Phenotypic Heterogeneity Enables Uropathogenic *Escherichia coli* To Evade Killing by Antibiotics and Serum Complement

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Uropathogenic strains of *Escherichia coli* (UPEC) are the major cause of bacteremic urinary tract infections. Survival in the bloodstream is associated with different mechanisms that help to resist serum complement-mediated killing. While the phenotypic heterogeneity of bacteria has been shown to influence antibiotic tolerance, the possibility that it makes cells refractory to killing by the immune system has not been experimentally tested. In the present study we sought to determine whether the heterogeneity of bacterial cultures is relevant to bacterial targeting by the serum complement system. We monitored cell divisions in the UPEC strain CFT073 with fluorescent reporter protein. Stationary-phase cells were incubated in active or heat-inactivated human serum in the presence or absence of different antibiotics (ampicillin, norfloxacin, and amikacin), and cell division and complement protein C3 binding were measured by flow cytometry and immunofluorescence microscopy. Heterogeneity in the doubling times of CFT073 cells in serum enabled three phenotypically different subpopulations to be distinguished, all of them being recognized by the C3 component of the complement system. The population of rapidly growing cells resists serum complement-mediated lysis. The dominant subpopulation of cells with intermediate growth rate is susceptible to serum. The third population, which does not resume growth upon dilution from stationary phase, is simultaneously protected from serum complement and antibiotics.

The immune system has several pathways for recognizing and killing pathogenic bacteria. However, some pathogenic bacteria can maintain infection in mammalian hosts despite inflammation, specific antimicrobial mechanisms, and a robust adaptive immune response and can therefore give rise to persistent infection (1). Uropathogenic *Escherichia coli* (UPEC) causes recurrent urinary tract infections that can progress from the lower to the upper urinary tract and can lead to the dissemination of bacteria into the bloodstream.

The complement system is part of the defense against invading pathogens, with an essential role in both innate and adaptive immunity (2). It is composed of more than 40 plasma and membrane proteins. It can be activated via three distinct routes: the classical (antibody dependent), lectin, and alternative pathways. Activation of complement cascades leads to the formation of the key component C3b on the bacterial surface, which stimulates phagocytosis. Late complement components (C5 to C9 proteins) are also activated via C3b, resulting in the formation of the membrane attack complex (MAC) causing cell lysis (2). The primary source of complement is blood, but complement proteins are also synthesized by a variety of other cell types and tissues (3).

Bacterial resistance to serum complement killing depends on the presence or absence of antigenic outer membrane proteins (4). Pathogens often resist recognition and subsequent complement activation owing to their surface capsular polysaccharide, which masks underlying structures and by itself activates complement poorly (5–7). In addition, secretion of the exopolysaccharide colanic acid protects UPEC from complement-mediated killing (8). Modification of lipopolysaccharide (LPS) is also important for complement evasion (9, 10). It has been shown that serum sensitivity depends on the bacterial growth phase; cells are more readily killed by serum during early logarithmic phase (11–14). However, there are examples showing that exponential-phase cells are more resistant to complement-mediated killing than stationary-phase cells (15, 16). That phenomenon is largely explained by the growth-phase-dependent expression of antigens, capsule, and LPS modification (14, 15). Cell size can also be an important determinant of complement-mediated killing, since larger or aggregated cells have more complement protein C3b on their membranes (17).

Some of the mechanisms that contribute to immune evasion can also help the bacteria survive antibiotic treatment. For example, the inability of many antibiotics to cross host membranes readily limits their effectiveness against intracellular bacteria (18). Also, changes in bacterial outer membrane composition can simultaneously influence the binding of complement proteins and the influx of antibiotics (4). Importantly, antibiotic-tolerant bacteria are not able to grow in the presence of drugs, but once the antibiotic concentration drops, surviving persister cells restore the population (19). Two clinical studies have demonstrated that patients with long-term infections had pathogen strains with elevated persister levels, indicating that persisters in a bacterial population might account for the failure of antibiotic treatment in chronic infections (20, 21). These studies investigated cystic fibrosis patients infected with *Pseudomonas aeruginosa* (20) and cancer
patients with oral thrush caused by *Candida albicans* (21). In UPEC, it has been previously shown that recurrent community-acquired urinary tract infections are mainly due to persisting UPEC clones (22) and the presence of intracellular bladder reservoirs for UPEC in humans (23). Furthermore, the persistence of UPEC in bladder has been shown in a mouse urinary tract model, where the combined effect of antibiotics and immune system was investigated (18).

