

Sds22 (S. pombe) antibody

Cat. No. GTX64165

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB, ICC/IF, IP
Reactivity	Schizosaccharomyces pombe

Package
100 µl

Applications

Application Note

Immunoblotting (dilution: 1:200-1:500)

Calculated MW	38 kDa. (Note)
Product Note	Specific to S. pombe

Properties

Form	Liquid
Buffer	Serum
Preservative	0.05% Sodium azide
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	Batch dependent (Please refer to the vial label for the specific concentration.)
Immunogen	Recombinant C-terminal region (1.8kb) of S. pombe Sds22
Conjugation	Unconjugated

Note

For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.



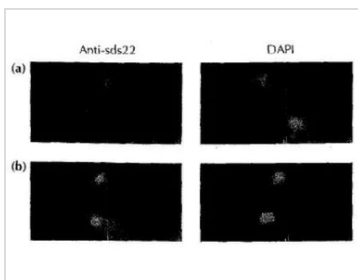
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DATA IMAGES



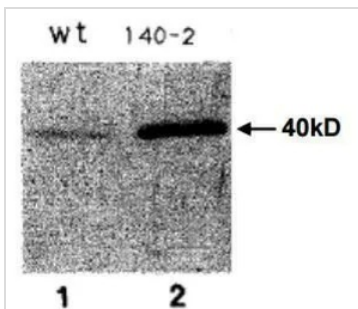
GTx64165 IP Image

Sds22 coprecipitates with Dis2 and Sds21. Yeast extracts of wild type (wt) strain HM123 (lane 1 and 2), *dis2::ura4+* deletion mutant ($\Delta d2$, lane 3), *sds21::ura4+* deletion mutant ($\Delta s21$, lane 4) were immunoprecipitated followed by immunoblotting with the indicated antiserum, to detect the Sds22 or Dis2/Sds21 proteins. Lane 1 was immunoprecipitated with the appropriate preimmune serum, lane 2-4 with the anti-Sds22 serum. (a) denotes anti-Sds22 immunoblot; (b) denotes anti-D2C immunoblot. Anti-D2C crossreacts with both Sds21 and Dis2. Anti-Sds22 antiserum coprecipitates both Dis2 and Sds21 proteins in the wild type strain (lane 2b). Consistently, Sds21 alone is precipitated in the *dis2* deletion mutant (lane 3b), and Dis2 alone is precipitated in the *sds21* deletion mutant (lane 4b).



GTx64165 ICC/IF Image

Sds22 subcellular localization Indirect immunofluorescence microscopy was performed by staining methanol fixed cells with (first column) anti-Sds22 antiserum, or (second column) DAPI to visualize chromosomal DNA. (a) wild type HM123; (b) HM123 carrying multicopy *sds22+* plasmid pHR140-2. Anti-Sds22 antibody stains the cytoplasm as well as the non-chromosomal domain of the nucleus of a wild type strain, as shown in (a). Nuclear staining increases in strains carrying a multicopy *sds22+* plasmid (b).



GTx64165 WB Image

Immunoblot with anti-Sds22 antiserum of yeast extracts from (1) wild type strain HM123, (2) *sds::ura4+* deletion mutant carrying pHR140-2. The 40kD protein band was identified by immunoblot analysis of wild-type strain using anti-Sds22 antisera (lane1). The 40 kD band is enhanced in the *sds22::ura4+* disruption mutant strain that is rescued by the multicopy *sds22+* plasmid pHR140-2 (lane2).



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