

TABLE S2. Primers used in this study.

Primer	Sequence* (5'—3')	Restriction enzyme† and use
P1	GTTGCCGTCCCTGGTATGG	PCR <i>ntrC</i> from the <i>A. vinelandii</i> genome
P2	GTGGACACGGCATAGATTTTC	PCR <i>ntrC</i> from the <i>A. vinelandii</i> genome
P3	<u>AGAGCTCAGCCATGCC</u> CAGGAG	<i>SacI</i> ; <i>ntrC</i> deletion mutation construction
P4	<u>AGAGCTCGGCTTGGG</u> CAGGTATTC	<i>SacI</i> ; <i>ntrC</i> deletion mutation construction
P5	AAAT <u>CTAGAAGCAGCGCTTC</u> GAAAC	<i>XbaI</i> ; PCR <i>nasAB</i> promoter region from the <i>A. vinelandii</i> genome
P6	<u>TGGATCCATGCCG</u> GCCAGCACCGGAGAG	<i>BamHI</i> ; PCR <i>nasAB</i> promoter region from the <i>A.</i>

		<i>vinelandii</i> genome
P8	AGCTGGGGCCGCAATTC	PCR tandem <i>rrnBI</i> from pPROBE-NT
P9	ATTCAGCTTCCGATC	PCR tandem <i>rrnBI</i> from pPROBE-NT
P10	CCGACAGAACGGGATAGGACAAAGGCGTC	<i>nasAB</i> leader small ORF mutation
P11	CCTATCCCGTTCTGTCAAAGGCGTC	<i>nasAB</i> leader small ORF mutation
P12	GCAGTGTTTCCCGACAGGAATGGACAAAGGCGTC	<i>nasAB</i> leader hairpin II deletion mutation
P13	CTGTCGGGAAACACTGC	<i>nasAB</i> leader hairpin II deletion mutation
P14	GGGGATCGGAAGCTGAATG	Confirmation of tested sequence in pVnflacZ(a/b)
P15	GTCTCATGAGCGGATAC	Confirmation of <i>vnf</i> in pWhite
P16	<u>AGTCGACAAGCTT</u> CCTCGCTCACTGACTC	<i>Sal</i> I, <i>Hind</i> III; PCR <i>bla</i> and pM1 of pBbluscript II KS

		(+)
P17	AGAGCTCAGGTGGCACTTTTCG	PCR <i>bla</i> and pM1 of pBbluscript II KS (+)
P18	GTATTACCGCCTTTGAGTG	Confirmation of <i>vnf</i> in pWhite
P19	GTGATCTGTCGGTTTTTC	<i>nasAB</i> leader hairpin I deletion mutation
P20	AAACCGACAGATCACATAAACGTGGAGGGCAGTG	<i>nasAB</i> leader hairpin I deletion mutation
P21	AAATCTAGAACCAAGACAGTGCAAG	<i>Xba</i> I; <i>nasAB</i> promoter deletion construction
P22	AAATCTAGAAACCCATAAGAGG	<i>Xba</i> I; <i>nasAB</i> promoter deletion construction
P23	AAATCTAGAGCTCTGCTCGCCCTGTTG	<i>Xba</i> I; <i>nasAB</i> promoter deletion construction
P24	AAATCTAGAGTCATCCGTGAAAC	<i>Xba</i> I; <i>nasAB</i> promoter deletion

		construction
P25	CAAGGCGATTAAGTTGGGTAAC	Confirmation of tested sequence in pVnflacZ(a/b)
P26	GTAAAACGACGGCCAGTG	pBluescript II KS(+) sequencing
P27	CAGGAAACAGCTATGAC	pBluescript II KS(+) sequencing
P28	TAACACAGCCCCTATATCGAGAAAACCGAC	σ^{54} binding site mutation
P29	TATAGGGGCTGTGTTAGTTTCACGGATGACTG	σ^{54} binding site mutation
P30	CAGAACGGGAATGGACAATTTTTGTTTTCCGCTTT C	Terminator hairpin deletion mutation
P31	AATTGTCCATTCCCGTTCTG	Terminator hairpin deletion mutation
P32	<u>AAGGATCCG</u> CTTATGTGATCTGTCTG	<i>Bam</i> HI; Transcriptional fusion construction
P33	AAAGAGCT <u>CATCATCTCG</u> CCAGTTC	<i>Sac</i> I; PCR <i>vnf</i> sequence
P34	AAACGCCT <u>CGAGA</u> AAGAGCATG	<i>Xho</i> I; PCR <i>vnf</i> sequence

P35	CGCCTGGCTCGACGAATG	<i>nasA</i> promoter region cloning confirmation
P36	GTCCATTCCCGTTCTGTC	<i>nasA</i> promoter region cloning confirmation
P37	<u>TCGAATTCC</u> ATGCTGCGCATCCTCCTG	<i>EcoRI</i> ; Clone <i>nasT</i> into pDK6
P38	ATA <u>GATCTC</u> AGCTTCCCAGCATGTCGTGCATG	<i>BglII</i> ; Clone <i>nasT</i> into pDK6
P39	ATAGA <u>ATTCC</u> ATGACAGACCACCACGCA ACTTC	<i>EcoRI</i> ; Clone <i>nasS</i> into pDK6
P40	ATA <u>AGATCTT</u> TAGGAGGATGCGCAG	<i>BglII</i> ; Clone <i>nasS</i> into pDK6
P41	TCCGTTGTGGGAAAGTTATC	Construct pBTW
P42	<u>AGCGGCCGCT</u> CACAATTCCACAC	Construct pBTW
P43	AGAG <u>CGGCCG</u> CCGACAGATCACATAAG	<i>NotI</i> ; PCR <i>nasA'</i> - ' <i>lacZ</i> fragment

* The underlined sequences are the restriction sites.

† Restriction enzymes that can digest the underlined sequences.