

Supplemental materials of

Distinct nitrite and nitric oxide physiology in *Escherichia coli* and *Shewanella oneidensis*

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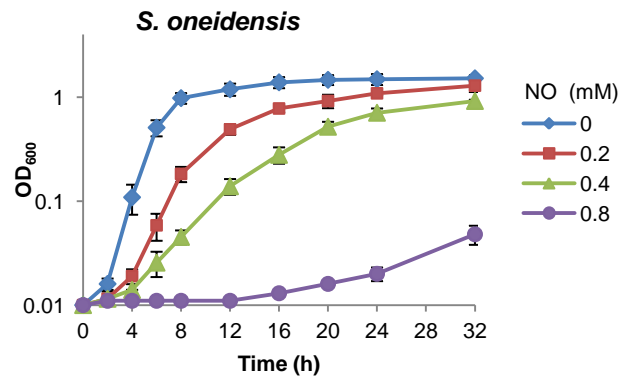
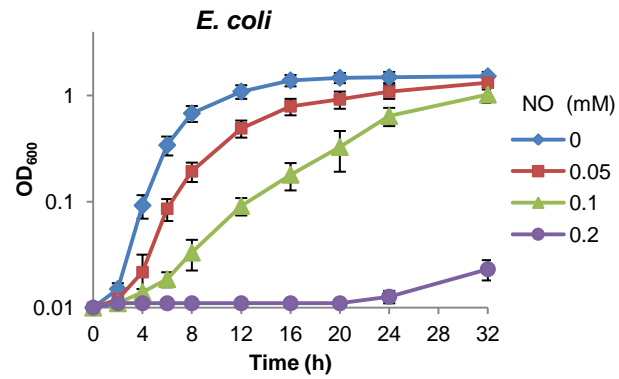
A**B**

FIG S1 Inhibition of NO released from DETA-NONOate at varying concentrations on growth in liquid media. **(A)** On *S. oneidensis*. **(B)** On *E. coli*. Data are shown as mean \pm SEM from at least three experiments.

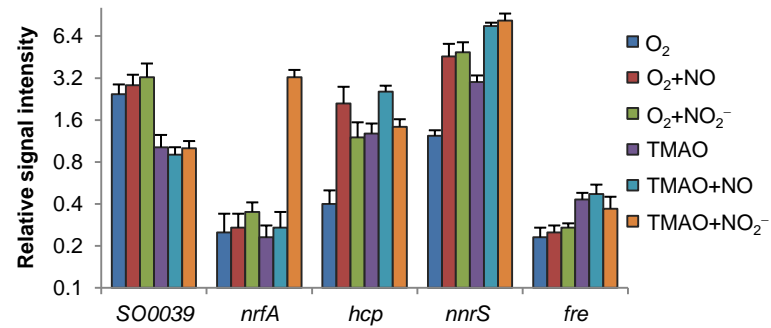


FIG S2 qRT-PCR analysis of RNA extracted from mid-log growing cells. Cells were prepared the same as in Fig. 3A. All data were normalized to expression of the *arcA* rRNA gene, which was constant during the exponential growth phase. Numbers reported are standardized to expression of the reference gene. Data are shown as mean \pm SEM from at least three experiments.

