

Supplementary Figure S6 -- Construction of the pC-LS plasmid for constitutive expression of the *Mentha spicata* 4S-limonene synthase encoding gene adapted to the *Synechocystis* PCC 6803 codon usage. The *NdeI* and *EcoRI* restriction sites are used for the cloning the *limonene synthase* gene from the pEX\_K4\_LS (provided by Eurofins Genomics) into the pC vector, generating the pC-LS plasmid. The *limonene synthase* gene (shown in yellow) is expressed constitutively from the strong  $\lambda$  phage pR promoter (shown in red). The genes are represented by colored arrows pointing into the direction of their transcription. Note that the  $Cm^R$  gene of pC-LS is truncated and that the  $Sp^R/Sm^R$  gene (pink) is flanked by the double terminator (TT, orange) to prevent readthrough of gene expression.

