

Involvement of Mismatch Repair in the Reciprocal Control of Motility and Adherence of Uropathogenic *Escherichia coli*

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Type 1 fimbriae and flagella, two surface organelles critical for colonization of the urinary tract by uropathogenic *Escherichia coli* (UPEC), mediate opposing virulence objectives. Type 1 fimbriae facilitate adhesion to mucosal cells and promote bacterial persistence in the urinary tract, while flagella propel bacteria through urine and along mucous layers during ascension to the upper urinary tract. Using a transposon screen of the *E. coli* CFT073 *fim* locked-ON (L-ON) mutant, a construct that constitutively expresses type 1 fimbriae and represses motility, we identified six mutants that exhibited a partial restoration of motility. Among these six mutated genes was *mutS*, which encodes a component of the methyl-directed mismatch repair (MMR) system. When complemented with *mutS* *in trans*, motility was again repressed. To determine whether the MMR system, in general, is involved in this reciprocal control, we characterized the effects of gene deletions of other MMR components on UPEC motility. Isogenic deletions of *mutS*, *mutH*, and *mutL* were constructed in both wild-type CFT073 and *fim* L-ON backgrounds. All MMR mutants showed an increase in motility in the wild-type background, and Δ *mutH* and Δ *mutS* mutations increased motility in the *fim* L-ON background. Cochallenge of the wild-type strain with an MMR-defective strain showed a subtle but significant competitive advantage in the bladder and spleen for the MMR mutant using the murine model of ascending urinary tract infection after 48 h. Our findings demonstrate that the MMR system generally affects the reciprocal regulation of motility and adherence and thus could contribute to UPEC pathogenesis during urinary tract infections.

Flagella, surface organelles that drive motility, and fimbriae, surface appendages that mediate adherence, perform opposing functions, and yet both are important for some bacterial pathogens to infect their host (5, 43). Specifically, in uropathogenic *Escherichia coli* (UPEC), type 1 fimbriae are highly expressed *in vivo* in the murine model of urinary tract infections (UTI) and are considered necessary for the bladder colonization by UPEC in this model (19, 52). Whereas type 1 fimbriae allow establishment of infection in the bladder by binding to mannose-containing glycoproteins, flagella allow UPEC to ascend from the bladder to the kidneys (31, 40, 47). One may argue that a single bacterium should not be both adherent and motile simultaneously, and we have provided evidence that the expression of fimbriae and flagella is reciprocally regulated (3, 9, 18, 30, 34).

The promoter that controls type 1 fimbriae expression resides upstream from the major structural subunit gene *fimA* and is located on an invertible element that alternates between two orientations, phase ON and phase OFF (1). When the promoter is facing the *fim* operon (*fimAICDFGH*), it is in the phase ON orientation and type 1 fimbriae are expressed. Alternatively, when the promoter faces in the opposite direction of the *fim* operon, it is phase OFF and the transcription of *fim* genes is repressed (6, 15, 26, 37). Alternatively, flagellin expression is regulated in a hierarchical manner consisting of three classes with more than 40 genes expressed (8, 53). Transcription of the class I flagellar genes, *flhDC*, is necessary for the transcription of the class II flagellar genes, including *fliA*, the sigma factor that promotes expression of class III flagellar genes. Reciprocal control of adherence and motility explains that when it is advantageous for the bacteria to be motile, flagella are expressed and fimbriae are repressed because adherence would impede flagellar motility. Conversely, when it is advantageous for bacteria to be adherent, fimbriae are expressed and motility is repressed. Examples of such regulation have been

observed in other pathogens, including *Vibrio cholerae* (18), *Salmonella enterica* serovar Typhimurium (9), *Bordetella pertussis* (3), and *Proteus mirabilis* (34).

A transposon screen of UPEC CFT073 mutants in which the *fim* promoter is locked in the phase ON orientation (CFT073 *fim* L-ON) identified six nonfimbrial genes involved in the repression of motility: one is *mutS*, a gene encoding a component of the methyl-directed mismatch repair (MMR) system (49). MMR is a well-characterized and widely conserved DNA repair system found in both eukaryotes and prokaryotes. The MMR pathway in *E. coli* is primarily composed of three proteins: MutS, MutL, and MutH (23, 24, 39). MutS recognizes mismatches in newly replicated chromosomal DNA; MutL forms a complex with MutS and recruits MutH in an ATP-dependent manner, allowing MutH to nick the unmethylated daughter strand at a hemimethylated GATC site (4, 21, 28, 56). Helicase II (UvrD), activated by the MutS/MutL complex, separates the daughter and the parent strands (12, 22, 57), while exonuclease Exo I, Exo VII, or RecJ degrades the mismatched containing DNA strand (10). After re-synthesis of the gapped region, DNA ligase seals the nick to complete the repair process (28). MMR primarily corrects single base nucleotide mismatches, and small insertion/deletion mismatches introduced during DNA replication (for reviews, see references 27

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and 46). MMR contributes ~1,000-fold to the overall fidelity of the DNA replication pathway. In addition to correcting base-pairing errors, the MMR system in *E. coli* also prevents the recombination between divergent DNA sequences (44).

The striking role that we identified for MutS in the regulation of motility was confirmed by the construction of a clean isogenic *mutS* deletion in CFT073 that demonstrated increased motility (49). Complementation with *mutS* *in trans* restored motility and flagellin production to wild-type levels (49). We hypothesized that the entire MMR system and not just MutS mediated this reciprocal control of adherence and motility.

To test this hypothesis, we examined the contribution of three components of the methyl-directed mismatch repair system to the motility of wild-type CFT073 and CFT073 *fim* L-ON bacteria. FliC and FimA expression was directly assessed by Western blot, by immunofluorescence microscopy, and by determining the orientation of the type 1 fimbrial promoter by the invertible element assay and indirectly assessed by quantitative PCR (qPCR). Using the murine model of ascending UTI, we demonstrated that the loss of MMR decreased UPEC fitness only when type 1 fimbriae are expressed constitutively, whereas deletion of *mutS* in wild-type CFT073 increased the dissemination of bacteria into the bloodstream.

MATERIALS AND METHODS

Bacterial strains and culture conditions. The prototype uropathogenic *E. coli* strain CFT073 was isolated from the urine and blood of a hospitalized patient suffering from acute pyelonephritis (38, 55). All *E. coli* strains and plasmids used in the present study are listed in Table 1. *E. coli* strains were cultured on Luria-Bertani (LB) agar or in LB broth and were incubated overnight at 37°C; liquid cultures were agitated at 200 rpm. Ampicillin (100 µg/ml), kanamycin (25 µg/ml), and chloramphenicol (20 µg/ml) were included as appropriate.

Construction and complementation of isogenic mutants of *E. coli* CFT073 and CFT073 *fim* L-ON. Deletion of MMR genes was accomplished in strains CFT073 and CFT073 *fim* L-ON using lambda red recombination technology (13). To construct CFT073 Δ *mutH*, CFT073 *fim* L-ON Δ *mutH* (Δ *mutH* L-ON), and CFT073 *fim* L-ON Δ *mutL* (Δ *mutL* L-ON), using primers homologous to sequences flanking the 5' and 3' ends of the target gene, the kanamycin resistance gene was PCR amplified from plasmid pKD4 (the primers are listed in Table 1). To create CFT073 Δ *mutL* the same technology was used; however, the chloramphenicol resistance gene from template plasmid pKD3 was amplified (Table 1). Mutants were verified by using diagnostic PCR with the primers listed in Table 1. CFT073 Δ *mutS* and CFT073 *fim* L-ON Δ *mutS* (Δ *mutS* L-ON) mutants were constructed and verified as previously described (49).

To complement mismatch repair mutants, target genes with associated promoters were PCR-amplified (Table 1). The resulting PCR fragments were digested with EcoRI and HindIII, followed by ligation into a digested pGEN-MCS plasmid, a derivative of pGEN222 (17), with EcoRI and HindIII. The newly constructed pGEN plasmids with the inserted target gene were used to transform their respective mutant strains. Antibiotic resistance to ampicillin, as well as antibiotic resistance of the mutant strain to kanamycin or chloramphenicol, was used to select for transformants. Constructs were confirmed by PCR using the primer sets listed (Table 1).

