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**ASC/TMS1 (D2W8U) Rabbit mAb (Alexa Fluor® 647 Conjugate)**

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

Applications:	Reactivity:	Sensitivity:	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
IF-F, IF-IC	M	Endogenous	Rabbit IgG	#Q9EPB4	66824
<b>Product Usage Information</b>	<b>Application</b>			<b>Dilution</b>	
	Immunofluorescence (Frozen)			1:50	
	Immunofluorescence (Immunocytochemistry)			1:50	
<b>Storage</b>	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.				
<b>Specificity/Sensitivity</b>	ASC/TMS1 (D2W8U) Rabbit mAb (Alexa Fluor® 647 Conjugate) recognizes endogenous levels of total ASC protein.				
<b>Source / Purification</b>	Monoclonal antibody is produced by immunizing animals with recombinant mouse ASC protein.				
<b>Description</b>	This Cell Signaling Technology antibody is conjugated to Alexa Fluor® 647 fluorescent dye and tested in-house for direct immunofluorescent analysis in mouse cells. The antibody is expected to exhibit the same species cross-reactivity as the unconjugated ASC/TMS1 (D2W8U) Rabbit mAb #67824.				
<b>Background</b>	TMS1 (target of methylation-induced silencing)/ASC (apoptosis-associated speck-like protein containing a CARD), also referred to as PYCARD and CARD5, is a 22 kDa pro-apoptotic protein containing an N-terminal pyrin domain (PYD) and a C-terminal caspase recruitment domain (CARD) (1-2). The <i>ASC/TMS1</i> gene was originally found to be aberrantly methylated and silenced in breast cancer cells (2), and has since been found to be silenced in a number of other cancers, including ovarian cancer (3), glioblastoma (4), melanoma (5), gastric cancer (6), lung cancer (7), and prostate cancer (8). Expression of ASC/TMS1 can be induced by pro-apoptotic/inflammatory stimuli (9). During apoptosis ASC/TMS1 is re-distributed from the cytosol to the mitochondria and associates with mitochondrial Bax to trigger cytochrome c release and subsequent apoptosis (10). ASC/TMS1 has also been found to be a critical component of inflammatory signaling where it associates with and activates caspase-1 in response to pro-inflammatory signals (11).				
<b>Background References</b>	<ol style="list-style-type: none"> <li>Masumoto, J. et al. (1999) <i>J Biol Chem</i> 274, 33835-8.</li> <li>Conway, K.E. et al. (2000) <i>Cancer Res</i> 60, 6236-42.</li> <li>Terasawa, K. et al. (2004) <i>Clin Cancer Res</i> 10, 2000-6.</li> <li>Stone, A.R. et al. (2004) <i>Am J Pathol</i> 165, 1151-61.</li> <li>Guan, X. et al. (2003) <i>Int J Cancer</i> 107, 202-8.</li> <li>Moriai, R. et al. (2002) <i>Anticancer Res</i> 22, 4163-8.</li> <li>Virmani, A. et al. (2003) <i>Int J Cancer</i> 106, 198-204.</li> <li>Das, P.M. et al. (2006) <i>Mol Cancer</i> 5, 28.</li> <li>Strong, R. et al. (1991) <i>Brain Res</i> 542, 23-8.</li> <li>Ohtsuka, T. et al. (2004) <i>Nat Cell Biol</i> 6, 121-8.</li> <li>Srinivasula, S.M. et al. (2002) <i>J Biol Chem</i> 277, 21119-22.</li> </ol>				

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

**Applications Key** **IF-F:** Immunofluorescence (Frozen) **IF-IC:** Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key** **M:** Mouse

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