

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

Ricky L. Ulrich\*

Bacteriology Division, United States Army Medical Research Institute of Infectious Diseases,  
Fort Detrick, Maryland

Received 1 April 2004/Accepted 13 May 2004

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 (AHSs) 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 (C<sub>10</sub>-HSL) and *N*-octanoylhomoserine lactone (C<sub>8</sub>-HSL), reduced *N*-hexanoylhomoserine lactone (C<sub>6</sub>-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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-HSL to nondetectable levels; altered both swarming and twitching motility; and caused fluctuations in carbon utilization. Individual disruption of the *B. thailandensis* AHSs, specifically disruption of the *btaI1* and *btaI3* genes, which encode the proteins that direct the synthesis of C<sub>8</sub>-HSL and C<sub>6</sub>-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 processes (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  $<10^1$  CFU of *B. pseudomallei* causes mortality in hamsters within 3 to 5 days, in contrast to a dose of  $>10^6$  CFU for *B. thailandensis* (3).

Several investigations have recently identified functional QS networks in *Burkholderia* species (2, 5, 17, 26, 27). In *Burk-*

*holderia cepacia*, QS negatively regulates ornibactin biosynthesis and positively induces protease and *N*-octanoylhomoserine lactone (C<sub>8</sub>-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 (C<sub>10</sub>-HSL) and with C<sub>8</sub>-HSL and *N*-hexanoylhomoserine lactone (C<sub>6</sub>-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*-butanoylhomoserine lactone (C<sub>4</sub>-HSL) and C<sub>8</sub>-HSL

\* Mailing address: Bacteriology Division, USAMRIID, 1425 Porter St., Fort Detrick, MD 21702. Phone: (301) 619-8332. Fax: (301) 619-8351. E-mail: Ricky.Ulrich@AMEDD.ARMY.MIL.

| Report Documentation Page                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           |                              | Form Approved<br>OMB No. 0704-0188       |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------|------------------------------------------|
| Public reporting burden for the collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington VA 22202-4302. Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to a penalty for failing to comply with a collection of information if it does not display a currently valid OMB control number.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |                              |                                          |
| 1. REPORT DATE<br><b>13 MAY 2004</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                | 2. REPORT TYPE<br><b>N/A</b> | 3. DATES COVERED<br><b>-</b>             |
| 4. TITLE AND SUBTITLE<br><b>Quorum quenching: enzymatic disruption of N-acylhomoserine lactone-mediated bacterial communication in Burkholderia thailandensis, Applied Environmental Microbiology, 70:6173- 6180</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |                              | 5a. CONTRACT NUMBER                      |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |                              | 5b. GRANT NUMBER                         |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |                              | 5c. PROGRAM ELEMENT NUMBER               |
| 6. AUTHOR(S)<br><b>Ulrich, RL</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   |                              | 5d. PROJECT NUMBER                       |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |                              | 5e. TASK NUMBER                          |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |                              | 5f. WORK UNIT NUMBER                     |
| 7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)<br><b>United States Army Medical Research Institute of Infectious Diseases, Fort Detrick, MD</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 |                              | 8. PERFORMING ORGANIZATION REPORT NUMBER |
| 9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |                              | 10. SPONSOR/MONITOR'S ACRONYM(S)         |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |                              | 11. SPONSOR/MONITOR'S REPORT NUMBER(S)   |
| 12. DISTRIBUTION/AVAILABILITY STATEMENT<br><b>Approved for public release, distribution unlimited</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               |                              |                                          |
| 13. SUPPLEMENTARY NOTES<br><b>The original document contains color images.</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      |                              |                                          |
| 14. ABSTRACT<br><b>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 (AHSs) 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 (C(10)-HSL) and N-octanoylhomoserine lactone (C(8)-HSL), reduced N-hexanoylhomoserine lactone (C(6)-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 C(6)-HSL, C(8)-HSL, and C(10)-HSL to nondetectable levels; altered both swarming and twitching motility; and caused fluctuations in carbon utilization. Individual disruption of the B. thailandensis AHSs, specifically disruption of the btaI1 and btaI3 genes, which encode the proteins that direct the synthesis of C(8)-HSL and C(6)-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.</b> |                              |                                          |

|                                                                                                                    |                                    |                                     |                               |                        |                                    |
|--------------------------------------------------------------------------------------------------------------------|------------------------------------|-------------------------------------|-------------------------------|------------------------|------------------------------------|
| 15. SUBJECT TERMS                                                                                                  |                                    |                                     |                               |                        |                                    |
| <b>Burkholderia thailandensis, lactone-mediated communication, characterization, Bacillus anthracis expression</b> |                                    |                                     |                               |                        |                                    |
| 16. SECURITY CLASSIFICATION OF:                                                                                    |                                    |                                     | 17. LIMITATION OF<br>ABSTRACT | 18. NUMBER<br>OF PAGES | 19a. NAME OF<br>RESPONSIBLE PERSON |
| a. REPORT<br><b>unclassified</b>                                                                                   | b. ABSTRACT<br><b>unclassified</b> | c. THIS PAGE<br><b>unclassified</b> |                               |                        |                                    |