The question remains: why are these antibiotic persisters not killed by the immune system? It is generally believed that the immune system kills planktonic persisters, but biofilm persisters cells are protected from the host defenses by the exopolymer matrix (24). It has also been proposed that dormant intracellular bacteria are simultaneously protected from antibiotics and the immune system (18). While the phenotypic heterogeneity of bacteria influences antibiotic tolerance, the possibility that it also makes cells refractory to the immune system has not been considered. We therefore sought to determine whether the heterogeneity of bacterial cultures manifested by the presence of persisters also influences bacterial targeting by the serum complement system. In addition, we determined whether the number of persisters is reduced by antibiotic treatment in human serum.

We demonstrate here that the complement system kills most dividing cells, but minor subpopulations of dormant and rapidly dividing cells evade complement killing. Importantly, stationary-phase dormancy protects intrinsically susceptible UPEC cells from killing by both antibiotics and the serum complement system.

**MATERIALS AND METHODS**

**Bacterial strains, plasmids, and growth media.** *E. coli* strains CFT073 (25) and BW25113 [F^−^ (Δ araBAD-araB)567 Δ leuZ4787 (::rrnB-3) X rph-1 Δ (ruaD-ruaB)568 hsdR514] (26) were used in the experiments. To study cell division, the CFT073 strain was transformed with the plasmid pET-gfp-mut2AGGAGG(3) carrying a green fluorescent protein (GFP) *mut2* gene under the control of an IPTG (isopropyl-β-d-thiogalactopyranoside)-inducible promoter and a kanamycin resistance marker (27). For plasmid selection, kanamycin (25 μg/ml; Amresco) was added to the growth medium. To induce GFP, 1 mM IPTG was added when indicated. To test antibiotic sensitivity, ampicillin sodium (Sandoz), norfloxacin (Sigma), or amikacin (Sigma) was prepared as an aqueous solution and used at the indicated concentrations. To inhibit cell growth the bacteriostatic antibiotics, ampicillin, norfloxacin, and amikacin for CFT073 strain were determined in 96-well microtiter plate. Growth was assessed after incubation for 18 h, and the MIC value was determined.

**Serum and antibiotic sensitivity.** Cells from DMSO stock were diluted 1:100 in filtered LB medium and grown aerobically to the stationary phase (20 h). Cells were harvested, washed, and diluted in 1× PBS or LB medium so that the final OD_{600} in the assays was 0.05. The final percentage of serum or HIS was 50% in PBS. Antibiotics were added at the indicated concentrations at time point zero. The cells were incubated at 37°C without shaking. Samples were withdrawn at the indicated times, diluted in sterile 1× PBS, and plated on LB plates. Colonies were counted on the following day after overnight incubation at 37°C.

**Cell division assessed by flow cytometry.** To measure cell division, the CFT073 strain carrying GFP encoding plasmid pETgfp-mut2AGGAGG(3) was used. Experiments were carried out as described above for the serum and antibiotic sensitivity assay, except that the overnight cultures were supplemented with kanamycin (25 μg/ml) and 1 mM IPTG to induce GFP expression. Samples were mixed with an equal volume of 30% glycerol in 1× PBS and frozen at −80°C until further analysis with an LSR II flow cytometer (BD Biosciences). Populations of bacterial cells were gated on GFP fluorescence (GFP-H) and side scatter (SSC-H) plots. GFP was excited with a 488-nm laser, and emission was measured using a 530/30-nm band-pass filter. The software package Flowing Software was used to visualize and analyze the data. The distribution of the fluorescence level of events (single cells) is presented as a histogram consisting of 376 repartition bins. The number of events with the respective GFP fluorescence levels in 1 ml of cell culture grown under particular conditions is shown.

**SEM of bacteria with or without serum treatment.** To investigate the morphology of intact and lysed bacteria, scanning electron microscopy (SEM) experiments were performed. The CFT073 strain carrying plasmid pETgfp-mut2AGGAGG(3) was used. Experiments were carried out as described above for the serum and antibiotic sensitivity assay, except that the overnight cultures were supplemented with kanamycin (25 μg/ml) and 1 mM IPTG to induce GFP expression. Bacterial suspensions (100 μl) were withdrawn at the indicated time points, and slides were prepared with Cytospin 4 (Thermo Scientific, USA). After a cytopsinning step at 2,000 rpm for 3 min, the bacteria were fixed with freshly prepared 4% paraformaldehyde for 30 min at room temperature. All samples were washed three times with 1× PBS, dehydrated with a series of graded ethanol solutions, and dried in vacuum oven for 30 min. Samples were sputtered with 5 nm of gold using a high vacuum coater (EM ACE600; Leica, Hitachi, Japan).

