Quorum Quenching: Enzymatic Disruption of N-Acylhomoserine Lactone-Mediated Bacterial Communication in *Burkholderia thailandensis*

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Many species of gram-negative bacteria communicate by synthesizing, secreting, and responding to N-acylhomoserine lactones (AHLs), a mechanism termed quorum sensing. Several investigations have characterized numerous AHL-degrading enzymes (AiiA lactonases) encoded by environmental isolates of *Bacillus* spp. The *Burkholderia thailandensis* quorum system is comprised of at least three AHL synthases (AHHs) and five transcriptional regulators belonging to the LuxIR class of proteins. Expression of the *Bacillus anthracis* (Ames strain) AiiA lactonase in *B. thailandensis* completely abolished the accumulation of N-decanoylhomoserine lactone (C₁₀-HSL) and N-octanoylhomoserine lactone (C₈-HSL), reduced N-hexanoylhomoserine lactone (C₆-HSL) levels, altered both swarming and twitching motility, caused a significant increase in generation time, and affected carbon metabolism. In contrast, heterologous expression of the *Bacillus cereus* strain A24 AiiA lactonase in *B. thailandensis* reduced the concentrations of C₈-HSL, C₆-HSL, and C₁₀-HSL to nondetectable levels; altered both swarming and twitching motility; and caused fluctuations in carbon utilization. Individual disruption of the *B. thailandensis* AHHs, specifically disruption of the btaI7 and btaI3 genes, which encode the proteins that direct the synthesis of C₈-HSL and C₆-HSL, respectively, caused the hyper-beta-hemolysis of sheep erythrocytes on blood agar plates. In contrast, AHL cleavage in *B. thailandensis* by the *Bacillus* AiiA lactonases failed to enhance beta-hemolytic activity. The results of this study demonstrate that heterologous expression of *Bacillus* sp. AiiA lactonases in *B. thailandensis* reduced AHL accumulation, affected both swarming and twitching motility, increased generation time, altered substrate utilization, and prevented the beta-hemolysis of sheep erythrocytes.

Many gram-negative bacteria utilize N-acylhomoserine lactones (AHLs) in a cell density-dependent manner, referred to as quorum sensing (QS), to control gene expression (14). When an adequate AHL concentration is reached, the signaling molecules are bound by LuxR proteins, which in turn either induces or represses gene expression. QS networks have been identified in dozens of bacterial species and have been shown to be involved in various physiological process (8, 12, 19, 36).

*Burkholderia thailandensis* is a gram-negative bacterium that is genetically and biochemically similar to *Burkholderia pseudomallei* (the causative agent of melioidosis) and *Burkholderia mallei* (the etiologic agent of glanders) (3, 16). Although the bacterium is genetically related to *B. pseudomallei*, *B. thailandensis* is an avirulent *Burkholderia* species that does not cause disease in humans. Comparative analysis of the *B. pseudomallei* and *B. thailandensis* 16S rRNAs revealed differences in 15 nucleotides, which justified the classification of a new *Burkholderia* species (3). The most striking difference between these organisms is their ability to cause acute disease in a Syrian hamster model. An intraperitoneal challenge of <1×1⁰⁵ CFU of *B. pseudomallei* causes mortality in hamsters within 3 to 5 days, in contrast to a dose of >1×1⁰⁶ CFU for *B. thailandensis* (3).

Several investigations have recently identified functional QS networks in *Burkholderia* species (2, 5, 17, 26, 27). In *Burkholderia cepacia*, QS negatively regulates ornibactin biosynthesis and positively induces protease and N-octanoylhomoserine lactone (C₈-HSL) production (26, 27). Recently, an additional pair of luxIR genes (cciIR) were identified in *B. cepacia* and shown to be involved in pathogenicity (2). *Burkholderia vietnamiensis* produces multiple AHL molecules, with the predominant AHL being N-decanoylhomoserine lactone (C₁₀-HSL) and with C₆-HSL and N-hexanoylhomoserine lactone (C₆-HSL) being produced to a lesser extent (5, 17). Our laboratory has recently identified a complex QS network encoded by *B. thailandensis* that is comprised of at least three AHL synthases and five putative transcriptional regulators homologous to LuxIR proteins, which were shown to affect swarming and twitching motility, lipase production, beta-hemolysis, and carbon metabolism (38).

