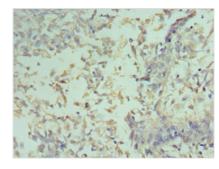




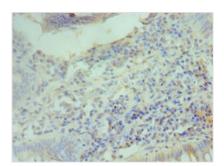


CCND2 Antibody

Product Code	CSB-PA004812ESR2HU
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P30279
Immunogen	Recombinant Human G1/S-specific cyclin-D2 protein (1-289AA)
Raised In	Rabbit
Species Reactivity	Human, Mouse
Tested Applications	ELISA, WB, IHC,IF; Recommended dilution: WB:1:500-1:5000, IHC:1:20-1:500, IF:1:50-1:200
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	PBS with 0.02% sodium azide, 50% glycerol, pH7.3.
Purification Method	Antigen Affinity purified
Isotype	IgG
Clonality	Polyclonal
Product Type	Polyclonal Antibody
Immunogen Species	Homo sapiens (Human)
Research Area	Cell Biology
Target Names	CCND2
Image	Immunohistochomistry of paraffin embodded



Immunohistochemistry of paraffin-embedded human cervical cancer using CSB-PA004812ESR2HU at dilution of 1:100



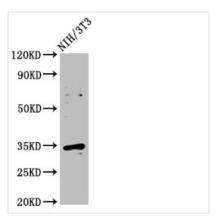
Immunohistochemistry of paraffin-embedded human colon cancer using CSB-PA004812ESR2HU at dilution of 1:100

CUSABIO TECHNOLOGY LLC









Western Blot

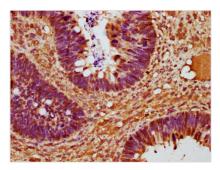
Positive WB detected in: NIH/3T3 whole cell

All lanes: CCND2 antibody at 2.25µg/ml

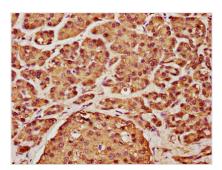
Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

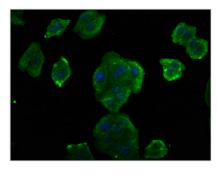
Predicted band size: 34 kDa Observed band size: 34 kDa



IHC image of CSB-PA004812ESR2HU diluted at 1:225 and staining in paraffin-embedded human ovarian 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.



IHC image of CSB-PA004812ESR2HU diluted at 1:225 and staining in paraffin-embedded human pancreatic 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 HepG2 cells with CSB-PA004812ESR2HU at 1:75, 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).