Assessment of Testing and Treatment of Asymptomatic Bacteriuria Initiated in the Emergency Department

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Background. Reducing antibiotic use in patients with asymptomatic bacteriuria (ASB) has been inpatient focused. However, testing and treatment is often started in the emergency department (ED). Thus, for hospitalized patients with ASB, we sought to identify patterns of testing and treatment initiated by emergency medicine (EM) clinicians and the association of treatment with outcomes.

Methods. We conducted a 43-hospital, cohort study of adults admitted through the ED with ASB (February 2018–February 2020). Using generalized estimating equation models, we assessed for (1) factors associated with antibiotic treatment by EM clinicians and, after inverse probability of treatment weighting, (2) the effect of treatment on outcomes.

Results. Of 2461 patients with ASB, 74.4% (N = 1830) received antibiotics. The EM clinicians ordered urine cultures in 80.0% (N = 1970) of patients and initiated treatment in 68.5% (1253 of 1830). Predictors of EM clinician treatment of ASB versus no treatment included dementia, spinal cord injury, incontinence, urinary catheter, altered mental status, leukocytosis, and abnormal urinalysis. Once initiated by EM clinicians, 79% (993 of 1253) of patients remained on antibiotics for at least 3 days. Antibiotic treatment was associated with a longer length of hospitalization (mean 5.1 vs 4.2 days; relative risk = 1.16; 95% confidence interval, 1.08–1.23) and Clostridioides difficile infection (CDI) (0.9% [N = 11] vs 0% [N = 0]; P = .02).

Conclusions. Among hospitalized patients ultimately diagnosed with ASB, EM clinicians commonly initiated testing and treatment; most antibiotics were continued by inpatient clinicians. Antibiotic treatment was not associated with improved outcomes, whereas it was associated with prolonged hospitalization and CDI. For best impact, stewardship interventions must expand to the ED.

Keywords: bacteriuria; emergency medicine; stewardship; urinary tract infection.

Urinary tract infections (UTIs) are the second most common infection treated in hospitalized patients [1, 2], but they are frequently misdiagnosed. Bacteriuria without signs or symptoms attributable to a UTI is defined as asymptomatic bacteriuria (ASB) [3]. Asymptomatic bacteriuria is frequent among hospitalized patients, and national guidelines recommend against antibiotic treatment in most patients. Nevertheless, ASB is often unnecessarily treated with antibiotics [2–5]. Antibiotic use is associated with increased adverse events, antimicrobial resistance, duration of hospitalization, cost, and Clostridioides difficile infection (CDI) [2, 6, 7]. Thus, ASB is a prime target for antimicrobial stewardship.

Unnecessary antibiotic use in the emergency department (ED) is common, but data on antibiotic stewardship in the ED are sparse [8–10]. Diagnostic uncertainty and time pressures may lead to antibiotic overuse by EM clinicians [11]. It is not yet known how often testing and treatment related to ASB in hospitalized patients is initiated by EM clinicians, or how often antibiotics started by EM clinicians are continued by the inpatient clinician. Thus, for hospitalized patients with ASB who were admitted through the ED, we sought to identify how often urine testing and antibiotic treatment is started by EM clinicians, patterns and predictors of antibiotic treatment, and the association of antibiotic treatment with patient outcomes.

METHODS

Study Setting and Inclusion and Exclusion Criteria
This study included 43 hospitals participating in the Michigan Hospital Medicine Safety (HMS) Consortium. HMS is a quality collaborative funded by Blue Cross Blue Shield of Michigan and Blue Care Network that aims to improve the care of hospitalized
For this study, patients were eligible for inclusion if they (1) were admitted through the ED, (2) had a positive urine culture (UC) collected within 2 days of hospitalization, and (3) had ASB. Patients were considered to have ASB if they had no documented signs or symptoms meeting UTI diagnostic criteria per Infectious Diseases Society of America (IDSA) Guidelines and National Healthcare Safety Network (NHSN) definitions [3, 12]. Specifically, patients could not have 1 of the following documented: dysuria, urinary frequency/urgency, suprapubic pain, fever (temperature > 38.0°C), costovertebral pain/tenderness, hematuria, autonomic dysreflexia, or increased spasticity in patients with a spinal cord injury. Because patients with acute alterations in mental status (AMS) cannot always describe their symptoms, patients with AMS and a systemic sign of possible infection (peripheral leukocytosis >10 000 cells/mm³, systolic blood pressure <90 mm Hg, or ≥2 criteria for systemic inflammatory response syndrome [SIRS]) were considered to potentially have UTI and thus were excluded.

