Community acquired urinary tract infections among adults in Accra, Ghana

Background: Urinary tract infection (UTI) is one of the most common bacterial infectious diseases encountered in clinical practice, and accounts for significant morbidity and high medical costs. To reduce its public health burden, there is the need for local research data to address aspects of prevention and management of UTI. The aim of this study was to investigate community-acquired UTI among adults in Accra, Ghana, including the risk factors, etiological agents, and antibiotic resistance.

Methods: This was a cross-sectional study involving 307 patients clinically diagnosed with UTI at the Korle Bu and Mamprobi polyclinics in Accra. Urine specimens were collected from the study participants and analyzed by culture, microscopy, and dipstick. The bacterial isolates were identified using standard microbiological methods and tested against a spectrum of antibiotics by the Kirby Bauer method. Multidrug resistant Enterobacteriaceae isolates were screened for Extended Spectrum β-lactamase (ESBL) production by the double disc method, and isolates that tested positive were analyzed by Polymerase Chain Reaction for ESBL genes. Demographic information and clinical history of study participants were collected.

Results: Based on the criteria for laboratory confirmed UTI, 31 (10.1%) of the 307 specimens were positive and the main risk factor of UTI among the study participants was pregnancy ($P=0.02$, OR=$2.43$). The most common uropathogen isolated was Escherichia coli (48.9%), followed by Klebsiella sp. (16.1%). Prevalence of resistance was highest for Piperacillin (87.1%) and Amoxicillin+Clavulanic Acid (87.1%) and lowest for Amikacin (12.9%). Prevalence of multidrug resistance among the uropathogens was 80.1% (25) and the most common ESBL gene detected was CTX-M-15.

Conclusion: Pregnant women constitute the key risk population of UTI in Accra, while Amikacin remains a suitable drug for the treatment of febrile UTI. The high prevalence of multidrug resistance among the uropathogens highlights the need for surveillance of antimicrobial resistance among these pathogens.

Keywords: extended spectrum β-lactamases, multidrug resistance, urinary tract infection

Introduction

Urinary tract infection (UTI) refers to microbial invasion of the urinary tract by one or more uropathogenic bacteria species, leading to significant bacteriuria and the presence of symptoms such as dysuria. It is one of the commonest diseases diagnosed in outpatients and affects approximately 150 million people yearly. The associated cost of healthcare is enormous, accounting for $659 million in direct costs for treating and $936 million in indirect costs, totaling $1.6 billion annually. Two main types of UTIs are known based on how the infection is acquired: hospital acquired UTI (Nosocomial UTI) and community acquired UTI.
UTI. Hospital acquired UTI is defined as the onset of UTI in patients, 48 hours after admission, while community acquired UTI refers to the development of infections before admission to the hospital and not within 10 days after the patient has been discharged. Escherichia coli is the commonest cause of both community and hospital acquired UTI. Other common uropathogens encountered in community acquired UTI include Staphylococcus saprophyticus, Klebsiella pneumoniae, and Citrobacter spp. The etiology of hospital acquired UTI is, however, more varied and includes a wide range of organisms such as Pseudomonas aeruginosa and Proteus sp., which are hardly encountered in community-acquired UTI.

Community-acquired UTI are usually uncomplicated, and the risk factors commonly include female sex, being sexually active, and use of spermicidal contraceptives. On the other hand, hospital acquired UTI is usually complicated and is associated with risk factors such as catheterization and recent antibiotic use.

The use of antimicrobial therapy has contributed a great deal to the management of UTI. However, accumulated evidence shows that treatment of these infections is increasingly becoming difficult due to the rapid emergence of antimicrobial resistance in hospitals and the community, a phenomenon attributed to overuse and misuse of antibiotics. Although a global problem, antibiotic resistance of uropathogens carries more significance in the developing world, where treatment options are limited. In Ghana, antimicrobial treatment of UTI is mainly empirical due to a relative lack of appropriate laboratory facilities for culture and susceptibility testing of bacteria in several health facilities. Unfortunately, few studies have reported on antimicrobial resistance patterns of uropathogens in Ghana, and without recent surveillance data of antimicrobial susceptibility, empirical treatment of UTI could be ineffective and expensive. To help address this problem and contribute to effective management and prevention of UTI, this study was carried out. The aim of the study was to investigate community-acquired UTI in Accra, Ghana, including the risk factors, etiological agents, and antibiotic resistance.

