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Alcohol attributable burden of incidence of cancer in eight European countries based on results from prospective cohort study

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ABSTRACT

Objective To compute the burden of cancer attributable to current and former alcohol consumption in eight European countries based on representative population based data on alcohol exposure.

Design Combination of prospective cohort study with representative population based data on alcohol consumption.

Setting Eight countries (France, Italy, Spain, United Kingdom, the Netherlands, Greece, Germany, Denmark) participating in the European Prospective Investigation into Cancer and Nutrition (EPIC) study.

Participants 109,118 men and 254,870 women, mainly aged 37-70.

Main outcome measures Hazard rate ratios expressing the relative risk of cancer incidence for former and current alcohol consumption among EPIC participants. Hazard rate ratios combined with representative information on alcohol consumption to calculate alcohol attributable fractions of causally related cancers by country and sex. Partial alcohol attributable fractions for consumption higher than the recommended upper limit (two drinks a day for men with about 24 g alcohol, one for women with about 12 g alcohol) and the estimated total annual number of cases of alcohol attributable cancer.

Results If we assume causality, among men and women, 10% (95% confidence interval 7 to 13%) and 3% (1 to 5%) of the incidence of total cancer was attributable to former and current alcohol consumption in the selected European countries. For selected cancers the figures were 44% (31 to 56%) and 25% (5 to 46%) for upper aerodigestive tract, 33% (11 to 54%) and 18% (3 to 38%) for liver, 17% (10 to 25%) and 4% (1 to 10%) for colorectal cancer for men and women, respectively, and 5.0% (2 to 8%) for female breast cancer. A substantial part of the alcohol attributable fraction in 2008 was associated with alcohol consumption higher than the recommended upper limit: 33,037 of 178,578 alcohol related cancer cases in men and 17,470 of 397,043 alcohol related cases in women.

Conclusions In western Europe, an important proportion of cases of cancer can be attributable to alcohol consumption, especially consumption higher than the recommended upper limits. These data support current political efforts to reduce or to abstain from alcohol consumption to reduce the incidence of cancer.

INTRODUCTION

Alcohol consumption is thought to account for a substantial number of deaths worldwide, with Europe and America showing the highest alcohol attributable fractions of 6.5% and 5.6%, respectively.1 Chronic diseases, especially cancer, contribute markedly to this burden. In 2007 the International Agency for Research on Cancer (IARC) added two of the most common cancers—female breast and colorectal cancer—to the list of cancers causally related to alcohol, which previously consisted of oral cavity, pharynx, larynx, oesophagus, and liver cancer.2 Although alcohol consumption is a major risk factor for cancer incidence,
and Europe is among the regions with the highest per capita alcohol consumption, detailed information on the fractions of cancer that are attributable to alcohol consumption based on direct empirical evidence for the different cancer sites is sparse, and systematic and comparable estimations across European countries are lacking. Moreover, previous estimates of the alcoholic attributable fractions refer to the burden from current alcohol consumption but do not consider the risk of former alcohol consumption. Also, in 2007 the World Cancer Research Fund/American Institute for Cancer Research published recommendations on the maximum recommended daily alcohol consumption. We do not know how much of the burden of incidence of cancer is attributable to alcohol and occurs because of consumption higher than the recommended upper limit.

We estimated the total (current and former alcohol consumption) and partial (alcohol consumption higher than the recommended upper limit) alcohol attributable fractions for the incidence of total and specific cancers related to alcohol in eight European countries based on hazard rate ratios from the European Prospective Investigation into Cancer and Nutrition (EPIC) study and linked those alcohol attributable fractions to incidences of cancer to estimate the annual absolute number of cancer cases attributable to alcohol in these countries.

METHODS

Study population

The EPIC study is a multicentre prospective cohort study that, from 1992 to 2000, recruited about 520,000 randomly selected men and women aged mainly 35-70 from 10 European countries. Eligible participants were selected from the general population, except in France, where selection was based on members of the health insurance system or state school employees, and in Utrecht (the Netherlands), where selection was based on women attending screening for breast cancer. Participants gave informed consent and completed questionnaires on diet and lifestyle. The present analyses included participants free from cancer at recruitment and who were not in the top or bottom 1% of the ratio of energy requirement to energy expenditure ($n=478,478$). Participants with incomplete information on alcohol consumption at recruitment or in the past ($n=114,481$) and missing dietary information ($n=9$) were excluded, leaving 363,988 men and women from France, Italy, Spain, the Netherlands, United Kingdom, Greece, Germany, and Denmark. France and Utrecht enrolled only women. As we wanted to consider the risk of cancer incidence associated with former alcohol consumption, we had to exclude the centres of Norway, Sweden, Bilthoven, and Naples because they did not have information on past consumption.