Considering the diverse microbial species that use QS for the regulation of numerous physiological processes, applications targeted at disrupting this mode of intraspecies communication would be of significant interest to the medical, environmental, and agricultural communities. In the marine environment, *Delisea pulchra* (a microalga) produces brominated furanones that are hypothesized to compete with bacterial AHLs for the binding sites within putative LuxR proteins (29). In some culture environments, AHL signaling molecules can be utilized as sole carbon and energy sources by species of *Pseudomonas*, *Variovorax*, and *Ralstonia* (22, 24, 28). Xu et al. recently demonstrated that porcine kidney acylase I deacylates both N-butanoxyhomoserine lactone (C₄-HSL) and C₆-HSL.
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15. SUBJECT TERMS
Burkholderia thailandensis, lactone-mediated communication, characterization, Bacillus anthracis expression

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TABLE 1. Bacterial strains and plasmids used in this study

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MATERIALS AND METHODS

Bacterial strains and plasmids. All bacterial strains and plasmids used in this study are described in Table 1. B. thailandensis, Agrobacterium tumefaciens A136, and Escherichia coli were cultured in Luria-Bertani (LB) broth or on LB agar plates at 30 or 37°C, as required.

Cloning of B. anthracis aiiA. Genomic DNA for PCR amplification of the B. anthracis aiiA (Ames strain) aiiA lactonase gene was purified by using the Puregene yeast and gram-positive bacterium kit according to the instructions of the manufacturer (Gentra Systems, Minneapolis, Minn.). The B. anthracis aiiA gene was PCR amplified by using the primer set (forward, 5'-ATG GGA TCC B. anthracis aiiA C for 5 min; 30 cycles of 94°C for 30 s, 50°C for 30 s, and 72°C for 30 s; and a final 7-min extension at 72°C. All PCRs were performed with the FastSafe PCR system with buffer E (Epitrend Technologies, Madison, Wis.), and amplicons were cloned into pCR2.1-TOPO (Invitrogen, Carlsbad, Calif.). Ligation were transformed into One Shot chemically competent E. coli (Invitrogen) and screened on LB plates containing 25 μg of kanamycin (Sigma, St. Louis, Mo.) per ml and 50 μg of 5-bromo-4-chloro-3-indolyl-β-D-galactoside (X-Gal) (Sigma) per ml. For heterologous expression in B. thailandensis, the B. anthracis aiiA gene was PCR amplified as described above with the forward primer 5'-CCG GAA TCT CCG ATG ACA GTA GAA AAG AAG CTI TAT TTC G-3' and reverse primer 5'-CAT GCC ATG GCA TGC TCA ACA AGA TAC TCC TAA TGA TGT-3' (Boldface represents the EcoRI and NcoI sites incorporated into the oligonucleotides for directional cloning.) The resulting amplicon was digested with EcoRI and NcoI and cloned in frame into similarly digested pBHR1 by using the Fast-Link DNA ligation kit (Epitrend Technologies). Plasmid DNA was purified using the Wizard Plus Miniprep DNA kit (Promega, Madison, Wis.), electrotransfected into E. coli S17-1, and mobilized into B. thailandensis, and transconjugants were selected on LB plates containing 25 μg of kanamycin per ml and 15 μg of polymyxin per ml (Sigma) as described previously (37). The expression vectors pME6000 and pME6863 containing the B. cereus strain A24 aiiA lactonase gene were generously forwarded to our laboratory from Clemson University and mobilized into B. thailandensis as described above.

Analysis of AHL-degrading activity by B. thailandensis expressing the AiiA lactonases. Extraction of AiiAs from culture supernatants and preparative thin-layer chromatography (TLC) were performed as described by Shaw et al. (34). Extracts were reconstituted in 1 ml of acetonitrile (Sigma) and separated (60:40 [vol/vol] methanol solution) by spotting 10 μl of a 1:10 dilution of each extract onto a reversed-phase C18 plate (Aliotech Inc., Newark, Del.). TLC plates were dried and overlaid with 30 μl of top agar (0.75% LB agar) containing 5 ml of A. tumefaciens A136 (from an overnight culture) and X-Gal. The plates were incubated at 30°C for 48 h or until adequate color development was achieved.

Motility and exoprotein secretion. Protease and lipase production were examined by using methods described by DeShazer et al. (7). To assay for beta-hemolysis, colonies were transferred onto 5% sheep blood agar plates and incubated for 48 to 72 h at 37°C. Twitching motility and swarming motility were examined by using methods described by Reimann et al. (30).

