

Letters to the Editor

RE: "VALIDATION OF A METHOD FOR RECONSTRUCTING HISTORICAL RATES OF SMOKING PREVALENCE"

In their recent paper, Bilal et al. (1) incorrectly assert that no researchers have assessed the validity of historical smoking prevalence derived from retrospectively reported data. A large body of literature validates this type of data in general (2–4) and smoking data in particular (5–12). In fact, at least 2 previously published studies applied an improved method (8, 9) with better data.

Bilal et al. correct retrospectively derived smoking prevalence rates for differential mortality rates of smokers and nonsmokers and validate their estimates against contemporaneously reported data. However, they validate prevalence rates in just 5 prior years using paired surveys that differ by, at most, 20 years. The earlier studies validate prevalence rates in 11 (8) and 27 (9) prior years using paired surveys that differ by up to 40 years. More importantly, Bilal et al. correct for differential mortality bias with a method developed in 1983 (13), when age, sex, and cause-specific mortality data were unavailable. They ignore the updated method of Christopoulou et al. (9), which exploits these now widely available vital statistics. The new data allowed Christopoulou et al. to relax the assumption that differential mortality rates are constant within broad age categories. Following Peto et al. (14), these authors calculate the number of smoking-attributable deaths for each smoking-related disease by sex, year, and 5-year age category and apply the method of Harris (13) using these data. Thus, they reconstruct historical

smoking prevalence for disaggregated demographic groups (by sex and birth cohort) and formally test when, and for which groups, reconstructed prevalence rates deviate from "true" rates.

The more detailed data and method lead to substantively different research and policy implications. Bilal et al. (1) find that the mortality correction affects only the reconstructed smoking prevalence rates of men in the early decades of their study. For all other cases, they find no statistically significant differences. These results and their perception of the correction method's complexity lead Bilal et al. to advise against correcting for differential mortality rates for recent historical periods. Christopoulou et al. (9) showed that whether, when, and how the differential mortality adjustment affects smoking rates depends mostly on how old respondents were when interviewed, not on the time period of study.

To illustrate, we apply their method to the Spanish National Health Survey data used by Bilal et al. (for details, see the Web material available at <http://aje.oxfordjournals.org/>). In Figure 1, we plot the adjusted (dashed line) and unadjusted (solid line) smoking prevalence rates for Spanish men and women who were aged A) 60–69 years, B) 70–79 years, or C) 80 or more years in 2007. In Spain, as in other countries treated in the article by Christopoulou et al. (9), the 2 series overlap almost completely for the group of men aged 60–69 years and for all groups of women (whose

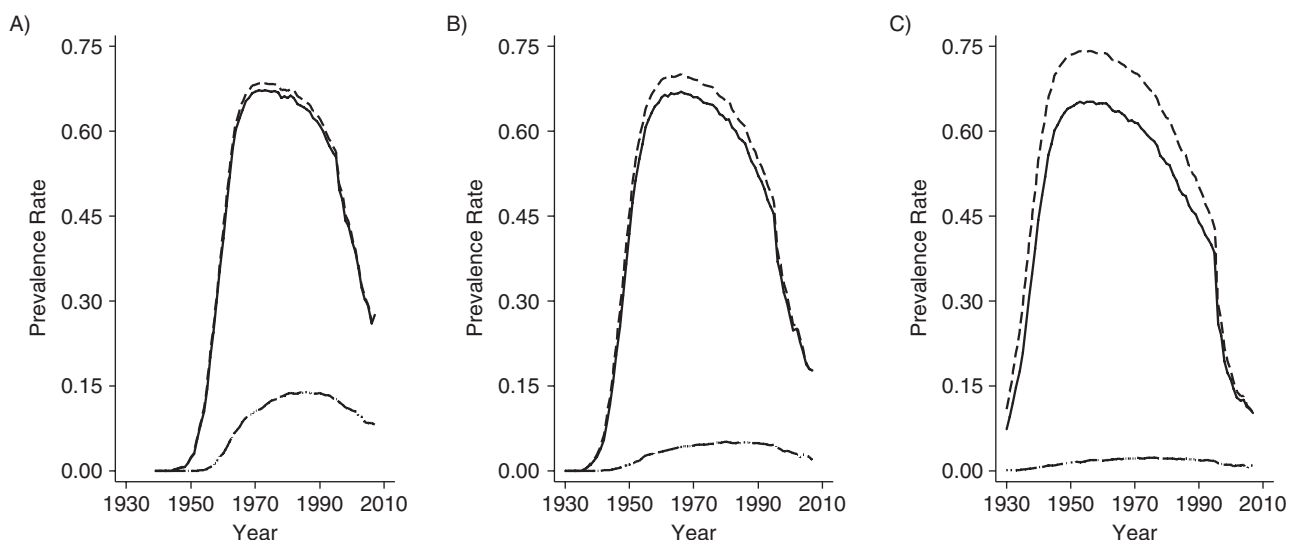


Figure 1. Correction of smoking prevalence for differential mortality rates of men (2 top lines) and women (2 bottom overlapping lines) at ages A) 60–69 years, B) 70–79 years, and C) 80 or more years in 2007. Retrospective smoking data are from the Spanish National Health Survey in 1995, 1997, 2001, 2003, and 2006.

