

Supplementary Information

Table S1 Primers used in this study

Target gene	Name	Sequence 5' to 3'	Purpose of primers	Product size (bp)	T _m (°C)/Cycles
<i>rbcL</i>	rbcL_F	GGACTAGTATGGTACAAGCCAAAGC	PCR	1413	58 °C/30
	rbcL_R	AAAACCTGCAGTTAGAGGGTATCCATGGC			
<i>rbcX</i>	rbcX_F	GCTCTAGATTACCGGGTAGAGTGTTCAT	PCR / RT-PCR	433	58 °C/26
	rbcX_R	GGACTAGTTAGGACGGGGAGAACGTT			
<i>rbcS</i>	rbcS_F	GGACTAAGTCATCAGCAAGGAAAACTTTA	PCR / RT-PCR	373	58 °C/26
	rbcS_R	AACTGCACTTAGTAACGGCCTGGTTTGG			
UppsbA2	UppsbA2_F	TGCCTGTCAGCAAAACAACTT	PCR	-	58 °C
DSpsbA2	DSpsbA2_R	CGAGGGCAATCATCAATTCCG			
CM	CM_F	GAGTTGATCGGGCACGTAAG	PCR	899	58 °C
	CM_R	CTCGAGGGCTTGGATTCTCTCAC			
16S rRNA	16S_F	AGTTCTGACGGTACCTGATGA	RT-PCR	521	59 °C/8
	16S_R	GTCAAGCCTTGGTAAGGTTCT			
RT_	rbcL_F	GGTATCACCATGGGCTTCGTTGACCT	RT-PCR	411	58 °C/30
	rbcL_R	AAAACCTGCAGTTAGAGGGTATCCATGGC			
RT_	phaA_F	CATGATGGTTGACGGACAG	RT-PCR	310	58 °C/30
	RTphaA_R	AGACTTTCCACGGTGGTGT			
RT_	phaB_F	GCTCCATTGTGGCCATTAGT	RT-PCR	255	58 °C/26
	RTphaB_R	CAATTCCCTCCGGTTACCA			
RT_	phaC_F	GGGCACATTAGCCTGTGTT	RT-PCR	267	58 °C/26
	RTphaC_R	CCCATAATATCGGGCACATC			
RT_	phaE_F	GGCCATGGCAGACTATCAAT	RT-PCR	384	58 °C/23
	RTphaE_R	TAGCCTGGGTTGCTCTGT			
RT_	plsX_F	AAGGGGTGGTGGAAATGGAA	RT-PCR	467	59 °C/28
	RTplsX_R	AAGTACGTCCTCCTTCGG			
RT_	plsC_F	TCTCTACCGGGCTTGAAATG	RT-PCR	508	58 °C/32
	RTplsC_R	CGCCTTACCAATGCGAATAGT			
RT_	glgX_F	GAGCTTCATCGAGGACGGAA	RT-PCR	360	58 °C/26
	RTglgX_R	GCCCCGAATTGGGGTTGCGGG			

Table S2 The band intensity data detected by Syngene® Gel Documentation (Syngene, Frederick, MD, USA) in Figures 8 and 9.

(A) The band intensity of each gene (transcript level)

Genes	Normal BG ₁₁			BG ₁₁ -N			BG ₁₁ -P			BG ₁₁ -NP		
	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>
<i>phaA</i>	0.1	0.7	0.7	5.0	6.2	5.5	1.0	4.0	5.0	2.0	4.0	5.5
<i>phaB</i>	0.3	0.3	0.3	3.0	3.5	3.5	0.5	0.5	0.5	1.2	4.0	3.8
<i>phaC</i>	0.2	0.2	0.1	2.5	4.0	3.2	0.8	0.9	1.0	1.0	3.0	3.0
<i>phaE</i>	0.1	0.2	0.2	0.3	0.3	0.3	0.3	0.4	0.4	0.1	0.1	0.1
<i>plsX</i>	0.2	0.2	0.1	4.0	4.0	0.8	0.4	0.8	1.8	0.4	0.4	0.4
<i>plsC</i>	1.2	0.8	2.0	2.5	3.0	2.0	3.1	3.4	3.6	1.2	1.3	1.6
<i>glgX</i>	0.2	0.2	0.2	0.6	2.7	2.5	1.0	1.5	1.6	1.2	2.0	2.5
<i>16S</i>	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	0.8	0.8

