Escherichia coli bacteriuria in female adults is associated with the development of hypertension


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1. Introduction

Although the urinary tract is normally sterile, asymptomatic bacteriuria is a common phenomenon, especially in women. Different studies report a prevalence of asymptomatic bacteriuria of approximately 5% among healthy young women, increasing to over 20% in the elderly.1,2 Escherichia coli is the most prevalent uropathogen.3 Several authors in the first half of the twentieth century have suggested a role of bacteriuria in the etiology of hypertension.4 Although more recent studies have also found a correlation, no prospective study has convincingly shown that bacteriuria itself leads to hypertension.5

The present study aimed to address the question of whether or not Escherichia coli bacteriuria is associated with an increased risk of the development of hypertension during a long-term follow-up period.

In addition, the risk of a heart attack or stroke in the presence of bacteriuria was studied.

2. Methods

2.1. Study population

A full cohort analysis was performed for women who participated in two population-based studies. Between 1974 and 1986, all women born between 1911 and 1945, who lived in Utrecht, the Netherlands, were invited to participate in a breast cancer screening program. The participants completed a questionnaire, underwent a medical examination, and collected a morning urine sample that remained stored. From 1993 to 1997 another population-based study was performed. We performed a full cohort analysis for 444 women who participated in both studies. E. coli bacteriuria was diagnosed by a real-time PCR. Hypertension was defined as the use of antihypertensive medication and/or a measured systolic blood pressure of at least 160 mmHg or a diastolic blood pressure of 95 mmHg or higher. The mean follow-up was 11.5 ± 1.7 years.

Results: Forty women (9%) had E. coli bacteriuria at baseline. Women who had bacteriuria at baseline had a mean blood pressure at study endpoint of 133 ± 20 mmHg systolic and 78 ± 11 mmHg diastolic, and women without bacteriuria had values of 129 ± 20 and 78 ± 11 mmHg, respectively (p-values for difference 0.33 and 0.88). Although E. coli bacteriuria was not associated with the blood pressure as a continuous variable, it was associated with the development of hypertension during follow-up (OR 2.8, 95% CI 1.4–5.5).

Conclusion: E. coli bacteriuria may increase the risk of future hypertension.
urine was stored in plastic polypropylene jars, without preserving agents, and stored at −20 °C for future analyses. All women gave oral consent to use their data and urine samples for future scientific research.

From 1993 to 1997, participants in the breast cancer screening program received an invitation by mail to join an additional study to assess the relationship between nutrition and cancer and other chronic diseases, the Prospect-EPIC study (the follow-up cohort). A total of 17 357 women living in Utrecht and the surrounding area agreed to take part (participation rate 34.5%). Participants were between 49 and 70 years of age at enrolment. Information was collected on the basis of two self-administered questionnaires and a medical examination including blood pressure. Non-fasting blood samples were successfully drawn from 97.5% of the women and stored under liquid nitrogen at −196 °C. The serum creatinine level was later measured from these samples. Approximately 88% of the women signed a detailed informed consent, enabling the researchers to use their blood samples for future analysis, and to obtain information on future morbidity and mortality.

In total, 506 women participated in both the baseline cohort and the follow-up cohort. Sixty-two women had to be excluded for the following reasons: a missing urine sample (n = 13), the use of antihypertensive medication at baseline (n = 45), kidney transplantation during follow-up (n = 1), or missing data on hypertension at study endpoint (n = 3). Finally, 444 women were included in the prospective study to assess the relationship between bacteriuria and the development of hypertension. The mean duration of follow-up was 11.5 ± 1.7 years, ranging from 8.3 to 18.6 years from baseline until participation in the follow-up study (and was not different for women with compared to women without bacteriuria).

This study was approved by the Medical Ethics Committee of the University Medical Center Utrecht, the Netherlands.

