Cas9 (7A9-3A3) Mouse mAb (PE Conjugate)		ell Signaling сн N о L о g Y®
	Orders:	877-616-CELL (2355) orders@cellsignal.com
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#35193	Web:	info@cellsignal.com cellsignal.com
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For Research Use Only. Not for Use in Diagnostic Procedures.		

Applications: Rea FC-FP	ctivity: Sensitivity: Source/Isotype: All Transfected Mouse IgG1 Only	UniProt ID:Entrez-Gene Id:#Q99ZW2901176
Product Usage Information	Application Flow Cytometry (Fixed/Permeabilized)	Dilution 1:50
Storage	Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 antibodies. Protect from light. Do not freeze.	mg/ml BSA. Store at 4°C. Do not aliquot the
Specificity / Sensitivity	Cas9 (7A9-3A3) Mouse mAb (PE Conjugate) recognizes trans	sfected levels of total Cas9 protein.
Source / Purification	Monoclonal antibody is produced by immunizing animals with terminus of Cas9 from <i>Streptococcus pyogene</i> .	recombinant protein specific to the amino
Product Description	This Cell Signaling Technology antibody is conjugated to phyc flow cytometry analysis in human cells. This antibody is expec reactivity as the unconjugated Cas9 (7A9-3A3) Mouse mAb #	cted to exhibit the same species cross-
Background	The CRISPR associated protein 9 (Cas9) is an RNA-guided D pyogenes CRISPR antiviral immunity system that provides ad genetic material (1). The CRISPR antiviral mechanism of action foreign DNA by host bacterium; (ii), synthesis and maturation formation of RNA-Cas nuclease protein complexes; and (iii), to foreign DNA by the complex and its cleavage by Cas nuclease antiviral immunity system provides a powerful tool for precise gene regulation and therapeutic applications (3). The Cas9 pr between a crRNA and a trans-activating crRNA (tracrRNA) mu nucleotide sequence at the 5' end of the guide RNA directs Ca Cas9 can be "programmed" to cut various DNA sites both <i>in v</i> CRISPR/Cas9 genome editing tools have been used in many (4,5). Research studies demonstrate that CRISPR can be use genes in rodents and primate embryonic stem cells (6-8).	laptive immunity against extrachromosomal on involves three steps: (i), acquisition of of CRISPR RNA (crRNA) followed by the arget interference through recognition of e activity (2). The type II CRISPR/Cas genome editing and has potential for specific otein and a guide RNA consisting of a fusion ust be introduced or expressed in a cell. A 20- as9 to a specific DNA target site. As a result, <i>ritro</i> and in cells and organisms. organisms, including mouse and human cells
Background Reference	 S 1. Horvath, P. and Barrangou, R. (2010) Science 327, 167-70. 2. Wiedenheft, B. et al. (2012) Nature 482, 331-8. 3. Singh, P. et al. (2015) Genetics 199, 1-15. 4. Cong, L. et al. (2013) Science 339, 819-23. 5. Mali, P. et al. (2013) Science 339, 823-6. 6. Li, D. et al. (2013) Nat Biotechnol 31, 681-3. 7. Shen, B. et al. (2013) Cell Res 23, 720-3. 8. Niu, Y. et al. (2014) Cell 156, 836-43. 	
Species Reactivity	Species reactivity is determined by testing in at least one appro	oved application (e.g., western blot).
Applications Key	FC-FP: Flow Cytometry (Fixed/Permeabilized)	
Cross-Reactivity Key	 H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: viru X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cere GP: Guinea Pig Rab: rabbit All: all species expected 	0

1 Cas9 (7A9-3A3) Mouse mAb (PE Conjugate) (#35193) Datasheet Without Images Cell Signaling Technology

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