A Systematic Evaluation of Factors Associated With Nocturia-The Population-based FINNO Study

Kari A. O. Tikkinen, HUCH Ltd
Anssi Auvinen, Tampere University Hospital
Theodore Johnson II, Emory University
Jeffrey P. Weiss, SUNY Downstate Medical School
Tapani Keraenen, Tampere University Hospital
Aila Tiitinen, Helsinki University Hospital
Olli Polo, Tampere University Hospital
Markku Partinen, Helsinki University Hospital
Teuvo L. J. Tammela, Tampere University Hospital

Proceedings Title: American Journal Of Epidemiology
Conference Name: 24th Congress of the Annual European-Association-of-Urology
Publisher: Oxford University Press Inc
Conference Place: Stockholm, Sweden
Volume/Issue: Volume 170 | Issue 3
Publication Date: 2009-08-01
Type of Work: Conference | Final Publisher PDF
Publisher DOI: 10.1093/aje/kwp133
Permanent URL: https://pid.emory.edu/ark:/25593/s6nh7

Final published version: http://dx.doi.org/10.1093/aje/kwp133

Copyright information:
American Journal of Epidemiology © 2009 The Authors
This is an Open Access work distributed under the terms of the Creative Commons Attribution NonCommercial License 2.0 Germany License (http://creativecommons.org/licenses/by-nc/2.0/de/).

Accessed April 7, 2018 5:20 AM EDT
Original Contribution

A Systematic Evaluation of Factors Associated With Nocturia—The Population-based FINNO Study

Kari A. O. Tikkinen, Anssi Auvinen, Theodore M. Johnson II, Jeffrey P. Weiss, Tapani Keränen, Aila Tiitinen, Olli Polo, Markku Partinen, and Teuvo L. J. Tammela

Initially submitted December 8, 2008; accepted for publication April 29, 2009.

In a case-control study with prevalence sampling, the authors explored the correlates for nocturia and their population-level impact. In 2003–2004, questionnaires were mailed to 6,000 subjects (aged 18–79 years) randomly identified from the Finnish Population Register (62.4% participated; 53.7% were female). Questionnaires contained items on medical conditions, medications, lifestyle, sociodemographic and reproductive factors, urinary symptoms, and snoring. Nocturia was defined as ≥2 voids/night. In age-adjusted analyses, factors associated with nocturia were entered into a multivariate model. Backward elimination was used to select variables for the final model, with adjustment for confounding. Although numerous correlates were identified, none affected ≥50% of nocturia cases of both sexes. The factors with the greatest impact at the population level were (urinary) urgency (attributable number/1,000 subjects (AN) = 24), benign prostatic hyperplasia (AN = 19), and snoring (AN = 16) for men and overweight and obesity (AN = 40), urgency (AN = 24), and snoring (AN = 17) for women. Moreover, correlates included prostate cancer and antidepressant use for men, coronary artery disease and diabetes for women, and restless legs syndrome and obesity for both sexes. Although several correlates were identified, none accounted for a substantial proportion of the population burden, highlighting the multifactorial etiology of nocturia.

coronary artery disease; diabetes mellitus; lifestyle; obesity; prostatic hyperplasia; prostatic neoplasms; sleep disorders; urinary bladder, overactive

Abbreviations: BPH, benign prostatic hyperplasia; FINNO, Finnish National Nocturia and Overactive Bladder.

Nocturia (waking at night to urinate) is a common cause of awakenings and may lead to sleep maintenance insomnia (1–3). Nocturia can be bothersome (4) and is associated with impaired mental and somatic health (5), impaired quality of life (4), and even increased mortality (6).

The etiology of nocturia is inadequately understood. Nocturia is frequently attributed to aging or childbirth in women and to benign prostatic hyperplasia (BPH) in men. Other related conditions include overactive bladder syndrome; nocturnal polyuria; obstructive sleep apnea; awakening for other reasons such as anxiety; primary sleep disorders; and use of diuretics, caffeine, or alcohol (7–9). Bedtime fluid intake correlates poorly with nocturia episodes (10).

Nocturia persists frequently following simple prostatectomy (11). In one study, 38% of men reported ≥2 voids/night 3 years after transurethral resection of the prostate (12). Several pharmacologic approaches have yielded limited nocturia reductions or significant side effects (7, 13–25).

