

Impact of Education and Data Feedback on Guideline-Concordant Prescribing for Urinary Tract Infections in the Outpatient Setting

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Background. Urinary tract infections (UTIs) are the most common outpatient indication for antibiotics and an important target for antimicrobial stewardship (AS) activities. With The Joint Commission standards now requiring outpatient AS, data supporting effective strategies are needed.

Methods. We conducted a 2-phase, prospective, quasi-experimental study to estimate the effect of an outpatient AS intervention on guideline-concordant antibiotic prescribing in a primary care (PC) clinic and an urgent care (UC) clinic between August 2017 and July 2019. Phase 1 of the intervention included the development of clinic-specific antibiograms and UTI diagnosis and treatment guidelines, presented during educational sessions with clinic providers. Phase 2, consisting of routine clinic- and provider-specific feedback, began ~12 months after the initial education. The primary outcome was percentage of encounters with first- or second-line antibiotics prescribed according to clinic-specific guidelines and was assessed using an interrupted time series approach.

Results. Data were collected on 4724 distinct patients seen during 6318 UTI encounters. The percentage of guideline-concordant prescribing increased by 22% (95% CI, 12% to 32%) after Phase 1 education, but decreased by 0.5% every 2 weeks afterwards (95% CI, -0.9% to 0%). Following routine data feedback in Phase 2, guideline concordance stabilized, and significant further decline was not seen (-0.6%; 95% CI, -1.6% to 0.4%). This shift in prescribing patterns resulted in a 52% decrease in fluoroquinolone use.

Conclusions. Clinicians increased guideline-concordant prescribing, reduced UTI diagnoses, and limited use of high-collateral damage agents following this outpatient AS intervention. Routine data feedback was effective to maintain the response to the initial education.

Keywords. antibiotics; fluoroquinolones; outpatient antimicrobial stewardship; urinary tract infections.

Approximately 10% of adult ambulatory care visits conclude with an antibiotic prescription, 30% of which are unnecessary [1–3]. Of those needed, nearly 50% are prescribed inappropriately [2, 3]. While outpatient antimicrobial stewardship (AS) improves prescribing without adversely affecting patient outcomes, lack of incentives and resources limits the implementation of formalized programs [4, 5]. The Joint Commission (TJC) recently recognized outpatient AS as a patient safety priority and outlined new requirements for accredited ambulatory health care organizations [6]. These new standards will drive development of outpatient AS programs. Several national organizations have published best practices for implementing outpatient AS [2, 5, 7, 8].

With >8.6 million annual ambulatory care visits, urinary tract infections (UTIs) are the most commonly diagnosed outpatient bacterial infection and an important target for outpatient AS [5, 9]. Infectious Diseases Society of America (IDSA) guidelines emphasize fluoroquinolone-sparing treatment for uncomplicated UTIs due to increasing antibiotic resistance and known associated collateral damage [10]. Additionally, the Food and Drug Administration updated boxed warnings suggest avoiding fluoroquinolones for indications where there are safer alternatives, such as uncomplicated UTIs [11]. Despite this guidance, overall concordance with IDSA guidelines is poor in ambulatory practices, and fluoroquinolones are still used in >40% of uncomplicated UTIs [12–15]. Our study aimed to estimate the impact of an AS intervention, including local guidelines, education, and clinic data feedback, on guideline-concordant prescribing rates and urinary tract infection diagnoses.

METHODS

We conducted a prospective, 2-phase, quasi-experimental study to estimate the effect of a multifaceted education and data feedback intervention on antibiotic prescribing for UTI at 1 primary

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care (PC) clinic and 1 urgent care (UC) clinic in Durham, North Carolina.

Study Outcomes

The primary outcome, rate of guideline concordance, was defined as percentage of UTI encounters where first- or second-line antibiotics were prescribed according to clinic-specific, antibiogram-based guidelines. Secondary outcomes included (1) UTI diagnosis rates, (2) return visits for UTI within 30 days, (3) new antibiotic prescriptions written for UTI within 30 days, (4) clinic encounters, emergency department (ED) visits, or inpatient admissions due to antibiotic adverse events within 30 days, (5) average duration of therapy prescribed for UTI, and (6) 4-factor guideline concordance rates (ie, antibiotic, dose, frequency, and duration all align with developed guidelines). Guideline concordance and UTI diagnosis rates were assessed on all patients, while all other secondary outcomes were evaluated through chart review on a random 4% sample of the population. Recognizing that standardized guidelines are not applicable in all clinical scenarios, we pursued an additional aim to assess the percentage of patients with UTI diagnoses eligible for application of the clinic-specific guideline and used this to establish a target for percent guideline concordance at each clinic.