**Immunofluorescent detection of serum complement protein C3 binding.** To identify complement C3b deposition, the same experiments and sample preparation methods were used as for SEM analysis. However, after fixation with 4% paraformaldehyde, the slides were washed with 1× PBS for 5 min and blocked in PBS containing 2% bovine serum albumin for 15 min at room temperature. The samples were then incubated with 1:100-diluted mouse monoclonal complement C3b antibody (6C9; Thermo Scientific) in a blocking buffer for 1 h at room temperature in closed containers. After three washes with 1× PBS, the slides were incubated with secondary goat anti-mouse IgG antibodies labeled with red-fluorescent Alexa Fluor 568 dye (1:1,000 dilution in blocking buffer) for 1 h at room temperature in closed containers. Finally, the slides were washed three times with 1× PBS, moistened with 3 μl of 1× PBS, covered with slips, and sealed with nail polish. Samples were stored in covered closed containers at 4°C.

**Fluorescence was observed, and images were acquired using the confocal fluorescence microscope LSM710 (Carl Zeiss, Munich, Germany) and Zen software (Zeiss). To detect GFP expression, a 488-nm laser and a 484- to 571-nm emission filter were used. To detect Alexa Fluor 568 (C3 signal) excitation, a 561-nm laser and a 575- to 708-nm emission filter were used. The image resolution was 52.72 nm per pixel.**

Photographed bacterial cells were examined using Cell Profiler (GPL,
The effect of serum on the bacteria was triphasic. The reaction started with a lag phase for 60 min during which the bacteria were not killed. The cells were then killed efficiently between 60 and 90 min and then at a slower rate between 120 and 150 min (Fig. 1). This raises the question of whether surviving cells are really more tolerant to serum killing or whether the complement proteins are exhausted and some cells survive only because there is not enough complement protein to kill all of the susceptible cells. To address this, we repeated our experiment with a 10-fold-lower bacterial CFU count and observed enhanced, but similarly reduced, killing at between 120 and 150 min (see Fig. S1A and B in the supplemental material). Furthermore, when serum-treated samples were supplemented with additional fresh serum after 90 min, the triphasic killing pattern remained (see Fig. S1A and B in the supplemental material). In addition, the nonpathogenic laboratory E. coli strain BW25113 (stationary-phase cells at a concentration of \(10^7\) CFU/ml) was readily killed (the CFU level was reduced by 6 orders of magnitude) by 20% serum over 30 min. These results suggest that in our experimental setup, the amount of serum protein per bacterial cell is somewhat limiting, but the CFT073 strain has a subpopulation of cells with enhanced serum tolerance.

Our results show that the CFT073 strain tolerates serum since the cells are not killed during lag phase. However, it is not strictly serum resistant because the cells can be killed during active growth. At the same time, a subpopulation of the cells is tolerant to serum killing.

The combined effect of antibiotics and serum depends on the antibiotic. The existence of a serum-tolerant subpopulation raises the questions: (i) can the complement system kill antibiotic persisters, and (ii) do antibiotics eradicate serum persisters? To answer these questions, we counted the surviving cells after 150 min of incubation in the presence or absence of active serum in combination with the cell wall-targeting antibiotic ampicillin, the DNA replication-inhibiting antibiotic norfloxacin, and the translation-inhibiting antibiotic amikacin. These antibiotics were chosen for analysis, since they are all bactericidal and represent different antibiotic classes. Furthermore, these antibiotics have been used in several important studies investigating persisters and persistent infections (34–38). The MICs for CFT073 strain in LB medium were as follows: amikacin, 8 µg/ml; ampicillin, 2 µg/ml; and norfloxacin, 0.125 µg/ml. Figure 2 shows the survival of bacterial cells at different concentrations of each antibiotic. The addition of ampicillin to the serum reduced the number of surviving cells by about 1 order of magnitude (Fig. 2A). This suggests that some serum-tolerant cells are killed by ampicillin. An additive effect of serum to ampicillin is apparent when the numbers of persisters in HIS and serum are compared, indicating that some ampicillin persisters are killed by serum (Fig. 2A). The bacteria are killed more efficiently in LB medium than in HIS (Fig. 2A). Growth resumption rates in different growth media affect the number of ampicillin persisters (39). Therefore, we can assume that the slower growth resumption in HIS compared to LB medium could result in more ampicillin persisters.

Next, we measured the number of persisters after we incubated the cells with norfloxacin (Fig. 2B). The number of norfloxacin persisters was similar under all growth-promoting conditions: LB medium, HIS, and serum (Fig. 2B). This finding suggests that the same subpopulation survives treatment by both serum and norfloxacin.
Finally, we measured the number of amikacin persisters in different media (Fig. 2C). Surprisingly, in HIS and in serum the number of CFU started to decrease at lower amikacin concentrations than in LB medium (Fig. 2C). This indicates that both serum and HIS sensitize the bacteria to amikacin. At high amikacin concentrations, the numbers of surviving bacteria are similar in HIS and serum, suggesting that bacteria surviving the antibiotic treatment are also refractory to serum. We also confirmed that the antibiotics do not kill nongrowing cells since bacteria incubated in PBS were refractory to ampicillin, norfloxacin, and amikacin (Fig. 2A to C).