Patients for whom treatment of ASB may be appropriate were excluded, including those with altered urinary tract anatomy, pregnancy, severe immune compromise (solid organ or bone marrow transplant, human immunodeficiency virus with CD4 <200 cells/mm³, neutropenia [absolute neutrophil count <0.5 cells/mm³]), or those who met criteria for severe sepsis (defined as ≥2 SIRS criteria + signs of organ dysfunction, including elevated lactate, creatinine >2 mg/dL, bilirubin >2 mg/dL, platelet <100 000/μL, international normalized ratio >1.5). Additional exclusions included (1) age <18 years, (2) intensive care unit admission within 3 days before or after UC, (3) entered hospice during hospitalization, (4) left against medical advice, (5) concomitant infection (documentation by provider of an additional bacterial infection during hospitalization, except CDI), (6) active treatment and/or prophylaxis for UTI on admission, (7) isolated candiduria, (8) within 30 days from a prior hospitalization that would have qualified the patient for inclusion in the study. Patients were also excluded from analysis if relevant data were missing or the antibiotic was ordered more than 1 day from the UC (ensuring that the decision was based on patient status around time of testing).

Study Design, Data Collection, and Patient Sampling

Hospital medicine safety procedures for patient selection, data collection, and quality assurance have been described [2, 6, 13]. In brief, from February 1, 2018 to February 25, 2020, trained abstractors at each hospital retrospectively screened consecutive patients 30 days after discharge and included the first patient each day with a positive UC (positive defined as a UC with any bacterial growth identified as abnormal by the hospital's microbiology policy). Data were collected from 90 days before admission until follow-up was terminated by a major complication (eg, death) or 30 days after discharge. Signs and symptoms were collected from 3 days before and after UC collection. An abnormal urinalysis (UA) was defined as presence of leukocyte esterase or >5 white blood cells per high-power field and/or nitrates. At 30 days postdischarge, outcome data were collected by medical record review and scripted telephone follow-up (3 attempts). A standardized data dictionary and random audits by quality coordinators ensured data integrity. Variables collected from the medical record included (1) patient demographics, (2) receipt of antibiotic (within 90 days prior), (3) nonspecific signs or symptoms not consistent with the definition of UTI, (4) severity of illness, and (5) laboratory results.

Outcomes and Exposures

The primary outcome was the percentage of patients who had antibiotic treatment (at least 1 dose) initiated by an EM clinician. Antibiotic treatment likely directed at CDI (eg, metronidazole) was not considered ASB treatment. We also assessed total antibiotic duration (inpatient plus discharge) in patients who received treatment. Secondary outcomes included 30-day mortality, 30-day readmission, 30-day ED visit, discharge to postacute care facility, CDI within 30 days, and duration of hospitalization after urine testing (UA or UC).

Statistical Analysis

Descriptive statistics were used to characterize the population. To evaluate predictors of antibiotic treatment by EM clinicians, we first used bivariable logistic general estimating equation (GEE) models accounting for hospital-level clustering then determined a multivariable model using stepwise selection based on the Schwarz criterion [14].

To evaluate the association of antibiotic treatment by EM clinicians with secondary outcomes, we first used logistic regression to create inverse probability of treatment weights [15] based on baseline covariates found to be significant in the bivariable and/multivariable analyses and other factors potentially associated with the outcome. These weights were then applied to GEE models (logistic or binomial, as appropriate) to assess differences in outcomes by treatment. Because there were no CDI events in patients with ASB who were not treated with antibiotics, we used Fisher’s exact test to compare CDI rates between treatment and no treatment. P values less than .05 were considered statistically significant. All analyses were performed using SAS, version 9.4 (SAS Institute Inc., Cary, NC).

