Anaerobic Respiration Using a Complete Oxidative TCA Cycle Drives Multicellular Swarming in *Proteus mirabilis*

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**ABSTRACT** *Proteus mirabilis* rapidly migrates across surfaces using a periodic developmental process of differentiation alternating between short swimmer cells and elongated hyperflagellated swarmer cells. To undergo this vigorous flagellum-mediated motility, bacteria must generate a substantial proton gradient across their cytoplasmic membranes by using available energy pathways. We sought to identify the link between energy pathways and swarming differentiation by examining the behavior of defined central metabolism mutants. Mutations in the tricarboxylic acid (TCA) cycle (*fumC* and *sdhB* mutants) caused altered patterns of swarming periodicity, suggesting an aerobic pathway. Surprisingly, the wild-type strain swarmed on agar containing sodium azide, which poisons aerobic respiration; the *fumC* TCA cycle mutant, however, was unable to swarm on azide. To identify other contributing energy pathways, we screened transposon mutants for loss of swarming on sodium azide and found insertions in the following genes that involved fumarate metabolism or respiration: *hybB*, encoding hydrogenase; *fumC*, encoding fumarase; *argH*, encoding argininosuccinate lyase (generates fumarate); and a quinone hydroxylase gene. These findings validated the screen and suggested involvement of anaerobic electron transport chain components. Abnormal swarming periodicity of *fumC* and *sdhB* mutants was associated with the excretion of reduced acidic fermentation end products. Bacteria lacking SdhB were rescued to wild-type pH and periodicity by providing fumarate, independent of carbon source but dependent on oxygen, while *fumC* mutants were rescued by glycerol, independent of fumarate only under anaerobic conditions. These findings link multicellular swarming patterns with fumarate metabolism and membrane electron transport using a previously unappreciated configuration of both aerobic and anaerobic respiratory chain components.

**IMPORTANCE** Bacterial locomotion and the existence of microbes were the first scientific observations that followed the invention of the microscope. A bacterium can swim through a fluid environment or coordinate motion with a group of bacteria and swarm across a surface. The flagellar motor, which propels the bacterium, is fueled by proton motive force. In contrast to the physiology of the microscope. A bacterium can swim through a fluid environment or coordinate motion with a group of bacteria and swarm across a surface. The flagellar motor, which propels the bacterium, is fueled by proton motive force. In contrast to the physiology that governs swimming motility, much less is known about the energy sources required for multicellular swarming on surfaces. In this study, we used *Proteus mirabilis* as a model organism to study vigorous swarming behavior and genetic and biochemical approaches to define energy pathways and central metabolism that contribute to multicellular motility. We found that swarming bacteria use a complete aerobic tricarboxylic acid (TCA) cycle but do not respire oxygen as the terminal electron acceptor, suggesting that multicellular cooperation during swarming reduces the amount of energy required by individual bacteria to achieve rapid motility.

*Proteus mirabilis* differentiates from a single short rod-shaped swimmer cell into a multinucleate, elongated, and hyperflagellated swarmer cell in response to extreme viscosity or solid surfaces. This periodic developmental process, known as swarming differentiation, requires multicellularity and results in a regular pattern of rapid migration across a surface. Specifically, swimmer cells associate with one another and move in “rafts” (1). As the advancement of the swarm front decreases in velocity (2) and pauses, each polyploid hyperelongated swarmer cell septates and divides into numerous individual swimmer cells in a process known as consolidation. After a period of consolidation, vegetative swimmer cells differentiate into swimmer cells, and this behavior repeats multiple times, resulting in the bull’s-eye appearance of concentric rings of growth on an agar plate (3, 4). This cyclical developmental process and multicellular swarming behavior on solid (1.5% [wt/vol]) agar is characteristic of *P. mirabilis* and distinguishes it from other bacteria (5). While it is clear that initiation of swarming involves signaling events that lead from surface sensing to upregulation of *flhDC* and flagellum production (6–9), the molecular and physiological basis for the periodic multicellular swarming behavior remains poorly understood.

*P. mirabilis* swarmer cells possess an outer membrane composition biochemically distinct from nonswarming bacteria, or swimmer cells. Swarmer cells possess an increased number of long

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**Received** 16 September 2012 **Accepted** 10 October 2012 **Published** 30 October 2012

**Citation** Alteri CJ, Himpsl SD, Engstrom MD, Mobley HLT. 2012. Anaerobic respiration using a complete oxidative TCA cycle drives multicellular swarming in *Proteus mirabilis*. mBio 3(6):e00365-12. doi:10.1128/mBio.00365-12.

**Editor** Roberto Kolter, Harvard Medical School

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O-antigen side chains on the outer membrane lipopolysaccharide (LPS) (10), which could be important for surface sensing during initiation of swarming (11). Indeed, disruptions in genes responsible for LPS and cell wall biosynthesis affect swarmer cell elongation (8, 11). In contrast, the surface capsular polysaccharide is required for surface migration but not differentiation into elongated swarmer cells (12). Additionally, a mutation in ccmA (curved cell morphology) causes irregularly curved swarmer cells that disrupt alignment into multicellular rafts and produces migration defects (13). Four umo genes, some of which encode membrane proteins, have been identified that upregulate the expression of the flhDC master operon, which in turn controls flagellar biosynthesis, cell division, and swarming (14). However, despite their effect on flhDC expression, the proteins encoded by umo genes localize to the cell membrane or periplasm; our study of P. mirabilis gene expression during swarming and consolidation identified umoD, flagellar genes, zapA, amino acid transport genes, peptide transport genes, and cell division genes as upregulated during consolidation (16), perhaps in preparation for the next round of differentiation and swarming.

