Short Communication

Toenail selenium concentrations and bladder cancer risk in women and men

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Prediagnostic selenium concentrations measured in archived toenails were inversely associated with bladder cancer risk in women (P for trend = 0.002), but not in men, in a nested case–control study of 338 cases and 341 matched controls. These findings may be due to chance and more studies are needed to determine whether associations between selenium and bladder cancer risk differ by sex.

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Data from animal studies, observational studies and clinical trials suggest that selenium is anticarcinogenic and can reduce the risk of a number of different cancers (Combs, 2005). Five studies, each with fewer than 36 bladder cancer cases, reported inconsistent findings for selenium measured in serum (Nomura et al., 1987; Helzlsouer et al., 1989), toenails (Knekt et al., 1990; Garland et al., 1995), or given as supplements in an intervention trial (Clark et al., 1996). In two large cohort studies with selenium measurements from prediagnostic toenail samples, no association was observed with bladder cancer risk among male smokers (Michaud et al., 2002), but a strong inverse association was reported in a study conducted in the Netherlands (Zeegers et al., 2002).

Of the two largest studies to date, only 59 women were available for analysis in one study (Zeegers et al., 2002) and the other study consisted only of men (Michaud et al., 2002). We examined the sex-specific association between prediagnostic toenail selenium and bladder cancer risk using data from two large cohort studies.

MATERIALS AND METHODS

The Health Professionals Follow-Up Study (HPFS) and the Nurses’ Health Study (NHS) cohorts provided data for our analyses of toenail selenium concentrations and bladder cancer risk. Details on these two cohorts have been previously published (Michaud et al., 2001). Toenail clippings were obtained from 33,737 HPFS participants in 1987 and from 68,213 NHS participants in 1983.

At baseline, participants of these cohorts provided information on disease history, history of cigarette smoking, medication use, weight and height. In addition, smoking status is updated biennially. The National Death Index was used to determine vital status for nonrespondents, and the remaining nonrespondents were assumed to be alive and at risk for bladder cancer incidence.

Participants report new medical conditions on a biennial basis. Self-reported bladder cancers are confirmed by review of medical records (87%) or with an additional letter or phone call. Similar procedures are followed when the National Death Index lists a previously unreported bladder cancer for a deceased participant.

Participants with bladder cancer were eligible for this analysis if the cancer diagnosis occurred after the toenail samples were obtained and they had no prior history of cancer (other than nonmelanoma skin cancer). From the time of the toenail collection through 2000, 338 incident cases with toenail samples were available for the selenium analyses (222 men; 116 women). Controls were matched (1:1) on gender, age (within 1 year), smoking status (current, past, never, unknown) and month of toenail collection. In addition, controls had to be cancer free (other than nonmelanoma skin cancer) at the time of the bladder cancer diagnosis in the matched case.

Selenium concentrations were measured in two batches by Dr Morris at the University of Missouri Research Reactor, Columbia, MO, USA, using neutron activation analysis (NAA) (Baskett et al., 1995). For quality control, selenium was measured in NIST SRM 1577, giving 1.088 ± 0.065 μg g−1 (n = 24, in the first batch) and 1.108 ± 0.031 μg g−1 (n = 8, in second batch) vs a certified value of 1.1 ± 0.1. The CV% for these quality controls was 6.0 and 2.8 for the first and second batches, respectively. In addition, we obtained a CV% of 5.7 for 24 internal controls (split toenail samples), which were blinded to the laboratory.

Odds ratios (OR) and 95% confidence intervals (CI) were estimated using conditional logistic regression models. Quartiles of selenium concentrations were created according to the distribution of the controls for each sex, separately. A continuous variable for pack-years of smoking was additionally included in all models. As a secondary analysis, we performed analyses with a 4-year lag to exclude preclinical cases at baseline. All P-values are based on two-sided statistical tests. We performed tests for trend by assigning
the median value to each category and modelling this variable as a continuous variable.

RESULTS

Mean selenium toenail concentrations were higher in men than women (0.896 vs 0.787 μg g⁻¹, among controls, respectively), but similar among cases and controls for each sex (Table 1). Even though cases and controls were matched on smoking status (never, past, current), pack-years of smoking (among past and current smokers) were higher among cases than controls. Among women, menopausal status and age at menopause were similar in cases and controls.

In a conditional regression model, higher toenail selenium concentrations were associated with significantly lower risk of bladder cancer in women (P for trend = 0.02), but not in men (Table 2). Median selenium concentrations varied by smoking status (HPFS (μg g⁻¹): never 0.825, past 0.814, current 0.739; NHS (μg g⁻¹): never 0.777, past 0.785, current 0.716). Among women, the strongest association between selenium and bladder cancer was observed among past smokers (RR = 0.17, 95% CI = 0.04 – 0.74 for the highest tertile of selenium, compared to the lowest); among men, smoking status did not significantly modify the associations (data not shown).

Removing cases diagnosed with bladder cancer within the first 4 years of follow-up strengthened the inverse association in women (RR = 0.27, 95% CI = 0.10 – 0.74, highest vs lowest quartile), but resulted in a positive association in men (RR = 1.53, 95% CI = 0.71 – 3.28, highest vs lowest quartile).