Sensitivity Analysis

Because some patients with AMS are unable to report symptoms, we conducted a sensitivity analysis assessing factors and outcomes associated with ASB treatment by EM clinicians after excluding patients with AMS.
RESULTS

Baseline Demographics

Of 2461 patients admitted through the ED and ultimately determined to have ASB (Supplementary eFigure 1), the median age was 78 years (interquartile range [IQR], 67–87) and 73.9% (N = 1818) were women (Table 1). The most common comorbidities included chronic kidney disease (41.2%,

Table 1. Demographics of Hospitalized Patients Presenting to the Emergency Department Ultimately Identified as Having Asymptomatic Bacteriuria, N = 2461

| Characteristic                           | N (%)  |
|-----------------------------------------|--------|
| Age, median [IQR]                       | 78 [67–87] |
| Women                                   | 1818 (73.9%) |
| Length of stay [IQR]                    | 5 [3–6]  |
| Comorbidities*                         |        |
| Moderate to severe chronic kidney disease | 942 (38.3%) |
| Diabetes                                | 1013 (41.2%) |
| Hemodialysis                            | 35 (1.4%) |
| Liver disease                           | 138 (5.6%) |
| Congestive heart failure                | 619 (25.2%) |
| Cerebrovascular disease                 | 646 (26.2%) |
| History of cancer                       | 510 (20.7%) |
| Spinal cord injury                      | 50 (2.0%) |
| Immunosuppressed*                      | 71 (2.9%) |
| Dementia                                | 622 (25.3%) |
| Urinary Catheter                        |        |
| Indwelling                              | 353 (14.3%) |
| Other*                                  | 67 (2.7%) |
| Urinalysis                               |        |
| Urinalysis obtained                     | 2421 (98.4%) |
| Urinalysis Result                       |        |
| Abnormal urinalysis*                    | 2313 (94.0%) |
| Positive LE and/or >5 WBC/hpf           | 2264 (92.0%) |
| Positive nitrite                        | 966 (39.3%) |
| Documentation of reason for culture*    | 1830 (74.4%) |
| Abnormal urinalysis                     | 543 (22.1%) |
| Altered mental status                   | 223 (9.1%) |
| Nausea, vomiting, abdominal pain        | 199 (8.1%) |
| Changes in urine characteristics         | 173 (70.0%) |
| Urine Pathogens                         |        |
| *Escherichia coli*, n (%)               | 1180 (47.9%) |
| *Klebsiella spp*, n (%)                 | 383 (15.6%) |
| *Enterococcus spp*, n (%)               | 259 (10.5%) |
| *Proteus spp*, n (%)                    | 162 (6.6%) |
| *Pseudomonas aeruginosa*, n (%)         | 108 (4.4%) |
| *Enterobacter spp*, n (%)               | 77 (3.1%)  |
| *Citrobacter spp*, n (%)                | 69 (2.8%)  |

N = 1013) and diabetes (38.3%, N = 942). Indwelling urinary catheters were present in 14.3% (N = 353) of patients. Almost all UAs were abnormal (94.0%, N = 2313). When documented (74.4%, 1830 of 2461), the most common indication for the UC was an abnormal UA (22.1%, 543 of 2461). The most common bacteria isolated were *Escherichia coli* (47.9%, N = 1180), *Klebsiella* spp (15.6%, N = 383), and *Enterococcus* spp (10.5%, N = 259).