Serum preferentially kills cells with average growth rates. The growth of bacteria in HIS demonstrated that this serum serves as a growth substrate (Fig. 1). Active serum has both killing- and growth-promoting effects, which are indistinguishable in experiments where only CFU counts are determined. The final CFU count in serum is the outcome of both the increase of CFU due to cell division and the decrease due to killing by complement. Therefore, we investigated the causes of the final CFU count to determine whether a random population survived serum treatment or whether these were nondividing cells as in the case of antibiotic persisters. We used CFT073 cells carrying plasmid pETgfp-mut2AGGAGG(3), which encodes GFP under the control of the IPTG-inducible promoter. This strain allowed us to monitor cell division using a previously established GFP dilution method (39, 40). The induction of GFP production using IPTG was performed only during the initial growth to the stationary phase.

Stationary-phase cells carrying pETgfp-mut2AGGAGG(3) and filled with GFP were diluted in fresh IPTG-free medium and grown as in the previous experiment. The cells were analyzed at the beginning of the experiment, which allowed us to determine the initial GFP and side scatter (SSC) levels to gate the cells before they started dividing (P1 on Fig. 3A to E). When cells were kept in PBS for 150 min, most of the events identified by flow cytometry had similar high green fluorescence levels and low SSC, as at the zero time point (Fig. 3A and B). Flow cytometry measurements of cells grown in LB medium for 150 min showed high-SSC and low-green-fluorescence (GFP-H) events, indicating that most of the cells grown in LB medium had divided (Fig. 3C). These measurements allowed us to set the second region, P3, for actively dividing cells. Flow cytometry measurements of cells grown in 50% HIS showed somewhat less division than in LB medium, and the extent of SSC was lower (Fig. 3D). Most cells grown in HIS had GFP and SSC values intermediate between those of the nondividing (P1) and actively dividing cells (P3) and were in the region P2 (Fig. 3D). Samples treated with serum showed events in regions P1 and P3; events with very low GFP fluorescence and low SSC also appeared, most likely representing dead cells and cell debris (Fig. 3E). The combined number of events detected by flow cytometry was comparable with CFU in all samples, indicating that all three regions (P1, P2, and P3) contain live cells (see Fig. S2 in the supplemental material).

FIG 2 Subpopulation of cells surviving antibiotics and complement killing. Cells were grown to stationary phase in LB medium and diluted in fresh LB medium or PBS supplemented or not with HIS or serum (final concentration, 50%). The number of surviving cells (CFU/ml) was determined after 150 min by plating. The averages and standard deviations of at least three independent experiments are presented.
integrity and were dented; (ii) cells that had lost their normal shape and were ruptured; and (iii) cells that were misshaped and completely lysed (Fig. 3H). These modified and lysed cells were not observed in HIS samples (Fig. 3G).

Nondividing cells preferentially survive combined treatment with serum and antibiotics. We also analyzed the cell division profile after antibiotic treatment. Stationary-phase cells carrying the pETgfp-mut2AGGAGG(3) and filled with GFP were diluted in fresh IPTG-free medium and incubated in the presence or absence of antibiotics for 150 min. Antibiotics were used at the following concentrations: ampicillin, 200 μg/ml; norfloxacin, 5 μg/ml; and amikacin, 5 μg/ml. After the incubation, the numbers and fluorescence levels of cells were determined by flow cytometry, and the CFU counts were determined by plating.

First, we inspected cells incubated in PBS. As expected, these cells were not dividing, and the addition of antibiotics did not

FIG 3 A bacterial population that survives serum treatment is enriched in rapidly dividing and nondividing cells. Stationary-phase CFT073 cells [carrying plasmid pETgfp-mut2AGGAGG(3)] were diluted in PBS, LB medium, 50% HIS, or 50% serum. The cells were incubated at 37°C without shaking for 150 min, and their fluorescence was measured by using flow cytometry. Density plots of GFP fluorescence (GFP-H) and the SSC parameter (SSC-H) are shown for cells in PBS at zero time (A), in PBS after 150 min (B), in LB medium (C), in HIS (D), or in serum (E). Each dot represents the fluorescence of a single event (particle). (F) The distribution of the fluorescence level of events (single cells) analyzed by flow cytometry is presented as a histogram consisting of 376 repartition bins. The number of events with the respective GFP fluorescence levels in 1-ml cell cultures grown under particular conditions is shown. (G and H) Scanning electron micrographs illustrate the morphology of the *E. coli* cells after 150 min of incubation in 50% HIS (G) and in 50% serum (H). Arrows indicate the different bacterial cells during lysis.
change the fluorescence level distribution or the number of events per volume of sample (see Fig. S3A in the supplemental material). The CFU counts and the numbers of GFP-positive cells (as determined by flow cytometry) were approximately equal in PBS in the presence or absence of antibiotics, indicating that all cells were viable and able to form colonies after the antibiotics were removed (see Fig. S3A and B in the supplemental material).

