INTRODUCTION

Directed motility in response to environmental chemical cues, a process called chemotaxis, is an important trait for many bacterial pathogens. *Helicobacter pylori*, a Gram-negative gastric pathogen, is proposed to situate itself within the gastric mucosa in response to specific chemical cues such as pH (Schreiber et al., 2004). Dysregulation of chemotaxis compromises the ability of *H. pylori* to colonize the murine and gerbil stomach, and results in its abnormal distribution and modulation of inflammation in this tissue (Foynes et al., 2000; McGee et al., 2005; Terry et al., 2005; Williams et al., 2007).

The chemotactic response, which has been studied most extensively in *Escherichia coli*, is mediated by chemoreceptors termed methyl-accepting chemotaxis proteins (MCPs) (Armitage, 1999). The *H. pylori* chemosensory machinery is similar but not identical to that of *E. coli*. The core signal-transduction apparatus (CheW, CheA and CheY) is present in both bacteria, but *H. pylori* also possesses three CheV proteins composed of both CheW and response-regulator motifs (Lowenthal et al., 2009; Pittman et al., 2001). The *H. pylori* genome encodes four chemoreceptors, TlpA, TlpB, TlpC and TlpD (originally called HylB or HlyB), although strain G27 does not express TlpC due to a frameshift mutation in the gene. Unlike peritrichous *E. coli* cells, which exhibit forward swimming and tumbling behaviour, we and others have observed that *H. pylori* cells swim forward, reverse and stop (Lowenthal et al., 2009; Schweinitzer et al., 2008). Once the cells stop, they usually resume movement in a different direction, thus allowing them to explore new spaces, similar to the tumbling behaviour of *E. coli*.

Only a handful of chemotactic signals have been demonstrated for *H. pylori* chemoreceptors. In strain 26695 the chemoreceptor TlpA has been shown to respond to

*Helicobacter pylori* perceives the quorum-sensing molecule AI-2 as a chemorepellent via the chemoreceptor TlpB

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*Helicobacter pylori* moves in response to environmental chemical cues using a chemotaxis two-component signal-transduction system. Autoinducer-2 (AI-2) is a quorum-sensing signal produced by the LuxS protein that accumulates in the bacterial environment in a density-dependent manner. We showed previously that a *H. pylori* luxS mutant was defective in motility on soft agar plates. Here we report that deletion of the luxS gene resulted in swimming behaviour with a reduced frequency of stops as compared to the wild-type strain. Stopping frequency was restored to wild-type levels by genetic complementation of the luxS mutation or by addition of synthetic 4,5-dihydroxy-2,3-pentanedione (DPD), which cyclizes to form AI-2. Synthetic DPD also increased the frequency of stops in wild-type *H. pylori*, similar to the behaviour induced by the known chemorepellent HCl. We found that whereas mutants lacking the chemoreceptor genes tlpA, tlpC or tlpD responded to an exogenous source of synthetic DPD, the chemoreceptor mutant tlpB was non-responsive to a gradient or uniform distribution of the chemical. Furthermore, a double mutant lacking both tlpB and luxS exhibited chemotactic behaviour similar to the tlpB single mutant, whereas a double mutant lacking both tlpB and the chemotransduction gene cheA behaved like a nonchemotactic cheA single mutant, supporting the model that tlpB functions in a signalling pathway downstream of luxS and upstream of cheA. We conclude that *H. pylori* perceives LuxS-produced AI-2 as a chemorepellent via the chemoreceptor TlpB.

Abbreviations: AI-2, autoinducer-2; DPD, 4,5-dihydroxy-2,3-pentanedione; iPCR, inverse PCR; SAM, S-adenosylmethionine.
arginine and bicarbonate (Cerda et al., 2003), and in strain SS1 the chemoreceptor TlpB has been demonstrated to respond to acid (Croxen et al., 2006). The intracellular chemoreceptor TlpD has been described as an energy sensor, although the molecular nature of the cues it perceives is not known (Schweinitzer et al., 2008). There are no known ligands for TlpC.

One environmental condition that *H. pylori* experiences is the endogenously produced quorum-sensing molecule autoinducer-2 (AI-2) (Forsyth & Cover, 2000; Joyce et al., 2000; Rader et al., 2007). Autoinducers (AIs) are bacterially produced extracellular signalling molecules that trigger quorum sensing, a form of bacterial cell–cell communication (Ng & Bassler, 2009). AI concentration increases as a function of bacterial cell density, and at critical concentration thresholds it initiates coordinated gene regulation. Several types of quorum-sensing systems have been characterized. Typically, Gram-positive and Gram-negative bacteria use oligopeptides and acylated homoserine lactones as AIs, respectively. Many Gram-positive and Gram-negative bacteria possess a quorum-sensing system that produces the furanone signal AI-2, which functions as a signal for interspecific communication. AI-2 is produced as a metabolic byproduct of the reaction carried out by LuxS, which cleaves 3-ribosylhomocysteine, producing homocysteine and 4,5-dihydroxy-2,3-pentanedione (DPD). DPD undergoes rapid dehydration and cyclization, existing in equilibrium as several molecules collectively termed AI-2 (Schauer et al., 2001). In many bacteria, LuxS is a key enzyme in the production of the activated methyl donor, S-adenosylmethionine (SAM); however in *H. pylori*, the homocysteine produced by LuxS is metabolized to produce cysteine (Doherty et al., 2010).