To our knowledge, no earlier study comprehensively covered the possible associations of medical conditions; medications; lifestyle; and anthropometric, reproductive, and sociodemographic factors with nocturia. We explored correlates for nocturia and assessed their population-level impact in a large population-based study.

Correspondence to Dr. Kari A. O. Tikkinen, Clinical Research Institute HUCH Ltd./Tutkijatilat H3011, Haartmaninkatu 4, P.O. Box 105, FI-00029 HUS, Helsinki, Finland (e-mail: kari.tikkinen@fimnet.fi).
MATERIALS AND METHODS

Finnish National Nocturia and Overactive Bladder (FINNO) Study

In 2003–2004, questionnaires were mailed to 6,000 subjects aged 18–79 years randomly identified from the Population Register Centre. Age stratification was used with oversampling of younger age groups to ensure precise estimates even in age groups experiencing lower nocturia frequency (3, 26). More information on the FINNO Study (including characterization of nonrespondents) has been published previously (3, 26, 27).

Outcome

Nocturia cases were defined as subjects reporting 2 or more voids/night because this frequency involves clinically significant bother (4). All subjects without nocturia were considered controls in unconditional logistic regression analyses. Self-reported nocturia frequency was determined by using a previously described algorithm (27) combining responses to the Danish Prostatic Symptom Score (28) and the American Urological Association Symptom Index (29) (Web Table 1; this information is described in the first of 4 supplementary tables, each referred to as “Web table” in the text and posted on the Journal’s website (http://aje.oupjournals.org/)).

Correlate assessment

Self-reported information on physician-diagnosed conditions, prescribed medication, specific symptoms, and lifestyle factors was obtained by using questions modified from surveys conducted by the National Public Health Institute (30). Comorbidity indicators were formulated for 36 conditions deemed common or previously hypothesized as determinants of lower urinary tract symptoms (Web Tables 1 and 2). Medication use was classified into 27 groups by using the Anatomical Therapeutic Chemical Classification (31) (Web Tables 1 and 3). Lifestyle factors included body mass index, smoking, and alcohol and coffee consumption (Web Tables 1 and 4). The Danish Prostatic Symptom Score questions were used to evaluate (urinary) urgency and stress urinary incontinence (28) and the Basic Nordic Sleep Questionnaire to evaluate snoring (32) (Web Tables 1 and 4). These symptoms have been specifically shown to be risk factors for nocturia (26, 33, 34).

With the exception of the alcohol consumption question (response rate: men, 86%; women, 76%), data on potential correlates were highly available (men, 97%–100%; women, 95%–100%). Alcohol consumption was not associated with nocturia.

Potential confounders

Age, sociodemographic factors (marital status, education, employment, urbanization) (3), and female reproductive/gynecologic factors (parity, postpartum period, menopausal status, hormone therapy, hysterectomy, stress urinary incontinence surgery) (27) were treated as potential confounders.

Data on urbanization, parity, and delivery date(s) were obtained from the population register. Information on each potential confounder was available for at least 99% of subjects.

Statistical analysis

Logistic regression was used for the analyses stratified by sex, with nocturia as the outcome. All potential correlates and confounders associated (P < 0.10) with nocturia in the age-adjusted analyses (basic analysis population) were entered into the multivariate model (Web Figure 1, also posted on the Journal’s website (http://aje.oupjournals.org/)). In these analyses, potential correlates and confounders numbered 16 and 2 for men and 18 and 6 for women, respectively. Backward elimination techniques were used to select variables for the final model, with likelihood ratio tests used to determine significance (P < 0.05). Finally, cofactors that were not actual confounders (they did not change any estimate by ≥10%) were eliminated. Confirmed confounders were age for men, menopausal status for women, and employment for both sexes. The final analysis included subjects with information on nocturia, correlates, and confounders (Web Figure 1). For the correlates identified, age-standardized sensitivity, positive predictive value, attributable fraction in the exposed, population attributable fraction, and attributable number were calculated (35).