Motility assay on soft agar. Soft agar plates were used to evaluate the motility of bacterial strains and constructs, as previously described (32). Briefly, a sterile inoculating needle was dipped into a standardized (optical density at 600 nm [OD₆₀₀] of 1.0), late-log-phase liquid culture, stabbed into the middle of a soft agar (0.25% [wt/vol]) plate, where touching the bottom of the plate was avoided. Motility diameters were measured after 16 h of incubation at 30°C. This was repeated at least in triplicate, and

wild-type CFT073 and CFT073 *fim* L-ON were always included as controls. Ampicillin was added to the cultures and soft-agar plates when determining the motility of complemented mutants.

Capillary chemotaxis assay. An alternative standardized quantitative measurement of random motility was conducted by using a modified capillary chemotaxis assay (2, 29, 42). Strains cultured on motility agar and then used to inoculate LB medium were incubated to mid-exponential phase (OD₆₀₀ = 0.3 to 0.4), harvested by centrifugation to remove all spent medium, and resuspended and diluted (1:10) in phosphate-buffered saline (PBS). Samples of diluted, resuspended cultures (100 µl) were serially diluted and plated where CFU/ml was determined after an overnight incubation at 37°C. Bacterial suspensions (500 µl) were added to the wells of the chemotaxis chambers, followed by microcapillaries filled with sterile PBS. Chemotaxis chambers were incubated at 30°C for 90 min, and the contents of the capillaries were diluted and plated. The CFU/ml was determined after an overnight incubation at 37°C, and results are presented as the percentage of input bacteria present in the capillary. Assays were conducted in triplicate, and wild-type CFT073 and CFT073 *fim* L-ON were always included as controls.

Detection of flagellum and type 1 fimbrial protein production. Western blotting was used to detect flagellin and type 1 fimbrial protein production, while qPCR was used to detect *fliC* and *fimA* transcript levels. For Western blotting, 15 µl of standardized (OD₆₀₀ = 1.0), mid-log-phase cultures were added to 4 µl of 6× sodium dodecyl sulfate (SDS) loading buffer and boiled for 10 min. Lysates were electrophoresed on a 12.5% denaturing SDS-polyacrylamide (SDS-PAGE) gel and transferred to a polyvinylidene difluoride (PVDF) membrane (Immobilon-P; Millipore Corp.). The membrane was blocked with 2% milk in Tris-buffered saline plus 0.01% Tween 20 (TBST). FliC was detected with rabbit anti-FliC (1:40,000 dilution with TBST), followed by anti-rabbit-horseradish peroxidase (HRP) (1:20,000 dilution with TBST) (Sigma) and Amersham ECL Plus detection reagents (GE Healthcare Life Sciences).

Standardized whole-cell lysates were used to detect the FimA subunit of type 1 fimbriae by Western blotting. Bacterial pellets from 100 µl of culture were resuspended in 100 µl of acidified double-distilled H₂O (pH 1.8) and boiled for 7 min. 15 µl of resuspended lysates were added to 10 µl of 2.5× SDS loading buffer, neutralized with 1 N NaOH (~2.0 µl), electrophoresed on a SDS–12.5% PAGE gel, and transferred to a PVDF membrane blocked as previously described. FimA was detected using rabbit anti-FimA (1:10,000 dilution with TBST), followed by anti-rabbit-HRP (1:20,000 dilution with TBST), and Amersham ECL Plus detection reagents.

qPCR. To extract total *fliC* and *fimA* RNA for qPCR, 500 µl of a standardized (OD₆₀₀ = 1.0), late-log-phase culture was added to 65 µl of ice-cold phenol-ethanol stop solution (5% phenol in ethanol) and was collected by centrifugation. Using lysozyme, cells were lysed and RNA was extracted using the RNeasy kit (Qiagen) following the manufacturer's recommended protocol. Nucleic acid concentrations were determined by spectrophotometry (NanoDrop), and any DNA contamination was removed by incubation of the samples with 4 U of Turbo DNase (Ambion). RNA was recovered, quantified, and used as a template for PCR to confirm inactivation of any trace DNA. PCR-negative RNA was used for first-strand cDNA synthesis using SuperScript II reverse transcriptase (Invitrogen) according to manufacturer's protocol and using 50 ng of random hexamers. cDNA was purified on a QIAquick column (Qiagen), quantified, and diluted to 6 ng/µl with nuclease-free water. RNA transcripts were quantified on a MX3000P real-time PCR instrument (Stratagene) using Brilliant SYBR green qPCR mix (Stratagene) in 25-µl volumes containing 30 ng of cDNA. Optimal primer concentrations were determined empirically and are listed in Table 1.

For comparative, quantitative analysis, transcript levels were normalized to the level of *gapA* (glyceraldehyde 3-phosphate dehydrogenase A). Fold changes were determined using wild-type CFT073 and CFT073 *fim* L-ON as calibrators and the MXPro v 3.00 software package (Stratagene). Any transcript level comparisons greater than a 2-fold change (log₂ = 1)

TABLE 1 Relevant bacterial strains, plasmids, and primers used in this study

Strain, plasmid, or primer	Description ^a or sequence (5'–3')	Source or reference
<i>E. coli</i> strains		
CFT073	Wild-type pyelonephritis isolate	38
<i>fim</i> L-ON	CFT073 <i>fim</i> IE phase locked ON; constitutive type 1 fimbrial expression	19
<i>fim</i> L-OFF	CFT073 <i>fim</i> IE phase locked OFF; blocking type 1 fimbrial expression	19
Δ <i>mutS</i> mutant	CFT073 Δ <i>mutS</i> :Kan ^r	49
Δ <i>mutH</i> mutant	CFT073 Δ <i>mutH</i> :Kan ^r	This study
Δ <i>mutL</i> mutant	CFT073 Δ <i>mutL</i> :Cam ^r	This study
<i>fim</i> L-ON Δ <i>mutS</i>	CFT073 <i>fim</i> L-ON Δ <i>mutS</i> :Kan ^r	49
<i>fim</i> L-ON Δ <i>mutH</i>	CFT073 <i>fim</i> L-ON Δ <i>mutH</i> :Kan ^r	This study
<i>fim</i> L-ON Δ <i>mutL</i>	CFT073 <i>fim</i> L-ON Δ <i>mutL</i> :Kan ^r	This study
Plasmids		
pGEN-MCS	Derivative of pGEN222 with a multiple cloning site replacing <i>gfpuv</i> (Amp ^r)	31
pGEN- <i>mutS</i>	CFT073 <i>mutS</i> cloned into BamHI-SphI sites of pGEN-MCS (Amp ^r)	49
pGEN- <i>mutH</i>	CFT073 <i>mutH</i> cloned into EcoRI-HindIII sites of pGEN-MCS (Amp ^r)	This study
pGEN- <i>mutL</i>	CFT073 <i>mutL</i> cloned into EcoRI-HindIII sites of pGEN-MCS (Amp ^r)	This study
Primers		
Invertible element		
IE-P1 forward	AGTAATGCTGCTCGTTTTGC	
IE-P2 reverse	GACAGAGCCGACAGAACAAC	
Mutant construction		
<i>mutS</i> H1P1	TGAAAGCCCAGCATCCCGAGATCCTGCTGTTTTACCGGATGTGTAGGCTGGAGCTGCTTC	
<i>mutS</i> H2P2	TGCGGTGCGCGCTTAATAACCTCTTTTGGCACACCCGCCAATGGGAATTAGCCATGGTCC	
<i>mutH</i> H1P1	ATGTCCCAACCTCGCCCACTGCTCTCTCCTCCCGGTGTAGGCTGGAGCTGCTGCTTC	
<i>mutH</i> H2P2	ATCCGTTCCGCCCCGGGCACCAATAGCTTCGGTACGATGGGAATTAGCCATGGTCC	
<i>mutL</i> L-ON H1P1	GGGTGCAACGGCACAACCGCCAGTACGGTGACGACGGTGTAGGCTGGAGCTGCTTC	
<i>mutL</i> L-ON H2P2	GCGGACATAACCGTTCCACGTCCGCCAGCAGGGATGGGAATTAGCCATGGTCC	
<i>mutL</i> H1P1	ACCGCCACAACCTGGCGAACAGATTGCCGAGGTGTAGGCTGGAGCTGCTTC	
<i>mutL</i> H2P2	TGGCTGTGCCATTGACCACTGCGCATGTTCCGATGGGAATTAGCCATGGTCC	
Mutant screen		
<i>mutS</i> forward	CCCGAGATCCTGCTGTTTTA	
<i>mutS</i> reverse	TGCGCGCTTAATAACCTCTT	
<i>mutH</i> forward	GTTACACTGCGAATATTCAGC	
<i>mutH</i> reverse	CAATCCTAGCCAATAGGTACG	
<i>mutL</i> forward	GATGCAATCTGCGCCGACG	
<i>mutL</i> reverse	CAGCTCAATGGCTAACGCCG	
Mutant complementation		
pGEN-MCS forward	ACTTGCTCACGCTCTGTCAT	
pGEN-MCS reverse	AGAAGGACCATGTGGTCACG	
<i>mutS</i> forward	CAGCATGCGGACATGCTGCCTCCATACT	
<i>mutS</i> reverse	TGGATCCAGCAAAAGGCTATCGGGAAT	
<i>mutH</i> forward	ATATATGAATTCCGATAACATGGTGTAAAGATCCTGTT	
<i>mutH</i> reverse	GTGCTGTGGGAAGCTTTGCCACCAC	
<i>mutL</i> forward	ATATATGAATTCTGTTCTGCGTTCCGCCGATATTCGG	
<i>mutL</i> reverse	ATATATAAGCTTCTGCGACGGATCGCGAATATTCAG	
Sanger sequencing		
<i>mutH</i> forward	GGCCGATACAAAACCGGCTTCGTGTC	
<i>mutH</i> reverse	CGCCATGACGTAATCACTGGAACAC	
<i>mutL</i> forward	GTACCTTGCTCAAT	
<i>mutL</i> reverse	CGAACTGATGCAC	
Pap promoter forward	GCGTGAAGAGTATTTCCGGGGCC	
Pap promoter reverse	CCATAGCTACCGCACCGGCA	
FlhDC promoter forward	TCTGTGAACCTTCAGGTGAC	
FlhDC promoter reverse	TCAGCAACTCGGAGGTATGC	
qPCR		
<i>gapA</i> forward	AAGTTGGTGTGACGTTGTGCG	
<i>gapA</i> reverse	AGCGCCTTTAACGAACATCG	
<i>fliC</i> forward	ACAGCCTCTCGCTGATCACTCAA	
<i>fliC</i> reverse	GCGCTGTTAATACGCAAGCCAGA	
<i>fimA</i> forward	ACTCTGGCAATCGTTGTTCTGTGCG	
<i>fimA</i> reverse	ATCAACAGAGCCTGCATCAACTGC	