Growth kinetics and substrate utilization of B. thailandensis expressing the B. anthracis and B. cereus AiiA lactonases. From a glycerol stock of RU01, -02, -03, -04, 50 μl was inoculated into 3 ml of LB broth containing the appropriate antibiotics and incubated for 24 h at 37°C. For growth curve analysis, the optical density at 600 nm from the 3-mI overnight cultures was measured, and 25 μl was used to inoculate 250 ml of LB broth. The optical density at 600 nm was monitored hourly over a 15-h time course. At each time point, serial 10-fold
dilutions were prepared and appropriate dilutions were plated onto LB plates containing selective antibiotics.

To examine whether carbon utilization was affected by AHL cleavage, BiOLOG (Hayward, Calif.) phenotypic arrays were employed. Each derivative strain (with the exception of RU02, which was grown for 48 h) was cultured on R2A agar plates (BiOLOG) with the appropriate antibiotics for 24 h at 37°C. Cellular densities were adjusted in IF-0 fluid (BiOLOG), and plates were inoculated with 100 µl of the cell suspension. Plates were incubated at 37°C for 24 to 48 h or until sufficient color development was reached. For growth in M9 minimal medium (31), 50 µl of wild-type B. thailandensis and RU04 from an overnight culture in LB broth was pelleted (14,000 rpm for 2 min with a Spectrafuge 16M [National Labnet Co., Edison, N.J.]), washed three times in phosphate-buffered saline, resuspended in 50 µl of phosphate-buffered saline, and inoculated into M9 minimal medium with and without D-arabinose as the sole carbon source.

RESULTS

Cloning of the B. anthracis aiiA gene and expression in B. thailandensis. The B. anthracis aiiA lactonase gene was PCR amplified by using primers described by Dong et al. (9) and directionally cloned into the broad-host-range expression vector pBHR1 to produce pBA01. With pBA01 as the template, the B. anthracis aiiA lactonase gene was reamplified and sequenced to ensure fidelity during initial PCR amplification for subcloning. Nucleotide and amino acid sequence alignments between the B. anthracis aiiA and other closely related lactonase genes were performed (data not shown), and homologies were identical to the findings of Lee et al. (25). The B. cereus strain A24 aiiA lactonase gene was cloned and genetically characterized as described by Reimmann et al. (30).

Lactonase expression in B. thailandensis reduces AHL accumulation. To examine the effect of aiiA lactonase expression on AHL accumulation in B. thailandensis, a semiquantitative approach was employed, incorporating soft-agar TLC overlays containing the bioreporter strain A. tumefaciens A136. Heterologous expression of the B. cereus AiiA lactonase in B. thailandensis reduced C6-HSL, C8-HSL, and C10-HSL to undetectable levels, while the B. anthracis AiiA lactonase reduced the level of C6-HSL but completely prevented the accumulation of C8-HSL and C10-HSL (Fig. 1, lanes 4 and 5). The concentrations of C6-HSL, C8-HSL, and C10-HSL in overnight culture supernatants have not been determined for wild-type B. thailandensis. However, the limits of detection for C6-HSL, C8-HSL, and C10-HSL by A. tumefaciens A136 are approximately 300, 2.4, and 100 pmol, respectively (34), indicating that the expression of these Bacillus sp. AiiA lactonases in B. thailandensis reduces the concentration of these signaling molecules below the threshold of detection for A. tumefaciens A136 (Fig. 1, lanes 4 and 5). Maintenance of each broad-host-range expression vector (pME6000 and pBHR1) in wild-type B. thailandensis had no effect on AHL production (Fig. 1, lanes 2 and 3).

Swarming and twitching motilities in B. thailandensis are affected by aiiA expression. Several investigations have shown that QS regulates motility in gram-negative bacteria (1, 15, 23, 30). Expression of pBA01 (RU02) in B. thailandensis reduced swarming motility, while heterologous expression of pME6863 (RU04) resulted in a hyperswarming phenotype (Fig. 2, row 1, panels B and C). RU04(pME6863) translocation on swarm plates began at 12 h, and after 24 to 30 h the entire plate was colonized (Fig. 2, row 2, panel C). RU04(pME6863) grew in a rhizoid fashion with semiumbonlyte colony elevations. In contrast, RU02(pBA01) exhibited a reduction in swarming motility and displayed a colony morphology similar to that of wild-
type *B. thailandensis*. Disruption of the *btaI1* gene (*luxI* homologue) resulted in a similar phenotype observed by expression of the *B. cereus* AiiA lactonase in *B. thailandensis* (Fig. 2, row 1, panel D). Maintenance of the broad-host-range expression vector pBHR1 or pME6000 in *B. thailandensis* had no affect on swarming motility (data not shown).