Alcohol consumption at recruitment (in grams per day) was measured with a validated dietary questionnaire assessing frequency and portion size of beer, cider, wine, spirits, and fortified wine covering the 12 months before recruitment. Consumption in the past was assessed as self reported consumption of beer, wine, and spirits at the ages of 20, 30, 40, and 50. Based on consumption in the past and at recruitment we distinguished between never (no consumption in the past and no consumption at recruitment), former (consumption in the past but no consumption at recruitment), and lifetime consumers (consumption in the past and at recruitment). For lifetime consumers, consumption as applied in this analysis reflects the past years’ consumption before recruitment.

We obtained information on incidence of cancer through record linkage with regional cancer registers in countries with passive follow-up (Denmark, Italy, the Netherlands, Spain, UK) or by a combination of methods including medically verified self reports of the participant or the next of kin, cancer or pathology registers, health insurance records, or death certificates in countries with active follow-up (France, Germany, Greece). The follow-up ended between 2002 and 2005, and loss to follow-up was relatively low, with <2% in all countries irrespective of active or passive follow-up. We investigated cancers with a causal association to alcohol consumption (colorectal (C18-21), ICD-O (international classification of diseases-oncology, 2nd revision), upper aerodigestive tract (C00-10, C12-15, C32), liver (C22), female breast (C50)), as well as total cancer (C00-C80, except C44 skin cancer) and alcohol related cancers combined (upper aerodigestive tract, colorectal, liver, and, for women, female breast cancer). ICD codes of cancer end points were in accordance with the GLOBOCAN-2008 cancer definitions.

Statistical analysis

We combined hazard rate ratios derived from the EPIC study with representative data on alcohol consumption from the general population. Cox proportional hazard regressions were applied to compute hazard rate ratios during a mean follow-up time of 8.8 years for alcohol consumption among lifetime consumers per 12 g/day increment (equivalent to one drink of any alcoholic beverage) and for former compared with never consumers and incidence of first primary cancer. Age was used as the underlying time variable with entry and exit time defined as the participant’s age at recruitment and age at diagnosis of cancer or at censoring, respectively. Results on some single cancer outcomes have been published earlier. We used updated information and recomputed hazard rate ratios for these sites. To control for age and variations in study procedures across the EPIC centres we stratified the analyses by age (in 1 year categories) and centre.

All models were run separately for men and women, and included the following potential confounders, which were measured at recruitment: smoking (never; past <10 years ago, ≥10 years ago; current <15, 15-25, or ≥25 cigarettes/day, other (cigars, pipe, cigarettes with missing dose)) and smoking duration (<10, 10-<20, 20-<30, 30-<40, ≥40) years, missing
was examined by the meta-analytic approach and (P<0.1).

Liver cancer in men and for colorectal cancer in women

Heterogeneity of hazard rate ratios across centres was examined by the meta-analytic approach and (P<0.01 for non-linearity). We therefore used regression coefficients (β) as risk functions to express the risk for cancer incidence per 1 g/day increment in alcohol consumption among current lifetime consumers.

We tested effect modification by smoking for the site specific cancers by including product terms of smoking status (never, former, current smoker) with alcohol (g/day) and performing the likelihood ratio test between nested models. There was indication for an effect modification by smoking for upper aerodigestive tract and liver cancer in men and for colorectal cancer in women (P<0.01 for non-linearity). We therefore used this approach for liver cancer in men.

