Gliding motility and polarized slime secretion

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Summary
Myxococcus leaves a trail of slime on agar as it moves. A filament of slime can be seen attached to the end of a cell, but it is seen only at one end at any particular moment. To identify genes essential for A motility, transposon insertion mutations with defective A motility were studied. Fifteen of the 33 mutants had totally lost A motility. All these mutant cells had filaments of slime emerging from both ends, indicating that bipolar secretion prevents A motility. The remaining 18 A motility mutants, also produced by gene knockout, secreted slime only from one pole, but they swarmed at a lower rate than A+ cells and are called ‘partial’ gliding mutants, or pgl. For each pgl mutant, the reduction in swarm expansion rate was directly proportional to the reduction in the coefficient of elasticotaxis. The pgl mutants have a normal reversal frequency and normal gliding speed when they move. But their probability of movement per unit time is lower than pgl+ cells. Many of the pgl mutants are produced by transposon insertions in glycosyltransferase genes. It is proposed that these glycosyltransferases carry out the synthesis of a repeat unit polysaccharide that constitutes the slime.

Introduction
Many bacteria glide; they translocate across surfaces without the aid of flagella, and they cannot swim (Henrichsen, 1972). How is gliding propelled? Some bacteria glide by retracting polar type IV pili, as reviewed (Nudleman and Kaiser, 2004). Other gliders lack type IV pili: members of the cytophaga-flavobacterium group and the mycoplasmas (for example, McBride, 2001). Moreover, mutants of Myxococcus xanthus that have lost their ability to produce type IV pili, and hence their S motility, continue to glide. They use what has been called adventurous or A motility, the subject of this report (Hodgkin and Kaiser, 1979a,b).

The molecular basis of A motility has long been sought. Observing trails of slime, Jahn suggested that myxobacteria might be propelled by slime secretion, but he questioned whether slime was capable of pushing cells along (Jahn, 1924). Hodgkin approached the problem by isolating mutants specifically deficient in A motile gliding as opposed to S motile gliding (Hodgkin and Kaiser, 1979a). When the amino acid sequences of A motility genes became available, they revealed many proteins: outer membrane lipoproteins (Rodriguez and Spormann, 1999), transport proteins, proteases and peptidases (Youderian et al., 2003). But the ensemble of proteins failed to suggest a mechanism.

An electron microscopic examination of slime secretion associated with gliding was undertaken in Phormidium, a filamentous gliding cyanobacterium (Hoiczyk and Baumeister, 1998). Hoiczyk and Baumeister observed the extrusion of slime from junctional pore complexes in the cell wall, and correlated the rate of slime extrusion with the speed of movement of the cyanobacterial filaments. The junctional pore complexes were pore-shaped organelles embedded in the cell wall near the sites of slime extrusion (Hoiczyk and Baumeister, 1998). Recently, ribbons of an amorphous material, taken to be slime, were photographed emerging from the ends of M. xanthus cells (Wolgemuth et al., 2002). Next to the cell, the ribbons were thin, while farther from the cell they coalesced into a single ribbon having the width of the cell (Wolgemuth et al., 2002). Strikingly, the ribbons emerged uniquely from one pole of the cell; the other pole was devoid of ribbons.

A+ cells (both A+S+ and A+S−) are found either depositing slime as they glide on agar, or gliding on a trail left by another cell (Wolgemuth et al., 2002). A+ gliding cells seem to prefer to follow a slime trail previously laid down than to lay a new trail (Burchard, 1982). Many examples of individual myxobacterial cells turning to follow a slime trail are shown in Reichenbach’s time lapse movies (Reichenbach et al., 1965). The deposition and the following of slime trails are independent of the presence or absence of pili or S motility in M. xanthus (Wolgemuth et al., 2002). Burchard showed that Myxococcus can also follow the slime trails of other gliders like Beggiatoa and Oscillatoria (Burchard, 1982). Elasticotaxis denotes the ability of Myxococcus cells to orient their movement along lines of elastic stress in an agar gel that has been
stretched or compressed (Stanier, 1942). Elasticotaxis is related to slime trails by the fact that it is restricted to A motile cells. Moreover, A motile cells display less elasticotaxis when they also exhibit S motility (Fontes and Kaiser, 1999).

In this report, we suggest that A motility, the following of slime trails, and elasticotaxis are different manifestations of the secretion, swelling and alignment of polymeric slime. In support of that suggestion, we report two new classes of mutants with defects in A motility. One class produces slime from both ends of the cell and is non-motile. The other class of gene knockout mutants diminishes A motility to different degrees, but does not abolish it.

Results

Isolating A motility mutants

To explore the connection between A motility and slime secretion, we isolated a random set of A motility mutants and examined their ability to secrete slime. Two A’S- strains were exposed to the Himar transposon, and drug-resistant strains that formed small colonies were isolated. Among 32 612 kanamycin-resistant transformants, 33 strains that reproducibly formed smaller colonies than the parental strains were confirmed to have defects in A motility by microscopic examination of the edge of their colonies. Fifteen of the mutants lacked the thin and crenellated edges indicative of swarm spreading. The remaining 18 mutants retained some A motility, but it was noticeably less than the parents.

Visualizing slime secretion

Many A+ cells depositing and following slime trails are shown in Fig. 1A and B. Figure 1C–H show several frames of a movie made by Lars Jelsbak of two well-separated gliding cells. Each leaves a slime trail, reverses several times, and with each reversal follows, extends and thickens its own trail. The trail elongates progressively at both ends of these reversing cells, clearly demonstrating a moment by moment correlation between movement and the deposition of slime. Slime has been stained with acridine orange, a non-specific fluorescent dye (Wolgemuth et al., 2002). In the present study, extrusions from A+ cells were visualized without staining by means of differential interference contrast (DIC) microscopy. The refractive index difference between slime and the glass of a microscope slide renders the slime visible, and is consistent with a polysaccharide gel. Figure 2 shows representative cells, each of them have a single ribbon of slime in line with the axis of the cell, and only at one end of each cell. None of the more than 200 wild-type (A+ S+) cells examined critically had ribbons at both ends, in agreement with the observations on

Fig. 1. M. xanthus lays down slime trails and follows them. Frames A and B show the swarm front of the A’S- strain DK10410 on agar photographed after 1 day at 32°C. Photographs taken with a Leitz 16×, phase contrast objective. Frames C–H show two cells of the A’S- strain, DK1622, gliding on agarose. Photographs taken with a Nikon 40×, phase contrast objective by Dr Lars Jelsbak.

