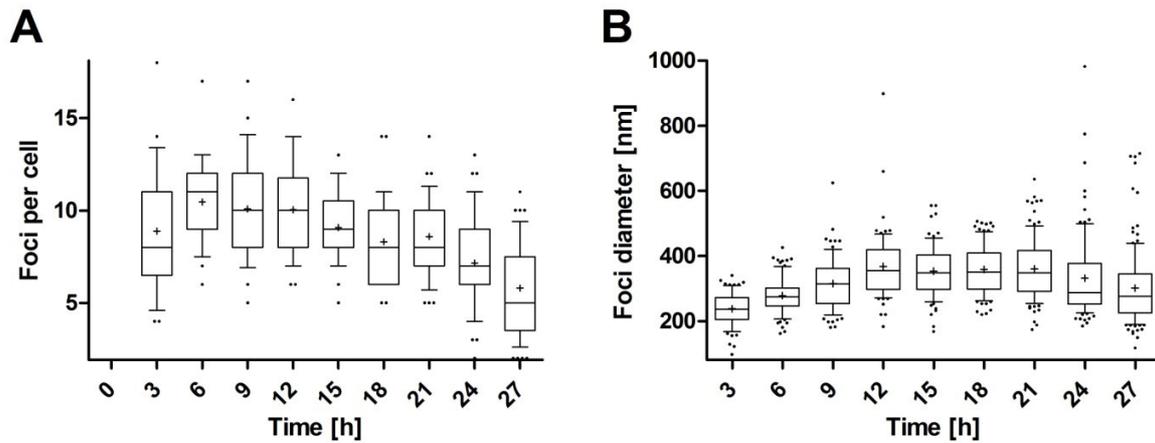
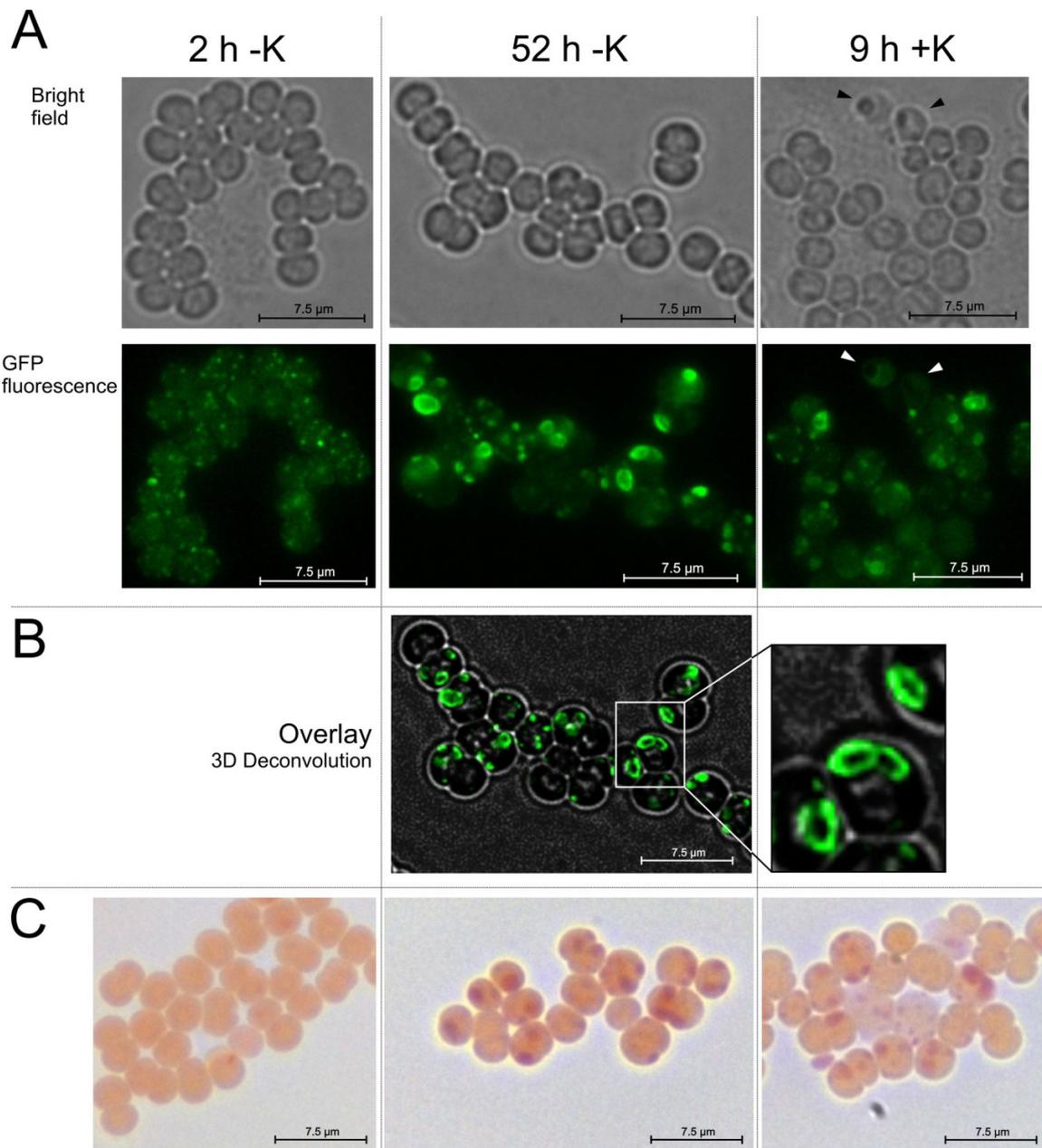


