## Topoisomerase IIα (D10G9) XP<sup>®</sup> Rabbit mAb (PE Conjugate)



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

<b>Applications:</b> FC-FP	Reactivity: H Mk	<b>Sensitivity:</b> Endogenous	Source/Isotype: Rabbit	UniProt ID: #P11388	Entrez-Gene Id: 7153
Product Usage Information		<b>Application</b> Flow Cytometry (Fixed/Pe	ermeabilized)		<b>Dilution</b> 1:50
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the antibody. Protect from light. Do not freeze.			
Specificity/Sensitivity		Topoisomerase II $\alpha$ (D10G9) XP $^{\otimes}$ Rabbit mAb (PE Conjugate) recognizes endogenous levels of total topoisomerase II $\alpha$ protein.			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human topoisomerase ${\rm II}\alpha$ protein.			
Description		This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometry analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated Topoisomerase IIα (D10G9) XP <sup>®</sup> Rabbit mAb #12286.			
Background		DNA topoisomerases I and II are nuclear enzymes; type II consists of two highly homologous isoforms: topoisomerase II $\alpha$ and II $\beta$ . These enzymes regulate the topology of DNA, maintain genomic integrity, and are essential for processes such as DNA replication, recombination, transcription, and chromosome segregation by allowing DNA strands to pass through each other (1). Topoisomerase I nicks and rejoins one strand of the duplex DNA, while topoisomerase II transiently breaks and closes double-stranded DNA (2). Topoisomerases are very susceptible to various stresses. Acidic pH or oxidative stress can convert topoisomerases to DNA-breaking nucleases, causing genomic instability and cell death. DNA-damaging topoisomerase targeting drugs (e.g., etoposide) also convert topoisomerases to nucleases, with the enzyme usually trapped as an intermediate that is covalently bound to the 5+ end of the cleaved DNA strand(s). Research studies have shown that this intermediate leads to genomic instability and cell death. Thus, agents that target topoisomerases are highly sought after cancer chemotherapeutic drugs (3). Ca <sup>2+</sup> -regulated phosphorylation of topoisomerase II $\alpha$ at Ser1106 modulates the activity of this enzyme and its sensitivity to targeting drugs (4).			
Background References		1. Wang, J.C. (2002) <i>Nat. Rev. Mol. Cell. Biol.</i> 3, 430-40. 2. Pulleyblank, .E. (1997) <i>Science</i> 277, 648-9. 3. Li, T.K. and Liu, L.F. (2001) <i>Annu. Rev. Pharmacol. Toxicol.</i> 41, 53-77. 4. Chikamori, K. et al. (2003) <i>J. Biol. Chem.</i> 278, 12696-702.			

**Species Reactivity** 

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

**Applications Key** FC-FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key H: Human Mk: Monkey

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