Previous reports have shown that several *H. pylori* strains produce AI-2 in a luxS-dependent fashion, as detected through a *Vibrio harveyi* luminescence assay (Forsyth & Cover, 2000; Joyce et al., 2000; Lee et al., 2006; Rader et al., 2007; Shen et al., 2010). Mutation of the luxS gene has been associated with reduction of the flagellin flaA transcript and protein, enhanced biofilm formation, decreased motility on soft agar plates, and colonization defects in mouse and gerbil infection models (Cole et al., 2004; Lee et al., 2006; Loh et al., 2004; Osaki et al., 2006; Rader et al., 2007). We have previously reported that LuxS-produced AI-2 functions as a signal molecule that modulates transcript levels of the flagellar regulator FlhA, thereby influencing global flagellar regulation (Rader et al., 2007). Although it is clear that AI-2 signalling affects flagellar gene expression, we found that the flagella in luxS mutants are morphologically normal and thus the altered flagellar gene regulation in this strain may not explain fully its motility defect on soft agar plates. Because colonial expansion in these plates requires both motility and chemotaxis, we sought to examine how AI-2 affects the latter process. In this study we identify a novel function for AI-2 as a chemorepellent that is sensed by the chemoreceptor TlpB.

### METHODS

**Bacterial strains and culture conditions.** *H. pylori* strain G27 and its isogenic mutants were used in this study and are listed in Table 1. All *H. pylori* strains were maintained on blood agar plates consisting of Columbia agar (Difco) and 5% defibrinated horse blood (Hemostat) (CHBA), or CHBA plates supplemented with 0.02 mg β-cyclodextrin ml⁻¹ (Sigma), 8 mg amphotericin B ml⁻¹ (Sigma) and 20 μg vancomycin ml⁻¹ (Sigma), and incubated at 37 °C in 10% CO₂. Selective plates were supplemented with 10–15 μg kanamycin ml⁻¹ (Fisher), 18 μg metronidazole ml⁻¹, 10 μg chloramphenicol ml⁻¹ or 80 mM sucrose. *H. pylori* liquid medium (BB10) consisted of filtered sterilized Brucella broth (Difco) supplemented with 10% fetal bovine serum (Gibco) and 20 μg vancomycin ml⁻¹ (Sigma). Liquid cultures were grown in 50 ml conical tubes (BD Falcon) with loosened lids shaking at 37 °C in anaerobic jars (Oxoid) with CampyGen microaerobic sachets (Oxoid) or in glass test tubes (20 x 148 mm) shaking in a 37 °C/8% CO₂ incubator.

### Table 1. Strains used in this study

<table>
<thead>
<tr>
<th><em>H. pylori</em> strain</th>
<th>Description</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>G27 (KG01)</td>
<td>Wild-type</td>
<td>Kovac et al. (1993)</td>
</tr>
<tr>
<td>luxS (BR08)</td>
<td>G27, luxS::kan sacB</td>
<td>Rader et al. (2007)</td>
</tr>
<tr>
<td>ΔluxS (BR09)</td>
<td>G27, deletion of luxS</td>
<td>Rader et al. (2007)</td>
</tr>
<tr>
<td>luxS* (BR10)</td>
<td>G27, deletion of luxS and rdx::loxS</td>
<td>Rader et al. (2007)</td>
</tr>
<tr>
<td>mG27 (KO625)</td>
<td>Mouse-passaged strain G27</td>
<td>Castillo et al. (2008)</td>
</tr>
<tr>
<td>ΔtlpA (KO1002)</td>
<td>mG27, deletion of tlpA</td>
<td>This study</td>
</tr>
<tr>
<td>tlpB (KO1003)</td>
<td>mG27, tlpB::kan sacB</td>
<td>This study</td>
</tr>
<tr>
<td>ΔtlpB (KO1004)</td>
<td>mG27, deletion of tlpB</td>
<td>This study</td>
</tr>
<tr>
<td>tlpB* (BR31)</td>
<td>mG27, deletion of tlpB and rdx::tlpB,Δss1-aphA-3</td>
<td>This study</td>
</tr>
<tr>
<td>luxS tlpB (CW30)</td>
<td>mG27, tlpB::cat and luxS::kan sacB</td>
<td>Williams et al. (2007)</td>
</tr>
<tr>
<td>tlpC (KO1005)</td>
<td>mG27, tlpC::aphA-3</td>
<td>This study</td>
</tr>
<tr>
<td>tlpD (KO1006)</td>
<td>mG27, tlpD::cat (made using pL30A2cat12)</td>
<td>Williams et al. (2007)</td>
</tr>
<tr>
<td>cheA (KO629)</td>
<td>mG27, cheA::cat reverse (made using pKT22)</td>
<td>Terry et al. (2005)</td>
</tr>
<tr>
<td>tlpB cheA (CW31)</td>
<td>G27, tlpB::kan sacB and cheA::cat reverse</td>
<td>This study</td>
</tr>
</tbody>
</table>
Construction of \textit{H. pylori} mutants. All transformations were performed using natural transformation. PCR for cloning purposes was done with either \textit{Pi}ta Turbo (Stratagene) or Phusion (NEB), and all DNA manipulation enzymes were from New England Biolabs. The \textit{G27} lux\textit{S},lux\textit{S} and \textit{lux}+\textit{S} isogenic strains (strains BR08, BR09 and BR10, respectively) were constructed as previously described (Rader et al., 2007). Throughout, we use ‘\textit{A}’ to designate deletion mutations, in which most of a gene’s coding sequence is removed by allelic exchange, and the superscript ‘\textit{a}’ to designate complemented strains in which a wild-type copy of a mutated gene is supplied at another genomic locus from the mutated gene. The \textit{lux}+\textit{S} strain contains a wild-type copy of the \textit{luxS} gene and its promoter introduced in the \textit{rdxA} locus of the \textit{lux}+\textit{S} mutant chromosome. For most chemotaxis mutants, a mouse-selected variant of \textit{G27}, \textit{mG27}, was used as the parent strain (Castillo et al., 2008).