^a Cam^r, chloramphenicol resistance; Amp^r, ampicillin resistance; Kan^r, kanamycin resistance.

were considered significant changes. Assays were conducted in duplicate. Wild-type CFT073 and CFT073 *fim* L-ON were always included as controls.

Indirect fluorescent antibody staining. Sequential staining was used to quantify surface flagella. Standardized bacterial cultures ($OD_{600} = 1.0$) were diluted with LB (1:2), spotted, and dried onto glass microscope coverslips (Fisher), and fixed with 4% formaldehyde in PBS. Coverslips were washed with PBS three times, and flagella were detected with rabbit anti-FliC (1:1,000 dilution with 1% fetal bovine serum [FBS]) and anti-rabbit IgG (1:5,000 dilution with 1% FBS) (Alexa Fluor 488 conjugate; Molecular Probes). To detect nucleic acid, the slides were washed once with a dilution of 1.0 mg/ml propidium iodide (Sigma) in PBS (1:10,000), washed twice with PBS, and mounted using *p*-phenylenediamine onto glass microscope slides (25 by 75 by 1 mm; VWR). Slides were viewed on an Olympus BX60 upright microscope; images were quantified, and backgrounds were corrected using Metamorph Premier software (Molecular Devices, Inc.). Ratios of parental flagellar counts were calculated by dividing the ratio of flagella per mutant cell by the ratio of flagella per parental strain, quantified over three microscopic fields.

Invertible element PCR assay. The orientation of the *fim* invertible element was determined using a PCR assay as previously described (35). Briefly, the primers IE-P1 and IE-P2 (Table 1) were used to amplify the 601-bp invertible element using bacteria from standardized ($OD_{600} = 1.0$) cultures. CFT073 and CFT073 *fim* L-ON were included as controls. Asymmetric digestion of the PCR products with SnaBI allowed determination of the orientation of the invertible element when electrophoresed on a 2% agarose-1× Tris-acetate-EDTA buffer (TAE) gel.

Measurement of growth rate. Growth curves of the mutant constructs were generated by inoculating 3 ml of LB broth with 50 μ l overnight culture, in duplicate, using μ Quant (Bio-Tek Instruments, Inc. [KCJunior Software]). The OD_{600} of a culture sample (200 μ l) was recorded every hour for 7 h. Wild-type CFT073 and CFT073 *fim* L-ON were always included as controls. Growth rates were determined using the equation $\mu = [(\log_{10} N - \log_{10} N_0) 2.303] / (t - t_0)$, where N is the concentration of cells (number of cells per unit volume) and t is time.

Measurement of spontaneous mutation frequency. Spontaneous mutation frequencies were determined as described previously (48, 51). Briefly, 1.5 ml of late-log-phase cultures were standardized ($OD_{600} = 1.0$), centrifuged, and resuspended in 100 μ l of 0.85% saline. The suspension was serially diluted and plated onto LB agar plates. The CFU/ml was determined after an overnight incubation at 37°C. The remainder of the suspension was plated onto LB agar plates containing rifampin (100 μ g/ml). CFU of rifampin-resistant mutants were scored after an overnight incubation at 37°C, and spontaneous mutation frequencies were calculated by dividing the number of rifampin-resistant colonies by total viable cells.

CBA/J mouse model of ascending UTI. Female CBA/J mice, 6 to 8 weeks old, were inoculated transurethrally as previously described (20). Briefly, pelleted overnight cultures (50 ml) were resuspended to an OD_{600} of ~ 4.0 in sterile PBS. Diluted bacterial suspensions were mixed in a 1:1 ratio, and 50 μ l containing 10^8 CFU was used to transurethrally inoculate each mouse. At 48 h or 7 days postinoculation, urine was collected, and the bladders, kidneys, and spleens were aseptically removed, weighed, and homogenized (Omni International). Serial dilutions were plated onto LB agar plates with or without kanamycin using a spiral plater (Autoplate 4000; Spiral Biotech) to quantify CFU of wild-type CFT073 (Kan^s) and mutant (Kan^r), respectively. Competitive indices were calculated by dividing the ratio of wild-type to mutant at an endpoint CFU/ml by the ratio of wild-type to mutant input CFU/ml. A competitive index < 1 indicates a fitness defect of the mutant strain.

Statistical analysis. A paired Student *t* test was used to determine significant differences in motility between strains in motility assays (InStat; GraphPad Prism Software). For the murine model, statistically significant differences of *in vivo* colonization ($P < 0.05$) were calculated using a two-tailed, Wilcoxon signed-rank test where column medians

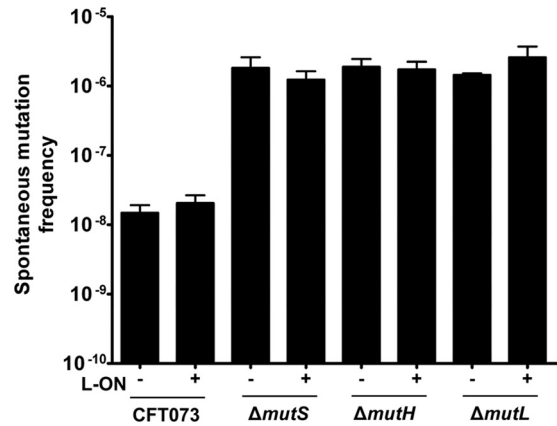


FIG 1 Frequency of rifampin-resistant mutants per number of isolates of motile CFT073 mismatch repair mutants. The “+” and “-” signs represent the *fim* L-ON status of the CFT073 background; “+” represents CFT073 *fim* L-ON, and “-” represents CFT073 wild type. Error bars represent the standard errors of the mean (SEM) of three independent cultures.

were compared to a hypothetical value of zero (InStat; GraphPad Prism Software). Prism Software (GraphPad) was also used in linear regression analyses of motility data.

