

bs-0573R**[Primary Antibody]**

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Cyclin E1 Rabbit pAb

DATASHEET

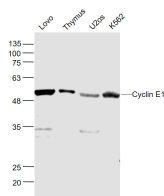
Host: Rabbit**Isotype:** IgG**Clonality:** Polyclonal**GeneID:** 25729**SWISS:** P39949**Target:** Cyclin E1**Immunogen:** KLH conjugated synthetic peptide derived from rat Cyclin E: 375-411/411.**Purification:** affinity purified by Protein A**Concentration:** 1mg/ml**Storage:** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

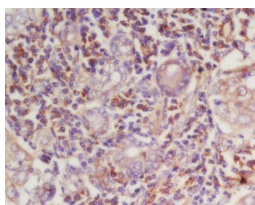
Background: Cyclin E is a regulatory subunit of Cdk2 and controls G1 / S transition during the mammalian cell cycle. Multiple isoforms of Cyclin E are only expressed in tumors but not in normal tissue, suggesting a post transcriptional regulation of Cyclin E. In vitro analyses indicated that these truncated variant isoforms of Cyclin E are able to phosphorylate histone H1. Alterations in the Cyclin E protein have been implicated as indicators of worse prognosis in various cancers.

Applications: WB (1:500-2000)**IHC-P** (1:100-500)**IHC-F** (1:100-500)**IF** (1:100-500)**Flow-Cyt** (1µg/Test)**Reactivity:** Human, Mouse, Rat**Predicted MW.:** 45 kDa**Subcellular Location:** Nucleus

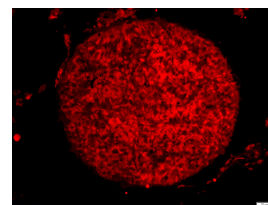
VALIDATION IMAGES



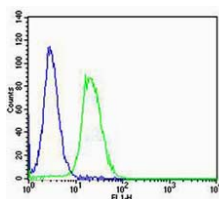
Sample: Lovo (Human) Cell Lysate at 30 ug
Thymus (Mouse) Lysate at 40 ug U2os (Human)
Cell Lysate at 30 ug K562 (Human) Cell Lysate at
30 ug Primary: Anti- Cyclin E1 (bs-0573R) at
1/1000 dilution Secondary: IRDye800CW Goat
Anti-Rabbit IgG at 1/20000 dilution Predicted
band size: 45 kD Observed band size: 50 kD



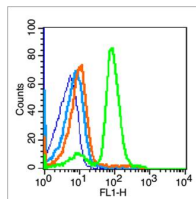
Tissue/cell: human laryngocarcinoma; 4%
Paraformaldehyde-fixed and paraffin-
embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block
endogenous peroxidase by 3% Hydrogen
peroxide for 30min; Blocking buffer (normal goat
serum, C-0005) at 37°C for 20 min; Incubation:
Anti-Cyclin-E Polyclonal Antibody,
Unconjugated(bs-0573R) 1:200, overnight at 4°C,
followed by conjugation to the secondary
antibody(SP-0023) and DAB(C-0010) staining



Tissue/cell: rat testis tissue; 4%
Paraformaldehyde-fixed and paraffin-
embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min;
Blocking buffer (normal goat serum, C-0005) at
37°C for 20 min; Incubation: Anti-Cyclin E
Polyclonal Antibody, Unconjugated(bs-0573R)
1:200, overnight at 4°C; The secondary antibody
was Goat Anti-Rabbit IgG, Cy3
conjugated(bs-0295G-Cy3) used at 1:200 dilution
for 40 minutes at 37°C.



Cell: NIH/3T3 Concentration: 1:100
Host/Isotype: Rabbit/IgG Flow cytometric
analysis of primary antibody (Cat#: bs-0573R) on
NIH/3T3 (green) compared with Rabbit IgG
isotype control in the absence of primary
antibody (blue) followed by Alexa Fluor 488-



Blank control (blue line): Mouse spleen cells
(blue). Primary Antibody (green line): Rabbit
Anti-Cyclin E1 antibody (bs-0573R) Dilution: 1µg
/10⁶ cells; Isotype Control Antibody (orange
line): Rabbit IgG . Secondary Antibody (white
blue line): Goat anti-rabbit IgG-FITC Dilution: 1µg

Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

conjugated goat anti-rabbit IgG(H+L) secondary antibody .

/test. Protocol The cells were fixed with 70% ethanol (overnight at 4°C) and then permeabilized with 0.1% PBS-Tween for 20 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The cells were then incubated in 1 X PBS/2%BSA/10% goat serum to block non-specific protein-protein interactions followed by the antibody for 15 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

— SELECTED CITATIONS —

- **[IF=11.205]** Cong Lan. et al. Inhibition of DYRK1A, via histone modification, promotes cardiomyocyte cell cycle activation and cardiac repair after myocardial infarction. EBIOMEDICINE. 2022 Aug;82:104139 WB ;Rat. 35810562
- **[IF=8.74]** Chen et al. STAT1 inhibits human hepatocellular carcinoma cell growth through induction of p53 and Fbxw7. (2015) Cancer.Cell.Int. 15:111 WB ;Human. 26617467
- **[IF=9.1]** Fan Yang. et al. A Novel ceRNA Axis LOC121818100/Novel-miR-400/SSRP1 Regulated Muscle Growth and Injury Repair in Sheep. J CACHEXIA SARCOPENI. 2025 Jun;16(3):e13836 WB ;Sheep. 40468947
- **[IF=5.652]** Haijun Sun. et al. WD Repeat Domain 43 promotes malignant progression of non-small cell lung cancer by regulating CDK2. INT J BIOCHEM CELL B. 2022 Aug;;106293 WB ;Human. 10.1016/j.biocel.2022.106293
- **[IF=6.1]** Cuifang Chang. et al. The orphan GPR50 receptor interacting with TβRI induces G1/S-phase cell cycle arrest via Smad3-p27/p21 in BRL-3A cells. BIOCHEM PHARMACOL. 2022 Aug;202:115117 WB ;Rat. 35671788