TABLE 1. Bacterial strains and plasmids used in this study

| Strain or plasmid                             | Description                                                                                                                | Reference or source |
|-----------------------------------------------|----------------------------------------------------------------------------------------------------------------------------|---------------------|
| <i>E. coli</i>                                |                                                                                                                            |                     |
| SM10                                          | Mobilizing strain; RP4 <i>tra</i> genes; Km <sup>r</sup>                                                                   | 35                  |
| TOP10                                         | Used for cloning and blue-white screening                                                                                  | Invitrogen          |
| S17-1                                         | <i>thi pro hsdR recA</i> ; chromosomal RP4; Tra <sup>+</sup> ; Sm <sup>r</sup> Sp <sup>r</sup>                             | 35                  |
| <i>A. tumefaciens</i> bioreporter strain A136 | Lacks the Ti plasmid                                                                                                       | 13                  |
| <i>B. thailandensis</i>                       |                                                                                                                            |                     |
| DW503                                         | Derived from E264; Δ( <i>amrR-oprA</i> )(Km <sup>s</sup> Gm <sup>s</sup> Sm <sup>s</sup> ); <i>rpsL</i> (Sm <sup>r</sup> ) | 4                   |
| RU01                                          | DW503 containing pBHR1; Km <sup>r</sup>                                                                                    | This study          |
| RU02                                          | DW503 containing pBA01; Km <sup>r</sup>                                                                                    | This study          |
| RU03                                          | DW503 containing pME6000; Tc <sup>r</sup>                                                                                  | This study          |
| RU04                                          | DW503 containing pME6863; Tc <sup>r</sup>                                                                                  | This study          |
| BTRJ1                                         | DW503 derivative; <i>11::pRU11</i> ; Gm <sup>r</sup>                                                                       | 38                  |
| BTRJ3                                         | DW503 derivative; <i>13::pRU13</i> ; Gm <sup>r</sup>                                                                       | 38                  |
| BTRJ6                                         | DW503 derivative; <i>R3::pRUR3</i> ; Gm <sup>r</sup>                                                                       | 38                  |
| Plasmids                                      |                                                                                                                            |                     |
| pPCR2.1-TOPO                                  | TA cloning vector; Km <sup>r</sup> Ap <sup>r</sup>                                                                         | Invitrogen          |
| pBHR1                                         | Mobilizable broad-host-range vector; Km <sup>r</sup> Cm <sup>r</sup>                                                       | MoBiTee             |
| pME6000                                       | Broad-host-range cloning vector; Tc <sup>r</sup>                                                                           | 30                  |
| pME6863                                       | pME6000 carrying the <i>aiaA</i> gene from <i>B. cereus</i> strain A24; Tc <sup>r</sup>                                    | 30                  |
| pBA01                                         | pBHR1 containing the <i>B. anthracis</i> (Ames strain) <i>aiaA</i> gene                                                    | This study          |

into L-homoserine (41). This acylase I-mediated deacylation of C<sub>4</sub>-HSL and C<sub>8</sub>-HSL also reduced biofilm formation in an aquarium water model (41). Enzymatic cleavage, specifically lactone ring hydrolysis, of signaling molecules by numerous *Bacillus* species has also been reported (9–11). These enzymes, termed AiiA lactonases, hydrolyze the lactone bond within the AHL moiety, thus changing the relative conformational structure of the signaling molecule, which prevents binding to the LuxR transcriptional regulator. Heterologous expression of the *Bacillus cereus* strain A24 AiiA lactonase in *Pseudomonas aeruginosa* PA01 reduced AHL accumulation, altered swarming motility, and decreased the expression and secretion of numerous virulence factors, including elastase, rhamnolipid, hydrogen cyanide, and pyocyanin (30). In *Erwinia carotovora*, a plant pathogen, *aiaA* expression reduced the synthesis of pectolytic enzymes and reduced pathogenicity in several plant models of infection (11).

This study analyzes the effect of C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-HSL cleavage on twitching and swarming motility, beta-hemolysis, and carbon utilization by the expression of *Bacillus* sp. AiiA lactonases in *B. thailandensis*.