Three quarters (74.4%, 1830 of 2461) of patients were treated with antibiotics (Supplementary eFigure 2 and eFigure 3) with a median treatment duration of 6 days (IQR, 3–9) and median hospital duration of 5 days (IQR, 3–6). The UC was ordered by EM clinicians in 80.0% (N = 1970 of 2461) of patients (Supplementary eFigure 2), and antibiotic treatment was started by EM clinicians in 68.5% (1253 of 1830) of those treated (Supplementary eFigure 3). When antibiotic therapy was started by EM clinicians, 79.2% (993 of 1253) of patients remained on antibiotics for 3 or more days (Figure 1). Likewise, when antibiotic therapy was started by inpatient clinicians, 82.0% (473 of 577) remained on antibiotics for 3 days or more. The most common initial antibiotic was ceftriaxone (78.2%, N = 1418), and at discharge the most common antibiotic was a fluoroquinolone (30.2%, N = 290). Hospitals with higher rates of urine testing by EM clinicians also had higher rates of
Variables Associated With Asymptomatic Bacteriuria Treatment

Variables associated with ASB treatment by EM clinicians included patient comorbidities such as dementia or spinal cord injury, patient symptoms such as AMS, or laboratory results, in particular an abnormal UA (Table 2).

In the multivariable model (Table 3), patient characteristics associated with treatment by EM clinicians included the following: dementia (odds ratio [OR], 1.43; 95% confidence interval [CI], 1.11–1.84), spinal cord injury (OR, 5.92; 95% CI, 1.36–25.72), presence of urinary catheter (OR, 1.54; 95% CI, 1.17–2.03), incontinence (OR, 1.81; 95% CI, 1.40–2.33), and AMS (OR, 2.34; 95% CI, 1.82–3.00). Laboratory characteristics associated with ASB treatment by EM were peripheral leukocytosis (OR, 1.42; 95% CI, 1.21–1.68) and abnormal UA (OR, 9.68; 95% CI, 5.34–17.54).

Patient Outcomes

Within 30 days of discharge, 77% (1895 of 2461) of patients were evaluated by record review and/or telephone or had follow-up terminated by a major complication. After adjustments, there were no differences in mortality, hospital readmission, ED visit, or discharge to postacute care facility among patients treated by EM clinician versus never treated with antibiotics (Table 4). Patients treated with antibiotics by an EM clinician were more likely to develop CDI within 30 days (0.9% [N = 11] vs 0% [N = 0]; P = .02) and have a longer duration of hospitalization after urine testing (mean 5.1 vs 4.2 days; RR, 1.16; 95% CI, 1.08–1.23).

