LETTERS TO THE EDITOR 1433

Formula version	Incidence (measured: 1.9)	Prevalence (measured: 3.2)
Published ^a	0.56	7.60
Adjusted 1 ^b	2.29	2.66
Adjusted 2 ^c	1.14	5.32

^a The formula published by Rhame and Sudderth.¹

^b Using the minimum antibiotic treatment duration of

7 days as the length of infection.

^c Using the maximum antibiotic treatment duration of

14 days as the length of infection.

allowed for cases of HCA UTI to be diagnosed on the basis of clinical symptoms only, meaning that we are not necessarily comparing like with like. However, in this scenario, the assumption is that the incidence would be underestimated in a laboratory system. Using the formula as it is presented by Rhame and Sudderth¹ for converting the prevalence to incidence instead shows the laboratory-based incidence to be an overestimation.

Earlier studies have commented on the questionable accuracy of this formula for converting incidence and prevalence data.⁴⁻⁶ There are important arguments for the poor performance of the conversion. First, the types of data being compared represent different diagnostic criteria. Second, reflecting the argument put forward by Hoare et al,⁵ patients who contract HCA UTI are generally older and have comorbidities. Therefore, using their time in the hospital from diagnosis to discharge to reflect duration of infection is unreasonable.

Using antibiotic treatment as a proxy for infection duration greatly improved the comparability of the prevalence and incidence values, and we would recommend amending this formula to the following: $I = P \times (LA/\Delta Rx)$ and $P = I \times (\Delta Rx/LA)$, in which *I* is incidence, *P* is prevalence, LA is mean length of stay for all patients, and ΔRx is the duration of infection treatment.

ACKNOWLEDGMENTS

Financial support. The UK Clinical Research Collaboration funds the National Centre for Infection Prevention and Management at Imperial College London, which is also supported by the UK National Institute for Health Research Biomedical Research Centre funding scheme. The Dr Foster Unit at Imperial College London is largely funded by a research grant from Dr Foster Intelligence (an independent health service research organization) and is affiliated with the Centre for Patient Safety and Service Quality at Imperial College Healthcare National Health Service Trust, funded by the National Institute for Health Research.

Potential conflicts of interest. All authors report no conflicts of interest relevant to this article. All authors submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest, and the conflicts that the editors consider relevant to this article are disclosed here.

Carina King, PhD, MSc;^{1,2} Paul Aylin, MBChB, FFPHM;^{2,3} Alison Holmes, MD, MPH^{2,4}

Affiliations: 1. Institute of Global Health, University College London, United Kingdom; 2. The National Centre for Infection Prevention and Management, Imperial College, London, United Kingdom; 3. Dr Foster Unit at Imperial, School of Public Health, Imperial College London, London, United Kingdom; 4. Infection Prevention and Control, Imperial College Healthcare National Health Service Trust, London, United Kingdom.

Address correspondence to Carina King, Institute for Global Health, 30 Guilford Street, London, WC1 1NE, United Kingdom (c.king@ucl.ac.uk). Infect Control Hosp Epidemiol 2014;35(11):1432-1433

© 2014 by The Society for Healthcare Epidemiology of America. All rights reserved. 0899-823X/2014/3511-0020\$15.00. DOI: 10.1086/678435

REFERENCES

- 1. Rhame FS, Sudderth WD. Incidence and prevalence as used in the analysis of the occurrence of nosocomial infections. *Am J Epidemiol* 1981;113(1):1–11.
- 2. European Centre for Disease Prevention and Control (ECDC). Point prevalence survey of healthcare-associated infections (HAI) and antimicrobial use in European acute care hospitals–protocol version 4.3. Stockholm: ECDC, 2012.
- Health Protection Agency. English national point prevalence survey of healthcare-associated infections and antimicrobial use, 2011: preliminary data. London: Health Protection Agency, 2012.
- Gastmeier PMD, Bräuer H, Sohr DP, et al. Converting incidence and prevalence data of nosocomial infections: results from eight hospitals. *Infect Control Hosp Epidemiol* 2001;22(1):31–34.
- Haore HG, Muller A, Daniel Talon P, Xavier Bertrand P. Estimation of the cumulative incidence of hospital-acquired bacteremia from prevalence data: a formula. *Infect Control Hosp Epidemiol* 2005;26(4):415–417.
- 6. Berthelot P, Garnier F, Fascia P, et al. Conversion of prevalence survey data on nosocomial infections to incidence estimates: a simplified tool for surveillance? *Infect Control Hosp Epidemiol* 2007;28:633–636.

Redefining the National Healthcare Safety Network's Definition of Catheter-Associated Urinary Tract Infections: The Hazard of Including *Candida* Species

To the Editor—Catheter-associated urinary tract infections (CAUTIs) are a complication of urinary catheterization in hospitalized patients and represent a threat to patient safety.¹ CAUTI surveillance is a key tool to track the prevalence of this complication and measure the impact of prevention interventions. One key component of an effective surveillance program is valid measures of infection. *Candida* species are commonly isolated from the urine of hospitalized patients and may not represent urinary tract infection.^{2,3} We believe including *Candida* species as part of the National Healthcare Safety Network (NHSN) surveillance definition introduces

inaccuracy into CAUTI surveillance efforts, making CAUTI surveillance measures imprecise and improvement efforts more difficult to assess.