#### 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* *aiaA*.** Genomic DNA for PCR amplification of the *Bacillus anthracis* (Ames strain) *aiaA* 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* *aiaA* gene was PCR amplified by using the primer set (forward, 5'-ATG GGA TCC ATG ACA GTA AAG AAG CTT TAT-3'; reverse, 5'-GTC GAA TTC CTC AAC AAG ATA CTC CTA ATG-3') reported by Dong et al. (9) and amplified as follows: one cycle of 94°C for 5 min; 30 cycles of 94°C for 30 s, 50°C for 30 s, and 72°C for 1 min; and a final 7-min extension at 72°C. All PCRs were performed with the FailSafe PCR system with buffer E (Epicentre Technologies,

Madison, Wis.), and amplicons were subcloned into pCR2.1-TOPO (Invitrogen, Carlsbad, Calif.). Ligations 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* *aiaA* gene was PCR amplified as described above with the forward primer 5'-CCG GAA TTC CGC ATG ACA GTA AAG AAG CTT 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 (Epicentre Technologies). Plasmid DNA was purified using the Wizard Plus Miniprep DNA kit (Promega, Madison, Wis.), electroporated 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 *aiaA* 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 AHLs 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 C<sub>18</sub> plate (Analtech Inc., Newark, Del.). TLC plates were dried and overlaid with 30 ml 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 exoproduct 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 Reimmann 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, or -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-ml 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

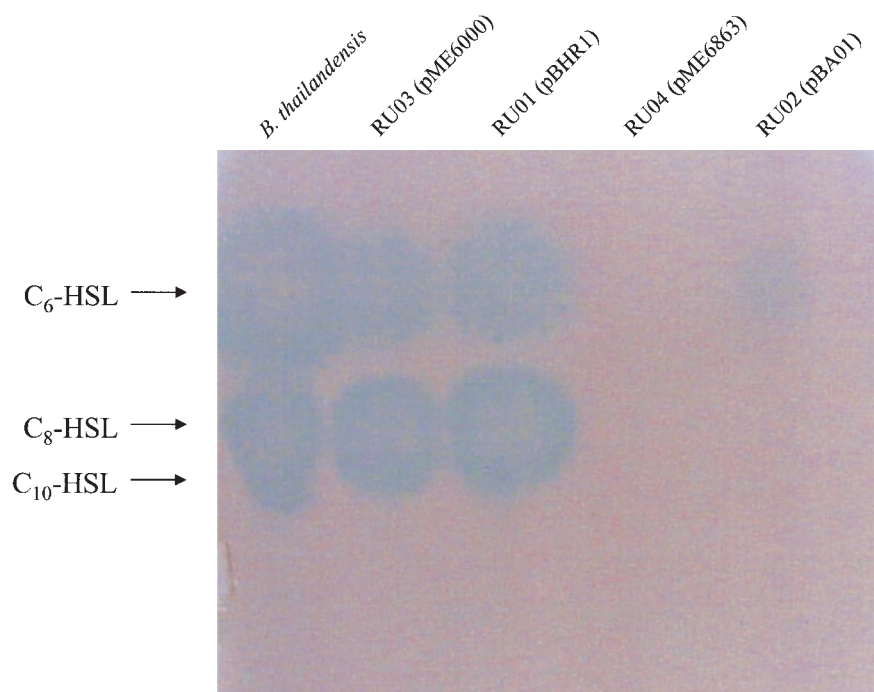


FIG. 1. Expression of the *B. anthracis* and *B. cereus* AiiA lactonases reduces AHL accumulation in *B. thailandensis* culture supernatants. TLC plates were overlaid with the bioreporter strain *A. tumefaciens* A136 and incubated at 30°C for 48 h. Each lane contains 10  $\mu$ l (1:10 dilution) of the corresponding AHL extract. Strain genotypes are listed in Table 1.

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  $\mu$ l 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  $\mu$ l 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  $\mu$ l 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* *aiaA* gene and expression in *B. thailandensis*.** The *B. anthracis* *aiaA* 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* *aiaA* 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* *aiaA* 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 *aiaA* lactonase gene was cloned and genetically characterized as described by Reimann et al. (30).