Sensitivity Analysis

Results were similar after excluding patients with AMS (Supplementary eTable 1). Specifically, patients treated with antibiotics by an EM clinician were more likely to develop CDI within 30 days (0.9% [N = 8] vs 0% [N = 0]; P = .03) and have a longer duration of hospitalization after urine testing (mean
| Variable                                      | Antibiotic Treatment by EM (n = 1253) | No Antibiotics (n = 631) | Odds Ratio (95% CI) | P Valuea |
|-----------------------------------------------|--------------------------------------|--------------------------|---------------------|----------|
| Baseline Characteristics                      |                                      |                          |                     |          |
| Age (median, IQR)                             | 80 (69–87)                           | 75 (62–85)               | 1.02 (1.01–1.03)    | <.0001   |
| Gender (female)                               | 911 (72.7%)                          | 471 (74.6%)              | 0.99 (0.78–1.25)    | .92      |
| Race (white)                                  | 921 (74.0%)                          | 495 (78.6%)              | 0.96 (0.79–1.13)    | .54      |
| Charlson comorbidity index,0                  | 132 (10.5%)                          | 84 (13.3%)               | REF                 | .16      |
| 1–2                                           | 407 (32.5%)                          | 200 (31.7%)              | 1.28 (0.91–1.79)    | .29      |
| 3–4                                           | 396 (31.6%)                          | 175 (27.7%)              | 1.42 (1.04–1.94)    | .10      |
| ≥5                                            | 318 (25.4%)                          | 172 (27.3%)              | 1.27 (0.96–1.67)    |          |
| Diabetes                                      | 452 (36.1%)                          | 250 (39.5%)              | 0.89 (0.73–1.10)    | .29      |
| Moderate or severe chronic kidney disease     | 520 (41.5%)                          | 247 (39.1%)              | 1.24 (0.96–1.61)    | .10      |
| History of cancer                             | 261 (20.8%)                          | 132 (20.9%)              | 1.04 (0.85–1.28)    | .68      |
| Spinal cord injury                            | 35 (02.8%)                           | 3 (0.5%)                 | 5.17 (1.42–18.77)   | .01      |
| Dementia                                      | 384 (30.6%)                          | 87 (13.8%)               | 2.27 (1.83–2.83)    | <.0001   |
| Immunosuppressed                              | 37 (03.0%)                           | 18 (2.9%)                | 1.24 (0.81–1.62)    | .55      |
| IV chemotherapy in preceding 30 days          | 10 (0.8%)                            | 6 (1.0%)                 | 0.96 (0.20–4.58)    | .96      |
| Hemodialysis                                  | 20 (01.6%)                           | 9 (1.4%)                 | 0.84 (0.51–1.37)    | .48      |
| Transfer from postacute carec                | 88 (07.0%)                           | 27 (4.3%)                | 1.74 (1.27–2.39)    | .0006    |
| Nonambulatory                                 | 224 (179%)                           | 55 (8.7%)                | 1.92 (1.46–2.53)    | <.0001   |
| Hospitalization in past 90 days               | 373 (29.8%)                          | 195 (30.9%)              | 0.92 (0.76–1.10)    | .34      |
| Antibiotics in preceding 90 days              | 263 (21.0%)                          | 92 (14.6%)               | 1.31 (1.06–1.62)    | .01      |
| Indwelling catheter                           | 215 (172%)                           | 58 (9.2%)                | 1.55 (1.21–1.98)    | .0006    |
| Any urinary catheterd                         | 249 (19.9%)                          | 68 (10.8%)               | 1.55 (1.22–1.96)    | .0003    |
| Signs and Symptoms                            |                                      |                          |                     |          |
| Abdominal pain                                | 248 (19.8%)                          | 158 (25.0%)              | 0.88 (0.73–1.05)    | .16      |
| Incontinence                                  | 445 (35.5%)                          | 138 (21.9%)              | 2.20 (1.80–2.68)    | <.0001   |
| Functional decline                            | 93 (07.4%)                           | 28 (44.4%)               | 1.79 (0.88–3.64)    | .11      |
| Acutely altered mental status                 | 393 (31.4%)                          | 86 (13.6%)               | 2.51 (2.02–3.13)    | <.0001   |
| Fatigue, malaise, lethargy                    | 396 (31.6%)                          | 175 (27.7%)              | 1.53 (1.16–2.01)    | .003     |
| Nausea or vomiting                            | 267 (21.3%)                          | 179 (28.4%)              | 0.88 (0.69–1.13)    | .33      |
| Change in color, sediment, or malodorous urine| 202 (16.1%)                          | 71 (11.3%)               | 1.93 (1.23–3.03)    | .004     |
| Urinary retention or postvoid residual > 200 cc| 143 (11.4%)                          | 54 (8.6%)                | 1.34 (0.96–1.88)    | .08      |
| Severity of Illness                           |                                      |                          |                     |          |
| qSOFAa (≥2 vs <2)                             | 152 (12.1%)                          | 65 (10.3%)               | 1.33 (1.06–1.68)    | .01      |
| ≥ 2 SIRSb criteria                            | 300 (23.9%)                          | 191 (30.3%)              | 0.80 (0.66–0.97)    | .02      |
| Laboratory Results                            |                                      |                          |                     |          |
| Peripheral leukocytosisa                      | 356 (28.4%)                          | 186 (29.5%)              | 1.01 (0.87–1.16)    | .93      |
| Abnormal urinalysisb                         | 1232 (96.3%)                         | 528 (83.7%)              | 9.42 (5.30–16.75)   | <.0001   |
| Hospital Characteristics                      |                                      |                          |                     |          |
| Type of control                               |                                      |                          |                     |          |
| Not-for-profit                                | 1141 (91.1%)                         | 606 (96.0%)              | REF                 | .04      |
| For profit                                    | 112 (8.9%)                           | 25 (4.0%)                | 2.57 (1.03–6.40)    |          |
| Bed size (10 bed increase)                    | 327 (203–443)                        | 310 (189–443)            | 1.01 (0.99–1.01)    | .62      |
| Teaching hospital                             | 1165 (93.0%)                         | 555 (88.0%)              | 1.19 (0.59–2.40)    | .62      |