RESULTS

Deletions in the mismatch repair system increase the mutation rate of CFT073. Deletion mutants were constructed for *mutS*, *mutH*, and *mutL* in both the wild-type CFT073 and CFT073 *fim* L-ON backgrounds. To verify that mutants were MMR defective, we measured the spontaneous mutation frequency by determining the occurrence of rifampin-resistant bacteria in culture (54). Rifampin resistance can be conferred by missense mutations in the *rpoB* (β subunit) gene of RNA polymerase. Defects in MMR cause an increase in spontaneous mutation frequency, which can be measured by an increase in the occurrence of rifampin-resistant bacteria (54).

Deletion of *mutS*, *mutH* and *mutL* resulted in an ~ 100 -fold increase in spontaneous mutation frequency in both the wild-type CFT073 and CFT073 *fim* L-ON genetic backgrounds. The frequency of mutant occurrence was nearly indistinguishable between the *mutS*, *mutL*, and *mutH* deletion constructs (Fig. 1). These results confirm that the MMR pathway is inactive in constructs deleted for *mutS*, *mutH*, and *mutL* (33).

Deletion of mismatch repair genes in both wild-type CFT073 and *fim* L-ON backgrounds increases motility. A previous study in our lab reported that MutS plays a role in the reciprocal control of motility and adherence in CFT073 (49). Deletion of *mutS* partially restored motility of a *fim* locked phase ON strain of CFT073 and increased motility of wild-type CFT073. Thus, we hypothesized that an intact MMR system was required to properly regulate motility and adherence. We predicted that mutation of *mutH* or *mutL*, two other genes encoding MMR components, would also result in increased motility. To test whether MutS alone or the entire MMR system plays a role in the reciprocal control of motility and adherence, the motilities of MMR deletion mutants *mutS*, *mutH*, and *mutL*, constructed in both CFT073 and CFT073 *fim* L-ON strains, were assessed on soft agar plates (Fig. 2A). CFT073 $\Delta mutS$ exhibited an average motility diameter increase of $9.0 \text{ mm} \pm 3.6 \text{ mm}$ compared to the wild-type CFT073 parental

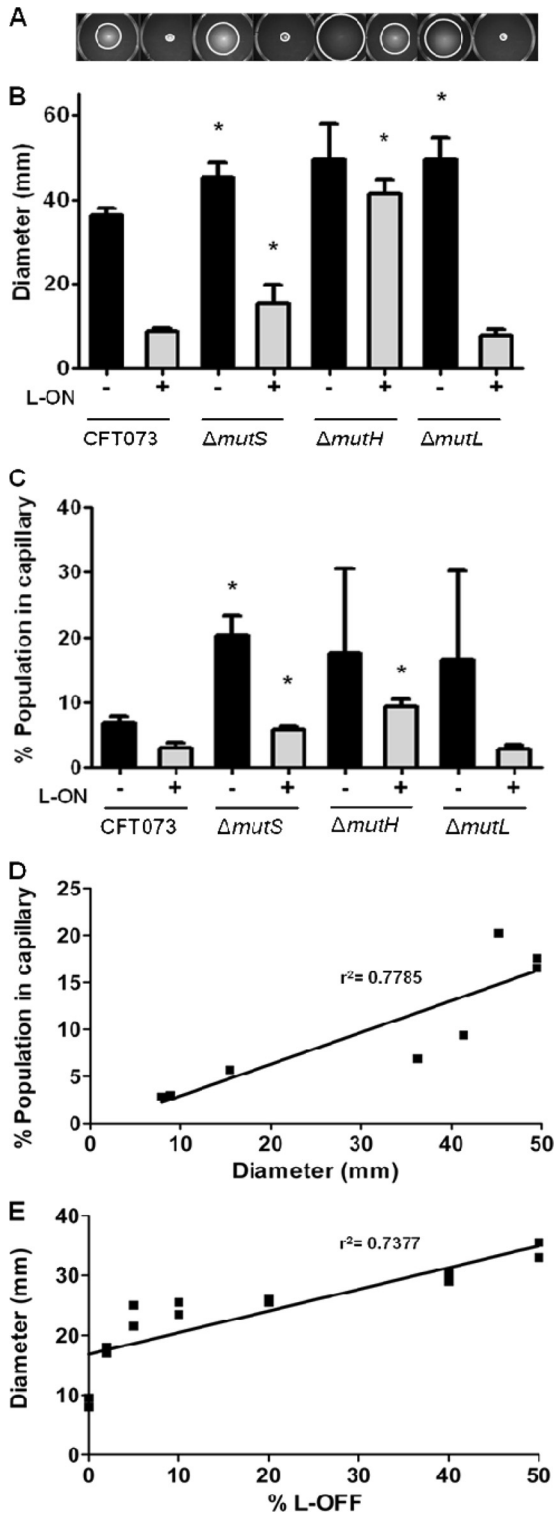


FIG 2 Analysis of motility, percentage of motile bacteria within a population, correlation analysis between soft-agar motility, and percentage of motile bacteria within a population of CFT073 MMR mutants in wild-type and *fim* L-ON backgrounds and analysis of motility of mixed cultures with CFT073 *fim* L-ON and CFT073 *fim* L-OFF. (A) Motility of MMR mutants in soft agar are shown. (B) Bar graph representation of swimming motility diameters of motile deletion mutants. (C) Graphical representation of the percentage of motile bacteria within the populations of MMR mutants using the modified capillary assay. (D) Correlation between the average soft-agar swimming diameter and the

strain ($P = 0.034$) (Fig. 2B). A significant increase in motility ($6.6 \text{ mm} \pm 2.8 \text{ mm}$) was also observed by $\Delta mutS$ L-ON compared to the parental CFT073 *fim* L-ON strain ($P = 0.039$) (Fig. 2B). Deletion of *mutH* increased motility in both the wild-type CFT073 and the CFT073 *fim* L-ON, $13.2 \text{ mm} \pm 7.0 \text{ mm}$ compared to wild-type CFT073 ($P = 0.085$) and $32.5 \text{ mm} \pm 3.1 \text{ mm}$ compared to CFT073 *fim* L-ON ($P < 0.001$) (Fig. 2B), respectively. Interestingly, the motility increase of $\Delta mutH$ L-ON constitutes a full restoration of motility to CFT073 wild-type levels where only partial restoration of motility had been observed with $\Delta mutS$ L-ON (Fig. 2B). This unexpected result is examined below. Strain CFT073 $\Delta mutL$ also showed a significant increase in motility ($13.2 \text{ mm} \pm 4.8 \text{ mm}$) compared to wild-type CFT073 ($P = 0.017$); however, no motility difference was observed between $\Delta mutL$ L-ON and CFT073 *fim* L-ON ($0.9 \text{ mm} \pm 1.6 \text{ mm}$; $P = 0.589$) (Fig. 2B).

An increase in the proportion of motile and flagellated CFT073 in a population correlates with increased motility. Modified capillary assays were also used to verify the motility phenotypes of MMR deletion mutants where we determined the percentage of the population within a culture that would enter a capillary in 90 min (Fig. 2C). From the assay, ca. $6.9\% \pm 1.0\%$ of the wild-type CFT073 population entered the capillary, while only $3.0\% \pm 0.8\%$ of the CFT073 *fim* L-ON population did so (Fig. 2C). As expected, the $\Delta mutS$, $\Delta mutH$, and $\Delta mutL$ mutants all demonstrated a higher average motility increase compared to wild-type CFT073: $13.2\% \pm 4.0\%$ ($P = 0.014$), $10.7\% \pm 14.0\%$ ($P = 0.459$), and $9.6\% \pm 14.8\%$ ($P = 0.523$), respectively (Fig. 2C). Also, as predicted, $\Delta mutS$ L-ON and $\Delta mutH$ L-ON strains demonstrated a significant increase in motility compared to CFT073 *fim* L-ON: $0.9\% \pm 0.5\%$ ($P = 0.041$) and $6.4\% \pm 2.0\%$ ($P = 0.011$), respectively, whereas $\Delta mutL$ L-ON did not show a significant difference in the percentage of motile population $-0.2\% \pm 1.4\%$ ($P = 0.872$) (Fig. 2C).

