

## 15016

## Rad54 (D4W3Z) Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 84	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q92698	Entrez-Gene Id: 8438
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation		<b>Dilution</b> 1:1000 1:100		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Rad54 (D4W3Z) Rabbit mAb recognizes endogenous levels of total Rad54 protein. Based on sequence, this antibody is not expected to recognize Rad54B. This antibody also cross-reacts with an unidentified protein of 200 kDa.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly246 of human Rad54 protein.				
Background		DNA double-strand breaks (DSBs) are potentially hazardous lesions that can be induced by ionizing radiation (IR), radiomimetic chemicals, or DNA replication inhibitors. Cells sense and repair DSBs via two distinct but partly overlapping signaling pathways, nonhomologous end joining (NHEJ) and homologous recombination (HR). Defects in both pathways have been associated with human disease, including cancer (1). The DNA repair and recombination protein RAD54-like (Rad54, RAD54L) is a Swi2/Snf2 family DNA helicase that is involved in homologous recombination DNA repair. Rad54 is a double-stranded DNA-dependent ATPase that translocates in a processive manner along double-stranded DNA. The Rad54 helicase interacts with the Rad51 recombinase to regulate its DNA binding and strand exchange activities during homologous recombination (2-4). Mutations in the corresponding <i>RAD54L</i> gene are associated with multiple forms of human cancer, including non-Hodgkin's lymphoma, breast cancer, and parathyroid adenoma (5-8).				
Background References		<ol> <li>Hartlerode, A.J. and Scully, R. (2009) Biochem J 423, 157-68.</li> <li>Sung, P. et al. (2003) J Biol Chem 278, 42729-32.</li> <li>Wright, W.D. and Heyer, W.D. (2014) Mol Cell 53, 420-32.</li> <li>Ceballos, S.J. and Heyer, W.D. (2011) Biochim Biophys Acta 1809, 509-23.</li> <li>Smirnova, M. et al. (2004) J Biol Chem 279, 24081-8.</li> <li>Gonzalez, R. et al. (1999) Br J Cancer 81, 503-9.</li> <li>Carling, T. et al. (1999) Int J Cancer 83, 80-2.</li> <li>Matsuda, M. et al. (1999) Oncogene 18, 3427-30.</li> </ol>				
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).
Western Blot Buffer		IMPORTANT: For western blots insubstantian manhrane with diluted primary antibody in FOV w/v perfet				

**Western Blot Buffer** 

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween\$ 20 at 4°C with gentle shaking, overnight.

**Applications Key** 

W: Western Blotting IP: Immunoprecipitation

**Cross-Reactivity Key** 

**H:** Human

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