

Anti-p53 acetyl-K382 antibody, monoclonal (2B7E4)

71-133 100 µg

p53 mutants are found in more than half of human cancers and are considered as the most important human cancer related gene. *p53* is detected at 53kD position by electrophoresis and is composed of 393 amino acids. In the unstressed normal cells, the *p53* level is low and it is inactive. However, with stress, especially with DNA damage, it is activated to promote arrest of cell cycle and repair of DNA damage, or induction of apoptosis. The functions and stability of *p53* are regulated by phosphorylation of serine and threonine, and acetylation of lysine at various sites in the molecule.

Acetylation of lysine 382 (acetyl-K382) of *p53* occurs after DNA damage and is catalyzed by the p300/CBP acetyltransferase, which stabilizes *p53* protein (ref 1).

Applications

1) Western blotting (~1 µg/ml) Other applications have not been tested.

Immunogen: Synthetic peptide containing acetyl-Lys382 of human *p53*

Specificity: Reacts with human *p53* acetylated at Lys382. Other species have not been tested.

Isotype: Mouse IgG1 (κ)

Form: Purified IgG 1mg/ml in PBS (pH 7.4), 50% glycerol, sterilized by filtration

Storage: Sent at 4°C or -20°C. Upon arrival, briefly centrifuge and store at -20°C

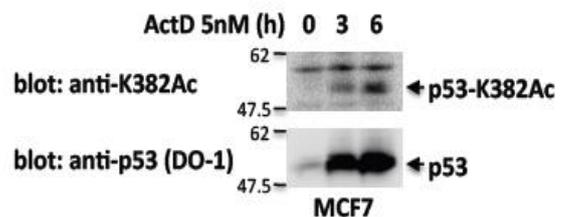
Data Link UniProtKB/Swiss-Prot [P04637](#) (P53_HUMAN)

Reference

1. Bode AM & Dong Z "Post-translational modification of *p53* in tumorigenesis" *Nature Rev Cancer* 4:793-805 (2004) PMID:[15510160](#)

Fig.1 Identification of *p53* protein, whose Lys382 is acetylated, by Western blotting with 2B7E4 antibody.

MCF7 cells in culture were treated with actinomycin D at 5 nM for the indicated periods and the cell extracts were analyzed by Western blotting with anti-*p53* acetyl-K382 antibody (2B7E4) and omnipotent anti-*p53* antibody (DO-1). Acetylation of *p53* at K382 was induced by the DNA damaging treatment.



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