Using linear regression analysis (Fig. 2D), a strong positive correlation ($r^2 = 0.7785$, $P = 0.0037$) between average diameters of motility determined by the soft agar motility assay (Fig. 2B) and average percentage of population expressing motility determined by the modified capillary assay (Fig. 2C) was observed.

To further verify whether the proportion of flagellated bacteria in a population correlates with the motility phenotype, CFT073 *fim* L-OFF (highly motile) and CFT073 *fim* L-ON (nonmotile) were mixed together in various ratios and motility was assessed in soft agar as shown in Fig. 2A. As predicted, an increase in the ratio of motile, non-type 1-fimbriated cells (CFT073 *fim* L-OFF) to nonmotile, type 1 fimbriated cells (CFT073 *fim* L-ON) within a population led to an increase in the diameter of swimming motility on soft agar ($r^2 = 0.7377$, $P = 0.0013$) (Fig. 2E).

average percentage of the population in a capillary. There is a positive correlation between soft agar motility and capillary motility, where the slope has a significant deviation from zero ($P = 0.0037$). (E) Measurement of swimming motility diameters from soft agar plates of mixed cultures starting with 100% nonmotile *fim* L-ON and 0% motile *fim* L-OFF strains and going up to 50% *fim* L-ON and 50% *fim* L-OFF. The line represents the best-fitting regression line. The slope has a significant deviation from zero ($P = 0.0013$). (B and C) The “+” and “-” signs indicate the *fim* L-ON status of the CFT073 background; “+” represents CFT073 *fim* L-ON, and “-” represents CFT073 wild type. The data represent the averages of at least three separate experiments. Error bars represent the SEM. Significant differences between MMR strains compared to parental strains were determined using a paired Student *t* test. *, $P < 0.05$.

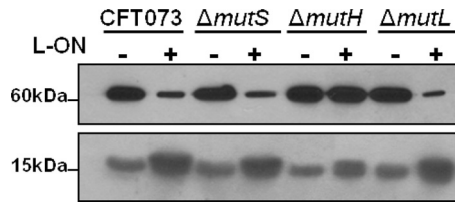


FIG 3 Flagellin and fimbrial protein production of CFT073 MMR mutants in wild-type and *fim* L-ON backgrounds. Western blot analysis of flagellin production, FliC (60 kDa), and fimbrial production, FimA (15 kDa). The “+” and “-” signs indicate the *fim* L-ON status of the CFT073 background; “+” represents CFT073 *fim* L-ON, and “-” represents CFT073 wild type.

From these findings we conclude that motility diameters observed on soft agar plates have a strong positive correlation with motility measured in the standardized capillary assay conducted in Palleroni chambers (Fig. 2D). Furthermore, the observed motility of MMR mutants could be explained by the increased frequency of type 1 fimbrial promoter switching; as the increase in the population switches from *fim* ON to *fim* OFF, the population exhibits an increased motility (Fig. 2E).

Flagellin and type 1 fimbrial protein expression in CFT073 MMR deletion mutants constructed in wild-type and *fim* L-ON backgrounds correlate with motility studies. To determine whether the increased diameter of motility correlates with increased flagellin production, FliC expression was measured using Western blotting with antibodies against FliC protein. As predicted, the levels of flagellin expression of motile CFT073 wild-type, $\Delta mutS$, $\Delta mutH$, $\Delta mutL$, and $\Delta mutH$ L-ON strains were notably higher compared to the levels of nonmotile CFT073 *fim* L-ON, $\Delta mutS$ L-ON, and $\Delta mutL$ L-ON strains (Fig. 3, 60 kDa). Again, the motility behavior of $\Delta mutH$ L-ON was examined further (see below).

To determine whether the concept of reciprocal control holds true for the MMR mutants, the level of type 1 fimbrial structural subunit was measured using Western blotting against FimA protein. As predicted, the expression of FimA in motile CFT073 wild-type, $\Delta mutS$, $\Delta mutH$, $\Delta mutL$, and $\Delta mutH$ L-ON strains was notably lower compared to the expression of nonmotile CFT073 *fim* L-ON, $\Delta mutS$ L-ON, and $\Delta mutL$ L-ON strains (Fig. 3, 15 kDa). Surprisingly, the hypermotile strain $\Delta mutH$ L-ON exhibited higher levels of FimA than comparably motile strains (see below).

qPCR was used to quantify the levels of *fliC* and *fimA* transcripts of MMR mutants constructed in both wild-type and *fim* L-ON backgrounds where transcript levels were then compared to parental strain levels (Fig. 4). As predicted, CFT073 *fim* L-ON, compared to the wild-type strain, had a 27.1-fold decrease of *fliC* transcript (\log_2 of -4.76) and a 172.5-fold increase (\log_2 of 7.43) in *fimA* transcript (Fig. 4). Motile $\Delta mutS$ and $\Delta mutL$ MMR mutants showed similar *fliC* and *fimA* transcript levels compared to the wild type; 1.6- and 1.6-fold (\log_2 of 0.66 and 0.64) and -2.0 - and 1.0-fold (\log_2 of -0.99 and 0.05), respectively (Fig. 4). The $\Delta mutH$ strain, compared to the wild type, exhibited similar *fliC* transcript levels (1.2-fold, \log_2 of 0.23) and a slight increase in *fimA* transcript levels (2.8-fold, \log_2 of 1.48) (Fig. 4).

$\Delta mutS$ L-ON exhibited similar quantities of *fliC* and *fimA* transcripts compared to CFT073 *fim* L-ON (1.9 [\log_2 of 0.92] and -1.2 [\log_2 of -0.24], respectively), whereas highly motile $\Delta mutH$ L-ON showed a 128.4-fold (\log_2 of 7.01) increase in *fliC* transcript

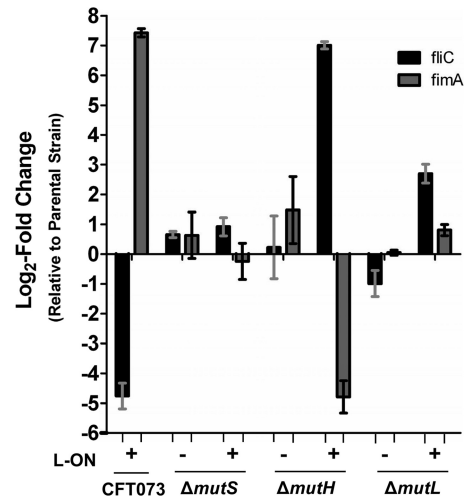


FIG 4 Quantification of flagellar and type 1 fimbrial RNA transcript levels by qPCR of CFT073 MMR mutants in wild-type and *fim* L-ON backgrounds. RNA was isolated from wild-type CFT073, CFT073 *fim* L-ON, and MMR mutants cultured in LB. Transcript levels were measured in cDNA preparations from each strain and normalized to the *gapA* level. The results are shown as the \log_2 fold change relative to parental strain levels. Dashed lines represent a significant cutoff 2-fold change ($\log_2 = 1$), and bars indicate the SEM. The “+” and “-” signs represent the *fim* L-ON status of the CFT073 background; “+” represents CFT073 *fim* L-ON, and “-” represents CFT073 wild type.

levels compared to CFT073 *fim* L-ON (Fig. 4) and even a 4.4-fold increase (\log_2 of 2.14) compared to the wild type, as expected (data not shown) (see below). Also as predicted based on motility data, as well as Western blot data (Fig. 3, 15 kDa), the *fimA* transcript levels of $\Delta mutH$ L-ON were decreased 27.7-fold (\log_2 of -4.79) compared to the *fim* L-ON strain (Fig. 4) and increased 6.6-fold compared to the wild-type strain (\log_2 2.72) (data not shown) (see below). Lastly, nonmotile $\Delta mutL$ L-ON exhibited an unpredicted increase in *fliC* transcript levels of 6.5-fold (\log_2 of 2.70) and similar levels of *fimA* transcript (1.1, \log_2 of 0.81) (Fig. 4).