As an example, at our medical center, an otherwise healthy 28-year-old woman presented with 1 week of productive cough, dyspnea, fevers, myalgias, and diarrhea after frequenting a steam room at her local gym. At admission to the hospital, she was intubated for hypoxemic respiratory failure with septic shock. Computed tomography of the chest revealed extensive, multilobar consolidations with moderate pleural effusions. Blood and sputum cultures obtained at admission showed no growth; however, a urine specimen had antigen test results positive for *Legionella*. The patient remained critically ill and febrile during the first week of hospitalization, but her condition gradually improved.

At admission, the patient had a urinary catheter inserted. On the fourth day of hospitalization, because of the patient's persistent fever, a urine culture was obtained that grew *Candida albicans*. Although both the infectious diseases consult service and the intensive care unit team attributed the patient's fever to severe *Legionella* pneumonia and did not treat the *Candida* colonization, the patient met the NHSN criteria for CAUTI.⁴

In 2013, the NHSN changed the definition of CAUTI such that fever is no longer allowed to be attributed to another cause in patients with a urinary catheter and positive urine culture.⁴ Although this change increases the sensitivity of diagnosing CAUTIs for surveillance purposes, the specificity is substantially reduced, as in the case described above, and is less clinically relevant. Catheter-associated bacteriuria and funguria are common (5%–8% daily and 100% by 2 weeks), and national guidelines require that other etiologies of findings such as fever be ruled out before attributing them to a CAUTI and initiating treatment.^{1,2}

As Fakih et al⁵ and others have pointed out, the definitional change has serious financial implications for hospitals, which are mandated to report CAUTI rates, because Medicare reimbursement is affected by these data. Additionally, physicians may be more likely to inappropriately prescribe antibiotics to patients who, in reality, have asymptomatic bacteriuria but are now classified as having a CAUTI by the new NSHN criteria.⁶

To further illustrate the problematic nature of this new definition in terms of *Candida* species, following the 2013 definitional change, 40% of CAUTIs in our hospital were attributed to *Candida*. This is well above the nationally reported rates of candiduria (3%–32%), and the majority of these cases likely represent asymptomatic candiduria in patients who had alternative explanations for their fever (eg, sepsis from another source, neurologic injury, and trauma).²

We suggest that the NHSN revise the current CAUTI definition by removing *Candida* from the list of potential pathogens causing CAUTIs. Although there may be some true CAUTIs related to *Candida*, these are likely uncommon, and exclusion of these cases will have little impact on overall CAUTI rates. The current definitions threaten to create a system in which more than one-third of CAUTIs represent organisms of dubious pathogenicity in the urinary tract and undermine the validity of a tool that is critical for ongoing efforts to improve patient safety among hospitalized persons.

ACKNOWLEDGMENTS

Potential conflicts of interest. All authors report no conflicts of interest relevant to this article. All authors submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest, and the conflicts that the editors consider relevant to this article are disclosed here.

Darcy Wooten, MD, MS;¹ Jenifer Ramsey, RN, BSN, PHN;² Loren G. Miller, MD, MPH^{1,2}

Affiliations: 1. Division of Infectious Diseases, Harbor–University of California, Los Angeles (UCLA) Medical Center, Los Angeles, California; 2. Division of Hospital Epidemiology, Harbor-UCLA Medical Center, Los Angeles, California.

Address correspondence to Darcy Wooten, MD, MS, Infectious Diseases Fellow, David Geffen School of Medicine at the University of California, Los Angeles (UCLA), Division of Infectious Diseases, Harbor-UCLA Medical Center, 1000 West Carson Street, Box 466, Torrance, CA 90509 (darcy .wooten@gmail.com).

Infect Control Hosp Epidemiol 2014;35(11):1433-1434

© 2014 by The Society for Healthcare Epidemiology of America. All rights reserved. 0899-823X/2014/3511-0021\$15.00. DOI: 10.1086/678431

REFERENCES

- Hooton TM, Bradley SF, Cardenas DD, et al. Diagnosis, prevention, and treatment of catheter-associated urinary tract infection in adults: 2009 International Clinical Practice Guidelines from the Infectious Diseases Society of America. *Clin Infect Dis* 2010; 50(5):625–663.
- Kauffman CA. Diagnosis and management of fungal urinary tract infection. *Infect Dis Clin North Am* 2014;28(1):61–74.
- Al-Qas Hanna F, Sambirska O, Iyer S, Szpunar S, Fakih MG. Clinician practice and the National Healthcare Safety Network definition for the diagnosis of catheter-associated urinary tract infection. *Am J Infect Control* 2013;41(12):1173–1177.
- Centers for Disease Control and Prevention. National Healthcare Safety Network surveillance for urinary tract infections. http:// www.cdc.gov/nhsn/acute-care-hospital/CAUTI/index.html. Published 2013. Accessed June 2014.
- Fakih MG, George C, Edson BS, Goeschel CA, Saint S. Implementing a national program to reduce catheter-associated urinary tract infection: a quality improvement collaboration of state hospital associations, academic medical centers, professional societies, and governmental agencies. *Infect Control Hosp Epidemiol* 2013;34(10):1048–1054.
- Fakih MG, Greene MT, Kennedy EH, et al. Introducing a population-based outcome measure to evaluate the effect of interventions to reduce catheter-associated urinary tract infection. *Am J Infect Control* 2012;40(4):359–364.