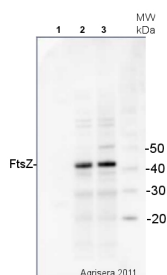


Product no **AS10 715****Anti-FtsZ | Prokaryotic cell division GTPase****Product information**

<b>Immunogen</b>	KLH-conjugated synthetic peptide derived from known bacterial sequences of FtsZ including <i>E.coli</i> UniProt: <a href="#">P0A9A6</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Serum
<b>Format</b>	Lyophilized
<b>Quantity</b>	50 µl
<b>Reconstitution</b>	For reconstitution add 50 µl of sterile water
<b>Storage</b>	Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tube.

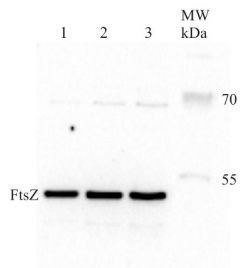
**Application information**

<b>Recommended dilution</b>	1 : 100-1 : 200 (IF), 1 : 1000 (WB)
<b>Expected   apparent MW</b>	40   42 kDa
<b>Confirmed reactivity</b>	<i>Candidatus Moranella endobia</i> PCIT, <i>Escherichia</i> DH5a, <i>Escherichia coli</i> BW 25113, <i>Shigella flexneri</i>
<b>Predicted reactivity</b>	<i>Candidatus sp.</i> , <i>Citrobacter</i> sp. 30_2, <i>Dickeya</i> sp., <e> Species of your interest not listed? <a href="#">Contact us</a> > <i>Enterobacter</i> sp., <i>Klebsiella pneumoniae</i> subsp. <i>pneumoniae</i> MGH, <i>Salmonella</i> sp., <i>Shigella sonnei</i> Ss046, <i>Vibrio</i> sp., <i>Yersinia pestis</i> D182038, <i>Xanthomonas oryzae</i>
<b>Not reactive in</b>	<i>Allysiella filiformis</i> , <i>B. subtilis</i> , <i>Haloflex mediterranei</i> ( <i>Listeria</i> sp., <i>Neisseria meningitidis</i> , <i>Pseudomonas aeruginosa</i> , <i>Staphylococcus aureus</i> (strain MRSA252), cyanobacteria
<b>Selected references</b>	<a href="#">Chakraborty</a> et al. (2024). Dynamics of interdomain rotation facilitates FtsZ filament assembly. JBC, 7 May 2024, 107336. <a href="#">Vedyaykin</a> et al. (2020). SulA is able to block cell division in <i>Escherichia coli</i> by a mechanism different from sequestration. Biochem Biophys Res Commun. DOI: 10.1016/j.bbrc.2020.03.012 <a href="#">Ranjit</a> et al. (2020). Chlamydial MreB Directs Cell Division and Peptidoglycan Synthesis in <i>Escherichia coli</i> in the Absence of FtsZ Activity. mBio. 2020 Feb 18;11(1). pii: e03222-19. doi: 10.1128/mBio.03222-19. (Immunofluorescence) <a href="#">Sekar</a> et al. (2018). Synthesis and degradation of FtsZ quantitatively predict the first cell division in starved bacteria. Mol Syst Biol. 2018 Nov 5;14(11):e8623. doi: 10.15252/msb.20188623. <a href="#">Mückl</a> et al. (2018). Filamentation and restoration of normal growth in <i>Escherichia coli</i> using a combined CRISPRi sgRNA/antisense RNA approach. PLoS One. 2018 Sep 11;13(9):e0198058. doi: 10.1371/journal.pone.0198058. eCollection 2018. <a href="#">Pende</a> et al. (2014). Size-independent symmetric division in extraordinarily long cells. Nat Commun. 2014 Sep 15;5:4803. doi: 10.1038/ncomms5803. <a href="#">Söderström</a> et al. (2014). Disassembly of the divisome in <i>Escherichia coli</i> : Evidence that FtsZ dissociates before compartmentalisation. Mol Microbiol. 2014 Feb 7. doi: 10.1111/mmi.12534. (western blot and immunofluorescence)

**Application example**

**5 µg of total protein** from *Synechocystis* sp. (1), *E.coli* DH5a (2), *E. coli* (3), extracted with Agrisera [PEB extraction buffer](#) were separated on **4-12% SDS-PAGE** and blotted 1h to **PVDF**. Blots were blocked with Advance blocking reagent for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 10 000 for 1h at RT with agitation. The antibody solution was decanted and the blot

was rinsed briefly twice, then washed once for 15 min and 3 times for 5 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, from Agrisera, [AS09\\_602](#)) diluted to 1:25 000 in for 1 h at RT with agitation. The blot was washed as above and developed for 5 min with ECL according to the manufacturers instructions. Exposure time was 120 seconds.



Total protein extract from *E. coli* CFT073: 8µg (1), 12µg (2), 16µg (3). Proteins were separated on 10% SDS-PAGE and blotted to PVDF. Blocked with 5 % non-fat milk in TBS-T for 1 hour. Blot was incubated in the primary antibody at a dilution of 1 :10 000 for 1 h at RT with agitation. Secondary antibody (anti-rabbit IgG, HRP conjugated, Agrisera, [AS09\\_602](#)) were diluted to 1 : 50 000 and blot was incubated for 1h at RT with agitation. Immunodetection was performed using chemiluminescent detection method for 3 min. Scan was made after 30 sec.

Courtesy of Dr. Marta Kicia, Wroclaw Medical University, Poland