Inactivation of the Pta-AckA Pathway Causes Cell Death in Staphylococcus aureus

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During growth under conditions of glucose and oxygen excess, Staphylococcus aureus predominantly accumulates acetate in the culture medium, suggesting that the phosphotransacetylase-acetate kinase (Pta-AckA) pathway plays a crucial role in bacterial fitness. Previous studies demonstrated that these conditions also induce the S. aureus CidR regulon involved in the control of cell death. Interestingly, the CidR regulon is comprised of only two operons, both encoding pyruvate catabolic enzymes, suggesting an intimate relationship between pyruvate metabolism and cell death. To examine this relationship, we introduced ackA and pta mutations in S. aureus and tested their effects on bacterial growth, carbon and energy metabolism, cid expression, and cell death. Inactivation of the Pta-AckA pathway showed a drastic inhibitory effect on growth and caused accumulation of dead cells in both pta and ackA mutants. Surprisingly, inactivation of the Pta-AckA pathway did not lead to a decrease in the energy status of bacteria, as the intracellular concentrations of ATP, NAD+, and NADH were higher in the mutants. However, inactivation of this pathway increased the rate of glucose consumption, led to a metabolic block at the pyruvate node, and enhanced carbon flux through both glycolysis and the tricarboxylic acid (TCA) cycle. Intriguingly, disruption of the Pta-AckA pathway also induced the CidR regulon, suggesting that activation of alternative pyruvate catabolic pathways could be an important survival strategy for the mutants. Collectively, the results of this study demonstrate the indispensable role of the Pta-AckA pathway in S. aureus for maintaining energy and metabolic homeostasis during overflow metabolism.

Staphylococcus aureus is a dangerous human pathogen, asymptptomatically colonizing over 30% of healthy individuals worldwide yet capable of causing invasive opportunistic infections (1). It is a leading cause of both nosocomial and community-associated infections, ranging from mild skin infections to life-threatening diseases, such as severe sepsis, necrotizing pneumonia, endocarditis, and bacteremia (2, 3). S. aureus infections represent an enormous challenge to physicians, due to the ability of the bacterial pathogen to survive in diverse host environments and the emergence of multidrug-resistant strains (1, 3). The versatility of the bacterium is dependent on its proficiency in sensing and utilizing nutrients from different sources and responding appropriately to rapid environmental changes. This is achieved by maximizing virulence factor and energy production via modulation of the expression of genes involved in different metabolic pathways (4–9).

During growth, staphylococci catabolize glucose and other carbohydrates primarily through glycolysis (the Embden-Meyerhof-Parnas pathway) and the pentose phosphate pathway (10). Carbon flux through glycolysis produces two molecules of pyruvate per molecule of glucose consumed, generates two molecules of ATP, and reduces two molecules of NAD+ to NADH (8). During anaerobic growth, pyruvate undergoes mixed-acid fermentation, leading to the generation of lactate, formate, acetate, and ethanol, with the concomitant oxidation of reducing equivalents during lactate and ethanol fermentation (10–12). Under aerobic conditions, however, pyruvate is decarboxylated to acetyl-coenzyme A (Ac-CoA) and CO2 by the pyruvate dehydrogenase complex (PDHC), with reduction of one molecule of NAD+ to NADH (8, 13). Depending on the growth conditions, acetyl-CoA is then oxidized by the tricarboxylic acid (TCA) cycle and/or is hydrolyzed to acetate by the phosphotransacetylase-acetate kinase (Pta-AckA) pathway, which generates one molecule of ATP through substrate level phosphorylation. For example, it was shown that, under conditions of excess glucose and oxygen (overflow metabolism), when carbon flow into the TCA cycle is limited by carbon catabolite repression (6, 14–16), S. aureus predominantly excreted acetate into the culture medium (10, 17). This implies a critical role for the Pta-AckA pathway during overflow metabolism, and its activity may be essential for the growth and fitness of S. aureus, as has been reported for other bacteria (18–22). To date, however, the Pta-AckA pathway in staphylococci has not been characterized.