1  
2 Fig. S1. *Synechocystis* sp. cells after 4 days of nitrogen starvation and during resuscitation. Cells  
3 were resuscitated by adding 17.3 mM nitrate. Photos were taken 0, 18 and 24 h after the addition  
4 of nitrogen. Bright-field images and the associated GFP fluorescence images showing the  
5 localization of CphA-eGFP are shown. The Sakaguchi reaction was used to specifically stain the  
6 cyanophycin granules in cells at the same time points.



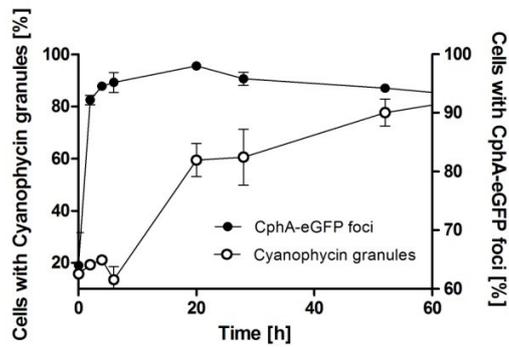
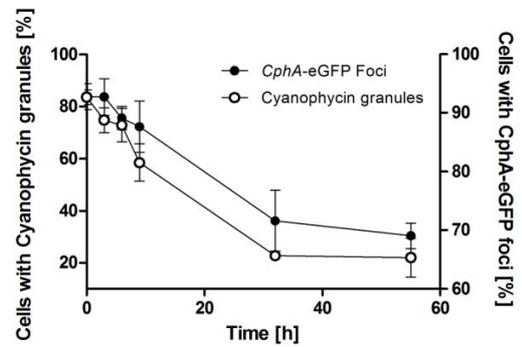
7  
 8 Fig. S2. Number of CphA-eGFP foci and foci diameter during 27 h of resuscitation of  
 9 *Synechocystis* sp. cultures nitrogen starved for 4 days. Cells were resuscitated by adding 17.3  
 10 mM nitrate to the starved cultures. (A) Number of CphA-eGFP foci per cell (n = 25–50 cells per  
 11 time point). (B) Diameter of foci in nm (n =  $\geq 70$  foci per time point). Values are the means of  
 12 three biological replicates. Whiskers range from 10% to 90% of the values. +, arithmetic mean;  
 13 dots, outliers.