Numerous differences in physiology occur in P. mirabilis swarmer cells compared to nonswarmer cells. For example, swarmer cells have reduced amino acid uptake and have increased sensitivity to hydrophobic antibacterial agents, suggesting that reorganization of the outer membrane might affect active transport processes and protective properties (10, 17). During swarming, the rates of uptake and incorporation of precursors into DNA, RNA, and protein are also reduced (18). While ATP concentrations remain constant, the rate of oxygen uptake is also reduced during P. mirabilis swarming (18). At the end of the swarm period, during consolidation, macromolecule synthesis and oxygen uptake are restored to levels equivalent to preswarming levels (18). Furthermore, a study of Proteus vulgaris swarming showed that membrane vesicles from swarm cells have reduced rates of NADH, malate, and succinate respiration (19). Interestingly, swarm cells were also found to contain diminished levels of cytochrome b as well as being deficient in cytochrome a and d (19). Since swarming can occur during both anaerobic and aerobic conditions, it has been proposed that the energy required during swarming is generated by fermentation (19). This is surprising, because using fermentation, rather than membrane respiration, would be much less energetically favorable for fueling the flagellar motor.

The proton gradient that is created during respiration contributes to the gradient of both pH and charge across the membrane and collectively determines proton motive force (PMF). It is during aerobic conditions with O₂ as a terminal electron acceptor that the largest number of protons can be pumped, thereby producing the largest PMF for the cell. During fermentation, however, bacteria are unable to respire and must generate PMF by reversing the direction of the F₁F₀ ATPase rotation to consume ATP to pump protons out of the cytoplasm. Consequently, generation of PMF during anaerobic conditions is substantially more difficult for the cell compared to aerobic conditions, because the rotational speed of the flagellum is dependent on the magnitude of the PMF (20), which would be very limited in the absence of membrane respiration during fermentation.

We previously identified two P. mirabilis transposon mutants disrupting genes encoding proteins in the oxidative tricarboxylic acid (TCA) cycle that each displayed an aberrant swarming phenotype (21). These mutations resided in aceE, which encodes pyruvate dehydrogenase, and sdhC, which encodes the cytochrome b-556 subunit of succinate dehydrogenase. Those findings suggested a link between PMF, the TCA cycle, and swarming motility. Surprisingly, however, Western blot analysis using antibodies to the flagellum subunit, FlaA, showed that both mutants produce similar levels of flagellin despite a dramatic difference in the swarming periodicity (21). Pyruvate dehydrogenase converts pyruvate, the end product of glycolysis, to acetyl coenzyme A (acetyl-CoA), which enters a complete oxidative TCA cycle during aerobic conditions. Succinate dehydrogenase functions in the complete TCA cycle and is a component of the aerobic electron transport chain. Because mutations in both genes resulted in aberrant swarming phenotypes, we hypothesized that a complete oxidative TCA cycle is required for swarming. However, previous findings also indicated that aerobic cytochromes are absent in swarmer cells, and fermentation was proposed as an energy source during P. mirabilis swarming (19), despite the notable requirement for PMF to drive flagellar rotation.

In this current study, we used genetic and biochemical approaches to define central metabolic and energy pathways that contribute to multicellular swarming motility. Consistent with previous studies (19), we found that swarming is resistant to sodium azide. However, characterization of swarming for defined central metabolism mutants indicated that the aerobic TCA cycle and fumarate metabolism, independent of fumarate reductase, controls swarming periodicity in P. mirabilis. Here, we present evidence for an intimate connection between energy metabolism and bacterial multicellular behavior that supports a model for anaerobic respiration using a complete oxidative TCA cycle to generate PMF to drive swarming.

RESULTS

P. mirabilis central metabolism mutants have altered periods of swarming. From a previous signature-tagged mutagenesis study (21), we noted that transposon insertions in genes encoding enzymes of the TCA cycle displayed defects in the ability of P. mirabilis HI4320 to swarm but did not affect flagellin production. To better understand the connection between central metabolism and swarming differentiation, we systematically generated defined mutants deficient in specific central pathways and tested their ability to swarm on lysogeny broth (LB) medium solidified with 1.5% (wt/vol) agar. These mutations resided in classes of altered swarming resulting from disruption of the TCA cycle that each displayed an aberrant swarming phenotype (21). These mutations resided in aceE, which encodes pyruvate dehydrogenase, and sdhC, which encodes the cytochrome b-556 subunit of succinate dehydrogenase. Those findings suggested a link between PMF, the TCA cycle, and swarming motility. Surprisingly, however, Western blot analysis using antibodies to the flagellum subunit, FlaA, showed that both mutants produce similar levels of flagellin despite a dramatic difference in the swarming periodicity (21). Pyruvate dehydrogenase converts pyruvate, the end product of glycolysis, to acetyl coenzyme A (acetyl-CoA), which enters a complete oxidative TCA cycle during aerobic conditions. Succinate dehydrogenase functions in the complete TCA cycle and is a component of the aerobic electron transport chain. Because mutations in both genes resulted in aberrant swarming phenotypes, we hypothesized that a complete oxidative TCA cycle is required for swarming. However, previous findings also indicated that aerobic cytochromes are absent in swarmer cells, and fermentation was proposed as an energy source during P. mirabilis swarming (19), despite the notable requirement for PMF to drive flagellar rotation.

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TABLE 1 P. mirabilis metabolism mutants examined in this study

| Pathway        | Gene | Description                  |
|----------------|------|------------------------------|
| Glycolysis     | pfkA | Phosphofructokinase          |
|                | pgi  | Phosphoglucone isomerase     |
|                | tpiA | Triosephosphate isomerase    |
| Pentose phosphate | gnd  | 6-Phosphogluconate dehydrogenase |
|                | talB | Transaldolase                |
| Entner-Doudoroff | edd  | 6-Phosphogluconate dehydratase |
| TCA cycle      | sdhB | Succinate dehydrogenase      |
|                | frdA | Fumarate reductase           |
|                | fumC | Fumarate                     |
| Gluconeogenesis | pckA | Phosphoenolpyruvate carboxykinase |

All mutants were constructed for this study using the TargeTron system as described in Materials and Methods.