Interestingly, the LysR-type transcriptional regulator CidR, which is involved in control of the molecular components mediating cell death in S. aureus, exerts its control during aerobic growth under carbon excess (23). Inactivation of the cidR gene reduces transcription of the cid operon, encoding a putative holin that affects murein hydrolase activity and cell viability (23). Intriguingly, CidR function during overflow metabolism also affects the transcription of genes involved in diverse pathways of pyruvate catabolism (i.e., cidC and alsSD operons), leading to the generation of acetate and acetoain (24). It has been speculated that activation of the CidR regulon during overflow metabolism may result from the accumulation of intracellular pyruvate (25, 26).
TABLE 1 Strains and plasmids used in this study

<table>
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<th>Strains or plasmid</th>
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<td>RN4220</td>
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In the present study, we highlight the contribution of the Pta-AckA pathway to staphylococcal metabolism. We demonstrate that disruption of either pta or ackA in S. aureus has a drastic inhibitory effect on growth during overflow metabolism and leads to significant accumulation of dead cells during the exponential phase of growth. Our results show that the detrimental effects on growth were not caused by the decrease in the energy status of bacteria, as intracellular concentrations of ATP, NAD⁺, and NADH were found to be higher in the mutants but appear to be associated with a metabolic block at the pyruvate node that globally alters the intracellular metabolic status and activates CidR-dependent pathways.

**MATERIALS AND METHODS**

**Bacterial strains, plasmids, and growth conditions.** The bacterial strains and plasmids used in this study are listed in Table 1. Escherichia coli strains were grown in LB broth (Novagen, EMD) or on LB agar. S. aureus strains were grown in tryptic soy broth (TSB) without dextrose (BD Biosciences) supplemented with 0.25% glucose (Sigma-Aldrich) or on TSB containing agar. S. aureus cultures were inoculated to 0.06 optical density at 600 nm (OD₆₀₀) units from overnight cultures, incubated at 37°C, and aerated at 250 rpm with a flask-to-medium ratio of 10:1. Bacterial growth was assessed by measuring the optical density at 600 nm or by determining the number of CFU ml⁻¹. Antibiotics were purchased from Fisher Scientific and were used at the following concentrations: ampicillin, 100 μg/ml; chloramphenicol, 10 μg/ml; and erythromycin, 10 μg/ml.

**Construction of the ackA and pta mutants in S. aureus.** All primers (see Table S1 in the supplemental material) used for construction and confirmation of the ackA and pta mutations were generated based on the sequence of S. aureus strain MRSA252 (NC_002952.2). The ackA mutant was constructed by replacing a 1.2-kb region, which included the ackA gene, with an erythromycin resistance gene (ermB), using the gene splicing by overlap extension (SOE) technique (27). The ermB antibiotic resistance cassette was obtained from pEC4 (28), using ackA-ermB-f and ackA-ermB-r primers, which contain sequences homologous to the ackA gene. The primers SAR1790-f and ermB-ackA-r were used for amplification of a 1.4-kb region upstream of the ackA gene, while a 1.3-kb region downstream of the ackA gene was amplified using ermB-ackA-f and SacI-ald1-r primers. All three amplified fragments were mixed in equimolar ratios (1:1:1) and amplified using SAR1790-f and SacI-ald1-r primers. The resulting 3.7-kb PCR product consisted of the 1-kb ermB gene flanked by sequences upstream and downstream of the ackA gene. Following digestion with the restriction endonucleases SacI and Smal, the 3.7-kb product was cloned into pTSI-d (29). The resulting plasmid (pMRS78) was used to construct the UAMS-1-ackA:ermB mutant through standard allelic-exchange methodology described in an earlier study (30). Replacement of the internal region of the ackA gene by the ermB cassette was verified by PCR using the primers ermB-f, ermB-r, ald1-r, and SAR1790-f.

The pta mutant was constructed by in-frame deletion of the pta gene, using the gene SOE technique (27). Primers SacI-SAR0593-f and pLapl-eutD-r were used for amplification of a 0.9-kb region upstream of the pta gene. Primers eutD-lplA-ermB and lplA-ram were used to amplify a 1.1-kb region downstream of the pta gene containing the IplA gene. Importantly, primers pLapl-eutD-r and eutD-lplA-f contain a ribosome binding site (RBS) of the pta (eutD) gene combined with the start codon (ATG) of the IplA gene. The amplified fragments were mixed in equimolar ratios (1:1) and amplified using SacI-SAR0593-f and IplA-r primers. The resulting 2.0-kb PCR product containing a single SacI site was cloned into pTSI-d (29) digested with SacI and SmaI to generate the plasmid pWS2. Plasmid pWS2 was used to construct the UAMS-1-Δpta mutant by allelic-exchange methodology (30), and the in-frame deletion of the pta gene in the KB8001 strain was verified by PCR and DNA sequencing, using primers SAR0592-f, SAR0596-r, IplA-r1, and SacI-P-eutD-f.