Abbreviations: CI, confidence interval; EM, emergency medicine clinician; IQR, interquartile range; IV, intravenous; qSOFA, quick sequential organ failure assessment; REF, Reference; SIRS, systemic inflammatory response syndrome.

aP < .05 is considered significant.

bDefined as chemotherapy administered within 30 days, human immunodeficiency virus with CD4 ≥200, >10 mg/day prednisone for at least 30 days (or equivalent steroid dose), on biologic agents such as tumor necrosis factor inhibitors or other immunosuppressant agents, congenital or acquired immunodeficiency.

cIncludes transfer from the following: subacute rehabilitation center, skilled nursing home, acute rehabilitation center, assisted living, other hospital. Also includes if patient had been admitted or resided in a nursing home, subacute rehabilitation center, or extended care facility in the prior 30 days.

dIncludes Foley catheter, intermittent straight catheterization, and suprapubic catheter present on day of urine culture collection or 1 day before urine culture collection.

eQuick SOFA score: systolic blood pressure <100 mmHg = 1, respiratory rate ≥22 breaths per minute = 1; Glasgow coma score <15 = 1.

fSIRS (temperature <36°C [96.8°F] or >38.0°C [100.4°F], heart rate >90 beats per minute, respiratory rate >20 breaths per minute, white blood cell count <4000/mm³ or >12 000/mm³).

gDefined as white blood cell count >10 per high-power field.

hDefined as presence of leukocyte esterase or nitrite, or white blood cells >5 per high-power field.
Table 3. Multivariable Model of Patient Factors Associated With Treatment by Emergency Medicine Clinicians of Patients Ultimately Diagnosed With Asymptomatic Bacteriuria, N = 1884

| Variable                          | Odds Ratio (95% CI) | P Value |
|----------------------------------|---------------------|---------|
| Age                              | 1.01 (1.00–1.02)    | .006    |
| Dementia                         | 1.43 (1.11–1.84)    | .006    |
| Urinary catheter                 | 1.54 (1.17–2.03)    | .002    |
| Incontinence                     | 1.81 (1.40–2.33)    | <.0001  |
| Spinal cord injury               | 5.92 (3.66–25.72)   | .02     |
| Acutely altered mental status    | 2.34 (1.62–3.00)    | <.0001  |

Table 4. Outcomes for Treatment by Emergency Medicine vs No Antibiotic Treatment for Asymptomatic Bacteriuria, N = 1884

| Outcome                          | Treated by EM (N = 1253) | No Antibiotics (N = 631) | Unadjusted Odds Ratio (95% CI) | P Value | Adjusted Odds Ratio (95% CI) | P Value |
|----------------------------------|---------------------------|--------------------------|-------------------------------|---------|-------------------------------|---------|
| Death                            | 41 (3.3%)                 | 12 (1.9%)                | 1.76 (0.90–3.44)              | .10     | 1.83 (0.86–3.92)              | .12     |
| Readmission                      | 131 (10.5%)               | 69 (10.9%)               | 1.21 (0.96–1.54)              | .10     | 1.19 (0.81–1.77)              | <.0001  |
| ED visit                         | 160 (12.8%)               | 71 (11.3%)               | 1.21 (0.89–1.63)              | .22     | 1.19 (0.86–1.63)              | <.0001  |
| Discharge to postacute care facility | 47 (3.7%)                | 45 (7.1%)                | 1.03 (0.68–1.57)              | .90     | 1.02 (0.67–1.55)              | .90     |
| Clostridioides difficile infection | 11 (0.9%)                | 0 (0%)                   | N/A                           | N/A     | N/A                           | N/A     |
| Duration of hospitalization (mean (SD)) | 5.1 (2.6)              | 4.2 (2.4)                | 1.21 (1.12–1.27)              | .02     | 1.16 (0.98–1.33)              | <.0001  |

Abbreviations: CI, confidence interval. NOTE: Odds ratios >1 indicate factors associated with treatment of asymptomatic bacteriuria; P < .05 is considered significant.