Fig. 2. Visualization of slime extruded from individual wild-type M. xanthus cells. Panel A, A’S- DK1622; panel B, A’S- DK10410. Cells that had been deposited on the bottom of a plastic tissue culture plate were incubated overnight. After gentle harvest, the cells were transferred to a microscope slide. They were photographed at 100× under oil immersion using a DIC objective.
acridine-stained cells (Wolgemuth et al., 2002). At any instant, slime secretion from normal cells was always polar and restricted to one end of each cell.

By contrast, the seven non-motile mutants isolated from DK10410 and one from DK8615 extruded slime from both poles (Fig. 3). These strands of slime were thinner, more curved than the A\(^+\) slime, and the slime filaments tended to angle away from the long axis of the cell. We suggest that bipolar extrusion causes the loss of A motility. Finally, the mutants with partial A motility showed unipolar extru-

Fig. 3. Slime distributions in the non-motile (A\(^+\)S\(^-\)) mutants. Cells were prepared and examined as described in the legend to Fig. 2.
A. agmK::TF (DK13001).
B. agmA::TF (DK13002).
C. mglA::TF (DK13003).
D. agmX::TF (DK13004).
E. agnB::TF (DK13005).
F. agnC::TF (DK13006).
G. aglU::TF (DK13007).
H. cglB::TF (DK13008).

Fig. 4. Slime distributions in the partial A motile mutants. Cells were prepared and examined as described in the legend to Fig. 2.
A. pglB::TF (DK13010).
B. pglC::TF (DK13011).
C. pglE::TF (DK13013).
D. pglF::TF (DK13014).
E. pglH::TF (DK13016).
F. pglU::TF (DK13018).
G. mglB::TF (DK13020).
H. pglK::TF (DK13021).
sion (Fig. 4). We refer to these mutants as *pgl*, to indicate their partial gliding motility.

**Quantifying partial A motility**

To investigate why the *pgl* mutants form smaller colonies, their rates of swarm expansion were measured. Swarm expansion rates summarize the net outward movement of thousands of cells, mostly in a single layer, which constitutes the edge of a colony of motile cells. Figure 5 shows that swarms of 14 *pgl* mutants expanded at rates from just slightly greater than DK11316 (A−S−) to almost as large as DK10410 (A+S−). (Four of the 18 total mutants were unavailable for testing at the time of this experiment.) DK10410, one of the parental A+S− strains, possesses full A motility. DK11316, an A−S− strain that carries null mutations in *cglB* and in *pilA* is non-motile and a negative control. It had the slowest rate, which measures colony expansion solely due to growth and cell division. All strains had similar growth rates measured in liquid culture (data not shown). Each *pgl* swarm expanded at a roughly constant rate (Fig. 5A and B), but each rate was lower than the A+ and higher than the A− controls, and the rates spread continuously across the available range.

To test the possibility that partial A motility might arise from a polar effect of the transposon insertion on a downstream motility gene, an in-frame deletion mutant was constructed for comparison. The *pglJ* gene, which is adjacent to genes that might be co-transcribed with it, was deleted and the rate of swarm expansion was measured. The Δ*pglJ* strain had a swarm rate that is 63% of a fully A-motile strain (DK10410), compared with the 76% of the *pglJ*::TF (the pMiniHimar-*lacZ* insertion mutant) (Fig. 5C). These data show that this mutant’s partial motility is not caused by polarity on a downstream gene but is more likely the direct consequence of the total inactivation of *pglJ*. The possibility that the transposon had inserted into the C-terminal region of each of the *pgl* genes leaving some residual function was examined through sequencing the myxobacterial DNA adjacent to the transposon in each mutant. It is indicated in the sequence section below that each of the insertions would have been expected to inactivate their target gene.

Cells at the leading edge of a swarm have different orientations which direct their movement, and that movement is sporadic: cells move, stop and start again (Jelsbak and Søgaard-Andersen, 1999). Frequently after stopping, a cell reverses its gliding direction (Kuhlwein and Reichenbach, 1968; Blackhart and Zusman, 1985). To see how *pgl* function contributes to swarm expansion, time-lapse movies were made of *pgf* and *pgf* strains at low cell density in order that individual cells could be tracked. Three parameters of movement were measured on individual cells: the fraction of cells in a given microscopic field that moved in 15 min, termed the movement probability; the reversal frequency; and the cell speed when they were moving. Comparing A+ with *pgl* discriminated between several possible explanations for a reduction in swarm rate. First, *pgl* cells might pause for a longer time between moves than A+ cells. Indeed, all five *pgl* mutants examined had half or less the probability of movement than the A+ strain, indicating longer pause times (Table 1).

Fig. 5. Expansion of A+S− and *pgl* mutant swarms. The radius of the swarm on CTT agar plates incubated at 32°C is shown as a function of time.

A and B. Expansion rates of the partial A motile *M. xanthus* strains are compared with the A+S− strain DK10410 and the non-motile A−S− strain DK11316. Only the partial A motile mutants generated in DK10410 background were used in this experiment.

C. Swarm expansion rates of the in-frame deletion mutant Δ*pglJ* (DK13024) and the transposon-inserted mutant *pglJ*::TF (DK13018) are compared with DK10410 as a control.
Second, a lower swarm expansion rate might result from a change in the number of reversals per unit time (reversal frequency). Cells with a very high reversal frequency would be expected to have a lower swarm rate because they would spend more time reversing than moving. Reversal frequencies were measured on isolated cells of A⁺ and five pgl mutants in the time-lapse movies. The mutants were found to have the same reversal frequency, within experimental error, as the A⁺ in Table 1, except for pglH, which had twice the reversal frequency of A⁺. A third possibility is that each cell might move more slowly when it does move, but no significant speed differences were detected (Table 1). The average speed of A⁺ (DK10410) cells might be slightly greater than either the pglB or pglE mutant, which could be an indirect consequence of their lower movement probabilities.

