

**S11 Table. Bacterial strains and plasmids used in this study.**

Strain/Plasmid	Relevant Characteristics	Source or reference
<i>S. meliloti</i>		
Rm2011	Nx <sup>r</sup> , Sm <sup>r</sup> , <i>expR</i> <sup>+</sup> ; Sm resistant derivative of <i>S. meliloti</i> SU47. Referred to as wild type in this study.	[90]
2011 <i>ecpR1</i>	<i>ecpR1</i> deletion mutant	This work
2011Pσ70 <i>ecpR1</i>	2011 carrying changes in the <i>ecpR1</i> σ <sup>70</sup> -dependent promoter -10 region	This work
2011 <i>rpoN::Tn5</i>	2011 carrying a mini-Tn5 inserted in <i>rpoN</i> Km <sup>r</sup>	[91]
2011 <i>mcherry</i>	2011 labeled with <i>mCherry</i> by a single chromosomal integration of pKOSm	This work
2011 <i>egfp</i>	2011 labeled with <i>egfp</i> by a single chromosomal integration of pKOSe	This work
2011 <i>ecpR1 egfp</i>	2011 <i>ecpR1</i> labeled with <i>egfp</i> by a single chromosomal integration of pKOSe	This work
2011 <i>relA</i>	<i>relA</i> deletion mutant	[38]
Sm2B6015	Rm2011 <i>visN</i> mutant	[42]
<i>gcrA</i> -eGFP	2011 carrying the fusion protein GcrA-EGFP	[46]
2011 <i>gcrA</i> -P <sub>lac</sub> <i>gcrA</i>	<i>gcrA</i> deletion mutant carrying P <sub>lac</sub> <i>gcrA</i>	This work
2011 <i>hfq</i>	<i>hfq</i> insertion mutant, Gm <sup>r</sup>	[92]
Rm4011	2011 derivative strain, <i>expR/sinI</i> double mutant. Nx <sup>r</sup> , Sm <sup>r</sup>	[81]
4011 <i>ecpR1</i>	<i>ecpR1</i> deletion mutant	This work
4011 <i>ecpR1 rne675</i>	2011 carrying pK18mobII inserted in the 675th codon of <i>rne</i> , Km <sup>r</sup>	[59]
<i>S. medicae</i> WSM419	Acid <sup>t</sup> , sardinian isolate; Nx <sup>r</sup> , Cm <sup>r</sup>	[93]
<i>S. fredii</i> NGR234	Broad-host-range bacterium isolated from <i>Lablab purpureus</i> , Rf <sup>r</sup>	[94]
<i>A. tumefaciens</i> C58	Wild-type, isolated from a cherry tree ( <i>Prunus</i> ) tumor Nx <sup>r</sup>	[95]
<i>A. radiobacter</i> ATCC19358 <sup>T</sup>	<i>A. radiobacter</i> type strain, isolated from soil. Nx <sup>r</sup>	[96]
<i>R. etli</i> CFN42 <sup>T</sup>	<i>R. etli</i> type strain, isolated from <i>Phaseolus</i> Nx <sup>r</sup> , Sm <sup>r</sup>	[97]
<i>R. tropici</i> CIAT899 <sup>T</sup>	<i>R. tropici</i> type strain, isolated from <i>P. vulgaris</i> . Sm <sup>r</sup>	[98]
<i>E. coli</i>		
DH5α	F- <i>endA1 supE44 thi-11-recA1 gyrA96 relA1 deoRD(lacZYA-argF)U169</i>	[99]
S17-1	<i>E. coli</i> 294 Thi RP4-2-Tc::Mu-Km::Tn7 integrated into the chromosome	[100]
Plasmids		
pPHUtrap	pPHU231 containing <i>sinI</i> 5'UTR fused to <i>egfp</i> , Tc <sup>r</sup>	Matthew McIntosh
p <i>PecpR1</i> _5'1-204	pPHUtrap with <i>ecpR1</i> promoter- <i>egfp</i> transcriptional fusion of the whole IGR	This work
p <i>PecpR1</i> _5'1	pPHUtrap with <i>ecpR1</i> promoter- <i>egfp</i> transcriptional fusion until 1 <sup>st</sup> 5'-end	This work
p <i>PecpR1</i> _5'2	pPHUtrap with <i>ecpR1</i> promoter- <i>egfp</i> transcriptional fusion until 2 <sup>nd</sup> 5'-end	This work
p <i>PecpR1</i> _5'2-Pσ70	pTSS2- <i>egfp</i> carrying changes in the σ <sup>70</sup> -dependent promoter -10 region	This work
