

Product no **AS07 217****Anti-FtsZ | Procaryotic cell division GTPase (cyanobacterial)****Product information****Immunogen** | Whole cyanobacterial (*Anabaena* PCC 7120) FtsZ protein, UniProt: [Q3MC27](#) overexpressed in *E.coli*.**Host** | Rabbit**Clonality** | Polyclonal**Purity** | Serum**Format** | Lyophilized**Quantity** | 50 µl**Reconstitution** | For reconstitution add 50 µl of sterile water**Storage** | 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.**Additional information** | To detect *E.coli* FtsZ protein we recommend a following product: [AS10 715](#) | anti-FtsZ procaryotic cell division GTPase (bacterial), rabbit antibody

**To detect FtsZ protein in higher plants** following antibodies are recommended:  
[AS09 413](#) | Anti-FtsZ1 and 2 | Plant cell division protein FtsZ1 and FtsZ2, rabbit antibodies  
[AS13 2651](#) | Anti-FtsZ2 | Plant cell division protein ftsZ2, rabbit antibodies

**Application information****Recommended dilution** | 1 : 200 (IL) Immunogold-TEM, 1 : 500 (IF), 1 : 2000-1 : 5000 (WB)**Expected | apparent MW** | 44.5 | 50 kDa**Confirmed reactivity** | *Cylindrospermopsis raciborskii* CS-505, *Listeria monocytogenes* (weak reaction), *Synechococcus elongatus***Predicted reactivity** | *Phaeodactylum tricornutum*, *Prochlorococcus* sp.**Not reactive in** | higher plants**Additional information** | This antibody can be used as a loading control antibody in cyanobacteria. Immunofluorescence has been done by labelling *Synechococcus elongatus* cells at 30°C for 2 hours with FtsZ antibodies diluted to 1: 500 in blocking buffer. Detection images can be found in [Kabeya](#) et al (2010).

This product can be sold containing ProClin if requested.

**Selected references**

[Kurmayer](#) et al. (2020). Chemically labeled toxins or bioactive peptides show a heterogeneous intracellular distribution and low spatial overlap with autofluorescence in bloom-forming cyanobacteria. *Sci Rep.* 2020 Feb 17;10(1):2781. doi: 10.1038/s41598-020-59381-w. (Immunofluorescence)

[Zhan](#) et al. (2018). Photobleaching Enables Super-resolution Imaging of the FtsZ Ring in the Cyanobacterium *Prochlorococcus*. *J Vis Exp.* 2018 Nov 6;(141). doi: 10.3791/58603.

[MacCready](#) et al. (2016). Robust Min-System Oscillation in the Presence of Internal Photosynthetic Membranes in Cyanobacteria. *Molecular Microbiology* November 5 2016. doi: 10.1111/mmi.13571

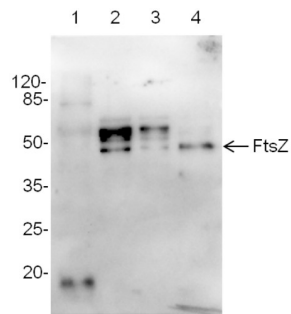
[Probst](#) et al. (2014). Biology of a widespread uncultivated archaeon that contributes to carbon fixation in the subsurface. *Nat Commun.* 2014 Nov 26;5:5497. doi: 10.1038/ncomms6497.

[Miyagishima](#) et al. (2014). DipM is required for peptidoglycan hydrolysis during chloroplast division. *BMC Plant Biol.* 2014 Mar 6;14(1):57. (immunofluorescence)

[Plominsky](#) et al. (2013). Dinitrogen Fixation Is Restricted to the Terminal Heterocysts in the Invasive Cyanobacterium *Cylindrospermopsis raciborskii* CS-505. *PLOS ONE, Open Access.*

[Kabeya](#) et al (2010). The YlmG protein has a conserved function related to the distribution of nucleoids in chloroplasts and cyanobacteria. *BMC Plant Biology* 10:57.

## Application example



Total protein samples (5 or 10 µg) from: *Arabidopsis thaliana*, leaf (**1**), *Synechocystis* 6803 motile (**2**), *Synechocystis* 6803 GT (glucose tolerant strain) (**3**), *Synechococcus elongates* 7942 (**4**), Marker - Pierce™ Prestained Protein MW Marker (kat #26612) were extracted with buffer (10mM Tris HCl, pH 8.0, 0.5% LDS, 4% glycerol, 0.1 mM EDTA ) were mixed with sample buffer and denatured for 5 min at 95 °C. Samples were separated on 10% SDS -PAGE and blotted 1h to nitrocellulose membrane (Amersham Protran) using sem i-dry transfer (Bio -Rad) in standard transfer buffer in presence of 10% methanol. Transfer of proteins to the membrane was checked using 0,5% Ponceau S staining before the blocking step. Blots were blocked in buffer (2% low -fat milk in 1xPBS, 0,1% Tween) for 1h at room temperature (RT) with agitation. Blots were incubated in the primary antibody at a dilution of 1: 1000 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 PBS -T at RT with agitation. Blot was incubated in secondary antibody ( goat anti -rabbit IgG, [AS09 602](#), Agrisera) diluted to 1:30 000 for 1h at RT with agitation. The blot was washed as above and developed for 5 min with chemiluminescence detection reagent and ChemiDoc detection system.

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