(B) The ratios of band intensity of each gene/*16s*

Genes	Normal BG ₁₁			BG ₁₁ -N			BG ₁₁ -P			BG ₁₁ -NP		
	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>
<i>phaA</i>	0.1	0.7	0.7	5.0	6.2	5.5	1.0	4.0	5.0	2.0	5.0	6.9
<i>phaB</i>	0.3	0.3	0.3	3.0	3.5	3.5	0.5	0.5	0.5	1.2	5.0	4.8
<i>phaC</i>	0.2	0.2	0.1	2.5	4.0	3.2	0.8	0.9	1.0	1.0	2.8	2.9
<i>phaE</i>	0.1	0.2	0.2	0.3	0.3	0.3	0.3	0.4	0.4	0.1	0.1	0.1
<i>plsX</i>	0.2	0.2	0.1	4.0	4.0	0.8	0.4	0.8	1.8	0.4	0.5	0.5
<i>plsC</i>	1.2	0.8	2.0	2.5	3.0	2.0	3.1	3.4	3.6	1.2	1.6	2.0
<i>glgX</i>	0.2	0.2	0.2	0.6	2.7	2.5	1.0	1.5	1.6	1.2	2.5	3.1

(C) Fold change value of “band intensity ratio of each gene/*16s* under stress” divided by “band intensity ratio of each gene/*16s* under normal condition”, shown in Figure 9.

Genes	BG ₁₁ -N			BG ₁₁ -P			BG ₁₁ -NP		
	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>	WTc	OX <i>rbcL</i>	OX <i>rbcLXS</i>
<i>phaA</i>	50.0	8.9	7.9	10.0	5.7	7.1	30.0	7.1	9.8
<i>phaB</i>	9.7	10.9	12.5	1.6	1.6	1.8	3.9	15.6	17.0
<i>phaC</i>	12.5	20.0	32.0	4.0	4.5	10.0	5.0	14.0	29.0
<i>phaE</i>	3.0	1.5	1.5	3.0	2.0	2.0	1.0	0.6	0.6
<i>plsX</i>	20.0	20.0	8.0	2.0	4.0	18.0	2.0	2.5	5.0
<i>plsC</i>	2.1	3.8	1.0	2.6	4.3	1.8	1.0	2.0	1.0
<i>glgX</i>	3.0	13.5	12.5	5.0	7.5	8.0	6.0	12.5	15.6

Figure S1 : Agarose gel electrophoresis of RT-PCR product of *rbcL* transcript in *Synechocystis* sp. PCC 6803 grown under normal BG₁₁ condition, shown in **Figure 2D**. The *16s* rRNA transcript was used as the reference.

