

AsCpf1/Cas12a (Strain *BV3L6*) Antibody

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

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:
W	All	Transfected Only	151	Rabbit	#U2UMQ6

Product Usage Information	Application Western Blotting	Dilution 1:1000
Storage	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at -20°C. Do not aliquot the antibody.	
Specificity/Sensitivity	AsCpf1/Cas12a (Strain <i>BV3L6</i>) Antibody recognizes transfected levels of total AsCpf1/Cas12a (Strain <i>BV3L6</i>) protein. This antibody does not cross-react with Cas9 (<i>S. pyogenes</i>), Cas9 (<i>S. aureus</i>), and FnCpf1/Cas12a (Strain <i>U112</i>) proteins.	
Source / Purification	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ile57 of <i>Acidaminococcus sp.</i> Cpf1/Cas12a (Strain <i>BV3L6</i>) protein. Antibodies are purified by protein A and peptide affinity chromatography.	
Background	CRISPR-Cas (clustered regularly interspaced short palindromic repeats and CRISPR-associated proteins) are RNA-guided nuclease effectors that are utilized for precise genome editing in mammalian systems (1). Cpf1/Cas12a (CRISPR from <i>Prevotella</i> and <i>Francisella</i>) proteins are members of the Class 2 CRISPR system (2). Class 2 CRISPR systems, such as the well characterized Cas9, rely on single-component effector proteins to mediate DNA interference (3). Cpf1/Cas12a endonucleases, compared to Cas9 systems, have several unique features that increase the utility of CRISPR-based genome editing techniques: 1) Cpf1/Cas12a-mediated cleavage relies on a single and short CRISPR RNA (crRNA) without the requirement of a trans-activating crRNA (tracrRNA), 2) Cpf1/Cas12a utilizes T-Rich protospacer-adjacent motif (PAM) sequences rather than a G-Rich PAM, and 3) Cpf1/Cas12a generates a staggered, rather than a blunt-ended, DNA double-stranded break (2). These features broaden the utility of using CRISPR-Cas systems for specific gene regulation and therapeutic applications. Several Cpf1/Cas12a bacterial orthologs have been characterized for CRISPR-mediated mammalian genome editing (2,4). AsCpf1 (Strain <i>BV3L6</i>)/Cas12a is a Cpf1/Cas12a enzyme derived from <i>Acidaminococcus sp. BV3L6</i> (5,6).	
Background References	<ol style="list-style-type: none"> Cong, L. et al. (2013) <i>Science</i> 339, 819-23. Zetsche, B. et al. (2015) <i>Cell</i> 163, 759-71. Horvath, P. and Barrangou, R. (2010) <i>Science</i> 327, 167-70. Zhang, Y. et al. (2017) <i>Sci Adv</i> 3, e1602814. Zetsche, B. et al. (2015) <i>Cell</i> 163, 759-71. Zhang, Y. et al. (2017) <i>Sci Adv</i> 3, e1602814. 	
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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