2.2. Escherichia coli bacteriuria

E. coli bacteriuria was defined as the presence of 10^5 colony forming units (cfu) of E. coli per ml of urine. It was diagnosed by a real-time PCR that we developed and validated beforehand, a technique that has also been used by others. Briefly, PCR primers and probe complementary to regions of the gapA gene specific for E. coli were designed for the real-time PCR assay. The laboratory sensitivity and specificity of the real-time PCR tested with 50 E. coli strains and with 41 non-E. coli strains (including the most prevalent uropathogens and many members of the vaginal and anal flora) were 100% (50/50) and 98% (40/41), respectively. For clinical evaluation, 42 clinical urine specimens (12 with and 30 without E. coli) were tested and the results were compared to those of a clinical conventional urine culture. The sensitivity and specificity of the real-time PCR in these clinical samples were 92% and 87%, respectively. The test results were quantitative and allowed distinguishing between significant bacteriuria (i.e., 10^5 cfu/ml) and low-count bacteriuria that might have been due to contamination.

To test a study urine sample, 1 ml of urine was centrifuged at 16 250 × g for 5 min. The pellet was washed twice, suspended in 1 ml of sterile injection water, and 100 μl of the suspension was heated for 2 min at 1000 Watt in a microwave oven for DNA preparation. Five microliters was added to the real-time PCR reaction volume as DNA template. Each 25 μl reaction volume consisted of 12.5 μl 2 × TaqMan Universal PCR Master Mix (Applied Biosystems, Branchburg, New Jersey, USA) that contains AmpliTaq Gold DNA polymerase, 300 nM forward primer (5′-ACCCACATCGTGTGATGC-3′), 300 nM reverse primer (5′-AGCAA-CAGTTGAAAGTCCA-3′), and 175 nM probe (5′-CAT-TATGTCGTCTCCCGGTCTCA-3′); the DNA template was the last ingredient added. The ABI PRISM 7700 Sequence Detection System (PE Biosystems, Nieuwerkerk aan de IJssel, the Netherlands) was used for the real-time PCRs. Cycling parameters were first the uracil-N-glycosylase (UNG) reaction at 50 °C for 2 min, then AmpliTaq Gold activation at 95 °C for 10 min, followed by 45 cycles of denaturation at 95 °C for 15 s – combined annealing and extension at 60 °C for 1 min. Emitted fluorescence from each well was measured during both the denaturation and annealing/extension steps in every cycle. Amplification plots were constructed using the ABI PRISM 7700 Sequence Detection System software, version 1.7 (PE Biosystems).

2.3. Blood pressure, heart attack, stroke

Blood pressure was measured with a standard mercury sphygmomanometer after the subject had been seated for 5 min. Hypertension was defined as the previous use of antihypertensive medication (assessed at follow-up by the question: “Have you ever been treated with drugs for high blood pressure?”) and/or a measured systolic blood pressure of at least 160 mmHg or a diastolic blood pressure of 95 mmHg or higher. A history of having had a heart attack or stroke was assessed at follow-up by the two additional questions: “Have you ever had a heart attack?”, “Have you ever had a stroke?”.

2.4. Data analysis

The present results are based on a cohort study of 444 women who were followed for the development of hypertension in relation to E. coli bacteriuria at baseline.

Baseline characteristics were compared between women with and without bacteriuria. Comparisons between means were performed with the Student’s t-test and comparisons between nominal or categorical data with the Chi-square test. Linear regression analysis was used to calculate the adjusted difference in blood pressure between women with versus women without bacteriuria. The relative risk of hypertension in the presence of bacteriuria was estimated by logistic regression and quantified as odds ratios (OR) and 95% confidence intervals (CI). Adjustment was done for potential confounding factors, i.e., age, weight, and serum creatinine level.

Table 1

<table>
<thead>
<tr>
<th>Age, years</th>
<th>Body mass index, kg/m²</th>
<th>Postmenopausal</th>
<th>Married or living with partner</th>
<th>Given birth to living child(ren)</th>
<th>Antibiotics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total study group (N = 444)</td>
<td>44.9 ± 3.2</td>
<td>24.0 ± 3.2</td>
<td>98 (22%)</td>
<td>303 (90%)</td>
<td>401 (90%)</td>
</tr>
<tr>
<td>No. E. coli bacteriuria (n = 404)</td>
<td>44.9 ± 3.2</td>
<td>23.9 ± 3.2</td>
<td>88 (22%)</td>
<td>273 (90%)</td>
<td>363 (90%)</td>
</tr>
<tr>
<td>E. coli bacteriuria (n = 40)</td>
<td>45.3 ± 3.4</td>
<td>24.6 ± 3.6</td>
<td>10 (25%)</td>
<td>30 (97%)</td>
<td>38 (95%)</td>
</tr>
<tr>
<td>p-Value</td>
<td>0.51</td>
<td>0.26</td>
<td>0.64</td>
<td>0.20</td>
<td>0.29</td>
</tr>
</tbody>
</table>

Values are given as mean ± standard deviation or as number of patients (percentage).

