

Table S1. Strains Used in This Study, Related to Experimental Procedures

Name	Genotype	Construction	Reference/Source
<i>C. crescentus</i>			
CB15N	Synchronizable variant of wild-type strain CB15		Evinger & Agabian, 1977
CS606	CB15N Δ blaM		West et al., 2002
EK61	CB15N P _{xyi} ::P _{xyi} -mcherry-CC1398		Werner et al., 2009
EK363	CB15N P _{xyi} ::P _{xyi} -pstS-mcherry	Transformation of CB15N with pEK301	This work
EK389	CB15N Δ stpAB P _{xyi} ::P _{xyi} -pstS-mcherry	Transformation of SW51 with pEK301	This work
EK392	CB15N P _{xyi} ::P _{xyi} -yfp	Transformation of CB15N with pXYFPN-5	This work
EK393	CB15N Δ stpAB P _{xyi} ::P _{xyi} -mcherry	Transformation of SW51 with pXCHYN-5	This work
EK416	CB15N P _{xyi} ::P _{xyi} -mcherry	Transformation of CB15N with pXCHYN-5	This work
EK417	CB15N Δ stpAB P _{xyi} ::P _{xyi} -yfp	Transformation of SW51 with pXYFPN-5	This work
EK424	CB15N pstS::miniTn5LacZ P _{xyi} ::P _{xyi} -pstS-mcherry	Transformation of EK425 with pEK301	This work
EK425	CB15N pstS::miniTn5LacZ	Transduction of kan ^R from YB2811	This work
EK486	CB15N Δ stpCD P _{xyi} ::P _{xyi} -yfp	Transformation of SS250 with pXYFPN-5	This work
EK487	CB15N Δ stpCD P _{xyi} ::P _{xyi} -mcherry	Transformation of SS250 with pXCHYN-5	This work
GB255	CB15N popZ:: Ω		Bowman et al., 2008
JK5	CB15N Δ bacAB		Kühn et al., 2010
MT304	CB15N Δ pbpC		Kühn et al., 2010
NR4042	CB15N Δ stpX		Viollier, P. H. (unpubl.)
PV5064	CB15N stpX::stpX-gfp		Hughes et al., 2010
SS141	CB15N Δ stpA P _{xyi} ::P _{xyi} -stpB-mcherry	Transformation of SW49 with pSW32	This work
SS142	CB15N Δ stpB P _{xyi} ::P _{xyi} -stpA-mcherry	Transformation of SW50 with pSW35	This work
SS160	CB15N stpB-mcherry	Gene replacement in CB15N with pSS109	This work
SS165	CB15N Δ bla P _{xyi} ::P _{xyi} -stpB-bla	Transformation of CS606 with pSS120	This work
SS167	CB15N Δ pbpC P _{xyi} ::P _{xyi} -stpB-mcherry	Transformation of MT304 with pSW32	This work
SS172	CB15N Δ bla P _{xyi} ::P _{xyi} -stpA-bla	Transformation of CS606 with pSS119	This work
SS191	CB15N ftsZ::P _{xyi} -ftsZ P _{van} ::P _{van} -stpB-mcherry	Transformation of YB1585 with pSS142	This work
SS213	CB15N Δ stpX P _{xyi} ::P _{xyi} -stpB-mcherry	Transduction of kan ^R from SW30 to NR4042	This work
SS214	CB15N stpB::stpB-mcherry pP _{xyi} -stpAB	Transformation of SS160 with pSW64	This work
SS216	CB15N Δ stpAB pP _{xyi} -TAT-dimer2	Transformation of SW51 with pEJ216	This work
SS220	CB15N Δ stpA stpB::stpB-His	Transformation of SW49 with pSS187	This work