**Lactonase expression in *B. thailandensis* reduces AHL accumulation.** To examine the effect of *aiaA* 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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-HSL to undetectable levels, while the *B. anthracis* AiiA lactonase reduced the level of C<sub>6</sub>-HSL but completely prevented the accumulation of C<sub>8</sub>-HSL and C<sub>10</sub>-HSL (Fig. 1, lanes 4 and 5). The concentrations of C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-HSL in overnight culture supernatants have not been determined for wild-type *B. thailandensis*. However, the limits of detection for C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-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 *aiaA* 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 1, panel C). RU04(pME6863) grew in a rhizoid fashion with semiumbonate colony elevations. In contrast, RU02(pBA01) exhibited a reduction in swarming motility and displayed a colony morphology similar to that of wild-

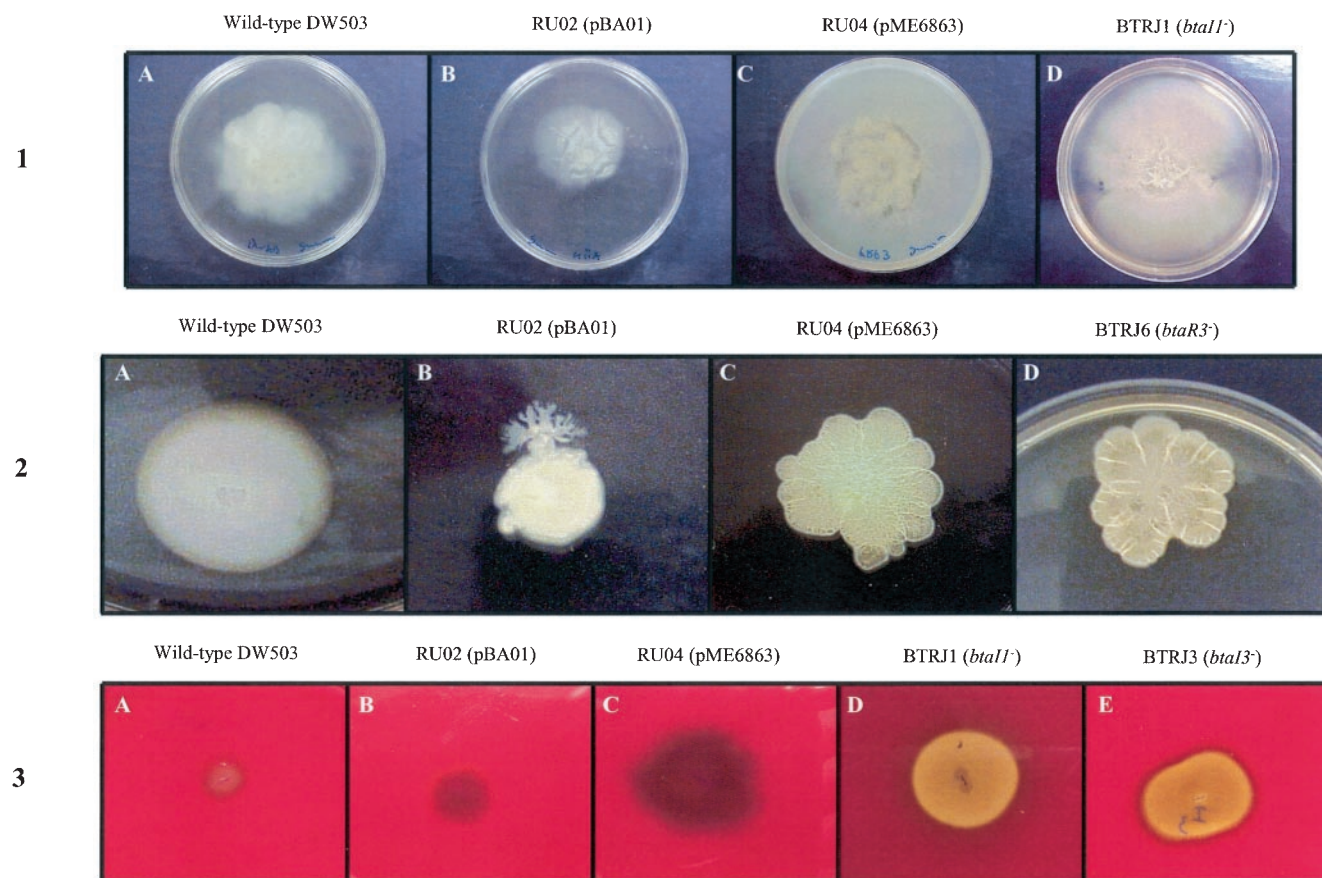


FIG. 2. AHL cleavage in *B. thailandensis* affects swarming and twitching motility in addition to the beta-hemolysis of sheep erythrocytes. Row 1, swarming motility; row 2, twitching motility; row 3, beta-hemolytic activity. DW503 represents wild-type *B. thailandensis*. Strain genotypes are listed above each image and are described in Table 1. The *B. thailandensis luxI* genes are listed as *btaI*, while the *luxR* alleles are listed as *btaR*.