* The total number of study subjects is 447 unless otherwise stated.
The limitations include the fact that we had to rely on only one urine sample to define bacteriuria. However we, and others, have validated this before. We made the assumption that however bacteriuria might be transient in a proportion of the bacteriuric study subjects, bacteriuria at one point reflects a higher susceptibility to recurrent and persistent bacteriuria in general, even after antimicrobial therapy. Previous findings are supportive of this assumption. Some of the urine samples may have become contaminated before storage, however this will be equally divided in the total study group. Moreover, contamination usually leads to the growth of more pathogens, often non-E. coli, with lower colony counts, which are not picked up by our real-time PCR. E. coli is the causative microorganism found to be most prevalent, and the prevalence of E. coli bacteriuria of 9% among middle-aged women reported here is in the range of what could be expected beforehand. There are no published data that can be cited to support the test characteristics of real-time PCR on urine stored for more than 10 years. However, it seems plausible that the gadA gene for E. coli should still be present also after a long time period. We did test urine samples that were stored for up to 5 years and compared the results to those of the conventional urine culture that was performed at the time the samples were fresh (unpublished data). The results were similar to those from the fresh samples. The baseline data included the use of any drugs, which allowed us to exclude women who used antihypertensive medication. However, blood pressure was not measured at baseline, and therefore the study cohort will include some women with undiagnosed hypertension. At follow-up we had to rely on a single blood pressure measurement. But we assume that the incidence of increased blood pressure due to other reasons will be equally divided among women with and without bacteriuria at baseline. Moreover, the increased prevalence of hypertension in the group of bacteriuria was mainly due to more women who started antihypertensive drugs in this group compared to the group of bacteriuria. Unfortunately, data on the prevalence of diabetes mellitus at baseline were lacking, and therefore adjustment for diabetes was not performed.

Several authors in the first half of the twentieth century have suggested a role of bacteriuria in the etiology of hypertension, as reviewed before. For instance, Kass showed small differences in blood pressure between bacteriuric and non-bacteriuric women aged 15 to 64 years. Although more recent studies have also found a correlation, no prospective study has convincingly shown that bacteriuria itself leads to hypertension. In our cohort study, we found a higher prevalence of hypertension in the bacteriuric group after 12 years of follow-up. The underlying mechanism of this finding is not clear. Hypertension is a lasting increase in blood pressure with a heterogeneous etiology consisting of both genetic and environmental factors. Patients share the inability to excrete sodium at a normal arterial pressure. If bacteriuria leads to hypertension, the most attractive explanation would be that hypertension arises secondary to renal scarring caused by the type 1 fimbriae of the uropathogens. In the multivariate analysis, correction for creatinine did not change the results, but hypertension can occur before the reduction in creatinine clearance becomes apparent (for example in chronic glomerulonephritis). The absence of a change in odds ratio after adjustment for age, weight, and creatinine confirms the notion that these factors were not related to bacteriuria. An alternative explanation is that both bacteriuria and hypertension are found more frequently among individuals with co-morbidity or that they share a same (currently
unknown) cause. This is supported by the higher prevalence of bacteriuria among women who used antihypertensive drugs at baseline. It remains unclear why *E. coli* bacteriuria was associated with the development of hypertension but not with the blood pressure as a continuous variable. However, it seems plausible that part of it is due to the success of antihypertensive drugs.

The percentage of women who developed hypertension during follow-up was in agreement with what could be expected. A large population-based study performed in the Netherlands also between 1993 and 1997, showed a prevalence of hypertension in women in the same age group of 15%.17

In conclusion, in this prospective study a strong correlation was found between *E. coli* bacteriuria and hypertension after 12 years of follow-up. Given the importance of hypertension the nature of this correlation needs to be studied further.

Conflict of interest: No conflict of interest to declare.

Acknowledgement

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References