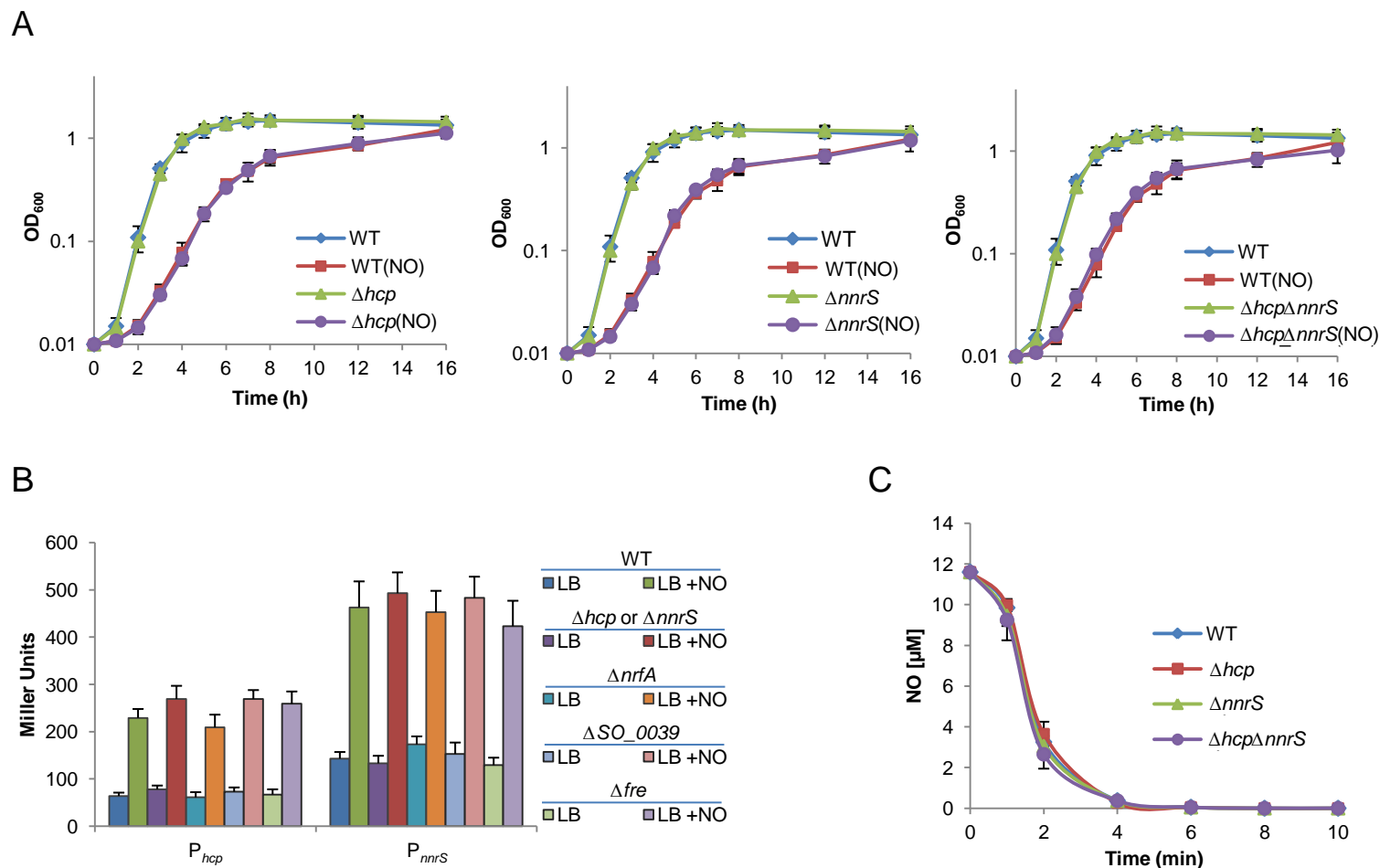
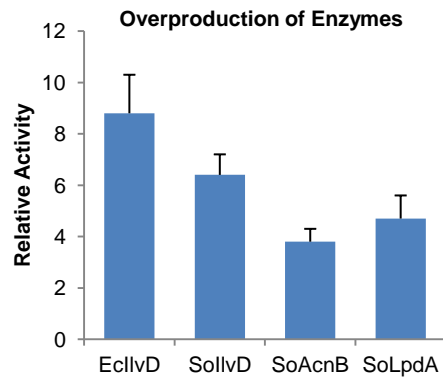


FIG S3 Impacts of Hcp, NnrS, SO_0039, and Fre on NO resistance and removal. (A) Effects of indicated proteins on NO-induced growth inhibition. Growth of WT, Δhcp , ΔSO_0039 , $\Delta nnrS$, Δfre , and $\Delta hcp\Delta nnrS$, in LB without or with 0.4 mM DETA-NONOate was compared. In all cases, difference was insignificant. Only Δhcp , $\Delta nnrS$, and $\Delta hcp\Delta nnrS$ were shown because these two genes are responsive to NO. (B) Promoter activity measurement of P_{nrfA} , P_{SO0039} , P_{hcp} , P_{nnrS} , and P_{fre} by an integrated *lacZ* reporter in strains lacking one of these genes. Mutation of a single gene did not significantly affect expression of the other genes. Only data for *hcp* and *nnrS* genes were shown. (C) Effects of indicated proteins on NO consumption. Data are shown as mean \pm SEM from at least three experiments.

A



B

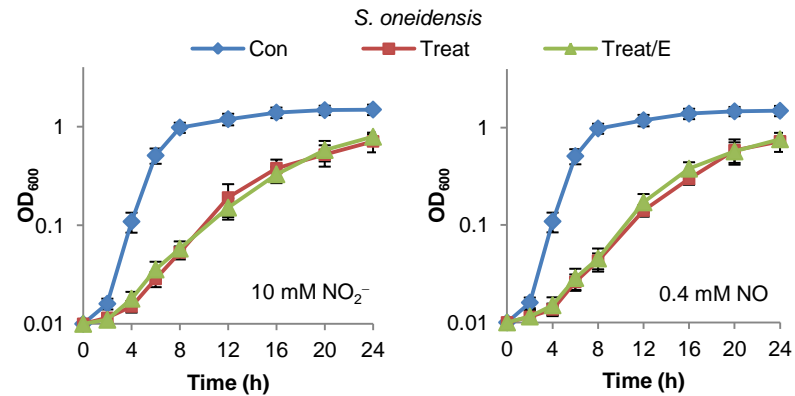


FIG S4 Validation of enzyme overproduction. Controlled expression of indicated genes was carried out by using IPTG-inducible promoter *Ptac*. The expression system is effective, validated previously and here by measuring respective enzyme activities. Relative activity is given by normalizing to the wild-type containing empty vector. (B) Effects of overproduction of *EcllVD* on growth of *S. oneidensis*. Con, grown in LB; Treat, grown in LB with either 0.4 mM NO or 10 mM nitrite; Treat/E, cells with overproduced *SollVD* in the presence of 1 mM IPTG. Data are shown as mean \pm SEM from at least three experiments.