type *B. thailandensis*. Disruption of the *btaII* 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 effect 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 *btaR3* 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 puri-

fied 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 C<sub>6</sub>-HSL and C<sub>8</sub>-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 C<sub>10</sub>-

TABLE 2. Phenotypes affected by heterologous expression of *aiiA* lactonases in *B. thailandensis*

| Substrate                          | Phenotype for strain <sup>a</sup> : |      |      |      |      |       |       |       |
|------------------------------------|-------------------------------------|------|------|------|------|-------|-------|-------|
|                                    | DW503                               | RU01 | RU02 | RU03 | RU04 | BTRJ1 | BTRJ3 | BTRJ6 |
| C <sub>6</sub> -HSL                | D                                   | D    | D    | D    | ND   | D     | ND    | D     |
| C <sub>8</sub> -HSL                | D                                   | D    | ND   | D    | ND   | ND    | D     | D     |
| C <sub>10</sub> -HSL               | D                                   | D    | ND   | D    | ND   | D     | D     | D     |
| Swarming motility                  | NA                                  | NA   | M    | NA   | M    | M     | NA    | NA    |
| Twitching motility                 | NA                                  | NA   | M    | NA   | M    | NA    | NA    | A     |
| β-Hemolysis                        | NA                                  | NA   | NA   | NA   | NA   | A     | A     | A     |
| Generation time                    | NA                                  | NA   | A    | NA   | NA   | NA    | NA    | NA    |
| α-Hydroxybutyric acid              | G                                   | NG   | NG   | NG   | NG   | G     | G     | G     |
| L-Threonine                        | G                                   | NG   | NG   | NG   | NG   | G     | G     | G     |
| L-Alanylglycine                    | G                                   | NG   | NG   | NG   | NG   | G     | G     | G     |
| D-Xylose                           | G                                   | G    | G    | NG   | NG   | G     | G     | G     |
| P-Hydroxy phenyl acetic acid       | G                                   | G    | NG   | G    | NG   | G     | G     | G     |
| D-Trehalose                        | G                                   | NG   | G    | NG   | NG   | G     | G     | G     |
| D-Glucosaminic acid                | G                                   | G    | G    | G    | NG   | G     | G     | G     |
| α-Ketobutyric acid                 | G                                   | G    | G    | G    | NG   | G     | G     | G     |
| L-Aspartic acid                    | G                                   | G    | NG   | G    | NG   | G     | G     | G     |
| L-Fucose                           | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| Acetic acid                        | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| α-Ketoglutaric acid                | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| Glucose-1-phosphate                | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| Citric acid                        | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| L-Serine                           | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| Gelatin                            | G                                   | NG   | G    | NG   | G    | G     | G     | G     |
| L-Isolucine                        | G                                   | NG   | NG   | NG   | G    | G     | G     | G     |
| DL-Camitine                        | G                                   | NG   | NG   | NG   | G    | NG    | G     | G     |
| D-Arabinose                        | NG                                  | NG   | NG   | NG   | G    | NG    | G     | NG    |
| D-Arabitol                         | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| Xylitol                            | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| β-Hydroxybutyric acid              | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| Malonic acid                       | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| L-Phenylalanine                    | G                                   | G    | NG   | G    | G    | G     | G     | G     |
| M9 minimal medium with D-arabinose | NG                                  | NG   | NG   | NG   | G    | NT    | NG    | NT    |

<sup>a</sup> D, detected; ND, not detected; M, colony morphology altered; NA, not affected; A, affected; G, growth; NG, no growth; NT, not tested. Strain genotypes are listed in Table 1.

HSL and C<sub>8</sub>-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 C<sub>8</sub>-HSL and C<sub>10</sub>-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 (Ta-

ble 2). As with PM1 plates, fluctuations in carbon metabolism were observed for RU02(pBA01) on the PM2 carbon source plates, including D-arabitol, xylitol, 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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-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.*