RESULTS

Of the 6,000 subjects approached, 3,727 (62.4%) participated; 23 were unavailable, and 130 were excluded because of pregnancy, puerperium, or urinary tract infection (Web Figure 1). (The response rate was approximately 32% after the first round, 50% after the second round, and 62.4% after the final, third round.) Of the 3,597 subjects included, 98% provided nocturia information (basic analysis population). The 3,307 subjects (92%) who responded to all nocturia, correlate, and confounder questions formed the final analysis population. Prevalence of nocturia was 12.5% (95% confidence interval: 10.7, 14.3) among men and 12.9% (95% confidence interval: 11.0, 14.9) among women (age standardized to match Finland’s age structure (36)). Excluding subjects with missing information on any correlate or confounder (final analysis population) did not change these estimates. For more detail, refer to Table 1 and to Web Tables 2–4. Correlates for nocturia included (urinary) urgency, snoring, restless legs syndrome, and obesity for both sexes; BPH, antidepressant use, and prostate cancer for men; and overweight, diabetes, and coronary artery disease for women (Table 1).

At the population level (Table 2), urgency and snoring (both sexes), BPH (men only), and overweight/obesity (women only) accounted for the largest proportion of nocturia. At the individual level, the strongest correlate for both sexes was urgency, although odds ratio differences between correlates were mainly statistically nonsignificant (Table 1).

No correlate affected 50% or more of men with nocturia (Figure 1): BPH, urgency, and snoring had the highest sensitivity for nocturia (31%–49%). Of the women with nocturia, 71% were overweight or obese; other correlates were reported by 50% or more of women with nocturia (Figure 2).
A majority of men with prostate cancer or urgency reported nocturia, yielding positive predictive values of 74% and 59%, respectively. Half of the men with BPH, and a minority of men with other correlates, reported nocturia (Figure 1). Among women, no correlates were associated with a 50% or greater probability of nocturia (Figure 2).

Generally, questionnaire mailing round did not affect correlate prevalence. However, 4 exceptions emerged (age adjusted). First-round responders reported more nocturia and urgency than responders in subsequent rounds ($P$ for trend $= 0.01$ for both nocturia and urgency; sexes combined). Moreover, first-round male responders reported more antidepressant use ($P$ for trend $= 0.04$), and first-round female responders were slightly less obese than those in subsequent rounds ($P$ for trend $= 0.05$). However, the odds ratio estimates for these factors were similar for each round, suggesting absence of systematic error.

**DISCUSSION**

In this large, population-based study, numerous factors associated with nocturia were identified. However, no single correlate accounted for more than half of the cases of nocturia, highlighting its multifactorial etiology. At the population level, urgency, BPH, and snoring for men and overweight/obesity, urgency, and snoring for women accounted for the largest proportion of nocturia.

Our FINNO Study population is representative of Finnish adults in terms of sociodemographic, anthropometric, and female reproductive factors (3, 27, 37). The frequencies of comorbidities, medications, and lifestyle factors were similar to those in large-scale population surveys conducted by the National Public Health Institute (38–40). Concordant with earlier reports (26, 41–44), urinary urgency was strongly associated with nocturia in both sexes, yet only 1 in 3 with nocturia reported urgency. Among men,
BPH had the second highest population impact, but only a third with nocturia reported BPH. Lower urinary tract symptoms suggestive of benign prostatic obstruction constitute a well-recognized risk factor for nocturia (45). However, the impact of BPH may be overestimated. In Japanese studies, nocturia was the lower urinary tract symptom least related to prostatic obstruction, and treatment to relieve obstruction had less of an effect on nocturia than on other symptoms (15, 46). In a lower urinary tract symptoms/BPH study, patients receiving doxazosin experienced very modest reductions in nocturia, whereas finasteride had no effect (23). In the current study, prostate cancer was associated with nocturia. More than 70% of men with prostate cancer reported at least 2 voids/night, yet only 7% of men with nocturia reported prostate cancer.

Concurring with earlier findings (37, 47–49), nocturia was associated with obesity in both sexes and, among women, also with overweight. Indeed, overweight/obesity had the greatest population impact among women.

Snoring had a strong population impact because of its high prevalence, yet the strength of association was relatively weak (odds ratio = 1.5–1.8). Snoring has been associated with nocturia (34). The severity of obstructive sleep apnea predicted nocturia frequency, and continuous positive airway pressure treatment decreased nocturia (50). In a home sleep study, the prevalence of obstructive sleep apnea was double among urology patients with nocturia compared with those without (51). In our study, reported obstructive sleep apnea was not associated with nocturia after adjustment, which may be due to correlation with snoring (three-quarters of subjects with obstructive sleep apnea reported snoring, and snoring was 10 times more prevalent than obstructive sleep apnea).

