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<th><strong>Title</strong></th>
<th>The fraction of ischaemic heart disease and stroke attributable to smoking in the WHO Western Pacific and South-East Asian regions</th>
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<tr>
<td><strong>Author(s)</strong></td>
<td>Martiniuk, ALC; Lee, CMY; Lam, TH; Huxley, R; Suh, I; Jamrozik, K; Gu, DF; Woodward, M</td>
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</table>
The fraction of ischaemic heart disease and stroke attributable to smoking in the WHO Western Pacific and South-East Asian regions

A L C Martiniuk, C M Y Lee, T H Lam, R Huxley, I Suh, K Jamrozik, D F Gu, M Woodward and for the Asia Pacific Cohort Studies Collaboration

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RESEARCH PAPER

The fraction of ischaemic heart disease and stroke attributable to smoking in the WHO Western Pacific and South-East Asian regions

A L C Martiniuk, C M Y Lee, T H Lam, R Huxley, I Suh, K Jamrozik, D F Gu, Woodward M, for the Asia Pacific Cohort Studies Collaboration

Background: Tobacco will soon be the biggest cause of death worldwide, with the greatest burden being borne by low and middle-income countries where 8/10 smokers now live.

Objective: This study aimed to quantify the direct burden of smoking for cardiovascular diseases (CVD) by calculating the population attributable fractions (PAF) for fatal ischaemic heart disease (IHD) and stroke (haemorrhagic and ischaemic) for all 38 countries in the World Health Organization Western Pacific and South East Asian regions.

Design and subjects: Sex-specific prevalence of smoking was obtained from existing data. Estimates of the hazard ratio (HR) for IHD and stroke with smoking as an independent risk factor were obtained from the ~600 000 adult subjects in the Asia Pacific Cohort Studies Collaboration (APCSC). HR estimates and prevalence were then used to calculate sex-specific PAF for IHD and stroke by country.

Results: The prevalence of smoking in the 33 countries, for which relevant data could be obtained, ranged from 28–82% in males and from 1–65% in females. The fraction of IHD attributable to smoking ranged from 13–33% in males and from <1–28% in females. The percentage of haemorrhagic stroke attributable to smoking ranged from 4–12% in males and from <1–9% in females. Corresponding figures for ischaemic stroke were 11–27% in males and <1–22% in females.

Conclusions: Up to 30% of some cardiovascular fatalities can be attributed to smoking. This is likely an underestimate of the current burden of smoking on CVD, given that the smoking epidemic has developed further since many of the studies were conducted.
in the region is still currently lower than in “the West”, mainly because the full effects of the large recent increases in male smoking are not yet evident. Yet, even at this early stage, because of the size of its population, China has more deaths from tobacco than the United States or Russia.

**Trends**

Whereas current mortality from past smoking patterns can serve as an early warning, greater concern should be directed toward the much larger number of deaths that are expected in coming decades as a result of current smoking, especially for LMIC. Unfortunately, knowing what to expect is difficult as detailed age- and sex-specific trends in smoking by country in the WHO Western Pacific and South East Asian regions are not consistently available. However, it is known that overall tobacco consumption increased in all LMIC in the past two decades by about 3.4% per year, compared with declines in consumption in high-income countries. In selected countries in the Western Pacific and South East Asian regions, percentage increases in per capita cigarette consumption over the period 1970–1990 range from 30% for Thailand and Cambodia to more than 130% for Nepal, China and Indonesia. For example, the average consumption of cigarettes per Chinese man per day was 1 in 1952, 4 in 1972, 10 in 1992, and 11 in 1996. Cases of CVD are increasing simultaneously, with anticipated increases of 120% for women and 137% for men in LMIC between 1990 and 2020, compared with age-related increases of between 30–60% in high-income countries. These are conservative estimates, as they do not take into account the likely increase in the prevalence of other CVD risk factors over time.

**Rationale**

This paper addresses a major area of uncertainty, as indicated by the World Bank Report on Smoking, the lack of knowledge regarding the burden of tobacco-attributable disease, as well as estimates for the prevalence and impact of smoking on women in LMIC. It quantifies the direct burden of smoking for cardiovascular diseases (CVD) by calculating the population attributable fractions (PAFs) for ischaemic heart disease (IHD) and stroke (haemorrhagic and ischaemic) for countries in the WHO Western Pacific and South-East Asian regions. The PAF provides estimates of the proportion of heart disease and stroke cases in a country that could be avoided if smoking were eliminated.

**METHODS AND POPULATIONS**

Calculation of the PAFs required estimates of the prevalence of smoking by country, as well as estimates of hazard ratio (HR) for CVD associated with smoking.

