

## Anti-MCM7 antibody, rabbit polyclonal IgG

70-120 100 μg

**Key words:** DNA replication licensing factor, MCM complex, DNA replication initiation, G1/S transition, DNA damage response, DNA helicase,

**Function:** MCM7 (human; 718 aa, 80 kDa) acts as component of the MCM2-7 complex (MCM complex) which is the putative replicative helicase essential for 'once per cell cycle' DNA replication initiation and elongation in eukaryotic cells. The active ATPase sites in the MCM2-7 ring are formed through the interaction surfaces of two neighboring subunits such that a critical structure of a conserved arginine finger motif is provided in trans relative to the ATP-binding site of the Walker A box of the adjacent subunit. The six ATPase active sites, however, are likely to contribute differentially to the complex helicase activity. Required for S-phase checkpoint activation upon UV-induced damage.

## Applications

- 1) Western blotting (1/1,000~1/5,000 dilution).
- 2) Immunoprecipitation (assay dependent)
- Immunofluorescence staining (1/200~1/1,000 dilution).
   Other applications have not been tested.

**Immunogen:** Purified His6-tagged human MCM7 protein encompassing 562 -719 amino acids.

**Reactivity:** Reacts with human mouse, rat and hamster. Not tested in other species. **Product:** Purified IgG from the rabbit antiserum. 1 mg/ml in PBS, 50% glycerol, filter-sterilized. Azide- and carrier-free.

Storage: Shipped at 4°C. Upon arrival, spin-down, aliquot and store at -20°C.
Data Link UniProtKB/Swiss-Prot P33993 MCM7\_HUMAN

References: This antibody was described and used in the following publications.

- Fujita M et al. (1996) hCDC47, a human member of the MCM family. Dissociation of the nucleus-bound form during S phase. J Biol Chem. 271:4349-54. <u>PMID 8626784</u> Free Article. WB, IP, IF
- Fujita M. et al. (1997) In vivo interaction of human MCM heterohexameric complexes with chromatin. Possible involvement of ATP. J Biol Chem. 272:10928-35. <u>PMID</u> <u>9099751</u> Free Article. WB, IP
- Fujita M. et al. (2002) Nuclear organization of DNA replication initiation proteins in mammalian cells. J Biol Chem. 277:10354-61. <u>PMID 11779870</u> Free Article. WB, IP, IF.



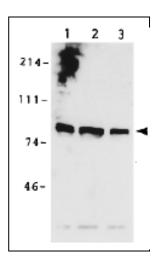
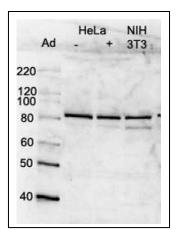
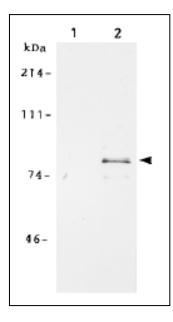


Fig. 1. Identification of MCM7 protein in whole cell extracts of human cells by western blotting using anti-MCM7 antibody. Lane 1; SiHa cells Lane 2; C33A cells Lane3; WI38 cells All cell lines are cervical cancer derived. Samples are obtained from approximately 10<sup>5</sup> cells.



## Fig. 2 Identification of MCM7 protein in whole cell extracts of human and mouse cells by western blotting using anti-MCM7 antibody.

- Lane 1. Size marker proteins in kDa.
- Lane 2. Extract of HeLa cells untreated (-).
- Lane 3. Extract of HeLa cells treated with 100 nM adriamycin for 24 hr (+)
- Lane 4. Extract of NIH3T3 (mouse) cells.
- Anti-MCM7 antibody was used at 1/1,000 dilution.
- \* Indicates the band of MCM7 protein



## Fig. 3. Immunoprecipitation of MCM7 protein from crude extract of human fibroblast cell line WI38 by using anti-MCM7 antibody.

Lane 1; Immunoprecipitation with pre-immune serum

Lane 2; Immnoprecipitation with anti-MCM7 antiserum.

Cells were labeled with  $S^{35}$  methionine and MCM7 was immunoprecipitated with the anti-MCM7 antibody followed by SDS-PAGE and autoradiography.



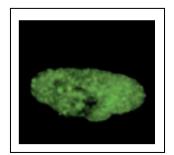


Fig. 4. Immunofluorecence staining and confocal microscopic analysis of MCM7 in  $G_1$  phase HeLa cell nucleus by using anti-MCM7 antibody after treatment with protein cross-linking reagent, DSP and chromatin extraction. The processed cells were fixed with formaldehyde before staining.