The pta-cidC double mutant was generated by bacteriophage Φ11-mediated transduction (31) of the cidC:ermC allele from KB1058 and confirmed by PCR using sets of primers, SAR0592-f and SAR0596-r, and cidA-f and cidC DN. The UAMS-1 background in all mutants was confirmed by pulsed-field gel electrophoresis (PFGE) and PCR using primers cna-f and cna-r.

**Complementation of the ackA and pta mutants.** For complementation of the ackA mutation, a 1.2-kb PCR product containing the promoterless wild-type ackA gene was amplified using primers Sall-RBS-ackA-f and Sall-ackA-r (see Table S1 in the supplemental material). Following digestion with the restriction endonucleases Sall and SacI, the PCR product was ligated into the plasmid pBK123 under the control of the cadC gene (32). The resulting recombinant plasmid was designated pWS5. For complementation of the pta mutation, a 1.2-kb PCR product containing the wild-type pta allele with its promoter region was amplified using primers Sall-P-eutD-f and Sall-eutD-r (see Table S1 in the supplemental material). The PCR product containing Sall and SacI sites was cloned into pLI150 (33) digested with Sall and SacI to generate the plasmid pMRS102. The plasmids pWS5 and pMRS102 were transformed into strain RN4220 by electroporation and then introduced into UAMS-1-ackA:ermB and UAMS-1-Δpta strains, respectively, by phage Φ11-mediated transduction (31).

**Measurement of glucose, acetic acid, lactate, pyruvate, and acetoin concentrations in the culture medium.** Aliquots of bacterial cultures (1 ml) were centrifuged for 3 min at 14,000 rpm at 4°C. The supernatants were removed and stored at −20°C until use. Acetate, glucose, and D- and L-lactate concentrations were determined using kits purchased from R-Biopharm, according to the manufacturer’s protocol. Pyruvate concentrations were determined using the Pyruvate Assay Kit (MBL) according to the manufacturer’s protocol. Acoutin concentrations were determined at 560 nm as described previously (34).

**Determination of intracellular ATP, NAD⁺, NADH, Ac-CoA, and pyruvate concentrations.** Intracellular ATP concentrations were determined using the BacTiter-Glo kit (Promega) according to the manufac-
turier’s protocol and normalized to the corresponding viable-cell counts at the time of harvest.

Intracellular NAD⁺ and NADH concentrations were determined using a Fluoro NAD/NADH kit (Cell Technology). Aliquots of bacterial cultures (10 ml) were harvested by centrifugation at 4°C for 10 min at 4,000 rpm. The bacterial pellets were washed twice with 1 ml of phosphate-buffered saline, pH 7.4 (PBS). Then, the bacterial pellets were resuspended in 0.2 ml of the NAD/NADH extraction buffer and 0.2 ml of the lysis buffer. The cells were lysed using lysing matrix B tubes in a FastPrep instrument (Qbiogene), and then the lysates were incubated at 60°C for 15 min. The lysates were centrifuged at 4°C for 3 min at 14,000 rpm. NAD⁺ and NADH concentrations in the lysates were determined according to the manufacturer’s protocol and normalized to the corresponding viable-cell counts determined at the time of harvest.

Intracellular Ac-CoA concentrations were determined using a PicoProbe Acetyl-CoA Assay Kit (BioVision). Aliquots of bacterial cultures (25 ml) were harvested by centrifugation at 4°C for 10 min at 4,000 rpm. The bacterial pellets were washed twice with 1 ml of PBS and resuspended in 0.5 ml of PBS, followed by the addition of 0.1 ml of ice-cold 3 M perchloric acid. The cells were lysed using lysing matrix B tubes in a FastPrep instrument (Qbiogene). The lysates were centrifuged at 4°C for 3 min at 14,000 rpm. Subsequently, 300 µl of supernatants was neutralized with 75 µl of saturated solution of potassium bicarbonate and centrifuged at 4°C for 3 min at 14,000 rpm. Ac-CoA concentrations were determined according to the manufacturer’s protocol and normalized to the corresponding colony counts.