Expression of pBA01 (RU02) and pME6863 (RU04) in *B. thailandensis* also resulted in altered colony morphologies on twitching motility plates. The uniform concentric ring produced by wild-type *B. thailandensis* was not observed with RU02(pBA01) and RU04(pME6863) (Fig. 2, row 2, panels B and C). The colony morphology of RU02(pBA01) was semi-round with a protruding lobate region extending at the peripheral colony margin (Fig. 2, row 2, panel B). In contrast, RU04(pME6863) grew in an irregular and spreading pattern with hilly elevations, displayed an extensive wrinkling phenotype, and produced a diffusible pale orange pigment (Fig. 2, row 2, panel C). Similarly, mutagenesis of the *btaI3* allele (*luxR* homologue) caused a similar twitching phenotype observed with RU04(pME6863). Twitching motility was not altered by the maintenance of pBHR1 or pME6000 (broad-host-range expression vectors) in *B. thailandensis* (data not shown).

**Lactonase expression in *B. thailandensis* prevents hyper-hemolysis of sheep erythrocytes.** Haussler et al. recently purified and characterized a hemolytic rhamnolipid synthesized by *B. pseudomallei* (20). Interestingly, *B. thailandensis* produces a rhamnolipid that is structurally and biochemically analogous to the *B. pseudomallei* cytotoxic and hemolytic exolipid (38). To determine whether expression of the *Bacillus* sp. AiiA lactonases in *B. thailandensis* affected beta-hemolytic activity, each derivative strain was tested on sheep blood agar plates. Expression of pBA01 (RU02) and pME6863 (RU04) in *B. thailandensis* had no effect on beta-hemolytic activity, whereas individual mutagenesis of the *btaI1* and *btaI3* genes (*luxI* mutant strains), which encode the proteins that direct the synthesis of C6-HSL and C8-HSL, resulted in hyperhemolytic phenotypes on sheep blood agar plates (Fig. 2, row 3, panels D to E). Even following prolonged incubation periods (5 to 7 days), no visible beta-hemolysis was observed for RU02(pBA01) or RU04(pME6863) (data not shown).

**Expression of the *B. anthracis* and *B. cereus* aiiA in *B. thailandensis* affects growth and carbon utilization.** During analysis of the effect of AHL cleavage on swarming and twitching motility in *B. thailandensis*, a notable AHL-dependent growth phenotype was observed for RU02(pBA01) (Table 2). To further confirm this growth reduction, the generation times of wild-type *B. thailandensis* and RU02(pBA01) were calculated throughout exponential growth in LB broth. Cleavage of C10-
HSL and C₈-HSL by expression of pBA01 (RU02) in *B. thailandensis* caused a pronounced increase in generation time (243 min) in comparison to that of wild-type *B. thailandensis* (48 min) (Table 2). In addition, the duration of logarithmic growth of RU02(pBA01) in comparison to wild-type *B. thailandensis* was reduced, as were the relative cell yields (data not shown). Further, the transition from lag-phase to logarithmic growth for RU02(pBA01) compared to wild-type *B. thailandensis* was increased (data not shown). Maintenance of the *B. cereus* AiiA lactonase (RU04) or the broad-host-range expression vectors pBHRI (RU01) and pME6000 (RU03) imposed no inhibitory effect on *B. thailandensis* growth (data not shown).

To further assess this growth characteristic, BiOLOG phenotypic microarrays were utilized to determine if carbon acquisition or metabolism was affected by AiiA lactonase expression in *B. thailandensis*. On BiOLOG PM1 plates, heterologous expression of the *B. cereus* (RU04) and *B. anthracis* (RU02) AiiA lactonases in *B. thailandensis* prevented growth on L-aspartic acid and P-hydroxy phenyl acetic acid (Table 2). Numerous substrates unique to RU02(pBA01), which heterologously expresses the *B. anthracis* AiiA lactonase and does not accumulate C₈-HSL and C₁₀-HSL (Fig. 1, lane 5), were identified on PM1 plates, including L-fucose, acetic acid, glucose-1-phosphate, citric acid, L-serine, and α-ketoglutaric acid (Table 2). As with PM1 plates, fluctuations in carbon metabolism were observed for RU02(pBA01) on the PM2 carbon source plates, including D-arabitol, xyitol, β-hydroxybutyric acid, malonic acid, and l-phenylalanine (Table 2). Of the 192 substrates analyzed, only D-arabinose appears to be negatively regulated [RU04(pME6863)]; that is, QS in *B. thailandensis* confers the ability to utilize a carbon source (Table 2). To further confirm this phenotype, wild-type *B. thailandensis* and RU04(pME6863) were cultured in M9 minimal broth containing D-arabinose as the sole carbon source. Consistent with the results obtained on PM2 plates, only RU04(pME6863) was able to metabolize D-arabinose as a sole carbon source (Table 2). A summary of the relevant phenotypes analyzed in this study is given in Table 2.

