

**Table S2 – Characteristics of the bacterial strains and plasmids used in this study**

Strain/Plasmid	Relevant features	Reference
<b><i>Escherichia coli</i></b>		
<b>CM404</b>	<i>E. coli</i> strain harboring the self-transmissible plasmid pRK2013 enabling the conjugative transfer of RSF1010-derived pC plasmids into cyanobacteria	(Mermet-Bouvier and Chauvat, 1994)
<b>TOP10</b>	<i>E. coli</i> strain for cloning	Invitrogen
<b>NEB 5-alpha</b>	<i>E. coli</i> strain for cloning	New England Biolabs
<b>NEB 10-beta</b>	<i>E. coli</i> strain for cloning	New England Biolabs
<b>Cloning vectors</b>		
<b>pTwist</b>	Amp <sup>R</sup> plasmid vector	Twist Biosciences
<b>pUC4K</b>	Source of the Km <sup>R</sup> marker with no transcription terminator	Pharmacia
<b>pC</b>	RSF1010-derived plasmid vector (Sp <sup>R</sup> /Sm <sup>R</sup> , Cm <sup>R</sup> ) harboring the strong p <sub>R</sub> promoter for constitutive gene expression in <i>E. coli</i> and cyanobacteria	(Veaudor et al., 2018)
<b>Replicative plasmids for strong constitutive expression of terpene synthase genes in cyanobacteria</b>		
<b>pCLS</b>	pC derived plasmid (Sp <sup>R</sup> /Sm <sup>R</sup> , Cm <sup>S</sup> ) harboring the <i>Mentha spicata</i> 4S-limonene synthase gene (LS) cloned between the <i>NdeI</i> and <i>EcoRI</i> restriction sites	(Chenebault et al., 2020)
<b>pCFS</b>	pC derivative (Sp <sup>R</sup> /Sm <sup>R</sup> , Cm <sup>S</sup> ) carrying the <i>Picea abies</i> farnesene synthase gene (FS) cloned between <i>NdeI</i> and <i>EcoRI</i>	(Blanc-Garin et al., 2022)
<b>Replicative plasmids for strong constitutive expression of terpene synthase encoding genes and <i>crtE</i> genes in cyanobacteria</b>		
<b>pCLS-CrtE<sub>6803</sub></b>	pCLS derived plasmid (Sm <sup>R</sup> /Sp <sup>R</sup> ) harboring the <i>crtE</i> gene from S.6803 cloned downstream the LS gene between the <i>EcoRI</i> and <i>AclI</i> sites	This study
<b>pCLS-CrtE<sub>7002</sub></b>	pCLS derived plasmid (Sm <sup>R</sup> /Sp <sup>R</sup> ) harboring the <i>crtE</i> gene from S.7002 cloned downstream the LS gene, between the <i>EcoRI</i> and <i>AclI</i> sites	This study
<b>pCFS-CrtE<sub>6803</sub></b>	pCFS derived plasmid (Sm <sup>R</sup> /Sp <sup>R</sup> ) harboring the <i>crtE</i> gene from S.6803 cloned downstream the FS gene, between the <i>EcoRI</i> and <i>AclI</i> sites	This study
<b>pCFS-CrtE<sub>7002</sub></b>	pCFS derived plasmid (Sm <sup>R</sup> /Sp <sup>R</sup> ) harboring the <i>crtE</i> gene from S.7002 cloned downstream the FS gene, between the <i>EcoRI</i> and <i>AclI</i> sites	This study
<b>Plasmids used for deletion of <i>cruF crtR</i>, <i>phaAB</i>, <i>ccmK</i> or <i>crtE</i> in <i>Synechocystis</i> PCC 6803</b>		
<b>pTwist_CrtR</b>	pTwist harboring an <i>EcoRV</i> restriction site flanked by two platforms of homology for deletion of the <i>crtR</i> open reading frame from the ATG start codon to the 245 amino acids of <i>CrtR</i> from <i>Synechocystis</i> PCC 6803. The 67 last amino acids are conserved to avoid the truncation of <i>slr1543</i> .	This study & Twist Biosciences
<b>pCrtR_Km</b>	pTwist_CrtR harboring a Km <sup>R</sup> cassette cloned in the <i>EcoRV</i> site	This study
<b>pTwist_CruF</b>	pTwist harboring an <i>EcoRV</i> restriction site flanked by two platforms of homology for deletion of the <i>cruF</i> open reading frame from the amino acid 100 to the amino acid 271 from <i>Synechocystis</i> PCC 6803 to avoid truncation of genes <i>sll0813</i> and <i>sll0815</i>	This study and Twist Biosciences
<b>pCruF_Km</b>	pTwist_CruF harboring a Km <sup>R</sup> cassette cloned in the <i>EcoRV</i> site	This study
<b>pTwist_phaAB</b>	pTwist harboring an <i>EcoRV</i> restriction site flanked by two platforms of homology for deletion or <i>phaA</i> and <i>phaB</i> genes	This study and Twist Biosciences

<b>pphaAB_Km</b>	pTwist_phaAB harboring a Km <sup>R</sup> cassette cloned in the <i>EcoRV</i> site	This study
<b>pTwist_CcmK3K4_Km</b>	pTwist_CcmK3K4 harboring a Km <sup>R</sup> cassette cloned in the <i>SwaI</i> et <i>BamHI</i> site	This study and Twist Bioscience
<b>Replicative plasmids for strong constitutive expression of <i>rbc</i> operon or <i>prk</i> genes from various cyanobacteria</b>		
<b>pC-rbcLXS_Scy6803</b>	pC derivative plasmid (Sp <sup>R</sup> /Sm <sup>R</sup> ) harboring the ribulose-1,5-bisphosphate carboxylase/oxygenase (RuBisCO) encoding operon ( <i>rbcLXS</i> ) from <i>Synechocystis</i> PCC 6803 cloned in between the <i>NdeI</i> and <i>EcoRI</i> restriction sites	This study
<b>pC-rbcLS_Sco7942</b>	pC derivative plasmid (Sp <sup>R</sup> /Sm <sup>R</sup> ) harboring the ribulose-1,5-bisphosphate carboxylase/oxygenase (RuBisCO) encoding operon ( <i>rbcLS</i> ) from <i>Synechococcus</i> PCC 7942 cloned in between the <i>NdeI</i> and <i>EcoRI</i> restriction sites	This study
<b>pC-rbcLXS_Sco7002</b>	pC derivative plasmid (Sp <sup>R</sup> /Sm <sup>R</sup> ) harboring the ribulose-1,5-bisphosphate carboxylase/oxygenase (RuBisCO) encoding operon ( <i>rbcLXS</i> ) from <i>Synechococcus</i> PCC 7002 cloned in between the <i>NdeI</i> and <i>EcoRI</i> restriction site	This study
<b>pC- rbcLXS_Cya7425</b>	pC derivative plasmid (Sp <sup>R</sup> /Sm <sup>R</sup> ) harboring the ribulose-1,5-bisphosphate carboxylase/oxygenase (RuBisCO) encoding operon ( <i>rbcLXS</i> ) from <i>Cyanothece</i> PCC 7425 cloned in between the <i>NdeI</i> and <i>EcoRI</i> restriction site	This study
<b>pC-prk_Cya7425</b>	pC derivative plasmid (Sp <sup>R</sup> /Sm <sup>R</sup> ) harboring the phosphoribulokinase encoding gene ( <i>prk</i> ) from <i>Cyanothece</i> PCC 7425 cloned in between the <i>NdeI</i> and <i>EcoRI</i> restriction sites	This study

References:

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