The \textit{cheA} mutant (strain K0629) was created by transforming \textit{H. pylori} \textit{mG27} with pKT22 (Terry et al., 2005) and selecting for resistance on chloramphenicol CHBA.

\textbf{\Delta \textit{tlpB} (strain KO1004) was made in two steps. First, \textit{mG27} was transformed to kanamycin resistance (Km\textsuperscript{b}) by the plasmid pTC-B111, which bears a \textit{tlpB::kan-sacB} allele. The correct nature of this mutation was verified by PCR using primers that flank the insertion site. pTC-B111 was made starting with pTCB101, which has \textit{tlpB} from strain S11 cloned into pBluescript (McGee et al., 2005). pTCB101 was then subjected to IPCR using primers tlpB-30 and tlpB-40 (sequence in McGee et al., 2005). The \textit{kan-sacB} construct was obtained from pKSFII (Copass et al., 1997) by digesting with \textit{SmaI} and \textit{XhoI}. The \textit{kan-sacB} and \textit{tlpB} flanking pieces were ligated using T4 ligase. The correct nature of the plasmid was verified by digestion. In this construct, \textit{tlpB} is transcribed in the same direction as \textit{sacB} (opposite to \textit{aphA-3}). This strain is called KO1003. KO1003 was then transformed with a plasmid bearing a large deletion of \textit{tlpB}, pTC-B113, followed by selection for sucrose resistance (Suc\textsuperscript{c}). pTC-B113 was created using IPCR with primers tlpB-30 and tlpB-40, followed by intramolecular ligation of the plasmid. The deletion removes all of \textit{tlpB} except the first 27 and the last 17 amino acids, although the deletion is +1 out of frame, resulting in a 3’ end that encodes a different sequence with three additional amino acids. The \textit{tlpB} region from Suc\textsuperscript{c}, kanamycin-sensitive (Km\textsuperscript{c}) colonies was amplified using PCR with primers that flank \textit{tlpB}. This PCR product was sequenced to verify that \textit{tlpB} was deleted.

To create the \textit{tlpB\textsuperscript{c}} complemented strain (strain BR31), \textit{mG27} \textit{\DeltatlpB} mutant bacteria were naturally transformed with a construct encoding the \textit{rdxA} locus with an insertion composed of the full-length \textit{tlpB} gene, expressed from the \textit{cheY} promoter, and cloned upstream of the non-polar \textit{aphA-3} cassette. Disruption of the \textit{rdxA} locus confers metronidazole resistance, and the non-polar \textit{aphA-3} cassette contains two internal ribosome-binding sites flanking the kanamycin-resistance gene conferring that resistance. Transformants were first screened on kanamycin plates, further screened on metronidazole-supplemented plates, and verified by PCR amplification and sequencing of the genomic locus.

