Treatment of Asymptomatic Bacteriuria Might Be Harmful

Florian M. E. Wagenlehner¹ and Kurt G. Naber²

¹Clinic for Urology, Pediatric Urology and Andrology, Justus-Liebig-University, Giessen, and ²Technical University of Munich, Germany

(See the Major Article by Cai et al on pages 1655-61.)

Keywords. asymptomactic bacteriuria; recurrent urinary tract infections; prophylaxis.

In this issue of Clinical Infectious Diseases, Cai et al present a follow-up study [1] to their previously published prospectively randomized cohort study in which female patients with recurrent urinary tract infection (UTI) were followed clinically, but also investigated microbiologically at regular visits up to 1 year [2]. One group (A) was not treated, and the other group (B) was treated if asymptomatic bacteriuria was diagnosed, with the result that more symptomatic UTIs occurred in group B than in group A. The present study started immediately after the end of the first study. The patients remained in their groups and were followed every 6 months up to about 3 years. However, in the follow-up study [1], patients received antibiotic therapy only in case of a symptomatic UTI. Nevertheless, group B experienced statistically significantly more symptomatic UTIs

Clinical Infectious Diseases[®] 2015;61(11):1662–3 © The Author 2015. Published by Oxford University Press on behalf of the Infectious Diseases Society of America. This is an Open Access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs licence (http://creativecommons.org/licenses/by-nc-nd/4.0/), which permits non-commercial reproduction and distribution of the work, in any medium, provided the original work is not altered or transformed in any way, and that the work is properly cited. For commercial re-use, please contact journals.permissions@oup. com.

DOI: 10.1093/cid/civ698

than group A, although the therapeutic strategy was the same. In addition, the resistance rates of isolated *Escherichia coli* against amoxicillin-clavulanic acid, cotrimoxazole, and ciprofloxacin were significantly higher in group B than in group A. This finding is especially interesting, as these antibiotics were used only in a few cases during the follow-up study.

In 2 previous studies, continuous antibiotic prophylaxis with cotrimoxazole was compared to prophylaxis with cranberry [3] or lactobacilli [4]. Both studies showed higher resistance rates of commensal E. coli to cotrimoxazole in urine and feces in the antibiotic arms compared with the nonantibiotic arms. Thus, the question arises whether the higher antibiotic resistance in the studies by Cai et al occurred already during the first study [2], because in the current study [1], both groups were treated with the same antibiotic strategy. Considering both studies together, a somewhat higher antibiotic consumption was found in group B, which apparently translated also into higher antibiotic resistance rates of the urinary pathogens.

The results of all these studies confirm the current guidelines [5] to preferably incorporate nonantibiotic strategies for prevention of these very frequent, but generally benign infections, if these strategies are confirmed to be effective in well-designed clinical studies [6]. Such strategies may also be important to decrease the general antibiotic consumption in the population and thus to slow down emergence of antibiotic resistance; as shown in an interventional comparative study, antibiotic resistance, once established, has a low probability to be reversed, at least for trimethoprim and cotrimoxazole [7].

Note

Potential conflict of interest. F. M. E. W. has received grants from Deutsche Forschungsgemeinschaft and Deutsches Zentrum für Infektionsforschung, and personal fees from Achaogen, Astellas, AstraZeneca, Bionorica, Cubist/MSD, Leo-Pharma, Medpace, Merlion, Vifor Pharma, Rempex, and Rosen Pharma. K. G. N. has received personal fees from Basilea, Bionorica, Boehringer Ingelheim, Cubist/MSD, Daiichi Sankyo, Enteris Biopharm, Galenus, Helperby, Leo Pharma, Melinta, MerLion, OM Pharma/Vifor, Paratek, Pierre Fabre, Rempex Pharma, Shionogi, and Zambon.

Both authors have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest. Conflicts that the editors consider relevant to the content of the manuscript have been disclosed.

References

- Cai T, Nesi G, Mazzoli S, et al. Asymptomatic bacteriuria treatment is associated with higher prevalence of antibiotic resistant strains in women with urinary tract infections. Clin Infect Dis 2015; 61:1655–61.
- Cai T, Mazzoli S, Mondaini N, et al. The role of asymptomatic bacteriuria in young women with recurrent urinary tract infections: to treat or not to treat? Clin Infect Dis 2012; 55:771–7.
- Beerepoot MA, ter Riet G, Nys S, et al. Cranberries vs antibiotics to prevent urinary tract infections: a randomized double-blind noninferiority trial in premenopausal women. Arch Intern Med 2011; 171:1270–8.

Received 13 July 2015; accepted 18 July 2015; electronically published 12 August 2015.

Correspondence: Kurt G. Naber, MD, PhD, Assoc. Professor of Urology, Technical University of Munich, Karl Bickleder Strasse 44c, Straubing, D-94315, Germany (kurt@nabers.de).

- Beerepoot MA, ter Riet G, Nys S, et al. Lactobacilli vs antibiotics to prevent urinary tract infections: a randomized, double-blind, noninferiority trial in postmenopausal women. Arch Intern Med 2012; 172:704–12.
- 5. Grabe M, Bjerklund Johansen TE, Cai T, et al. EAU guidelines on urological infections. Part

 pp 1–78. In: European Association of Urology, ed. Arnhem, The Netherlands, 2015. Available at: http://uroweb.org/guideline/ urological-infections/. Accessed 12 July 2015.
Beerepoot MA, Geerlings SE, van Haarst EP,

van Charante NM, ter Riet G. Nonantibiotic prophylaxis for recurrent urinary tract infections: a systematic review and meta-analysis of randomized controlled trials. J Urol **2013**; 190:1981–9.

 Sundqvist M, Geli P, Andersson DI, et al. Little evidence for reversibility of trimethoprim resistance after a drastic reduction in trimethoprim use. J Antimicrob Chemother **2010**; 65:350–60.