SS224	CB15N divJ:: Ω P _{xyi} ::P _{xyi} -stpB-mcherry	Transduction of spec ^R from YB3202 to SW30	This work
SS228	CB15N P _{xyi} ::P _{xyi} -stpC-mcherry	Transformation of CB15N with pSS204	This work
SS233	CB15N stpB::stpB-His	Transformation of CB15N with pSS187	This work
SS234	CB15N Δ stpAB P _{xyi} ::P _{xyi} -stpD-gfp	Transformation of SW51 with pSS202	This work
SS236	CB15N Δ stpAB P _{xyi} ::P _{xyi} -stpC-mcherry	Transformation of SW51 with pSS204	This work
SS239	CB15N Δ stpC	Gene replacement in CB15N using pSS209	This work
SS240	CB15N Δ stpC P _{xyi} ::P _{xyi} -stpD-gfp	Transformation of SS239 with pSS202	This work
SS243	CB15N stpD::stpD-gfp P _{xyi} ::P _{xyi} -stpA-mcherry	Transformation of SW33 with pSS205	This work
SS244	CB15N stpD::stpD-His	Transformation of CB15N with pSS206	This work
SS247	CB15N stpC::stpC-His	Transformation of CB15N with pSS210	This work
SS248	CB15N stpD::stpD-gfp	Transformation of CB15N with pSS205	This work
SS250	CB15N Δ stpCD	Gene replacement in SS239 using pSS208	This work
SS252	CB15N Δ stpD	Gene replacement in CB15N using pSS208	This work
SS258	CB15N stpB::stpB-mcherry pBXMCS-2	Transformation of SS160 with pBXMCS-2	This work
SS263	CB15N Δ stpD P _{xyi} ::P _{xyi} -stpC-mcherry	Transformation of SS252 with pSS204	This work
SS264	CB15N Δ stpB P _{xyi} ::P _{xyi} -stpD-gfp	Transformation of SW50 with pSS202	This work
SS265	CB15N Δ stpB P _{xyi} ::P _{xyi} -stpC-mcherry	Transformation of SW50 with pSS204	This work
SS269	CB15N stpD::stpD-gfp pP _{xyi} -TATdimer2	Transformation of SS248 with pEJ216	This work
SS272	CB15N Δ stpAB P _{xyi} ::P _{xyi} -gspG-mcherry	Transformation of SW51 with pJK86	This work
SS273	CB15N Δ bla P _{xyi} ::P _{xyi} -stpC-blaM	Transformation of CS606 with pSS220	This work
SS274	CB15N Δ bla P _{xyi} ::P _{xyi} -stpD-blaM	Transformation of CS606 with pSS221	This work
SS277	CB15N stpD::stpD-gfp P _{xyi} ::P _{xyi} -gspG-mcherry	Transformation of SS248 with pJK86	This work
SS281	CB15N Δ bacAB P _{xyi} ::P _{xyi} -stpB-mcherry	Transformation of JK5 with pSW32	This work
SS283	CB15N stpD::stpD-gfp P _{xyi} ::P _{xyi} -elpS-mcherry	Transformation of SS248 with pSW67	This work
SS284	CB15N Δ stpAB P _{xyi} ::P _{xyi} -elpS-mcherry	Transformation of SW51 with pSW67	This work
SS292	CB15N popZ:: Ω P _{xyi} ::P _{xyi} -stpB-mcherry	Transformation of GB255 with pSW32	This work
SS294	CB15N Δ stpAB P _{xyi} ::P _{xyi} -malA-mcherry	Transformation of SW51 with pSS227	This work