Interventions

Clinic-specific urinary-source antibiograms were created for the PC and UC clinics using urine cultures obtained from patients seen at either clinic between January 1, 2015, and December 31, 2016. UTI diagnosis and treatment guidelines were developed for each clinic based on these antibiograms and consensus guidelines (Supplementary Figures 1–4) [10, 16–19]. Clinic guidelines emphasized reducing fluoroquinolone use, avoiding asymptomatic bacteriuria (ASB) treatment, and using appropriate durations of therapy. PC guidelines promoted nitrofurantoin, sulfamethoxazole-trimethoprim, and oral beta-lactams for cystitis and fluoroquinolones for pyelonephritis. Due to increased resistance, UC guidelines recommended nitrofurantoin and oral beta-lactams for cystitis and adjunctive intramuscular ceftriaxone in combination with fluoroquinolones or sulfamethoxazole-trimethoprim for pyelonephritis.

Phase 1 began with a 1-hour educational session during mandatory quarterly provider meetings at PC on August 15, 2017, and UC on November 7 and November 14, 2017. Educators focused on the importance of antibiotic stewardship, reviewed the appropriate diagnosis, treatment, and duration of therapy for UTIs, and discussed clinic-specific guidelines. Digital copies of the guidelines and a link to CustomID, an online Duke infectious diseases resource, were provided to all clinicians [20]. “Commitment to Patients” posters, adapted from those created by the CDC, were provided to clinics, though unforeseen restrictions limited the intended high visibility [21]. Data were

emailed to the clinic once during Phase 1 and included monthly trends in UTI diagnoses, percentage of guideline concordance, percentage of mixed-growth urine cultures, antibiotic prescribing data, and a copy of the UTI guidelines to serve as reinforcement of the initial education session. The baseline period was 12 months before the education session, and the Phase 1 intervention included encounters from education to September 2018 for PC and November 2018 for UC.

A second in-person educational session in September 2018 for PC and November 2018 for UC started Phase 2 and was delivered during mandatory provider meetings. Original content presented in 2017 was reviewed along with data from Phase 1. In total, 11 of 14 (79%) PC providers and 72 of 96 (75%) UC providers attended education sessions, and 100% of providers were emailed education following sessions. There was low clinician turnover between the study periods. In Phase 2, 4 routine data feedback emails and 1 in-person feedback session were provided to clinicians at PC and 3 emails were sent to UC, with the last sent in April 2019. Data included trends in UTI diagnoses, guideline concordance, antibiotics prescribed, and actionable recommendations to improve prescribing based on findings from chart review. Additionally, peer comparison reports were emailed directly to PC providers in January and March 2019. In these reports, fluoroquinolone prescribing rates for each provider were compared with the clinic average and the “top performers” with the lowest rates of fluoroquinolone use, an approach used in other outpatient AS studies [22]. A detailed timeline of study interventions is provided in Figure 1, and a sample of data feedback reports can be found in Supplementary Figures 5 and 6.

Patient Selection and Eligibility

Patients were identified using the Duke Enterprise Data Unified Content Explorer (DEDUCE), a web-based clinical research tool [23]. DEDUCE was queried monthly for adult patients with a diagnosis code of acute cystitis or acute pyelonephritis between August 1, 2016, and July 30, 2019. Chart review validated the use of nonspecific codes such as “dysuria” and “UTI site not specified” for a diagnosis of cystitis. A list of included *International Classification of Diseases*, Ninth and Tenth Revision, Clinical Modification (ICD-9-CM) and (ICD-10-CM) diagnosis codes is located in Supplementary Table 1. Encounters without antibiotics prescribed within 5 days of the visit date were excluded from data analysis.

