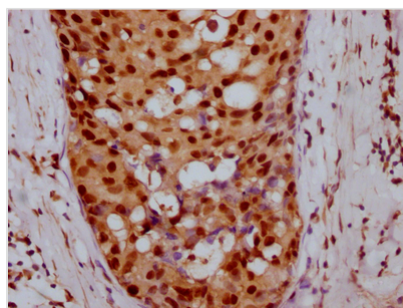




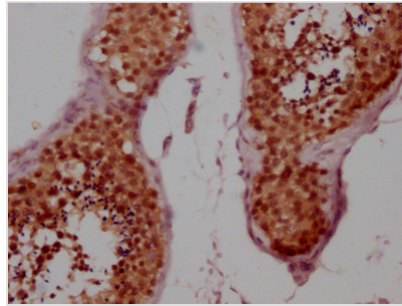
# CCNE1 Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA968740A0HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P24864
<b>Immunogen</b>	A synthesized peptide derived from human Cyclin E1
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	ELISA, IHC, IF; Recommended dilution: IHC:1:50-1:200, IF:1:20-1:200
<b>Relevance</b>	Essential for the control of the cell cycle at the G1/S (start) transition.
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	Rabbit IgG
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Recombinant Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Epigenetics and Nuclear Signaling; Cancer; Cell biology
<b>Gene Names</b>	CCNE1
<b>Clone No.</b>	3A5

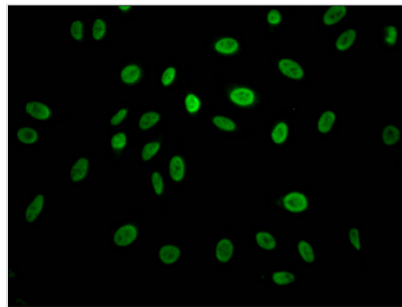
## Image



IHC image of CSB-RA968740A0HU diluted at 1:100 and staining in paraffin-embedded human breast cancer performed on a Leica Bond™ 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? overnight. The primary is detected by a Goat anti-rabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.



IHC image of CSB-RA968740A0HU diluted at 1:100 and staining in paraffin-embedded human testis tissue performed on a Leica Bond<sup>TM</sup> 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<sup>o</sup> overnight. The primary is detected by a Goat anti-rabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.



Immunofluorescence staining of Hela Cells with CSB-RA968740A0HU at 1:50, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeated by 0.2% TritonX-100, and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4<sup>o</sup>. Nuclear DNA was labeled in blue with DAPI. The secondary antibody was FITC-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L).

## Description

The CCNE1 recombinant antibody production process involves four key steps: sequencing the CCNE1 monoclonal antibody gene, inserting the gene into a plasmid vector, transfecting the recombinant vector into a host cell line, and purifying the CCNE1 recombinant monoclonal antibody from the cell culture supernatant via affinity chromatography. The CCNE1 monoclonal antibody is derived from CCNE1 antibody-producing hybridomas, and the production process involves using a synthesized peptide from human CCNE1 as the immunogen. This CCNE1 recombinant monoclonal antibody is recommended for use in detecting human CCNE1 protein through ELISA, IHC, and IF applications.

The CCNE1 protein plays a critical role in the regulation of the cell cycle. It is a regulatory protein that binds to and activates cyclin-dependent kinase 2 (CDK2) to form a complex that is required for the progression of cells from the G1 to the S phase of the cell cycle. This complex phosphorylates various substrates, leading to DNA replication and cell division. The expression of CCNE1 is tightly regulated, with elevated levels of expression associated with various types of cancer. In addition to its role in the cell cycle, CCNE1 has also been implicated in DNA repair and apoptosis.