Central carbon pathways that generate pyruvate are required for swarming differentiation. Mutations within pfkA and tpiA that encode glycolytic enzymes resulted in a diminished capacity to migrate; in particular, the diameter of swarming was reduced (class IV) compared to the wild-type HI4320 swarming diameter.
mutations within NADPH and sugars, such as ribose-5-phosphate, G3P, and catabolism is the oxidative and nonoxidative PPP, during which NADPH and sugars, such as ribose-5-phosphate, G3P, and fructose-6-phosphate, are interconverted, respectively. Distinct swelling phenotypes were also observed for the PPP (class III); mutations within gnd of the oxidative PPP and talB of the nonoxidative PPP resulted in a swelling phenotype consisting of dense elongated swelling zones following consolidation (Fig. 1, arrow). Interestingly, mutations of the ED pathway (edA mutant) and gluconeogenesis (pckA mutant) result in swelling phenotypes identical to that of the wild-type strain HI4320 (class I). Glycolysis, the ED pathway, and the PPP are all generally involved in the catabolism of 6-carbon sugars to pyruvate, yet only mutations in tpiA, pkaA, talB, and gnd have swelling defects on LB-1.5% agar. Since LB medium lacks a carbohydrate carbon source like glucose, these findings suggest that their swelling defects could result from the inability to produce sufficient pyruvate for energy metabolism or specific macromolecular precursors; DHAP, for cell membranes; fructose-6-P, for peptidoglycan (PfkA); ribose-5-P (TalB), for nucleobases; or NADPH (Gnd), which powers most biosynthetic pathways.

Multicellular swarming requires energy pathways dependent on the TCA cycle. During aerobic respiration, pyruvate generated by glycolysis, the PPP, or the ED pathway is converted to acetyl-CoA by pyruvate dehydrogenase before entering the TCA cycle, where it condenses with oxaloacetate to form citrate. Previously, we found that pyruvate dehydrogenase mutants (aceE mutant) produce wild-type levels of flagellin (FlaA) yet are unable to swim (21), suggesting that an aerobic TCA cycle is required for swimming. The TCA cycle operates during aerobic and anaerobic respiration or fermentation by running in an oxidative cycle (when respiring oxygen) or in an incomplete, reductive, and branched pathway, respectively. Mutants in TCA cycle genes have distinct swelling periodicity phenotypes (class II) compared to wild-type HI4320 (Fig. 2A). The class II phenotype was highly reproducible (see Fig. S1 in the supplemental material) and demonstrated a consistent 50% reduction (4 mm for class II compared to 8 mm for the wild type) in the distance between the consolidation rings that immediately follow the inoculation point compared to the wild type at both 30°C and 37°C (Fig. 3A and B). Bacteria with a mutation within furmarase, the fumC mutant, which functions in both the aerobic and anaerobic TCA cycle, produce shorter distances between swelling and consolidation raf (Fig. 3). The altered swelling phenotype of the fumC mutant bacteria was restored to wild-type swelling when complemented by the wild-type fumC allele on a plasmid (Fig. 2D). The same aberrant swelling phenotype is observed for the aerobic TCA cycle mutant, the sdbB mutant, which encodes succinate dehydrogenase (Fig. 2A) and was also observed previously for a strain with a mutation in sdbC (21) (see Fig. S1). In contrast, a mutation within frdA, which encodes furmarate reductase, a branched TCA pathway enzyme that is important during anaerobic respiration, causes swelling like in wild-type HI4320 (Fig. 2A). Together, the swelling phenotypes of these three TCA cycle mutants indicate that _P. mirabilis_ swelling is dependent on the complete oxidative pathway since mutation of frdA, which disrupts only the incomplete, reductive (branched) TCA pathway, did not affect swelling periodicity (Fig. 3A and B).

Addition of TCA cycle intermediates alters swelling differentiation of _P. mirabilis_. Because disruption of the aerobic oxidative TCA cycle (aceE, sdbB, sdbC, and fumC mutant bacteria) resulted in aberrant swelling phenotypes (class II), we sought to determine whether swelling periodicity of the wild-type parent strain or these TCA cycle mutants could be manipulated or chemically complemented by the addition of exogenously supplied TCA cycle intermediates. We tested _P. mirabilis_ HI4320 and the TCA cycle mutants on agar replete with furmarate, succinate, or malate. The biochemical intermediate for which a specific mutant was deficient in generating was added to the agar to test for restoration of the mutant’s swelling cycles to wild-type periodicity. For example, during the aerobic TCA cycle (Fig. 4B), succinate is oxidized to furmarate by succinate dehydrogenase, SdhB; if the altered swelling phenotype of the sdbB mutant is due to its inability to generate furmarate, then swelling of the sdbB mutant should be restored to the wild-type phenotype by the addition of furmarate. The furmarate mutant (fumC), which is unable to convert furmarate to malate during a complete oxidative TCA cycle (malate to furmarate in the branched reductive pathway [Fig. 4C]), was also examined on agar containing each of the tested intermediates (Fig. 4A). Indeed, furmarate, succinate, and malate all allow the sdbB mutant to revert to wild-type swelling periodicity, while succinate and malate rescue swelling periodicity in the fumC mutant (Fig. 4A). It was also possible to alter wild-type HI4320 swelling with malate, which caused a more pronounced secondary raft (Fig. 4A, arrow). During the complete oxidative TCA cycle (Fig. 4B), furmarate catalyzes the oxidation of furmarate to malate during aerobic respiration; however, under anaerobic respiration, the TCA cycle operates in a reduced branched pathway during which the intermediate malate is reduced to furmarate by furmarase (Fig. 4C). Collectively, these findings also indicate that swelling differentiation relies on a complete, oxidative TCA cycle because succinate dehydrogenase is part of the aerobic pathway and the furmarase mutant was not affected by the addition of furmarate (Fig. 4A).

_P. mirabilis_ swelling is resistant to sodium azide, a poison of aerobic respiration. Mutation of the oxidative TCA cycle genes fumC and sdbB resulted in shorter periods of swelling. Succinate dehydrogenase can participate in membrane electron transfer during respiration. This predicts that swelling of wild-type strain HI4320 would be affected by adding sodium azide (Na3N) to poison aerobic respiration by irreversibly binding and inhibiting quinones involved in the transfer of electrons to oxygen. Despite the genetic requirement for the complete oxidative (aerobic) TCA cycle, we found that _P. mirabilis_ HI4320 swarms on growth inhibitory concentrations (0.005% [wt/vol]) of NaN3 at 37°C (Fig. 5A). This suggests that _P. mirabilis_ swelling is resistant to sodium azide and does not require aerobic respiration, supporting the notion that anaerobic or fermentation pathways can provide energy for swelling (18, 19).