A random sample from at least 4% of all encounters was chart-reviewed to evaluate all secondary outcomes except for UTI diagnosis rates. To determine goal guideline adherence rates, patients evaluated during chart review were defined as guideline eligible if they lacked the following features: recurrent UTI (2 unique UTI episodes within the prior 6 months or 3 episodes within the prior 12 months), received antibiotics within the 30 days before UTI diagnosis, had allergies

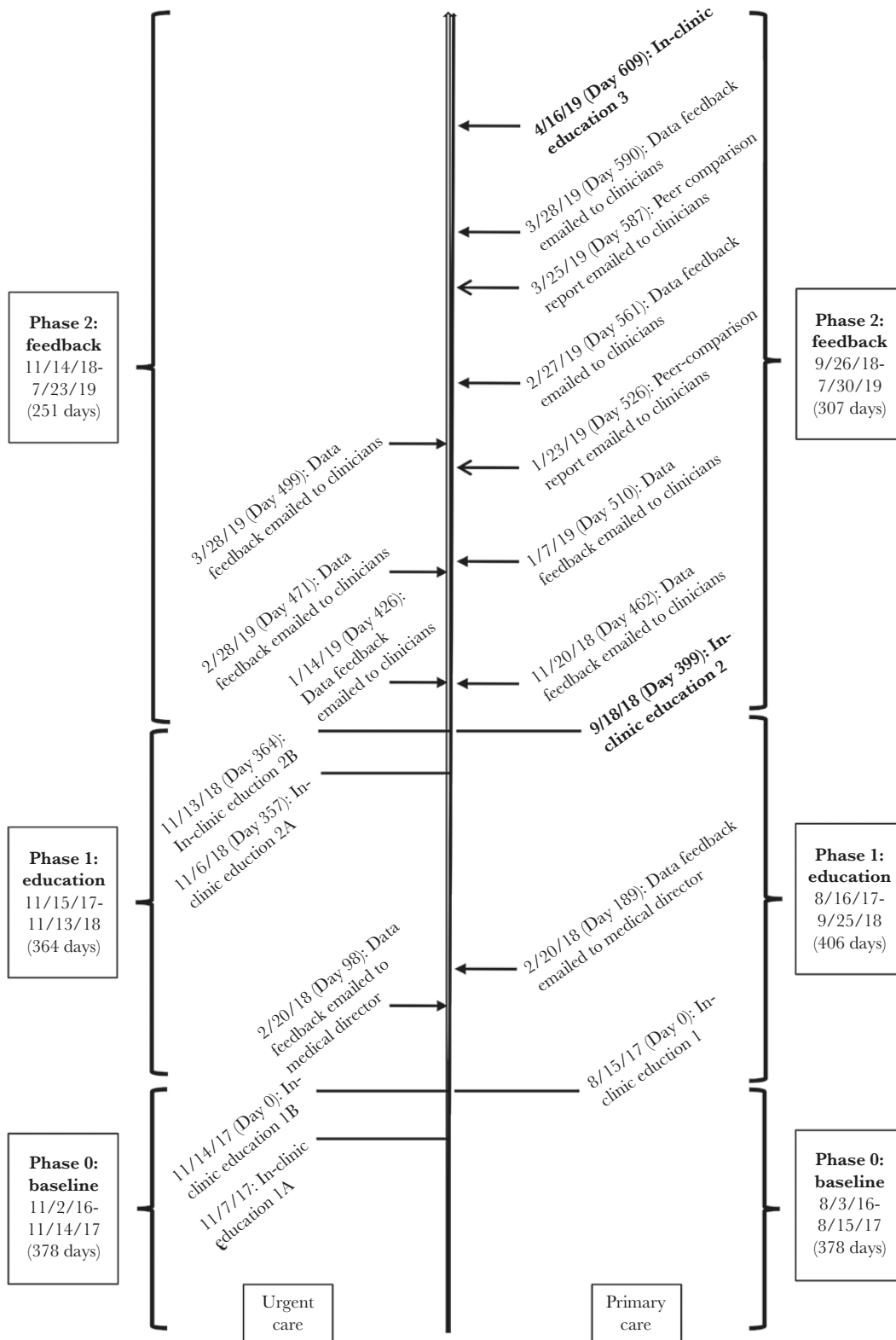


Figure 1. Study timeline. Timelines for primary care and urgent care are separate; timelines at either clinic have been either truncated or expanded to align the beginning of each phase of the intervention.

or intolerances to all first- and second-line antibiotics, had a second bacterial infection that warranted antibiotics, had urine cultures within the last year that were resistant to all first- and second-line antibiotics, or had indications that may support the treatment of ASB (pregnant, immunosuppressed, or patients undergoing genitourinary procedures associated with mucosal bleeding) [16].

