



**Category:** Monoclonal Antibodies

**Product Name:** Mouse Monoclonal Antibody to CHK2

**Catalog Number:** MAB-606020223

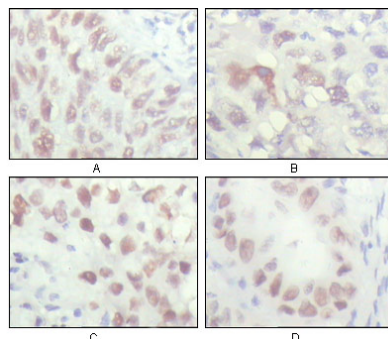
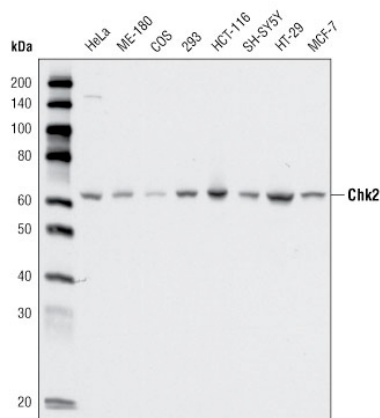


Figure 2: Immunohistochemical analysis of paraffin-embedded human lung carcinoma (A), liver carcinoma (B), breast carcinoma (C) and kidney carcinoma (D), showing nuclear localization with DAB staining using CHK2 mouse mAb.

Lot#:  
Clone#: 1C12B8  
Host and isotype: Mouse IgG2b  
Size: 0.1ml  
MW: 61kDa  
Aliases: CDS1; LFS2; CHEK2  
Entrez Gene: 11200  
Species reactivity: Human

Figure 1: Western blot analysis using CHK2 mouse mAb against cell lysate from various cell types.

**Description** CHK2: CHK2 checkpoint homolog (*S. pombe*). In response to DNA damage and replication blocks, cell cycle progression is halted through the control of critical cell cycle regulators. The protein encoded by this gene is a cell cycle checkpoint regulator and putative tumor suppressor. It contains a forkhead-associated protein interaction domain essential for activation in response to DNA damage and is rapidly phosphorylated in response to replication blocks and DNA damage. When activated, the encoded protein is known to inhibit CDC25C phosphatase, preventing entry into mitosis, and has been shown to stabilize the tumor suppressor protein p53, leading to cell cycle arrest in G1. In addition, this protein interacts with and phosphorylates BRCA1, allowing BRCA1 to restore survival after DNA damage. Mutations in this gene have been linked with Li-Fraumeni syndrome, a highly penetrant familial cancer phenotype usually associated with inherited mutations in TP53. Also, mutations in this gene are thought to confer a predisposition to sarcomas, breast cancer, and brain tumors. This nuclear protein is a member of the CDS1 subfamily of serine/threonine protein kinases. Three transcript variants encoding different isoforms have been found for this gene.

**Immunogen** Purified recombinant fragment of human CHK2 (aa481-531) expressed in *E. coli*.

**Application** Western Blotting: 1/500 - 1/2000.  
Immunohistochemistry: 1/200 - 1/1000.  
Immunofluorescence: 1/200 - 1/1000.  
ELISA: Propose dilution 1/10000.  
Not yet tested in other applications.  
Determining optimal working dilutions by titration test.

**Formulation** Ascitic fluid containing 0.03% sodium azide.

**Storage** Store at 4°C, for long term storage, store at -20°C.

**Related product**

**References** 1. Int J Cancer. 2007 Dec 15;121(12):2661-7.  
2. Nat Rev Cancer. 2007 Dec;7(12):925-36.  
3. Carcinogenesis. 2008 Apr;29(4):762-5.

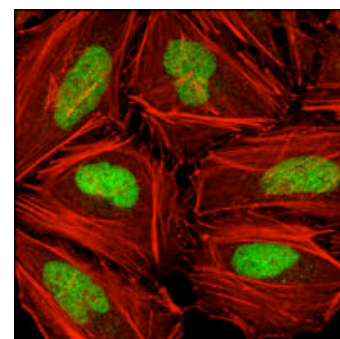


Figure 3: Confocal immunofluorescence analysis of HeLa cells using CHK2 mouse mAb (green), showing nuclear localization. Red: Actin filaments have been labeled with DY-554 phalloidin.

**For Research Use Only**

**Contact:** Antagene, Inc. | Tel: 1 (866) 964-2589 | Fax: 1 (888) 225-1868 | Email: [Info@antageneinc.com](mailto:Info@antageneinc.com)