pK18 <i>mobsacB</i>	Suicide plasmid in <i>S. meliloti</i> , <i>sacB</i> , <i>oriV</i> , Km <sup>r</sup>	[80]
pK <i>delecpR1</i>	pK18 <i>mobsacB</i> with <i>ecpR1</i> flanking regions for geneSOEing	This work
pKPσ70 <i>ecpR1</i>	pK18 <i>mobsacB</i> with <i>ecpR1</i> locus region carrying changes in the σ <sup>70</sup> -dependent promoter -10 region for double recombination	This work
pK <i>delgcrA</i>	pK18 <i>mobsacB</i> with <i>gcrA</i> flanking regions for geneSOEing	This work
pK18mobII	Suicide vector; mob lacZ Km <sup>r</sup>	[101]
pKOSm	pK18mobII with P <sub>T5</sub> : <i>mCherry</i> cassette fused to <i>recG</i>	Oliver Schauer
pKOSe	pK18mobII with P <sub>T5</sub> : <i>egfp</i> cassette fused to <i>recG</i>	Pornsri Charoenpanich
pK18ins <i>gcrA</i> <sub>300</sub>	pK18mobII carrying an internal fragment of <i>gcrA</i> , nt 4–297; Km <sup>r</sup>	This work
pSRKKm	pBBR1MCS-2 derivative with a Plac promoter, <i>lacIq</i> , <i>lacZa</i> <sup>+</sup> , Km <sup>r</sup>	[102]
pRel <sub>Sm</sub>	pSRKKm carrying the Rel <sub>Sm</sub> coding sequence	[38]
pSRKKm <i>divK</i>	pSRKKm carrying the <i>divK</i> coding sequence	[32]
pSKControl <sup>+</sup>	pSRKKm carrying the <i>smel812</i> coding sequence fused to <i>sinR</i> -P <sub>sinI</sub>	This work
pSKEcpR1 <sup>+</sup>	pSRKKm carrying the <i>ecpR1</i> coding sequence starting from 5'1 fused to <i>sinR</i> -P <sub>sinI</sub>	This work
pSKEcpR1 <sub>5'2</sub> <sup>+</sup>	pSRKKm carrying the <i>ecpR1</i> coding sequence starting from 5'2 fused to <i>sinR</i> -P <sub>sinI</sub>	This work
pSKEcpR1-1 <sup>+</sup>	pSKEcpR1 <sup>+</sup> carrying 1 nt change in the first conserved loop	This work
pSKEcpR1-2 <sup>+</sup>	pSKEcpR1 <sup>+</sup> carrying 2 nt changes in the first conserved loop	This work

pSKEcpR1-3 <sup>+</sup>	pSKEcpR1 <sup>+</sup> carrying 3 nt changes in the first conserved loop	This work
pSRKGm	pBBR1MCS-2 derivative with a Plac promoter, <i>lacIq</i> , <i>lacZa</i> <sup>+</sup> , Gm <sup>r</sup>	[102]
P <sub>lac</sub> <i>gcrA</i>	pSRKGm carrying the <i>gcrA</i> <sub>Sm</sub> coding sequence	This work
pSGControl <sup>+</sup>	pSRKGm with <i>smel812</i> coding sequence fused to <i>sinR</i> -P <sub>sinI</sub>	Jan Philip Schlüter
pSGEcpR1 <sup>+</sup>	pSRKGm with the <i>ecpR1</i> coding sequence fused to <i>sinR</i> -P <sub>sinI</sub>	Lars-Ole Loehr
pWBT	pSRKGm carrying the T5 promoter sequence downstream the <i>lac</i> promoter	Matthew McIntosh
pWBT <i>pleD</i>	pWBT carrying the <i>pleD</i> <sub>Sm</sub> coding sequence	Simon Schäper
pR_EGFP	Reporter fusion plasmid for cloning of sRNA targets, Tc <sup>r</sup> , Ap <sup>r</sup>	[44]
<i>pgcrA</i> <sub>.122+3-</sub> <i>egfp</i>	pR_EGFP expressing the <i>gcrA</i> :: <i>egfp</i> translational fusion from TSS (-130)	This work
<i>pPgcrA</i> <sub>.122+3-</sub> <i>egfp</i>	pR_EGFP expressing the <i>gcrA</i> :: <i>egfp</i> translational fusion from its own promoter (-223, relative to the AUG)	This work
<i>pdnaA</i> <sub>.17+30-</sub> <i>egfp</i>	pR_EGFP expressing the <i>dnaA</i> :: <i>egfp</i> translational fusion from TSS1	This work
<i>pdnaA</i> <sub>.56+30-</sub> <i>egfp</i>	pR_EGFP expressing the <i>dnaA</i> :: <i>egfp</i> translational fusion from TSS2	This work