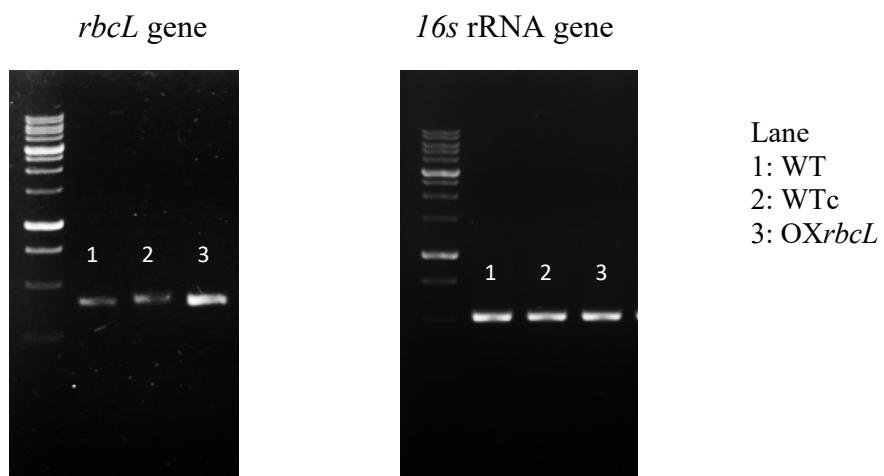


Figure S2 : Agarose gel electrophoresis of RT-PCR product of *rbcL*, *rbcX*, *rbcS* transcripts in *Synechocystis* sp. PCC 6803 grown under normal BG₁₁ condition, shown in **Figure 2D**. The *16s* rRNA transcript was used as the reference.