Table S1. Strains Used in This Study (continued)

Name	Relevant Genotype/Description	Construction	Reference/Source
SS297	CB15N <i>stpD::stpD-gfp</i> P _{xyl} ::P _{xyl} - <i>malA-mcherry</i>	Transformation of SS248 with pSS227	This work
SS299	CB15N <i>stpD::stpD-gfp</i> P _{xyl} ::P _{xyl} - <i>pstS-mcherry</i>	Transformation of SS248 with pJK101	This work
SS302	CB15N Δ <i>stpAB</i> P _{xyl} ::P _{xyl} - <i>pstS-mcherry</i>	Transformation of SW51 with pJK101	This work
SS304	CB15N Δ <i>stpCD</i> pP _{xyl} - <i>TAT-dimer2</i>	Transformation of SS250 with pEJ216	This work
SS388	CB15N <i>stpD::stpD-gfp stpB::stpB-mcherry</i>	Transformation of SS160 with pSS205	This work
SS389	CB15N <i>stpD::stpD-gfp stpC::stpC-mcherry</i>	Transformation of SS248 with pSS310	This work
SS412	CB15N <i>stpB::stpB-mcherry</i>	Transformation of CB15N with pSS309	This work
SS413	CB15N <i>stpC::stpC-mcherry</i>	Transformation of CB15N with pSS310	This work
SS414	CB15N Δ <i>stpA stpB::stpB-mcherry</i>	Transformation of SW49 with pSS309	This work
SS415	CB15N Δ <i>stpAB stpC::stpC-mcherry</i>	Transformation of SW51 with pSS310	This work
SW30	CB15N P _{xyl} ::P _{xyl} - <i>stpB-mcherry</i>	Transformation of CB15N with pSW32	This work
SW33	CB15N P _{xyl} ::P _{xyl} - <i>stpA-mcherry</i>	Transformation of CB15N with pSW35	This work
SW49	CB15N Δ <i>stpA</i>	Gene replacement in CB15N using pSW51	This work
SW50	CB15N Δ <i>stpB</i>	Gene replacement in CB15N using pSW52	This work
SW51	CB15N Δ <i>stpAB</i>	Gene replacement in CB15N using pSW53	This work
YB1585	CB15N <i>ftsZ::P_{xyl}-ftsZ</i>		Wang et al., 2001
YB2811	NY111d1 <i>pstS::miniTn5LacZ</i>		Ireland et al., 2002
YB3202	CB15N <i>divJ::Ω</i>		Pierce et al., 2006
YB5058	CB15N <i>stpX::StpX-gfp</i> P _{xyl} ::P _{xyl} - <i>stpB-mCherry</i>	Transduction of spec ^R from PV5064	This work
YB5059	CB15N Δ <i>stpAB stpX::stpX-gfp</i>	Transduction of spec ^R from PV5064	This work
YB5231	CB15N Δ <i>stpX</i>		Hughes et al., 2010
A. excentricus			
CB48	Wild-type strain		Poindexter, 1964
SS309	CB48 <i>Astex_0987::Astex_0987-mcherry</i>	Transformation of CB48 with pSS229	This work
E. coli			
TOP10	General cloning strain		Invitrogen

Table S2. General Plasmids Used in This Study, Related to Experimental Procedures

Name	Description	Reference/Source
pBXMCS-2	Replicating plasmid for the inducible overexpression of genes in CB15N, Kan ^R	Thanbichler et al., 2007
pCHYC-1	Plasmid for integrating genes encoding C-terminal fusions to the red fluorescent protein mCherry at the native gene locus, Spec/Str ^R	Thanbichler et al., 2007
pEJ216	Replicating plasmid carrying TAT-dimer2 under control of P _{xyl} , Cam ^R	Judd et al., 2005
pJAMY31	Kan ^R ColE1 replicon, β -lactamase translational fusion vector	Alley, M. R. K. (unpubl.)
pNPTS138	<i>sacB</i> -containing suicide vector used for double homologous recombination, Kan ^R	Alley, M. R. K. (unpubl.)
pTCYC-2	Plasmid for integrating C-terminal fusions to the tetracycline tag at the native gene locus, Spec/Str ^R	Thanbichler et al., 2007
pVCHYC-1	Plasmid for integrating genes encoding C-terminal fusions to the red fluorescent protein mCherry at the chromosomal <i>vanA</i> locus, Spec/Str ^R	Thanbichler et al., 2007
pXCHYC-2	Plasmid for integrating genes encoding C-terminal fusions to the red fluorescent protein mCherry at the chromosomal <i>xylX</i> locus, Kan ^R	Thanbichler et al., 2007
pXCHYC-5	Plasmid for integrating genes encoding C-terminal fusions to the red fluorescent protein mCherry at the chromosomal <i>xylX</i> locus, Tet ^R	Thanbichler et al., 2007
pXCHYN-5	Plasmid for integrating genes encoding N-terminal fusions to the red fluorescent protein mCherry at the chromosomal <i>xylX</i> locus, Tet ^R	Thanbichler et al., 2007
pXGFPC-2	Plasmid for integrating genes encoding C-terminal fusions to the green fluorescent protein GFP at the chromosomal <i>xylX</i> locus, Kan ^R	Thanbichler et al., 2007
pXYFPC-5	Plasmid for integrating genes encoding C-terminal fusions to the yellow fluorescent protein eYFP at the chromosomal <i>xylX</i> locus, Tet ^R	Thanbichler et al., 2007
pXYFPN-5	Plasmid for integrating genes encoding N-terminal fusions to the yellow fluorescent protein eYFP at the chromosomal <i>xylX</i> locus, Tet ^R	Thanbichler et al., 2007