From these data, we can conclude that motile CFT073 wild-type, $\Delta mutS$, $\Delta mutL$, $\Delta mutH$, and $\Delta mutH$ L-ON strains have a significantly greater quantity of flagellin transcripts and a reduced quantity of type 1 fimbrial transcripts compared to the nonmotile strains. The *fliC* and *fimA* transcript levels in these strains were also found to be relatively similar to one another, with the exception of $\Delta mutH$ L-ON, where increased levels of *fliC* and *fimA* transcripts were detected (see below), and $\Delta mutH$, where a slight increase in *fimA* transcripts was detected; neither of these strain’s *fimA* transcript levels however, reached CFT073 *fim* L-ON quantities. Nonmotile CFT073 *fim* L-ON, $\Delta mutS$ L-ON, and $\Delta mutL$ L-ON strains showed similar levels of *fliC* and *fimA* transcription with the exception of $\Delta mutL$ L-ON, which exhibited a rather large increase in *fliC* transcript levels; this increase, however, did not reach wild-type quantities.

Quantification of surface flagella correlates with motility. Since whole-cell lysates were used to determine whether deletions in the mismatch repair system affected flagellin production, knowledge of the length, number, or presence of surface flagella was still unknown. Immunofluorescence microscopy was used to visualize and quantify flagella, and determine whether the increase in motility of mutants in both wild-type CFT073 and in the CFT073 *fim* L-ON backgrounds could be attributed to flagellar

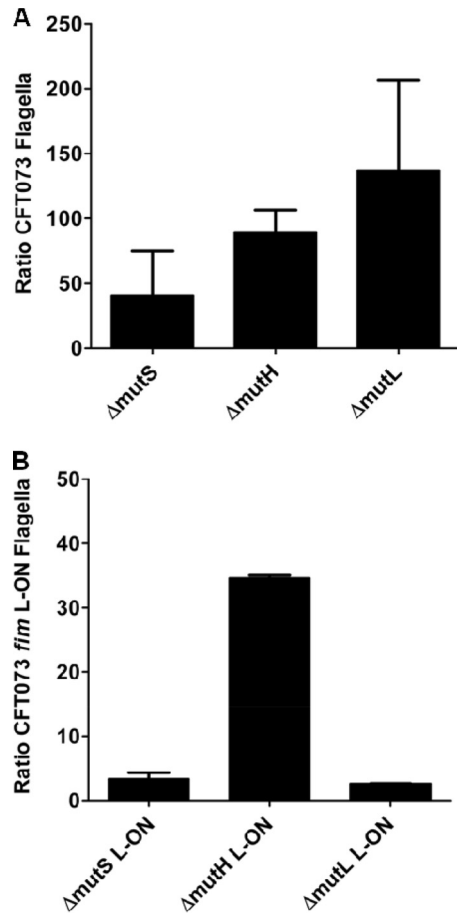


FIG 5 Immunofluorescence analysis of surface flagellum expression of motile MMR mutants in wild-type CFT073 and CFT073 *fim* L-ON backgrounds as a ratio of the parental strain. (A) Bar graphs represent the average counts of flagella per MMR mutant bacterium in a CFT073 wild-type background as a ratio of average wild-type flagella per bacterium. (B) Graphical representation of average counts of flagella per mismatch repair mutant bacterium in CFT073 *fim* L-ON background as a ratio of average CFT073 *fim* L-ON flagella per bacterium is also shown.

expression (Fig. 5). Individual bacteria were stained with propidium iodide, and surface flagella were detected with anti-FliC serum. When comparing CFT073 wild type to the $\Delta mutS$, $\Delta mutH$, and $\Delta mutL$ mutants, we observed flagella that appeared similar in size and morphology; however, we found that the MMR mutant UPEC produced a greater number of flagellated cells (Fig. 5A). When quantified, the average ratio of flagella/ $\Delta mutS$ bacterium was (40.3 ± 48.9) -fold higher than the ratio of flagella/wild-type CFT073 bacterium, the $\Delta mutH$ strain was (89.5 ± 24.0) -fold higher than the wild type, and the $\Delta mutL$ strain was (137.1 ± 97.9) -fold higher than the wild type (Fig. 5A).

We expected to observe few flagella on CFT073 *fim* L-ON because the strain constitutively expressed type 1 fimbriae and lacked motility. Indeed, few, if any, flagella were detected on the surface of CFT073 *fim* L-ON. When comparing $\Delta mutS$ L-ON and CFT073 *fim* L-ON, a slight increase in surface flagella was observed [(3.4 ± 1.3) -fold more; Fig. 5B], which was expected because $\Delta mutS$ L-ON showed a partial restoration of motility compared to CFT073 *fim* L-ON. Due to the increased motility of $\Delta mutH$ L-ON, we expected to observe more flagella than with

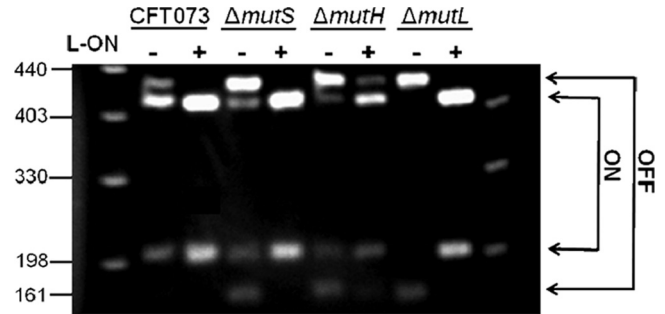


FIG 6 Orientation of the invertible *fim* promoter in CFT073 MMR mutants. The promoter orientation was determined by λ red PCR amplification of the invertible element, followed by digestion using SnaB1. Asymmetric cutting of SnaB1 revealed the orientation of the invertible element; phase ON is indicative of the expression of type 1 fimbriae, and phase OFF is indicative of blocked type 1 fimbrial expression. The “+” and “-” signs represent the *fim* L-ON status of CFT073 background; “+” represents CFT073 *fim* L-ON, and “-” represents CFT073 wild type.

CFT073 *fim* L-ON. Indeed, $\Delta mutH$ L-ON produced a greater number of flagella than CFT073 *fim* L-ON [(34.6 ± 0.6) -fold more; Fig. 5B] and (48.0 ± 41.6) -fold more flagella per bacterium than wild-type CFT073. Lastly, $\Delta mutL$ L-ON expressed (2.6 ± 0.01) -fold more flagella than CFT073 *fim* L-ON (Fig. 5B), which was unexpected since deletion of *mutL* failed to show a partial restoration of motility to the *fim* L-ON strain (Fig. 2B).

Deletion of *mutH* in the *fim* L-ON background reverts the “locked” invertible element back to the wild-type sequence. The $\Delta mutH$ L-ON construct yielded numerous unexpected findings including hypermotility compared to other *fim* L-ON strains, increased levels of FimA and FliC from Western blot compared to other motile strains, and increased levels of *fimA* and *fliC* transcript levels from qPCR data compared to other motile strains. To understand the mechanism underlying these changes, the invertible element assay was used to determine the orientation of the type 1 fimbrial promoter to verify that the mutant remained “locked” in *fim* phase ON orientation (review construction in reference 19). We found, however, that the orientation of the type 1 *fim* promoter of $\Delta mutH$ L-ON was capable of switching, whereas all other *fim* L-ON mutants remained locked in the phase ON orientation (Fig. 6). The *fim* promoters of three independent $\Delta mutH$ L-ON constructs were sequenced and found to have reverted to the wild-type sequence in the left inverted repeat, which had originally been mutated to create the parental locked-ON strain for all MMR *fim* L-ON mutants (19). This result clearly explains the anomalous behavior of this mutant, including the large increase in motility and the increase in flagellin production. In addition, we sequenced the promoter of the electrocompetent cells prepared from CFT073 *fim* L-ON that were used to introduce the *mutH* knockout mutation and confirmed that this sequence was “locked” ON. We believe that the ability to revert to wild-type switching capability and reversion of the left repeat sequence can be attributed to the inadvertent introduction of a methylation site (GATC) to mutate one repeat within the invertible element during the construction of this strain, although the exact mechanism is not known. Therefore, with the exception of $\Delta mutH$ “L-ON”, the deletions of mismatch repair genes do not affect the orientation of the invertible *fim* promoter and is not the source of increased motility when type 1 fimbriae are constitutively expressed.

