

94402

Cas9 *(S. pyogenes)* (E7M1H) XP[®] Rabbit mAb (Biotinylated)



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Applications: W	Reactivity: All	Sensitivity: Transfected Only	MW (kDa): 150	Source/Isotype: Rabbit IgG	UniProt ID: #Q99ZW2	Entrez-Gene Id 901176
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 136 mM NaCl, 2.6 mM KCI, 12 mM sodium phosphate (pH 7.4) dibasic, 2 mg/ml BSA, and 50% glycerol. Store at –20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		Cas9 <i>(S. pyogenes)</i> (E7M1H) XP [®] Rabbit mAb (Biotinylated) recognizes transfected levels of total Cas9 (<i>S. pyogenes</i>) protein. This antibody does not cross-react with Cas9 (<i>S. aureus</i>), FnCpf1, and AsCpf1 proteins.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Leu833 of Cas9 (<i>S. pyogenes</i>) protein.				
Description		This Cell Signaling Technology antibody is conjugated to biotin under optimal conditions. The biotinylated antibody is expected to exhibit the same species cross-reactivity as the unconjugated Cas9 <i>(S. pyogenes)</i> (E7M1H) XP [®] Rabbit mAb #19526.				
Background		The CRISPR associated protein 9 (Cas9) is an RNA-guided DNA nuclease and part of the <i>Streptococcus pyogenes</i> CRISPR antiviral immunity system that provides adaptive immunity against extrachromosomal genetic material (1). The CRISPR antiviral mechanism of action involves three steps: (i), acquisition of foreign DNA by host bacterium; (ii), synthesis and maturation of CRISPR RNA (crRNA) followed by the formation of RNA-Cas nuclease protein complexes; and (iii), target interference through recognition of foreign DNA by the complex and its cleavage by Cas nuclease activity (2). The type II CRISPR/Cas antiviral immunity system provides a powerful tool for precise genome editing and has potential for specific gene regulation and therapeutic applications (3). The Cas9 protein and a guide RNA consisting of a fusion between a crRNA and a trans-activating crRNA (tracrRNA) must be introduced or expressed in a cell. A 20-nucleotide sequence at the 5' end of the guide RNA directs Cas9 to a specific DNA target site. As a result, Cas9 can be "programmed" to cut various DNA sites both <i>in vitro</i> and in cells and organisms. CRISPR/Cas9 genome editing tools have been used in many organisms, including mouse and human cells (4,5). Research studies demonstrate that CRISPR can be used to generate mutant alleles or reporter genes in rodents and primate embryonic stem cells (6-8).				
Background References		1. Horvath, P. and Barrangou, R. (2010) <i>Science</i> 327, 167-70. 2. Wiedenheft, B. et al. (2012) <i>Nature</i> 482, 331-8. 3. Singh, P. et al. (2015) <i>Genetics</i> 199, 1-15. 4. Cong, L. et al. (2013) <i>Science</i> 339, 819-23. 5. Mali, P. et al. (2013) <i>Science</i> 339, 823-6. 6. Li, D. et al. (2013) <i>Nat Biotechnol</i> 31, 681-3. 7. Shen, B. et al. (2013) <i>Cell Res</i> 23, 720-3. 8. Niu, Y. et al. (2014) <i>Cell</i> 156, 836-43.				
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 nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting				
Cross-Reactivity Key		All: All Species Expected				

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