Table S3. Plasmids Generated in This Study, Related to Experimental Procedures

Name	Description	Construction
pEK301	pXCHYC-2 carrying <i>pstS</i>	a) amplification of the <i>pstS</i> gene using primers EK225/EK226, followed by restriction with NdeI and EcoRI b) ligation of the digested PCR product into pXCHYC-2 cut with NdeI and EcoRI
pJK86	pXCHYC-2 carrying <i>gspG</i> (CCNA_00175)	a) amplification of <i>gspS</i> from genomic DNA using primers CC0176-NdeI-for/CC0176-EcoRI-rev, followed by restriction with NdeI and EcoRI b) ligation of the digested PCR product into pXCHYC-2 cut with NdeI and EcoRI
pJK101	pXCHYC-2 carrying <i>pstS</i>	a) amplification of <i>pstS</i> from genomic DNA using primers pstS-for/pstS-rev-EcoRI, followed by restriction with NdeI and EcoRI b) ligation of the digested PCR product into pXCHYC-2 cut with NdeI and EcoRI
pSS98	pSW32 carrying <i>stpB</i> downstream region	a) amplification of the <i>stpB</i> downstream sequence from genomic DNA using primers CC2476-1/-2, followed by restriction with BsrGI and NheI b) ligation into pSW32 cut with BsrGI and NheI
pSS109	pNTPS138-based plasmid for replacing native <i>stpB</i> with <i>stpB-mcherry</i>	a) isolation of <i>stpB-mcherry</i> from pSS98 by restriction with NdeI and NheI, followed by blunting of the fragment with T4 DNA polymerase b) ligation into pNTPS138 cut with EcoRV
pSS119	pXBlaMC-2 carrying <i>stpB</i>	a) isolation of <i>stpB</i> from pSW32 by restriction with NdeI and EcoRI b) ligation into pXBlaMC-2 cut with NdeI and EcoRI
pSS120	pXBlaMC-2 carrying <i>stpA</i>	a) isolation of <i>stpA</i> from pSW35 by restriction with NdeI and EcoRI b) ligation into pXBlaMC-2 cut with NdeI and EcoRI
pSS123	pTCYC-2 carrying <i>stpB</i>	a) isolation of <i>stpB</i> from pSW32 by restriction with NdeI and EcoRI b) ligation into pTCYC-2 cut with NdeI and EcoRI
pSS142	pVCHYC-1 carrying <i>stpB</i>	a) isolation of <i>stpB</i> from pSW32 by restriction with NdeI and EcoRI b) ligation into pVCHYC-1 cut with NdeI and EcoRI
pSS187	pTCYC-2-based plasmid carrying <i>stpB-His</i>	a) in-vitro assembly of a phosphorylated His ₁₀ -linker from AM_299f/AM_300r b) ligation into pSS123 cut with EcoRI
pSS202	pXGFPC-2 carrying <i>stpD</i>	a) amplification of <i>stpD</i> from genomic DNA using primers CCNA_2271-for/-rev, followed by restriction with NdeI and SacI b) ligation of the digested PCR product into pXGFPC-2 cut with NdeI and SacI
pSS204	pXCHYC-2 carrying <i>stpC</i>	a) amplification of <i>stpC</i> from genomic DNA using primers CCNA_02560-for/-rev, followed by restriction with NdeI and SacI b) ligation of the digested PCR product into pXCHYC-2 cut with NdeI and SacI
