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## Cleaved PARP (Asp214) (D64E10) XP<sup>®</sup> Rabbit mAb (Alexa Fluor<sup>®</sup> 488 Conjugate)



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

Applications: IF-IC, FC-FP	Reactivity: H Mk	Sensitivity: Endogenous	Source/Isotype: Rabbit IgG	UniProt ID: #P09874	Entrez-Gene Id: 142	
Product Usage Information	In	<b>pplication</b> nmunofluorescence ( low Cytometry (Fixed	(Immunocytochemistry) I/Permeabilized)		<b>Dilution</b> 1:50 1:50	
Storage		pplied in PBS (pH 7. tibody. Protect from I	. Do not aliquot the			
Specificity / Sensit	lev	Cleaved PARP (Asp214) (D64E10) XP <sup>®</sup> Rabbit mAb (Alexa Fluor <sup>®</sup> 488 Conjugate) detects endogenous levels of the large fragment (89 kDa) of human PARP1 protein produced by caspase cleavage. The antibody does not recognize full length PARP1 or other PARP isoforms.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp214 of human PARP protein.				
Product Descriptic	ho	This Cell Signaling Technology antibody is conjugated to Alexa Fluor <sup>®</sup> 488 fluorescent dye and tested in- house for direct flow cytometry and immunofluorescent analysis in human cells. The antibody is expected to exhibit the same species cross-reactivity as the unconjugated Cleaved PARP (Asp214) (D64E10) XP <sup>®</sup> Rabbit mAb #5625.				
Background	to on As cal	PARP, a 116 kDa nuclear poly (ADP-ribose) polymerase, appears to be involved in DNA repair in response to environmental stress (1). This protein can be cleaved by many ICE-like caspases <i>in vitro</i> (2,3) and is one of the main cleavage targets of caspase-3 <i>in vivo</i> (4,5). In human PARP, the cleavage occurs between Asp214 and Gly215, which separates the PARP amino-terminal DNA-binding domain (24 kDa) from the carboxy-terminal catalytic domain (89 kDa) (2,4). PARP helps cells to maintain their viability; cleavage of PARP facilitates cellular disassembly and serves as a marker of cells undergoing apoptosis (6).				
Background Refer	2.   3. ( 4.   5. <sup>-</sup>	Lazebnik, Y. A. et al. Cohen, G.M. (1997) Nicholson, D. W. et a Tewari, M. et al. (199	<ul> <li>Iahl, T. (1992) Nature 356, 35 (1994) Nature 371, 346-347.</li> <li>Biochem. J. 326, 1-16.</li> <li>al. (1995) Nature 376, 37-43.</li> <li>95) Cell 81, 801-809.</li> <li>98) J. Biol. Chem. 273, 33533</li> </ul>			
Species Reactivity	spe	ecies reactivity is det	ermined by testing in at least (	one approved application (e.g., we	stern blot).	
Applications Key	IF-	IC: Immunofluoresce	ence (Immunocytochemistry)	FC-FP: Flow Cytometry (Fixed/Per	meabilized)	
Cross-Reactivity K	X: >	<ul> <li>H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster</li> <li>X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse</li> <li>GP: Guinea Pig Rab: rabbit All: all species expected</li> </ul>				
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