Next, we inspected the division profile of cells incubated in growth-supporting media. In HIS there was a decrease of ∼2 orders of magnitude in the number of dividing cells (events with low GFP fluorescence) when antibiotics were added (Fig. 4A). Ampicillin reduced the number of dividing cells in HIS, indicating that these cells have been lysed (Fig. 4A), although the number of non-dividing (high GFP content) cells was not affected (Fig. 4A). In contrast to the ampicillin-treated cultures, the number of cells with high GFP content was increased after incubation in HIS, together with amikacin or norfloxacin (Fig. 4A). This indicates that norfloxacin and amikacin inhibited growth resumption.

Treatment with a bactericidal antibiotic in growth-supporting media has been shown to leave a large fraction of cells in a non-recoverable state; therefore, the number of events detected by flow cytometry can exceed the number of persister cells that form colonies after plating on a solid medium (41). In the case of norfloxacin, the ratio of CFU to GFP-positive events was almost 1 in HIS (Fig. 4B). We can therefore say that norfloxacin inhibited cell growth, but the cells retained their ability to form colonies. When they were incubated in the presence of amikacin in HIS, there was a difference of >2 orders of magnitude between CFU and GFP-positive events detected by flow cytometry (Fig. 4B). This means that amikacin killed most of the cells in HIS, but these were still visible by flow cytometry as cells containing high levels of GFP because they were not lysed as they were after ampicillin treatment (Fig. 4A). The division profiles of cells incubated with antibiotics in LB medium were similar to those in HIS (see Fig. S3C and D in the supplemental material).

Finally, the cell division profile in serum was determined. In
serum alone there were two surviving subpopulations: nondividing and rapidly dividing cells (Fig. 3E and 4C). When serum and antibiotic treatment were applied simultaneously, the rapidly dividing subpopulation disappeared while the nondividing subpopulation survived (Fig. 4C). In serum, the number of nondividing cells was increased after treatment with norfloxacin or amikacin (Fig. 4C). Importantly, comparison of CFU and events detected by flow cytometry shows that in the presence of amikacin most of the cells detected by flow cytometry are not viable (Fig. 4D). In contrast, the nondividing cells induced by norfloxacin remained viable and were also refractory to serum (Fig. 4C and D). This result indicates that growth inhibition by an antibiotic can make bacteria resistant to lysis by serum. We conclude that the nondividing antibiotic persisters are not cleared by the complement system as efficiently as growing cells.

Differences in C3 deposition are not responsible for selective lysis of bacteria. From the antibiotic and serum cotreatment it became evident that antibiotic persisters are not efficiently killed by human serum complement. It is not clear whether bacterial cells are refractory to complement killing because they are not recognized and subsequently opsonized, or whether they avoid complement-mediated lysis. In order to determine whether serum complement recognizes all of the CFT073 strain cells equally, C3 deposition was measured using sequential C3 antibody staining with Alexa 568 and immunofluorescence microscopy.

First, C3 deposition on CFT073 cells in serum, HIS, and PBS without antibiotics was investigated. Complement C3 deposition was observed on cells incubated in human serum after only 30 min compared to the cells incubated in HIS (data not shown). In order to discriminate between dividing and nondividing subpopulations, a 150-min time point was chosen for further analysis. The presence of GFP inside the intact cells enabled us to visualize nondividing cells with high GFP intensity, dividing cells with low GFP intensity, and lysed cells with no detectable GFP (Fig. 5A). In negative controls, cells incubated in HIS, or cells incubated in PBS, no specific C3 binding was detected (Fig. 5B), and the relative fluorescence of Alexa 568 was less than 0.05 U, which was taken as the threshold for determining C3-binding cells. After 150 min, almost all GFP-containing cells (99%) were recognized by human serum complement, and C3 protein was bound to them (Fig. 5A and B). In addition, C3-bound cells or cell membranes with no detectable GFP fluorescence were observed (Fig. 5A). This is consistent with CFU counts showing that most of the cells were lysed by serum (Fig. 1). In order to determine whether this C3 deposition level was sufficient for killing, we compared the C3 levels of lysed (no GFP) and intact cells. Figure 5C shows that the Alexa 568 intensities of lysed and intact cells were comparable; hence, unequal C3 deposition is not responsible for selective killing.