Finally, the possibility of a pgl effect on cell flexibility or cell–cell cohesion that would indirectly affect movement was considered. Gliding cells bend and change their gliding direction when they strike another cell or other impediment on the agar surface. If pgl mutants bent more easily or cohered differently to each other than the A⁺, a qualitatively different distribution of cells at the edge of the swarm might be expected. To explore these and related possibilities, the time-lapse movies of the edge of swarm zones of a partially motile mutant were compared with those of A⁺ S⁻ and A⁻ S⁻ strains. The swarm zone of DK10410 cells, the parental strain with full A motility, is shown in Fig. 6A. A⁻ S⁻ mutant cells did not move beyond the sharp edge of the colony at any time during the 60 min recording (Fig. 6D). Expansion of a non-swarming A⁻ S⁻ colony is due to cell growth and cell division, motility is not involved; consequently the A⁻ S⁻ strain expands at the lowest rate in Fig. 5. If there is no change in the distribution of pgl mutant cells, as the swarm expansion rate of the pglU mutant is 60% that of the A⁺ strain, one might expect that the cell and slime trail distribution of pglU at 90 min would resemble the A⁺ at 54 min (60% of 90 min). Figure 6B shows the 54 min frame from the 90 min movie of the A⁺ S⁻, and indeed, the 90 min frame of the pglU mutant (Fig. 6C) shows a pattern of slime trails, trail curvatures, trail length and cell distributions that qualitatively resemble Fig. 6A and B. In sum, the major effect of the loss of pgl function appears to be a lower probability of movement, resulting from longer pauses.

**Elasticotaxis**

Partial motility is also manifest in elasticotaxis. Elasticotaxis, first described by Stanier in 1942, measures the preference of myxobacterial cells to glide along lines of stress in an agar gel on which they are moving (Stanier, 1942). Elasticotaxis can be measured by the ratio of the rate of swarm spreading in the direction of stress in the agar to the rate perpendicular to the stress (Fontes and Kaiser, 1999). On unstressed agar, the rate of spreading

### Table 1. Movement probability, reversal frequency and gliding speed.

<table>
<thead>
<tr>
<th>Strain</th>
<th>Movement probability ± SD (n)</th>
<th>Reversal frequency (reversals h⁻¹ ± SD)</th>
<th>Gliding speed (µm min⁻¹± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>DK10410 (A⁺)</td>
<td>0.63 ± 0.007 (563)</td>
<td>6.00 ± 2.41</td>
<td>1.42 ± 0.32</td>
</tr>
<tr>
<td>DK13010 (pglB::TF)</td>
<td>0.34 ± 0.01 (406)</td>
<td>5.27 ± 2.81</td>
<td>1.13 ± 0.23</td>
</tr>
<tr>
<td>DK13013 (pglE::TF)</td>
<td>0.18 ± 0.01 (559)</td>
<td>6.01 ± 3.28</td>
<td>1.24 ± 0.26</td>
</tr>
<tr>
<td>DK13014 (pglF::TF)</td>
<td>0.21 ± 0.01 (431)</td>
<td>4.24 ± 2.82</td>
<td>N/A</td>
</tr>
<tr>
<td>DK13016 (pglH::TF)</td>
<td>0.18 ± 0.01 (386)</td>
<td>12.13 ± 3.74</td>
<td>N/A</td>
</tr>
<tr>
<td>DK13020 (mglB::TF)</td>
<td>0.36 ± 0.01 (525)</td>
<td>5.31 ± 2.50</td>
<td>N/A</td>
</tr>
</tbody>
</table>

Fig. 6. Distribution of *M. xanthus* cells in the spreading zone. Individual frames from the time-lapse movies whose preparation is described in Experimental procedures for photomicroscopy.

A. DK10410 (A⁺ S⁻) at 90 min.
B. DK10410 (A⁺ S⁻) at 54 min.
C. Partially A motile pglU::TF (DK13018) at 90 min.
D. DK11316 (A⁻ S⁻) at 60 min.
is the same in all radial directions and the swarm is circular, while on stressed agar the swarm is elliptical. The major axis of the resulting ellipse coincides with the direction of the stress. The elasticotactic responses of all the A motility mutants isolated in the DK10410 background and described above were measured as shown in Fig. 7. The eight non-swarming (A–S–) mutants tested showed no elasticotaxis; their colonies were circular on stressed agar (Fig. 7A). Evidently, the direction of cell growth shows little or no response to substrate stress. Elasticotaxis was evident in all 15 mutants that have partial A motility (Fig. 7B). Their swarms were elliptical, and in every case the major axis of the ellipse was coincident with the direction of stress (Fig. 7B). The elasticotaxis coefficients of the \( pgl \) mutants were less than an A\(^+\)S– strain but greater than an A–S– strain. Strikingly, the elasticotaxis coefficients of the partially motile mutant strains are directly proportional to their swarm rates. The numerical correspondence shown in Fig. 8 strongly suggests the swarm expansion rates and the elasticotaxis coefficients are different expressions of a common cause. Despite a superficial similarity between the assays for elasticotaxis and swarm expansion, their proportionality is not trivial; swarm expansion averages the outward movement of many cells and has the dimensions of a velocity; the elasticotaxis coefficient measures the asymmetry of movement direction and is dimensionless. Thus the two measures of motility are fundamentally different; their correlation is not trivial.

Functions of genes whose destruction gives partial motility

With data from the TIGR/Monsanto sequence of \( M. xanthus \) (GenBank CP000113), the coding sequences (CDS) disrupted by \( Himar \) were identified by cloning and sequencing a short segment adjacent to the transposon insertion in each mutant. All disrupted CDS could be rec-
ognized unambiguously in the published sequence. The complete CDS of each (uninterrupted) gene was then compared with the public protein databases, using Protein–Protein BLAST, to identify the most likely biochemical function of each A motility gene product by sequence similarity to known proteins. The highest-scoring hits to proteins that have catalytic functions corresponding to all the mutants isolated in this study are listed in Table 2. Some genes were hit more than once: agmK was hit six times (putatively it is a large gene, encoding 3822 amino acids), agmX, cglB, pglA, pglF and mglB were each hit twice. Deletion of mglB had previously been shown to markedly decrease the stability of MglA protein (Stephens and Kaiser, 1987; Stephens et al., 1989; Hartzell and Kaiser, 1991). Because swarming depends on MglA (Hodgkin and Kaiser, 1979a,b; Kaiser, 2007), mglB was, in fact, the first gene knockout mutant established to exhibit partial A motility. Isolation of new mglB mutants demonstrates the sensitivity of the mutant screen that was employed for reductions in A motility. The other genes listed in Table 2 will be examined below in light of their putative functions.