The anaerobic branched TCA pathway mutant was also resistant to NaN3, and had a swelling phenotype identical to that of wild-type HI4320 on azide (Fig. 5A). In the absence of FrdA, a
Specific *P. mirabilis* central metabolism mutants cause aberrant swarming phenotypes. (A) Swarming phenotypes of wild-type HI4320, glycolysis mutants (A) Swarming phenotypes of wild-type HI4320, *pfkA* (phosphofructokinase) and *tpiA* (triose phosphate isomerase), glycolysis mutants, *fumC* (fumarase) and *sdhB* (succinate dehydrogenase) TCA cycle mutants, *frdA* (fumarate reductase) the anaerobic TCA cycle mutant. Mutations in *pfkA* and *tpiA* result in reduced swarming phenotype diameters, while *fumC* and *sdhB* TCA cycle mutants have reduced distances between swarming and consolidation which result in shorter consolidation rafts or rings. The anaerobic TCA cycle mutant has a similar swarming phenotype to that of wild-type HI4320. Complementation of *tpiA* (B), *pfkA* (C) and *fumC* (D) swarming phenotypes. *P. mirabilis* HI4320 containing pGEN empty vector and each mutant containing pGEN empty vector were compared to mutant-complemented strains. The complemented *tpiA* (pGEN+*tpiA*), *pfkA* (pGEN+*pfkA*), and *fumC* (pGEN+*fumC*) mutants all exhibit a restored swarming phenotype to that of wild-type HI4320. The vertical lines in panels A to D indicate the outer edge of the swarm.
component of the incomplete reductive TCA pathway, either a fermentative pathway or an aerobic pathway that uses succinate dehydrogenase in the complete TCA cycle, would likely operate, because the frdA mutant is unable to assemble an anaerobic respiratory chain using fumarate reductase (Fig. 4C). However, our findings demonstrate that an otherwise aerobic pathway is not using oxygen as a terminal electron acceptor during membrane respiration, because mutation of frdA does not render the bacteria susceptible to sodium azide during swarming. Mutation of fumC, which can function in both the anaerobic and aerobic pathways (Fig. 4B and C), severely inhibits swarming when respiration is poisoned by NaN₃, while swarming of the aerobic TCA cycle mutant sdhB was modestly affected by NaN₃ (Fig. 5A). Interestingly, swarming of wild-type strain HI4320 was inhibited when 0.02% (wt/vol) glucose was added to LB-1.5% agar containing NaN₃ (Fig. 5B), suggesting that in the presence of glucose, which promotes aerobic respiration when oxygen is present (22), P. mirabilis is unable to use its apparently preferred anaerobic-like electron acceptor and energy pathway required for swarming. The activity of azide on swarming was specific for membrane respiration with oxygen rather than acting as an uncoupler, because P. mirabilis was unable to swarm when the proton gradient was collapsed by adding the conventional uncoupler carbonyl cyanide m-chlorophenylhydrazone (CCCP) to LB-1.5% agar (data not shown).

Genetic screen to identify genes required for P. mirabilis to swim on sodium azide. To identify genes that encode or are required for the expression of alternative anaerobic-like respiratory chains, we tested P. mirabilis transposon mutants on LB-1.5% agar containing 0.005% (wt/vol) NaN₃ for the loss of swarming only when aerobic respiration is poisoned. Eighteen of 1,920 P. mirabilis transposon mutants consistently demonstrated decreased swarming diameters compared to that of wild-type HI4320 (Fig. 6B and E) on LB-1.5% agar containing NaN₃ (Fig. 6A). To eliminate motility-defective mutants, for example, those with disruptions in flagellar genes, all mutants were simultaneously tested for swarming on agar with and without NaN₃. Mutants with deficient swarming phenotypes on agar without NaN₃ were removed from the screen.

Transposon insertions were identified that disrupted genes encoding proteins involved in energy metabolism that are required during swarming (Table 2). Transposon mutant 37B5 has a disruption in gene argH, which encodes argininosuccinate lyase that catalyzes the breakdown of argininosuccinate to arginine and fumarate (Table 2). Mutant 33H5 has an insertion that disrupts fumC, which encodes the TCA cycle enzyme fumarase (Fig. 6A). Identification of fumarase from these experiments validated our azide swarming screen since the de novo fumC mutant (Fig. 6C) and 33H5 (Fig. 6D) displayed the same aberrant swarming phenotype (see Fig. S2 in the supplemental material), and both are unable to swarm on NaN₃, a poison of aerobic respiration (Fig. 6F and G). The transposon insertion in mutant 25H1 (Table 2) disrupts the hybB gene, which encodes hydrogenase-2, an anaerobic cytochrome that catalyzes the oxidation of hydrogen into two protons and two electrons during fermentation. Mutant 11F4 disrupts P. mirabilis transposon HI4464, which encodes a protein that has 75% identity to a quinone hydroxylase of Providencia rettgeri (Table 2). Interestingly, the azide-defective swarming mutant 31H4 has a disruption in rnh, which encodes an exonuclease II. This gene, however, is located approximately 1 kb downstream of genes encoding a predicted electron transport system, RnfABCDGE.

In the azide swarming screen of P. mirabilis transposon mutants, genes encoding respiratory enzymes and genes unrelated to energy metabolism were both identified as a result of NaN₃ sensitivity (Table 2). Although we expected to identify genes related to alternative respiratory chains or fermentative functions (discussed above), we were not surprised to also find disrupted genes that increased susceptibility to azide but are unrelated to swarming. In addition, we expected to find insertions in genes encoding cell wall machinery, LPS biosynthesis, and capsule production, because an altered permeability barrier might increase susceptibility to sodium azide. Mutants with decreased swarming diameters identified on NaN₃ (Fig. 6A) included transposon mutant 25A2 in yadF, which encodes an oxidation subunit involved in fatty