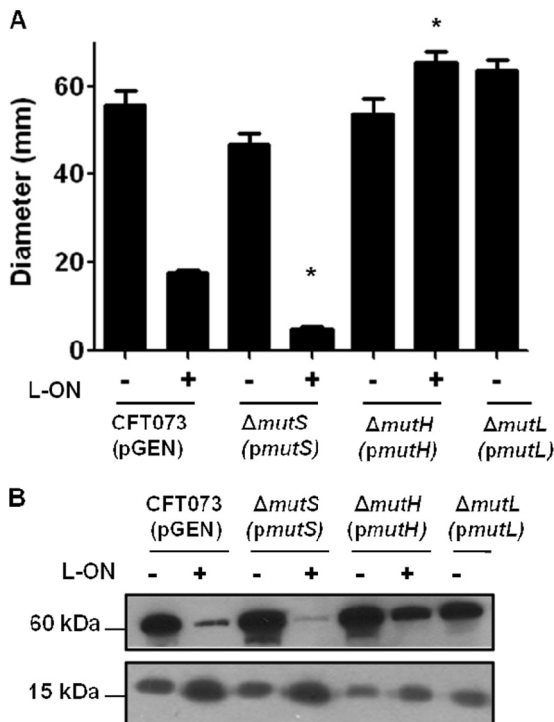


FIG 7 Analysis of motility, flagellin, and fimbrial production of CFT073 mismatch repair mutants in wild-type and *fim* L-ON backgrounds transformed with the complementing plasmid. (A) Graphical representation of swimming motility diameters of complemented motile deletion mutants. The “+” and “-” signs represent the *fim* L-ON status of the CFT073 background; “+” represents CFT073 *fim* L-ON, and “-” represents CFT073 wild type. The data represent averages of at least three separate experiments. Error bars represent the SEM. Significant differences in motility from parental strain motility were determined by using a paired Student *t* test. *, $P < 0.05$. (B) Western blot analysis of flagellin production, FliC (60 kDa), and fimbrial production, FimA (15 kDa). The “+” and “-” signs again represent the CFT073 *fim* L-ON status. The size in kilodaltons of both proteins is given.

Increased motility cannot be attributed to increased growth rate. Since an increase in surface flagella was observed in strains exhibiting increased motility and, with the exception of $\Delta mutH$ “L-ON”, the “locked” *fim* invertible element was demonstrated to be stable; we sought to determine whether changes in motility could be attributed to an increased or decreased growth rate. The growth rate in LB medium for each mutant was compared to its parental strain to assess any growth deficiencies. No notable variations in growth rate were calculated, with the exception of the $\Delta mutH$ strain, where a slight growth deficiency was observed compared to its parental strain, wild-type CFT073 (data not shown). Therefore, we do not attribute the increased motility of the MMR mutant strains to *in vitro* growth.

The motility of MMR mutants can be restored to wild-type levels by complementation. The motility of the complemented MMR mutants was assessed using soft agar and was compared to the motility of MMR mutants transformed with the vector alone and to the motility of parental wild-type CFT073 and CFT073 *fim* L-ON containing vector alone (Fig. 7A). The motility of the complemented $\Delta mutS$ strain was not different from wild-type CFT073 levels ($P = 0.115$), and complementation of the $\Delta mutS$ L-ON strain resulted in decreased motility to CFT073 *fim* L-ON levels ($P < 0.001$) (Fig. 7A). Complemented $\Delta mutH$ strain’s motility

was not different from wild-type CFT073 motility ($P = 0.696$); however, complemented $\Delta mutH$ “L-ON” was not able to reduce motility to CFT073 *fim* L-ON levels ($P < 0.001$) (Fig. 7A). This finding was expected due to the $\Delta mutH$ “L-ON” strain’s ability to unlock itself from the phase ON orientation, as seen in Fig. 6. Therefore, as expected, the motility of the *mutH* “L-ON” mutant was not different from wild-type CFT073 levels ($P = 0.096$) (Fig. 7A). The motility of the complemented $\Delta mutL$ strain was also not different from wild-type CFT073 levels ($P = 0.1466$) (Fig. 7A). Complementation of $\Delta mutL$ L-ON motility was not measurable because the motilities of $\Delta mutL$ L-ON and CFT073 *fim* L-ON were indistinguishable.

Flagellin and FimA levels in the complemented mutant strains correlated well with motility; bacteria that produced more flagellin were more motile than bacteria that produced more FimA. As expected, the flagellin levels of CFT073 and the complemented $\Delta mutS$, $\Delta mutH$, $\Delta mutH$ “L-ON”, and $\Delta mutL$ strains were much higher than the nonmotile CFT073 *fim* L-ON and complemented $\Delta mutS$ L-ON strains (Fig. 7B, 60 kDa). The FimA levels of CFT073 and the complemented $\Delta mutS$, $\Delta mutH$, $\Delta mutH$ “L-ON”, and $\Delta mutL$ strains were much lower than the FimA levels of CFT073 *fim* L-ON and the complemented strain $\Delta mutS$ L-ON (Fig. 7B, 15 kDa). We conclude that the motilities of MMR deletion strains can be reduced to wild-type CFT073 levels by complementation, with the exception of $\Delta mutH$ “L-ON,” whose ability to unlock its type 1 fimbrial promoter has made full complementation a challenge.

The $\Delta mutS$ strain outcompetes the wild-type strain in the bladder and spleen during a 48-h ascending UTI. Flagella and type 1 fimbriae are known to be important in the colonization of the murine urinary tract by UPEC (19, 31, 47, 52). From our study, we have determined that deletions in the MMR system play a surprising role in the reciprocal control of these two functions. To identify any contribution of MMR *in vivo*, we assessed host colonization using the murine model of ascending UTI. Using the $\Delta mutS$ strain as a representative mutant of the MMR system, *in vivo* colonization of the $\Delta mutS$ strain was measured in both a 48-h and 7-day cochallenge of the mutant with wild-type CFT073 (Fig. 8A and B). The $\Delta mutS$ mutant outcompeted wild-type CFT073 in both the bladder ($P = 0.040$) and the spleen ($P < 0.001$) during the 48-h infection (Fig. 8A). The ability to disseminate to the bloodstream and reach the spleen can possibly be attributed to the increase in motility of the $\Delta mutS$ mutant bacteria. A cochallenge of $\Delta mutS$ L-ON with CFT073 *fim* L-ON was also tested, and we found that the MMR mutant had a fitness defect, where the parental strain outcompeted the mutant in the bladder during the 48-h infection ($P = 0.011$) (Fig. 8C). Differences noted were subtle but nonetheless statistically significant. We also measured the fitness of the constructs over a longer period. No significant differences were detected from the 7-day infection of either of the cochallenges (Fig. 8B and D).

The loss of MMR does not cause mutations in the *flhDC* or *pap* operon promoters. To identify a mechanism of MMR’s involvement in the regulation of motility and adherence, we examined the most likely candidate genes that might affect these functions. The first of these was the promoter region for class I flagellar genes *flhDC*, which initiates the complex cascade of gene expression leading to flagellar expression (25). The second was the promoter region for the *pap* (pyelonephritis-associated pili) genes, which encode P fimbriae (7, 50). P fimbriae have also been shown

