

# Regulation of tricarboxylate transport and metabolism in *Acinetobacter baylyi* ADP1

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**ABSTRACT** Despite the significant presence of plant-derived tricarboxylic acids in some environments, few studies detail the bacterial metabolism of *trans*-aconitic acid (Taa) and tricarballic acid (Tcb). In a soil bacterium, *Acinetobacter baylyi* ADP1, we discovered interrelated pathways for the consumption of Taa and Tcb. An intricate regulatory scheme tightly controls the transport and catabolism of both compounds and may reflect that they can be toxic inhibitors of the tricarboxylic acid cycle. The genes encoding two similar LysR-type transcriptional regulators, TcuR and TcIR, were clustered on the chromosome with *tcuA* and *tcuB*, genes required for Tcb consumption. The genetic organization differed from that in *Salmonella enterica* serovar Typhimurium, in which *tcuA* and *tcuB* form an operon with a transporter gene, *tcuC*. In *A. baylyi*, *tcuC* was not cotranscribed with *tcuAB*. Rather, *tcuC* was cotranscribed with a gene, designated *pacl*, encoding an isomerase needed for Taa consumption. TcuC appears to transport Tcb and *cis*-aconitic acid (Caa), the presumed product of Pacl-mediated periplasmic isomerization of Taa. Two operons, *tcuC-pacl* and *tcuAB*, were transcriptionally controlled by both TcuR and TcIR, which have overlapping functions. We investigated the roles of the two regulators in activating transcription of both operons in response to multiple effector compounds, including Taa, Tcb, and Caa.

**IMPORTANCE** Ingestion of Taa and Tcb by grazing livestock can cause a serious metabolic disorder called grass tetany. The disorder, which results from Tcb absorption by ruminants, focuses attention on the metabolism of tricarboxylic acids. Additional interest stems from efforts to produce tricarboxylic acids as commodity chemicals. Improved understanding of bacterial enzymes and pathways for tricarboxylic acid metabolism may contribute to new biomanufacturing strategies.

**KEYWORDS** ADP1, *Acinetobacter baylyi*, TcuR, aconitate isomerase, LTTR, LysR, transcriptional regulator, tricarboxylic acids, tricarballic acid, aconitate

Tricarboxylic acids, produced by grasses and legumes, can be relatively abundant in plant-based forage. These compounds, such as tricarballic acid (Tcb) and *trans*-aconitic acid (Taa), have been of interest because of problems that occur when grazing ruminants ingest them. Tcb in the rumen gets absorbed by the animal wherein the chelation properties of Tcb can cause a serious magnesium deficiency that manifests as a malady called grass tetany (1). This problem, which can be fatal to livestock, was at one time attributed to Taa but later shown to result from the conversion of Taa to Tcb by microbial fermentation in the rumen (2, 3).

The biochemical basis for the bacterial uptake and use of Tcb as a sole carbon source has, to date, been reported for only *Salmonella enterica* subspecies *enterica* serovar Typhimurium LT2 (hereafter LT2; 4–7). Fewer studies have been published on the microbial consumption of Taa as a growth substrate. The only biochemical studies center

**Editor** Isaac Cann, University of Illinois Urbana-Champaign, Champaign, Illinois, USA

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The authors declare no conflict of interest.

See the funding table on p. 17.

**Received** 27 November 2023

**Accepted** 13 December 2023

**Published** 30 January 2024

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on a constitutively expressed periplasmic aconitate isomerase involved in Taa assimilation by *Pseudomonas* sp. WU-0701 (8, 9). Further information about Taa assimilation or transport has not been reported. Interest in this aconitate isomerase stems from biomanufacturing because tricarboxylic acids are important commodity chemicals that have the potential to be synthesized by bacteria (10, 11). Heterologous expression of the aconitate isomerase was used in a two-step bioconversion of citric acid (Cit) to *trans*-aconitic acid (Taa) by whole bacterial cells (12).

As current understanding of these pathways remains incomplete, we characterized the consumption of *cis*-aconitic acid (Caa), Taa, Tcb, and Cit by a soil bacterium, *Acinetobacter baylyi* ADP1. This metabolically versatile bacterium was chosen because its environmental habitat is replete with plant-derived growth substrates. Moreover, ADP1 has a uniquely powerful genetic system that facilitates investigation using methods to exploit its exceptionally high efficiency of natural transformation and homologous recombination. Hypotheses concerning metabolism can be readily tested with mutational approaches. This soil bacterium is gaining traction as a model organism for metabolic engineering (13, 14).

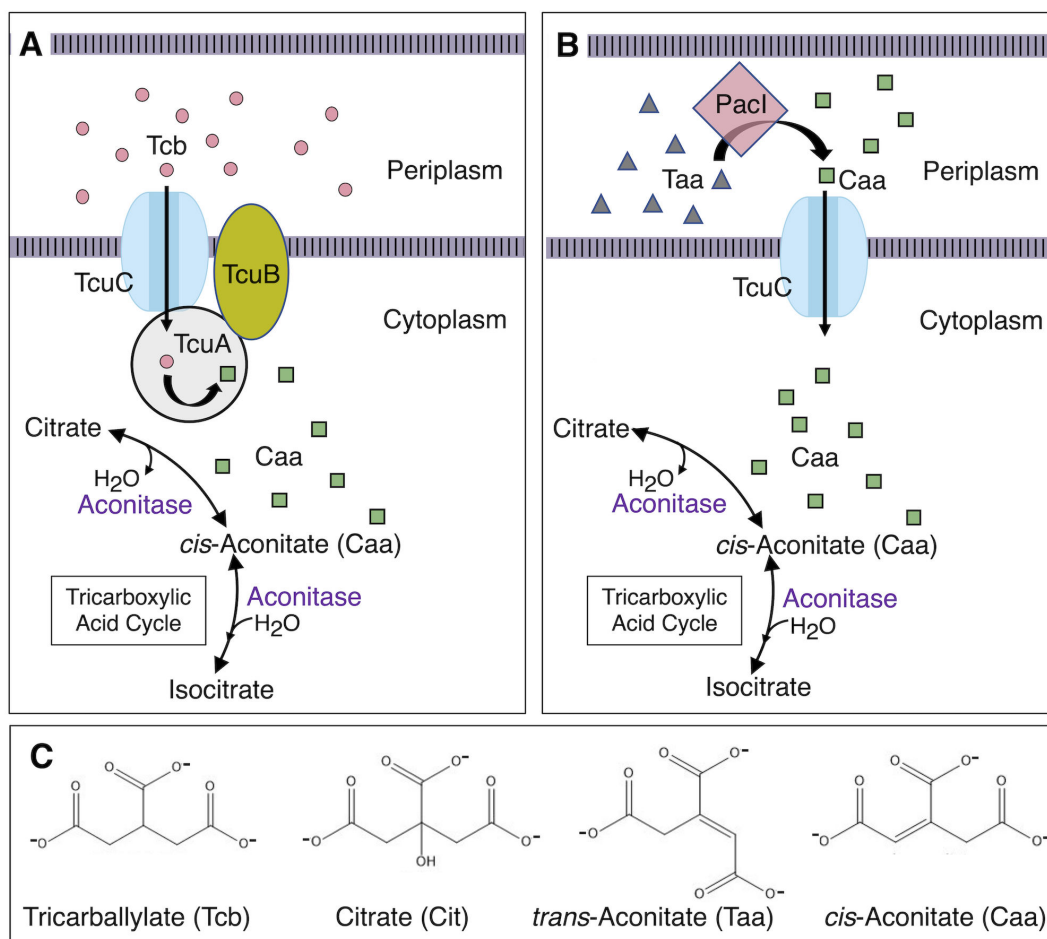
Genome sequence and a global metabolic model of ADP1 (15) identified genes for Tcb catabolism based on homology to genes characterized in LT2, *tcuABC*, and *tcuR* (4–7). Studies of the Tcu proteins in LT2 demonstrate that TcuA is an FAD-dependent Tcb dehydrogenase and TcuB is a transmembrane protein that re-oxidizes the flavin cofactor of the dehydrogenase (Fig. 1; 6, 7). In LT2, *tcuC*, which encodes a Tcb transporter, is cotranscribed with *tcuA* and *tcuB*. An adjacent regulatory gene, *tcuR*, encodes a LysR-type transcriptional regulator (LTTR) that activates the *tcuABC* operon in response to Tcb (5). LT2 does not use Taa as a sole carbon source, and the consumption of this compound by ADP1 was not previously determined.

The *tcu* gene order in ADP1 is similar but not identical to that of LT2 (Fig. 2). Notably, *tcuC* is not adjacent to *tcuA* and *tcuB*. As described in this report, we investigated the roles of two *tcuR* paralogs near *tcuA* in the regulation of tricarboxylate metabolism in ADP1 and renamed one *tc1R* (TcuR-like regulator). Studies of genetic regulation and function revealed that the uncharacterized gene immediately downstream of *tcuC* (locus tag ACIAD\_RS07095) in ADP1 enables the use of Taa as a sole carbon source. Based on the homology of its gene product to the aconitase of *Pseudomonas* sp. WU-0701 (8, 9), we designated this gene *pacI* (for periplasmic aconitate isomerase). Here, we present a novel metabolic and regulatory scheme whereby the growth of ADP1 on Taa involves the periplasmic PacI-mediated conversion of Taa to Caa followed by the TcuC-mediated transport of Caa into the cytoplasm (Fig. 1B). The TcuC transport protein appears to have dual specificity for Caa and Tcb.