Discussion

New evidence is presented here that links A motility with the biosynthesis of, and gel formation by, a polysaccharide in M. xanthus. The new evidence complements high resolution electron micrographs which revealed several hundred thick walled rings, 80% of which were located at the cell poles (Wolgemuth et al., 2002). The rings are thought to be end views of secretory nozzles. Near the rings, narrow ribbons of an amorphous material, interpreted as a polysaccharide gel, were observed in the course of extrusion from cell ends (Wolgemuth et al., 2002). Evidently several narrow ribbons fuse laterally to form the single unipolar ribbon seen by light microscopy (Fig. 2). In an aqueous environment, ribbons of a polysaccharide gel would be expected to fuse. Both electron and light microscopy showed that the gel ribbons were to be found only at one cell pole at any particular moment; they were always absent from the opposite pole of the cell. Unipolarity of the extruded ribbons parallels unidirectional cell movement. As a consequence of gel extrusion, A'S' cells leave a phase bright trail of slime when they move (Reichenbach et al., 1965; Burchard, 1982; Wolgemuth et al., 2002 and Fig. 1). Slime extrusion at the back ends of many cells is shown in that figure. Fig. 1A and B show that every cell is on a slime trail. Figure 1C–H show how the trail grows as slime is deposited. It has been calculated that the swelling of a polysaccharide (slime) gel as it hydrates could produce a force sufficient for cells to glide at the speed observed (Wolgemuth et al., 2002; Wolgemuth, 2005).

The major new finding of this study is that all the new mutant strains identified because they had lost some or all of their A motility prove to have defects in their secretion of slime. Almost half (15/33) of the new mutants have completely lost A motility. They are non-motile because they are secreting slime and therefore push from both of their ends simultaneously. Because the force is comparable at both ends, they are unable to make progress in either direction. The majority (18/33) of the A motility mutants secrete slime only from one end of the cell. However, they are only partially A motile, they have a lower rate of swarm expansion and a proportionately lower coefficient of elasticotaxis than pglf+. They have defects in slime secretion that lowers their movement probability. The two sets of mutants show a perfect correlation between A motility and unipolar slime secretion. These data in themselves are a strong argument that slime secretion drives A motility. But, having visualized slime and seen its effects on cell movement, we need to understand its chemistry to see how force might be developed. The pgl mutants connect us to the enzymology of A motility: six pgl mutants carry null mutations in genes annotated as glycosyltransferases or other enzymes of polysaccharide biosynthesis.

Like pglB, almost all the partially motile mutants have wild-type reversal frequency, wild-type gliding speed, but a decreased movement probability compared with wild type (Table 1). According to Table 2, PglB protein is expected to have glycosyltransferase activity because it strongly resembles a glycosyltransferase of Leptospira interrogans (the value of the expectation, E = 3e-81). Enzymes of this type are needed to transfer an activated sugar (a UDP, ADP, GDP or GMP linked sugar) to a variety of substrates (Campbell et al., 1997). PglB, a glycosyltransferase of group 1, is a member of pfam00534. The second Pgl protein in Table 2, PglF is predicted to have two different glycosyltransferase domains: one in the N-terminal half that resembles RfaG, COG0438 and the glycosyltransferases of group 1. These enzymes are expected to transfer an activated sugar to an oligosaccharide acceptor. The more C-terminal glycosyltransferase domain of PglF belongs to group 2 that transfer sugar from UDP-glucose, UDP-N-acetyl-galactosamine, GDP-mannose or GDP-abequose to a range of acceptors like dolichol phosphate, teichoic acid and cellulose (Table 2). Growing polysaccharide chains in bacteria are generally anchored to a membrane by undecaprenylphosphate (Raetz and Whitfield, 2002), and the group 2 transferase of PglF may catalyse formation of Und-PP-oligosaccharides. The third Pgl protein in Table 2, PglD, has a nucleosidyl transferase domain (PF00483) and a mannose-1-phosphate isomerase domain (PF01050). This suggests that it may catalyse the synthesis of GDP-mannose for the synthesis of a mannose containing polysaccharide. PgL1 encodes a bifunctional ADP
Table 2. Inferred function of disrupted A motility genes.