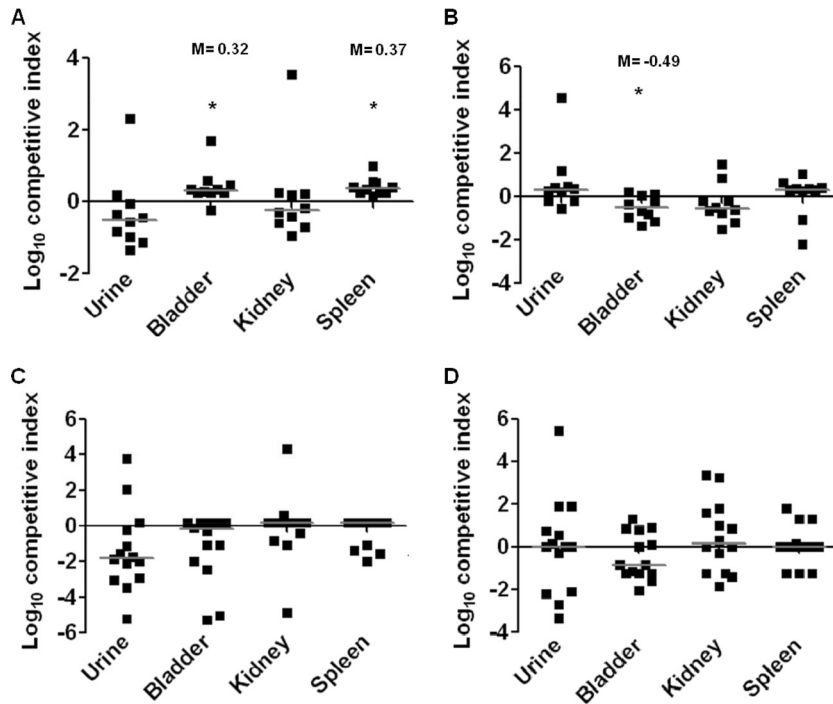


FIG 8 Mice were infected transurethrally with $\sim 10^8$ CFU of a 1:1 mixture of a $\Delta mutS$ mutant with its appropriate parental strain. After a 48 h or 7 days postinfection, urine, bladders, kidneys, and spleens were quantitatively cultured. Each symbol represents a datum from an individual mouse. Horizontal bars represent the median from each tissue. The limit of detection is 100 CFU/ml of urine or g of tissue. Significant differences in strain isolates cultured from organs were determined using a Wilcoxon signed-rank test. *, $P < 0.05$. (A) Cochallenge at 48 h with a 1:1 mix of CFT073 wild-type and $\Delta mutS$ strains in a wild-type background ($n = 10$). (B) Cochallenge at 7 days with a 1:1 mix of CFT073 wild-type and $\Delta mutS$ strains in a wild-type background ($n = 14$). (C) Cochallenge at 48 h with a 1:1 mix of CFT073 *fim* L-ON and $\Delta mutS$ *fim* L-ON ($n = 10$). (D) Cochallenge at 7 days with a 1:1 mix of CFT073 *fim* L-ON and $\Delta mutS$ *fim* L-ON ($n = 14$).

to play a role in the reciprocal control of motility and adherence and are regulated by methylation of GATC sites in the *pap* promoter (7, 50). Primers were used to amplify the promoter regions of the MMR deletion strains in both *fim* L-ON and wild-type backgrounds (Table 1). The resulting PCR fragments were purified and sequenced for all bacterial constructs used in the present study. Sequences of the *flhDC* and *pap* promoters of all constructs were identical to the wild-type CFT073 sequence and thus cannot explain the observed increase in motility in the absence of MMR (data not shown).

DISCUSSION

In our previous study, we observed that motility was repressed when type 1 fimbrial genes (*fim*) were constitutively expressed in UPEC strain CFT073 (30). When the entire *fim* region was deleted in the CFT073 *fim* L-ON mutant, only a partial restoration of motility to wild-type levels was observed (30). This suggested that the repression of UPEC motility was not solely based on the expression or assembly of these surface organelles and additional factors must be considered. A screen of 12,000 transposon mutants in CFT073 *fim* L-ON identified six nonfimbrial genes that reversed the repression of motility (49). This suggested that these six genes play a role in the reciprocal control of adherence and motility. Among these six genes was *mutS*, which encodes a component of the mismatch repair (MMR) system. In the present study, we demonstrate that the MMR system itself, including MutS, MutH, and MutL, plays a role in the reciprocal regulation of motility and adherence in UPEC.

Deletion of the MMR genes *mutS*, *mutH*, and *mutL* significantly increased motility in both CFT073 wild-type and *fim* L-ON backgrounds (with the exception of strain $\Delta mutL$ L-ON, in which no increase in motility was observed). Along with motility, the FimA and FliC levels, in whole cells and expressed on the bacterial surface, were analyzed using Western blots and immunofluorescence, respectively. These assays revealed that the levels of the two proteins correlate well with motility; expressing more flagellin results in more motility and more FimA production occurs when motility is reduced: qPCR was used to confirm and quantify these observations. Motility, the levels of FimA, and the levels of flagellin, could be fully complemented for all strains except for $\Delta mutH$ "L-ON"; this exception was due to the unique ability of this strain to unlock the *fim* promoter in the *fim* L-ON background of CFT073.

The *fim* promoter, located on an invertible element, was analyzed to determine its orientation in the six MMR mutants. We found that the *fim* promoters of more motile strains were primarily in the phase OFF orientation. This could suggest that in the absence of MMR, the rate of recombination at the *fim* promoter repeats increases and thus, ON-OFF or OFF-ON switching is increased. Keeping in mind the theory of reciprocal control of adherence and motility, increasing the frequency at which recombination of the *fim* promoter occurs, switching from the phase ON orientation to the phase OFF orientation, results in less type 1 fimbriae being synthesized and, consequently, allows for increased motility. To further support this theory, the motility of a

mixed population of CFT073 *fim* L-ON and CFT073 *fim* L-OFF was assessed and revealed a strong, positive correlation ($r^2 = 0.74$) between the amount of CFT073 *fim* L-OFF in a culture and the diameter of swimming motility. These findings support our capillary assay data that demonstrate an increase in the percentage of motile bacteria within a MMR mutant population that also exhibit an increased swimming diameter on soft agar plates.

As a representative of MMR, the $\Delta mutS$ mutation was used to assess *in vivo* fitness during a cochallenge with wild-type CFT073 using the murine model of ascending UTI to determine whether the MMR effect on motility affects UPEC pathogenesis. We found that wild-type CFT073 was outcompeted by bacteria with *mutS* deleted in both the bladder and the spleen 48 h postinoculation. This relationship between MMR and UPEC pathogenesis may be supported, in part, by evidence that UTI isolates exhibit a higher occurrence of mutator strains than commensal *E. coli* or any other *E. coli* pathotype (14). This is remarkable given that >1% of food-borne *E. coli* pathogens exhibit high mutation rates as a result of defective MMR systems (33). Defects in the MMR system have also been identified and linked to virulent isolates of other pathogenic bacteria, including *Salmonella enterica* serovar Typhimurium (33), *Neisseria meningitidis* (45) and *Pseudomonas aeruginosa* (41). Overall, these associations are correlative, and some caution should be used in their interpretation.

To explain how MMR deficiencies could lead to increased virulence, we must explain how MMR assists in avoiding genetic change. It does this by two main pathways: first, by limiting genetic variation through the correction of errors in newly replicated DNA strands (23, 24, 39) and, second, by acting as a barrier between interspecies gene exchange (11, 16, 36, 44). Therefore, when an organism exhibits defects in its MMR system, it is more prone to genetic variations and interspecies horizontal gene transfer and, consequently, has an increased capacity to adapt to the host environment or acquire new virulence genes, respectively (11, 16, 36, 44).

The mutator phenotype allows bacteria to rapidly respond via genetic change when faced with challenges, such as the presence of antibiotics, attack from host immune cells, altered nutritional availability, and competition in the environment. In the urinary tract, an organism is exposed to many environmental changes as it ascends from the urethral opening, to the bladder, up the ureters, to the kidneys and eventually, if left untreated, to the bloodstream. This includes the exposure to iron-limited urine, neutrophil influx, diverse antibacterial defense molecules, and host cell types. The ability to survive such changes is essential for the organism's ability to successfully colonize the host. This survival strategy is consistent with our finding that the loss of MMR increases fitness during ascending UTI. It should be noted, however, that the motility phenotypes of MMR mutant UPEC cannot be directly explained by inheritable genetic mutations, other than the loss of MMR genes, since the mutant phenotypes can be restored to wild type by complementation.

In summary, our findings show a direct relationship between methyl-directed MMR systems and the reciprocal control of motility and adherence in UPEC. Deletion of three genes involved in the MMR system—*mutS*, *mutH*, and *mutL*—increased motility in wild-type CFT073 and in CFT073 *fim* L-ON, with the latter constitutively expressing type 1 fimbriae. A mechanism that satisfactorily explains this association remains to be determined; however, our study strengthens the connection between mutator

isolates and virulence and suggests that MMR's effect on reciprocal control of motility and adherence contributes to UPEC pathogenesis.

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