<i>pdnaA</i> <sub>.70+30-</sub> <i>egfp</i>	pR_EGFP expressing the <i>dnaA</i> :: <i>egfp</i> translational fusion from TSS3	This work
<i>pdnaA</i> <sub>.154+30-</sub> <i>egfp</i>	pR_EGFP expressing the <i>dnaA</i> :: <i>egfp</i> translational fusion from -154	This work
<i>pdnaA</i> <sub>.198+30-</sub> <i>egfp</i>	pR_EGFP expressing the full <i>dnaA</i> :: <i>egfp</i> translational fusion from putTSS4	This work
<i>pdnaA</i> <sub>.198+162-</sub> <i>egfp</i>	pR_EGFP expressing the full <i>dnaA</i> :: <i>egfp</i> translational fusion to +162	This work
<i>pctrA</i> <sub>.26+93-</sub> <i>egfp</i>	pR_EGFP expressing the <i>ctrA</i> :: <i>egfp</i> translational fusion from TSS1	This work
<i>pctrA</i> <sub>.69+93-</sub> <i>egfp</i>	pR_EGFP expressing the <i>ctrA</i> :: <i>egfp</i> translational fusion from TSS2	This work
<i>pminD</i> <sub>.105+3-</sub> <i>egfp</i>	pR_EGFP expressing the <i>minD</i> :: <i>egfp</i> translational fusion	This work
<i>ppleC</i> <sub>.153+132-</sub> <i>egfp</i>	pR_EGFP expressing the <i>pleC</i> :: <i>egfp</i> translational fusion	This work
<i>pftsZ</i> <sub>.99+108-</sub> <i>egfp</i>	pR_EGFP expressing the <i>ftsZ</i> :: <i>egfp</i> translational fusion	This work
<i>pdivJ</i> <sub>.112+126-</sub> <i>egfp</i>	pR_EGFP expressing the <i>divJ</i> :: <i>egfp</i> translational fusion	This work
<i>pSMc00888</i> <sub>.235+57-</sub> <i>egfp</i>	pR_EGFP expressing the <i>SMc00888</i> :: <i>egfp</i> translational fusion	This work
<i>pdivK</i> <sub>.85+45-</sub> <i>egfp</i>	pR_EGFP expressing the <i>divK</i> :: <i>egfp</i> translational	This work
<i>pgcrA</i> -BS- <i>egfp</i>	<i>pgcrA</i> <sub>.122+3-</sub> <i>egfp</i> carrying 2 nt compensatory changes in <i>ecpR1</i> interaction region	This work
<i>pgcrA</i> -ATT- <i>egfp</i>	<i>pgcrA</i> <sub>.122+3-</sub> <i>egfp</i> carrying 3 nt changes in <i>ecpR1</i> interaction region	This work
<i>pdnaA</i> <sub>.154+162-</sub> <i>egfp</i>	pR_EGFP expressing the <i>dnaA</i> :: <i>egfp</i> translational fusion from -154 to +162	This work
<i>pdnaA</i> <sub>.154+162-</sub> BS5- <i>egfp</i>	<i>pdnaA</i> <sub>.154+162-</sub> <i>egfp</i> carrying 5 nt changes in <i>ecpR1</i> predicted binding site 5	This work
<i>pdnaA</i> <sub>.56+30-</sub> BS4- <i>egfp</i>	<i>pdnaA</i> <sub>.56+30-</sub> <i>egfp</i> carrying 5 nt changes in <i>ecpR1</i> binding site 4	This work
<i>pdnaA</i> <sub>.56+30-</sub> BS3 <i>egfp</i>	<i>pdnaA</i> <sub>.56+30-</sub> <i>egfp</i> carrying 3 nt changes in <i>ecpR1</i> binding site 3	This work
<i>pdnaA</i> <sub>.56+30-</sub> BS3+4- <i>egfp</i>	<i>pdnaA</i> <sub>.56+30-</sub> <i>egfp</i> carrying 8 nt changes in <i>ecpR1</i> binding sites 3 and 4	This work
<i>pdnaA</i> <sub>.154+162-</sub> BS3+4- <i>egfp</i>	<i>pdnaA</i> <sub>.154+162-</sub> <i>egfp</i> carrying 8 nt changes in <i>ecpR1</i> binding sites 3 and 4	This work
<i>pdnaA</i> -BSs- <i>egfp</i>	<i>pdnaA</i> <sub>.56+30-</sub> <i>egfp</i> carrying 3 nt compensatory changes in <i>ecpR1</i> binding sites 3 and 4	This work

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