

Table S1. Bacterial strains and plasmids used in this study.

Strain or plasmid	Genotype and/or relevant characteristics	Reference or source
<i>P. aeruginosa</i>		
PAO1 (ATCC15692)	Prototroph	American type culture collection
$\Delta lpxO1$	PAO1 derivative with an in-frame deletion in the <i>lpxO1</i> (PA4512) coding sequence	This work
$\Delta lpxO2$	PAO1 derivative with an in-frame deletion in the <i>lpxO2</i> (PA0936) coding sequence	This work
$\Delta lpxO1\Delta lpxO2$	$\Delta lpxO1$ derivative with an in-frame deletion in the <i>lpxO2</i> coding sequence	This work
<i>E. coli</i>		
S17.1 λ pir	<i>thi pro hsdRhsdM⁺ recA RP4-2-Tc::Mu-Km::Tn7 λpir, Sm^R</i>	Simon <i>et al.</i> 1983
Plasmid		
pBluescript II (pBS)	Sequencing vector; ColE1 replicon; Ap ^R	Stratagene
pBS <i>lpxO1</i> ↑ <i>lpxO1</i> ↓	pBS derivative containing the regions upstream and downstream of the <i>lpxO1</i> coding sequence	This work
pBS <i>lpxO2</i> ↑ <i>lpxO2</i> ↓	pBS derivative containing the regions upstream and downstream of the <i>lpxO1</i> coding sequence	This work
pDM4	Suicide vector; <i>sacBR</i> , <i>oriR6K</i> ; Cm ^R	Milton <i>et al.</i> 1996
pDM4 $\Delta lpxO1$	pDM4 derivative for <i>lpxO1</i> in-frame deletion, generated by sub-cloning the <i>lpxO1</i> ↑ <i>lpxO1</i> ↓ fragment of pBS <i>lpxO1</i> ↑ <i>lpxO1</i> ↓ into pDM4, Cm ^R	This work
pDM4 $\Delta lpxO2$	pDM4 derivative for <i>lpxO2</i> in-frame deletion, generated by sub-cloning the <i>lpxO2</i> ↑ <i>lpxO2</i> ↓ fragment of pBS <i>lpxO2</i> ↑ <i>lpxO2</i> ↓ into pDM4, Cm ^R	This work
pME6032	Shuttle vector for IPTG inducible expression in <i>P. aeruginosa</i> , Tc ^R	Heeb <i>et al.</i> 2002
pME <i>lpxO1</i>	pME6032 derivative carrying the <i>lpxO1</i> coding sequence under the control of the IPTG-inducible promoter	This work
pME <i>lpxO2</i>	pME6032 derivative carrying the <i>lpxO2</i> coding sequence under the control of the IPTG-inducible promoter	This work

Reference non included in the main text:

Simon, R.; Priefer, U; Pühler, A. A broad host range mobilization system for *in vivo* genetic engineering: transposon mutagenesis in Gram negative bacteria. *Bio/Technology*. **1983**, *1*, 784-790.