Patient Consent

This quality improvement study was deemed exempt by the Duke University Institutional Review Board, and waivers of informed consent were granted.

Statistical Analysis

For the primary outcome, a segmented, linear regression with piecewise linear spline for time was used to model the level of change and trends in both the baseline and postintervention periods, an approach similar in design to multiple prior evaluations [24–27]. Specifically, the model adjusted for intervention (Phase 0 vs Phase 1 vs Phase 2), clinic (PC vs UC), and allowed time (measured in 2-week intervals) to have different slopes at different phases. The

secondary outcome, number of UTI encounters, was assessed similarly using a segmented Poisson regression model. The remaining secondary outcomes, collected and managed using REDCap electronic data capture tools hosted at Duke University, were assessed in the random sample of encounters described above [28]. For the primary outcome, significance of the test was assessed at alpha = .05. For secondary outcomes, point estimates and their 95% CIs were reported. Data management and analyses were conducted using SAS 9.4 (SAS Institute Inc., Cary, NC, USA).

RESULTS

A total of 4724 patients and 6318 encounters met inclusion criteria of an acute UTI at PC or UC between August 1, 2016, and July 30, 2019. Twenty percent (n = 1296) of encounters were excluded from further analysis because either no antibiotics were prescribed within 5 days of the encounter or antibiotic data were invalid (Figure 2). Of the remaining 5022 encounters that resulted in antibiotic prescriptions, 4875 (97%) were associated with a diagnosis of cystitis, the majority (83%) of which were at UC. Encounter characteristics were similar throughout the study (Table 1).

