

# Heat Shock Protein DnaJ in *Pseudomonas aeruginosa* Affects Biofilm Formation via Pyocyanin Production

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

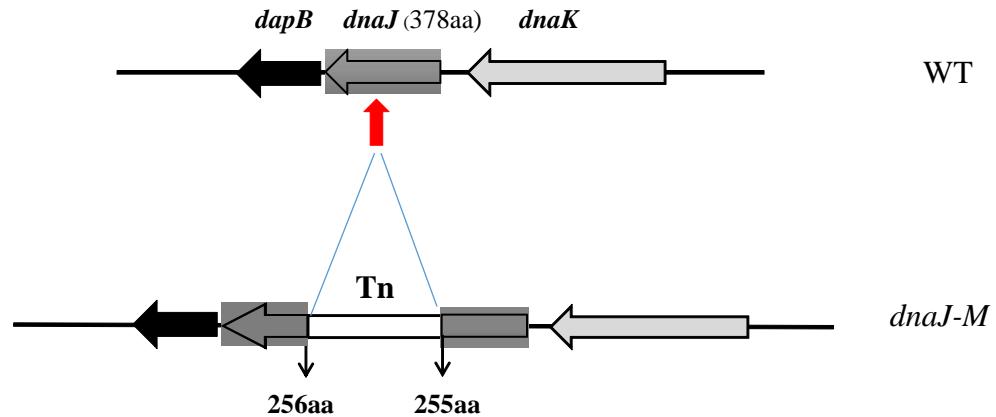
Stains	Relevant characteristic <sup>1</sup>	Source
<i>Pseudomonas aeruginosa</i> PAO1		[1]
DH10B	F- <i>mcrA</i> Δ( <i>mrr-hsdRMS-mcrBC</i> ) φ80lacZΔM15 Δ <i>lacX74</i> <i>recA1 endA1 araD139</i> Δ( <i>ara-leu</i> )7697 <i>galU galK λ-</i> <i>rpsL</i> (Str <sup>R</sup> ) <i>nupG</i>	Invitrogen
<i>danJ-M</i>	PA4760 transposon mutant of PAO1; Gm <sup>R</sup>	This study
<b>Plasmids</b>		
pMS402	Expression reporter plasmid carrying the promoterless <i>luxCDABE</i> gene; Kn <sup>R</sup> , Tmp <sup>R</sup>	[2]
pAK1900	<i>E. coli-P. aeruginosa</i> shuttle cloning vector carrying plac upstream of MCS; Amp <sup>R</sup> , Cb <sup>R</sup>	[3]
pKD- <i>phzM</i>	pMS402 containing <i>phzM</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[4]
pKD- <i>phzS</i>	pMS402 containing <i>phzS</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[4]
pKD- <i>phzA1</i>	pMS402 containing <i>phzA1</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[4]
pKD- <i>phzA2</i>	pMS402 containing <i>phzA2</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[4]
pKD- <i>pqsA</i>	pMS402 containing <i>pqsA</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[5]
pKD- <i>lasI</i>	pMS402 containing <i>lasI</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[6]
pKD- <i>lasR</i>	pMS402 containing <i>lasR</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[6]
pKD- <i>rhII</i>	pMS402 containing <i>rhII</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[6]
pKD- <i>rlIR</i>	pMS402 containing <i>rlIR</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	[6]
pKD- <i>flhF</i>	pMS402 containing <i>flhF</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	This study
pKD- <i>flhA</i>	pMS402 containing <i>flhA</i> promoter region; Kn <sup>R</sup> , Tmp <sup>R</sup>	This study
pAK- <i>dnaJ</i>	pAK1900 with a 1.1 kb fragment of <i>dnaJ</i> between XbaI and HindIII; Amp <sup>R</sup>	This study

<sup>1</sup> Tmp<sup>R</sup>, trimethoprim resistance; Gm<sup>R</sup>, gentamicin resistance; Kn<sup>R</sup>, kanamycin resistance; Amp<sup>R</sup>, ampicillin resistance; Cb<sup>R</sup>, carbencillin resistance.

**Table S2.** Primers used in this study.

Primer	Sequence (5'-3') <sup>1</sup>	Restriction site
PA4760 up	AATA <u>AAGCTT</u> GATGATCCGCCACGC	HindIII
PA4760 down	GG <u>CTCTAGA</u> CGGTCTGCTAACGGC	XbaI
ARB1	GGCCACGCGTCGACTACTACNNNNNNNNNGATAT	
P7-1	CTAACAA <u>TTCGTT</u> CAAGCCG	
ARB2	GGCCACGCGTCGACTAGTAC	
P7-2	GGATGCGTCTAAAGCCTGC	
<i>flhF</i> up	GC <u>ACTCGAGG</u> ACAAGGCTCCGAGG	XhoI
<i>flhF</i> down	AC <u>AGGATCCG</u> CAGCTCCACTCCAG	BamHI
<i>flhA</i> up	CGGT <u>CTCGAG</u> TAGTTGGTGATTGGCG	XhoI
<i>flhA</i> down	AAT <u>GGATCCG</u> ACCACGTAGTTGCCG	BamHI

<sup>1</sup> Restriction sites are underlined.



**Figure S1.** Schematic depiction of the genomic organization of *dnaJ*-*M*. Top: intact *dnaJ* gene on the chromosome in the wild type (WT). Bottom: *dnaJ* was disrupted by the transposon element (Tn). The insertion site is indicated with the DnaJ amino acid residues.

## References

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