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PathScan® Phospho-cdc2 (Thr161) Sandwich ELISA Kit

UniProt ID: #P06493
Entrez-Gene Id: #983

For Research Use Only. Not for Use in Diagnostic Procedures.

Product Includes	Product #	Quantity	Color	Storage Temp
TMB Substrate	7004	11 ml	Colorless	+4C
STOP Solution	7002	11 ml	Colorless	+4C
Sealing Tape	54503	2 ea		+4C
ELISA Wash Buffer (20X)	9801	25 ml	Colorless	+4C
Cell Lysis Buffer (10X)	9803	15 ml	Yellowish	-20C

Kit contents scale proportionally with size, except sealing tape.

Example: The V1 kit contains 5X the listed quantities above, but will exclude the sealing tape.

The microwell plate is supplied as 12 8-well modules - Each module is designed to break apart for 8 tests.

Description

CST's PathScan® Phospho-cdc2 (Thr161) Sandwich ELISA Kit is a solid phase sandwich enzyme-linked immunosorbent assay (ELISA) that detects endogenous levels of phospho-cdc2 (Thr161) protein. A Phospho-cdc2 (Thr161) Rabbit polyclonal Ab #9114* has been coated onto the microwells. After incubation with cell lysates, phospho-cdc2 (Thr161) protein is captured by the coated antibody. Following extensive washing, cdc2 Mouse mAb #2658* is added to detect the captured phospho-cdc2 protein. Anti-mouse IgG, HRP-linked Antibody #7076* is then used to recognize the bound detection antibody. HRP substrate, TMB, is added to develop color. The magnitude of optical density for this developed color is proportional to the quantity of phospho-cdc2 (Thr161) protein.

* Antibodies in kit are custom formulations specific to kit.

Specificity/Sensitivity

CST's PathScan® Phospho-cdc2 (Thr161) Sandwich ELISA Kit detects endogenous levels of phospho-cdc2 (Thr161). As shown in Figure 1, using the Phospho-cdc2 (Thr161) ELISA Kit #7184, a significant induction of phospho-cdc2 (Thr161) is detected in HeLa cells treated with Interleukin-4. This kit detects proteins from the indicated species, as determined through in-house testing, but may also detect homologous proteins from other species.

Background

The entry of eukaryotic cells into mitosis is regulated by cdc2 kinase activation, a process controlled at several steps including cyclin binding and phosphorylation of cdc2 at Thr161 (1). However, the critical regulatory step in activating cdc2 during progression into mitosis appears to be dephosphorylation of cdc2 at Thr14 and Tyr15 (2). Phosphorylation at Thr14 and Tyr15, resulting in inhibition of cdc2, can be carried out by Wee1 and Myt1 protein kinases (3,4). The cdc25 phosphatase may be responsible for removal of phosphates at Thr14 and Tyr15 and subsequent activation of cdc2 (1,5).

Background References

1. Atherton-Fessler, S. et al. (1994) *Mol Biol Cell* 5, 989-1001.
2. Norbury, C. et al. (1991) *EMBO J* 10, 3321-9.
3. McGowan, C.H. and Russell, P. (1993) *EMBO J* 12, 75-85.
4. Wells, N.J. et al. (1999) *J Cell Sci* 112 (Pt 19), 3361-71.
5. Hunter, T. (1995) *Cell* 80, 225-36.

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Revision 2

#7184

PathScan® Phospho-cdc2 (Thr161) Sandwich ELISA Kit