## RESULTS

### Organization and bioinformatic analysis of clustered *tcu* genes in *A. baylyi* ADP1

Database searches revealed many bacterial homologs of TcuA, TcuB, TcuC, and TcuR that shared sequence similarity comparable to that between the gene products in ADP1 and LT2 (Table S1). However, the gene downstream of *tcuC* in ADP1 has no counterpart in LT2, and we questioned its role. Genome annotation and homology searches suggested only that this gene, ACIAD\_RS07095, encodes a substrate-binding periplasmic transport protein of unknown function. We posited that if ACIAD\_RS07095 homologs clustered with *tcu* genes in diverse bacteria, it might signify that this gene has a function related to Tcb metabolism. To identify genomic synteny among bacteria with *tcu* genes near homologs of ACIAD\_RS07095, chromosomal neighborhoods near *tcuB* were searched with the online STRING program (20). Regions in diverse bacteria that might be of interest were examined individually, and several different genetic arrangements were found with *tcuAB* homologs in the vicinity of homologs of the gene of unknown function from ADP1 (Fig. 2).

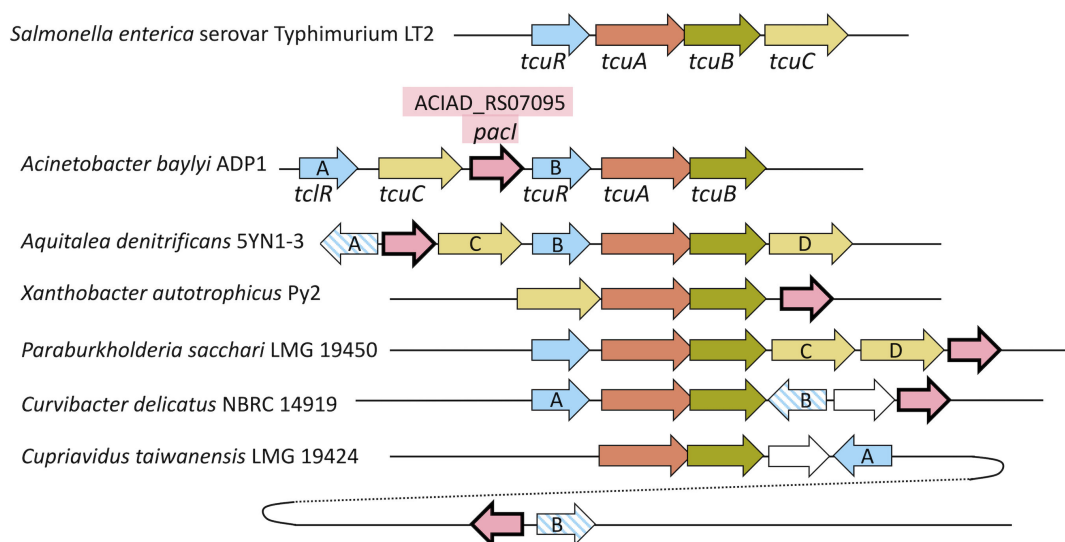


**FIG 1** Tcb, Taa, and Caa utilization. (A) The functions of TcuA and TcuB in converting Tcb (circles) to Caa (squares) and the role of TcuC in Tcb transport, which were demonstrated in strain LT2 (4–7), are predicted to be the same in ADP1. (B) In ADP1, our results support a model in which Pacl, which does not have a counterpart in LT2, converts Taa (triangles) to Caa (squares) in the periplasm, prior to transport into the cytoplasm by TcuC. (C) Chemical structures are shown for tricarboxylic acids investigated in this report.

The protein products of the homologs of unknown function were aligned, and conserved sequences guided additional homology searches. An iterative approach revealed homology to the aconitate isomerase of *Pseudomonas* sp. WU-0701 (9), as shown in Fig. S1. The genomic context in strain WU-0701 was unclear because surrounding DNA sequences are unavailable. However, a protein encoded by *Pseudomonas oryzae* was found to differ from that of WU-0701 by a single amino acid (P or S at position 202, Fig. S1). There were no *tcu* genes in the genome of *P. oryzae* and no LTR genes close to this putative aconitate isomerase gene. Nevertheless, we explored the possibility that Tcb and aconitate (Taa and Caa) metabolism were linked in ADP1. We hypothesized that ACIAD\_RS07095 encodes an aconitate isomerase, designated Pacl, that converts Taa to Caa. Periplasmic localization was predicted by SignalP analysis (21). This program identified the first 24 amino acids of Pacl as a cleavable signal used to export the enzyme. Therefore, our model depicts the production of Caa from Taa in the periplasm followed by transport into the cytoplasm (Fig. 1B).

### Effects of gene-specific mutation: growth on tricarboxylic acids as sole carbon sources

We tested whether genes in the *tcu* region of ADP1 affect the regulation and metabolism of tricarboxylic acids after constructing mutants with specific deletions (described in



**FIG 2** Chromosomal regions from diverse bacteria (drawn to scale), aligned relative to *tcuB*. Homologs, shown in the same color and pattern, were aligned using SyntTax and BioCyc programs (16, 17). Similarity searches based on *pacI* (ACIAD\_RS07095) revealed four bacteria in which homologs (in pink) were clustered in the vicinity of *tcuB* homologs. In *Cupriavidus taiwanensis* LMG 19424, a *pacI* homolog is adjacent and divergent to an LTR gene, but it is distal to the *tcu*-like region. In this case, *tcuA* is at map position 1,262,684–1,264,114, and the *pacI* homolog (RALTA\_RS24780) is at map position 2,072,478–2,073,302 (NCBI Reference Sequence: NC\_010530.1). Sequence identifiers for genomes are listed in Materials and Methods. Labels A and B (arbitrarily assigned) indicate two LTR paralogs. Striped patterns mark three LTR genes that are more like each other in sequence than they are to the solid-colored LTR genes (Fig. S2 and S3). Labels C and D (arbitrarily assigned) denote two *tcuC*-like paralogs in the same region predicted to encode transporters in the major facilitator superfamily (MFS), entry IPR011701 of InterPro (18); their sequences are aligned in Fig. S5. In each bacterium, the *tcuA* and *tcuB* genes have overlapping stop and start codons, a configuration that may couple translation to produce equal levels of both interrelated proteins (19).

Table 1; Table S2 through S4). These strains were compared with ADP1 for growth using Cit, Tcb, Taa, Caa, or pyruvate as the carbon source. ADP1 grows well on pyruvate, unlike sugars, and pyruvate is typically used as a growth substrate. The growth rates of ADP1 and all mutants on pyruvate were comparable (Table 1). Isocitrate was also tested, but no growth was observed.

Deletion of *pacI* (in ACN1388) prevented growth on Taa as the carbon source but not growth on Tcb or Caa (Taa<sup>-</sup>, Tcb<sup>+</sup>, and Caa<sup>+</sup>). In contrast, a *tcuA* mutant (ACN3123) was Taa<sup>+</sup> and Caa<sup>+</sup> but did not grow on Tcb (Table 1). These results support a model where TcuA is required to grow on Tcb (Fig. 1A), and the loss of Pacl only prevents growth on Taa (Fig. 1B). A  $\Delta$ *tcuC* mutant (ACN1456) was Tcb<sup>-</sup>, Taa<sup>+</sup>, and Caa<sup>+</sup> but grew on pyruvate and Cit (Table 1). These results are consistent with TcuC transporting Tcb, as in LT2 (4). Failure to grow on both compounds could be explained by a model wherein Taa is converted to Caa in the periplasm and TcuC transports Caa in addition to Tcb (Fig. 1B). Growth on Taa with Caa as an intermediate is supported by the observation that the four mutants that are Caa<sup>-</sup> are all also Taa<sup>-</sup> (Table 1).

Growth of the  $\Delta$ *tcuR* mutant (ACN1376) on Caa suggests that the TcuR regulator in ADP1 is not required for TcuC expression, unlike in LT2 where transcription of the *tcuABC* operon requires TcuR (5). Different gene organization also raises regulatory questions about the roles of two *tcuR* paralogs in ADP1, one designated *tcIR* (Fig. 2). The  $\Delta$ *tcIR* mutant (ACN3178) and the  $\Delta$ *tcIR* plus  $\Delta$ *tcuR* mutant (ACN3171) were both Caa<sup>-</sup> and Taa<sup>-</sup> (Table 1). Therefore, TcIR rather than TcuR may control TcuC expression in ADP1 during growth on these substrates.

Since growth on Tcb required *tcuR* (in ACN1376), TcuR may control the transcription of *tcu* genes, as does its counterpart in LT2. However, deletion of *tcIR* also affected growth on Tcb, leading to very slow growth and a long lag (ACN3178). When both *tcuR* and *tcIR* were deleted (ACN3171), no growth was observed with Tcb, Caa, or Taa as the carbon source. Furthermore, while the  $\Delta$ *tcuR* mutant (ACN1376) was Tcb<sup>-</sup>, it gave rise to