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**FIG 3** P. mirabilis HI4320 oxidative TCA cycle mutants display shorter swarming distances between consolidation rafts in comparison to wild-type HI4320. To quantify the aberrant swarming phenotype, swarm agar plates were inoculated with a 5-μl volume from one overnight culture of wild-type P. mirabilis HI4320 and fumC, sdhB, frdA, 33H5 (fumC), and 10E4 (sdhC) mutants. Plates were incubated for 18 h, and the width of each swarming raft was measured in millimeters. The white-shaded portion of the bar represents the averaged measurements of the first swarm raft (R1), the gray-shaded portion of the bar represents the averaged measurements of the second raft (R2), and the black-shaded portion of the bar represents the averaged measurements of the third raft (R3). In panel A, the bar graph displays the measurements from a total of 5 inoculated agar swarm plates for each strain following incubation at 30°C. In panel B, measurements were taken from a total of 10 inoculated agar plates of each strain following incubation at 37°C, and in panel B, measurements were taken from a total of 10 inoculated agar plates of each strain following incubation at 30°C. 33H5 was identified in the present study; in an earlier study (see reference 21), 10E4 was shown to produce aberrant swarming patterns on agar.
acid metabolism (Table 2); 30B6 in PMI3180, which is adjacent (11 kb downstream) to the LPS O-antigen biosynthetic gene cluster; 25B6 with an insertion in the colonic acid capsular biosynthesis activation protein encoded by *rcsA*; and 35F4 in *mrcA*, which encodes a peptidoglycan synthetase (Table 2). Insertions in a hemagglutinin encoded by PMI2914; an FKBP-type peptidyl-prolyl cis-trans-isomerase encoded by *fkpA*; a periplasmic peptide transport protein; encoded by *sapA*; and mutant 23D6, in which the transposon was located in an intergenic region between *acrB* and *ybaJ* (Table 2) were also found in the screen (Fig. 6A).

**Fermentation end products are not excreted during swarming.** By screening *P. mirabilis* transposon mutants for loss of swarming on NaN₃, a poison of aerobic respiration, we identified *hybB* and *argH*, which encode proteins involved in alternative anaerobic processes potentially required to generate energy for *P. mirabilis* swarming. However, the identification of an insertion in *fumC* supports our assertion that the enzyme is participating in the oxidative TCA cycle during *P. mirabilis* swarming. To test whether *P. mirabilis* swarming is also dependent on fermentation or is using a less canonical respiratory chain in conjunction with

FIG 4 LB-1.5% agar replete with TCA cycle intermediates alters the interval between swarming and consolidation of *P. mirabilis* TCA cycle mutants. (A) Wild-type *P. mirabilis* HI4320 and the *fumC* and *sdhB* aerobic TCA cycle mutants were examined on LB-1.5% agar alone or agar containing either 10 mM fumarate, 10 mM succinate, or 10 mM malate. All three TCA cycle intermediates cause the *sdhB* mutant to revert to a wild-type swarming phenotype, while only succinate and malate cause reversion to the wild-type swarming phenotype for the *fumC* mutant. The addition of malate causes the formation of a pronounced secondary raft for wild-type HI4320 (arrow). Complete oxidative (B) and incomplete reductive (C) TCA cycle pathways indicating the relevant enzymes, intermediates, and direction of reactions during aerobic and anaerobic processes, respectively.
the complete aerobic TCA cycle, we used the pH indicator phenol red in LB-1.5% agar to examine swarming of \textit{P. mirabilis} under aerobic and anaerobic conditions, because excretion of acidic end products is indicative of fermentation. A change in pH was determined by a visible color change of the agar from orange (neutral pH) to pink (alkaline pH) or to yellow (acidic pH). Under aerobic conditions at 30°C and 37°C, \textit{P. mirabilis} HI4320 swarming created an alkaline environment (Fig. 7A), raising the surrounding pH, suggesting that the bacteria are not excreting acidic fermentation end products during swarming differentiation.

Although no exogenous urea was added to the plates, to ensure that the increase in pH was not the result of urease hydrolyzing urea to CO₂ and ammonia, \textit{P. mirabilis} mutations in \textit{ureC} and \textit{argG}, required for urease activity and the urea cycle, respectively, were examined on the phenol red agar (see Fig. S3 in the supplemental material). Both the urease mutant and \textit{argG} mutant bacteria exhibited a similar color change of the agar from orange to pink, indicating alkaline conditions and suggesting that the formation of ammonia by urease was not responsible for the increase in pH by wild-type HI4320 (see Fig. S3).

Swarming of \textit{P. mirabilis} under anaerobic conditions on phenol red agar, however, resulted in agar turning yellow, indicating an acidic pH (Fig. 7B). These results suggest that under anaerobic conditions, \textit{P. mirabilis} is swarming using fermentation when a membrane electron acceptor is lacking. It is notable that both \textit{argH}, which was identified in the screen (Table 2), and \textit{argG} (see Fig. S3) are arginine auxotrophs, yet only \textit{argH} had a defect in swarming on sodium azide, suggesting that it is the production of fumarate rather than arginine biosynthesis itself that is important for the energetics that govern swarming.

To better understand how fermentation and respiration produce energy during swarming, carbon source glycerol or glucose, with and without fumarate as an energy source, was added to agar containing phenol red. Under aerobic conditions at 37°C, wild-type HI4320 and \textit{sdhB}, \textit{frdA}, and \textit{hybB} mutants alkalinized the phenol red agar to turn pink (Fig. 7A). Interestingly, when glycerol or glucose was added to the phenol red agar, the \textit{frdA} and \textit{sdhB} mutant bacteria acidified the agar, indicating fermentation (Fig. 7A). Addition of fumarate to the phenol red agar containing either glycerol or glucose rescued the pH of the \textit{sdhB} mutant to an alkaline pH like that of the wild type (Fig. 7A). Under aerobic conditions, the \textit{hybB} and \textit{frdA} mutants remained alkaline under all conditions (Fig. 7A). Under anaerobic conditions at 37°C, wild-type HI4320 and \textit{sdhB}, \textit{frdA}, and \textit{hybB} mutants were able to ferment and acidify the phenol red agar (Fig. 7B). After anaerobic incubation, plates were exposed to aerobic conditions for approximately 6 h, and wild-type HI4320 and the \textit{hybB} and \textit{frdA} mutants slowly turned the medium alkaline; however, the phenol red agar, plated with the \textit{frdA} and \textit{sdhB} mutants, remained acidic (Fig. 7C). While swarming of the \textit{sdhB} mutant could be rescued to wild-type pH by the addition of exogenous fumarate, independent of carbon source and dependent on oxygen, \textit{frdA} mutant bacteria were rescued to wild-type pH by glycerol, independent of fumarate, under anaerobic conditions (Fig. 7D). Taken together, these findings provide evidence that during swarming, the oxidative TCA cycle is required, because loss of fumarate reductase did not affect swarming under any condition tested, and fumarate was unable to rescue \textit{frdA} mutant bacteria pH or periodicity during swarming.