To our knowledge, no association has been reported between nocturia and restless legs syndrome. Increased nocturia among patients with restless legs syndrome may reflect sleep disturbance (52). Moreover, such patients use more medications (particularly antidepressants) than controls do (53). Nocturia has been previously linked to (untreated) depression (54, 55) and use of selective serotonin-reuptake inhibitors (54). In our study, only for men was nocturia associated with antidepressant use; depression itself was not associated with nocturia.

Diabetes and coronary artery disease were associated with nocturia in the age-adjusted analyses for both sexes but for only women in multivariate analysis. An association between diabetes and nocturia has been reported sometimes (43, 47, 49, 50, 56–60), but not always (10, 42, 45). In the BACH Survey (47), nocturia was associated with increasing body mass index, diabetes, and cardiac disease, whereas among Danes aged 60–80 years (49), increasing body mass index, diabetes, urinary incontinence, and recurrent cystitis were associated with 2 voids/night. In both surveys (47, 49), sex was used as a covariate, but results were not reported by gender. Some earlier reports (47, 59, 61), but not all (42, 43, 45, 49, 58), found cardiac/coronary disease a correlate for nocturia.

Coffee or alcohol consumption (10, 33, 41, 43, 49, 62, 63) and smoking (33, 49, 61, 63) have been shown elsewhere not to be associated with nocturia. Our findings were the same. Differences in our results from previous findings may be explained by differences in study procedures and samples (64). Several factors explored here were not assessed in earlier studies (10, 41–43, 45–47, 49–51, 55, 56, 58, 59, 61–63). In addition, several previous studies were not population based (3, 13, 43, 46, 50, 51, 59, 61). The association of numerous factors with nocturia in age-adjusted analyses,
partly differing by gender, highlights the importance of appropriate analysis, including controlling for confounders.

Given the multiple possible determinants (7–10, 34, 41–43, 45–51, 54, 56–63), we assessed numerous candidates. Because of inconsistencies in the literature, using existing evidence to choose the potential confounders was not justified. Hence, because of the exploratory nature of this analysis, we used stepwise methods for model building (65).

By our methodology, we avoided selection bias due to treatment seeking (reflecting both severity and health care service use). Our study’s strengths include 1) a representative sample of both sexes and all adult ages, 2) a high participation rate and completeness of questionnaire responses, 3) a large number of relevant factors, 4) systematic control for confounding, and 5) assessment of nocturia and related symptoms with validated instruments. Furthermore, determinant prevalences were largely similar by response round, indicating absence of selection bias; any trend in the risk estimates by response round was also lacking.

This study has some limitations. First, the validity of self-report has not been established for all characteristics we considered. Second, alcohol consumption reporting was incomplete, yet nocturia prevalence did not vary by alcohol consumption among those reporting this information. In addition, reported alcohol consumption was comparable with the national statistics (66). Third, we had no information on physical activity, although physical activity has not previously been related to nocturia (41). Finally, these results from the Finnish population may not be directly generalizable to other ethnicities because impact measures generally are context specific. There may be ethnic differences in the prevalence of nocturia. Socioeconomic status attenuated, but did not entirely remove, the effect of race/ethnicity on nocturia (67).

Nocturia has been classified as a symptom caused by 1) nocturnal polyuria, 2) low nocturnal bladder capacity, 3) diminished global bladder capacity, 4) a combination of nocturnal polyuria and low bladder capacity, 5) global polyuria, and/or 6) sleep disorders (7). We found several risk factors that may well cause these pathways and, finally, nocturia. However, the pathways are probably complex, and there may also be numerous other underlying causes for the associations, such as autonomic nervous system hyperactivity and/or metabolic syndrome (68, 69). At the population level, urgency, BPH, and snoring for men and overweight and obesity, urgency, and snoring for women explained the largest proportion of nocturia, whereas obesity, antidepressant use, and prostate cancer in men; diabetes and coronary artery disease in women; and restless legs syndrome in both sexes had less of an impact. Even though numerous correlates for nocturia were identified, none was associated with nocturia in more than half of the affected subjects of both sexes, highlighting the multifactorial etiology.