<table>
<thead>
<tr>
<th>Coding sequence</th>
<th>Gene</th>
<th>Knockout phenotype</th>
<th>Domain function</th>
<th>Sequence</th>
<th>E-value&lt;sup&gt;b&lt;/sup&gt;</th>
<th>Conserved domains&lt;sup&gt;c&lt;/sup&gt;</th>
</tr>
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<tbody>
<tr>
<td>MXAN2921</td>
<td>pglB</td>
<td>Partially A motile</td>
<td>Glycosyltransferase similar to YP 000604 of <em>Leptospira interrogans</em></td>
<td>3.00E-81</td>
<td>Pfam00534</td>
<td></td>
</tr>
<tr>
<td>MXAN4616</td>
<td>pglF</td>
<td>Partially A motile</td>
<td>Glycosyltransferase 1 domain and glycosyltransferase 2 domain</td>
<td>CDD 40621, 1e-18</td>
<td>COG0438</td>
<td></td>
</tr>
<tr>
<td>MXAN6501</td>
<td>pglD</td>
<td>Partially A motile</td>
<td>GDP-mannose synthesis</td>
<td>CDD 40622, 6e-18</td>
<td>PF0483, PF01050</td>
<td></td>
</tr>
<tr>
<td>MXAN4710</td>
<td>pglN</td>
<td>Partially A motile</td>
<td>ADP-heptose synthase, bifunctional sugar kinase/adenylyltransferase RfaE-like</td>
<td>cd01172, 9e-51</td>
<td>COG2870</td>
<td></td>
</tr>
<tr>
<td>MXAN2919</td>
<td>pglJ</td>
<td>Partially A motile</td>
<td>Integral membrane protein similar to <em>S. coelicolor</em> gi:1098142 with local similarity to Wzy_c polymerase</td>
<td>IPR007016, 6.00E-07</td>
<td>PF04932</td>
<td></td>
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<tr>
<td>MXAN7252</td>
<td>pglA</td>
<td>Partially A motile</td>
<td>Exopolysaccharide synthesis, ExoD</td>
<td>3.00E-12</td>
<td>PF05055</td>
<td></td>
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<tr>
<td>MXAN4148</td>
<td>pglK</td>
<td>Partially A motile</td>
<td>Predicted transmembrane transcriptional regulator</td>
<td>5.00E-60</td>
<td>NCBI cdd00189</td>
<td></td>
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<tr>
<td>MXAN5319</td>
<td>pglC</td>
<td>Partially A motile</td>
<td>TPR repeat, CheY-like receiver domain, and a winged-helix DNA-binding domain</td>
<td>5.00E-60</td>
<td>None</td>
<td></td>
</tr>
<tr>
<td>MXAN4867</td>
<td>pglI</td>
<td>Partially A motile</td>
<td>Hypothetical abductin-like protein (M. xanthus&lt;sup&gt;3&lt;/sup&gt;)</td>
<td>5.00E-60</td>
<td>None</td>
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<tr>
<td>MXAN5382</td>
<td>aspT</td>
<td>Partially A motile</td>
<td>tRNA-Asp</td>
<td>g13582, 6e-14</td>
<td>COG1807</td>
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<tr>
<td>MXAN5585</td>
<td>pglE</td>
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<td>Glycosyltransferase of PMT family</td>
<td>5.00E-14</td>
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<tr>
<td>MXAN6607</td>
<td>agmT</td>
<td>Partially A motile</td>
<td>Predicted periplasmic solute-binding protein (<em>Trichodesmium erythraeum</em>)</td>
<td>5.00E-14</td>
<td>COG1559</td>
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<td>MXAN7160</td>
<td>pglM</td>
<td>Partially A motile</td>
<td>Alanine racemase</td>
<td>5.00E-14</td>
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<tr>
<td>MXAN11926</td>
<td>mglB</td>
<td>Partially A motile</td>
<td>guanine nucleotide exchange factor for MglA</td>
<td>5.00E-14</td>
<td>COG1559</td>
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<tr>
<td>MXAN2050</td>
<td>pglH</td>
<td>Partially A motile</td>
<td>TPR repeat, CheY-like receiver domain, and a winged-helix DNA-binding domain</td>
<td>5.00E-14</td>
<td>COG1559</td>
<td></td>
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<tr>
<td>MXAN1125</td>
<td>mglA</td>
<td>Non-motile (A'S)</td>
<td>SAR1-like small GTPase</td>
<td>5.00E-14</td>
<td>COG1100</td>
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<tr>
<td>MXAN2541</td>
<td>agnA</td>
<td>Non-motile (A'S)</td>
<td>Unknown</td>
<td>5.00E-14</td>
<td>COG1100</td>
<td></td>
</tr>
<tr>
<td>MXAN3008</td>
<td>agnU</td>
<td>Non-motile (A'S)</td>
<td>WD-repeat lipoprotein, acylaminoacyl-peptidease</td>
<td>5.00E-14</td>
<td>COG1100</td>
<td></td>
</tr>
<tr>
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<td>Outer membrane lipoprotein</td>
<td>cd01056</td>
<td>COG1100</td>
<td></td>
</tr>
<tr>
<td>MXAN4862</td>
<td>cglX</td>
<td>Non-motile (A'S)</td>
<td>Outer membrane lipoprotein</td>
<td>cd01056</td>
<td>COG1100</td>
<td></td>
</tr>
<tr>
<td>MXAN4863</td>
<td>cglK</td>
<td>Non-motile (A'S)</td>
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<td></td>
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<tr>
<td>MXAN6403</td>
<td>cglB</td>
<td>Non-motile (A'S)</td>
<td>ABC-type transporter permease protein (<em>Vibrio fischeri</em>)</td>
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<td>COG1100</td>
<td></td>
</tr>
<tr>
<td>MXAN7296</td>
<td>cglC</td>
<td>Non-motile (A'S)</td>
<td>Unknown</td>
<td>5.00E-14</td>
<td>COG1100</td>
<td></td>
</tr>
</tbody>
</table>

<sup>a</sup> Gene number in the complete *M. xanthus* genome, GenBank CP000113.

<sup>b</sup> ID of the best match in the NCBI protein database, E-value is expected matching by chance.

<sup>c</sup> COG, Clusters of Orthologous Groups of proteins, NCBI; NOG, non-supervised orthologous group, STRING database (http://string.embl.de/).
heptose synthetase, RfaE (Table 2), which suggests that a heptose could be one of the sugars in the slime polymer. PglJ is similar to an integral membrane glycosyltransferase, PF04932 (Table 2). Because PglA has four predicted transmembrane domains, it is likely to be a transmembrane protein that aligns over 200 residues with ExoD (Table 2). ExoD is involved in exopolysaccharide production in Sinorhizobium meliloti and is required for nodule invasion (Reed and Walker, 1991; Cheng and Walker, 1998).

The Sinorhizobium meliloti ExoD product is a repeating heteropolymer – a repeat unit polysaccharide, or RUP for short. Due to their medical importance, most enzymatic studies of RUP biosynthesis have been carried out on the O-antigen-specific chains of lipopolysaccharide (Raetz and Whitfield, 2002) and the capsules of Escherichia coli (Whittfield and Roberts, 1999). In both lipopolysaccharide and capsules, biosynthesis of a repeat unit of the polysaccharide begins on the cytoplasmic face of the inner membrane and continues during translocation to the periplasmic face of the inner membrane where repeat units are joined together before secretion (Raetz and Whitfield, 2002). Typically each biosynthetic step requires a distinct membrane-localized glycosyltransferase that hands its products off to another one, as reviewed by Raetz and Whitfield (2002). A handing off action is clearly implied by the structure of PglF, which has two different glycosyltransferase domains. Thus, the findings that the structure of PglF, which has two different glycosyltransferase domains, and that pglJ is the functional homologue of one of the three integral membrane polymerases for RUP synthesis (Table 2) strongly suggest that the pgl enzymes catalyse the biosynthesis of an RUP whose unipolar secretion propels the cell.

PglC contains five tetra-trico peptide repeats (TPRs, NCBI cd00189, Table 2). A homologue has been found in Anaeromyxobacter dehalogenans with an E-value of 5e-60. TPR structural motifs are present in a wide range of proteins that assemble membrane-localized multiprotein complexes (D’Andrea and Regan, 2003). Just as Tgl with six TPR motifs is an assembly factor for the PilQ secretion in the outer membrane of M. xanthus (Nudleman et al., 2006), PglC could be an assembly factor for a putative RUP biosynthetic complex of Pgl proteins in the inner membrane. The entire length of PglK has weak similarity to a transmembrane anti-sigma factor, COG5662 (Table 2) and it is just downstream of the ECF sigma factor rpoE1 (Ward et al., 1998). Ward et al. reported that knockout mutants of pglK (their orf5) were able to swarm and to aggregate in two genetic backgrounds. That diagnosis is not inconsistent with partial A motility.