TABLE 1 Growth of ADP1 and mutants on different sole carbon sources

Strain	ADP1	ACN1388	ACN3225	ACN1456	ACN3168	ACN3123	ACN3178	ACN1376	ACN3171	
Genotype <sup>a</sup>	WT	$\Delta pacl$	$\Delta pacl$ $tcuC-gfp_{sf}$	$\Delta tcuC$	$\Delta tcuC$ $\Delta cltA$	$\Delta tcuA::gfp_{sf}$	$\Delta tclR$	$\Delta tcuR$	$\Delta tclR$ $\Delta tcuR$	
Carbon source	Parameter									
Pyr <sup>e</sup>	Growth rate <sup>b</sup>	1 ± 0.1	1.1 ± 0.1	1.2 ± 0.1	1 ± 0.1	1.2 ± 0.1	1.2 ± 0.2	1.1 ± 0.1	1.1 ± 0.1	1.2 ± 0.1
	Max OD <sup>c</sup>	1 ± 0.1	1.1 ± 0.2	1 ± 0.2	1 ± 0.1	1.1 ± 0.2	1.1 ± 0.2	.9 ± 0.2	1 ± 0.2	.9 ± 0.2
	Lag (h) <sup>d</sup>	2.25	2.75	3.25	2.75	2.75	3.25	2.25	2.25	2.75
Cit	Growth rate <sup>b</sup>	1.1 ± 0.2	1.1 ± 0.1	1.2 ± 0.2	1.4 ± 0.1	1.2 ± 0.2	1.2 ± 0.2	1.2 ± 0.1	1 ± 0.03	1.3 ± 0.2
	Max OD <sup>c</sup>	1 ± 0.2	1.1 ± 0.2	1.2 ± 0.2	1 ± 0.1	1 ± 0.2	1.1 ± 0.2	1 ± 0.1	1 ± 0.1	1 ± 0.1
	Lag (h) <sup>d</sup>	3.25	2.75	2.75	3.25	4.75	3.25	3.75	3.75	4.25
Tcb	Growth rate <sup>b</sup>	1 ± 0.1	1 ± 0.1	.2 ± 0.01	NG <sup>f</sup>	NG	NG	.2 ± 0.03	NG	NG
	Max OD <sup>c</sup>	.9 ± 0.1	.6 ± 0.1	.7 ± 0.1	NG	NG	NG	.5 ± 0.05	NG	NG
	Lag (h) <sup>d</sup>	7.75	10.25	18.75	NG	NG	NG	43.25	NG	NG
Taa	Growth rate <sup>b</sup>	.9 ± 0.1	NG	NG	NG	NG	1.1 ± 0.03	NG	1.1 ± 0.1	NG
	Max OD <sup>c</sup>	1 ± 0.2	NG	NG	NG	NG	1.1 ± 0.2	NG	1 ± 0.2	NG
	Lag (h) <sup>d</sup>	2.75	NG	NG	NG	NG	3.25	NG	2.75	NG
Caa	Growth rate <sup>b</sup>	1.3 ± 0.2	1.5 ± 0.1	1.26 ± 0.36	NG	NG	1.2 ± 0.1	NG	.9 ± 0.1	NG
	Max OD <sup>c</sup>	1.1 ± 0.2	1 ± 0.1	1.02 ± 0.14	NG	NG	1.1 ± 0.2	NG	1.1 ± 0.2	NG
	Lag (h) <sup>d</sup>	2.25	2.25	2.75	NG	NG	2.75	NG	2.25	NG

<sup>a</sup>Additional information on strain genotypes is provided in Table 2; Tables S2 to S4.

<sup>b</sup>Ratio of the growth rate of each strain on the indicated carbon source to that of ADP1 grown on pyruvate. A 95% CI calculated from three replicates is indicated by  $\pm$  values. The generation time of pyruvate-grown ADP1 is 1.3 h when grown in shaking tubes/flasks and 3.0 h when grown in the plate reader.

<sup>c</sup>Max growth assessed by turbidity (measured as OD<sub>595</sub>) in comparison to ADP1 grown on Pyr. A 95% CI calculated from three replicates is represented by  $\pm$  values.

<sup>d</sup>A graphing program (Prism, GraphPad Software) was used to calculate average growth rates across replicates at 30-min intervals. Lag time is here defined as the time taken for the growth rate to increase by  $\geq 0.01$  (absorbance unit/h), which corresponds to the beginning of the log phase. Standard deviations of lag times are <15% of the average values.

<sup>e</sup>Carbon sources were added to the minimal medium at the following final concentrations: 15-mM pyruvate (Pyr), 5-mM Cit, 5-mM Tcb, 5-mM Taa, and 5-mM Caa.

<sup>f</sup>No growth (NG) is defined as an increase in OD<sub>595</sub>  $\leq 0.05$  within 72 h following inoculation.

spontaneous Tcb<sup>+</sup> mutants. Two of these, which were isolated independently, ACN1445 and ACN1556, each encoded a TcIR variant, TcIR(R200Q, L216P) or TcIR(R200Q).

## Evaluation of *tcuA* regulation with a fluorescent transcriptional reporter

To clarify regulation, the *tcuA* coding sequence of ADP1 was replaced on the chromosome by the gene for a green fluorescent protein "super folder" (GFP<sub>sf</sub>; 24). Because this reporter strain (ACN3123) does not have TcuA, Tcb cannot be catabolized further and can be studied as a non-metabolizable inducer of GFP<sub>sf</sub> expressed from the *tcuA* promoter ( $P_{tcuA}$ ). Other reporter strains were similarly constructed with additional deletions of *tclR*, *tcuR*, or both (ACN3184, ACN3122, and ACN3173, respectively). The same chromosomal reporter was also studied in strains with *tcuR* deleted and the wild-type *tclR* coding sequence exactly replaced by a mutated allele encoding either TcIR(R200Q, L216P) or TcIR(R200Q), ACN3190, and ACN3191, respectively. Strains were grown on pyruvate with or without Tcb, Caa, or Taa as inducers. Compared to cultures grown without inducers, the fluorescence of the strain with both regulators (*tcuR*<sup>WT</sup>, *tclR*<sup>WT</sup>) increased when Tcb, but not Caa or Taa, was added (Fig. 3, solid dark bars).

With both regulators missing ( $\Delta tcuR$ ,  $\Delta tclR$ ), Tcb did not increase fluorescence. In contrast, TcuR without TcIR ( $\Delta tclR$ ) was sufficient for Tcb-increased fluorescence, albeit to a lesser extent than with both regulators. The loss of TcIR could reduce  $P_{tcuA}$ -controlled fluorescence if TcIR directly controls *tcuA* transcription. Alternatively, or in addition, loss of TcIR could affect fluorescence through effects on the transcription of *tcuC*, which could alter the transport of the inducer and affect transcription from  $P_{tcuA}$  indirectly. Unlike in a strain without both regulators ( $\Delta tcuR$ ,  $\Delta tclR$ ), there was a small increase in Tcb-dependent induction in a strain with TcIR alone ( $\Delta tcuR$ ; Fig. 3), suggesting that TcIR directly regulates Tcb-dependent *tcuA* transcription, but poorly. In strains encoding the TcIR variants ( $\Delta tcuR$ , *tclR*<sup>R200Q</sup> and  $\Delta tcuR$ , *tclR*<sup>R200Q, L216P</sup>), Tcb-increased fluorescence relative to no inducer and relative to the induction observed in the strain with only the wild-type

TABLE 2 *A. baylyi* strains<sup>a,b</sup>

Strain	Relevant characteristics	Source
ADP1	Wild-type strain (BD413)	(22, 23)
ACN1376	$\Delta tcuR51376$	This study
ACN1388	$\Delta pacI51388$	This study
ACN1445	$\Delta tcuR51376, tclR51445$	This study
ACN1456	$\Delta tcuC51456$	This study
ACN1556	$\Delta tcuR51376, tclR51556$	This study
ACN3122	$\Delta tcuR51376, \Delta tcuA::gfp_{sf}-\Omega K53122$	This study
ACN3123	$\Delta tcuA::gfp_{sf}-\Omega K53122$	This study
ACN3125	$\Delta cltA::sacB-\Omega S53125$	This study
ACN3168	$\Delta tcuC51456, \Delta cltA::sacB-\Omega S53125$	This study
ACN3171	$\Delta tcuR51376, \Delta tclR53171$	This study
ACN3173	$\Delta tcuR51376, \Delta tclR53171, \Delta tcuA::gfp_{sf}-\Omega K53122$	This study
ACN3178	$\Delta tclR53171$	This study
ACN3179	$\Delta tcuR51376, tclR51445$ [TclR(R200Q, L216P)]	This study
ACN3184	$\Delta tclR53171, \Delta tcuA::gfp_{sf}-\Omega K53122$	This study
ACN3187	$\Delta tcuR51376, tclR51556$ [TclR(R200Q)]	This study
ACN3190	$\Delta tcuR51376, tclR51445, \Delta tcuA::gfp_{sf}-\Omega K53122$ [TclR(R200Q, L216P)]	This study
ACN3191	$\Delta tcuR51376, tclR51556, \Delta tcuA::gfp_{sf}-\Omega K53122$ [TclR(R200Q)]	This study
ACN3222	$\Delta tcuR51376, tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$	This study
ACN3223	$\Delta tcuR51376, \Delta tclR53171, tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$	This study
ACN3224	$\Delta tclR53171, tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$	This study
ACN3225	$tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$	This study
ACN3226	$\Delta cltA::sacB-\Omega S53125, tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$	This study
ACN3227	$\Delta tcuR51376, tclR51445, tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$ [TclR(R200Q, L216P)]	This study
ACN3228	$\Delta tcuR51376, tclR51556, tcuC-gfp_{sf}-\Omega K53222, \Delta pacI53222$ [TclR(R200Q)]	This study

<sup>a</sup>For additional details about strain construction, see Tables S2–S4.

<sup>b</sup>*A. baylyi* strains were derived from ADP1, previously known as *Acinetobacter calcoaceticus* or *Acinetobacter* sp. (22, 23).