Materials and methods

Study site and sampling
The study was conducted at two clinics in Accra, namely Korle Bu Polyclinic and Mamprobi Polyclinic. Accra is the capital city of Ghana and has a population of about 1.8 million people. The two polyclinics provide primary healthcare to a large section of the adult population in Accra due to their strategic location in the city center. Korle Bu Polyclinic is a 42-bed facility, while Mamprobi Polyclinic is a 54-bed facility (Prince Horlortu, personal communication, 2016). In addition to general medical services, both clinics provide healthcare in areas of minor surgeries, medical laboratory service, radiography, and assorted body imaging services. This study was cross-sectional in design and was conducted between April and September 2016. Using a 95% confidence level, a 22.5% estimated UTI prevalence reported previously and a 5% allowable error, 307 patients who were clinically suspected of having a UTI were randomly recruited from both polyclinics. Patients recruited met the inclusion criteria of not been admitted into the hospital for not more than 48 hours and satisfying the age requirement of 13 years and above. Patients who had been on antibiotics 2 weeks or earlier prior to the study were excluded. Information on demographic and clinical features of the study participants was extracted from their clinical records, and these included age, gender, body mass index (BMI), previous UTI, presence of diabetes, frequency of sex, and pregnancy (in the case of females). A mid-steam urine sample was obtained from each of the study participants for analysis in the bacteriology laboratory of the School of Biomedical and Allied Health Sciences, University of Ghana, which is less than a kilometer from each of the study clinics.

Laboratory investigations

Analysis of urine samples
The urine samples were cultured on Cysteine Lactose Electrolyte Deficient (Oxoid Ltd., Basingstoke, UK) media and incubated at 37°C for 18–24 hours. Following incubation, colonies of bacteria were counted and counts of \( \geq 10^5 \) (cfu/mL) were regarded as significant bacteriuria. Isolated bacteria were identified based on their colonial morphology, Gram-stain reactions and biochemical tests including oxidase test, triple sugar iron (TSI) fermentation tests, indole test, citrate utilization test, urea utilization test, catalase test, coagulase test, and motility.

After the culture process, 5mL of the remaining urine samples were aseptically transferred into 15 mL falcon tubes for centrifugation. The macroscopic properties (color and appearance) of the urine samples were observed and recorded for each sample. Dipstick analysis of each sample was performed by immersing the urine test strip entirely into the urine samples for a brief period (2 seconds). The strips were removed, then examined for biochemical
parameters including glucose, proteins, pH, specific gravity, ketones, nitrites, bilirubin, urobilinogen, and the presence of cells such as leucocytes and erythrocytes. Attention was paid to the nitrite and leucocyte, indicating portions of the strip as they serve as highlights for possible bacterial infection. The urine transferred into the clean transparent falcon tubes were centrifuged at 2,500 rpm for 3 minutes, after which the supernatant was decanted and the residue used to prepare a wet mount and observed under the high dry power objective of the light microscope. Evidence of pus cells, red blood cells, parasites, yeasts cells, epithelial cells, crystals, casts, and bacteria were sought for and recorded. Urinary tract infections in the study participants were determined by significant bacteriuria with the presence of pyuria.26

Antibiotic susceptibility testing
Susceptibility testing of isolates was performed using the Kirby Bauer disc diffusion method with strict adherence to the 2016 Clinical Laboratory Standards Institute (CLSI) guidelines. The antibiotics tested included: Nalidixic acid (30 µg), Ciprofloxacin (5 µg), Norfloxacin (20 µg), Levofoxacin (5 µg), Amikacin (30 µg), Gentamicin (10 µg), Nitrofurantoin (300 µg), Pipercillin (20 µg), Oxacillin (1 µg), Amoxicillin+Clavulanic Acid (Augmentin) (30 µg), Cefuroxime (30 µg), Ceftazidime (20 µg), and Tetracycline (30 µg). Standardized inoculums were prepared in sterile saline to turbidity comparable to 0.5 McFarland standard solution of barium sulfate. Muller-Hinton agar plates were swabbed with the standardized inoculums and antimicrobial discs aseptically placed on it after drying the plate for 3–5 minutes. The plates were then incubated at 37°C for 18 hours.27,28 The zone sizes were measured and interpreted using the 2016 CLSI guidelines (www.clsi.org).28