The transposon insertions in the pgl mutants did not cluster in the carboxy end of the corresponding CDS, so they would not have been expected to leave partial enzyme activity. Is it possible that gene knockout engenders partial motility? Five pgl mutants tested in Table 1 pause twice as long between gliding movements as an A+ strain. We think it likely that the PglB, PglF, PglD, PglN, PglJ and PglA knockout mutants are also pausing in their synthesis of the propulsive polysaccharide for the lack of a glycosyltransferase. And because M. xanthus encodes many glycosyltransferases, needed for its multiple polysaccharides, it is possible that the lack of a particular transferase in an RUP biosynthetic pathway could be mitigated by a glycosyltransferase from another RUP pathway that happens to fit into the slime synthesizing enzyme complex. However, incorporating an alternate transferase into a multiprotein assembly complex would take time, thus pausing and interrupting polymerization. Moreover, a mitigated complex is likely to be less stable than the wild-type complex; it may have to be repeatedly reformed, introducing more pauses. Pausing clearly would be expected to decrease the swarm expansion rate. Because the incorporation of a mitigating sugar might change the sequence of sugars, it would be expected to affect interactions between the resulting polymer chains. If elasticotaxis is due to gelation of slime with the polysaccharides of agar, a reduction in the elasticotaxis coefficient would be expected as well.

pglH and mglB most likely have partial A motility for reasons other than RUP synthesis. PglH mutants increase the reversal frequency (Table 1); and pglH encodes a response regulator with a TPR repeat that is related to PleD; it is not a glycosyltransferase (Table 2). PglH is more likely to change the reversal frequency as a component of the reversal clock (Kaiser, 2007). The mutant phenotype suggests that the reversal frequency of wild type has evolved to maximize the rate of swarm expansion, so that an increase in the reversal frequency is likely to decrease the overall swarm rate. Although the mglB mutant has a normal reversal frequency (Table 1), MglB is a guanine nucleotide exchange protein (Table 2) that is expected to work with the MglA GTPase, to reverse cell polarity (Kaiser and Yu, 2005). MglB would catalyse release of GDP bound to the GTPase, promoting its replacement by GTP (Bourne et al., 1991). It has long been known that an mglB mutant swarms at a low rate and has little residual A motility (Stephen and Kaiser, 1987; Hartzell and Kaiser, 1991), and its movement probability is about half that of A+ (Table 1).

As mentioned above, approximately half the new mutants eliminated A motility; all of them secreted slime simultaneously from both ends of each cell. We suggested that these mutants are non-motile because they have lost the unipolarity of slime secretion that is essential for cell movement. The observation that bipolar mutants of M. xanthus are non-motile directly links a bipolar slime secretion morphology to the lack of movement. The pro-
totypic bipolar mutant is mglA, which was discovered by Hodgkin and Kaiser (1979a) who showed that it blocked both A motility and S motility. As mentioned, MglA is a GTPase switch that maintains the unipolarity of both the A and the S engines (Kaiser, 2007), and that ensures they have opposite polarity (Kaiser, 2003). MglA mutant cells had been observed to oscillate back and forth rapidly (Spormann and Kaiser, 1999). However, with each oscillation they move less than 1/5 of a cell length, and make no significant progress in either direction (Spormann and Kaiser, 1999). The rapid reversals of mglA mutants are not due to signals from the reversal generator because they lack an essential constituent of the generator. Instead, we suggest those reversals are a statistical consequence of active slime secretion from both ends. A calculation made by Charles Wolgemuth shows that the speed distribution of oscillating ∆mglAB cells can be deduced from the known speed distribution of ∆mglA cells (Spormann and Kaiser, 1999), assuming that the two ends of the same cell secrete independently (Kaiser, 2007).

All the proteins found in this study to be essential for unipolar slime secretion are listed at the bottom of Table 2. The second member of that list after MglA is CglB: the ∆cglB mutant also secretes slime from both ends (Fig. 3) and the cells oscillate 10-fold faster than A+ like ∆mglA (Spormann and Kaiser, 1999), which indicates a failure of the normal polarity switching mechanism. However, cglB mutants only affect A motility; their S motility is normal (Hodgkin and Kaiser, 1979a). Whereas MglA is cytoplasmic, CglB is a lipoprotein with a type II signal sequence (Rodriguez and Spormann, 1999) that resides in the outer membrane (Simunovic et al., 2003). In addition, CglB is transferred with high efficiency by stimulatory contact between cell ends (Hodgkin and Kaiser, 1977; White and Hartzell, 2000; Nudelman et al., 2005); this confirms its identification as a motility-related, outer membrane lipoprotein. Other non-motile mutants isolated in this study were also shown to have slime emerging from both poles by DIC microscopy (Fig. 3). Several had been identified as critical for A motility (White and Hartzell, 2000; Youderian et al., 2003). AgnA, AgnB and AgnC proteins are new. AgnB is homologous to a permease component of an ABC transporter from Vibrio fischeri which may be involved in localizing a lipoprotein (Table 2). Another bipolar mutant, AgnC, has a stretch of hydrophobic amino acids in its conceptually translated sequence suggesting that it is a transmembrane protein. Several old (AglU, AgmK, AgmX), and new (AgnB, AgnC) are either membrane proteins, or proteins involved in lipoprotein release. Their unknown roles could be related to that of CglB.

If M. xanthus is propelled by slime secretion, as proposed, then it is remarkable that none of the searches for A motility mutants have turned up mutants lacking slime. Despite extensive mutant hunts and frequent microscopic searches for locomotor organelles (Burchard et al., 1977; Pate and Chang, 1979; Burchard, 1981; 1984; Lunsdorf and Schairer, 2001), no alternative motility mechanism has survived initial experimental testing. There is, however, an alternative speculation based on Lapidus and Berg’s observation of energy-dependent movements of small particles adhering to Cytophaga U67 cells (Lapidus and Berg, 1982). After observing similar particle movement on the surface of Flexibacter johnsoniae, Dr M.J. McBride developed a mechanically complete model for gliding (McBride, 2000). The model proposes that mobile cell surface proteins in the F. johnsoniae outer membrane bind the substratum. Other proteins in the cytoplasmic membrane are proposed to harvest the proton motive force to propel yet other proteins that are embedded in the outer membrane along tracks that are attached to the peptidoglycan. Finally, processive attachment and detachment of the outer membrane proteins to the substratum are proposed to produce a walking motion. Might such a machine explain A motility? McBride’s mechanism predicts motorized binding proteins, perhaps AAA ATPases (Vale, 2000). On the one hand, none of the Mgl, Agm or Agn proteins have ATPase function, but a failure to find predicted proteins cannot be deemed a disproof. On the other hand, finding multiple glycosyltransferases, finding a strong preference to follow slime trails and finding sensitive elasticotactic orientation seem difficult to reconcile with the McBride mechanism of gliding. In our view, it makes more sense to hypothesize that slime secretion is vital, for reasons to be found. Perhaps the accumulation of intermediates in slime production is toxic to cells, or perhaps slime production facilitates survival after electroporation.

