglab.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

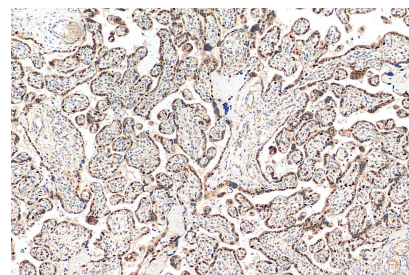
Selected Validation Data



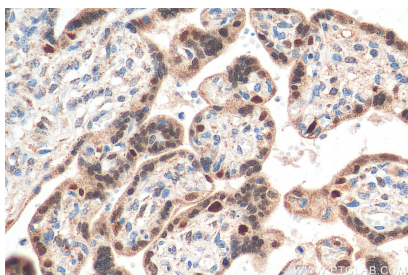
Various lysates were subjected to SDS PAGE followed by western blot with 11554-1-AP (Cyclin E1 antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours.



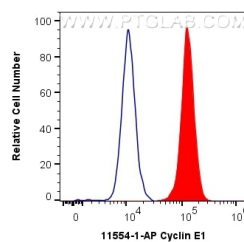
Immunohistochemical analysis of paraffin-embedded mouse testis tissue slide using 11554-1-AP (Cyclin E1 antibody) at dilution of 1:800 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



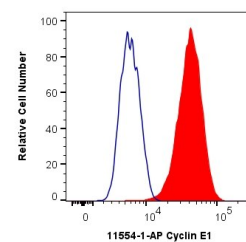
Immunohistochemical analysis of paraffin-embedded human placenta tissue slide using 11554-1-AP (Cyclin E1 antibody) at dilution of 1:800 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



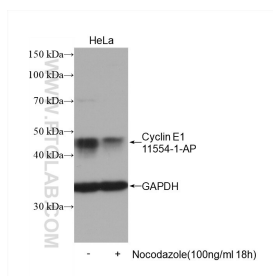
Immunohistochemical analysis of paraffin-embedded human placenta tissue slide using 11554-1-AP (Cyclin E1 antibody) at dilution of 1:800 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



1X10⁶ MCF-7 cells were intracellularly stained with 0.4 ug Anti-Human Cyclin E1 (11554-1-AP) and CoraLite®488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L) at dilution 1:1000 (red), or 0.4 ug Control Antibody. Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).



1X10⁶ HeLa cells were intracellularly stained with 0.4 ug Anti-Human Cyclin E1 (11554-1-AP) and CoraLite®488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L) at dilution 1:1000 (red), or 0.4 ug Isotype Control. Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).



Non-treated HeLa and nocodazole treated HeLa cells were subjected to SDS PAGE followed by western blot with 11554-1-AP (Cyclin E1 antibody) at dilution of 1:1200 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.