The \textit{tlpA} mutant (strain KO1002) was made in two steps by first transforming \textit{mG27} to Km\textsuperscript{b} with pTAl2, which bears a \textit{tlpA::kan-sacB} allele, to create strain KO1001, followed by transformation with a plasmid that bears a \textit{tlpA} deletion (pTAl4). pTAl2 was made from pTAl0, which has \textit{tlpA} from strain S11 cloned, with ~500 bp flanking sequence, by PCR amplification with primers TlpA6 (5’-ATTTAAGGGGGCCCG-3’) and TlpA7 (5’-TTTTCCTCTCAGGAAAGC-3’). This product was phosphorylated with T4 polynucleotide kinase, and then cloned into EcoRV-digested pBluescript KS+ (Stratagene) to create pTAl0. To create pTAl2, PCR was carried out with pTAl0 template plus primers TlpA12 (5’-CAGCACTATAGTTATACCG-3’) and TlpA13 (5’-CAAGAAATAGCAGACG-3’) to create an in-frame deletion that leaves a region coding for 14 amino acids at the 5’ end, and 19 amino acids at the 3’ end of \textit{tlpA}. This PCR product was then ligated to a \textit{kan-sacB} fragment that came from pKS2. pKS2 derived from pKS1, which was made via a three-piece ligation of the following pieces: (1) the \textit{aphA3} gene (also \textit{kan}) isolated from pBS-kan (Terry et al., 2005) by cutting with \textit{SmaI} + \textit{XhoI}, followed by dephosphorylation; (2) the \textit{sacB} gene from pKS1I (Copass et al., 1997) generated by cutting with BamHI, blunting with T4 polymerase followed by cutting with \textit{PstI}; (3) the vector backbone from pBluescript KS+ cut with \textit{PstI} and \textit{XhoI}, followed by phosphatase and gel purification of the 3.0 kb piece. These three pieces were ligated together to generate pKS1. pKS1 has the \textit{aphA3} promoter driving transcription of both \textit{aphA3} and the downstream \textit{sacB} gene. To create pKS2, pKS1 was digested with \textit{PvuI} to remove the \textit{bla} gene followed by self-ligation. For cloning of the \textit{aphA3-sacB} fragment from pKS2, PCR amplification using primers reverse (universal) and \textit{sacBend2} (5’-CTTTCGGTCTTTTTTTTTTGTTAAC-3’) was carried out, to amplify \textit{aphA3-sacB} without the \textit{sacB} transcriptional terminator, and the resulting fragment was ligated with the IPCR fragment of pTAl0 to create pTAl2. In pTAl2, \textit{tlpA}, \textit{aphA3} and \textit{sacB} are all in the same transcriptional orientation. To create pTAl4, the same pTAl0 IPCR fragment was self-ligated. \textit{H. pylori} KO1001 was then transformed with pTAl4, followed by selection for sucrose resistance and screening for kanamycin sensitivity. The \textit{tlpA} allele was PCR amplified from Suc\textsuperscript{c} Km\textsuperscript{b} colonies using primers tlpA10 (5’-CTTAAAGGTGGTTAGTATCGG-3’) and tlpA11 (5’-GCTGAATCTGAAACACTCTTTAACAGT-3’), and sequenced to verify the correct deletion. This \textit{tlpA} strain is also called \textit{mG27 tlpA} or KO1002.

The \textit{mG27 \textit{tlpC} mutant} is \textit{tlpC::aphA3} (strain KO1005). This mutant was constructed by transforming \textit{mG27} with plasmid pKO150. This plasmid was made using PCR of a plasmid containing \textit{tlpC} from S11 called pTC100 (Andermann et al., 2002) with primers tlpC31 (5’-TCAATCGAGATACATTTAAC-3’) and tlpC40 (5’-CTGTGCACAAAGATGAGTGAAGG-3’). The product, 3.5 kb, was ligated with the \textit{aphA3} gene prepared by PCR from pBS-kan with primers Kanup (5’-CACCACTGATCGATACAAATCCG-3’) and Kandown (5’-GGCCAGCTTTTAGCACATCTAAAAG-3’), and sequenced to verify the correct deletion. This \textit{tlpC} strain is also called \textit{mG27 tlpC} or KO1002.

Autoinducer-2 is a chemorepellent for \textit{H. pylori}.

\textbf{http://mic.sgmjournals.org}
Bioluminescence assay and synthetic DPD. Synthetic DPD, a kind gift from Martin Semmelhack and Bonnie Bassler (Princeton University), was prepared as described by Semmelhack et al. (2005). This DPD was synthesized with a protecting group that we removed by treatment with 15 mM H2SO4, followed by neutralization to pH 6.9 with potassium phosphate. As a mock DPD treatment, we neutralized 15 mM H2SO4 to pH 6.9 with potassium phosphate. Alternatively, we used DPD from OMM Scientific that did not contain a protecting group; BB10 medium was used to dilute this DPD solution was placed on the open side of the coverslip. After 5–10 min incubation, the slides were examined and still images were taken using Scion Image software at ×10 through a ×40 phase filter, conditions that mimic dark-field microscopy.

Western blotting. Total cell proteins were prepared from H. pylori cultured on CHBA plates for 2 days by resuspending and lysing the cells in 2× Laemmli sample buffer. Samples were separated on a 10% SDS-PAGE gel, transferred to Immunoblot PVDF membranes (Bio-Rad) and incubated with 1:5000 dilution of anti-GST_TlpA22 (Williams et al., 2007). This rabbit polyclonal antibody recognizes the conserved CheW-interacting methyl-accepting domain of all H. pylori chemoreceptors. For visualization, the blots were incubated with the secondary antibody goat anti-rabbit-HRP (Santa Cruz Biotech) at a dilution of 1:2000, followed by incubation with luminol, p-coumaric acid and hydrogen peroxide. Luminescent blots were visualized by exposure to Biomax Light film (Kodak).