Experimental procedures

Bacterial strains

Strains and plasmids employed are listed in Table 3. In addition E. coli strains DH5αpir and TOP10 were used as hosts for cloning. They were grown in Luria–Bertani broth (LB) or on 1.5% agar LB plates supplemented with kanamycin (50 μg ml⁻¹) or ampicillin (100 μg ml⁻¹) as needed. M. xanthus DK1622 and its S motility mutant strains DK10410 and DK8615 were routinely grown in CTT medium [1% Casitone, 10 mM Tris-HCl (pH 8.0), 8 mM MgSO4, 10 mM KPO4 (pH 7.6)] at 32°C or on 1.5% agar CTT plates containing 40 μg ml⁻¹ of kanamycin or 12.5 μg ml⁻¹ of oxytetracycline when required. Plasmid pMiniHimar-lacZ was the donor of the mini-mariner element Himar1. pMiniHimar-lacZ DNA was prepared using the BIO-RAD Plasmid Miniprep Kit.

Transposon mutagenesis

M. xanthus DK10410 and DK8615 cells were grown to a density of 5 × 10⁸ cells ml⁻¹ in CTT broth. A 1.8 ml aliquot of the
Table 3. *M. xanthus* strains and plasmids employed.

<table>
<thead>
<tr>
<th><em>M. xanthus</em> strain</th>
<th>Genotype(^a)</th>
<th>Phenotype(^b)</th>
<th>Reference/construction</th>
</tr>
</thead>
<tbody>
<tr>
<td>DK1622</td>
<td>Wild type</td>
<td>A(^+)S(^+)</td>
<td>Kaiser (1979)</td>
</tr>
<tr>
<td>DK8615</td>
<td>pBlO</td>
<td>A(^+)S(^+)</td>
<td>Wall et al. (1999)</td>
</tr>
<tr>
<td>DK10410</td>
<td>pBlA</td>
<td>A(^+)S(^+)</td>
<td>Wu and Kaiser (1997)</td>
</tr>
<tr>
<td>DK11316</td>
<td>ΔcglB (pia::tet)</td>
<td>A(^+)S(^+)</td>
<td>Fontes and Kaiser (1999)</td>
</tr>
<tr>
<td>DK13001</td>
<td>agmK::TF</td>
<td>A(^+)S(^+)</td>
<td>pMiniHimar-lacZ (×) DK10410, select Kan(^R), screen for A(^-)</td>
</tr>
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<td>A(^+)S(^+)</td>
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</tr>
<tr>
<td>DK13003</td>
<td>mgA::TF</td>
<td>A(^+)S(^+)</td>
<td>pMiniHimar-lacZ (×) DK10410, select Kan(^R), screen for A(^-)</td>
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<tr>
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<td>A(^+)S(^+)</td>
<td>pMiniHimar-lacZ (×) DK10410, select Kan(^R), screen for A(^-)</td>
</tr>
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<td>A(^+)S(^+)</td>
<td>pMiniHimar-lacZ (×) DK10410, select Kan(^R), screen for A(^-)</td>
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<td>A(^+)S(^+)</td>
<td>pMiniHimar-lacZ (×) DK10410, select Kan(^R), screen for A(^-)</td>
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</tr>
<tr>
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<td>pMiniHimar-lacZ (×) DK8615, select Kan(^R), screen for A(^-)</td>
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<td>A(^+)S(^+)</td>
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<tr>
<td>DK13024</td>
<td>(\Delta)pgU</td>
<td>A(^+)S(^+)</td>
<td>pRY101 (×) DK10410, select Kan(^R); select Gal(^R)Kan(^R); screen by PCR analysis</td>
</tr>
</tbody>
</table>

Plasmid

| pMiniHimar-lacZ | Encodes the Himar1 transposase, aph (kan\(^R\)), promoterless lacZ and oriR6K\(^\lambda\) | Kan\(^R\) | Duan and Kaplan, University of Texas Medical School at Houston. |

Plasmid

| pBJ113 | pUC118 containing kan\(^R\) and galK, used for constructing gene replacements | Kan\(^R\)Gal\(^\beta\) | Julien et al. (2000) |

| pRY101 | \(\Delta\)pgU, kan\(^R\) and galK | Kan\(^R\)Gal\(^\beta\) | PCR-generated \(\Delta\)pgU fragments cloned into pBJ113 |

\(\Delta\) TF represents the transposed fragment from pMiniHimar-lacZ in the *M. xanthus* genes.

\(\Delta\) A, full A motility; \(\Delta\) A, no A motility; \(\Delta\) A\(^\text{half}\), partial A motility as described in Results; S\(^+\), full S motility; S\(^-\), no S motility.

Culture was harvested by centrifugation, washed once with 1.8 ml of sterile water, then washed twice with 1 ml of sterile water, and resuspended in 40 μl of sterile water. Freshly washed cells were mixed with dialysed pMiniHimar-lacZ DNA and subjected to electroporation conditions of 0.65 kV, 400 Ω and 25 μF in a 0.1 cm gap cuvette. The electroporated cells were immediately added to 2.5 ml of CTT broth and incubated with shaking at room temperature for 12–16 h for recovery. Recovered cells were then mixed with CTT soft agar (0.7% agar) and plated on CTT plates (1.5% agar) supplemented with kanamycin. After 5–6 days of incubation at 32°C, individual colonies were picked and transferred with sterile toothpicks on to fresh CTT agar plates. After 2–3 days of incubation at 32°C, plates were screened visually to identify small colonies and colonies with smooth edges.