*anthracis* AiiA lactonase efficiently cleaved both C<sub>8</sub>-HSL and C<sub>10</sub>-HSL in *B. thailandensis*; however, trace amounts of C<sub>6</sub>-HSL were still detected by using TLC bioreporter overlays (Fig. 1, lane 5). In contrast, expression of the *B. cereus* AiiA lactonase in *B. thailandensis* completely abolished the accumulation of C<sub>10</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>6</sub>-HSL (Fig. 1, lane 4). These findings are consistent with the results reported by Lee et al., which demonstrated that numerous *Bacillus thuringiensis* strains produce biologically active AiiA lactonases that vary in enzymatic activity (25). Despite the detection of C<sub>6</sub>-HSL in RU02(pBA01) cell supernatants, these findings demonstrate that expression of AiiA lactonases in *B. thailandensis* reduces AHL accumulation (Fig. 1, lanes 4 and 5). The exact mechanism for AHL cleavage mediated by the *B. cereus* strain A24 and *B. anthracis* AiiA lactonases has not been deciphered and was not within the scope of this investigation. However, considering the extensive homology between these AHL-cleaving enzymes and other AiiA lactonases, it is likely that the reduction in AHL accumulation reported in this study is the result of enzymatic lactone ring hydrolysis within the AHL molecule (9–11).

Like *B. pseudomallei*, *B. thailandensis* is capable of bacterial motility, and disruption (individual gene mutagenesis) of the *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 C<sub>8</sub>-HSL and C<sub>10</sub>-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 C<sub>8</sub>-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 pillus 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 C<sub>8</sub>-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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-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 C<sub>8</sub>-HSL and C<sub>6</sub>-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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-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 C<sub>8</sub>-HSL and C<sub>10</sub>-HSL (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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-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* *aiaA* 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 C<sub>6</sub>-HSL, C<sub>8</sub>-HSL, and C<sub>10</sub>-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 multideletion 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.

#### ACKNOWLEDGMENTS

I extend a special thank you to Melanie Ulrich and Kathryn Kenyon for reviewing the manuscript and to Donald J. Chabot for providing purified *B. anthracis* genomic DNA for *aiaA* lactonase gene cloning. I additionally thank David DeShazer for scientific and technical discussions regarding this investigation.

The research described herein was sponsored by the Medical Biological Defense Research Program, U.S. Army Medical Research and Materiel Command.

Opinions, interpretations, conclusions, and recommendations are those of the author and are not necessarily endorsed by the U.S. Army.