DISCUSSION

In this nested case–control, a strong inverse association was observed between toenail selenium concentrations and bladder cancer risk in women, but no association was observed in men. The association, in women, was strongest among past smokers.

In the largest study to date on toenail selenium concentrations and bladder cancer, only 59 cases (out of 431 cases) were women and associations were not presented separately by sex (Zeegers et al, 2002). In that study, conducted in the Netherlands between 1986 and 1992, mean toenail selenium concentrations were substantially lower than ours, in both men and women (0.536 for men, 0.568 for women). A relative risk of 0.67 (95% CI = 0.46 – 0.97) was reported for the highest vs lowest quintile of selenium toenail level; the strongest association was among those who recently quit smoking (<10 years) and no associations were noted for never and current smokers. Our findings for women were similar to these as the inverse associations were also strongest among past smokers.

Data from animal studies, epidemiologic data and human prevention trials suggest that selenium is an important chemo-preventive agent (Combs, 2005). Although the exact mechanism by which selenium prevents cancer is unknown, multiple mechanisms are likely to be involved. At least part of the antimutagenic effect of

Table 1 | Prediagnostic toenail selenium level and baseline characteristics of nested bladder cancer cases and matched controls from the NH-S and HPFS cohort studies

| Mean (s.d.) or % | Cases | Controls |
|-----------------|-------|----------|
| **Men**         |       |          |
| No. of subjects | 222   | 224      |
| Toenail selenium level (μg g⁻¹) | 0.897 (0.53) | 0.896 (0.58) |
| Median          | 0.814 | 0.805    |
| Age (years)     | 62.3 (8.6) | 62.3 (8.6) |
| Smoking (%)     |      |          |
| Never           | 26.1  | 25.9     |
| Past            | 53.6  | 54.0     |
| Current         | 16.7  | 16.5     |
| Missing         | 3.6   | 3.6      |
| Pack-years      | 34.1 (22.9) | 29.6 (23.5) |

| **Women**       |       |          |
| No. of subjects | 116   | 117      |
| Toenail selenium level (μg g⁻¹) | 0.802 (0.50) | 0.787 (0.19) |
| Median          | 0.733 | 0.763    |
| Age (years)     | 53.7 (6.0) | 53.7 (6.1) |
| Smoking (%)     |      |          |
| Never           | 26.5  | 26.5     |
| Past            | 23.9  | 24.8     |
| Current         | 48.7  | 47.9     |
| Missing         | 0.9   | 0.9      |
| Pack-years      | 31.7 (22.9) | 24.6 (17.8) |
| Premenopausal   | 21    | 22       |
| Postmenopausal  | 74    | 71       |
| Dubious/missing | 5     | 7        |
| Age at menopause | 48.9 (4.2) | 49.4 (3.5) |

| **NHS = Nurses’ Health Study; HPFS = Health Professionals Follow-Up Study; s.d. = standard deviation. aBased on 221 cases and 223 controls. bMatched factors (age at time of toenail collection, smoking status in 1986 for men and 1980 for women). cAmong past and current smokers. dAmong natural postmenopausal women (55 cases and 49 controls). |
selenium is mediated by Se-dependent enzymes such as glutathione peroxidases and thioredoxin reductases, which protect DNA from oxygen radicals (Combs and Gray, 1998). In Se-adequate individuals, the anticarcinogenic effect of selenium may be mediated by selenium metabolites which can influence carcinogen metabolism, immune function, tumour growth, apoptosis induction (Combs and Gray, 1998) and DNA methylation (Schrauzer, 2000).

The different results observed in men and women may be due to a chance inverse association in women. Alternatively, a possible explanation for the lack of association in men in our study could be that selenium offers no additional protection within the observed range of selenium concentrations. Mean selenium concentrations were slightly higher in men than women in our study and substantially higher than those previously reported in the bladder study conducted in the Netherlands (Zeegers et al., 2002). The large sex differences in incidence rates that exist for this cancer suggest that important etiologic differences exist between men and women; thus, it is plausible that the association between selenium and bladder cancer varies by sex. An alternative possibility is that the difference is due to the timing of the toenail collection. On average, women in the NHS provided toenails when they were 8 years younger than when men in the HPFS provided toenails (Table 1), and consequently this difference may have provided a better measure of early exposure for women.

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A limitation to our study includes possible measurement error when assessing toenail selenium concentrations, which may have resulted in the null finding in men. However, because toenails from the two different cohorts were analysed at the same time for each of the two batches, and CV% were good for both batches, it is unlikely that this could explain the different results by sex.

Strengths of the study include the largest number of female bladder cancer cases to date with selenium data, pre-diagnostic measurements, use of toenails (where selenium is more time-integrated), detailed data on potential confounders, 1:1 matching by known confounders and high follow-up rates in the two cohorts used to select cases and controls.

In summary, we observed a statistically significant inverse association between prediagnostic selenium toenail concentrations and risk of bladder cancer in women, but not in men. To date, this is the largest study to report findings on selenium concentrations and bladder cancer risk in women. Future studies are needed to confirm our findings.

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