## SMG-1 (V72) Antibody



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

<b>Applications:</b> W, IP, IF-IC	Reactivity: H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 410	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #Q96Q15	Entrez-Gene Id: 23049
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation Immunofluorescence		nistry)		<b>Dilution</b> 1:1000 1:50 1:50
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		SMG-1 (V72) Antibody detects endogenous levels of total SMG-1 protein. The antibody cross-reacts with a 100 kDa band of unknown origin. By confocal immunofluorescence, the antibody labels the mitochondria of U-2 OS cells (7).				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human SMG-1.				
Background		SMG-1 is a member of the phosphoinositide 3-kinase-related kinase (PIKK) family, which includes ATM, ATR, mTOR, DNA-PKcs, and TRRAP (1,2). Activated by DNA damage, SMG-1 has been shown to phosphorylate p53 and hUpf1 (SMG-2) (1-4). hUpf1 is a subunit of the surveillance complex that allows degradation of messenger RNA species containing premature termination codons (PTCs). This process, known as nonsense-mediated mRNA decay (NMD), prevents the translation of truncated forms of proteins that may result in gain of function or dominant negative species. NMD occurs under normal cellular conditions as well as in response to damage (5,6). SMG-1 has also been shown to affect cell death receptor signaling and to protect cells from extrinsically induced apoptotic cell death (7).				
Background References		1. Denning, G. et al. (2001) <i>J Biol Chem</i> 276, 22709-14. 2. Yamashita, A. et al. (2001) <i>Genes Dev</i> 15, 2215-28. 3. Brumbaugh, K.M. et al. (2004) <i>Mol Cell</i> 14, 585-98. 4. Ohnishi, T. et al. (2003) <i>Mol Cell</i> 12, 1187-200. 5. Li, Z.Y. et al. (2006) <i>Curr Med Chem</i> 13, 1693-705. 6. Mendell, J.T. et al. (2004) <i>Nat Genet</i> 36, 1073-8. 7. Oliveira, V. et al. (2008) <i>J Biol Chem</i> 283, 13174-84.				
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**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 IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key** H: Human Mk: Monkey

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