Screening for Extended Spectrum β-lactamases (ESBLs) was performed on multi-drug resistant Enterobacteriaceae isolates using the double disc method.28 Antibiotic discs of Ceftazidime (30 µg), and Ceftazidime plus Clavulanic acid (30/10 µg) were placed on Mueller Hinton agar and incubated at 37°C for 18–24 hours. An organism was considered as an ESBL producer if there was a ≥5 mm increase in the zone diameter of Ceftazidime/Clavulanate disc and that of Ceftazidime disc alone.28 E. coli ATCC 25922 and a known in house ESBL producer were used as negative and positive controls, respectively. Positive ESBL isolates were further characterized using Polymerase Chain Reaction (PCR) (Qiagen Multiplex PCR kit) to determine the specific genes responsible for conferring resistance to the isolates.

For detection of ESBL genes, SHV, TEM, OXA-2, OXA-10, and the CTX-M group 1, 2, and 9 primers were used. All PCR protocols included an initial denaturation of 94°C for 15 minutes and a final extension at 72°C for 10 minutes. The PCR products were analyzed for nucleotide sequencing at Inqaba biotech™, South Africa, and compared with sequences in the NCBI database (http://www.ncbi.nlm.nih.gov/BLAST).

Data analysis
Data collected in the study were entered into Microsoft Excel version 2010 and analyzed using STATA version 12. Descriptive analyses were carried out on the study variables, which included computation of arithmetic means, percentages, and frequencies. Uni-variable associations were performed between UTI and all the other variables recorded: Analysis of variance was used for numeric variables, whereas Chi-square test was used for categorical variables. Variables significantly associated with urinary tract infection in the invariable analysis were used as independent variables in a logistic regression analysis to identify determinants of urinary tract infection. The statistical significance of the independent variables was evaluated by confidence intervals, and odds ratios including P-values; variables with P<0.05 were regarded as significant.

Ethical clearance
Ethical approval for the study was obtained from the ethical and protocol review committees of the Ghana Health Service and College of Health Sciences, University of Ghana (CHS-Et/M.4-P4.11/2015–2016). Written informed consent was also obtained from the study participants according to the Declaration of Helsinki.

Results
Demographic and clinical features of study participants
Demographic and clinical features of the 307 study participants are reported in Table 1. The mean age of the study participants was 37.2 (±17.6) years, and the majority (75%) of them were less than 50 years old. A high proportion of 79.8% (245) of the study participants were females, of which 27.4% (84) were pregnant. Diabetes occurred in 6.8% (21/307) of the study participants and 22.5% (69/307) had experience of previous UTI. About half of the study participants (56%) had a normal body mass index, 13.7% were obese, 27% were overweight and 3.3% were underweight. A proportion of 26.1% (80/307) did not
practice sex; 33.2% (102/307) practiced sex twice or less a month, while 40.7% (125/307) practiced it more than twice a month.

UTI and associated risk factors

Thirty-one (10%) of the 307 patients enrolled in the study had UTI, all of which were uncomplicated. Urinary tract infections were more common in females (93.6%, 31/29) than males (6.4%, 31/2) (P=0.06). In the multivariate analysis, pregnancy was the only factor that was significantly associated with UTI among the study participants (Table 2). Age, BMI, gender, diabetes, frequency of sex, and previous UTI did not show a significant association with UTI.

| Table 1 Demographic and clinical features of the study participants |
|-------------------|---|---|
| Feature              | N  | %   |
| **Gender**           |    |     |
| Male                 | 62 | 20.2|
| Female               | 245| 79.8|
| **Age (Mean=37.2±17.6)** |    |     |
| <50                  | 231| 75.2|
| ≥50                  | 76 | 24.8|
| **Frequency of sex** |    |     |
| No sexual activity   | 80 | 26.1|
| Twice or less a month | 102 | 33.2|
| More than twice a month | 125 | 40.7|
| **Diabetes**         |    |     |
| Diabetic             | 21 | 6.8 |
| Non-diabetic         | 286| 93.2|
| **Pregnancy**        |    |     |
| Yes                  | 84 | 27.4|
| No                   | 223| 72.6|
| **Body Mass Index**  |    |     |
| Underweight (<18)    | 10 | 3.3 |
| Normal (18–24.9)     | 172| 56  |
| Overweight (25.0–29.9) | 83  | 27  |
| Obese (>30)          | 42 | 13.7|
| **Previous UTI**     |    |     |
| Yes                  | 69 | 22.5|
| No                   | 238| 77.5|