Intracellular pyruvate concentrations were determined using the Pyruvate Assay Kit (MBL). Aliquots of bacterial cultures (10 ml) were harvested by centrifugation at 4°C for 10 min at 4,000 rpm. The bacterial pellets were washed twice with 1 ml of PBS and resuspended in 0.35 ml of pyruvate assay buffer, incubated for 20 min at 80°C, and lysed using lysing matrix B tubes in a FastPrep instrument (Qbiogene). The lysates were centrifuged at 4°C for 5 min at 14,000 rpm. Pyruvate concentrations were determined according to the manufacturer’s protocol and normalized to the corresponding viable-cell counts at the time of harvest. All assays were performed in duplicate for three independent experiments.

Measurement of oxygen consumption. Cells were cultured at 37°C in TSB supplemented with 0.25% glucose and aerated at 250 rpm with a flask-to-medium ratio of 10:1. After 3 h of growth, samples were collected and diluted in air-saturated TSB supplemented with 0.25% glucose to an OD₆₀₀ of 0.1 to 0.8. Relative oxygen consumption rates were determined for up to 120 min at 37°C by using a MitoXpress oxygen-sensitive probe (Qbiogene) according to the manufacturer’s instructions. The data were normalized to the corresponding viable-cell counts at the time of harvest. The results were recorded in duplicate for three independent experiments.

Flow cytometry. Bacterial cell viability was measured by using a LIVE/DEAD BacLight Viability Kit (Invitrogen), according to the manufacturer’s protocol, on a FACSCanto flow cytometer (BD Biosciences). Aliquots of bacterial cultures (1 ml) were harvested by centrifugation at 4°C for 10 min at 4,000 rpm. The bacterial pellets were washed twice with 1 ml of PBS. The bacteria (1 × 10⁷ to 5 × 10⁸ cells per ml) were then stained for 15 min with SYTO 9 and a propidium iodide dye mixture. Bacteria were distinguished from the background using a combination of forward-scatter and side-scatter parameters. In total, 50,000 events were collected for each sample, and fluorescent signals were measured using logarithmic amplifications. Flow cytometric measurements were performed with 488-nm excitation from a blue solid-state laser at 50 mW, and fluorescent emissions in the green and red spectra were detected using 530- and 610-nm band-pass filters, respectively. Data were analyzed with FlowJo software (Tree Star, Inc.). The results are representative of duplicates for three independent experiments.

NMR sample preparation. Samples for two-dimensional (2D) [¹H,¹³C] heteronuclear single quantum coherence (HSQC) spectra were prepared from three independent 50-ml cultures of S. aureus strains UAMS-1, UAMS-1-ackA, and UAMS-1-pta at the exponential growth phase (3 h) on TSB containing 0.25% [¹³C]glucose (Sigma-Aldrich). Bacteria (10 OD₆₅₀ units) were harvested by centrifugation, suspended in 700 µl of quenching solution (60% ethanol, 40% D₂O [Isotec], 50 mM phosphate buffer, pH 7.1), and lyced using lysing matrix B tubes in a FastPrep instrument (Qbiogene). To remove the cell debris, the lysates were centrifuged at 4°C for 5 min at 14,000 rpm. The samples were then lyophilized and suspended in a 99.8% D₂O (Isotec), 50 mM phosphate buffer and used for nuclear magnetic resonance (NMR) analysis.

NMR data collection and analysis. All cell lysates were stored in a −80°C freezer until the samples were utilized for NMR experiments. Each sample was thawed completely and allowed to equilibrate at 298.15 K for approximately 1 h. Then, 600 µl of deuterated 50 mM potassium phosphate buffer at pH 7.4 (uncorrected) with 500 µM 3(trimethylsilyl)propionic-2,2,3,3-d4 sodium salt (TMSp) was added to the cell lysates and transferred to an NMR tube. A Bruker Avance DRX 500-MHz spectrometer equipped with a 5-mm triple-resonance cryoprobe (H, ¹H, and ¹³C) with a z-axis gradient was utilized for all 2D [¹H-¹³C] HSQC NMR experiments. NMR spectra were acquired in random order, which included alternating between the three S. aureus strains. Also, a BACS-120 sample changer with Bruker Icon software was used to automate the NMR data collection and minimize instrument variations. The 2D [¹H-¹³C] HSQC spectra were collected at 298.15 K with 128 scans, 16 dummy scans, and a relaxation delay of 1.5 s. The spectra were acquired with a spectrum width of 5,000 Hz and 2,048 data points in the direct dimension and a spectrum width of 17,605.6 Hz and 64 data points in the indirect dimension. The NMR spectra were processed and analyzed using NMRPipe (35) and NMRView (36) and following the protocol described previously (37). Briefly, peaks were assigned to a metabolite by matching ¹H and ¹³C chemical shifts from the Human Metabolome Database (HMDB) (38), using error tolerances of 0.08 ppm and 0.25 ppm, respectively. The presence of metabolites and metabolic pathways within S. aureus was verified with the KEGG (39) and Metacyc (40) databases.