### DISCUSSION

This study analyzed the affect of heterologous expression of *Bacillus* sp. AiiA lactonases in *B. thailandensis* and found that these AHL-cleaving enzymes significantly reduced the accumulation of C₆-HSL, C₈-HSL, and C₁₀-HSL; affected bacterial motility; increased cellular generation time; and caused fluctuations in carbon metabolism and transport. The inhibition of AHL accumulation in *B. thailandensis* by maintenance of pBA01 (RU02) was similar to the cleavage pattern observed by expression of the *B. cereus* AiiA lactonase (RU04). The *B. thailandensis*...
B. thailandensis QS network resulted in defective motility phenotypes (38). As observed with the BTRJ1 (luxI) and BTRJ6 (luxR) QS mutants, expression of pBA01 (RU02) and pME6863 (RU04) in B. thailandensis also caused altered swarming and twitching motility phenotypes (Fig. 2, rows 1 and 2). The relative diameter of the swarm zone between RU02 (which does not accumulate C₈-HSL and C₁₀-HSL) and wild-type B. thailandensis was visibly different, possibly resulting from the growth phenotype observed by pBA01 expression (Fig. 2, row 1, panels A and B). However, this reduced swarm zone was still present after prolonged incubation periods (data not shown). In contrast, RU04(pME6863) demonstrated a hyperswarming phenotype as indicated by colonization of the entire swarm plate within 16 h. Interestingly, BTRJ1 (btaI1), which is a luxI mutant and deficient in C₈-HSL biosynthesis, produced a phenotype similar to that of RU04(pME6863) on swarm plates (Fig. 2, row 1, panel D). The exact mechanisms for these defects in swarming and twitching motility remain to be determined. Preliminary analysis of flagellum and pilus biosynthesis (by electron microscopy) implies that the motility defects are independent of these cellular structures required for swarming and twitching motility (data not shown). However, these findings suggest that the accumulation of C₈-HSL by B. thailandensis either directly or indirectly affects swarming motility.

As with swarming, alterations in twitching motility were observed by heterologous expression of pBA01 (RU02) and pME6863 (RU04) in B. thailandensis. Most notable was the phenotype observed for RU04(pME6863). Colony symmetry, morphology, and pigmentation of RU04 were similar to the phenotype observed for BTRJ6 (which contains an insertionally disrupted btaR3 gene, a luxR homologue) on twitching plates. Interestingly, BTRJ6 (btaR3) is a transcriptional regulator mutant, and previous analysis of AHL biosynthesis in B. thailandensis indicated that C₈-HSL, C₉-HSL, and C₁₀-HSL accumulation was not affected by disruption of any of the five encoded LuxR homologues (38). Curiously, disruption of the B. thailandensis btaI1, btaI2, and btaI3 luxI homologues failed to cause a phenotype similar to that of RU04(pME6863) on twitching plates (data not shown). These findings suggest that the altered colony morphologies observed for B. thailandensis on twitching motility plates are linked to the accumulation of multiple signaling molecules in conjunction with the BtaR3 (LuxR homologue) transcriptional regulator. However, further analysis of this phenotype at the genetic level will be necessary before any definitive conclusion can be reached.