pSS205	pGFPC-1 carrying <i>stpD</i>	a) isolation of <i>stpD</i> from pSS202 by restriction with NdeI and EcoRI b) ligation into pGFPC-1 cut with NdeI and EcoRI
pSS206	pTCYC-2-based plasmid carrying <i>stpD-His</i>	a) isolation of <i>stpD</i> from pSS202 by restriction with NdeI and EcoRI b) ligation into pSS187 cut with NdeI and EcoRI
pSS208	pNTPS138-based plasmid for constructing an in-frame deletion in <i>stpD</i>	a) amplification of the <i>stpD</i> 5' region from genomic DNA using primers CCNA_02271-3/-4 b) amplification of the <i>stpD</i> 3' region from genomic DNA using primers CCNA_02271-5/-6 c) overlap extension PCR using both PCR products and primers CCNA_02271-3/-6, followed by restriction with HindIII and EcoRI c) ligation of the final PCR fragment into HindIII/EcoRI-cut pNTPS138
pSS209	pNTPS138-based plasmid for constructing an in-frame deletion in <i>stpC</i>	a) amplification of the <i>stpC</i> 5' region from genomic DNA using primers CCNA_02560-3/-4 b) amplification of the <i>stpC</i> 3' region from genomic DNA using primers CCNA_02560-5/-6 c) overlap extension PCR using both PCR products and primers CCNA_02560-3/-6, followed by restriction with HindIII and EcoRI c) ligation of the final PCR fragment into HindIII/EcoRI-cut pNTPS138
pSS210	pTCYC-2-based plasmid carrying <i>stpC-His</i>	a) isolation of <i>stpC</i> from pSS204 by restriction with NdeI and EcoRI b) ligation into pSS187 cut with NdeI and EcoRI
pSS220	pXBlaMC-2 carrying <i>stpC</i>	a) isolation of <i>stpC</i> from pSS204 by restriction with NdeI and EcoRI b) ligation into pXBlaMC-2 cut with NdeI and EcoRI
pSS221	pXBlaMC-2 carrying <i>stpD</i>	a) isolation of <i>stpD</i> from pSS202 by restriction with NdeI and EcoRI b) ligation into pXBlaMC-2 cut with NdeI and EcoRI
pSS227	pXCHYC-2 carrying <i>malA</i> (CCNA_02370)	a) amplification of <i>malA</i> from genomic DNA using primers CC2287-1/-2, followed by restriction with NdeI and SacI b) ligation of the digested PCR product into pXCHYC-2 cut with NdeI and SacI
pSS229	pCHYC-2 carrying <i>Astex_0987</i>	a) amplification of <i>Astex_0987</i> from genomic DNA using primers Astex_0987-2f/-3r, followed by restriction with NdeI and EcoRI b) ligation of the digested PCR product into pCHYC-2 cut with NdeI and EcoRI
pSS309	pCHYC-2 carrying <i>stpB</i>	a) isolation of <i>stpB</i> from pSW32 by restriction with NdeI and EcoRI b) ligation into pXCHYC-2 cut with NdeI and EcoRI
pSS310	pCHYC-2 carrying <i>stpC</i>	a) isolation of <i>stpC</i> from pSS204 by restriction with NdeI and EcoRI b) ligation into pXCHYC-2 cut with NdeI and EcoRI

Table S3. Plasmids Generated in This Study (continued)