5.1 vs 4.1 days; RR, 1.16; 95% CI, 1.07–1.27) (Supplementary eTable 2).

**DISCUSSION**

In this study of 2461 patients with ASB admitted through the ED, three quarters were treated with antibiotics during their hospitalization. The majority of urine testing and antibiotic treatment was initiated by EM clinicians. Once started by EM clinicians, inpatient clinicians usually continued antibiotics during hospitalization. The ASB treatment by EM clinicians was not associated with clinical benefit, but instead it was associated with CDI and longer duration of hospitalization after urine testing. These findings identify the ED as a key target to reduce antibiotic use and improve outcomes in hospitalized patients with ASB.

Multiple prior studies have reported similarly high rates of ASB treatment in hospitalized patients ranging from 47% to 80% [2, 16–19]. However, data characterizing which clinicians order urine testing and initiate antibiotic therapy in hospitalized patients with ASB has been lacking. We identified that most initial urine testing is ordered by EM clinicians. It is notable that EM clinicians do not always initiate urine testing. To improve crowding, nurse-initiated order sets and testing protocols are common [20] and may contribute to unnecessary testing. Likewise, order sets may have prechecked or easily selected orders for urine testing for nonurinary complaints, such as stroke. Addressing these contributors through diagnostic stewardship efforts [21] can reduce urine testing sent to evaluate for possible UTI in asymptomatic patients. We also found that hospitals with higher rates of urine testing by EM clinicians were more likely to have higher rates of urine testing by the inpatient clinician, suggesting that an underlying “culture of culturing” may...
contribute to increased testing [22, 23]. Nevertheless, antibiotic stewardship should target the (1) ED and ED protocols and (2) order sets to reduce urine testing and ASB treatment.

We found the strongest predictor of unnecessary antibiotic treatment by EM clinicians to be an abnormal UA. Likewise, prior studies of ASB in hospitalized patients have found urine testing results, both the UA and UC, to be associated with treatment by clinicians [2, 18, 19]. Both internal medicine physicians and ED nurses have been found to have knowledge gaps regarding the interpretation of UAs and UTI diagnosis [24–27]. The misinterpretation of UAs (ie, a positive UA equates with a UTI) has been highlighted as a target for decreasing inappropriate ASB treatment [2, 18, 19, 28, 29]. Although the UA has a high negative predictive value and is useful in ruling out a UTI, it is not indicative of a UTI in the absence of symptoms. Qualitative studies have identified that this is poorly understand and that a “positive UA” is incorrectly cited as the reason for sending a UC, diagnosing UTI, and initiating antibiotics [24, 30].

One potential way to change UC ordering practices is through elimination of reflex UCs where the UC is automatically sent when the UA is abnormal [19]. Recent strategies in the ED un-linking UA and UC have been successful in reducing UC rates [31]. Our study reaffirms that interventions aimed at both physicians and nurses regarding the correct interpretation of the UA should be paired with nudges for when to send a UC based on evidence-based UTI signs and symptoms. Thus, creating system and culture changes targeting decreasing urine testing in patients without urinary symptoms is vital to reducing unnecessary testing and treatment of ASB.

We also identified common patient characteristics, dementia and AMS, to be associated with treatment of suspected UTI in patients without urinary symptoms by EM clinicians. Both factors have previously been associated with treatment of ASB for hospitalized patients [2, 18]. Both confusion and AMS are common in the elderly, and bacteriuria is also common in elderly patient populations, thus bacteriuria in an elderly patient with confusion can be seen often by chance alone [32]. A systematic review concluded that no strong evidence links AMS or confusion alone as a symptom of UTI [33]. Furthermore, inaccurately diagnosing a UTI in patients with ASB could lead to delays in another clinically significant diagnosis. Due to known harms of unnecessary antibiotics, the IDSA recommends observation and assessment for alternative etiologies (eg, dehydration, medication, hypoxia, sundowning) in elderly patients with AMS and/or dementia that are clinically stable.