Peak intensities were normalized by setting the maximal peak intensity to 100. Peak intensities for a specific metabolite were then averaged for each individual 2D [¹H-¹³C] HSQC spectrum and subsequently across the set of triplicate spectra. To account for differences in the numbers of viable cells between the S. aureus strains, the peak intensities were scaled by 2.5 for the UAMS-1-ackA and UAMS-1-pta strains. This corresponds to the ratio of CFU to that of the UAMS-1 wild-type strain. The peak intensities from the S. aureus UAMS-1-ackA and UAMS-1-pta mutant strains were then compared to the peak intensities from the UAMS-1 wild-type strain, where metabolite concentration changes were reported as a percentage change relative to the wild type.

Quantification of the mRNA transcripts. RNA isolation was carried out as described previously (29). Quantitative real-time PCR was performed using the sigA-, pfkA-, citZ-, cida-, ldhI-, ddh-, and als-specific primers listed in Table S1 in the supplemental material. Total RNA (500 ng) was converted to cDNA using the Quantitect Reverse Transcription Kit (Qiagen). The samples were then diluted 1:50, and the cDNA products were amplified using the LightCycler DNA Master SYBR green I kit (Roche Applied Science) following the manufacturer’s protocol. The relative transcript levels were calculated using the comparative threshold cycle (Ct) method (41) with normalization to the amount of sigA transcripts present in the RNA samples. The results were recorded in duplicate and are representative of three independent experiments. The statistical significance of changes between wild-type and mutant strains was assessed using several statistical tests. The Shapiro-Wilk test was used to check the normality of the data. Levene’s test was used to test the equality of variances among groups in comparison. Since the values were normally distributed, based on the Shapiro-Wilk test, and some groups in comparison have unequal variances, analysis of variance (ANOVA) with heterogeneous variance was used to compare the values between the ackA or pta mutants and the wild-type strain for experiments with certain primers.
separately. Dunnett’s method was used to account for multiple comparisons. A $P$ value of $\leq 0.05$ was considered statistically significant.

RESULTS

Inactivation of the Pta-AckA pathway inhibits growth of *S. aureus*. To examine the impact of the Pta-AckA pathway on *S. aureus* growth and carbon and energy metabolism, as well as CidR-mediated regulation, we constructed *ackA::ermB* and *pta* mutations in the *S. aureus* strain UAMS-1 (see Materials and Methods).

Inactivation of the Pta-AckA pathway by disrupting either the *ackA* or *pta* gene had a drastic inhibitory effect on growth rates during the exponential phase in both mutants compared to the isogenic wild-type strain (Fig. 1A). The observed growth defects were accompanied by lower acidification of the culture media (Fig. 1A) and a marked decrease in the rate and concentrations of acetic acid accumulation in both mutants (Fig. 1B). Additionally, the decreased growth rates of the *ackA* and *pta* mutants were also reflected in the reduced temporal depletion of glucose (Fig. 1C). However, when the concentrations of glucose in the culture media were plotted as a function of growth, the rates of glucose consumption during the exponential phase in both mutants compared to the isogenic wild-type strain (Fig. 1A). The observed growth defects were accompanied by lower acidification of the culture media (Fig. 1A) and a marked decrease in the rate and concentrations of acetic acid accumulation in both mutants (Fig. 1B). Additionally, the decreased growth rates of the *ackA* and *pta* mutants were also reflected in the reduced temporal depletion of glucose (Fig. 1C). However, when the concentrations of glucose in the culture media were plotted as a function of growth, the rates of glucose consumption during the exponential phase were found to be higher in the *ackA* and *pta* mutants (Fig. 1D; see Fig. S1 in the supplemental material), suggesting carbon is redirected from cell growth to other metabolic pathways and cellular processes. To exclude second-site mutations as the cause of these phenotypes, we performed a complementation study using plasmids pWS5 and pMRS102, containing the wild-type alleles of the *ackA* and *pta* genes, respectively, and observed restoration of the growth characteristics in the mutants to wild-type levels (see Fig. S2 in the supplemental material).