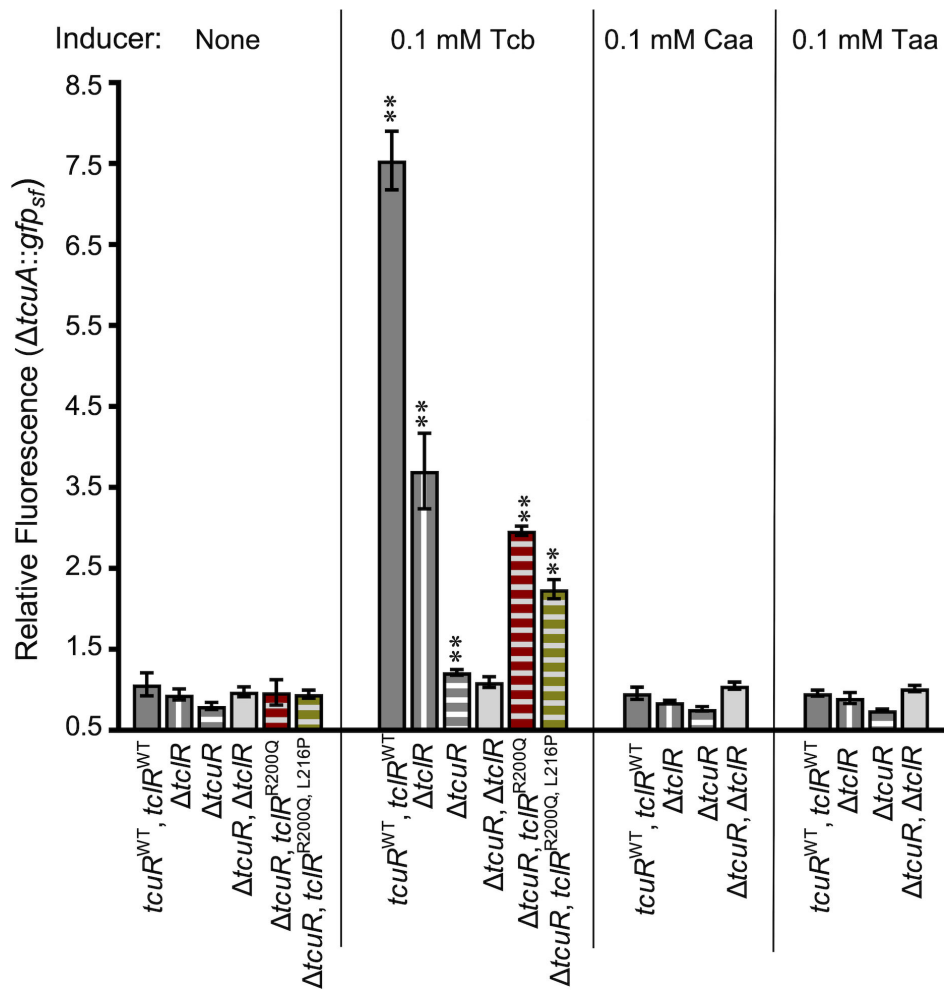
TclR (Fig. 3). This increase in Tcb-stimulated expression may account for the Tcb<sup>+</sup> phenotypes of the corresponding spontaneous mutants. These results indicate that TcuR is the primary transcriptional activator of *tcu* genes in response to Tcb, whereas TclR plays a lesser role.

### Evaluation of *tcuC* regulation with a fluorescent transcriptional reporter

To assess the regulation of the *tcuC* promoter,  $P_{tcuC}$ , the GFP<sub>sf</sub> gene was inserted in the chromosome immediately downstream of *tcuC*, close enough to ensure cotranscription. This location was designed to preserve the TcuC transport function. Additionally, *pacI* was deleted to prevent interconversion of Taa and Caa so their individual effects could be distinguished. The Tcb<sup>+</sup> phenotype of the resulting strain, ACN3225 (Table 1), indicated that TcuC was functional. However, this strain grew more slowly on Tcb and had a longer lag time than the wild type (ADP1). Additional strains were constructed with this *tcuC* transcriptional reporter and deletion of  $\Delta tclR$  or  $\Delta tcuR$  (ACN3224 and ACN3222) or deletions of both (ACN3223). Strains were also made with  $\Delta tcuR$  plus  $tclR^{R200Q}$  (ACN3228) or  $\Delta tcuR$  plus  $tclR^{R200Q, L216P}$  (ACN3227). With wild-type *tcuR* and *tclR*, but not without them ( $\Delta tcuR, \Delta tclR$ ), Tcb increased fluorescence (Fig. 4). Without TcuR, the TclR variants (with R200Q or with R200Q plus L216P) increased Tcb-dependent induction relative to wild-type TclR at both promoters. Yet, Tcb-dependent induction differed at  $P_{tcuA}$  (Fig. 3) and  $P_{tcuC}$  (Fig. 4). Response to Tcb at  $P_{tcuC}$  required higher concentrations than at  $P_{tcuA}$  (5 mM vs 0.1 mM) and caused a smaller increase in fluorescence (Fig. 3 and 4). The relative importance of TcuR and TclR for Tcb-regulated transcription appears to be reversed at  $P_{tcuC}$  and  $P_{tcuA}$ , with TclR being more important than TcuR for regulating  $P_{tcuC}$  (Fig. 4).

The addition of Caa and Taa increased fluorescence at  $P_{tcuC}$  (Fig. 4) but not  $P_{tcuA}$  (Fig. 3). With TclR and TcuR, but not without them, Taa (0.1 mM or 5 mM) or Caa (5 mM)





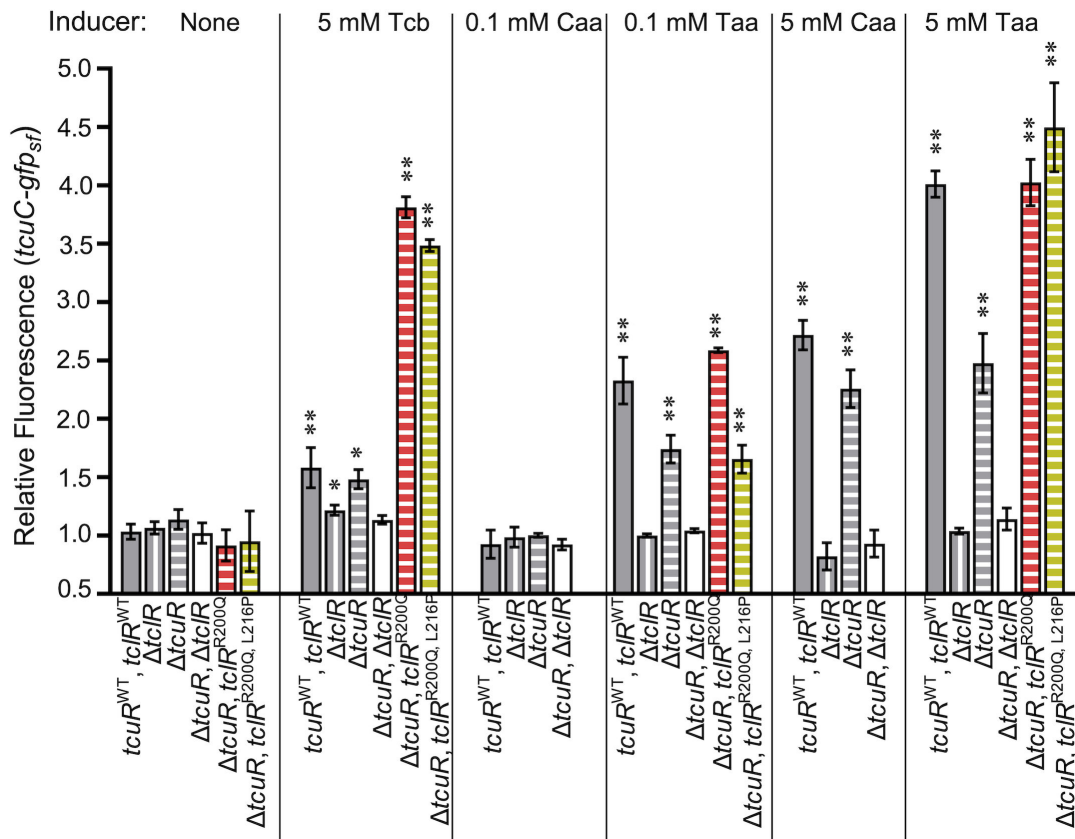
**FIG 3** Transcription of *gfp<sub>sf</sub>* controlled by  $P_{tcuA}$ . Each strain has a chromosomal *gfp<sub>sf</sub>* (allele  $\Delta tcuA::gfp_{sf}$ - $\Omega K53122$ ). Differences in genetic backgrounds are indicated. All strains were grown on pyruvate supplemented with inducer compounds as indicated. Relative fluorescence is the ratio of fluorescence/OD for each condition compared to that of ADP1, which does not contain *gfp<sub>sf</sub>*, grown on pyruvate. The error bars show the standard deviation ( $\sigma$ ) from at least three biological replicates. The relative fluorescence of each strain with an inducer was compared to that of the same strain with no inducer. Unless indicated, the relative fluorescence was not significantly different from uninduced conditions, as calculated by unpaired samples *t* test analyses. Significant increases in expression compared to uninduced conditions are indicated (\*\*,  $P < 0.01$ ). Strain names are indicated parenthetically: *tcuR*<sup>WT</sup>, *tcIR*<sup>WT</sup> (ACN3123);  $\Delta tcIR$  (ACN3184);  $\Delta tcuR$  (ACN3122);  $\Delta tcuR$ ,  $\Delta tcIR$  (ACN3173);  $\Delta tcuR$ , *tcIR*<sup>R200Q</sup> (ACN3191); and  $\Delta tcuR$ , *tcIR*<sup>R200Q, L216P</sup> (ACN3190).

increased fluorescence (Fig. 4). This induction did not occur without TcIR, suggesting that TcIR may be the only regulator that responds to Taa or Caa at  $P_{tcuC}$ . In strains with either TcIR or the TcIR variants, fluorescence increased in response to either Caa or Taa (Fig. 4).

### Transcriptional units and start sites

To assess which genes are transcribed in operons, RNA was isolated from Taa-grown ADP1. The cDNA synthesized from a *pacI*-specific primer was used as a PCR template with *tcuC* primers (Fig. 5A and B). Products of the expected sizes were observed that could only be generated if *tcuC* and *pacI* were cotranscribed. Similar experiments were conducted with Tcb-grown ADP1 to evaluate the cotranscription of *tcuA* and *tcuB*. The results showed that *tcuA* and *tcuB* form an operon (Fig. 5C and D).