Name	Description	Construction
pSW32	pXCHYC-2 carrying <i>stpB</i>	a) amplification of the <i>stpB</i> gene using primers CC2476-for/CC2476-rev, followed by restriction with NdeI and EcoRI
pSW35	pXCHYC-2 carrying <i>stpA</i>	b) ligation of the digested PCR products into pXCHYC-2 cut with NdeI and EcoRI a) amplification of the <i>stpA</i> gene using primers CC2477-uni/CC2477-rev, followed by restriction with NdeI and SacI
pSW51	pNTPS138-based plasmid for constructing an in-frame deletion in <i>stpA</i>	b) ligation of the digested PCR products into pXCHYC-2 cut with NdeI and SacRI a) amplification of the 5' region of <i>stpA</i> from genomic DNA using primers dCC2477-A-for/dCC2477-B-rev, followed by restriction with BamHI and EcoRI
pSW52	pNTPS138-based plasmid for constructing an in-frame deletion in <i>stpB</i>	b) amplification of the 3' region of <i>stpA</i> from genomic DNA using primers dCC2477-C-for/dCC2477-D-rev, followed by restriction with EcoRI and NheI c) ligation of both fragments into BamHI/NheI-cut pNTPS138
pSW53	pNTPS138-based plasmid for constructing an in-frame deletion in <i>stpAB</i>	a) amplification of the 5' region of <i>stpB</i> from genomic DNA using primers dCC2476-A-for/dCC2476-B-rev, followed by restriction with BamHI and EcoRI b) amplification of the 3' region of <i>stpB</i> from genomic DNA using primers dCC2476-C-for/dCC2476-D-rev, followed by restriction with EcoRI and NheI c) ligation of both fragments into BamHI/NheI-cut pNTPS138
pSW64 (<i>P_{xyI}-stpAB</i>)	pBXMCS-2 carrying <i>stpAB</i>	a) amplification of the 5' region of <i>stpAB</i> from genomic DNA using primers d2476d2477-A-for/d2476d2477-B-rev, followed by restriction with BamHI and EcoRI b) amplification of the 3' region of <i>stpB</i> from genomic DNA using primers d2476d2477-C-for/ d2476d2477-D-rev, followed by restriction with EcoRI and NheI c) ligation of both fragments into BamHI/NheI-cut pNTPS138
pSW67	pXCHYC-2 carrying <i>elpS</i> (CCNA_00169)	a) amplification of <i>stpAB</i> from genomic DNA using primers CC2477-uni/CC2476-rev2St, followed by restriction with NdeI and EcoRI b) ligation of the digested PCR product into pBXMCS-2 cut with NdeI and EcoRI
pXBlaMC-2	Plasmid for integrating genes encoding C-terminal TEM-1 β -lactamase fusions at the chromosomal <i>xylX</i> locus, Kan ^R	a) amplification of <i>elpS</i> from genomic DNA using primers CC0170-C-for/CC0170-C-rev, followed by restriction with NdeI and EcoRI b) ligation of the digested PCR product into pXCHYC-2 cut with NdeI and EcoRI a) amplification of the TEM-1 β -lactamase gene from pJAMY31 using primers blaM-C_for/blaM-C_rev, followed by restriction with AgeI and NheI b) ligation of the digested PCR product into pXGFPC-2 cut with AgeI and NheI

Table S4. Oligonucleotides Used in This Study, Related to Experimental Procedures

Note the different ORF numbers: CC2477 = CCNA_02562, CC2476 = CCNA_02561, CC0170 = CCNA_00169, CC0176 = CCNA_00175, CC2287=CCNA_02370.

