

## £11832

## USP8 (D18F6) Rabbit mAb



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

<b>Applications:</b> W, IP	Reactivity: H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 140	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P40818	Entrez-Gene Id: 9101
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:100	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		USP8 (D18F6) Rabbit mAb recognizes endogenous levels of total USP8 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Leu614 of human USP8 protein.				
Background		Ubiquitinating enzymes (UBEs) catalyze protein ubiquitination, a reversible process countered by deubiquitinating enzyme (DUB) action (1,2). Five DUB subfamilies are recognized, including the USP, UCH, OTU, MJD, and JAMM enzymes. The deubiquitinating enzyme ubiquitin-specific protease 8 (USP8/UBPy) is a cysteine protease belonging to the USP/UBP subfamily. Research studies have shown that USP8 is an essential growth-regulated enzyme indespensible for cell proliferation and survival (3,4). Indeed, conditional knock-out of murine USP8 was shown to promote a dramatic loss in expression of receptor tyrosine kinases, including EGFR, ErbB3, and c-Met (4). In agreement with these findings, USP8 inactivation leads to enhanced ubiquitination of ligand-activated EGFR (5,6). Furthermore, phosphorylation of USP8 at Ser680 results in its binding of 14-3-3, catalytic inactivation, and reduced EGFR deubiquitination (7). It appears as though USP8, in conjunction with components of the ESCRT-0 complex, plays an integral role in the early endosomal sorting machinery that functions to protect EGFR from lysosomal degradation (8,9).				
Background References		<ol> <li>Nijman, S.M. et al. (2005) <i>Cell</i> 123, 773-86.</li> <li>Nalepa, G. et al. (2006) <i>Nat Rev Drug Discov</i> 5, 596-613.</li> <li>Naviglio, S. et al. (1998) <i>EMBO J</i> 17, 3241-50.</li> <li>Niendorf, S. et al. (2007) <i>Mol Cell Biol</i> 27, 5029-39.</li> <li>Alwan, H.A. and van Leeuwen, J.E. (2007) <i>J Biol Chem</i> 282, 1658-69.</li> <li>Mizuno, E. et al. (2005) <i>Mol Biol Cell</i> 16, 5163-74.</li> <li>Mizuno, E. et al. (2007) <i>Exp Cell Res</i> 313, 3624-34.</li> <li>Row, P.E. et al. (2006) <i>J Biol Chem</i> 281, 12618-24.</li> <li>Berlin, I. et al. (2010) <i>J Biol Chem</i> 285, 34909-21.</li> </ol>				
Species Reactivi	ity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).
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**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

**Cross-Reactivity Key** 

H: Human Mk: Monkey

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