Disruption of the Pta-AckA pathway causes cell death. During growth experiments, we noticed that the viable-cell count differences observed for the wild-type and mutant strains seemed to be disproportionately high (unpublished data) compared to the differences observed for their optical densities (Fig. 1A). As the optical-density values reflect the total number of cells, including both live and dead cells, we reasoned that during growth the mutants may undergo increased cell death. Indeed, the number of viable cells (CFU) per OD$_{600}$ unit for the wild-type strain at the exponential phase of growth was approximately 2.5 times higher than that for the mutants (Fig. 2A). The observed accumulation of dead cells was not specific to the UAMS-1 strain, as inactivation of the *ackA* gene in strains JE2 and SA564 caused a similar decrease in the number of viable cells per OD$_{600}$ unit (see Fig. S3 in the supplemental material). To estimate the number of dead cells in the wild-type and mutant strains during exponential growth, we performed flow cytometry analysis of cells treated with the LIVE/DEAD staining dyes. As anticipated, the numbers of accumulated dead cells in the mutants were nearly 2 orders of magnitude higher than in the wild-type strain (Fig. 2B). These results indicate that the activity of the Pta-AckA pathway in *S. aureus* is critical for cell viability during overflow metabolism.

Pta-AckA pathway inactivation does not decrease intracellular ATP pools. During aerobic growth in the presence of excess glucose or other rapidly metabolizable carbon sources, the TCA cycle in staphylococci is subject to carbon catabolite repression (6, 10, 14–16). To support growth under these conditions, ATP is generated through substrate level phosphorylation via glycolysis.
and the Pta-AckA pathway (8, 17). The net yield of ATP directly generated by these pathways is four molecules per molecule of glucose consumed, where two molecules of ATP are produced by the Pta-AckA pathway. Hence, we speculated that the observed negative effect on bacterial growth and viability following inactivation of the Pta-AckA pathway may potentially result from a depleted energy status of the cell. To test this hypothesis, we determined the intracellular ATP concentrations in the ackA and pta mutants at the exponential phase of growth. Surprisingly, the intracellular concentrations of ATP were significantly higher in the mutants than in the wild-type strain (Fig. 3A), suggesting that the bacteria efficiently meet their cellular energy requirements by relying on other metabolic pathways.

**Inactivation of the Pta-AckA pathway alters the metabolic status of** **S. aureus.** We considered two main possibilities that might explain the increased ATP pools in the ackA and pta mutants: (i) the mutants have increased glucose uptake and carbon flow through the glycolytic machinery and (ii) the mutants have a partial redirection of carbon into the TCA cycle. Enhanced carbon flow through both glycolysis and the TCA cycle.

As a direct measure of the effect that disruption of the Pta-AckA pathway has on carbon flow through glycolysis and the TCA cycle, we performed an NMR-based metabolomic analysis of the wild-type and mutant strains at the exponential growth phase. Consistent with previous studies of the Pta-AckA pathway in other organisms (42, 43), this analysis revealed decreased and increased levels of acetyl phosphate in the mutants, as well as the relative concentrations of the TCA cycle intermediates, as well as metabolites associated with its activity (succinate, α-ketoglutarate, and glutamate concentrations were higher in the mutants) (Fig. 4A and B). In addition, evidence for carbon overflow into the pentose phosphate pathway (PPP) and various biosynthetic pathways, including amino acid, nucleotide, peptidoglycan, and polysaccharide biosynthesis (Fig. 4A and B), was detected. Overall, these results showed that the metabolic block caused by inactiva-
tion of the Pta-AckA pathway leads to redirection of carbon flux into diverse metabolic pathways and thus alters the metabolic status of bacteria.