The transcription initiation site (+1) for each operon was localized between two forward primers, one that gave a product (PCR 2) and the other that did not (PCR 3; Fig.

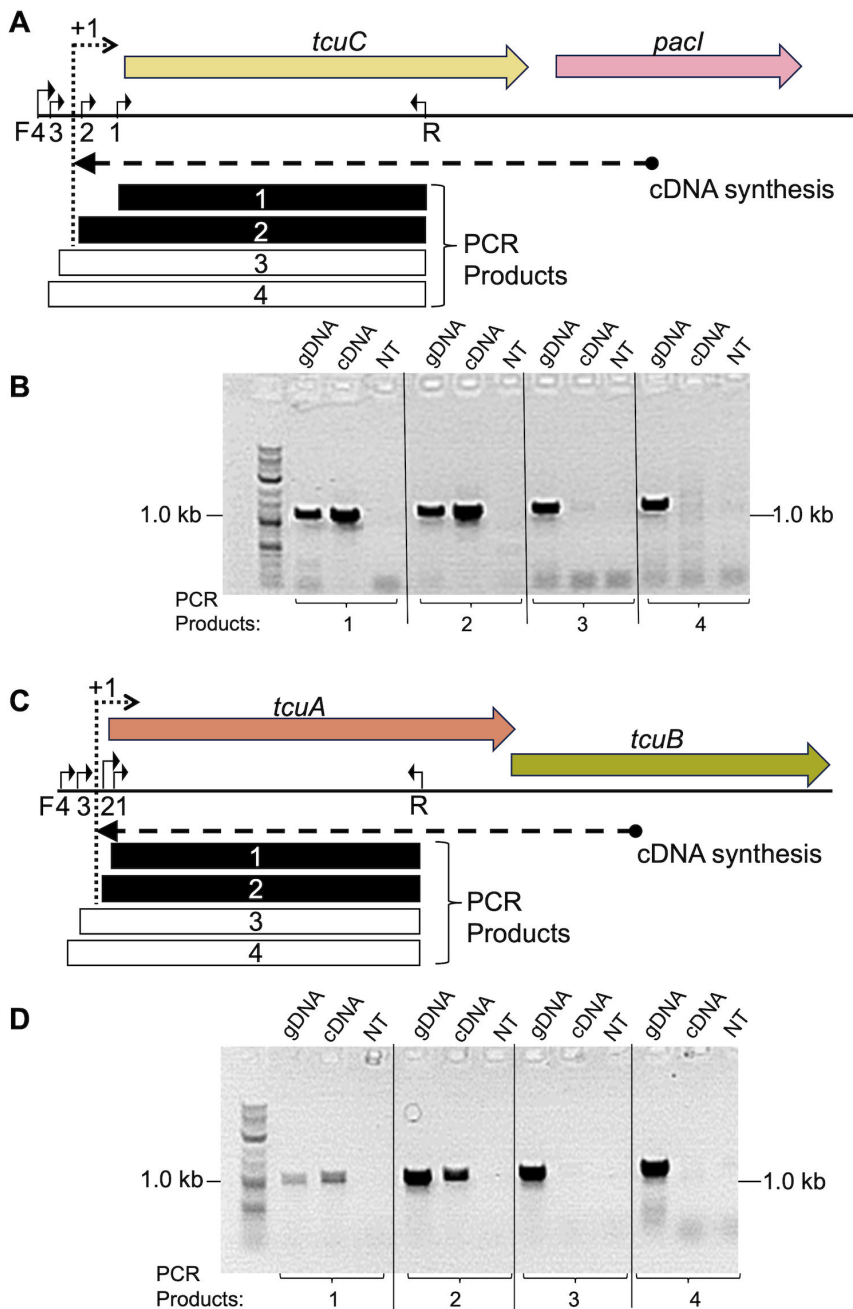


**FIG 4** Transcription of *gfp<sub>sf</sub>* controlled by  $P_{tcuC}$ . Each strain has a chromosomal *gfp<sub>sf</sub>* (allele *tcuC-gfp<sub>sf</sub>-ΩK53222; Δpacl53222*). Differences in genetic backgrounds are indicated. All strains were grown on pyruvate supplemented with inducer compounds as indicated. Relative fluorescence is the ratio of fluorescence/OD for each condition compared to that of ADP1, which does not contain *gfp<sub>sf</sub>*, grown on pyruvate. The error bars show the standard deviation ( $\sigma$ ) from at least three biological replicates. Unless indicated, fluorescence/OD is not significantly different from uninduced conditions, as calculated by unpaired samples *t* tests. Significant increases in expression compared to uninduced conditions are indicated (\* $P$  < 0.05; \*\* $P$  < 0.01). Strain names are indicated parenthetically: *tcuR*<sup>WT</sup>, *tcuR*<sup>WT</sup> (ACN3225);  $\Delta$ *tcuR* (ACN3224);  $\Delta$ *tcuR* (ACN3222);  $\Delta$ *tcuR*,  $\Delta$ *tcuR* (ACN3223);  $\Delta$ *tcuR*, *tcuR*<sup>R200Q</sup> (ACN3228); and  $\Delta$ *tcuR*, *tcuR*<sup>R200Q</sup>, L216P (ACN3227).

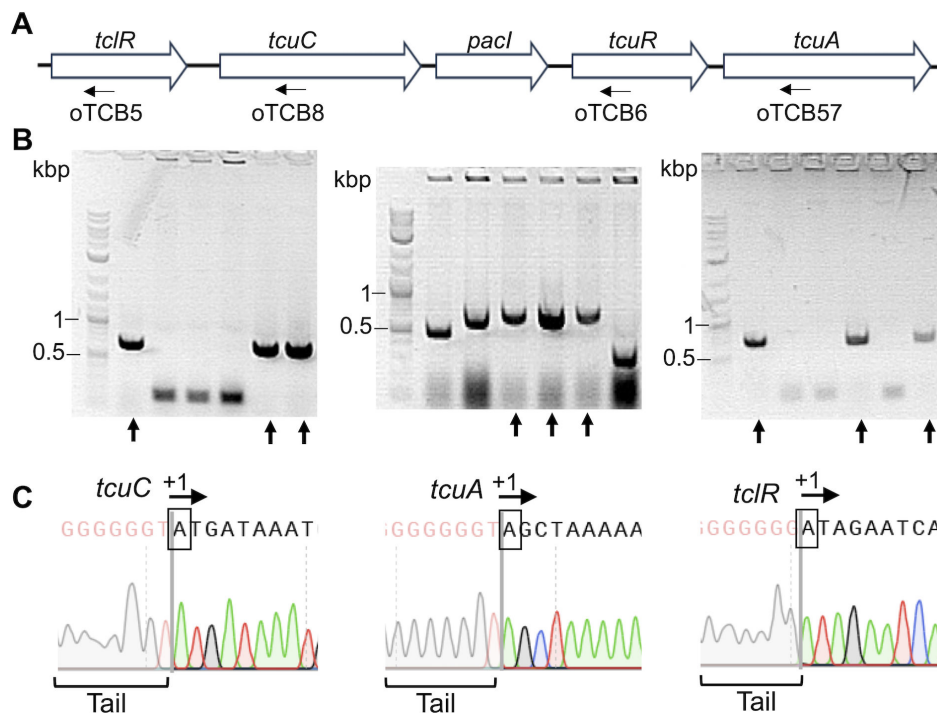
5). In both cases, PCR3 and PCR4 failed to yield a product with the cDNA template. The absence of these PCR products (3 and 4) indicates that *tcuR* is not cotranscribed with its downstream neighbor, *tcuA*, and that *tcuR* is not cotranscribed with its downstream neighbor, *tcuA*.

Transcription initiation sites were pinpointed by “Rapid Amplification of cDNA Ends” (5’ RACE; 25). For Taa-grown ADP1, mRNA was isolated, and cDNA was generated using primers for *tcuR* and *tcuA* in separate reactions. For Tcb-grown ADP1, mRNA was isolated, and cDNA was generated with primers for *tcuR* and *tcuA* in separate reactions. The 5’ RACE protocol modifies the cDNA by adding a known DNA sequence (a tail) to its 5’ end. This approach enables subsequent PCR amplification of the region. After cloning the PCR products, plasmids were checked for appropriately sized inserts (Fig. 6B). Individual clones were sequenced to find the junction between the tail and sequences identical to ADP1 DNA. These junctions, Fig. 6C, mark the transcription initiation sites (+1). The experimentally determined +1 site for *tcuC* was separated from its coding sequence by 187 nt. The transcription start site for *tcuA* was separated from its coding sequence by 22 nt. Both sites were within the regions localized by RT-PCR (Fig. 5). For *tcuR*, the +1 site was separated from its coding sequence by 40 nt, whereas multiple attempts to identify the start site for *tcuR* were unsuccessful.





**FIG 5** RT-PCR analysis of operons. (A) With RNA from Taa-grown ADP1, *pacl*-specific cDNA was made and used in four individual PCR reactions. The same reverse primer, R (oTCB113), was used with different forward primers, F1 (oTCB44), F2 (oTCB45), F3 (oTCB144), or F4 (oTCB145). With cDNA as a template, products were detected from reactions 1 and 2 (black rectangles), but not reactions 3 and 4 (white rectangles). (B). PCR products of the expected size, ~1 kbp, were observed on an agarose gel for the cDNA template using F1 or F2. With F3 or F4, no PCR products were observed. The reactions with genomic DNA (gDNA) as a template (positive controls) gave correctly sized products for all primer pairs, and the reactions with no template (NT) DNA (negative controls) gave no products. (C) Comparable studies of *tcuA* and *tcuB* were done with cDNA generated with a *tcuB*-specific primer and RNA from Tcb-grown ADP1. The reverse primer, R (oTCB154), was used with different forward primers, F1 (oTCB156), F2 (oTCB56), F3 (oTCB55), or F4 (oTCB155). (D) PCR products were assessed as indicated for panel B. The transcriptional start site (+1) for the *tcuC-pacl* operon and that for the *tcuAB* operon lies between forward primers 2 and 3, since the former but not the latter yielded a PCR product with the cDNA template.