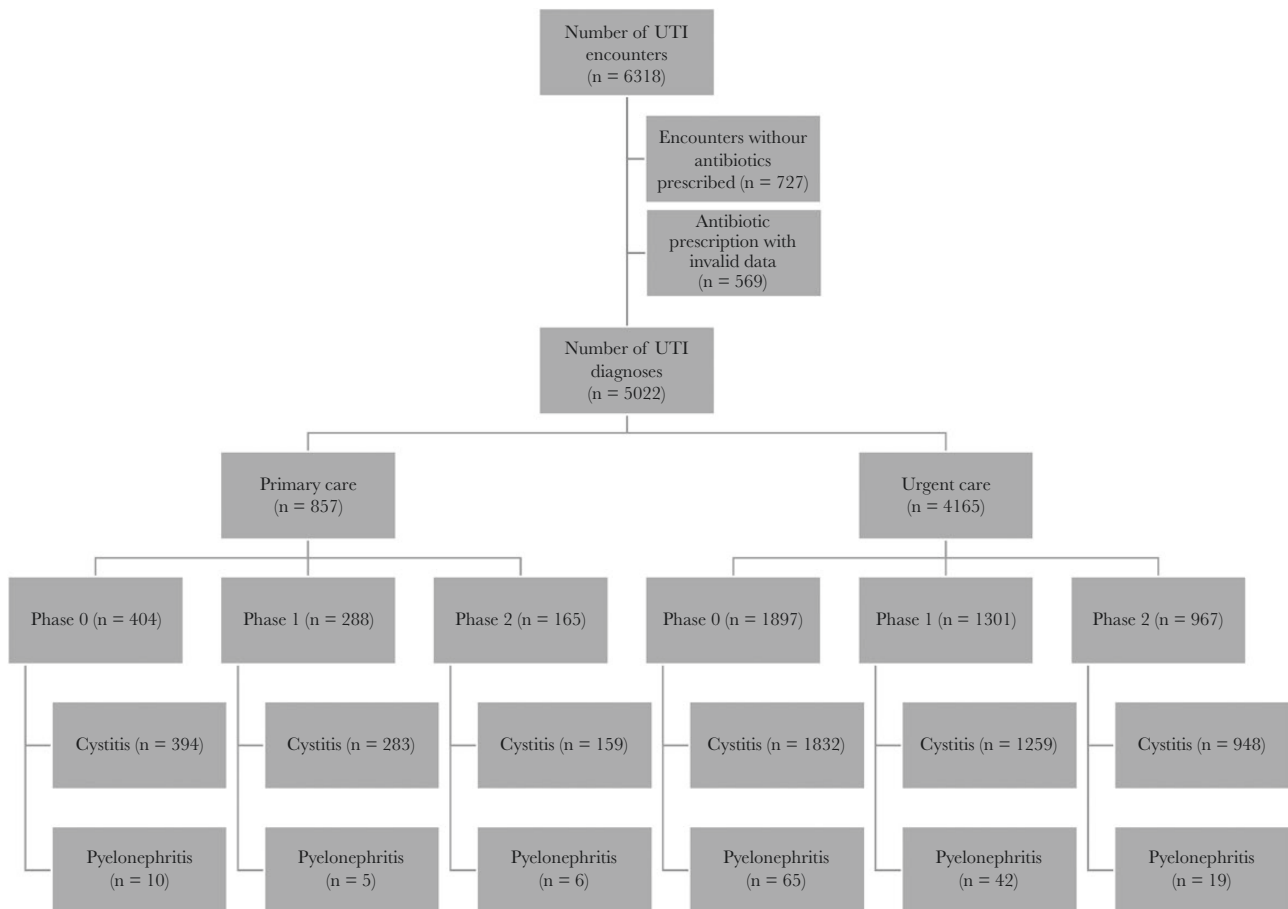


Figure 2. Flow diagram describing the patient encounters included in the analysis. Abbreviation: UTI, urinary tract infection.

Table 1. Patient Demographics for UTI Encounters at PC and UC Where Antibiotics Were Prescribed

	Primary Care			Urgent Care		
	Phase 0 (n = 404)	Phase 1 (n = 288)	Phase 2 (n = 165)	Phase 0 (n = 1897)	Phase 1 (n = 1301)	Phase 2 (n = 967)
Age, mean (SD), y	56.3 (18.1)	57.9 (18.5)	58.1 (18.1)	48.8 (20.4)	48.4 (20.8)	48.6 (20.1)
Female sex, No. (%)	373 (92.3)	268 (93.1)	146 (88.5)	1643 (86.6)	1127 (86.6)	835 (86.3)
Race, No. (%)						
Caucasian/White	226 (55.9)	172 (59.7)	90 (54.5)	1179 (62.2)	764 (58.7)	535 (55.3)
Black or African American	146 (36.1)	97 (33.7)	58 (35.2)	489 (25.8)	339 (26.1)	298 (30.8)
Other or unknown	32 (7.9)	19 (6.6)	17 (10.3)	229 (12.1)	198 (15.2)	134 (13.9)
Hispanic ethnicity, No. (%)	16 (4.0)	9 (3.1)	8 (4.8)	147 (7.7)	109 (8.4)	71 (7.3)
Payer group, No. (%)						
Private	230 (56.9)	138 (47.9)	83 (50.3)	1236 (65.2)	845 (65.0)	619 (64.0)
Medicare	157 (38.9)	131 (45.5)	71 (43.0)	517 (27.3)	353 (27.1)	258 (26.7)
Medicaid	15 (3.7)	18 (6.3)	11 (6.7)	132 (7.0)	97 (7.5)	84 (8.7)
Other or unknown	2 (0.5)	1 (0.3)	0	12 (0.6)	6 (0.5)	6 (0.6)
Encounter diagnosis, No. (%)						
Cystitis	394 (97.5)	283 (98.3)	159 (96.4)	1832 (96.6)	1259 (96.8)	948 (98.0)
Pyelonephritis	10 (2.5)	5 (1.7)	6 (3.6)	65 (3.4)	42 (3.2)	19 (2.0)

Demographics based on the number of unique UTI encounters, not the number of unique patients. Abbreviations: PC, primary care; UC, urgent care; UTI, urinary tract infection.

The segmented, linear regression analysis results on concordance over time are summarized in [Supplementary Tables 2 and 3](#). We observed no significant time-related effect on guideline concordance in the Phase 0 baseline period (−0.1%; 95% CI, −0.6% to 0.3%; *P* = .578). Immediately after Phase 1 education, an overall significant 21.8% increase in percentage of prescriptions for guideline-concordant antibiotics was observed (95% CI, 11.5% to 32%; *P* < .001). This effect diminished by 0.5% (95% CI, −0.9% to 0%; *P* = .049) for each 2-week period after the intervention. We observed a nonsignificant increase in the percentage of antibiotics that were guideline concordant upon

beginning Phase 2 when compared with the Phase 1 period (3.9%; 95% CI, −13.2% to 20.9%; *P* = .656). There was no significant change over time in the rate of guideline-concordant antibiotic prescriptions throughout the Phase 2 period (−0.6%; 95% CI, −1.6% to 0.4%; *P* = .232). The reversion back to baseline prescribing habits observed in the Phase 1 period halted with Phase 2 data feedback. Overall, the mean percentage of guideline concordance increased at PC from 65.8% at baseline to 72.6% in Phase 1 and further to 75% with routine data feedback. At UC, concordance increased from 35.8% at baseline to 57.3% in Phase 1 and 61% in Phase 2 ([Figure 3](#)).