Mutations in the btaI1 and btaI3 luxI homologues, which encode the proteins that direct the synthesis of C₈-HSL and C₉-HSL, result in elevated beta-hemolytic activity on sheep blood agar plates (Fig. 2, row 3, panels D and E). Likewise, disruption of numerous luxR homologues encoded by B. thailandensis (btaR1, btaR2, btaR3, and btaR5) also causes hyper-beta-hemolysis (38). Interestingly, and despite efficient AHL cleavage, expression of pBA01 (RU02) and pME6863 (RU04) failed to induce hyper-beta-hemolysis of sheep erythrocytes as observed by individual mutagenesis of the B. thailandensis btaI1 and btaI3 genes (Fig. 2, row 3, panels D to E). The mechanism(s), in relation to QS and this hyper-beta-hemolysis, remains to be determined; however, these findings suggest that enhanced beta-hemolytic activity observed with mutagenesis of the B. thailandensis btaR3 and btaR5 luxR homologues is independent of AHL accumulation. This hypothesis is further supported by the observation that disruption of the btaR3 and btaR5 genes has no effect on C₈-HSL, C₉-HSL, and C₁₀-HSL biosynthesis in B. thailandensis (38). The failure of RU02(pBA01) and RU04(pME6863) to lyse sheep erythrocytes may be the result of residual concentrations of AHL molecules, as seen for RU02, that are not detected by using the bioreporter strain A. tumefaciens A136 (Fig. 1, lane 5). It is also possible that B. thailandensis carries an additional luxI gene(s), which produces signaling molecules not recognized by A. tumefaciens A136, not recovered in our in silico analysis.

QS has been shown to affect the growth of various bacterial species belonging to the Rhizobium genus; however, the cellular mechanisms contributing to these reduction in growth remain to be determined (6, 18, 21, 32, 40). Interestingly, AHL cleavage in B. thailandensis, specifically C₈-HSL and C₁₀-HSL biosynthesis (Fig. 1, lane 5), by heterologous expression of pBA01 (RU02) caused an increase in generation time for RU02 (243 min) compared to wild-type B. thailandensis (48 min) and also extended the transition from lag-phase to logarithmic growth by 3 h (Table 2). Individual mutagenesis of the B. thailandensis btaI1, btaI2, and btaI3 genes, which encode the proteins that synthesize C₈-HSL, C₉-HSL, and C₁₀-HSL, respectively, failed to cause this increase in generation time, as did expression of the broad-host-range vector pBHR1 (data not shown). To further confirm these findings, and to eliminate the possibility of plasmid segregation despite antibiotic selection, serial dilutions were plated onto selective and nonselective (without kanamycin) LB plates during growth analysis (data not shown), in addition to performing colony PCR (10 isolates from each medium type) with the B. anthracis aiiA gene-specific primers. As anticipated, the proportions of recoverable bacteria on each medium type were identical, indicating stable maintenance of pBA01 (RU02). Several factors may contribute to this observed growth phenotype following AHL cleavage, with the most obvious being the metabolic burden of constitutive AiiA lactonase expression in RU02(pBA01). However, this increase...
in generation time is unique to RU02, and RU04(pME6863), which also constitutively expresses a homologous AiiA lactonase, exhibits an generation time analogous to that of wild-type B. thailandensis (data not shown). Additional parameters that may cause growth inhibition include defects in DNA translation, negative regulation (overproduction) of sigma factors, and, a focus of this study, substrate utilization. To determine if QS was linked to carbon acquisition and metabolism, BiOLOG phenotypic arrays were employed. Table 2 illustrates that AHL cleavage in B. thailandensis causes fluctuations in either the transport or catabolism of numerous carbon sources. Although to a lesser extent, individual mutagenesis of each of the B. thailandensis QS alleles also affected substrate utilization (38). Interestingly, expression of the B. cereus AiiA lactonase (RU04) in B. thailandensis demonstrated that AHL accumulation negatively regulates (confers the ability to utilize a carbon source) the protein(s) involved in the metabolism of d-arabinose (Table 2). Similar findings at the transcriptional level in P. aeruginosa PA01 have linked QS to substrate metabolism and transport (33, 39). In fact, Schuster et al. demonstrated that disruption of the P. aeruginosa PA01 lasIR and rhlIR QS networks both positively and negatively affected the transcription of various genes involved in cellular metabolism (33). The mechanism of this growth inhibition remains to be determined; however, these findings demonstrate that the B. thailandensis QS system is involved in the metabolism of various carbon sources. It is conceivable that B. thailandensis utilizes QS in the environment to reduce energy needed for substrate transport and catabolism when sufficient carbon sources (i.e., glucose versus arabinose) are present.

It seems that QS in B. thailandensis represents a complex gene regulatory network. This study demonstrates that AiiA lactonase expression in B. thailandensis reduces the accumulation of the signaling molecules C6-HSL, C8-HSL, and C10-HSL and bacterial motility; increases cellular generation time; and causes fluctuations in carbon metabolism. Stable heterologous expression of an AiiA lactonase(s) in Burkholderia species will possibly eliminate the need to generate multidilection mutants in a single organism that synthesizes numerous AHLs and may facilitate vaccine development for gram-negative bacterial pathogens that utilize QS to regulate virulence gene expression.

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