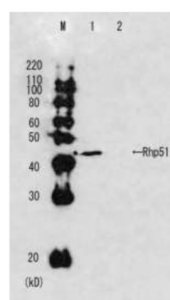


Product no **AS21 4553****Anti-Rhp51 | DNA repair protein rhp51 (*Saccharomyces pombe*)****Product information**

<b>Immunogen</b>	Purified, full-length, recombinant Rhp51 protein from <i>Saccharomyces pombe</i> , UniProt: <a href="#">P36601</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Serum. Contains 0.05% sodium azide.
<b>Format</b>	Liquid
<b>Quantity</b>	100 µl
<b>Storage</b>	Store at -20 °C; make aliquots to avoid repeated freeze-thaw cycles. Please remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tube.

**Application information**

<b>Recommended dilution</b>	Assay dependent (ChIP), 1:500 (IF), 1: 100-1:500 (IP), 1 : 2000 - 1: 5000 (WB)
<b>Expected   apparent MW</b>	39.8   42 kDa
<b>Confirmed reactivity</b>	<i>Saccharomyces pombe</i>
<b>Predicted reactivity</b>	Species of your interest not listed? <a href="#">Contact us</a>
<b>Selected references</b>	<p><a href="#">Akamatsu</a> et al. (2007) Fission yeast Swi5/Sfr1 and Rhp55/Rhp57 differentially regulate Rhp51-dependent recombination outcomes. EMBO J. 2007 Mar 7;26(5):1352-62. doi: 10.1038/sj.emboj.7601582. Epub 2007 Feb 15. PMID: 17304215; PMCID: PMC1817630.</p> <p><a href="#">Lambert</a> et al. (2005). Gross chromosomal rearrangements and elevated recombination at an inducible site-specific replication fork barrier. Cell. 2005 Jun 3;121(5):689-702. doi: 10.1016/j.cell.2005.03.022. PMID: 15935756. (Immunofluorescence)</p> <p><a href="#">Akamatsu</a> et al. (2003) Two different Swi5-containing protein complexes are involved in mating-type switching and recombination repair in fission yeast. Proc Natl Acad Sci U S A. 2003 Dec 23;100(26):15770-5. doi: 10.1073/pnas.2632890100. Epub 2003 Dec 8. PMID: 14663140; PMCID: PMC307643. (Immunoprecipitation, Western Blot)</p> <p><a href="#">Kibe</a> et al. (2003). Fission yeast Rhp51 is required for the maintenance of telomere structure in the absence of the Ku heterodimer. Nucleic Acids Res. 2003 Sep 1;31(17):5054-63. doi: 10.1093/nar/gkg718. PMID: 12930956; PMCID: PMC212814. (ChIP)</p>

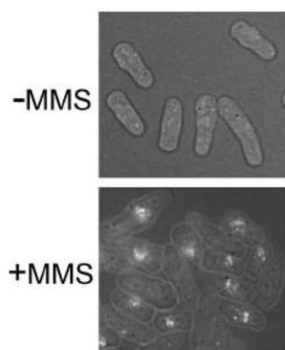


Whole cell extract of *Saccharomyces pombe* (**1**) and Rhp51 deletion mutant (**2**) was separated on SDS-PAGE and blotted to a membrane using wet transfer. Primary antibody was incubated at 1: 2000, followed by washes and incubation with a secondary goat anti-rabbit IgG HRP conjugated antibodies, used at 1: 10 000 1h/RT. Reaction was developed using chemiluminescence following manufacture's recommendations.

This product is **for research use only** (not for diagnostic or therapeutic use)

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*Saccharomyces pombe* cells without (-MMS) and with (+MMS at 0.025 % treatment for 1 h) were subjected to immunofluorescent staining. Primary antibody was used at 1 : 500 dilution.