Statistical analysis. The frequency of stops was analysed statistically using either one-way ANOVA (www.physics.csbsju.edu/cgi-bin/stats/anova_pnp) when strains were tested or two-way ANOVA (http://faculty.vassar.edu/lowry/anova2u.html) when both strains and chemical treatment were tested. Post-hoc multiple comparisons to identify significance within individual hypotheses were performed using t-tests, and the P-values were adjusted using Bonferroni correction to correct for an increase in type I error. P-values of <0.001 were considered significant and used for Bonferroni correction.

RESULTS

luxS is required for normal swimming behaviour of H. pylori

When examining the swimming behaviour of wild-type and an isogenic luxS deletion mutant (referred to as ΔluxS) of H. pylori strain G27 (Rader et al., 2007), we observed that the ΔluxS mutant displayed a propensity for swimming in a straight line (running), as opposed to the wild-type, which exhibited more typical run–stop–run behaviour. We quantified this behaviour in bacterial cultures using video microscopy to observe swimming behaviour. We found that the ΔluxS mutant exhibited significantly fewer stops per second than the wild-type strain (Fig. 1). Complementation of the luxS mutation with a wild-type copy of the luxS gene at the rdxA locus [referred to as luxS* (Rader et al., 2007)] significantly increased the number of stops per second as compared with the ΔluxS strain (Fig. 1). It should be noted that over the course of these experiments, we observed some variation between trials in the average number of stops per second exhibited by the strains, but the responses to chemicals and differences between genotypes were reproducible across experiments. We attribute these differences to factors that...
were difficult to control across all experiments, such as differences in the lots of serum used in the media.

**The LuxS product, AI-2, is a chemorepellent for *H. pylori***

To test whether the decrease in the number of stops in the ΔluxS mutant relative to the G27 wild-type and luxS* complemented isogenic strains was due to the loss of the AI-2 signal, we treated each strain with 0.1 mM synthetic DPD, corresponding to the approximate concentration of AI-2 in the cell-free supernatant of an early stationary-phase culture of wild-type *H. pylori* (Rader et al., 2007). We also tested each isogenic strain with sterile water, the DPD diluent, as a negative control. Bacterial cultures were incubated with video microscopy, and the number of stops that individual bacteria performed during 5 s was recorded. Each strain was treated with water, mock DPD, 0.1 M HCl and 0.1 mM synthetic DPD. The number of individual bacterial cells scored (n) for each condition is indicated below the column; bars indicate SE. * indicates each strain that is statistically significant different from all other strains within a treatment, + indicates chemorepellent treatments (HCl, synthetic DPD) that are significantly different from controls (water, mock DPD) within a strain (P<0.001 with Bonferroni correction).

![Fig. 1. LuxS is required for normal swimming behaviour of *H. pylori*. Swimming behaviours of G27 wild-type, ΔluxS mutant and luxS* complemented isogenic strains were observed by video microscopy, and the number of stops that individual bacteria performed during 5 s was recorded. Each strain was treated with water, mock DPD, 0.1 M HCl and 0.1 mM synthetic DPD. The number of individual bacterial cells scored (n) for each condition is indicated below the column; bars indicate SE. * indicates each strain that is statistically significant different from all other strains within a treatment, + indicates chemorepellent treatments (HCl, synthetic DPD) that are significantly different from controls (water, mock DPD) within a strain (P<0.001 with Bonferroni correction).](http://mic.sgmjournals.org)

and synthetic DPD, we incubated wild-type, ΔluxS and luxS* G27 with 0.1 M HCl and observed their swimming behaviour. The pH of the synthetic DPD solution was 6.9, in contrast to the pH of 1.0 of the 0.1 M HCl solutions. All three isogenic strains exhibited an increase in the number of stops when exposed to 0.1 M HCl, and this increase did not differ significantly from that in response to synthetic DPD (Fig. 1). To ensure that this similarity was not due to some component of the solution used to rehydrate and activate the synthetic DPD, we also performed the experiment with a solution identical to the synthetic DPD solution, to which the synthetic DPD was not added (mock DPD). This treatment did not alter the swimming behaviour of the strains relative to their behaviour in response to water (Fig. 1). All subsequent experiments employed mock DPD as the negative control. These experiments thus suggest that DPD functions as a chemorepellent for *H. pylori*, because it increases the frequency of bacterial stopping.