Next, we analyzed the C3 deposition on cells incubated in human serum in the presence of antibiotics. Intense C3 deposition was detected on cells incubated in serum in the presence of ampicillin, norfloxacin, or amikacin (Fig. 5A). Flow cytometry had revealed that in the presence of ampicillin, growing cells (expressing low levels of GFP) disappeared (Fig. 4C). As expected, we found no low-GFP-level cells in ampicillin-treated samples. Instead, the number of lysed cells (manifested by the absence of GFP) had increased (Fig. 5A). The opposite effect was seen in norfloxacin- and amikacin-supplemented serum cultures: fewer lysed cells were observed (Fig. 5A). These results indicate that C3 binding is not enough for serum-dependent cell lysis. Even cells that are killed by amikacin remain intact (retain high GFP fluorescence) despite the presence of C3 on the surface (Fig. 4C and 5A).

Exponentially growing cells are susceptible to serum and require active protein synthesis for survival. We observed that growth resumption coincides with serum-mediated killing (Fig. 1). To investigate how growth makes cells susceptible to complement-mediated lysis, the survival of exponential-phase cells was tested. We used the GFP dilution method as described in previous sections to follow cell division and the number of surviving cells simultaneously. Flow cytometry showed that after 60 min the cells in serum had divided as in HIS, but there were only half the number of cells (Fig. 6A). This indicated that some cells were lysed during cell division. The difference in the number of cells in HIS and serum was even more pronounced after 120 min; ca. 75% of cells were lysed in serum (Fig. 6A). Importantly, cell division profiles indicated that most cells surviving in serum had divided more actively than the population average in HIS (Fig. 6A).

Next, we used a bacteriostatic antibiotic, chloramphenicol, to inhibit growth and division without killing the cells. Parallel experiments were performed with stationary- and exponential-phase cells. Stationary-phase E.coli was transferred to fresh medium and chloramphenicol added at time point zero. After 120 min, the CFU were counted. As expected, the addition of chloramphenicol to HIS inhibited the growth of stationary-phase cells but did not kill them (Fig. 6B). Inhibition of cell growth with chloramphenicol slightly increased the number of cells surviving in the presence of serum (Fig. 6B). However, the effect of chloramphenicol on stationary-phase cells was not statistically significant (Fig. 6B). Next, we repeated the experiment with exponential-phase cells. Here, a 60-min time point was chosen because there is no lag phase, and the cells start to divide rapidly. Addition of chloramphenicol to exponential-phase cells growing in HIS stopped cell growth efficiently and even reduced the CFU count (Fig. 6B). When chloramphenicol was added to exponential-phase cells in serum, there was a decrease of >2 orders of magnitude in the number of CFU (Fig. 6B). This suggests that exponential-phase cells need active protein synthesis to counteract serum-mediated killing. The survival of nondividing cells in the presence of serum is characteristic only of stationary-phase persisters. Inhibition of cell growth in exponential-phase does not rescue them from complement-mediated lysis (Fig. 6B).

DISCUSSION

In nature, bacteria are constantly confronted with changes in nutrient availability and the presence of antimicrobials or other damaging agents. Stochastic differentiation of a genetically identical population into subpopulations with distinct phenotypes provides a strong advantage in an unpredictable and fluctuating environment (42, 43). One example of phenotypically different subpopulation is the formation of persisters, i.e., cells that survive treatment with bactericidal antibiotics (36, 44).

It is clear that during antibiotic treatment the immune system contributes heavily to the outcome of therapy. Therefore, in order to persist, bacteria have to evade killing by both antibiotics and the immune system. It has been proposed that biofilm-embedded persisters are the ones that survive antibiotic treatment, but planktonic persisters are killed by the immune system (24). We sought to determine whether the heterogeneity of cultures affects bacterial targeting by the immune system and, if so, what happens to the...
persisters when antibiotic treatment takes place in the presence of the serum complement system.

First, we determined the serum susceptibility of E. coli strain CFT073. Our results show that CFT073 is not strictly serum resistant as previously reported (6, 8, 9, 31, 32). Rather, only a fraction of cells survived serum treatment (Fig. 1 and Fig. 3H; see also Fig. S1 in the supplemental material). We observed that two subpopulations survived in serum, in contrast to cells grown in HIS: cells with low GFP levels (growing most actively) and cells with high GFP levels (dormant) (Fig. 3D and E). It is important to note that most cells, with moderate GFP levels, were lysed by serum.

