	Relevant characteristics	Source or reference
Strains		
M. extorquens		
AM1	Wild type	Lab strain
$\Delta phyR::kanR$	Deletion mutant of the <i>phyR</i> gene	(2)
$\Delta ecfGl$	Deletion mutant of the <i>ecfG1</i> gene using the vector pCM433-ecfG1	This study
E. coli		
DH5a	supE44 ∆lacU169 (ф80lacZDM15) hsdR17 recA1 endA1 gyrA96 thi-1 relA1	Invitroger
BL21(DE3)	$ompT hsdS_{B}(r_{B}m_{B}) gal dcm (DE3)$	Invitrogen
Plasmids		
pCM62	Tc^{R} ; <i>M. extorquens/E. coli</i> shuttle vector (P_{lac})	(1)
pCM62-phyR	Tc^{R} ; pCM62 derivative containing <i>phyR</i>	This study
pCM62-phyRD190A	Tc ^R ; pCM62 derivative containing <i>phyR</i> D190A	This study
pCM80	Tc ^R ; <i>M. extorquens/E. coli</i> shuttle vector for expression of genes under the control of the <i>mxaF</i> promoter (P_{lac} - P_{mxaF})	(1)
pCM80-phyR	Tc^{R} ; pCM80 derivative containing <i>phyR</i>	This study
pCM80-phyRNterm	Tc ^R ; pCM80 derivative containing the ECF sigma factor like domain- encoding part of $phyR$	This study
pCM80-nepR	Tc ^R ; pCM80 derivative containing <i>nepR</i>	This study
pCM80-nepR-his	Tc ^R ; pCM80 derivative for production of NepR with a C-terminal hexahistidine tag	This study
pCM433	Ap ^R , Cm ^R , Tc ^R ; broad-host-range <i>sacB</i> -based allelic exchange vector	
pCM433-ecfG1	Ap ^R , Cm ^R , Tc ^R ; pCM433 derivative used for the deletion of <i>ecfG1</i>	This study
pET16b	Amp ^R ; vector for production of recombinant protein in <i>E. coli</i> adding an N-terminal decahistidine tag	Novagen
pET16b-nepR	Amp ^R ; vector for production of NepR	This study
pET16b-ecfG1∆1-64	Amp ^R ; vector for production of $\sigma^{\text{EcfG1}}\Delta 1$ -64	This study
pET24b	Kan ^R ; vector for production of recombinant protein in <i>E. coli</i> adding a C-terminal hexahistidine tag	Novagen
pET24b-phyR	Kan ^R ; pET24b derivative for production of PhyR with a C-terminal hexahistidine tag	This study
pET101/D-TOPO	Amp ^R ; linearized, topoisomerase I-activated vector for production of recombinant protein with a C-terminal hexahistidine tag in <i>E. coli</i>	Invitrogen
pET101-phyRD190A	Amp ^R ; vector for production of the PhyR D190A mutant	This study
nFT101-nhvRNterm	Amp ^R , vector for production of the N-terminal ECE domain of PhyR	This study

Table S2. Bacterial strains an	l plasmids used in this study
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