To further explore whether *H. pylori* uses chemotaxis to respond to synthetic DPD, we employed a previously described wet-mount assay to monitor spatial chemotactic responses (Croxen et al., 2006). Wild-type G27 bacterial suspensions in BB10 medium were prepared as described earlier, seeded onto a glass microscope slide, and covered with a coverslip that was then sealed on three sides. Synthetic DPD (0.1 mM), HCl (0.1 M), mock DPD or water was then spotted onto the part of the coverslip that remained unsealed (Fig. 2a). Movement of the bacterial population was monitored before, immediately after and 5 min after addition of the chemical stimulus. Prior to the addition of treatment the bacteria were motile and uniformly distributed below the coverslip. Immediately after addition of treatment, the bacteria were flushed away from the liquid source; however after 5 min the hydrodynamic flow had ceased, and in the case of mock DPD or water, the bacteria returned to a uniform distribution. As previously reported (Croxen et al., 2006), in response to 0.1 M HCl, the percentage of motile bacteria immediately increased, with the majority of the bacteria exhibiting an increase in the frequency of stops, resulting in net movement away from the source of the acid. This increased movement proximal to the acid source ended in the formation of a visible barrier of bacterial cells toward the centre of the coverslip that accumulated presumably at a preferred concentration of the chemical treatment (Fig. 2C). Beyond this barrier, bacterial movement resembled that of the untreated controls. These observations show that *H. pylori* strain G27 responds to pH similarly to strain SS1 (Croxen et al., 2006). When treated with 0.1 mM synthetic DPD, the cultures exhibited a response that was similar to that to acid: the cells showed increased frequency of stops and formed a bacterial barrier at a similar location away from the chemical source (Fig. 2D). In response to mock DPD, the percentage of motile bacteria in the population increased, but there was no obvious net directional movement, and after 5 min no bacterial barrier
had formed and the bacteria were evenly distributed underneath the coverslip (Fig. 2B). Our data thus suggest that *H. pylori* cells respond chemotactically to synthetic DPD similarly to how they respond to HCl.

### The chemotaxis receptor TlpB is required for chemotaxis away from AI-2

Bacterial tactic movement toward or away from a chemical source is generally regulated by the chemotaxis system. Key to this system are the chemoreceptors which sense environmental signals and transmit the information through a transduction cascade, eventually affecting flagellar rotation (Armitage, 1999). To test whether the response to synthetic DPD was due to detection by any of the four *H. pylori* chemoreceptors, we repeated the wet-mount assay described above, using mouse-passaged *H. pylori* mG27 isogenic strains harbouring deletions or insertional mutations in the chemoreceptor genes *tlpA* (strain ΔtlpA), *tlpB* (strain ΔtlpB), *tlpC* (strain ΔlpC) or *tlpD* (strain ΔlpD). The wild-type strain mG27 does not produce TlpC as detectable by Western blotting (Fig. 3A), but we included the engineered *tlpC* mutant in our analysis for completeness. Each of these strains was challenged with 0.1 mM synthetic DPD, 0.1 M HCl or mock DPD. As a negative control for chemotaxis we employed a non-chemotactic mutant with a mutation in the chemotransduction gene *cheA*. The ΔtlpA, ΔlpC and ΔlpD mutants all formed bacterial barriers when challenged with synthetic DPD and HCl (Fig. 2F, G, L, M, O, P). As previously reported, the ΔtlpB mutant failed to form a bacterial barrier when challenged with HCl (Fig. 2I). In addition, this mutant failed to form a bacterial barrier when challenged with synthetic DPD (Fig. 2J). As expected, the *cheA* mutant did not form a bacterial barrier in response to HCl or synthetic DPD (Fig. 2R, S). None of the five strains produced bacterial barriers in response to mock DPD (Fig. 2E, H, K, N, Q). These data implicated TlpB in the sensing of AI-2.

To verify that the failure of the ΔtlpB mutant to form a bacterial barrier in response to DPD was due to the lack of the *tlpB* gene product, we constructed a complemented strain (referred to as *tlpB*+) by placing the full-length *tlpB* gene in the *rdxA* locus of the ΔtlpB mutant chromosome. Western blot analysis verified production of the TlpB protein in the *tlpB*+ strain (Fig. 3A). We then repeated the wet-mount assay with the *tlpB*+ strain. When challenged with 0.1 mM synthetic DPD or 0.1 M HCl, the *tlpB*+ strain responded by producing a bacterial barrier, in contrast to the mock DPD solution, which produced a uniform distribution of bacteria (Fig. 3B).

To further characterize TlpB as a chemoreceptor for synthetic DPD, we repeated the video taxis assay with the mG27 wild-type, ΔtlpB and *tlpB*+ isogenic strains. The average number of stops for each genotype was recorded for each strain grown in mock DPD, 0.1 M HCl or 0.1 mM synthetic DPD. As reported previously (Croxen et al., 2006), we observed that the ΔtlpB mutant displayed an increased stopping frequency as compared to its wild-type parent in medium without added chemorepellents (see below). To best represent the relative responsiveness of the different strains to HCl and DPD, we therefore show the normalized stopping frequencies for each strain in each condition relative to the average number of stops exhibited by that strain in the mock DPD medium (Fig. 3C). In accordance with the wet-mount assay, the ΔtlpB mutant exhibited significantly reduced responsiveness to DPD and HCl as compared with the wild-type strain. The complemented *tlpB*+ strain had restored responsiveness to DPD.