**FIG 6** Transcriptional start sites (+1) for *tcuC*, *tcuA*, and *tcIR*. (A) Genetic organization and locations of primers (arrows) used to generate cDNA for *tcIR*, *tcuC*, *tcuR*, and *tcuA*: oTCB5, oTCB8, oTCB6, and oTCB57, respectively. (B) Following the 5' RACE protocol (Materials and Methods), PCR products corresponding to the regions of transcriptional start sites were cloned into plasmids for *tcuC*, *tcuA*, and *tcIR*. Reactions for *tcuR* were unsuccessful. Individual plasmids were isolated, and PCR was used to detect cloned inserts shown on gels for *tcuC*, *tcuA*, and *tcIR* (left to right). Multiple DNA fragments (indicated by arrows) were sequenced. (C) DNA sequencing chromatograms used to identify the +1 sites (boxed and labeled with an arrow) that correspond to the junction between DNA added during the 5' RACE procedure (the tail) and ADP1 DNA sequence. Identical results were obtained for each gene using two or more independently isolated RNA samples and three or more sequenced clones.

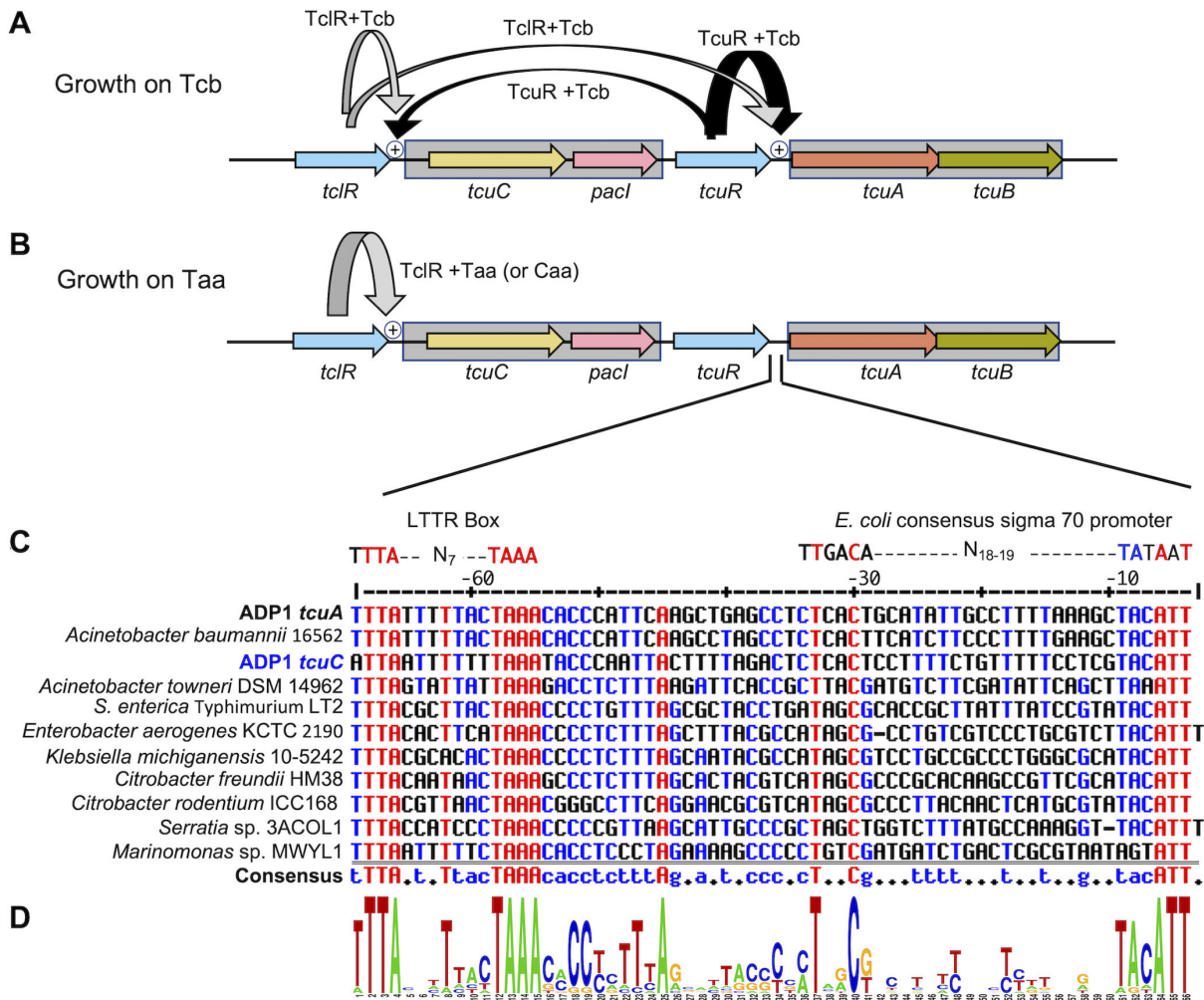
### Regulation of the *tcuC-pacI* and the *tcuAB* operons by TcuR and TcIR

A regulatory model based on the data is presented in Figure 7. The functional overlap of TcuR and TcIR at both operator-promoter regions is not surprising considering their sequence similarity. Sequences of TcuR, TcIR, and the other LTTRs encoded near *tcu* genes (Fig. 2) were aligned (Fig. S2). The three LTTRs encoded by genes (striped in Fig. 2) that are divergent from *pacI*-like genes are more like each other than they are to the TcuR-like homologs. The two LTTRs from ADP1 more closely resemble each other than the regulators from other bacteria (Fig. S2 and S3). The helix-turn-helix region (HTH), involved in DNA sequence recognition and binding by LTTRs (26, 27), is nearly identical in TcuR and TcIR (Fig. S2B). Thus, both proteins may recognize similar operator-promoter sequences, as has been shown for other paralogous LTTRs in ADP1 (28, 29).

Upstream of *tcuA*, a site matching a consensus for binding LTTRs (26), an LTTR box, was discovered using sequence alignments and pattern searches near *tcuA* homologs (Fig. 7C; 32). A putative binding site for TcuR was found to be TTTA-N<sub>7</sub>-TAAA. The sequence upstream of *tcuC* also aligns well with the *tcuA* regions (Fig. 7C). No LTTR box was found upstream of *tcIR*, but one potential binding site lies within the *tcIR* coding sequence (Fig. S4).

### Transport of tricarboxylic acids

TcuC, a member of the major facilitator superfamily (MFS), entry IPR011701 of InterPro (18, 33), affects Tcb consumption (Table 1). Its counterpart in LT2, originally identified



**FIG 7** Regulatory model. (A) During growth on Tcb, TcuR (black arrow) responds to Tcb to activate transcription of the *tcuAB* operon and to a lesser extent (thinner arrow) the *tcuC-pacli* operon. TcIR (gray arrow) responds to Tcb for low-level transcriptional activation from  $P_{tcuC}$  and  $P_{tcuA}$ . (B) During growth on Taa, TcIR (gray arrow) activates transcription from  $P_{tcuC}$  in response to Taa or Caa. (C) Alignment of the ADP1 *tcuA* promoter region with bacterial sequences upstream of *tcuA* from LT2 and other predicted *tcuA* genes. The rightmost T in the top line corresponds to position -5 relative to the transcriptional start site. The promoter region of ADP1 *tcuA* was aligned with the *tcuA* sequences. The rightmost T in the alignment (third line) corresponds to position -9 from the transcriptional start site. The alignment was generated using MultiAlin and default settings (30). Identical residues in all aligned positions are shown in red. Those that are identical in 50% or more of the sequences are shown in blue. (D) Sequence conservation represented with the WebLogo generator; sequence conservation at each position is indicated by the height of the letter (31). Conserved nucleotides in the -10 and -35 regions resemble the patterns of *E. coli* sigma 70 promoters.

as a Cit transporter, was initially designated CitA (34). In most databases, *tcuC* remains annotated as *citA*, as it is for ADP1. To clarify whether loss of *tcuC* affects Cit consumption, the *tcuC* mutant (ACN1456) was tested and found to be Cit<sup>+</sup> (Table 1). However, the impact of *tcuC* deletion could be masked by a paralog, also annotated as *citA* (ACIAD\_RS0183), which encodes a protein 57% identical to TcuC. We designated this gene *cltA* (for the *citA*-like gene) and compared strains with deletions in *tcuC* (ACN1456) or both *cltA* and *tcuC* (ACN3168). Since Cit<sup>+</sup> growth was comparable for both (Table 1), no role for TcuC in Cit transport was evident. Furthermore, the primary role of CltA remains unclear. Some bacteria have two nearby genes encoding TcuC-like transporters, labeled C and D (Fig. 2), which have sequences that align well with the other MFS transporters discussed here (Fig. S5).