**Prevalence of smoking**

Data on the prevalence of smoking for adult males and females were obtained for countries in the WHO Western Pacific and South-East Asian regions. Although the WHO does not recognise them as separate countries, Hong Kong, Macau and Taiwan were considered separately from China because these regions have unique health, political and administrative bodies that may influence the prevalence of smoking. Data on prevalence were obtained from studies compiled by the WHO and supplemented by data from national surveys, available either through Medline or via national government or non-governmental organisation websites. These other surveys were used only if they provided more recent data or where the WHO did not have sex-specific data on smoking for a particular country. Only those studies compiled by the WHO and judged by the WHO to be “methodologically sound and to provide reasonable reliable and comparable results”, or those national surveys that used random sampling techniques to obtain representative population estimates from countries in the WHO Western Pacific and South-East Asian regions, were used to obtain estimates of the prevalence of smoking.

**Relative risk of cardiovascular disease**

Age-adjusted estimates of the relative risk of smoking for cardiovascular death were obtained from the Asia Pacific Cohort Studies Collaboration (APCSC). The APCSC is an overview of cohort studies in the region. Details on the APCSC, including study identification, data collection, and event verification are described elsewhere. All studies included in APCSC were conducted prospectively in populations from the Asia-Pacific region, had at least 5000 person years of follow-up and recorded age, sex, and blood pressure at baseline and vital status at the end of the follow-up. Studies were excluded from the APCSC if entry was dependent upon the individual having a particular condition or risk factor.

All data on current cigarette smoking (yes/no) were self-reported at enrolment. Cardiovascular deaths were ascertained using monitoring, re-surveying and/or record linkage, and classified according to the ninth revision of the International Classification of Diseases (ICD); IHD (ICD 410–414); stroke (430–438); haemorrhagic stroke (431.0–432.9); ischaemic stroke (433.0–434.9). Deaths were ascribed solely to their underlying cause. Analyses used individual participant data, and were restricted to individuals aged 20 years and over at the time of the baseline survey for whom smoking status was recorded.

**Population attributable fraction**

Using the estimates of prevalence obtained from the literature and the HR from APCSC, population attributable fractions (%) for fatal ischaemic heart disease, haemorrhagic and ischaemic stroke caused by smoking were calculated for each country, for both males and females, using the formula: PAF = [100 × prevalence × (HR−1)]/[100 + prevalence × (HR−1)], where prevalence is in %.

**RESULTS**

Nationally representative data on the sex-specific prevalence of smoking were available for all 27 countries in the WHO Western Pacific Region except for the Marshall Islands and the Solomon Islands; and for all 11 countries in the WHO South-East Asia Region except for Bhutan, North Korea and Timor-Leste. Also, no data were available for Macau. For the Solomon Islands and Timor-Leste, data were available only for female smoking prevalence which, 33% and 1.1%, respectively, and the total prevalence of smoking (males and females) in Bhutan is reported to be 1%. The prevalence of smoking in the other 33 countries in the WHO Western Pacific and South-East Asian regions (plus Hong Kong and Taiwan) ranged from 28–82% in males and from 1–65% in females (fig 1).

A previous individual participant data analysis of the 40 cohort studies from APCSC provided estimates of HR for CVD. In APCSC there are no significant differences (p > 0.05) in the relative effect of smoking on IHD or stroke between countries, between Asia and Australia/New Zealand (ANZ) or between women and men. The HRs (95% confidence intervals (CI)) comparing current smokers with non-smokers are: 1.60 (95% CI 1.49 to 1.72) for ischaemic heart disease; 1.19 (95% CI 1.06 to 1.33) for haemorrhagic stroke; and 1.38 (95% CI 1.24 to 1.54) for ischaemic stroke.

Figures 2–4 present the population attributable fractions for males and females for IHD, haemorrhagic and ischaemic stroke caused by smoking for each country in the WHO Western Pacific and South-East Asian regions. Countries are
ranked by the total prevalence of smoking in fig 1 and this ranking remains for figs 2–4. The fraction of IHD attributable to smoking ranged from 13–33% in males and from 1–28% in females (fig 2). The fraction of haemorrhagic stroke attributable to smoking ranged from 4–12% in males and from 1–9% in females (fig 3). The fraction of ischaemic stroke attributable to smoking ranged from 11–27% in males and from 1–22% in females (fig 4).

**DISCUSSION**

We have quantified the direct burden of smoking on cardiovascular diseases (CVD) in the WHO Western Pacific and South East Asian regions by calculating the PAFs for IHD and stroke. Results from this study suggest that up to ~30% of fatal CVD in these populations can be attributed to smoking, depending on the outcome of interest. Much of the projected increase in mortality over the next 50 years could be avoided if adults quit smoking; however, quitting remains rare in LMIC.