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**Fig. 2.** Chemotaxis response of wild-type and mutant *H. pylori* to HCl and synthetic DPD. A wet-mount chemotaxis assay (illustrated in A) was used to analyse chemotactic behaviour of mG27 wild-type and the indicated isogenic mutant strains in response to mock DPD, the known chemorepellent 0.1 M HCl, and 0.1 mM synthetic DPD. White arrows indicate bacterial barrier formation.
and greater responsiveness to acid than the wild-type strain. We do not know the reason for this strain’s hyper-responsive-ness to acid, but it could be due to altered expression of the tlpB gene under acidic conditions when expressed from a different promoter and genomic locus than those of the native gene.

As noted above, the ΔtlpB mutant displayed an increased stopping frequency as compared to its wild-type parent when grown in medium in the absence of exogenously added chemorepellents (Fig. 4A). We wondered whether this increased frequency was due to a hyper-activation of the chemotaxis signal transduction pathway, or some other effect on swimming behaviour caused by the loss of tlpB. To distinguish these possibilities, we constructed a tlpB cheA double mutant and compared the swimming behaviour of this strain to the tlpB and cheA single mutants. As expected, the cheA single mutant, which lacks a functional chemotaxis signal transduction pathway, exhibited fewer stops than the wild-type strain in BB10 (Fig. 4A). The tlpB cheA double mutant was indistinguishable from the cheA single mutant, indicating that the increased stopping observed with the tlpB mutant was due to increased activation of the chemotaxis pathway, possibly as a result of increased signalling from other chemoreceptors in the absence of TlpB.

**TlpB functions downstream of LuxS in Al-2 responses**

Our data suggest that TlpB functions as the chemoreceptor for avoidance responses to Al-2, a molecule produced by the bacteria themselves using the enzyme encoded by luxS. If this model were correct, then we would predict that cells lacking both LuxS, the enzyme that makes Al-2, and TlpB, the chemoreceptor involved in its perception, should resemble cells lacking only the receptor. We generated a luxS tlpB double mutant and compared its swimming behaviour to that of luxS and tlpB single mutants. We observed that the luxS tlpB double mutant exhibited a high stopping frequency, statistically indistinguishable from the tlpB single mutant (Fig. 4A). Importantly, whereas the luxS single mutant formed barriers to both DPD and HCl in the wet-mount assay (Fig. 4F, G), the luxS tlpB double mutant failed to form any barriers (Fig. 4I, J), similar to the tlpB single mutant.
defect on soft agar (Rader et al., 2007), which was complemented by exogenous AI-2, a result confirmed by others (Shen et al., 2010). This assay cannot distinguish between dysfunction of the motility apparatus and dysregulation of chemotaxis. To better characterize the chemotactic response of H. pylori to the absence or presence of AI-2, we observed both swimming behaviour of individual bacterial cells and the response of populations of bacteria to chemical gradients. We found that luxS-deficient bacterial cells exhibited a decrease in stopping frequency as compared to wild-type cells. This behaviour was restored to wild-type levels by the addition of synthetic DPD. Synthetic DPD also increased the frequency of stops in the wild-type strain, and caused populations of wild-type cells to move away from a source of the chemical, suggesting that AI-2 is a chemorepellent for H. pylori. Our ability to modulate H. pylori swimming behaviour with exogenous AI-2 argues against a metabolic requirement for the luxS gene in chemotaxis. Additionally, although many bacteria regulate the sensitivity of their chemoreceptors via methylation, a mechanism which could be influenced by function of the SAM pathway, H. pylori lacks the methyl-utilizing enzymes required for this process (Szurmant & Ordal, 2004). We further demonstrated that an H. pylori strain deficient for the chemoreceptor TlpB failed to move away from a source of synthetic DPD, and did not display increased stopping behaviour upon addition of synthetic DPD. These behaviours were restored upon genetic complementation of the tlpB gene.

**TlpB and AI-2 perception**

Most reported responses to AIs involve regulation of transcription. Recently, however, E. coli was shown to respond to AI-2 as a chemoattractant through a mechanism that involves both the AI-2 periplasmic binding protein LsrB and the chemoreceptor Tsr (Bansal et al., 2008; Hegde et al., 2011). Here we show that H. pylori also perceives AI-2 as a chemical cue in response to which it directs its movement, in this case by moving away from the signal. We demonstrate that the H. pylori chemoreceptor TlpB is required for perception of AI-2 and confirm its role in negative pH taxis. Bacterial chemoreceptors can often sense more than one ligand and can elicit both positive and negative behavioural responses. For example, the E. coli chemoreceptor Tsr not only senses AI-2, but also mediates taxis toward the attractants serine and related amino acids, and taxis away from weak acids, indole and leucine, and transduces oxygen and redox signals (Boyd & Simon, 1982; Rebbapragada et al., 1997). We do not yet know whether TlpB senses AI-2 directly or via other binding proteins. TlpB shares no sequence similarity with the previously identified AI-2 binding proteins, V. harveyi LuxP (Chen et al., 2002), S. typhimurium LsrB (Miller et al., 2004) and Aggregatibacter actinomycetemcomitans RsbB and LsrB (Shao et al., 2007), and the H. pylori genome lacks homologues of any of these genes. We believe that the molecular mechanism of chemotaxis from AI-2 is distinct

**(Fig. 21, J).** These data demonstrate that tlpB is epistatic to luxS, consistent with our model that TlpB perceives a molecule produced by the luxS gene product.