Oligonucleotide	Sequence
AM_299f	AATCCCATCACCACCATCATCACCATCACCACCCTAGT
AM_300r	GGGTAGTGGTGGTAGTAGTGGTAGTGGTGGTGGTATCATTAA
Astex_0987-2f	ATTAATTCATATGTCGGCCAGACAGAAACCCGCTATG
Astex_0987-3r	ATGAATTCGATTCGAGGCGACCCAGACGACGCGG
blaM-C_for	TTTTACCGGTCGGCCACCATGCACCCAGAAACGCTGGTAAAAGTAAAAG
blaM-C_rev	TTTTGCTAGCTTACCAATGCTTAATCAGTGAGGCACCTATC
CCNA_02271-for	TTAATTCATATGCGTCATCAAATGGCGCGTCGCG
CCNA_02271-rev	TAGAGCTCCGTGATGGGCGGCGGCGGCTGCTTG
CCNA_02271-3	ATGAATTCGAACCAGACGACCTGAAGCGGCGCAG
CCNA_02271-4	CTTGTCCTTCACGCGACGCGCCATTTGATGAC
CCNA_02271-5	CGTCGCGTGAAGGACAAGCACGCCGCCGCCGC
CCNA_02271-6	TAAAGCTTCGGCGGTTTCCAGGTGATCGAGCA
CCNA_02560-for	TTAATTCATATGAGCAAGTCTGTTTCGTAGCCGGCTGG
CCNA_02560-rev	TAGAGCTCCGCATCCGACGAGGCCCGCGCCGACG
CCNA_02560-3	ATGAATTCGAGTCAAGGCGACCGGCACGATCATG
CCNA_02560-4	GAAATTACGGGAAACGGCCAGCCGGCTACGAAC
CCNA_02560-5	GCCGTTTCCCGTAATTTGTCGCGCGGGCCCTC
CCNA_02560-6	TAAAGCTTCTACGAGCAGGCGACGAAGCACCCG
CC0170-C-for	AATTCATATGAAGCTGTATAGAAACCTAATCCTCATGAGCTGCG
CC0170-C-rev	AATTGAATTCGTTGGTGGCGCAGCGAGCTG
CC0176-Ndel-for	ATATCATATGTCGACCGCAAACGCTGAAACGAAAC
CC0176-EcoRI-rev	ATATGAATTCGCTCCAGTTGCCAATGTCGG
CC2287-3	GCCAATGATCGCGTGATCGGGC
CC2287-4	CATCCCGCAGGAAGCCATCATCG
CC2476-for	TTAACATATGCGCCGCTCAGCCGCTGCTGG
CC2476-rev	TAGAATTCGATCGAGGAGCTCCCCCTTGTGAGGC
CC2477-uni	AAAACATATGCGCGAGGCGGGGACGCAATTGC
CC2477-rev	TAGAGCTCCGTAATTCCTTCGTTATACGGACGCCCGC
DCC2476-A-for	AATTGGATCCTCGGCCGTCGGAACACC
DCC2476-B-rev	AATTGAATTCGGCCAAAGCGCCAGC
DCC2476-C-for	AATTGAATTCGGTCCGCGCC
DCC2476-D-rev	AATTGCTAGCTTGAAGCAGCGGTTGTCGCC
CC2476-rev2St	AATTGAATTCATCGAGGAGCTCCCCCTTGT
CC2476-1	TATGTACAAGTAAGCAAGTCTGTTCTAGCCGGCTGGC
CC2476-2	TATAGCTAGCCGACAGCACCGTCTCAGCATCC
DCC2477-B-rev	TTAAGAATTCGCGAAGGCGCGCAATT
DCC2477-C-for	TTAAGAATTCGATCGGGGCGGGCGT
DCC2477-D-for	TTAAGCTAGCACCGGGGCTGGATCTTG
D2476d4277-A-for	TTAAGGATCCGGAGCTGGCCAATACGGC
D2476d4277-B-rev	TTAAGAATTCGCGAAGGCGCGCAATTG
D2476d4277-C-for	TTAAGAATTCGTTCCCGCGCTCAACAAG
D2476d4277-D-for	TTAAGCTAGCGGGGTGAAGATGCCGAG
EK225	TACTCATATGAACAAGCTCATCGGGC
EK226	TACTGAATTCGCTTCTCGGCGCCGGCATCGG
pstS-for	ATATACATATGAACAAGCTCATCGGCGGGTCCGC
pstS-rev-EcoRI	ATATGAATTCGCTTCTCGGCGCCGGCATC

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