Cloning of *M. xanthus* genomic DNA flanking pMiniHimar-lacZ insertions

Strains containing pMiniHimar-lacZ, described in Table 3, were grown overnight at 32°C in 10 ml of CTT broth. Bacteria were harvested by centrifugation and resuspended in 3 ml of sucrose-Tris-EDTA (25% sucrose, 10 mM Tris (pH 8.0), 1 mM EDTA (pH 8.0)). Cells were lysed by the addition of 0.6 ml of Lytic Mix (5% sodium dodecyl sulphate, 125 mM EDTA (pH 8.0), 0.5 M Tris (pH 9.4)) and incubated at 65°C for 60 min. Proteinase K was added to 100 μg ml\(^{-1}\) final concentration and the mixture was incubated at 37°C for 2 h. *Myxococcus* genomic DNA was prepared using standard phenol-chloroform extraction and ethanol precipitation. The precipitate was suspended in 60 μl of TE (10 mM Tris-HCl (pH 8.0), 0.1 mM EDTA (pH 8.0)) overnight at 4°C and stored at −20°C. One to two microlitres of the DNA suspension was digested with 2 μl SacII restriction endonuclease for 3–12 h in a total volume of 30 μl, followed by the addition of an additional 1 μl SacII and incubation for 2 h more, and finally heated to 65°C for 15 min to inactivate the endonuclease. The digested genomic DNA was purified by phenol-chloroform extraction, ethanol precipitation, and was resuspended in 20 μl of sterile water. Digested DNA was ligated with T4 DNA ligase in a total volume of 150 μl overnight at 16°C, and heated at 65°C for 20 min to inactivate the ligase. The DNA was cleaned by
mixing it with 500 μl of n-butanol, pelleting the DNA, and resuspending it in 10 μl water. Finally 2 μl DNA was electrotransformed into E. coli strain DH5α λpir, and the electroporated cells were plated on LB agar plates supplemented with kanamycin for selection. Plasmid DNA containing the origin of replication from pMiniHimar-lacZ and the flanking myxococcal genomic DNA was prepared using the BIO-RAD Plasmid Miniprep Kit and manufacturer’s instructions. The sequence of DNA adjacent to the transposon was obtained using a primer immediately upstream of the right inverted repeat of pMiniHimar-lacZ with the sequence 5′-GAA CTA TGT TGA ATA ATA AAA ACG A-3′.

Construction of an in-frame deletion mutant of pglJ (MXAN2919)

To create an in-frame deletion, two PCR fragments of 750–800 bp that correspond to the upstream and downstream regions of the target deletion were generated. Primers 5′-CGG AAT TCG GCG CGT GGA CGA AAT CA-3′ and 5′-CGG GAT CCG AGC GCG GAC ACG CTA TC-3′ with restriction sites EcoRI and BamH1, respectively were used to generate the upstream PCR product. Primers 5′-CGG GAT CCG GCC ACC TGG AGA TGG AA-3′ and 5′-AAC TGC AGC

Analysis of amino acid sequences

Complete CDS were compared with the public protein databases with Protein–Protein BLAST (BLASTP) to seek the possible biochemical function of the protein products by sequence similarity to known proteins (http://www.ncbi.nlm.nih.gov/BLAST/). The Conserved Domain Database (CDD) (http://www.ncbi.nlm.nih.gov/Structure/cdd/cdd.shtml) was also searched.

Visualizing slime

A small amount of exponentially growing M. xanthus, suspended in 50 μl of 1/2 CTT medium [0.5% Casitone, 10 mM Tris-HCl (pH 8.0), 8 mM MgSO4, 10 mM KPO4 (pH 7.6)] was added to a tissue culture well containing 1 ml of 1/2 CTT medium of a 24 well flat bottom plate. After overnight incubation at room temperature in the dark, the tip of a pipette was used to gently scrape the bottom of the culture well and to resuspend the cells that had settled and had moved over the bottom surface of the well. A 3–5 μl droplet of this culture was transferred to a microscope slide and a coverslip was placed on top. Cells and any slime they produced were observed through a 100× DIC objective in a Nikon Eclipse E800 microscope by DIC. Images were collected with a 5 M Hz Micromax 5600 cooled CCD camera controlled by Metamorph (Universal Imaging).

Swarm expansion

Swarm expansion was measured using a modified published procedure (Kaiser and Crosby, 1983). M. xanthus was grown to exponential phase and concentrated to a calculated density of 2.5 × 10⁹ in CTT medium. An aliquot of 5 μl of each concentrated culture was spotted on plates, prepared the day before use, containing 7 ml of CTT with 1.5% agar in 50 × 5 mm plastic Petri dishes with tightly fitting lids. After the liquid droplet had soaked into the agar depositing the cells, the plates were closed tightly to prevent further drying and they were incubated at 32°C. Three individual plates were made for each strain. The rate of swarm expansion was quantified by measuring the average width of the zone of spreading, which is the distance between the centre of the swarm and the outermost extents of the swarm. At each time point, two width measurements were made on each of three duplicate swarms. Thus, for each strain, an arithmetic mean of six radius measurements was recorded for that time. The average radius of each strain was plotted against time, and the slope of the resulting line was determined to give the rate of swarm expansion.

Elasticotaxis

The assay for elasticotaxis was performed as previously described (Fontes and Kaiser, 1999) with the following modifications. Square plastic Petri dishes 100 × 15 mm (Nunc) containing 35 ml of CTT with 1.5% agar were used. A sterile 7-cm-long piece of plastic tubing (3/32 ID, 5/32 OD) was inserted between the dish and the solidified agar to compress the agar by 1 part in 20. Compression squeezed a small amount of liquid from the agar, which was allowed to evaporate at 32°C. The compressed agar was inoculated with 5 μl aliquots of the concentrated (2.5 × 10⁷) M. xanthus cultures (prepared as described in Swarm expansion section), and incubated at 32°C. At each time point, the diameters of the swarms parallel and perpendicular to the compressing tube were measured, and the ratio between them was calculated to give the coefficient of elasticotaxis (E).

Time-lapse photomicroscopy

M. xanthus strains were grown to exponential phase and diluted to a calculated density of 2.5 × 10⁸ cells ml⁻¹ in CTT broth. An aliquot of 5 μl of each diluted culture was spotted on plates containing 7 ml of CTT with 1.5% agarose in
and divided by the number of frames spanned by the move-
for the back of the same cell. The two values were averaged,
obtained for both the path of the front of the cell and the path
measurement was repeated until a consistent value was
strains were grown to a calculated density of 1
ability, the same procedure was followed except that the
images were saved as QuickTime movies which were exam-
ized. This work was supported by a Postdoctoral Fel-
starting with the first frame of each QuickTime movie, every
number of reversals per hour. For movement probability,
individual cells were followed for each movie to determine the
movement of single cells.

To measure the reversal frequency and movement prob-
ability, the same procedure was followed except that the
strains were grown to a calculated density of 1 x 10^7 cells ml^{-1}
in CTT broth, and an aliquot of 10 µl of each diluted culture
was spotted on the plates. After the spot of cells had dried for
approximately 30 min, the plates were immediately examined
under the microscope. Individual cell movements were
tracked over the 120 frames of the 60 min movies manually.
Two movies were made for each strain, and approximately 30
individual cells were followed for each movie to determine the
number of reversals per hour. For movement probability,
starting with the first frame of each QuickTime movie, every
cell (roughly 350 cells) was tracked through 15 frames of the
movie. Each cell was scored as to whether it moved detect-
ably or not during those 15 frames to obtain the probability of
movement during a sample period that represented 12% of
the movie.

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