**DISCUSSION**

In this study we have shown that the quorum-sensing signal AI-2 acts as a chemorepellent for H. pylori, and demonstrated that the chemoreceptor TlpB is required for its perception. We previously reported that elimination of the luxS gene product in H. pylori exhibited a motility
from AI-2-mediated regulation of flhA transcription in H. pylori, because the latter does not require tlpB (B. A. R. and K. G., unpublished results). In addition, neither tlpB nor several other chemotaxis genes appear to be regulated transcriptionally by AI-2 because their transcript levels are the same in wild-type and ΔluxS mutant strains (Rader et al., 2007; B. A. R. and K. G., unpublished results).

**AI-2-regulated motility is likely to be important in the gastric environment**

For *H. pylori*, motility is essential for colonization of piglet, gerbil and mouse stomachs, where bacterial interactions with the mucosa promote gastric pathology (Eaton et al., 1992; McGee et al., 2002; O’Toole et al., 2000; Ottemann & Lowenthal, 2002). It is well established that non-chemotactic mutants (Che−) of *H. pylori* do not infect mice to the full wild-type level (Foynes et al., 2000; Terry et al., 2005). These mutants, lacking cheA, cheW or cheY, all engage almost exclusively in straight runs without stops or changes in direction. We found that the luxS mutant exhibits a similar swimming behaviour to the Che− mutants in broth, suggesting that LuxS is responsible for production of a significant proportion of chemorepellent signals present in *H. pylori* batch culture. We have confirmed previous reports that a luxS mutant is defective in colonization of the rodent stomach; however in these experiments it was not possible to determine whether the colonization defect was due to reduced AI-2 concentrations in the mouse stomach or metabolic deficiencies in the bacteria (Lee et al., 2006; Osaki et al., 2006), (B. A. R., K. G. and K. M. O., unpublished results). Whereas the ΔtlpB mutant colonizes wild-type mice and gerbils to normal levels (McGee et al., 2005; Williams et al., 2007), it is impaired in its ability to colonize the stomachs of IL-12-deficient mice (Croxon et al., 2006). There are likely to be additional chemical cues to which *H. pylori* responds within a mouse stomach (Schreiber et al., 2004). Indeed, non-chemotactic mutants are less closely associated with mouse gastric epithelia and induce a diminished inflammatory response as compared to wild-type *H. pylori*, suggesting that chemical cues from the epithelium are important for directing *H. pylori* localization within the stomach (Williams et al., 2007).

The fact that gastrointestinal pathogens have chemotactic responses to AI-2 raises the question of the role of this response in host colonization. In the case of enteropathogenic *E. coli*, the pathogen perceives AI-2 as an attractant, possibly using it as a cue to direct itself toward the bacteriagene colon. We hypothesize that *H. pylori* perceives AI-2 both as a repellent and as a cue to regulate flagellar gene expression as a means to coordinate its distribution and motility within the stomach. We imagine that at some threshold level of AI-2, the bacteria respond to this molecule as a chemorepellent and move away from the bulk bacterial population, thereby avoiding niche competition and promoting dispersal throughout the stomach. Consistent with this model, luxS mutants in several *H. pylori* strains have been reported to form biofilms more readily than their wild-type counterparts (Cole et al., 2004). With sustained AI-2 levels, indicative of high bacterial populations, the elevated AI-2 may alter FlhA levels to downregulate motility. Such a response might allow bacteria to avoid wasting energy on swimming and instead focus on adherence. Several bacterial species perform reciprocal regulation of motility and adherence (Holden & Gally, 2004), although this type of response has not been shown for *H. pylori*. Furthermore, our understanding of the gastric environment has changed in part due to 16S rRNA gene enumeration studies, which identified over 120 phylotypes of bacteria in the human stomach, many of which potentially produce AI-2 (Bik et al., 2006; Federle & Bassler, 2003). It is possible that *H. pylori* will move away from AI-2 produced by coincident bacteria, providing a further means of avoiding niche competition. As disease outcome is correlated with localization of *H. pylori* populations within the stomach (Blaser & Atherton, 2004), AI-2 may be an important environmental factor in the progression of disease in *H. pylori* infections. Further understanding of those factors that regulate *H. pylori* motility and chemotaxis within the gastric environment will undoubtedly enhance our ability to predict disease outcome and design therapies that can control *H. pylori*-caused chronic inflammation and subsequent progression to gastric cancer.

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