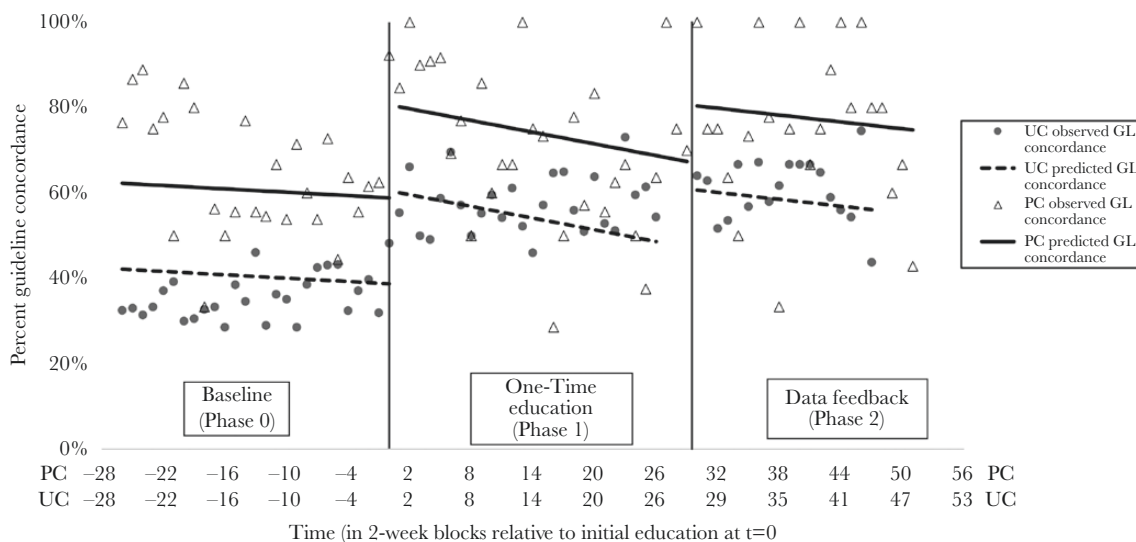


Figure 3. Time series analysis comparing rates of UTI visits with prescriptions for guideline-concordant antibiotics at the PC and UC clinics. Predicted concordance lines are plotted using predicted values for each clinic, obtained from the segmented regression analysis results. Abbreviations: GL, guideline; PC, primary care; UC, urgent care; UTI, urinary tract infection.

In the baseline period, the number of UTI diagnoses did not change over time (incidence rate ratio [IRR], 1 per 2 weeks; 95% CI, 0.99 to 1) (Supplementary Table 3). After Phase 1 education, UTI diagnoses decreased immediately by 21% (IRR, 0.79; 95% CI, 0.67 to 0.93). This immediate reduction was not seen again after education delivery in Phase 2 (IRR, 1.04; 95% CI, 0.8 to 1.36) (Supplementary Figure 7). Overall, the mean number of UTI diagnoses per 2-week period decreased at PC throughout the study period from 15 ± 3.5 at baseline to 9.9 ± 4.1 during Phase 1 and 7.5 ± 3.9 encounters per 2-week period during Phase 2. A similar trend was noted at UC with 2-week UTI diagnosis counts of 70.3 ± 11.1 at baseline, 50 ± 9.8 during Phase 1, and 53.7 ± 7.6 during Phase 2. Overall clinic visit volumes were stable throughout the study period.

At baseline, fluoroquinolones were prescribed during 26.7% of UTI visits at PC and 25% of UTI visits at UC (Table 2). Fluoroquinolone prescriptions decreased to 17.3% and 15.3% during Phase 1 and further to 16% and 11.5% during Phase 2 at PC and UC, respectively. Overall there was a 52.1% relative reduction in fluoroquinolone use for UTI and a 65.4% increase in nitrofurantoin (Table 2).