## DISCUSSION

We discovered interrelated metabolic pathways in *A. baylyi* ADP1 for the consumption of Tcb, Taa, and Caa (Fig. 1). Few studies have addressed the consumption of these compounds as growth substrates for soil bacteria. Nevertheless, the clustered genetic arrangement of *tcu* genes with an aconitate isomerase-like gene in diverse bacteria (Fig. 2) suggests that ADP1 is not unique in co-regulating Tcb and Taa consumption. Bacteria with these genes were isolated from different environments, ranging from wetland peat (*Aquitalea denitrificans* 5YN1-3) to root nodules (*Cupriavidus taiwanensis* LMG 19424). The wide distribution of bacteria capable of consuming plant-derived tricarboxylic acids presumably corresponds to the varied fates of plants themselves. In addition to the presence of Taa and Tcb in grasses and plant forage, these compounds have been detected in barley, maize, beets, and cane molasses, and they can be derived in animals from consuming carbohydrates (35). Therefore, Tcb utilization by gut microbes, such as LT2, most likely reflects exposure from the diet of the host.

In LT2, the regulation of Tcb utilization is tightly controlled at the transcriptional level by TcuR. The organization of the *tcuABC* genes in an operon may help to couple Tcb transport to its metabolism. Tethering TcuC-mediated transport to TcuAB-mediated conversion to Caa, as depicted in Figure 1A, appears to prevent the cellular accumulation of Tcb, a compound that can inhibit aconitase and isocitrate dehydrogenase (36). Similarly, the regulated cotranscription of *tcuC* and *pacI* could coordinate the Pacl-mediated conversion of Taa to Caa in the periplasm with the TcuC-mediated uptake of Caa, as depicted in Fig. 1B. This regulation might limit the cellular accumulation of Taa, a compound long known to inhibit aconitase (37).

The inference from growth patterns of ADP1-derived mutants (Table 1) that TcuC can transport both Caa and Taa is consistent with transport studies in *Salmonella* published 50 years ago. At that time it was shown that a Tcb-inducible transporter could also transport Caa (38). Although the associated genes/protein sequences were not identified at that time, the Tcb-inducible transporter presumably corresponds to TcuC in LT2.

### TcuC-like transporters

It seems unlikely that TcuC would import Taa, yet some of this compound must enter the cell for it to be an inducer in strains that do not express Pacl (Fig. 4). In its protonated form, Taa likely diffuses into the cell in amounts limited by pH and concentration. MFS family members are often stereospecific (33), so it is likely that TcuC is able to distinguish the *cis*- and *trans*-forms of aconitate. TcuC of LT2 also transports Cit and does so in preference to isocitrate (34). However, seven single amino acid replacements increase the transport of isocitrate (Fig. S5). ADP1 cannot use isocitrate as a sole carbon source, but it grows well on Cit, even in deletion mutants without *tcuC* and *cltA*. In *Pseudomonas aeruginosa*, deletion of a *citA* homolog did not affect the growth on Cit, even in conjunction with the inactivation of another putative transporter gene (39). Redundant paralogs with overlapping transport functions, often with different regulatory patterns, are frequently observed in bacteria, including ADP1 (29, 40). A few bacteria had two nearby *tcuC* paralogs (Fig. 2, labeled C and D), yet alignments failed to suggest any functional, phylogenetic, or regulatory significance to sequence differences, including at positions shown to affect the substrate specificity of TcuC from LT2 (Fig. S5).

### Regulation by TcuR and TcIR: effector compound recognition and response

The ten aligned LTTRs formed two branches in a phylogram (Fig. S2 and S3). TcuR and TcIR from ADP1 are more like each other and TcuR in LT2 (6) than to the three LTTRs encoded near the *pacI*-like genes of other bacteria. These differences raise questions about regulation mediated by the three distinct LTTRs. Nevertheless, the sequence similarity between TcuR and TcIR correlates with their regulatory overlap (Fig. 7A and B). Overall, these ADP1 paralogs appear to function similarly to members of the large LTTR family (26).

Typically, LTTRs respond to one or more small chemicals that serve as inducers, coinducers, and/or coactivators, collectively referred to as effectors to acknowledge that LTTRs can be transcriptional repressors, activators, or both (26). Binding an effector in the C-terminal region, the effector-binding domain (EBD), of an LTTR subunit causes a conformational change that propagates throughout the oligomer, usually a homotetramer (26). While both TcuR and TcIR responded to Tcb as an effector, two TcIR variants were independently selected that increased Tcb-dependent transcriptional activation (Fig. 3 and 4). In the absence of characterized atomic-level structures of any Tcb-binding LTTRs, we used models of TcIR-EBD and TcuR-EBD to compare with crystal structures of two Cit-responsive LTTRs, CcpE and CcpC (Fig. S6; 41, 42). In these structures, Cit binds between two EBD subdomains (Fig. S6A). Superimposition of the TcIR model indicates that the R200 side chain corresponds to the position of Cit. Therefore, the R200Q replacement in the TcIR variant may have a direct impact on the Tcb-binding site (Fig. S6C and D). The possible effect of the L216P replacement is less clear, based on its position in the protein and because it only appeared together with R200Q.

Positions of amino acids that render TcuR of LT2 able to activate transcription without Tcb (5) were also compared to the CcpC structure (Fig. S7). The R264 residue in the LT2 TcuR, which is conserved in all sequences in Fig. S2, is predicted to be centrally positioned in one of two beta-strands forming a hinge between the EBD subdomains (Fig. S7D). In LTTRs, an effector-binding pocket forms between these subdomains (26). The R264L replacement appears to lock the protein in an active conformation without Tcb. Other changes in constitutive TcuR variants (5) also cluster near the predicted effector-binding site (Fig. S7C).

While TcuR and TcIR responded to Tcb, TcIR additionally responded to Taa, under conditions where it was not metabolized. Caa may also activate transcription by TcIR, but this compound was only tested in strains that could metabolize it. Thus, Caa-dependent changes could be mediated by a downstream metabolite. This broader range of effectors for TcIR apparently helps induce TcuC expression for growth on Taa, Tcb, and Caa.

### Regulation by TcuR and TcIR: operator-promoter recognition

The transcriptional initiation sites (+1) for the *tcuAB* operon and the *tcuC-paI* operon were aligned with each other and comparable regions near other predicted *tcuA* genes (Fig. 7C). A conserved sequence, TTTA-N<sub>7</sub>-TAAA, matches the consensus sequence for LTTR binding (26, 27, 43). Based on its sequence, conservation, and position (−63 relative to the +1 site of *tcuA* of ADP1), we inferred that this site, called an LTTR box, anchors the binding of two subunits of a TcuR (or TcIR) tetramer. For transcriptional activation, interactions of the LTTR are also expected in the vicinity of the −35 region of the promoter. However, sequences that typically indicate such interactions (43) were not observed for *tcuA* or *tcuC*. There are many examples of atypical operator-promoter sequences regulated by LTTRs (26).

The *tclR* operator-promoter region was aligned with those of *tcuA* and *tcuC*, and no evident LTTR box was found upstream of the coding sequence (Fig. S4). Although a common method of negative autoregulation occurs when LTTR genes are divergent from the corresponding target, autoregulation sometimes occurs differently (26). The significance of an LTTR box sequence in the *tclR* coding region remains to be tested (Fig. S4). No potential LTTR box was identified for *tcuR* of ADP1, and studies in LT2 ruled out autoregulation in that microbe (5). The *tcuR* transcriptional start site was not identified. This failure of the 5′ RACE attempts may reflect that the levels of LTTR transcripts tend to be low and difficult to detect (44).

### Overlapping regulation by similar LTTR pairs

The nuanced control by TcuR and TcIR parallels that of other pairs of LTTR paralogs in ADP1. For example, DarR and AalR have interrelated but distinct functions in a shared regulon for aspartate metabolism (29). Each LTTR responds to a distinct enantiomer



of aspartate, but both recognize the same LTTR box, which controls several different genes and operons. Two other LTTR paralogs, BenM and CatM, jointly regulate multiple loci involved in aromatic compound metabolism (45, 46). Both recognize similar DNA sequences and the same effector molecule. An added level of fine-tuned control in this example is the integration of multiple cellular signals via the synergistic response of BenM to a second effector compound (28).

These elaborate regulatory schemes underscore the importance of balanced transport and consumption of growth substrates. For aromatic compound catabolism, there are very complicated multistep pathways in which each step may generate a toxic metabolite, and the accumulation of such intermediates is carefully regulated. For the consumption of Tcb, Taa, and Caa, there are few catabolic steps prior to entry to the tricarboxylic acid cycle. Nevertheless, the potential of Taa and Tcb to inhibit the tricarboxylic acid cycle, and other characteristics of these compounds, such as metal chelation, may account for multiple layers of transcriptional control governed by TcuR and TcIR.

## MATERIALS AND METHODS

### Media and growth conditions

Cultures were grown in minimal medium (MM; 47) or lysogeny broth (LB), also known as Luria-Bertani medium (10 g of Bacto Tryptone, 5 g of yeast extract, and 10 g of NaCl per L; 48), as previously described (29). Unless otherwise noted, carbon sources were added to MM at final concentrations of 15 mM for pyruvate or 5 mM for Tcb, Taa, Caa, or Cit. Antibiotics were added as needed at final concentrations of 25-mg/mL for kanamycin (Km), 12.5-mg/mL each for streptomycin (Sm) and spectinomycin (Sp), and 150-mg/mL for ampicillin (Ap).

Bacteria were incubated at 37°C or 30°C. Liquid cultures, except in microtiter plates, were aerated by shaking at 250 rpm, at 37°C. Culture volumes of 5 mL were grown in 15-mL tubes, and larger cultures of 25 mL were grown in 250-mL flasks. To inoculate, a dilution ratio of 1:100 was typical. For monitoring growth in a Synergy H1 plate reader (BioTek), 2  $\mu$ L of an inoculum (from a liquid culture that was grown for at least 12 h) was added to 198  $\mu$ L of medium in the well of a Costar 3603 96-well plate. The total volume of each well was 360  $\mu$ L. Readings were taken every 30 min ( $OD_{595}$ ) while the plate was incubated at 30°C, with orbital shaking at 282 rpm and a 3-mm orbit. All comparative data that we report used strains grown under identical conditions (e.g., Table 1). Strains grew approximately threefold slower in the plate reader, but the growth rates of different strains relative to each other were the same regardless of condition.