**Abbreviations:** N, number of study participants; UTI, Urinary Tract Infection.

| Table 2 Risk factors of urinary tract infection identified through logistic regression |
|------------------|---|---|---|
| Feature              | OR   | 95% CI | P-value |
| Gender               | 4.03 | 0.9–17.4 | 0.06 |
| Age (Mean=37.2±17.6) | 0.97 | 0.9–1.0 | 0.07 |
| Frequency of sex     | 1.55 | 0.9–2.5 | 0.08 |
| Diabetes             | 0.43 | 0.05–3.3 | 0.41 |
| Pregnancy            | 2.42 | 1.1–5.2 | 0.02 |
| Body Mass Index      | 1.01 | 0.6–1.6 | 0.97 |
| Previous UTI         | 0.81 | 0.3–2.1 | 0.66 |

**Abbreviations:** OR, odds ratio; CI, Confidence Interval; UTI, Urinary Tract Infection.

| Table 3 Bacteria isolated from urine specimens |
|-----------------------------|---|---|
| BACTERIA                   | N  | %   |
| *Escherichia coli*          | 15 | 48.4|
| *Klebsiella* sp.            | 5  | 16.1|
| *Staphylococcus aureus*     | 4  | 12.9|
| *Citrobacter* sp.           | 3  | 9.7 |
| *Pseudomonas aeruginosa*    | 1  | 3.2 |
| *Proteus mirabilis*         | 1  | 3.2 |
| *Klebsiella pneumoniae*     | 1  | 3.2 |
| *Enterobacter* sp.          | 1  | 3.2 |

**Abbreviation:** N, number of positive urine specimens.

Causative organisms of UTI and antibiotic resistance

Eight different bacterial species were isolated from urine specimens of the study participants; the most prevalent was *Escherichia coli* (48.4%) followed by *Klebsiella* sp. (16.1%) and *Staphylococcus aureus* (Table 3).

Overall prevalence of antibiotic resistance among the bacterial isolates ranged from Amikacin (12%) to Piperacillin (87%) (Table 4). Antibiotic resistance was particularly high for β-lactam antibiotics (Table 4). As shown in Figure 1 for *E. coli* (most common uropathogen isolated), the prevalence of antibiotic resistance decreased across Piperacillin/ Amoxicillin+Clavulanic Acid (93.4%), Nalidixic acid (73.4%), Tetracycline (53.4%) Norfloxacin (40.0%), Cefadizime (26.7%), Nitrofurantoin (26.7%), Gentamicin (26.7%), Ciprofloxacin (20.0%), Levofloxacin (20.0%), Amikacin (6.7%), and Cefuroxime (6.7%). The overall prevalence of MDR among the bacterial isolates was 77.4% (24/31); Multi-drug resistance (MDR) prevalence among *E. coli* isolates was 66.7% (10/15). The predominant ESBL gene detected among the sequenced isolates was CTX-M-15 type followed by TEM-3 (Table 5).
### Table 4 Overall prevalence of resistance among the antibiotics tested

| Class of antibiotics | Antibiotic tested                                                                 | N          | (%)             |
|----------------------|----------------------------------------------------------------------------------|------------|-----------------|
| Quinolones           | Nalidixic acid-NaI (30 µg), Ciprofloxacin-Cip (5 µg), Norfloxacin-Nor (20 µg), Levofloxacin-Lev (5 µg) | 22, 8, 13, 7 | 70.97, 25.81, 41.94, 22.58 |
| Fluoroquinolones     |                                                                                  |            |                 |
| Aminoglycosides      | Amikacin-Amk (30 µg), Gentamicin-Gen (10 µg)                                     | 4, 10      | 12.90, 32.26    |
| Furadantins          | Nitrofurantoin-Nit (300 µg)                                                      | 13         | 41.94           |
| β-lactam- β-lactamase inhibitors | Piperacillin-Pip (20 µg), Oxacillin-Ox (1 µg), amoxicillin+clavulanic acid-Aug (30 µg) | 27, 3, 27  | 87.10, 75.00, 87.10 |
| Cephalosporin 2nd generation, Cephalosporin 3rd generation | Cefuroxime-Cef (30 µg), Ceftazidime-Cft (20 µg)                                 | 7, 15      | 22.58, 48.39    |
| Tetracyclines        | Tetracycline-Tet (30 µg)                                                         | 18         | 58.06           |

Abbreviation: N, Number of resistant isolates.