#### REFERENCES

- Atkinson, S., J. P. Throup, G. S. Stewart, and P. Williams. 1999. A hierarchical quorum-sensing system in *Yersinia pseudotuberculosis* is involved in the regulation of motility and clumping. *Mol. Microbiol.* **33**:1267–1277.
- Baldwin, A., P. A. Sokol, J. Parkhill, and E. Mahenthalingam. 2004. The *Burkholderia cepacia* epidemic strain marker is part of a novel genomic island encoding both virulence and metabolism-associated genes in *Burkholderia cenocepacia*. *Infect. Immun.* **72**:1537–1547.
- Brett, P. J., D. DeShazer, and D. E. Woods. 1998. *Burkholderia thailandensis* sp. nov., a *Burkholderia pseudomallei*-like species. *Int. J. Syst. Bacteriol.* **48**:317–320.
- Burtnick, M., A. Bolton, P. Brett, D. Watanabe, and D. Woods. 2001. Identification of the acid phosphatase (*acpA*) gene homologues in pathogenic and non-pathogenic *Burkholderia* spp. facilitates TnphoA mutagenesis. *Microbiology* **147**:111–120.
- Conway, B. A., and E. P. Greenberg. 2002. Quorum-sensing signals and quorum-sensing genes in *Burkholderia vietnamiensis*. *J. Bacteriol.* **184**:1187–1191.
- Daniels, R., D. E. De Vos, J. Desair, G. Raedschelders, E. Luyten, V. Rosemeyer, C. Verreth, E. Schoeters, J. Vanderleyden, and J. Michiels. 2002. The *cin* quorum sensing locus of *Rhizobium etli* CNPAF512 affects growth and symbiotic nitrogen fixation. *J. Biol. Chem.* **277**:462–468.
- DeShazer, D., P. J. Brett, M. N. Burtnick, and D. E. Woods. 1999. Molecular characterization of genetic loci required for secretion of exoproducts in *Burkholderia pseudomallei*. *J. Bacteriol.* **181**:4661–4664.
- Donabedian, H. 2003. Quorum sensing and its relevance to infectious diseases. *J. Infect.* **46**:207–214.
- Dong, Y. H., A. R. Gusti, Q. Zhang, J. L. Xu, and L. H. Zhang. 2002. Identification of quorum-quenching *N*-acyl homoserine lactonases from *Bacillus* species. *Appl. Environ. Microbiol.* **68**:1754–1759.
- Dong, Y. H., L. H. Wang, J. L. Xu, H. B. Zhang, X. F. Zhang, and L. H. Zhang. 2001. Quenching quorum-sensing-dependent bacterial infection by an *N*-acyl homoserine lactonase. *Nature* **411**:813–817.
- Dong, Y. H., J. L. Xu, X. Z. Li, and L. H. Zhang. 2000. AiiA, an enzyme that inactivates the acylhomoserine lactone quorum-sensing signal and attenuates the virulence of *Erwinia carotovora*. *Proc. Natl. Acad. Sci. USA* **97**:3526–3531.
- Fuqua, C., M. R. Parsek, and E. P. Greenberg. 2001. Regulation of gene expression by cell-to-cell communication: acyl-homoserine lactone quorum sensing. *Annu. Rev. Genet.* **35**:439–468.
- Fuqua, C., and S. C. Winans. 1996. Conserved *cis*-acting promoter elements are required for density-dependent transcription of *Agrobacterium tumefaciens* conjugal transfer genes. *J. Bacteriol.* **178**:435–440.
- Fuqua, W. C., S. C. Winans, and E. P. Greenberg. 1994. Quorum sensing in bacteria: the LuxR-LuxI family of cell density-responsive transcriptional regulators. *J. Bacteriol.* **176**:269–275.
- Glessner, A., R. S. Smith, B. H. Iglewski, and J. B. Robinson. 1999. Roles of *Pseudomonas aeruginosa* *las* and *rhl* quorum-sensing systems in control of twitching motility. *J. Bacteriol.* **181**:1623–1629.
- Godoy, D., G. Randle, A. J. Simpson, D. M. Aanensen, T. L. Pitt, R. Kinoshita, and B. G. Spratt. 2003. Multilocus sequence typing and evolutionary relationships among the causative agents of melioidosis and glanders, *Burkholderia pseudomallei* and *Burkholderia mallei*. *J. Clin. Microbiol.* **41**:2068–2079.
- Gotschlich, A., B. Huber, O. Geisenberger, A. Togl, A. Steidle, K. Riedel, P. Hill, B. Tummler, P. Vandamme, B. Middleton, M. Camara, P. Williams, A. Hardman, and L. Eberl. 2001. Synthesis of multiple *N*-acylhomoserine lactones is wide-spread among the members of the *Burkholderia cepacia* complex. *Syst. Appl. Microbiol.* **24**:1–14.
- Gray, K. M., J. P. Pearson, J. A. Downie, B. E. Boboye, and E. P. Greenberg. 1996. Cell-to-cell signaling in the symbiotic nitrogen-fixing bacterium *Rhizobium leguminosarum*: autoinduction of a stationary phase and rhizosphere-expressed genes. *J. Bacteriol.* **178**:372–376.
- Hardman, A. M., G. S. Stewart, and P. Williams. 1998. Quorum sensing and the cell-cell communication dependent regulation of gene expression in pathogenic and non-pathogenic bacteria. *Antonie Leeuwenhoek* **74**:199–210.
- Haussler, S., M. Nimtz, T. Domke, V. Wray, and I. Steinmetz. 1998. Purification and characterization of a cytotoxic exolipid of *Burkholderia pseudomallei*. *Infect. Immun.* **66**:1588–1593.
- He, X., W. Chang, D. L. Pierce, L. O. Seib, J. Wagner, and C. Fuqua. 2003. Quorum sensing in *Rhizobium* sp. strain NGR234 regulates conjugal transfer (*tra*) gene expression and influences growth rate. *J. Bacteriol.* **185**:809–822.
- Huang, J. J., J. I. Han, L. H. Zhang, and J. R. Leadbetter. 2003. Utilization of acyl-homoserine lactone quorum signals for growth by a soil pseudomonad and *Pseudomonas aeruginosa* PAO1. *Appl. Environ. Microbiol.* **69**:5941–5949.
- Kohler, T., L. K. Curty, F. Barja, C. van Delden, and J. C. Pechere. 2000. Swarming of *Pseudomonas aeruginosa* is dependent on cell-to-cell signaling and requires flagella and pili. *J. Bacteriol.* **182**:5990–5996.
- Leadbetter, J. R., and E. P. Greenberg. 2000. Metabolism of acyl-homoserine lactone quorum-sensing signals by *Variovorax paradoxus*. *J. Bacteriol.* **182**:6921–6926.
- Lee, S. J., S. Y. Park, J. J. Lee, D. Y. Yum, B. T. Koo, and J. K. Lee. 2002. Genes encoding the *N*-acyl homoserine lactone-degrading enzyme are widespread in many subspecies of *Bacillus thuringiensis*. *Appl. Environ. Microbiol.* **68**:3919–3924.
- Lewenza, S., B. Conway, E. P. Greenberg, and P. A. Sokol. 1999. Quorum sensing in *Burkholderia cepacia*: identification of the LuxRI homologs CepRI. *J. Bacteriol.* **181**:748–756.
- Lewenza, S., and P. A. Sokol. 2001. Regulation of ornibactin biosynthesis and

