

## POLYCLONAL ANTIBODY

For research use only. Not for clinical diagnosis.

## Catalog No. BAM-63-135-EX

## Anti-Ppa2 (S. pombe) antibody, rabbit serum

## **BACKGROUND**

Schizosaccharomyces pombe Ppa2 is a type 2A-like serine/threonine-protein phosphatase catalytic subunit whose polypeptide sequence has ~80% identity to those of mammalian type 2A phosphatases. Ppa2 determines the sensitivity to okadaic acid, which is an inhibitor of protein serine/threonine phosphatases. The loss of the ppa2 gene causes cells to be hypersensitive to the okadaic acid. Ppa2 plays important roles in cell cycle control. It may be involved in controlling the entry into mitosis, possibly acting as an inhibitor (ref.1). Ppa2 is abundant in the cytoplasm, in contrast to the type 1-like phosphatase Dis2, which is enriched in the nucleus. Thus Ppa2 may perform major functions outside the nucleus.

**Product type** Primary antibodies

Host Rabbit Source serum **Form** Liquid

Rabbit antiserum added with 0.05 % sodium azide

Volume 100 µl

Concentration

Specificity Ppa2

**Antigen** Recombinant C-terminal polypeptide (26kDa) of S. pombe Ppa2 (Ref. 1)

Isotype

**Application notes** Immunoblotting, Immunofluorescence microscopy, Immunoprecipitation

Recommended use

Recommended dilutions

Immunoblotting (dilution: 1/1000)

Optimal dilutions/concentrations should be determined by the end user.

Data Link: Swiss-Prot P23636

Staining Pattern

**Cross reactivity** The antibody recognized both Ppa1 and Ppa2 polypeptides in S. pombe because of their

high amino acid similarity (~80% identity) (Fig.1 and ref. 1).

Storage -20°C (long period, -70°C)

References 1) Kinoshita N et al "Negative regulation of mitosis by the fission yeast protein phosphatase ppa2." Genes Dev 7:

1059-1071 (1993) PMID: 8389306

2) Kinoshita K et al "The regulatory subunits of fission yeast protein phosphatase 2A (PP2A) affect cell

morphogenesis, cell wall synthesis and cytokinesis." Genes Cell 1:29-45 (1996) PMID: 9078365



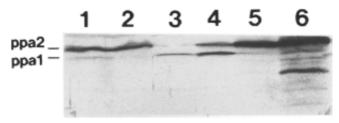


Fig.1 Identification of Ppa1 and Ppa2 proteins. An immunoblot with anti-ppa2 antibody is shown (ref.1).

lane 1: Wild-type S. pombe

lane 2: ppa1 lane 3: ppa2

lane 4: Wild-type carrying a multicopy plasmid with ppa1 gene lane 5: Wild-type carrying a multicopy plasmid with ppa2 gene

lane 6:Wild-type carrying a multicopy plasmid with ADH promoter ligated with ppa2 gene

The positions of ppa1 (36 kDa) and ppa2 (39 kDa) polypeptide bands are indicated.

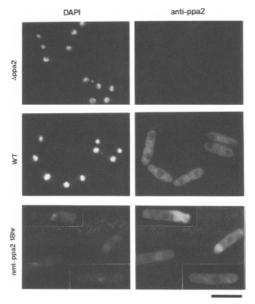


Fig.2 Cellular location of Ppa1 and Ppa2 (ref.1).

Indirect immunofluorescence microscopy of *∆ppa2* deletion, wild type (WT), and wild-type overexpressing *ppa2* (*nmt-ppa2*, 18hr) was done, using anti-ppa2 antibody (right);

The same cells stained by DAPI are also shown (left).

Immunofluorescence was hardly detected in  $\Delta ppa2$  cells, whereas cytoplasmic immunofluorescence was abundant in wild-type cells. Wild-type cells carrying nmt-ppa2 plasmid overexpress Ppa2 protein in the absence of thiamine for 18 hr. Immunofluorescence was enhanced further in the cytoplasm, often accumulated at the nuclear periphery or within restricted domains. The deformation of chromosomal DNA was also visible in overexpressed cells. Bar, 10um.

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