Manual chart reviews to assess the remaining secondary end points were performed on 71 (8.3%) and 166 (4%) encounters at PC and UC, respectively. The demographics of this sample were similar to the overall population. Twenty-five of 71 (35.2%) PC encounters and 39 of 166 (23.5%) UC encounters were identified as scenarios where the developed guidelines would not apply. Therefore, an estimated target goal for guideline concordance was 65% for PC and 76% for UC. The most common reasons for exclusion from the developed guidelines were antibiotics prescribed within the last 30 days or patients with recurrent UTIs, which were the case for 20 (28.2%) PC encounters and 29 (17.5%) UC encounters. Secondary end points were assessed using the remaining 173 total encounters at PC and UC.

The number of encounters meeting the 4-factor guideline-concordant criteria increased throughout the study from 15 of 79 (19%) at baseline to 16 of 69 (23.2%) during Phase 1 and 7 of 25 (28%) during Phase 2 (Supplementary Table 4). Inappropriate duration of therapy, defined as differing from guideline recommendations, was the most common reason for divergence. Despite this, the mean duration of therapy decreased from 7.6 ± 2.4 days at baseline to 7.3 ± 2.1 days and 6.6 ± 1.6 days among all sampled encounters during Phases 1 and 2, respectively (Supplementary Table 5). Treatment failure and antibiotic adverse effects were infrequent, and tests for statistical significance were not conducted due to the small sample size (Supplementary Table 6). Unnecessary treatment of asymptomatic bacteriuria was identified in 4 (5%), 4 (6%), and 4 (16%) patients during Phases 1, 2, and 3, respectively.

DISCUSSION

Inappropriate fluoroquinolone prescribing for UTI is a potential target for outpatient stewardship programs [10, 14]. In this study, we developed a multifaceted intervention with the aim of improving management of UTI in 2 outpatient clinics. The provision of education and guidelines during Phase 1 significantly increased rates of guideline-directed antibiotics and use of nonfluoroquinolone therapies for UTI. Despite the initial increase in guideline-concordant prescribing, prescribing patterns trended back toward baseline without continued reinforcement, a challenge reported by other investigators [29, 30]. In the second phase of this study, we added routine feedback with the aim of enhancing durability of guideline concordance. While this additional AS intervention did not significantly change rates of guideline-concordant antibiotic prescriptions, as occurred in Phase 1, the significant decline in concordance rates seen during Phase 1 halted. This sustained

Table 2. Antibiotics Prescribed for UTIs at PC and UC Before and After the Education

	Primary Care				Urgent Care				Combined PC and UC % Change ^a
	Phase 0 (n = 416), No. (%)	Phase 1 (n = 301), No. (%)	Phase 2 (n = 175), No. (%)	% Change ^a	Phase 0 (n = 2072), No. (%)	Phase 1 (n = 1440), No. (%)	Phase 2 (n = 1048), No. (%)	% Change ^a	
Nitrofurantoin	122 (29.3)	125 (41.5)	64 (36.6)	+ 24.7	573 (27.7)	586 (40.7)	501 (47.8)	+ 72.9	+ 65.4
Fluoroquinolone	111 (26.7)	52 (17.3)	28 (16.0)	- 40.0	518 (25.0)	220 (15.3)	120 (11.5)	- 54.2	- 52.1
TMP-SMX	116 (27.9)	53 (17.6)	33 (18.9)	- 32.4	460 (22.2)	155 (10.8)	125 (11.9)	- 46.3	- 44.2
PO cephalosporin	42 (10.1)	45 (15.0)	28 (16.0)	+ 58.5	185 (8.9)	237 (16.5)	157 (15.0)	+ 67.8	+ 65.8
IM ceftriaxone	4 (1.0)	9 (3.0)	7 (4.0)	+ 316	202 (9.7)	173 (12.0)	100 (9.5)	- 2.1	+ 5.7
Other ^b	21 (5.0)	17 (5.6)	15 (8.6)	+ 69.8	134 (6.5)	69 (4.8)	45 (4.3)	- 33.6	-21.3

Abbreviations: IM, intramuscular; PC, primary care; PO, oral; TMP-SMX, trimethoprim-sulfamethoxazole; UC, urgent care; UTI, urinary tract infection.

^aDefined as the relative change in antibiotic use throughout the entire study period, from Phase 0 to Phase 2.