Despite this, deeply held beliefs persist and varying practice patterns exist making it difficult to reduce antibiotic treatment in patients with AMS and ASB. Patients with AMS, no systemic signs of infection, and no other urinary symptom account for only ~25% of all ASB patients admitted through the ED, and thus they may not be the best to target for stewardship efforts. When patients with AMS were excluded from the analysis, the same potential harms remain associated with ASB treatment (increased duration of hospitalization and CDI). These data could be used as a common ground between antibiotic stewards and EM clinicians, to prevent harm by first focusing on those patients who can report symptoms (without AMS), because they remain a majority (75%) of the patients tested in the ED.

Similar to a prior study, we found that antibiotics started for ASB by EM clinicians are not typically discontinued [34]. Most patients received 3 or more days of antibiotics. If antibiotics are started by EM clinicians for a suspected UTI in a patient without urinary symptoms, the next critical point to intervene is on admission, when the inpatient clinician can reassess and stop the antibiotic. Each additional dose results in increasing risk of harm, including adverse drug events and CDI [7, 35].

Treatment of ASB has been associated with CDI in patients undergoing neurosurgery [36], and we identified the same association in hospitalized medicine patients. This demonstrates potential harm from antibiotics in hospitalized patients with ASB who are unnecessarily treated. In addition, similar to a prior study of hospitalized medical patients with ASB started on antibiotics by any clinician, we found that patients with ASB who were treated with antibiotics had a longer duration of hospitalization [2]. Because ASB is common, antibiotics are harmful, and diagnosis momentum makes discontinuing antibiotics challenging [37], stewardship interventions must target both the initial testing by EM clinicians and the continuation of antibiotics by inpatient clinicians.

Our study has limitations. First, as an observational retrospective study, we are limited in assessment of symptoms and signs of UTI by documentation, and therefore we may have overestimated the frequency of ASB. Although we attempted to exclude patients with a possible alternate source of infection, the retrospective nature of our study limits our ability to determine the reason for antibiotic prescribing with absolute certainty. Second, excluding patients with concomitant infections may have underestimated the true rate of testing and treatment in patients with ASB. Third, we cannot fully attribute the orders to a particular provider, because it is possible that EM clinicians were asked to order a urine test or start antibiotics by the admitting clinician. Fourth, by excluding patients who had antibiotics started more than 1 day from the culture date, we were biased toward higher rates of antibiotic starts by EM clinicians compared with inpatient clinicians. However, excluding this subset of patients was necessary to compare the decision to treat suspected UTI in an asymptomatic patient, avoiding clinical changes or new diagnostic information. This does not change the overall trend of higher antibiotic starts by EM clinicians. Fifth, given the retrospective nature of the study, we cannot determine how many patients would have been started on antibiotics by the inpatient clinician if the EM clinician had not started antibiotics. Last, although we adjusted for potential
con founding, residual confounding may still exist, including for the associated outcomes identified.

Our study has strengths. This cohort represents a diverse group of hospitals, improving generalizability. Our data were collected by trained abstractors who underwent quality control with excellent reliability, ensuring data accuracy. In addition, we used phone calls in conjunction with record review to capture higher rates of adverse events. We also attempted to minimize bias when evaluating secondary outcomes by using inverse probability of treatment weighting.

CONCLUSIONS

Emergency medicine clinicians often order urine testing and treat for a presumed UTI in patients who do not have urinary symptoms. Inpatient clinicians often continue unnecessary antibiotic therapy. Risk factors for unnecessary antibiotic initiation by EM clinicians include an abnormal UA and nonspecific symptoms. Antibiotic treatment of patients with ASB, also when excluding those with AMS, was not associated with improved outcomes but was associated with an increased risk of CDI and longer duration of hospitalization after urine testing. For efforts to be successful in curbing ASB treatment, stewardship should begin in the ED before the initiation of the testing and treatment cascade.

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 copyrighted and are the sole responsibility of the authors, so questions or comments should be addressed to the corresponding author.

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