

TABLE S1. Oligonucleotide primers used in this study

Primer	Target gene or plasmid	Sequence (5'- 3') <sup>a</sup>	Cohesive end
HisF	pNVHis18	GCTTGCGGGTAGATCCCCT	
HisR	pNVHis18, His-tag	TTAC <u>GAATTCTAGTGGTGGTGGTGGTGGT</u>	EcoRI
		GAGCTCGGTACCCGGGGAT	
TacF	<i>tac</i> promoter	AAG <u>CAAGCTT</u> CTTATCGACTGCACGGTG	HindIII
TacR	<i>tac</i> promoter	CGCG <u>CTGCAG</u> CATATGATATCTCCT	PstI
RHA1-xylB1F	<i>xylB1</i> from <i>R. jostii</i> RHA1	CGAT <u>GAATTCT</u> GATCACCGACAAGGCAC	EcoRI
RHA1-xylB1R	<i>xylB1</i> from <i>R. jostii</i> RHA1	ACTAA <u>AGCTT</u> ACGCCGCCTCGACTACA	HindIII
RHA1-xylB2F	<i>xylB2</i> from <i>R. jostii</i> RHA1	ATGC <u>GAATTCT</u> TCTATCAGCCGACGGTC	EcoRI
RHA1-xylB2R	<i>xylB2</i> from <i>R. jostii</i> RHA1	CTAAA <u>AGCTT</u> CGGT CGATGTCGGTGTCA	HindIII
XylA1	<i>xylA</i> from <i>S. lividans</i> TK23	ACG <u>ACTGCAG</u> ATGA ACTACCAGCCCA	PstI
XylA2	<i>xylA</i> from <i>S. lividans</i> TK23	TAAT <u>GGTACC</u> AGCGGGCGCCGAGGAG	KpnI
xylB1	<i>xylB</i> from <i>S. lividans</i> TK23	ATT <u>ACTGCAG</u> ATGTCAGCAGCCGAGGG	PstI
xylB2	<i>xylB</i> from <i>S. lividans</i> TK23	AATT <u>GGTACC</u> AGCGGGAGGCCCGTCC	KpnI
xyl-tac1	Expression cassette of <i>xylB</i>	GGGTTTCCCAGTCACGACGTTGTA	
xyl-tac2	Expression cassette of <i>xylB</i>	CTCG <u>AAGCTT</u> CGGCTCGTATGTTGTGT	HindIII

<sup>a</sup> The generated restriction sites used in the cloning procedure are underlined.

Figure S1. Map of plasmid pXYLAB. The plasmid contained both *xylA* and *xylB*, which expression was controlled by the *tac* promoter individually. The *repA* and *repB* encoded the region and proteins related to DNA replication in *Rhodococcus* strains. Other abbreviations of the plasmid as shown: kan, kanamycin resistance; neo, neomycin resistance phenotype; ori, origin of replication and P, promoter.