Interestingly, this study demonstrated that Australia and New Zealand, the two higher-income countries in the WHO Western Pacific and South-East Asian regions, with predominantly white populations, have relatively low smoking prevalence and therefore low PAFs for CVD. As expected, the attributable fractions for all outcomes are greater for males because their prevalence of smoking is currently higher. PAFs for CVD in women will almost certainly increase in the future because smoking by women...
in LMIC in these regions is expected to increase in the coming years. The attributable fractions for CVD for Australia and New Zealand from this study are similar to estimates previously published by Peto et al in their monograph providing PAFs for various outcomes caused by smoking in developed countries. However, the methods used here are more direct and robust.

While this paper addresses only the PAF for heart disease and stroke associated with smoking, not only is the prevalence of smoking increasing in the Western Pacific and South East Asian regions, but the countries in these regions are also experiencing increases in other risk factors for CVD, including overweight/obesity, diabetes, hypertension and dyslipidaemia. Ominously, these trends are likely to work together, further amplifying observed death and disability from IHD and stroke.

**Strengths and limitations**

This study has the strength of being able to calculate PAFs using HRs from the region based on the largest-ever study of CVD in the Asia-Pacific, and one of the largest anywhere in the world. The size of the APCSC means that the overall estimates of the relative effects of smoking are more precise than those employed in most previous calculations and offer an improvement over previous indirect methods for estimation.

However, the APCSC does have a number of disadvantages, particularly with respect to the assessment of smoking. First, this study needed to define smoking as a simple
self-reported, dichotomous variable (yes/no) reflecting status at baseline, since few variables on smoking habits are included in the Collaboration. Thus PAFs could not be calculated using duration of smoking, although duration is recognised as more important than daily consumption of cigarettes in determining the effects of smoking on some health outcomes such as cancer.\(^{16}\) However, the relative importance of duration versus daily consumption smoking measures is less clear for cardiovascular outcomes.\(^{19}\) If the measurement of the duration of smoking is important for CVD, as it is for cancer, then the inclusion of ex-smokers in the reference group will have led to an underestimation of the RRs associated with current versus never smoking and a concomitant underestimation of PAFs.

A second limitation is that no objective measures of smoking status were available in the APCSC database, nor in studies used to estimate the prevalence of smoking. Any misclassification of current smokers as former or never-smokers will have served to underestimate the HRs and hence the PAFs.

A third limitation is that, although there were no statistically significant differences in APCSC between countries in HR, for IHD or stroke, the same measure of HR has been used to calculate the PAFs for countries not represented in APCSC (for example, India). A fourth limitation is that although the estimates of risk were adjusted for age, data on prevalence of smoking were not. A final limitation is that the individual studies included in the APCSC used different
methods to verify CVD outcomes, and the lack of standardisation could have some effect on the estimation of the HR. Of particular importance for the current study is that only 57% of fatal strokes were classified by subtype and, while pathological diagnoses were verified by imaging or autopsy, limited information was available from many studies.

In summary, all of the limitations in this study are likely to have resulted in conservative estimates of the PAFs. This only further underscores the impact of smoking on CVD in these regions.

**Conclusion**

There is convincing evidence that, even on conservative assumptions, tackling smoking could prevent millions of deaths over the next few decades. A decrease in the

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**Figure 4** Population attributable fraction (%) of ischaemic stroke caused by smoking.

<table>
<thead>
<tr>
<th>Country</th>
<th>Year</th>
<th>Male PAF (%)</th>
<th>Female PAF (%)</th>
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<td>2001</td>
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</tr>
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<td>Singapore</td>
<td>2001</td>
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</tr>
<tr>
<td>Hong Kong</td>
<td>2000</td>
<td>10</td>
<td>2</td>
</tr>
<tr>
<td>Palau</td>
<td>1997</td>
<td>9</td>
<td>3</td>
</tr>
<tr>
<td>India</td>
<td>1999</td>
<td>11</td>
<td>1</td>
</tr>
<tr>
<td>Australia</td>
<td>2004</td>
<td>8</td>
<td>7</td>
</tr>
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<td>Thailand</td>
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<td>10</td>
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**Future**

CVD and its risk factors are moving not only from high-income to LMIC, but also from the more affluent in LMIC to the less affluent. Yet, a recent assessment by the WHO has indicated that the capacity of many countries to prevent and treat chronic diseases remains extremely weak. There is a need to develop integrated approaches to prevention, surveillance and control of chronic diseases. There is also a need to continue monitoring the state of smoking and CVD in order to ensure country-specific policies and programmes for the prevention and treatment of CVD are appropriate, and to gauge the effectiveness of current interventions. The present study is a useful step, and future studies with country-specific HRs are warranted.
The prevalence of smoking would not only address the increasing rates of CVDs and their risk factors but will also contribute substantially to reducing the incidence and impact of other chronic diseases, such as cancer and chronic respiratory diseases, that share smoking as a common risk factor. There are existing tobacco control policies that have been shown to be effective. Implementation of the Framework Convention on Tobacco Control and following through with its protocols is likely the best first step toward addressing the large fraction of cardiovascular disease that is attributable to smoking.

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