To test the hypothesis that active growth makes cells more vulnerable to serum, the survival of exponential-phase cells was tested. CFU counting and flow cytometry revealed the number of GFP-positive cells in serum to be half that in HIS after 60 min (Fig. 6A and B). However, analysis of the GFP levels showed that the cells had divided as rapidly in serum as in HIS, indicating that most of them had been lysed during active growth (Fig. 6). It has been shown previously by Miajlovic et al. that when exponential-phase CFT073 cells were incubated in the presence of 50% serum in LB medium, the CFU count did not decrease over 180 min (8). In that study, ∼50 times more cells were incubated with serum than in our experiment (Fig. 6). We repeated the experiment with the same initial cell concentration as in the study by Miajlovic et al. (3 × 10⁸ CFU/ml), and no killing of cells by serum was detected (data not shown). This suggests that complement pro-

**FIG 5** Recognition and opsonization of CFT073 strain bacterial cells by serum complement. C3 deposition was measured by sequential C3 antibody staining and immunofluorescence microscopy. Cells were grown to stationary phase in LB medium and diluted in fresh medium (in PBS supplemented with either 50% HIS or 50% serum or in PBS alone) and incubated for 150 min at 37°C. (A) Immunofluorescence micrographs of GFP and C3-bound Alexa 568 fluorescence. Cells were grown in serum in the presence of ampicillin (200 μg/ml), norfloxacin (5 μg/ml), or amikacin (25 μg/ml), and C3 binding was detected by using immunofluorescence. From left to right is shown the GFP fluorescence inside the cells (visible GFP signal indicates intact cells) and the C3 deposition on cells (together with no GFP signal enables lysed cells to be identified); figure enlargements for GFP and C3 tile figures are also shown. Red dashed boxes show the exact locations of the figure enlargements on the tile figures. (B) Dot plot of C3-bound Alexa 568 and GFP fluorescence of cells incubated in HIS, PBS, or serum. Each dot represents the mean intensity (GFP and C3-bound Alexa 568) per cell measured. The dashed horizontal line indicates threshold value for cells with specifically bound C3. The dashed vertical line separates subpopulations of nondividing (high GFP levels) and dividing (low GFP levels) cells. (C) Relative distribution of cells by mean C3-bound Alexa 568 fluorescence. The objects examined were distinguished by the presence or absence of GFP. Objects with a mean GFP fluorescence >0.01 (includes all cells shown in panel B) were considered live (GFP-positive cells). Objects with a GFP fluorescence <0.01 were considered lysed.
teins can become limiting when large numbers of exponential-phase cells are treated with serum. However, clinical data from adult bacteremic patients indicated that bacterial cell numbers in blood are relatively low, typically fewer than 10 CFU/ml (45).

We also determined the serum tolerance of exponential-phase cells in the presence of the bacteriostatic antibiotic chloramphenicol. The addition of chloramphenicol to HIS efficiently stopped cell growth with >3-fold killing of the exponential-phase cells (Fig. 6B). Chloramphenicol increased the killing of the exponential-phase cells by serum by >2 orders of magnitude (Fig. 6B). Thus, the survival of nondonciding cells in the presence of serum is characteristic only of stationary-phase persisters, and inhibition of growth does not rescue exponential-phase cells from complement-mediated lysis. A transcriptome study of serum-incubated exponential-phase CFT073 cells demonstrated that many RcsB-regulated genes are induced (8). RcsB, either alone or in combination with other regulators, activates transcription of a wide range of genes, including those for colanic acid capsule synthesis, cell division, and some membrane proteins, and it regulates flagellum synthesis negatively (46). The importance of the RcsB regulon in serum survival has been confirmed by the observation that an rcsB mutant survives less well than wild-type CFT073 in serum (8). It has been shown that E. coli K1 cells can resist complement attack by OmpA-dependent binding of complement protein C4bp, which subsequently prevents MAC formation (15). Interestingly, exponential-phase E. coli K1 cells were more resistant to complement attack than stationary-phase cells, suggesting an additional role of growth-phase-dependent expression of the capsule and LPS (15). Our results, together with literature data, suggest that active adaptation mechanisms are needed for the survival of rapidly dividing (exponential phase) cells in serum.

Our first characterization of serum-surviving cells revealed both dormant and dividing cells. Since antibiotic persisters are usually nondonciding cells, they might not be killed by serum either. We measured the survival of CFT073 cells in the presence of different antibiotics in a standard laboratory medium, as well as in human serum.

