

## TDP1 (D8D1B) Rabbit mAb



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

<b>Applications:</b> W, IF-IC	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 70	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q9NUW8	Entrez-Gene Id: 55775
Product Usage Information		<b>Application</b> Western Blotting Immunofluorescence	(Immunocytochem	istry)		<b>Dilution</b> 1:1000 1:50
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		TDP1 (D8D1B) Rabbit mAb recognizes endogenous levels of total TDP1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro587 of human TDP1 protein.				
Background		Topoisomerases are ubiquitous, conserved enzymes that remove DNA supercoils resulting from processes such as chromosome segregation, DNA replication, transcription, and repair (1). Topoisomerase inhibitors such as camptothecin and etoposide trap the enzyme as a DNA-bound intermediate, and these drugs are used to treat multiple human cancers (1,2). Tyrosyl-DNA-phosphodiesterases TDP1 and TDP2 function in the base excision repair (BER) and nonhomologous end joining (NHEJ) DNA repair pathways, respectively, and function in part in the repair of stalled topoisomerase-DNA complexes (3). Research has shown that inhibitors of tyrosyl-DNA-phosphodiesterases may act synergistically with topoisomerase inhibitors, allowing the potential for a more robust treatment of cancer (4,5). In small cell lung cancer, research suggests that TDP1 and topoisomerase 1 levels can predict sensitivity to topoisomerase 1 inhibitors (6).				
Background Re	eferences	1. Pommier, Y. (2013) <i>ACS Chem Biol</i> 8, 82-95. 2. Saijo, N. (2000) <i>Ann N Y Acad Sci</i> 922, 92-9. 3. Pommier, Y. et al. (2014) <i>DNA Repair (Amst)</i> 19, 114-29. 4. Nguyen, T.X. et al. (2012) <i>J Med Chem</i> 55, 4457-78. 5. Huang, S.N. et al. (2011) <i>Expert Opin Ther Pat</i> 21, 1285-92. 6. Meisenberg, C. et al. (2014) <i>J Cancer Sci Ther</i> 6, 258-67.				

**Species Reactivity** Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X

TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key H: Human M: Mouse R: Rat

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