**DISCUSSION**

Swarming is a coordinated and multicellular behavior that is dependent upon respiration to generate a proton motive force that...
drives the rotation of bacterial flagella. Remarkably, *P. mirabilis* swarm cells are deficient in aerobic cytochromes (19) and swarm on inhibitory growth concentrations of sodium azide, an inhibitor of aerobic respiration. A reduction in membrane cytochromes in swarm cells is surprising, because bacterial propulsion, mediated by rotating flagella, is dependent on proton motive force. Mutations that disrupt genes encoding the TCA cycle enzymes succinate dehydrogenase and fumarase cause an aberrant swarming phenotype; however, disruption of the anaerobic, incomplete reductive TCA cycle enzyme fumarate reductase has no effect on the swarming process. This finding is unexpected, because, in most bacteria, succinate dehydrogenase functions in the complete TCA cycle when oxygen is available as a terminal electron acceptor, while fumarate reductase catalyzes the reverse reaction in the incomplete, reductive TCA cycle in the absence of aerobic respiration. Since we and others (19) observed that swarming is resistant to sodium azide, which inhibits the electron transfer to oxygen, this suggests that *P. mirabilis* is oxidizing pyruvate using the complete TCA cycle in conjunction with membrane respiration that lacks transfer of electrons to oxygen. Collectively, these findings and our results suggest that *P. mirabilis* employs an alternative anaerobic-like electron transport chain to reoxidize NADH from reduced NADH that is produced during the oxidation of pyruvate to CO₂ in the complete TCA cycle and to generate proton motive force to drive flagellar rotation (Fig. 8).

To elucidate potential components of an alternative anaerobic-like electron transport chain, we screened a collection of *P. mirabilis* transposon mutants to test for the loss of swarming only on sodium azide, a poison of the aerobic respiration chain. We found 18 mutants with a wild-type swarming phenotype on LB-1.5% agar alone and swarming phenotypes of wild-type HI4320 (B), fumC mutant (C), and 33H5 mutant (D) on agar containing 0.005% (wt/vol) NaN₃. The gene disrupted in 33H5 (fumC), has the same swarming phenotypes as the de novo fumC mutant (C). The vertical lines in panels B to G indicate the outer edge of the swarm.

**FIG 6** Identification of *P. mirabilis* genes specifically required for swarming on sodium azide. (A) Eighteen out of 1,920 *P. mirabilis* transposon mutants were sensitive to 0.005% (wt/vol) sodium azide (NaN₃) and displayed decreased swarming phenotypes during aerobic conditions at 37°C (black bars). The mutants selected in this screen swarm at 37°C in the absence of NaN₃ (gray bars). A previously identified transposon mutant, 8B5, was included as a nonswarming control. Swarming phenotypes of wild-type HI4320 (B), fumC mutant (C), and 33H5 mutant (D) on LB-1.5% agar alone and swarming phenotypes of wild-type HI4320 (E), fumC mutant (F), and 33H5 mutant (G) on agar containing 0.005% (wt/vol) NaN₃. The gene disrupted in 33H5 (fumC), has the same swarming phenotypes as the de novo fumC mutant (C).
TABLE 2 Identification of *P. mirabilis* genes required for swarming on sodium azide at 37°C

| Mutant  | ORF       | Gene   | Annotation                               |
|---------|-----------|--------|------------------------------------------|
| 37B5    | PMI3240   | argH   | Argininosuccinate lyase                  |
| 33H5    | PMI1296   | fumC   | Fumarase                                 |
| 25H1    | PMI0033   | hybB   | Hydrogenase-2b cytochrome subunit        |
| 25A2    | PMI1807   | fadF   | Fatty acid oxidation complex subunit alpha|
| 11F4    | PMI2646   |        | Ubiquinone biosynthesis hydroxylase      |
| 31H4    | PMI1311   | rnb    | Exoribonuclease II                       |
| 31F4    | PMI2801   | fkpA   | FKBP-type peptidyl-prolyl cis-trans-isomerase|
| 28D2    | PMI2495   | holD   | DNA polymerase subunit psi               |
| 11D3    | PMI143    |        | DNA modification methyltransferase       |
| 6C5     | PMI2669   | pacB   | Putative periplasmic member of the SGNH-family of hydrolases |
| 6C6     | PMI548    | citC   | TonB-dependent receptor                  |
| 30B6    | PMI180    |        | M23B family outer membrane metalloprotease|
| 34A1    | PMI378    | sapA   | Peptide transport periplasmic protein    |
| 35F4    | PMI3021   | mrcA   | Peptidoglycan synthetase                 |
| 39H1    | PMI2993   | idsD   | Hypothetical protein                     |
| 23D6    | Intergenic|        | 400-bp 3' end of acrB, 200-bp 5' PMI0130/31 biofilm regulator ybal/hemolysin modulating protein |
| 25B6    | PMI1680   | rcsA   | Colanic acid capsular biosynthesis activation protein |
| 26B3    | PMI2914   |        | Hemagglutinin                            |