Our results demonstrate an additive effect of serum to ampicillin if the numbers of persisters in HIS and serum are compared, indicating that some ampicillin persisters are killed by serum complement (Fig. 2A). However, norfloxacin persisters were not killed by serum complement (Fig. 2B). Amikacin had the strongest effect on the killing of bacteria in HIS and serum (Fig. 2). Nevertheless, at high (>2 μg/ml) amikacin concentrations the numbers of persisters were similar in serum and HIS, indicating that amikacin persisters were also not killed by serum complement (Fig. 2C). Thus, antibiotic persisters are considerably more resistant to serum complement than the rest of the population. Our study therefore reveals that planktonic antibiotic persisters are not as efficiently cleared by the immune system as the antibiotic-susceptible population. This is different from the current view that antibiotic persisters need to be embedded in biofilm in order to escape the immune system (47).

Several studies have shown that stationary-phase CFT073 cells are resistant to serum complement-mediated killing (6, 9, 31, 32). Importantly, there was no significant growth of cells in HIS during the treatment time (6, 31) (or the data on growth of bacteria in HIS were not presented [9, 32]). This suggests that stationary-phase dormancy protects CFT073 cells from serum complement-mediated killing. A study by Buckles et al. (6) demonstrated that most CFT073 wild-type cells were phagocytosed in the presence of human serum within 90 min. Thus, the results of Buckles et al. support our observation that CFT073 cells are not killed in the absence of growth but are marked with the C3 protein (Fig. 4C and Fig. 5).

The presence of C3 on the bacterial cell membrane and resistance to lysis might constitute an important mechanism for establishing infection. Some UPEC strains have been shown to use C3 opsonization to facilitate uptake by uroepithelial cells (48). Importantly, UPEC can also survive within mouse and human macrophages (49). It has been suggested that chronically infected epithelial cells could thus provide a constant source for reinfection of macrophages, which are unable to eliminate the bacteria completely, thus helping the bacteria to spread (49).

A simplified model based on our results is shown in Fig. 7. We
propose that heterogeneous growth resumption and different growth rates of CFT073 cells (in serum) incorporate three phenotypically different subpopulations. Rapidly growing cells are susceptible to antibiotics but resist serum complement-mediated lysis. These cells might be important during the acute phase of infection. Cells that do not resume growth are simultaneously protected from serum complement and antibiotics. They might be important in establishing persistent infection. The dominant subpopulation of cells, with intermediate growth rate, is susceptible to both antibiotics and serum. Although the medium growth-rate population is killed by the complement system, it might still have a role in establishing infection by synthesizing and secreting virulence factors. In Salmonella, the virulence factors needed for host manipulation are expressed in a bistable fashion, leading to a slowly growing subpopulation that expresses virulence genes and a rapidly growing subpopulation that is phenotypically avirulent.

Several studies have focused on the interplay between antibiotic and serum susceptibility. For example, it has been shown that overexpression of antibiotic efflux pump MexCD-OprJ reduces P. aeruginosa virulence by increasing its susceptibility to complement-mediated killing (51). At the same time, growth in serum has been shown to induce expression of antibiotic efflux pumps in P. aeruginosa and Acinetobacter, suggesting enhanced antibiotic tolerance during infection (52). Beside induction of efflux pumps, the growth in serum has been shown to enhance expression of OprD porin in P. aeruginosa (53). Recent study has shown the mutation in oprD gene in P. aeruginosa conferred resistance to carbapenem antibiotics, as well as enhanced resistance to killing by human serum (54). The relationship between antibiotic and serum sensitivity can be even more complex when population heterogeneity during infection is considered. Recent work on persistent Salmonella infection in a mouse model has revealed that internalization by macrophages can induce phenotypic heterogeneity, leading to either bacterial growth or the formation of non-dividing persisters that could provide a reservoir for relapsing infection (55). Another study has shown that nondividing Salmonella survived antibiotic treatment the best, but overall clearance of infection was delayed primarily by abundant subsets of moderately growing Salmonella with partial tolerance (56).

The complex phenotypic heterogeneity described in the present study could be regulated by molecular mechanisms, which could become new targets for treating acute and persistent infections caused by UPEC. Further work is needed to establish the potential role of the phenotypic heterogeneity of UPEC described here in in vivo infection.

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FIG 7 Model of phenotypic heterogeneity of UPEC cells during infection. Phenotypic heterogeneity leads to the formation of three different subpopulations. Dormant nondividing cells are simultaneously protected from serum and antibiotics. The dominant subpopulation of cells with an intermediate growth rate is susceptible to antibiotics and serum. Rapidly dividing cells are susceptible to antibiotics but can resist serum complement-mediated lysis. Bacteria are depicted as dark gray ovals, and lighter gray ovals indicate dilution of the color by cell division.