ACKNOWLEDGMENTS

Author affiliations: Department of Urology, Helsinki University Central Hospital, Helsinki, Finland (Kari A. O. Tikkinen); Clinical Research Institute HUCH Ltd., Helsinki, Finland (Kari A. O. Tikkinen); Department of Urology, Tampere University Hospital, Tampere, Finland (Kari A. O. Tikkinen, Teuvo L. J. Tammela); Medical School, University of Tampere, Tampere, Finland (Kari A. O. Tikkinen, Teuvo L. J. Tammela); School of Public Health, University of Tampere, Tampere, Finland (Anssi Auvinen); Birmingham/Atlanta VA Geriatric Research, Education, and Clinical Center, Decatur, Georgia (Theodore M. Johnson); Division of Geriatric Medicine and Gerontology, Emory University School of Medicine, Atlanta, Georgia (Theodore M. Johnson); Department of Urology, SUNY Downstate Medical School, Brooklyn, New York (Jeffrey P. Weiss); Division of Neurology and Rehabilitation, Tampere University Hospital, Tampere, Finland (Tapani Keränen); Department of Obstetrics and Gynecology, Helsinki University Central Hospital, Helsinki, Finland (Aila Tiitinen); Department of Pulmonary Diseases, Tampere University Hospital, Tampere, Finland (Olli Polo); Department of Neurology, University of Helsinki, Helsinki, Finland (Markku Partinen); and Vitalmed Research Center, Helsinki, Finland (Markku Partinen).

This work was funded by unrestricted grants from the Competitive Research Funding of the Pirkanmaa Hospital District (Tampere, Finland) and Pfizer Inc. (New York, New York). The work of the corresponding author was funded by unrestricted grants from the Emil Aaltonen Foundation and the Pirkanmaa Regional Fund of the Finnish Cultural Foundation. The funding sources had no role in the design and conduct of the study; the collection, management, analysis, and interpretation of the data; or the preparation, review, or approval of the manuscript.

The authors thank Dr. Aila Rissanen for her valuable comments on earlier versions of the manuscript and Virginia Mattila for language revision.

K. A. O. T. had full access to all the data used in the study and takes responsibility for the integrity and the accuracy of the data. K. A. O. T., A. A., and T. L. J. T. designed the study. K. A. O. T. collected the data. K. A. O. T. and A. A. performed statistical analysis of the data. All authors contributed to the interpretation of the findings. K. A. O. T. drafted the manuscript. All authors contributed to revising the manuscript.

These results were presented, in part, at the 103rd Annual Meeting of the American Urological Association, Orlando, Florida, May 17–22, 2008; the 2nd North Eastern European Meeting of the European Association of Urology, Vilnius, Lithuania, September 12–13, 2008; and the 24th Annual Meeting of the European Association of Urology, Stockholm, Sweden, March 17–21, 2009.

K. A. O. T., A. A., T. K., and A. T. declare no conflicts of interest. K. A. O. T. served as a speaker at a Pfizer-sponsored overactive bladder symposium, but his honorarium was donated to the Public Library of Science. T. M. J. is a consultant for Ferring (Saint-Prex, Switzerland), Johnson & Johnson (New Brunswick, New Jersey), and Pfizer. J. P. W. is a consultant for Ferring, Pfizer, and Watson Pharma (Salt Lake City, Utah). O. P. is a consultant for Actelion (Allschwil, Switzerland), AstraZeneca (London, United Kingdom), Boehringer-Ingelheim (Ingelheim, Germany),
Cephalon (Frazer, Pennsylvania), Eli Lilly (Indianapolis, Indiana), GlaxoSmithKline (London, United Kingdom), Lundbeck (Copenhagen, Denmark), Merck Sharp & Dohme (Readington Township, New Jersey), Organon (Oss, The Netherlands), Orion Pharma (Espoo, Finland), Pfizer, ResMed (Espoo, Finland), Respironics (Murrysville, Pennsylvania), Sanofi-Aventis (Paris, France), and Servier (Neuilly sur Seine, France). M. P. is a consultant for Actelion, Boehringer-Ingelheim, GlaxoSmithKline, IST Technology (Helsinki, Finland), Leiras (Helsinki, Finland), Organon, Orion Pharma, ResMed, Sanofi-Aventis, Servier, Somnomedics GmbH (Randersacker, Germany), and UCB (Brussels, Belgium) and received honoraria from GlaxoSmithKline, Leiras, Sanofi-Aventis, and UCB. T. L. J. T. is a consultant for AstraZeneca, GlaxoSmithKline, Orion Pharma, and Pfizer and received honoraria from Astellas Pharma (Tokyo, Japan), GlaxoSmithKline, Leiras, and Pfizer.

REFERENCES