### Strains, plasmids, and PCR primers

*A. baylyi* strains were derived from the wild type, ADP1 (22, 23), as listed in Table 2 with additional details in Tables S2 to S4. *Escherichia coli* strains XL1-Blue (Agilent Technologies) and DH5 $\alpha$  (49) were used as plasmid hosts. Plasmids are listed in Table S3 and PCR primers in Table S4. Drug resistance cassettes were obtained from pUI1637 ( $\Omega$ K, confers Km<sup>R</sup>) and pUI1638 ( $\Omega$ S, confers Sm<sup>R</sup> and Sp<sup>R</sup>; 50). Standard vectors were used for cloning in *E. coli*, such as pUC18/pUC19 (51). The *gfp<sub>sf</sub>* transcriptional reporter gene encodes a green fluorescent protein (24). A selectable/counter-selectable cassette (*sacB*-Km<sup>R</sup>) was obtained from pRMJ1 (52).

Strains and plasmids were constructed by standard molecular biology methods (48) and techniques developed for ADP1 (13, 14). High-fidelity polymerases were used for PCR, PrimeSTAR Max (Takara Biosciences) and Phusion (New England Biolabs), with primers shown in Table S4. Plasmids were constructed by overlapping sequence assembly (53), by restriction digest and ligation (Quick Ligation Kit; New England Biolabs), with the NEBuilder kit (New England Biolabs) and/or by overlap extension PCR (54). Plasmids were confirmed by restriction mapping and/or regional DNA sequencing



(Eurofins Genomics). *A. baylyi* genomic changes were made by allelic replacement using linear DNA to transform naturally competent recipients (13, 14, 55, 56). When PCR products were used as donor DNA, digestion by DpnI (New England Biolabs) was used to degrade template DNA. Genotypes of the recipient strains were confirmed by PCR with LongAmp polymerase (New England Biolabs).

### Fluorescence assays of transcriptional fusions

Growth and fluorescence were assessed for strains with a chromosomal copy of *gfp<sub>sf</sub>* under the transcriptional control of *P<sub>tcuC</sub>* or *P<sub>tcuA</sub>* using a Synergy H1 plate reader (BioTek). Cultures were grown with pyruvate as the carbon source and diluted (1:100) in the same medium with and without an exogenous inducer (Tcb, Taa, or Caa at concentrations of 0.1 mM or 5 mM). Green fluorescence measurements (to assess GFP<sub>sf</sub> expression) were taken, at excitation 479 nm and emission 520 nm, concurrently to growth measurements at OD<sub>595</sub>. Fluorescence values, corresponding to cultures that reached an OD<sub>595</sub> of 0.35–0.50, were recorded relative to cell density (OD<sub>595</sub>). To account for a high background fluorescence in ADP1, the data in Fig. 3 and 4 are reported in comparison to that of the wild-type strain, which has no *gfp<sub>sf</sub>* gene.

### Transcript analysis

For RNA isolation, cultures were grown and harvested at the mid-log phase with an OD<sub>595</sub> of approximately 0.4–0.6. The cells were diluted to a concentration of approximately 10<sup>8</sup> cells. The cells were lysed using 200  $\mu$ L of lysozyme into 100  $\mu$ L of liquid culture. RNA was preserved using RNeasy Protect (Qiagen) and isolated using the RNeasy Mini Kit (Qiagen). RNA was further treated with dsDNase (Thermo Fisher) and purified (Monarch) or RQ1 DNase (Promega). RNA was confirmed to be free of DNA using PCR. cDNA was generated using the Maxima H Minus First Strand cDNA Synthesis Kit (Thermo Scientific). A 5' RACE DNA purification system kit (Invitrogen) was used for cDNA purification and to add a homopolymeric tail of nucleotides (dCTP) with terminal deoxynucleotidyltransferase. Nested PCR was performed on the tailed DNA as described in the kit protocol using a Taq polymerase and primers from the kit along with gene-specific primers (Table S4). Fragments were visualized on agarose gels, and the largest observed fragments were cloned in the pCR2.1-TOPO vector using a TOPO TA cloning kit (Invitrogen). Plasmids were isolated, after the transformation of *E. coli*, and DNA sequencing was used to identify sequences that correspond to transcriptional start sites. Multiple plasmids were sequenced and shown to indicate the same start site. All transcript analyses were confirmed with repetitions using two or more independently isolated RNA samples.

### Selection of spontaneous mutants that grow on Tcb without TcuR

A  $\Delta$ *tcuR* mutant, ACN1376, did not grow on Tcb as the sole carbon source. Pyruvate-grown cultures of ACN1376 were concentrated approximately 10-fold and spread on plates with Tcb as the carbon source. Tcb<sup>+</sup> colonies arose (at a frequency of 10<sup>-9</sup>). Two independently isolated Tcb<sup>+</sup> derivatives of ACN1376, ACN1445 and ACN1556, were streak-purified and further characterized. Localized DNA sequencing revealed mutations in *tclR*.

### DNA sequence analysis

The reference sequences, from the National Center for Biotechnology Information (NCBI; 57), for bacteria discussed in this report or referenced in figures are indicated parenthetically: *A. baylyi* ADP1 (NC\_005966), *Acinetobacter baumannii* 1656–2 (NC\_017162), *Acinetobacter towneri* DSM 14962 (NZ\_KB849689), *Aquitalea denitrificans* 5YN1-3 (NZ\_CP047241), *Citrobacter freundii* HM38 (NZ\_CP024672), *Citrobacter rodentium* ICC168 (NC\_013716), *Cupriavidus taiwanensis* LMG 19424 (NC\_010530), *Curvibacter delicatus* NBRC 14919 (NZ\_BCWP01000009), *Enterobacter aerogenes* KCTC 2190

(NC\_015663), *Klebsiella michiganensis* strain 10–5242 (NZ\_JH603153), *Marinomonas* sp. MWYL1 (NC\_009654), *Paraburkholderia sacchari* LMG 19450 (NZ\_JTDB02000009), *S. enterica* LT2 (NC\_003197), *Serratia* sp. 3ACOL1 (CP033055.1), and *Xanthobacter autotrophicus* Py2 (NC\_009720). For homology searches, we used the Basic Local Alignment Search Tool (BLAST; 58) and the sequence similarity database (SSDB) with associated search tools of the Kyoto Encyclopedia of Genes and Genomes (59). Protein sequences were aligned, and data were exported using programs on the UniProt website (30).

In the *A. baylyi* ADP1 genome, our new gene designations correspond to the following locus tags: *cltA* (ACIAD\_RS01830), *tcIR* (ACIAD\_RS07105), *tcuC* (ACIAD\_RS07100), *pacI* (ACIAD\_RS07095), *tcuR* (ACIAD\_RS07090), *tcuA* (ACIAD\_RS07085), and *tcuB* (ACIAD\_RS07080). Protein sequence alignments (in Fig. S1, S2 and S5) were generated using a Sequence Manipulation Suite program, Multiple Align Show (60). Promoter sequences shown in Fig. 7 and Fig. S4 were aligned and displayed using the MultAlin program (61). The Signal P program (21) was used online: <https://biolib.com/DTU/SignalP-6/>. The STRING database program (20) was used online: <https://string-db.org>.

## ACKNOWLEDGMENTS

Some of these studies were conducted during several semesters of a University of Georgia Microbiology laboratory course. The following students contributed to generating and characterizing plasmids and strains: Diana Ambrocio, John Hunter Bowen, Lindsey Brock, Gary Kevin Davis, Nicholas Evans, Destiny Finnell, John Patrick Goldun, Mark Goodwin, Grigsby Gordy, Amanda Helton, Maliha Ishaq, Brandy Le, Connor Lombardi, Mengzhuo (Jen) Ma, Kiel Mills, Alex Moosariparambil, Maria Najera, Nneka Okoro, Destiny Okpomo, Rashidatu Olorunsola, Emma Osborne, Cathryn Paul, Quynh Phan, Christopher Perez, Mason Pyles, Delaney Ragsdale, Liam Richardson, Tonya Brook Ringle, Emma Grace Turner, Keiana Watkins, Kaitlin Whitely, Winsen Wijaya, Domenic Won, and Christofer Zepeda. Emily McIntyre and Andrew Wiggins contributed as teaching assistants for the course. Additional students in the Neidle laboratory helped to design and conduct experiments: Stephanie Thurmond, Silke Andresen, Stacy Bedore, and Hannah Toutkoushian.

We thank Drs. Cory Momany and Jorge Escalante-Semerena for their helpful discussions and advice.

This work was supported by grants from the National Science Foundation (MCB2225858 to E.L.N. and DBI-1757720 and DBI-1062589 for Research Experience for Undergraduate Site programs for two participants, Kayla Figatner and Hannah Toutkoushian, respectively). Funding was also provided from the U.S. Department of Energy, Office of Science, Office of Biological and Environmental Research, Genomic Science Program (DE-SC0022220 to E.L.N.). Additional funding was provided by the UGA Microbiology Department.

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## FUNDING

Funder	Grant(s)	Author(s)
National Science Foundation (NSF)	MCB2225858, DBI-1757720, DBI-1062589	Ellen L. Neidle Chantel V. Duscent-Maitland Kayla Figatner
U.S. Department of Energy (DOE)	DE-SC0022220	Ellen L. Neidle Chantel V. Duscent-Maitland Alyssa C. Baugh Kayla Figatner Justin B. Defalco

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## ADDITIONAL FILES

The following material is available [online](#).

### Supplemental Material

**Supplemental figures and tables (AEM02111-23-s0001.pdf).** Tables S1 to S4 and Fig. S1 to S7.

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