## ASC/TMS1 (D2W8U) Rabbit mAb (Alexa Fluor® 488 Conjugate)



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

Applications: Reactivity IF-IC, FC-FP M	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q9EPB4	Entrez-Gene Id: 66824	
Product Usage Information	Application Immunofluorescence (Immunocytochemistry) Flow Cytometry (Fixed/Permeabilized)			<b>Dilution</b> 1:50 - 1:100 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	ASC/TMS1 (D2W8U) Rab ASC/TMS1 protein.	ASC/TMS1 (D2W8U) Rabbit mAb (Alexa Fluor $^{\$}$ 488 Conjugate) recognizes endogenous levels of total ASC/TMS1 protein.			
Source / Purification	Monoclonal antibody is	Monoclonal antibody is produced by immunizing animals with recombinant mouse ASC/TMS1 protein.			
Description	This Cell Signaling Technology antibody is conjugated to Alexa Fluor <sup>®</sup> 488 fluorescent dye and tested in-house for direct immunofluorescence analysis in mouse cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated ASC (D2W8U) Rabbit mAb #67824.				
Background	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 ASC/TMS1 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).				
Background References	1. Masumoto, J. et al. (1999) <i>J Biol Chem</i> 274, 33835-8.  2. Conway, K.E. et al. (2000) <i>Cancer Res</i> 60, 6236-42.  3. Terasawa, K. et al. (2004) <i>Clin Cancer Res</i> 10, 2000-6.  4. Stone, A.R. et al. (2003) <i>Int J Cancer</i> 107, 202-8.  6. Moriai, R. et al. (2002) <i>Anticancer Res</i> 22, 4163-8.  7. Virmani, A. et al. (2003) <i>Int J Cancer</i> 106, 198-204.  8. Das, P.M. et al. (2006) <i>Mol Cancer</i> 5, 28.  9. Strong, R. et al. (1991) <i>Brain Res</i> 542, 23-8.  10. Ohtsuka, T. et al. (2004) <i>Nat Cell Biol</i> 6, 121-8.  11. Srinivasula, S.M. et al. (2002) <i>J Biol Chem</i> 277, 21119-22.				

**Species Reactivity** 

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

**Applications Key** 

IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)

**Cross-Reactivity Key** 

M: Mouse

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