Supplementary Figure Legends

Fig. S1. Effect of *hns* and *cyaA* mutations on novobiocin and erythromycin resistance phenotype of the parental ($\Delta acrABE$) strain. LBA, lysogenic broth agar; LBA+NE, lysogenic broth agar plus novobiocin (1.25 or 2.5 µg/ml) and erythromycin (1.25 or 2.5 µg/ml). Plates were incubated for 24h at 37°C. Relevant genotypes are shown.

Fig S2. Effect of an in-frame deletion of *mdtA* on novobiocin resistance of a $\Delta acrABE$ strain containing either the *baeS51* or *rpoB58* mutation. LBA, lysogenic broth agar; LBA+N, lysogenic broth agar plus novobiocin (3.0 µg/ml). Plates were incubated for 24h at 37°C. Relevant genotypes are shown.



LBA+NE



LBA+NE







- 1. $\Delta acrABE$
- 2. ∆acrABE
- 3. *ΔacrABE ΔcyaA*
- 4. ΔacrABE ΔcyaA

- 1. ∆acrABE
- 2. ΔacrABE Δhns
- ΔacrABE Δhns
- 4. $\triangle acrABE \triangle hns$



LBA+Nov





- 1. baeS51
- 2. baeS51 ∆mdtA
- 3. rpoB58
- 4. rpoB58 ∆mdtA

Strain genotype	Novobiocin Minimal Inhibitory Concentration (µg/ml)
ΔacrABE	≤1.0
ΔacrABE ΔrpoS	≤1.0
∆acrABE ∆dksA	≤1.0
∆acrABE rpoB58	4.0
ΔacrABE rpoB58 ΔrpoS	4.0
∆acrABE rpoB58 ∆dksA	≤1.0
WT	64
ΔrpoS	64
ΔdksA	32

Table S3. Effect of *rpoS* or *dksA* deletion on minimal inhibitory concentration of novobiocin

MIC was conducted by a two-fold serial dilution method, as described in the main text