14

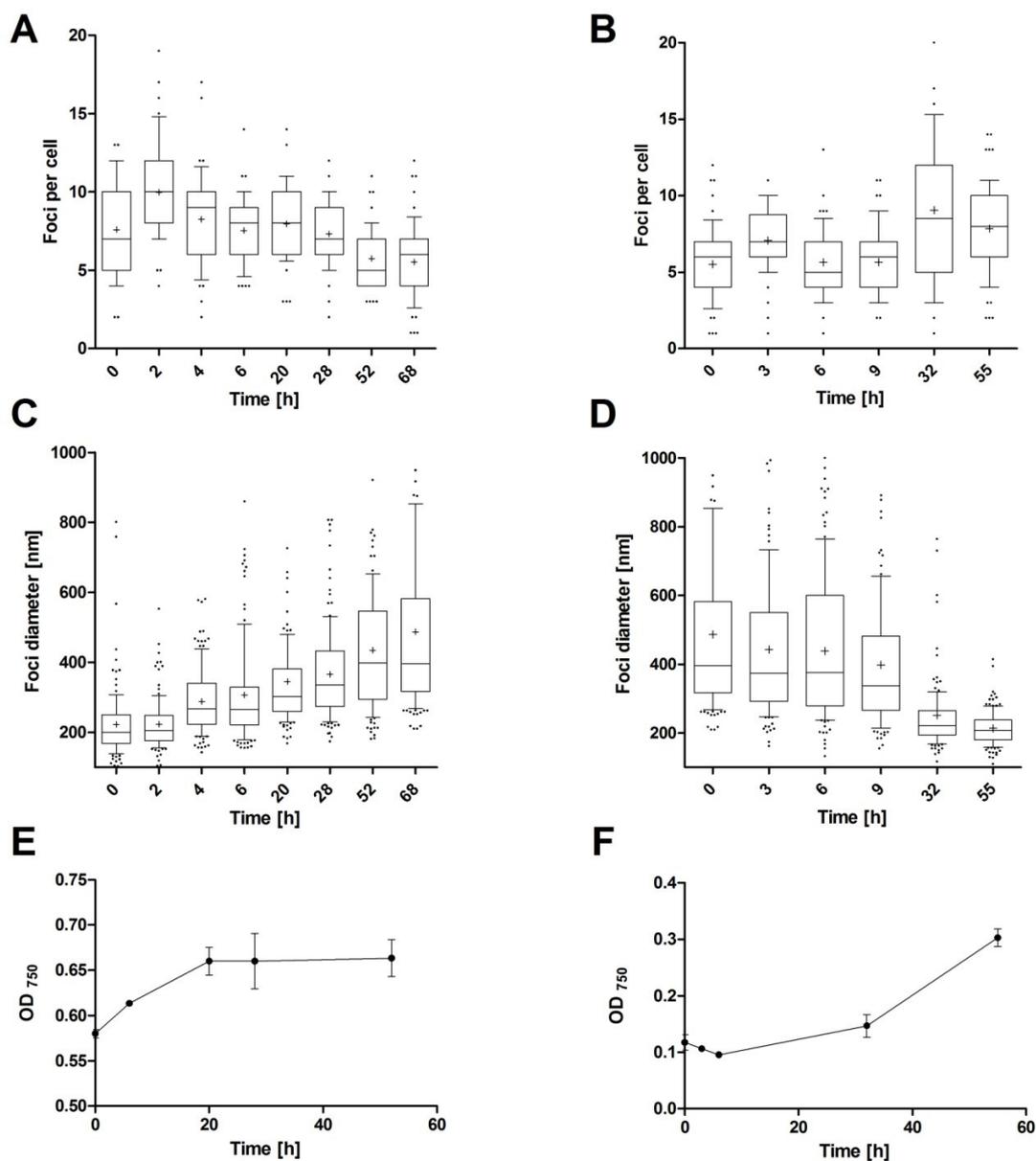


15  
 16 Fig. S3. Microscopy images of *Synechocystis* sp. during potassium starvation and regeneration.  
 17 Potassium starvation, which leads to growth arrest and cyanophycin accumulation, was induced  
 18 by transferring exponentially growing cells to BG-11 medium lacking potassium. To regenerate  
 19 starved cultures, cells were washed and resuspended in BG-11 medium containing potassium.  
 20 Images were taken 2 h and 52 h after induction of starvation and 9 h after potassium was  
 21 provided. (A) Bright-field images and GFP-fluorescence images showing CphA-eGFP

