

Product code	71-161
Size	100 µg
Storage	-20°C
Concentration	1.0 mg/ml
Buffer	PBS- with 50% glycerol
Purity	Purified IgG fraction with protein A from hybridoma cell culture medium.
Immunogen	Synthetic peptide containing phospho-Ser62 of cMyc
Isotype	Mouse IgG2b к
Reactivity	Human and mouse.
Special notes	Specificity of this antibody have been independently validated by siRNA
Application	 Western blotting (~1µg/ml, Fig.1) Immunofluorescence staining (0.25~1 µg/ml) Immunohistochemistry (5 µg/ml), Paraffin-embedded Flow cytometry (1 µg for 10⁶ cells) Indirect ELISA (Assay dependent concentration)
Background	cMyc is a proto-oncogene, which is overexpressed in a wide range of human cancers. Myc gene encodes a transcription factor that regulates a great number of genes through binding on Enhancer Box sequences (E-boxes) and recruiting histone acetyltransferase. It can also act as a transcriptional repressor. It regulates cell growth, apoptosis, differentiation and stem cell self- renewal. Previous studies on the phosphorylation of c-Myc have suggested functional association between phosphorylation at Thr58/Ser62 by glycogen synthase kinase 3, cyclin dependent kinase, ERK2 and C-Jun N terminal Kinase (JNK), cell proliferation and cell cycle regulation. Phosphorylation at Ser62 is required for Ras-induced stabilization and is prerequisite for phosphorylation at Thr58 for its degradation (ref.1).
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Anti-cMyc phospho-Ser62 antibody, mouse monoclonal (33A12E10), Validated

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Data Images: 71-161 Anti-cMyc phospho-Ser62 antibody, mouse monoclonal (33A12E10), Validated



Fig.1. Identification of cMyc protein whose Ser62 is phosphorylated by Western blotting.

Samples:Crude cell extracts of AGS (gastric adenocarcinoma) cells.

Scr; Scrambled siRNA was introduced into the cells as a negative control.

Neg.Contol; Negative control siRNA was transfected.

Myc1; siRNA for cMyc was transfected. (The data was provided by Drs.A. Khanna and J.Westermarck of University of Tampere)



Fig.2. Immunofluorescence staining of cMyc phosho-Ser62 in HeLa cells Left: Cells stained with anti-cMyc pS62 antibody (green) and DAPI (blue) Right: Proximity Ligation Analysis with anti-cMyc pS62 and CIP2A antibodies, association of cMyc pS62 with CIP2A (red) in nuclei (DAPI, blue) Images kindly provided by Prof Westermarck J and Dr. Qiao X. For details refer to Ref.1



DAPI

Merge



Fig.3. Immunofluorescence staining of cMyc phosho-Ser62 in nuclei of HeLa cells.

- 1. HeLa cells were fixed with 4% paraformaldehyde overnight, permealized with 0.25% Triton X-100 in PBS for 10 min.
- 2. Incubate cells with 1.5% BSA in PBS for 30 min to block non-specific binding of the antibodies. Incubate the cells with 1/4,000 diluted anti-cMyc p62 antibody in 1% BSA in PBS at 4° C overnight.
- 3. Incubate cells with a secondary antibody, goat anti-mouse IgG conjugated with Alex 488, at 1/1,000 dilution in 1% BSA for 1 hr at room temperature.
- 4. Nucleus (DNA) was stained with DAPI

References: This product was used in the following Publications.

- Myant K et al. Serine 62-Phosphorylated MYC Associates with Nuclear Lamins and Its Regulation by CIP2A Is Essential for Regenerative Proliferation. <u>Cell Rep.</u> 2015 Aug 11;12(6):1019-31. PMID: <u>26235622</u>. WB, IF (human, mouse)
- 2. Tibbitts DC *et al.* Studying c-Myc serine 62 phosphorylation in leukemia cells: concern over antibody cross-reactivity. *Blood* 119:5334-5 (2012).<u>PubMed: 22653959</u> **WB, IP (human)**
- Mathiasen DP. Identification of a c-Jun N-terminal kinase-2-dependent signal amplification cascade that regulates c-Myc levels in ras transformation. <u>Oncogene.</u> 2012 Jan 19;31(3):390-401. <u>PubMed: 21706057</u> WB (mouse)
- 4. Wang X. *et al.* Phosphorylation regulates c-Myc's oncogenic activity in the mammary gland. Cancer Res. 2011 Feb 1; 71(3): 925–936. <u>PubMed: 3077809</u> **WB (human)**
- 5. Khanna A. MYC-dependent regulation and prognostic role of CIP2A in gastric cancer. <u>J Natl Cancer Inst.</u> 2009 Jun 3;101(11):793-805. <u>PubMed: 19470954</u> WB (human)