- N*-acyl-L-homoserine lactone production by CepR in *Burkholderia cepacia*. J. Bacteriol. **183**:2212–2218.
28. Lin, Y. H., J. L. Xu, J. Hu, L. H. Wang, S. L. Ong, J. R. Leadbetter, and L. H. Zhang. 2003. Acyl-homoserine lactone acylase from *Ralstonia* strain XJ12B represents a novel and potent class of quorum-quenching enzymes. Mol. Microbiol. **47**:849–860.
29. Rasmussen, T. B., M. Manefield, J. B. Andersen, L. Eberl, U. Anthoni, C. Christophersen, P. Steinberg, S. Kjelleberg, and M. Givskov. 2000. How *Delisea pulchra* furanones affect quorum sensing and swarming motility in *Serratia liquefaciens* MG1. Microbiology **146**:3237–3244.
30. Reimmann, C., N. Ginot, L. Michel, C. Keel, P. Michaux, V. Krishnapillai, M. Zala, K. Heurlier, K. Triandafillu, H. Harms, G. Defago, and D. Haas. 2002. Genetically programmed autoinducer destruction reduces virulence gene expression and swarming motility in *Pseudomonas aeruginosa* PAO1. Microbiology **148**:923–932.
31. Sambrook, J., E. F. Fritsch, and T. Maniatis. 1989. Molecular cloning: a laboratory manual, 2nd ed. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y.
32. Schripsema, J., K. E. de Rudder, T. B. van Vliet, P. P. Lankhorst, E. de Vroom, J. W. Kijne, and A. A. van Brussel. 1996. Bacteriocin small of *Rhizobium leguminosarum* belongs to the class of *N*-acyl-L-homoserine lactone molecules, known as autoinducers and as quorum-sensing cotranscription factors. J. Bacteriol. **178**:366–371.
33. Schuster, M., C. P. Lostroh, T. Ogi, and E. P. Greenberg. 2003. Identification, timing, and signal specificity of *Pseudomonas aeruginosa* quorum-controlled genes: a transcriptome analysis. J. Bacteriol. **185**:2066–2079.
34. Shaw, P. D., G. Ping, S. L. Daly, C. Cha, J. E. Cronan, Jr., K. L. Rinehart, and S. K. Farrand. 1997. Detecting and characterizing *N*-acyl-homoserine lactone signal molecules by thin-layer chromatography. Proc. Natl. Acad. Sci. USA **94**:6036–6041.
35. Simon, R., U. Priefer, and A. Pühler. 1983. A broad host range mobilization system for in vivo genetic engineering: transposon mutagenesis in Gram-negative bacteria. Biotechnology **1**:784–791.
36. Swift, S., J. A. Downie, N. A. Whitehead, A. M. Barnard, G. P. Salmond, and P. Williams. 2001. Quorum sensing as a population-density-dependent determinant of bacterial physiology. Adv. Microb. Physiol. **45**:199–270.
37. Ulrich, R. L., and D. DeShazer. 2004. Type III secretion: a virulence factor delivery system essential for the pathogenicity of *Burkholderia mallei*. Infect. Immun. **72**:1150–1154.
38. Ulrich, R. L., H. B. Hines, N. Parthasarathy, and J. A. Jeddelloh. Mutational analysis and biochemical characterization of the *Burkholderia thailandensis* DW503 quorum-sensing network. J. Bacteriol. **186**:4350–4360.
39. Wagner, V. E., D. Bushnell, L. Passador, A. I. Brooks, and B. H. Iglewski. 2003. Microarray analysis of *Pseudomonas aeruginosa* quorum-sensing regulators: effects of growth phase and environment. J. Bacteriol. **185**:2080–2095.
40. Wilkinson, A., V. Danino, F. Wisniewski-Dye, J. K. Lithgow, and J. A. Downie. 2002. *N*-Acyl-homoserine lactone inhibition of rhizobial growth is mediated by two quorum-sensing genes that regulate plasmid transfer. J. Bacteriol. **184**:4510–4519.
41. Xu, F., T. Byun, H. J. Deussen, K. R. Duke, and H. J. Dussen. 2003. Degradation of *N*-acylhomoserine lactones, the bacterial quorum-sensing molecules, by acylase. J. Biotechnol. **101**:89–96.