### Table 5 Distribution of ESBL genotypes and genes among multi-drug resistant Enterobacteriaceae isolates from urine

| Specimen number | Bacteria        | ESBL genotype | ESBL sequence type |
|-----------------|-----------------|---------------|--------------------|
| 62              | Escherichia coli| TEM           | TEM-1              |
| 108             | Escherichia coli| TEM, CTX-M    | TEM-116, CTX-M-15  |
| 134             | Citrobacter sp. | CTX-M         | CTX-M-15           |
| 136             | Klebsiella sp.  | TEM, CTX-M    | TEM-116, CTX-M-15  |
| 149             | Citrobacter sp. | TEM, CTX-M    | TEM-1, CTX-M-15    |
| 284             | Klebsiella sp.  | TEM, SHV      | TEM-3, SHV-1       |
| 189             | Proteus sp.     | CTX-M         | CTX-M-15           |

Abbreviation: ESBL, extended spectrum β-lactamases.
Discussion

In this study, we investigated community-acquired UTI among adults in Accra with the goal of providing the necessary epidemiological information to reduce the burden of these infections. In the analysis of risk factors, only pregnancy emerged as a risk factor of UTI. The association of pregnancy with UTI concurs with several studies, and is probably due to the series of structural and functional urinary tract changes that occur during the course of pregnancy. During pregnancy, there is a usually high level of circulating progesterone, which causes urethral sphincter relaxation and a reduction in smooth muscle tone with slowing of ureteral peristalsis. Simultaneously, the enlarged uterus compresses the urinary bladder, thus increasing the intravesical pressure, which may result in vesico-ureteral reflux and urine retention in the bladder after micturition. Urinary stasis and impairment of the physiological anti-reflux mechanism create conditions favorable for bacterial growth and ascending infection.

Uropathogens isolated from the study participants were mainly members of the Enterobacteriaceae which concurs with previously published data. E. coli, which is the leading causative organism of both community and hospital, was the most predominant organism isolated from the study participants. In Ghana, E. coli was the most common uropathogen isolated from stroke patients, sickle cell disease patients, and pregnant women. In the pathogenesis of UTI, E. coli employs factors such iron acquisition systems, fimbrae that mediate attachment to host tissues, toxins (hemolysins and autotransporter toxins), flagella, and special proteins that help to weaken and aid evasion of the host innate and adaptive immune system. Furthermore, E. coli is known to form bacterial communities within the tissue of the bladder wall to enable it to proliferate without the influence of antibacterial molecules and host inflammatory cells and also away from the flow of urine. Two pathogenic mechanisms allow microorganisms to reach the urinary tract: the ascending and the hematogenous routes. In the ascending route, which is more common, microorganisms of the intestinal microbiota such as E. coli colonize the periurethral space and ascend through the urethra to the bladder and eventually the kidneys. The hematogenous route occurs in patients with bacteremia or endocarditis, mostly caused by S. aureus, which spreads to the kidneys.