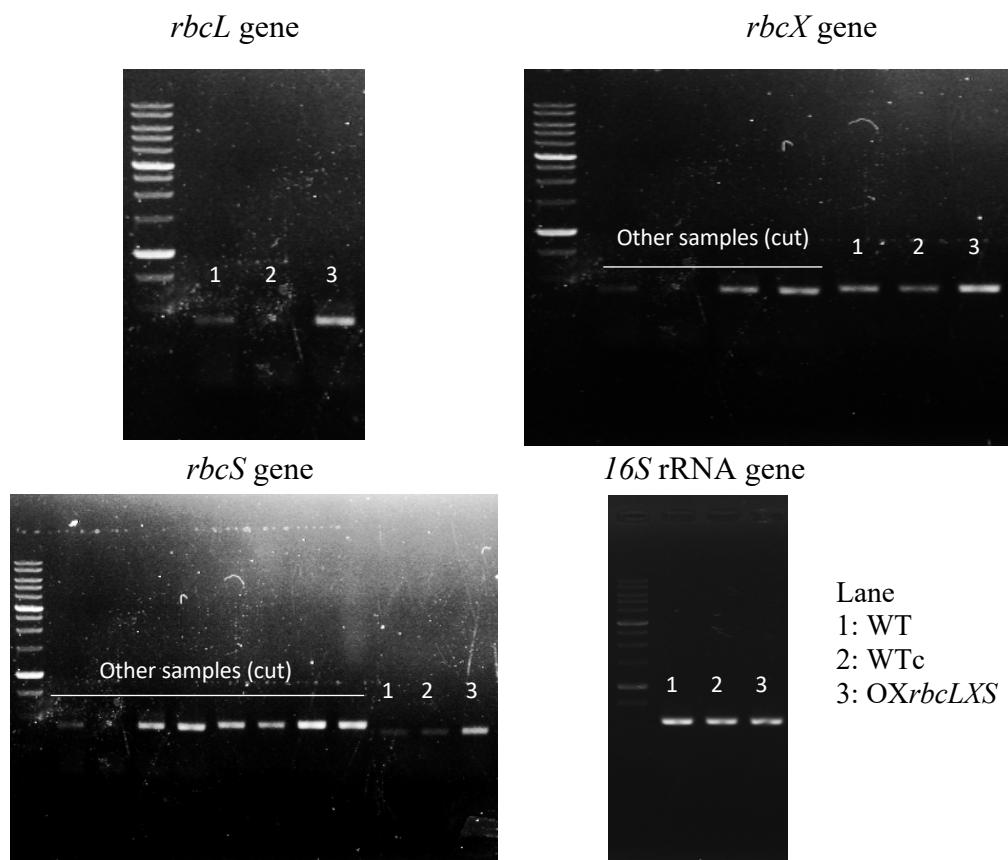
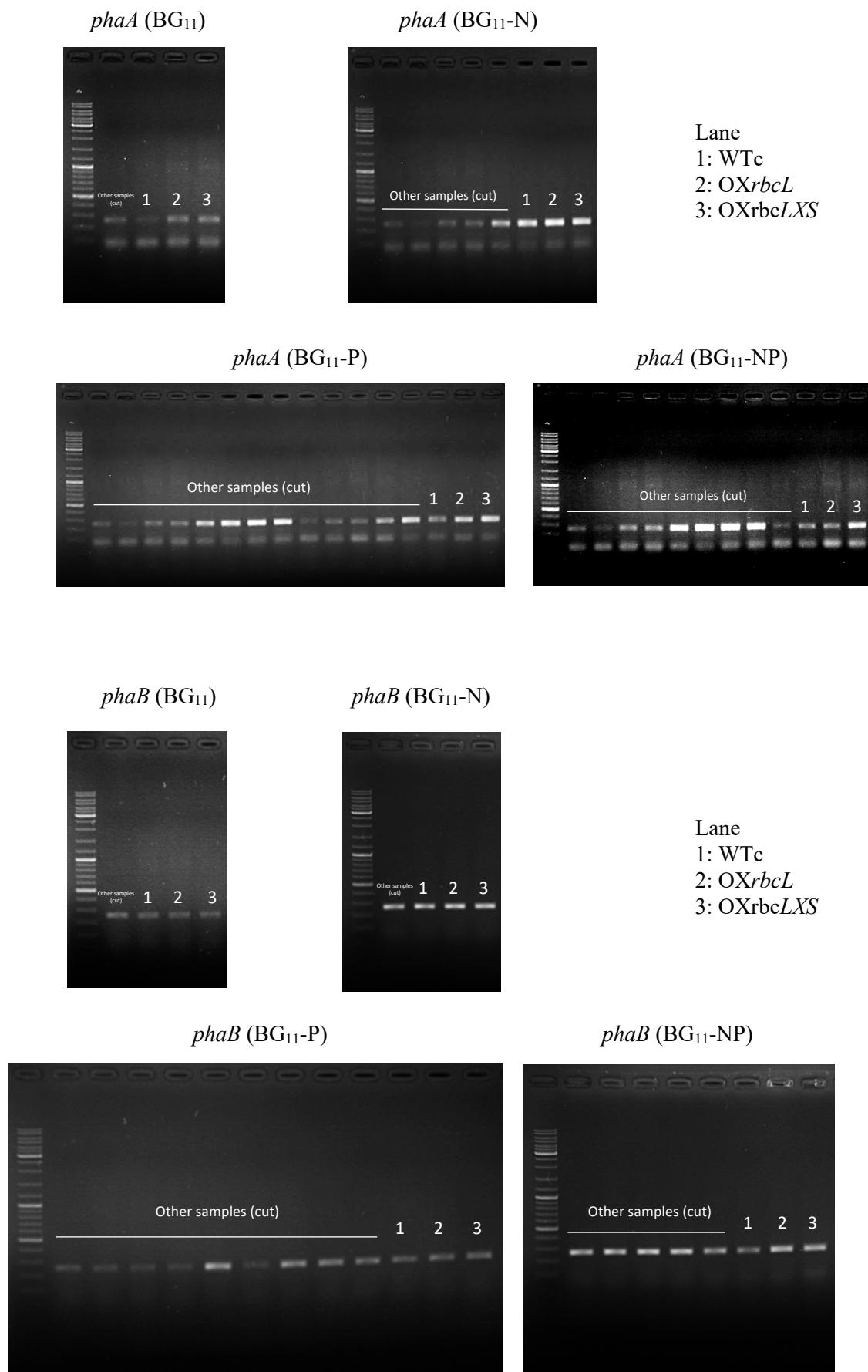
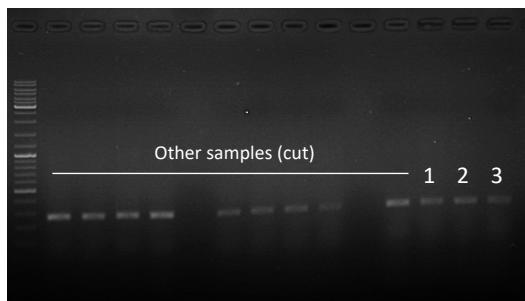