^bUncommonly prescribed antibiotics and those that may have been chosen for an alternative diagnosis. Category includes amoxicillin, clarithromycin, clindamycin, doxycycline, fosfomicin, and moxifloxacin. Fosfomicin prescribed in <1% of all encounters.

response to the intervention is highlighted by the steady reduction in fluoroquinolone prescribing across all phases of this study.

Upon evaluation of secondary outcomes, we found that UTI diagnoses declined after the start of Phase 1. A focus of the education throughout both phases of the intervention was appropriate diagnosis of UTI; thus a lower number of UTI diagnoses may indicate that more patients were identified as ASB and not treated. Despite an overall reduction in antibiotic durations and UTI diagnoses, chart review data suggest further opportunity to target durations of therapy and ASB. No evidence of harm following the initiative was detected. Management of UTIs after the intervention resulted in decreased exposure to broad-spectrum antibiotics and improved adherence to national guidelines.

Our findings are consistent with prior outpatient AS research for other infections, which suggests that displaying patient-centered AS posters, educating clinicians, and providing data feedback are effective strategies to promote stewardship [31, 32]. Data supporting the use of these interventions for UTI are limited. One French study assessed the impact of regional UTI guidelines in combination with provider education on antibiotic prescribing. While there were statistically significant reductions in norfloxacin prescribing, the study assessed antibiotic prescribing across all indications and did not directly analyze trends in other antibiotics commonly (but not specifically) prescribed for UTIs, including beta-lactams and sulfamethoxazole-trimethoprim [27]. Our study adds to these findings and provides an analysis of antibiotics that are specifically prescribed during UTI encounters.

Our study is not without limitations. Identification of our cohort relies on the accurate use of diagnosis codes by clinicians. Often, patients treated for UTI are coded with nonspecific diagnoses such as “dysuria.” In our review of a random subset of patients, we confirmed intent to treat a UTI even among those coded with nonspecific diagnoses. Diagnosis code shifting, a practice by which a provider intentionally miscodes a visit to justify an antibiotic prescription, has been seen in outpatient AS studies [2]. We assessed diagnostic accuracy by performing chart reviews on a random subset of all clinic visits and verified that the diagnosis code matched the clinical intent in almost all cases. Additionally, diagnosis rates for pyelonephritis, an allowable code for broader agents, were consistent throughout the study.

We excluded encounters without antibiotic prescriptions within 5 days of our analysis. Based on chart review, these visits were often found to be unrelated to UTI (eg, dysuria resulting from vulvovaginal candidiasis). While we hypothesize that a reduced number of UTI encounters treated with antibiotics is partially a result of decreased treatment of asymptomatic patients, we are unable to confirm this.

Lastly, we may not have seen the maximal effects of our intervention due to inherent implementation challenges. While we

intended for commitment posters to be posted in high-visibility areas, restrictions set by our health system required they be placed in alternative areas and likely limited their impact. There was limited provider turnover; thus our findings may not be reproducible in settings with more frequent staff changes, such as clinics staffed by medical residents. Our data feedback intervention was emailed at routine intervals, and there is no guarantee that all clinicians reviewed the data feedback. Secondary outcomes such as antibiotic duration of therapy, treatment failure, or adverse effects were only evaluated on a random subset chart review. Future studies that leverage the electronic health record to evaluate these outcomes on a larger scale would be beneficial.

The present study suggests that the provision of clinic-specific urinary antibiograms, along with treatment guidelines, clinician education, and data feedback to clinicians, was successful at increasing guideline concordance, including a robust shift away from fluoroquinolones toward other agents with a narrower spectrum and lower collateral damage, and reducing UTI diagnoses. While routine data feedback helped to maintain the initial improvement in guideline-concordant antibiotic selection, significant additional benefit was not realized. The provision of routine data feedback and peer comparison reports was time-intensive to implement and may be impractical for institutions to maintain. Future studies are warranted to further understand the sustainability and scalability of AS interventions in order to optimize outpatient antibiotic use, given the new TJC standards. The success of these stewardship strategies could be employed in other infectious syndromes or other non-infectious disease states where outpatient prescribing diverges from national guideline recommendations.

Supplementary Data

Supplementary materials are available at Open Forum Infectious Diseases online. Consisting of data provided by the authors to benefit the reader, the posted materials are not copyedited and are the sole responsibility of the authors, so questions or comments should be addressed to the corresponding author.

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