We observed very high percentage resistance for Piperacillin (87.1%), Amoxicillin+Clavulanic Acid (87.1%), and Nalidixic Acid (71%), which concur with previous studies on antibiotic resistance of uropathogens in Ghana. This raises concerns about the suitability of these antibiotics for empirical treatment of UTI in Ghana. In line with the trend of increasing MDR, especially in the developing world, we observed a prevalence of 77.4% overall and 66.7% for E. coli. This, coupled with ESBLs detected in Enterobacteriaceae isolates, poses a major challenge to the treatment of UTI in Ghana. ESBL producing uropathogenic E. coli appear to be associated with more serious UTI resulting in septicemia. CTX-M, the most common ESBL harbored by the Enterobacteriaceae in this study, is a new family of plasmid-mediated ESBLs that preferentially hydrolyzes cefotaxime. These ESBLs, particularly CTX-M-15 and CTX-M-14, are known to be associated with community-acquired infections, which concurs with this study. The problem of ESBLs is the fact that they could be transferred from one organism to another through plasmids and by conjugation among bacteria. Plasmid profiling of MDR E. coli by Baral et al. identified the presence of plasmids with varying sizes ranging from 2–51 kilobases coupled with a high frequency of conjugation. Some studies have attributed the emergence and spread of MDR to clonal groups of E. coli which have common antimicrobial sensitivity patterns and virulence. Our data depicts the situation of alarming high levels of antibiotic resistance in Ghana, which is due to several factors, including the prescription of antibiotics by poorly trained health workers and also weak government regulations and law enforcement. These allow for the sale of sub-standard drugs, which are mostly from unlicensed outlets resulting in self-medication practices that are now a common trend in Ghana. Additionally, it is common for unqualified health practitioners to offer antibiotics in small quantities to ignorant individuals with the motive of cutting cost and quickly treating suspected infections, which results in sub-inhibitory concentrations within the body’s tissues, thereby facilitating selection of drug resistant strains. In Ghana, antibiotics can be obtained without prescription from many pharmacies, and this now seems to be the mainstay of treating common infections in the community, including uncomplicated UTI. With the high incidence of antibiotic resistance of uropathogens observed in the current study, this mode of treating UTI should be discouraged as it is likely to result in treatment failures. There is the need for patients to report to licensed clinics where proper clinical
investigations including antibiotic susceptibility testing of 
uropathogens can be done to ensure effective treatment.

In contrast to Piperacillin, Amoxicillin+Clavulanic acid, and Nalidixic Acid, there was a generally high sus-
sceptibility of the urinary isolates to Amikacin (>85%),
which makes it a suitable option for treatment of UTI in 
Ghana. A recent study in Ghana also reported that a high 
percentage (93.6%) of urinary isolates were susceptible to 
Amikacin. It is important to note that Amikacin is used 
treating cases of febrile UTI and is regarded as an 
efficient option before carbapenem treatment, especially 
in patients with infections caused by ESBL-producing 
uropathogens that are extensively antibiotic resistant. 
Amikacin is a relatively expensive antibiotic in Ghana, 
and the high cost is likely to protect the efficacy of this 
drug in the country for a long period of time.

Although all the study subjects were clinically diagnosed 
with UTI, the prevalence of culture-positive urine specimens 
was as low as 10%. By comparison, similar studies in India 
and the US reported that 45% and 50%, respectively, of 
patients clinically diagnosed with UTI had culture-positive 
urine. The disparity between clinical and laboratory 
(culture) diagnosis of UTI can be partly attributed to non-
specific clinical symptoms, which is a common problem in 
the diagnosis of the infection. It could also be due to medica-
tion with antibiotics among the patients prior to specimen 
collection, although we excluded patients who had taken 
antibiotics 2 weeks or earlier prior to the study. Our data 
probably shows a situation of high prevalence of over-
diagnosis of UTI, which requires attention and further inves-
tigation. Inconsistency between clinical and laboratory 
diagnosis of UTI could lead to inappropriate use of antibiotics 
resulting in antimicrobial resistance, higher healthcare costs, 
increased antibiotic exposure, a greater number of adverse 
reactions, and other unintended outcomes, such as 
Clostridium difficile infection.

The main limitation of the study is that we obtained few 
urine culture positive samples, which limited the numbers of 
isolates for antimicrobial susceptibility testing. This could have 
also affected identification of some risk factors of UTI in the study.

Conclusion

E. coli is the most common etiological agent of commu-
nity acquired UTI in the study area, while pregnancy is the 
main risk factor. While antibiotics such as Amoxicillin 
+Clavulanic Acid and Piperacillin are unsuitable for 
empirical treatment of any form of UTI in the study area,
Amikacin remains a suitable drug for the treatment of febrile UTI. The high prevalence of MDR and occurrence of 
ESBLs among the uropathogens highlights the need for 
surveillance of antimicrobial resistance among these 
pathogens. In this regard, there could be a national system 
of pooling isolates of uropathogens from various hospitals. 
The study has exposed a serious problem with UTI diag-
nosis in Ghana which needs to be addressed. There is the 
need for further studies.

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Disclosure

The authors report no conflicts of interest in this work.

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