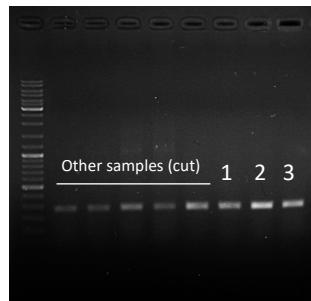
Figure S3 : Agarose gel electrophoresis of RT-PCR product of *phaA*, *phaB*, *phaC*, *phaE*, *plsX*, *plsC*, *rbcS*, *glgX*, and *16s* transcripts in *Synechocystis* sp. PCC 6803 grown under stresses and normal BG₁₁ condition, shown in **Figure 8**. The *16s* rRNA transcript was used as the reference.



phaC (BG₁₁)

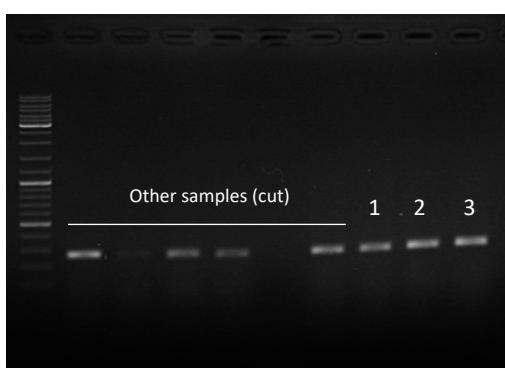


phaC (BG_{11-N})

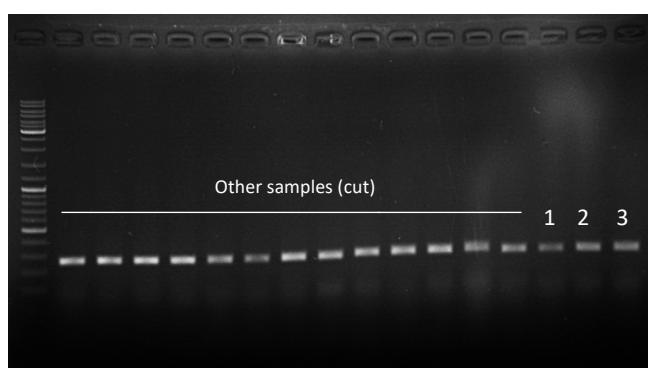


Lane
1: WTc
2: OXrbcL
3: OXrbcLXS

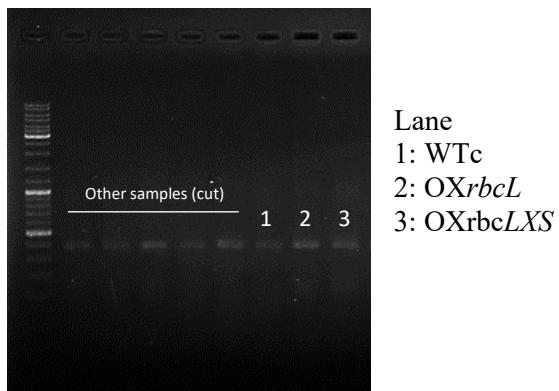
phaC (BG_{11-P})



phaC (BG_{11-NP})

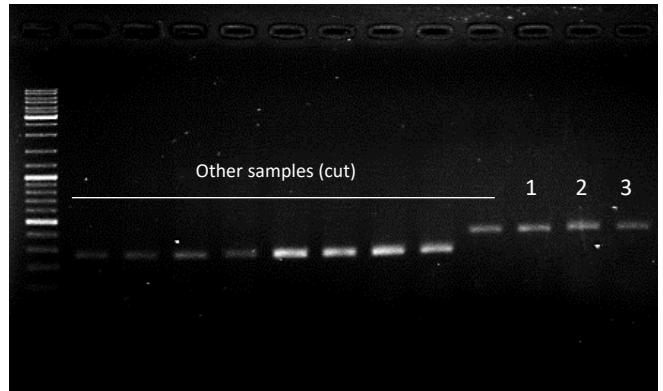


phaE (BG₁₁)

