

Supporting material

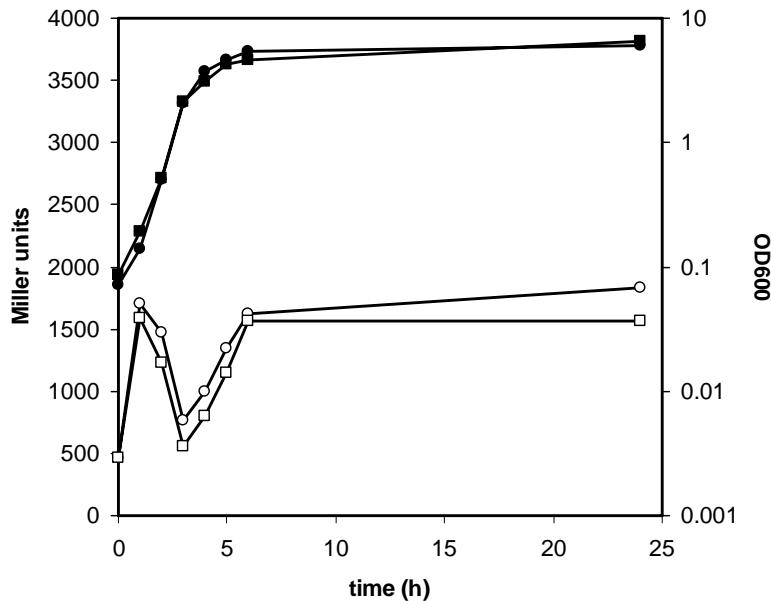


Figure S1. Growth (closed symbols) and β-galactosidase activity (open symbols) of KT2440 (circles) and the *rpoS* mutant C1R1 (squares) harboring the *lapA::lacZ* translational fusion in pMMGA.

<i>lapA</i>	GATGGCATGTTATAAGGTCAATAGTTGGCA-G--TCAGGCAATTCCAAAAAGTTATAG
<i>flhA</i>	TTTCAATCAATGAATTGAAAAGTT-GGAACGGTTCTGCAAT--GACCCGCCGAC
<i>lapA</i>	ACCGAATATT GACGTCAAAACGTCAAGAGATCATCGACATAGTTCCGCCTGAAGT-G
<i>flhA</i>	GC-GCATTCCGG CGTCAAAAGTTGTTGTCGGCCACCCACGGG-CGCCATACGCAG
<i>lapA</i>	GCTAGCAAGCGCGCTCTGGCAGGGAAGTACCCCATCGGGTCACACGGAGAGTCCAATG
<i>flhA</i>	GGCAGGGAGTCGAGAGTC

Figure S2. Alignment between the promoter regions of *lapA* of *P. putida* and *flhA* of *P. aeruginosa*. Shown in bold is the sequence corresponding to the FleQ box (obtained from the PRODORIC database: <http://prodoric.tu-bs.de/>) and the corresponding putative site in the *lapA* promoter region. Translation start codons are shown in blue.

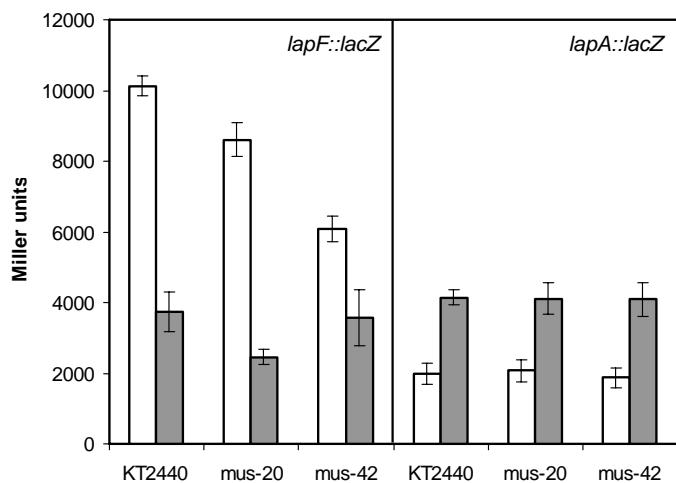


Figure S3. Analysis of β -galactosidase activity of the *lapF::lacZ* and *lapA::lacZ* fusions in KT2440, mus-20 (*lapF* mutant) and mus-42 (*lapA* mutant) harboring (grey bars) or not (white bars) plasmid pMAMV1, which confers high levels of c-di-GMP. Results are averages and standard deviations from three experiments with duplicate samples, in cultures after overnight growth in LB.