Disruption of the Pta-AckA pathway causes a metabolic block at the pyruvate node and activates the CidR regulon. As described above, we observed significant perturbations in the catabolic fate of glucose in the mutants relative to the wild-type strain. In addition to the increased concentrations of glycolytic and TCA cycle intermediates in the mutants, and the down arrows correspond to a relative decrease in the metabolite concentration. Statistical significance at the 90% confidence level (P < 0.10) is denoted by asterisks above the arrows. Acetyl-P, acetyl-phosphate; (1,3)BP-G, 1,3-bisphosphoglycerate; F6-P, fructose 6-phosphate; FB-P, fructose 1,6-bisphosphatase; DHAP, dihydroxyacetone phosphate; G6-P, glucose 6-phosphate; G1-P, glucose 1-phosphate; GlcN-1-P, glucosamine-1-phosphate; GlcN-6-P, glucosamine-6-phosphate; GlcN-1-P, glucosamine-1-phosphate; GA3-P, glyceraldehyde 3-phosphate; GlcN-1-P, glucosamine-1-phosphate; UDP-GlcNAc, UDP N-acetylglucosamine; X3-P, xylulose 5-phosphate.

Unlike E. coli, where a pta mutation led to excretion of D-lactate (20), inactivation of the Pta-AckA pathway in S. aureus has only a...
minor effect on the generation of \( L \)- and \( D \)-lactate (Fig. 4; see Fig. S4A in the supplemental material). Moreover, quantitative real-time PCR analysis revealed a decrease in the relative \( ldh1 \) and \( ddh \) transcript levels (see Fig. S4B in the supplemental material), suggesting that inactivation of the Pta-AckA pathway in \( S. aureus \) did not redirect carbon into these fermentative pathways. An alternate route of pyruvate redirection is through the CidR-regulated AlsSD and CidC pathways (24). Given the observation that the \( S. aureus \) CidR regulator responds to growth under conditions of excess glucose, it is possible that the \( pta \) and \( ackA \) mutations affect the \( cid \) and \( als \) operons. Thus, to determine if inactivation of the Pta-AckA pathway has an effect on \( cidABC \) and \( alsSD \) expression, we performed quantitative RT-PCR analysis, using primers specific to \( cidA \) and \( alsS \). Indeed inactivation of either \( pta \) or \( ackA \) resulted in a robust increase of both \( cidA \) and \( alsS \)-specific transcripts (Fig. 5D). The transcriptional upregulation of the \( alsSD \) and \( cidABC \) operons in the \( pta \) and \( ackA \) mutants was also reflected by a significant increase in the levels of accumulated acetoin in the medium (Fig. 5E) and by increased generation of acetate via pyruvate oxidase, CidC (Fig. 5F).

**DISCUSSION**

Overflow metabolism in \( S. aureus \) occurs during aerobic growth in the presence of rapidly metabolizable carbon sources, when glycolytic flux exceeds the capacity of the TCA cycle and results in the production of acetate. It has been proposed that substrate level phosphorylation during acetate metabolism (catalyzed by AckA) is an important secondary energy-yielding pathway, since under carbon overflow conditions, TCA cycle activity is limited by carbon catabolite repression (the Crabtree effect) (8, 22, 45–48). Indeed, introduction of a metabolic block (mutation of either \( pta \) or \( ackA \)) that hinders substrate level phosphorylation during overflow metabolism gave rise to growth defects and increased the rate of cell death during exponential growth. However, these deleterious effects on growth and viability did not stem from depleted energy reserves of the cell, as both of the isogenic \( pta \) and \( ackA \) mutants had comparable or even elevated levels of ATP relative to that of the wild-type strain. Interestingly, similar results were also reported for \( E. coli \), where inactivation of the \( pta \) gene resulted in a large growth defect but did not lead to a substantial loss in the amounts of generated ATP (90.5% of the total energy generated by the wild-type strain) (20).

How do the \( S. aureus pta \) and \( ackA \) mutants replenish their ATP pools during overflow metabolism? Several lines of evidence suggest that the energy derived from an increased rate of glycolysis combined with an increased flux through the TCA cycle may compensate for the loss of ATP following inactivation of the Pta-AckA pathway. First, the rate of glucose uptake was higher for both the \( pta \) and \( ackA \) mutants than for the wild-type strain, supporting a higher glycolytic flux in these mutants. Second, quantitative RT-
PCR analyses confirmed the increased expression of both phosphofructokinase (a critical glycolytic enzyme) and citrate synthase (the first enzyme of the TCA cycle that catalyzes the production of citrate). Finally, metabolomic profiles of both the pta and ackA mutants not only demonstrated an abundance of multiple glycolytic and TCA cycle intermediates, but also indicated carbon overflow into peripheral pathways that are directly linked to central metabolism.