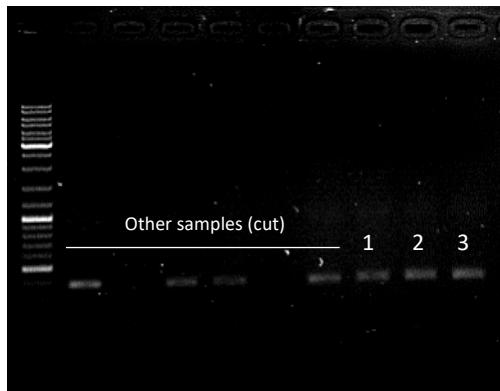


Lane
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2: OXrbcL
3: OXrbcLXS

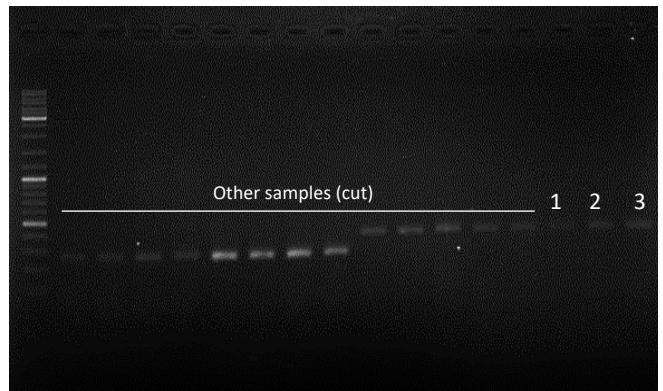
phaE (BG_{11-N})



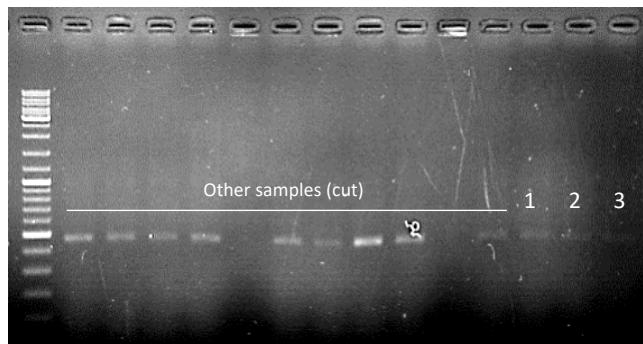
phaE (BG_{11-P})



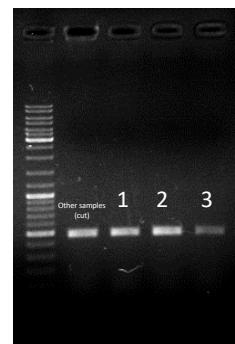
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plsX (BG₁₁)

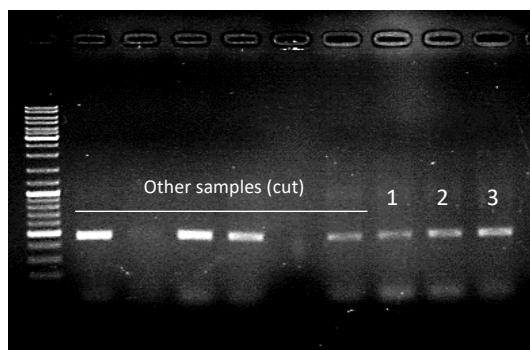


plsX (BG_{11-N})