22 localization of the same cells. White and black arrows point to cells without CphA-eGFP  
23 localization on the cyanophycin granule surface. (B) 3-D deconvolution overlay of GFP  
24 fluorescence and bright-field images after 52 h of potassium depletion. The enlarged area shows  
25 the localization of CphA on the granule surface, which forms a halo-like structure. (C) Cells  
26 stained with the arginine-specific Sakaguchi stain at the same time points. Cyanophycin granules  
27 appear as dark red dots in the cell.

**A****B**

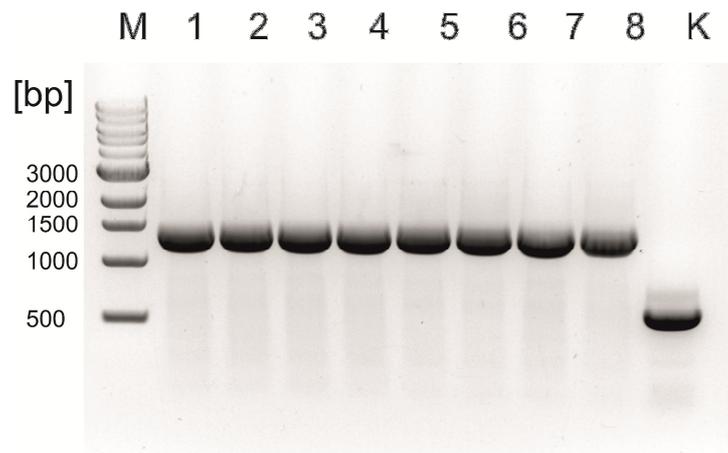
28 Fig. S4 (A) Percentage of cells with visible CphA-eGFP foci ( $n = \geq 120$  cells per time point; filled  
 29 circles) and cyanophycin granules ( $n = \geq 80$  cells per time point; open circles) during potassium  
 30 starvation and (B) during regeneration from potassium starvation. Potassium starvation was  
 31 induced by transferring exponentially growing cells to BG-11 medium lacking potassium. For  
 32 regeneration, the starved cells were washed and resuspended in BG-11 medium containing  
 33 potassium. Results shown are means of three biological replicates.  
 34



35

36 Fig. S5. Number of CphA-eGFP foci per cell and foci diameter during potassium starvation and  
 37 regeneration. Exponentially growing *Synechocystis* sp. cells were starved by washing and  
 38 resuspension in BG-11 medium lacking potassium. Regeneration was induced by washing  
 39 starved cells and resuspension in BG-11 medium containing potassium. Cyanophycin  
 40 accumulated during potassium depletion and degraded during regeneration. (A) Number of

41 CphA-eGFP foci per cell ( $n = \geq 40$  cells per time point) during potassium starvation and (B)  
42 during resuscitation from potassium starvation. (C) Diameter of foci in nm ( $n = \geq 110$  foci per  
43 time point) during potassium starvation and (D) during resuscitation from potassium starvation.  
44 Whiskers range from 10% to 90% of the values; +, arithmetic mean; black dots, outliers. (E)  
45 Growth curve of *Synechocystis* sp. during potassium starvation and (F) during resuscitation. All  
46 measurements are means of three biological replicates.



47

48 Fig. S6. Screening for the complete replacement of all genomic copies of *slr2002* (*cphA*) with the  
 49 kanamycin resistance cassette by PCR. Isolated genomic DNA of *Synechocystis* sp. wild-type (K)  
 50 and eight putative  $\Delta cphA$  clones (1–8) were used as template. In a three-primer approach  
 51 (primers *seg\_for*, *kan\_for*, and *down\_rev*), the wild-type *slr2002* yields a band of around 550 bp  
 52 (K) and the knockout construct yields a band of around 1300 bp. In clones 1–8, no wild-type  
 53 background band was observed. Clone 8 was used in the following studies.