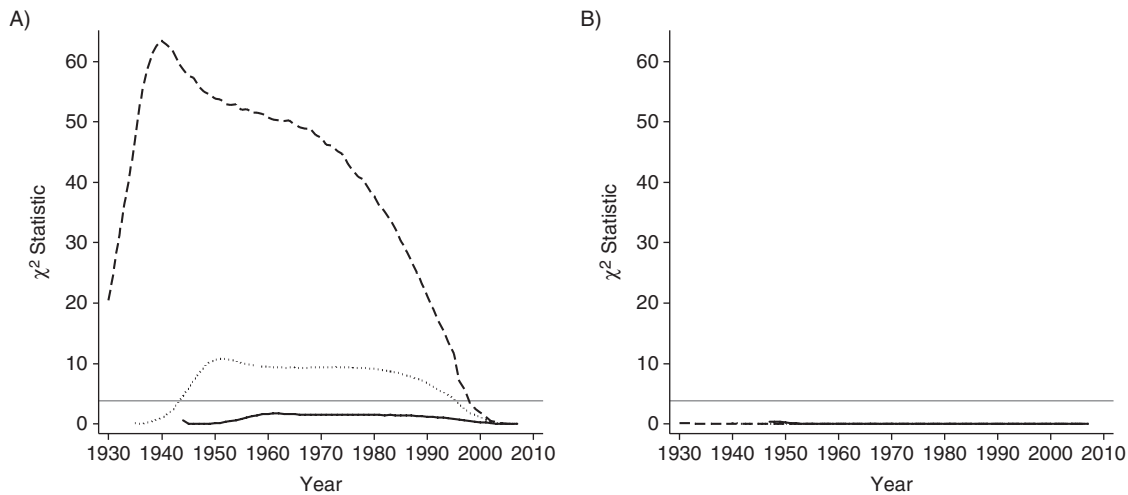


Figure 2. χ^2 test of differences between adjusted and unadjusted smoking prevalence rates for A) men and B) women at ages 60–69 years (solid line), 70–79 years (dotted line), and 80 or more years (dashed line) in 2007. The horizontal gray line marks the critical value at the 5% level of statistical significance.

smoking prevalence rates never exceed 14%). Figure 2A (for men) and Figure 2B (for women) plot for each year of each cohort’s life the χ^2 statistic derived from the standard Pearson test of independence for binary variables, which tests whether adjusted and unadjusted prevalence rates differ statistically. As in the study by Christopoulou et al. (9), rates differ statistically only for men who, when surveyed, were 70 or more years of age and differ statistically in years quite close to the survey year. Table 1 summarizes the results.

The findings are particularly pertinent to users of the growing number of surveys that interview individuals aged 50 years or more. Aging studies in the US Health and Retirement Study (15), the Survey of Health, Ageing, and Retirement in Europe (16), the English Longitudinal Study of Ageing (17), the Korean Longitudinal Study of Ageing (18), the China Health and Retirement Longitudinal Study (19), and the Longitudinal Aging Study of India (20) ask respondents to

retrospectively report their lifetime smoking behaviors. Especially because these samples include many older respondents, researchers need to account for differential smoking-related mortality rates.

More broadly, the findings highlight the point that, to identify the degree of smoking diffusion in a particular country and year, as proposed by the cigarette epidemic model (21), the population smoking prevalence rate masks important heterogeneity across cohorts, sexes, and countries. This point, recently acknowledged in an updated “epidemic” model (22), was already made by Christopoulou et al. (9), not only with respect to sex, but also with respect to age.

An interesting contribution of Bilal et al. (1) is that they follow the cigarette epidemic model (21, 22) and estimate the correlation between reconstructed smoking prevalence and future lung cancer mortality rates (as a proxy for smoking-attributable mortality rate). Of course, the underlying structural

Table 1. Summary of Indicators of Unadjusted and Adjusted Peak Smoking Prevalence Rates for 3 Cohorts of Spanish Men and Women

| Age in 2007, years, by Sex | Mean Sample Size ^a | Peak Unadjusted Prevalence | Peak Adjusted Prevalence | Difference at Peak Rate | Years With Statistically Significant Differences |
|----------------------------|-------------------------------|----------------------------|--------------------------|-------------------------|--|
| Men | | | | | |
| 60–69 | 2,235 | 67.26 | 68.53 | 1.27 | |
| 70–79 | 3,856 | 67.03 | 70.07 | 3.04 | 1944–1995 |
| ≥80 | 3,637 | 65.27 | 74.21 | 8.94 | 1924–1997 |
| Women | | | | | |
| 60–69 | 4,989 | 13.99 | 14.01 | 0.02 | |
| 70–79 | 5,858 | 5.15 | 5.15 | 0 | |
| ≥80 | 3,706 | 2.32 | 2.32 | 0 | |

^a Mean sample size is the average number of individuals over all available years.

relationships between smoking and disease are not captured in this reduced-form model, which weakens the policy relevance of the results. However, the more immediate point is that the exercise does not test their article's central claim—whether reconstructed smoking rates are valid.

Bilal et al. (1) also fail to acknowledge a broader body of literature that addresses other types of bias that are likely present in retrospective smoking data. For example, Kenkel et al. (23) treated an indicator of smoking initiation constructed from retrospective data as a standard binary variable measured with error and used an established technique to correct it (24). They showed that some of the bias occurs because people who smoke few cigarettes in early life do not identify themselves as smokers later in life (i.e., light-smoker bias). Further, Bar and Lillard (25) documented the bias that arises when people “heap” their retrospective reports of smoking cessation on units evenly divisible by 5 (i.e., heaping bias) and proposed a method to mitigate it. Of course, more work remains. The correction method described by Bar and Lillard (25) can be extended and enriched to control for factors that systematically predict recall errors. One might also refine the method of Christopoulou et al. (9) by relaxing the assumption that smokers who survive to answer a survey start and stop smoking at the same rate as smokers who die before they are surveyed. These and other issues are at the forefront of research that attempts to validate historical smoking rates generated with retrospective data.

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