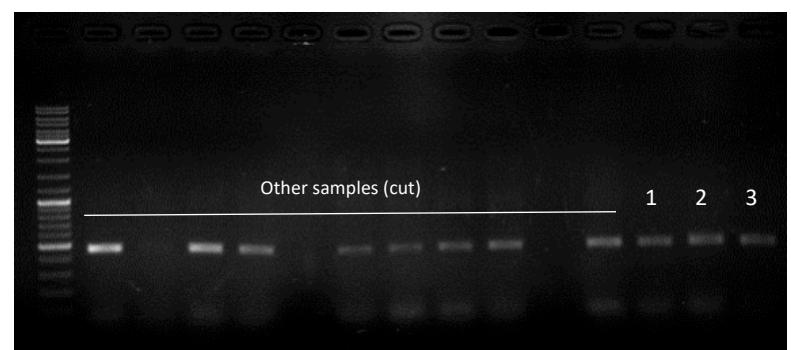


Lane
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2: OXrbcL
3: OXrbcLXS

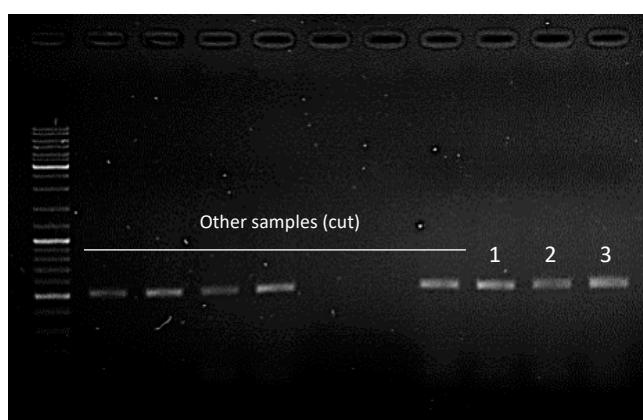
plsX (BG_{11-P})



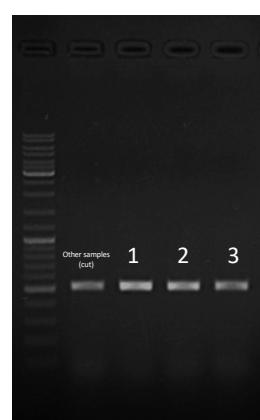
plsX (BG_{11-NP})



plsC (BG₁₁)

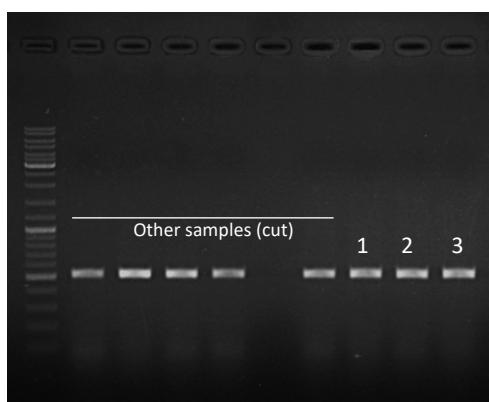


plsC (BG_{11-N})

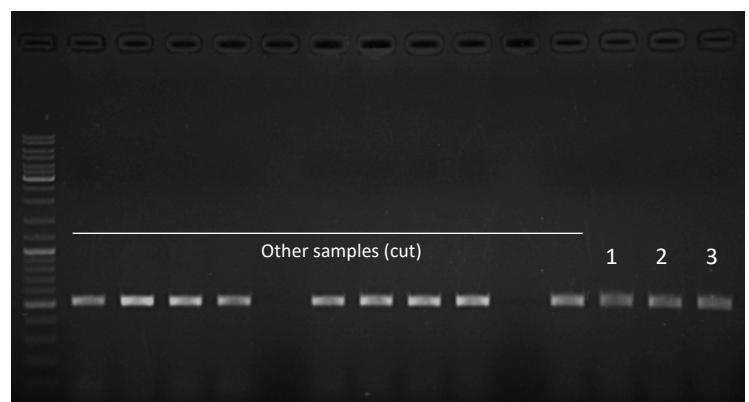


Lane
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2: OXrbcL
3: OXrbcLXS

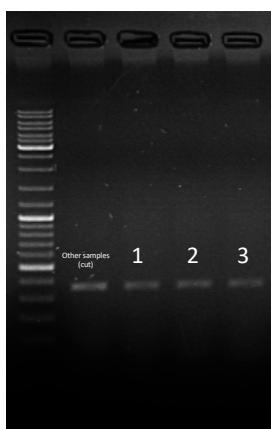
plsC (BG_{11-P})