Despite achieving ATP homeostasis, the observation that the pta and ackA mutants still display a growth defect or undergo increased cell death remains a paradox. However, based on our analyses, several possibilities exist. Foremost among them is the drastic increase in the expression of CidR-regulated operons, i.e., cidABC and allsD. Although CidC and AlsSD represent overflow metabolic enzymes that may redirect excess pyruvate, their transcriptional coupling to the CidAB-holin-like proteins that have been implicated in *S. aureus* cell death is noteworthy. Although the effects of CidAB overexpression in the pta and ackA mutants are yet to be characterized, their proposed role as effectors of cell death by oligomerizing and localizing to the *S. aureus* cytoplasmic membrane (49) makes this an intriguing possibility.

A second possibility that may account for the observed growth and viability defects of pta and ackA mutants involves intracellular metabolic toxicity that may arise from increased pyruvate and/or acetyl-CoA levels (Fig. 5A and B). Chang et al. proposed that the growth defect observed for the *E. coli* pta mutant resulted from the unbalanced flux of acetyl-CoA (20). Accumulation of acetyl-CoA caused inhibition of the pyruvate dehydrogenase complex, leading to increased concentrations of intracellular pyruvate and excretion of D-lactate and pyruvate into the medium (20). Recently, it was shown that triphenylbismuthdichloride (TPBC) suppresses growth and blocks pyruvate catabolism in *S. aureus* by specific inhibition of the PDHC (50). Using exometabolome analysis, Birkenstock et al. showed that the treatment of cells with TPBC led to an accumulation of pyruvate, reduced temporal depletion of glucose, and decreased acetate excretion (50). Although, the growth defect observed after TPBC treatment may result from decreased intracellular pools of acetyl-CoA and carbon flux into the Pta-AckA pathway and TCA cycle or associated effects on protein acetylation (51–53), a negative impact of the high levels of accumulated pyruvate on bacterial growth cannot be excluded from consideration.

Finally, fluctuations in the redox status or the generation of reactive oxygen species (ROS) as a result of enhanced oxidative phosphorylation in the pta and ackA mutants may yet constitute another reason for the observed deleterious growth characteristics. In support of this, determination of the oxygen consumption rates revealed increased levels of respiration in the mutants (Fig. 3D). Moreover, quantification of intracellular NAD⁺ and NADH demonstrated significantly higher concentrations of both species in the mutants (Fig. 3C and 4A), suggesting enhanced de novo biosynthesis of NAD⁺. Consistent with this observation, NMR-based metabolomics detected carbon redirection toward the nucleotide biosynthesis and pentose phosphate pathway that could generate precursors for NAD⁺ synthesis. Increased NADH levels could potentially result in the increased respiration observed for the mutants and generation of ROS (54–56).

Considering that inactivation of the Pta-AckA pathway alters the metabolic status of bacteria and has deleterious effects on bacterial growth and cell viability, our findings provide evidence that the activity of this pathway in *S. aureus* is critical under conditions of overflow metabolism. Although the exact mechanisms leading to cell death in the pta and ackA mutants still remain to be clarified, our data suggest that perturbations in the concentration of intracellular acetyl phosphate (Fig. 4A), which can affect two-component signal transduction systems and other cellular processes (22, 57–60), are not a primary cause of growth inhibition and cell death in the mutants, as disruption of either the pta or ackA gene had similar effects on bacterial growth and cell viability. Furthermore, inactivation of the Pta-AckA pathway did not decrease the energy status of bacteria; however, it did alter the metabolic state by causing a metabolic block at the pyruvate node, as well as an increase in carbon flux through both glycolysis and the TCA cycle.

Overall, the results of this study provide important insight into the Pta-AckA pathway in *S. aureus*, which is critical for acetate dissimilation (22). This pathway could also affect biofilm development, considering previous evidence suggesting acetyl phosphate functions as a global signal within a biofilm (59). Thus, further studies of this pathway will expand our understanding of the physiological processes that underlie the colonization and invasive mechanisms critical for staphylococcal infection.

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