

***Pseudomonas fluorescens*: identification of Fur-regulated proteins and evaluation of their contribution to pathogenesis**

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Table S1. Primers used for qRT-PCR.

Primer	Sequences (5'→3')	Gene name
AcaRT-F	ACCAAGAGCGAAGCGATTGC	Acetyl-CoA carboxylase subunit alpha
AcaRT-R	CCAGAACCAACCTTCACCGATC	
AdRT-F	CACAACCTGCTCACCGACATC	Arginine deiminase
AdRT-R	CAGCCACGAACCGACTTCATC	
AtRT-F	TGGTGGCGGGCGATGAC	ABC transporter
AtRT-R	GATGGTGTGGAGGAGCAGAAG	
ClpPRT-F	CAAGGAGCGAGTGATCTTCTGG	ATP-dependent Clp protease proteolytic subunit
ClpPRT-R	GGCGAGTTGATGTAGAGATGGATG	
DipRT-F	CGGTGACGACTATGCCAAGC	Dipeptidase
DipRT-R	CGGAGGAGTGAGCGGATACG	
FlaRT-F	CGCCAACGCCAACGAGAC	Flagellin
FlaRT-R	GTAGCAGCCAAGCCTGTCATC	
GbgRT-F	AGCCGTCCAAGGTCTCAAC	Glucan biosynthesis protein G
GbgRT-R	CCAGATCCACTCGCCATTGC	
GsRT-F	CTGGTGAAGAAGGCTGACGAAG	Glutamine synthetase
GsRT-R	GCGGCTTAGGCATGAAGGTAG	
HpRT-F	CCTGCTACTCAACTGCGACATC	Hypothetical protein
HpRT-R	GCAATCAATGAACGGCATGACTTC	
IcdRT-F	CAAGCACCCGCACAAGATGG	Isocitrate dehydrogenase
IcdRT-R	CCGTAGAAGTCGCCGTTGTT	
IclRT-F	AAGTGGCAGCGGACAACAAAC	Isocitrate lyase
IclRT-R	GGCACGACGGAACGAGTTG	
KarRT-F	CGTTCTGTGCGGCCGTAC	Ketol-acid reductoisomerase
KarRT-R	TCAGTTCGTGCAGGCATTG	
L25RT-F	CGTCACCCTTCCAAGCAGTTC	50S ribosomal protein L25
L25RT-R	AGGCACGATGGCGGTCA	
NupRT-F	AAGTGGACAACCTGCTCAAGTATG	Purine nucleoside permease, NUP family/hypothetical protein
NupRT-R	ACGCCGACCAGTTCTAACAAATC	
OprERT-F	TACCTGTACGGCGGCTACC	Outer membrane porin OprE3
OprERT-R	GAAGTCGGACCAGTAGGAGTTG	
SodRT-F	GCCTTGCCTTACGCCCTACG	Superoxide dismutase
SodRT-R	CTGGTGGTGCTTGGTAGTG	
StpRT-F	AAGGCAGCAGCAACCAGTTG	Sugar ABC transporter substrate-binding protein/putative iron-binding periplasmic protein
StpRT-R	GGCGGTGATTCTCGGTGTAG	
TsspRT-F	GCAGCCGCCGAAGTGATC	EvpB family type VI secretion protein
TsspRT-R	CGATACCAAACGCAACGACAC	
UhRT-F	CGCCTGCTCGCTGAATATCG	Urocanatehydratase
UhRT-R	GCCTTGCCTCCTGGTGTAC	
XthRT-F	TGGATGGCACGCTTGAAAGAAC	Exodeoxyribonuclease III
XthRT-R	CCACGGCTGCGGTAATCG	

Figure S1. Growth profiles of *Pseudomonas fluorescens* TSS Δ *oprF*, TSS Δ *oprFC*, and TSS. *P. fluorescens* TSS Δ *oprF*, TSS Δ *oprFC*, and TSS were cultured in LB medium, and cell density was determined at various time points by measuring absorbance at OD₆₀₀. Data are the means of three independent assays and presented as means \pm SEM.

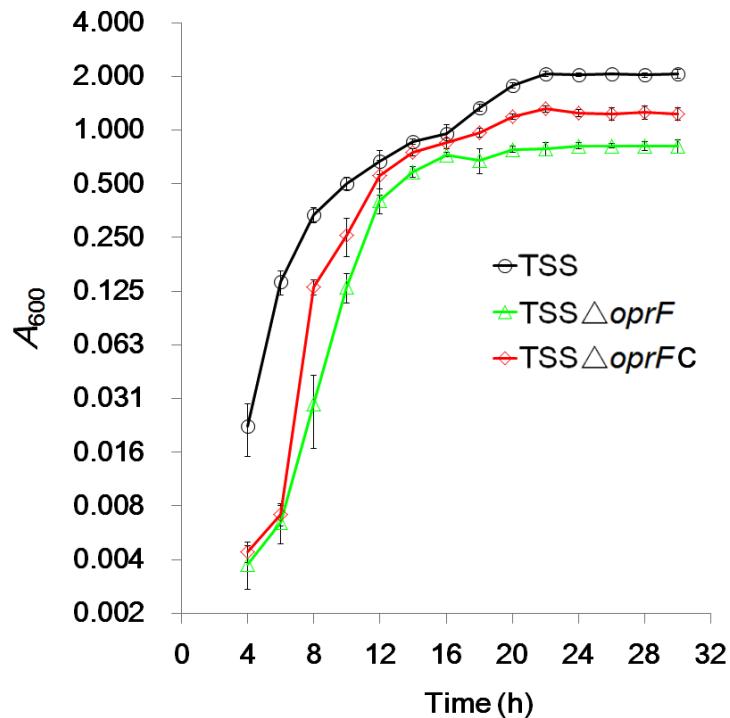


Figure S2. SDS-PAGE analysis of purified recombinant protein. Purified rOprF (lane 2) was analyzed by SDS-PAGE and viewed after staining with Coomassie brilliant blue R-250. Lane 1, protein markers.