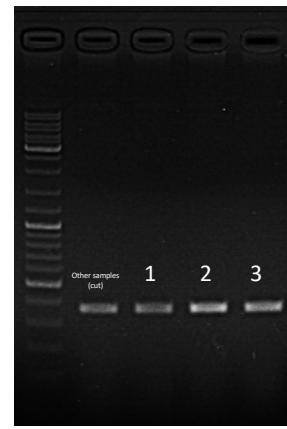
plsC (BG_{11-NP})



glgX (BG₁₁)

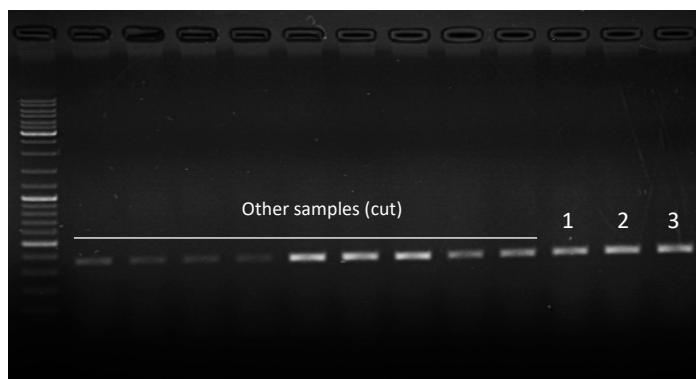


glgX (BG_{11-N})

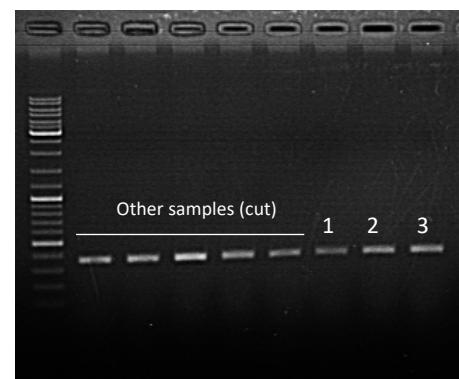


Lane
1: WTc
2: OXrbcL
3: OXrbcLXS

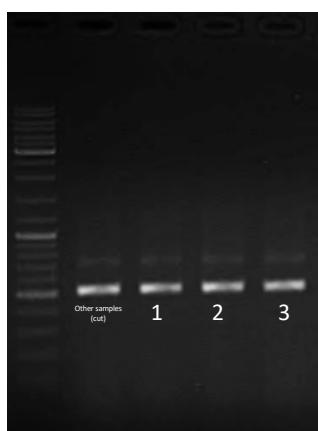
glgX (BG_{11-P})



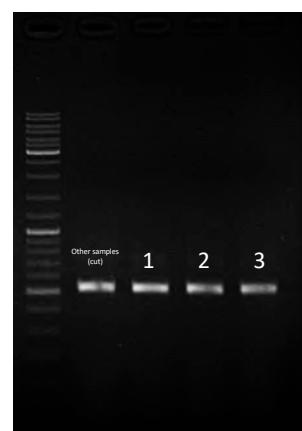
glgX (BG_{11-NP})



16S rRNA (BG₁₁)

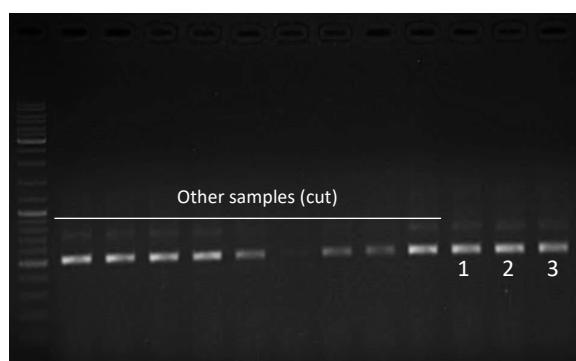


16S rRNA (BG_{11-N})



Lane
1: WTc
2: OXrbcL
3: OXrbcLXS

16S rRNA (BG_{11-P})



16S rRNA (BG_{11-NP})