agar but that displayed reduced swarming diameters on sodium azide; five mutants (33H5, 25H1, 11F4, 37B5, and 25A2) have disruptions in genes having an obvious association with respiration. The identification of fumarase, encoded by *fumC*, validated the azide swarming screen, because we previously noted that our isogenic *fumC* mutant was unable to swarm on sodium azide. Fumarase catalyzes the conversion of fumarate to malate during aerobic conditions and malate to fumarate during anaerobic conditions. Since succinate dehydrogenase is operating in the complete TCA cycle, converting fumarate to proton motive force, we believe that fumarase is most likely also operating in the complete TCA cycle, converting fumarate to malate. In addition, the enzyme argininosuccinate lyase, encoded by *argH*, was identified in the azide swarming screen and is required for the production of arginine and fumarate. Since a mutant was unable to swarm on sodium azide, we previously noted that our isogenic *fumC* mutant was unable to swarm on sodium azide. Fumarase catalyzes the conversion of fumarate to malate during aerobic conditions and malate to fumarate during anaerobic conditions. Since succinate dehydrogenase is operating in the complete TCA cycle, converting fumarate to proton motive force, we believe that fumarase is most likely also operating in the complete TCA cycle, converting fumarate to malate. In addition, the enzyme argininosuccinate lyase, encoded by *argH*, was identified in the azide swarming screen and is required for the production of arginine and fumarate. Since a mutation in *argG*, which encodes argininosuccinate synthase, is also an arginine auxotroph but does not have a defective swarming phenotype on azide, the activity of argininosuccinate lyase encoded by *argH* is for production of the TCA cycle intermediate fumarate required for respiration.

*hybB*, which encodes hydrogenase-2, a known anaerobic respiratory chain component, catalyzes the extracytoplasmic oxidation of dihydrogen into two protons and two electrons, often during fermentation (23). The protons remain in the periplasm of the bacterium, while the electrons are shuttled to a quinone in the electron transport chain of the inner membrane. Simultaneously, two protons located in the bacterial cytoplasm are consumed by hydrogenase-2. The conservation of protons mediating by hydrogenase-2 is consistent with the hypothesis that *P. mirabilis* swarming is supported by an anaerobic-like respiratory chain, and the identification of hydrogenase-2 in the azide screen suggests that extracytoplasmic energy pathways are important for the physiology of *P. mirabilis* swarming differentiation (24). Our findings do not address the possibility that *P. mirabilis* reverses the F$_{1}$F$_{0}$ ATP synthase to pump protons during swarming; however, expending ATP generated by substrate-level phosphorylation during fermentations would generate proton motive force (Fig. 8).

An essential component of electron transport chains are hydrophobic quinones that carry electrons and protons during membrane respiration and act as redox mediators linking dehydrogenases and terminal reductases within the electron transport chain. Interestingly, a transposon mutant was identified in the *P. mirabilis* azide swarming screen that has 75% identity to a quinone hydroxylase of *P. rettgeri*. *Escherichia coli* and related enteric bacteria are known to synthesize three different quinones: ubiquinone, menaquinone, and demethylmenaquinone. During aerobic respiration, ubiquinone is preferred, while during anaerobic conditions, menaquinone is used. Our findings suggest that *P. mirabilis* could use this predicted quinone hydroxylase to maintain pools of substrates to transfer electrons using the proposed anaerobic-like respiratory chain during swarming.

The fate of NADH, generated from glycolysis and primarily the TCA cycle, is dependent on which terminal electron acceptor is present: oxygen under aerobic conditions or an organic compound under anaerobic conditions. During glycolysis, triose phosphate isomerase, encoded by *tpiA*, reversibly isomerizes the ketone sugar dihydroxyacetone phosphate (DHAP) to the aldehyde sugar glycerol-3-phosphate (G3P). We have shown that the isogenic *tpiA* mutant results in a swarming-deficient phenotype, suggesting that both of these glycolysis intermediates, DHAP and G3P, are important during *P. mirabilis* swarming. Although not identified in the swarming azide screen, another known anaerobic respiratory chain component, glycerol-3-phosphate dehydrogenase, converts G3P to DHAP. It is possible that *P. mirabilis* utilizes glycerol-3-dehydrogenase to generate energy, because the *tpiA* mutant is unable to swarm and cannot reversibly isomerize DHAP and G3P. During anaerobic conditions, glycerol can be used as an energy source when fumarate is present as a terminal electron acceptor for fumarate reductase. Our findings show that fumarate reductase is not important for *P. mirabilis* swarming differentiation; however, future studies may reveal that succinate dehydrogenase fulfills this role.

In respiring cells, NADH and FADH$_{2}$ are generated during metabolism during oxidation steps of the TCA cycle. To reoxidize these molecules, protons and electrons are shuttled via the electron transport chain within the bacterial cytoplasmic membrane.
As electrons are transported through a series of membrane flavoprotein carriers, to a terminal electron acceptor, protons are translocated across the membrane. The proton gradient that is created during respiration contributes to the gradient of both pH and charge across the membrane and collectively determines proton motive force. It is during aerobic conditions with O₂ as a terminal electron acceptor that the largest number of protons can be pumped, thereby producing the greatest proton motive force for the cell. During fermentation, however, bacteria are unable to respire and can generate a proton gradient by reversing the rotation of the F₀F₁ ATPase to consume ATP to pump protons out from the cytoplasm. Consequently, generation of proton motive force is

**FIG 7** Oxidative TCA cycle mutants excrete acidic fermentation end products under aerobic conditions. The *hybB*, *frdA*, *fumC*, and *sdhB* mutants and wild-type HI4320 were observed on phenol red LB-1.5% agar with and without 0.2% (wt/vol) glycerol (gly) or 0.2% (wt/vol) glucose (glc) as carbon sources and 0.2% (wt/vol) fumarate (F) as an energy source where indicated. Inoculated plates were incubated at 37°C under aerobic conditions (A) and anaerobic conditions (B). In addition, fully developed swarm plates from anaerobic experiments were observed over time following exposure to aerobic conditions. Anaerobic plates were then maintained under aerobic conditions at room temperature and viewed at 6 (C) and 48 (D) h. Following approximately 6 h under aerobic conditions, the phenol red agar of wild-type HI4320 and the *hybB* and *frdA* mutants slowly turned alkaline; however, the *fumC* and *sdhB* mutants were still visibly acidic. Following approximately 48 h under aerobic conditions, all of the phenol red agar plates returned to alkaline conditions except for the *fumC* mutant when glucose was present as a carbon source. A color map depicting the color range from alkaline pH (pink) to acidic pH (yellow) is displayed bottom left. To the right of the color map is a reference image of an uninoculated agar plate following incubation under aerobic conditions at 37°C.
force during anaerobic conditions is substantially more difficult for the cell compared to that during aerobic conditions (25).

Interestingly, TCA cycle intermediates also can affect patterns formed by motile colonies of *E. coli* and *Salmonella* (26, 27). The ability of succinate and fumarate to affect patterns in these bacteria has been suggested to result from the buildup of chemoattractant or chemorepellent (26, 28); however, we propose an alternative explanation: that manipulating concentrations of TCA cycle intermediates affects periodic swarming due to their effect on energy metabolism. This is likely true for *P. mirabilis* multicellular swarming, since active swarms transplanted onto fresh agar maintain their same periods (3). The connection between energy pathways that generate proton motive force and swarming has broad implications, because surface motility is a behavior shared by many bacteria (28, 29). Our findings suggest that these universal patterns of swarming are dictated by membrane energetics and imply that bacterial cooperation during the development of multicellular swarming decreases the energy required by individual bacteria to travel across a surface.

** MATERIALS AND METHODS **

** Bacterial strains.** *P. mirabilis* HI4320 was cultured from the urine of a patient presenting with bacteriuria during long-term catheterization (30, 31). *P. mirabilis* HI4320 mutants (Table 1) were generated using the Tar-geTron gene knockout system (Sigma). The *P. mirabilis* tpiA mutant was complemented with a wild copy of the tpiA gene cloned into pGEN-MCS. The ureC (urease) and argG (argininosuccinate synthase) mutants were used as controls for phenol red experiments (32).

** Culture media and swarming phenotype testing.** *P. mirabilis* was routinely cultured in LB medium. Swarming phenotypes of *P. mirabilis* mutants were compared with the wild-type HI4320 swarming phenotype on LB-1.5% (wt/vol) agar (10.0 g/liter NaCl) by spotting 5 μl of overnight culture onto the center of a plate, incubating under aerobic conditions at 37°C, and examining after 18 h. *P. mirabilis* HI4320 and TCA cycle mutants were examined under the same conditions on LB-1.5% agar containing a TCA intermediate: 10 mM succinate, 10 mM fumarate, or 10 mM malate. *P. mirabilis* HI4320, the TCA cycle mutants, and *P. mirabilis* transposon mutants were also examined on LB-1.5% agar containing inhibitory concentrations of sodium azide, 0.01% (wt/vol) and 0.005% (wt/vol), and incubated under aerobic conditions at 30°C and 37°C. Strain HI4320 was also tested on LB-1.5% agar containing sodium azide and 0.2% (wt/vol) glucose. *P. mirabilis* HI4320, the ureC mutant, and other metabolism mutants were examined on LB-1.5% agar containing 0.04 g/liter phenol red following incubation under aerobic and anaerobic (BD GasPak EZ Anaerobe) conditions at 30°C. LB-1.5% agar containing phenol red and 0.2% (wt/vol) glucose. *P. mirabilis* HI4320, the ureC mutant, and other metabolism mutants were examined on LB-1.5% agar containing 0.04 g/liter phenol red following incubation under aerobic and anaerobic (BD GasPak EZ Anaerobe) conditions at 30°C. LB-1.5% agar containing phenol red and 0.2% (wt/vol) glucose. *P. mirabilis* HI4320, the ureC mutant, and other metabolism mutants were examined on LB-1.5% agar containing 0.04 g/liter phenol red following incubation under aerobic and anaerobic (BD GasPak EZ Anaerobe) conditions at 30°C. LB-1.5% agar containing phenol red and 0.2% (wt/vol) glucose. *P. mirabilis* HI4320, the ureC mutant, and other metabolism mutants were examined on LB-1.5% agar containing 0.04 g/liter phenol red following incubation under aerobic and anaerobic (BD GasPak EZ Anaerobe) conditions at 30°C. LB-1.5% agar containing phenol red and 0.2% (wt/vol) glucose.

** FIG 8 ** Proposed model for *P. mirabilis* energy metabolism during swarming. The rotation of flagellum and oxidative phosphorylation are dependent on the proton gradient that is generated by membrane respiration to oxidize NADH to NAD⁺. The majority of NADH is formed during the complete oxidation of pyruvate to CO₂ in the TCA cycle. The decreased interval between swarming and consolidation for aerobic TCA cycle mutants suggests that the capacity to reoxidize NAD⁺ and establish a proton gradient is due to inappropriate fermentation caused by loss of the complete TCA cycle and by the diminished capacity to metabolize fumarate. Extracytoplasmic substrate oxidation of dihydrogen and cytoplasmic reduction of glycerol could maintain the proton gradient and allow for reoxidation of NAD⁺ in the absence of electron transfer to oxygen. The ability to support motility without respiring oxygen may reflect the reduced energy demand caused by multicellular cooperation during swarming.
Sodium azide screen. Swarming phenotypes of 1,920 *P. mirabilis* transposon mutants, previously created by us (21), were compared with the wild-type HI4320 swarming phenotype on LB-1.5% agar with and without 0.005% (wt/vol) sodium azide (NaN₃), an inhibitor of electron transport. Swarming diameters were measured and documented from triplicate experiments performed on separate days. Transposon mutants unable to swarm on LB-1.5% agar alone were removed from further testing in the NaN₃ screen. Arbitrary PCR and molecular cloning into pCR2.1-TOPO (Invitrogen) was carried out as previously described (21, 33) on *P. mirabilis* transposon mutants with a reduced production diameter or inhibited swarming phenotype on LB-1.5% agar containing NaN₃ compared to HI4320 following incubation at 37°C under aerobic conditions. For these experiments, transposon mutant 8B5 with a disruption in aceF (pyruvate dehydrogenase), which is swarming deficient, was included as a nonswarming control.

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at http://mbio.asm.org/lookup/suppl/doi:10.1128/mBio.00365-12/-/DCSupplemental.

Figure S1, TIF file, 2.3 MB. Figure S2, TIF file, 3.1 MB. Figure S3, TIF file, 1.6 MB.

ACKNOWLEDGMENTS

This work was supported in part by Public Health Service grants AI43363 and AI59722 from the National Institutes of Health.

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