For adjusted risk estimates, the prevalence of exposure among cases rather than among the general popula-

Alcohol attributable fractions

The computation of alcohol attributable fractions requires not only the information on relative risks for alcohol consumption (such as hazard rate ratios) but also information on the distribution of alcohol consumption within the general population. We computed alcohol exposure data from the general population following an algorithm (triangulation) that combined information of alcohol consumption from survey data as reported by the World Health Organization and per capita consumption for each country and separately for men and women aged ≥15. These sex and country specific data on alcohol exposure were modelled as gamma function in current consumers of alcohol by applying a formula based on the triangulated population mean alcohol intake (combining alcohol survey information with data on per capita alcohol consumption). These gamma functions were shown to fit and best model the right skewed distribution of alcohol consumption on the population level. Furthermore, this exhibits the clear advantage of using alcohol consumption continuously, as the estimation of alcohol attributable fractions based on alcohol categories might lose valuable information. We also obtained information on the proportions of never and former consumers of alcohol from WHO, which used the GENACIS survey as source of information, except for Greece, for which data were derived from a national survey on licit and illicit drug use. We then calculated the country and sex specific alcohol attributable fractions reflecting the burden of cancer incidence associated with total alcohol consumption on the population level (equation A, fig 1).

We also computed the part of the alcohol attributable fraction (partial alcohol attributable fraction) that reflects the burden of cancer incidence associated with alcohol consumption higher than the recommended upper limit of two standard drinks a day (>24 g/day) for men and one standard drink a day (>12 g/day) in women (equation B, fig 1).

For adjusted risk estimates, the prevalence of exposure among cases rather than among the generalpopu-

Equations for computation of alcohol attributable fractions

**Equation A**

$$\text{AAF}_{\text{total}} = \frac{250 \text{ g/day}}{0.0001 \text{ g/day}} \int P_{NC}(x) \cdot HRR(x) \, dx - 1$$

$$\text{AAF}_{\text{partial}} = \frac{250 \text{ g/day}}{250 \text{ g/day}} \int P_{LC}(x) \cdot HRR(x) \, dx - 1$$

where $P_{NC}$, $P_{LC}$, and $P_{FC}$ = prevalence of never (%), former (%), or lifetime consumers (% and gamma distribution), respectively. $HRR(x) = \text{risk of cancer per consumed gram of alcohol a day for lifetime consumers}$. $HRR_{FC} = \text{risk of cancer incidence in former compared with never consumers}$.

**Equation B**

$$\text{AAF}_{\text{partial}} = \frac{250 \text{ g/day}}{250 \text{ g/day}} \int P_{LC}(x) \cdot HRR(x) \, dx - P_{LC} \geq 24 \text{ g/day}$$

where $P_{LC} \geq 24 \text{ g/day}$ = proportion of lifetime consumers with consumption larger than recommended upper limit. Counterfactual scenario for total and partial alcohol attributable fraction was complete elimination of alcohol consumption in population.
Table 1 | Proportions of never, former, and lifetime consumers of alcohol and mean alcohol consumption in lifetime consumers in general adult population aged 15 years or older

<table>
<thead>
<tr>
<th>Country</th>
<th>Never consumers (%)</th>
<th>Former consumers (%)</th>
<th>Lifetime consumers (%)</th>
<th>Mean* (SD) g/day</th>
<th>% drinking over daily recommended upper limit†</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Men</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Denmark</td>
<td>0.6</td>
<td>2.5</td>
<td>96.9</td>
<td>32.9 (38.7)</td>
<td>42.6</td>
</tr>
<tr>
<td>Germany</td>
<td>1.3</td>
<td>2.6</td>
<td>96.1</td>
<td>34.9 (41.0)</td>
<td>43.8</td>
</tr>
<tr>
<td>Greece</td>
<td>6.7</td>
<td>9.3</td>
<td>84.0</td>
<td>27.4 (32.3)</td>
<td>32.4</td>
</tr>
<tr>
<td>Italy</td>
<td>5.8</td>
<td>3.9</td>
<td>90.3</td>
<td>30.1 (35.3)</td>
<td>37.3</td>
</tr>
<tr>
<td>Spain</td>
<td>9.5</td>
<td>23.4</td>
<td>67.1</td>
<td>33.0 (38.8)</td>
<td>29.5</td>
</tr>
<tr>
<td>UK</td>
<td>8.9</td>
<td>1.5</td>
<td>89.7</td>
<td>35.2 (41.4)</td>
<td>41.1</td>
</tr>
<tr>
<td>Total‡</td>
<td>5.6</td>
<td>6.2</td>
<td>88.2</td>
<td>33.2 (39.0)</td>
<td>39.0</td>
</tr>
<tr>
<td><strong>Women</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Denmark</td>
<td>0.9</td>
<td>7.0</td>
<td>92.1</td>
<td>17.5 (21.5)</td>
<td>41.0</td>
</tr>
<tr>
<td>France</td>
<td>3.3</td>
<td>7.1</td>
<td>89.6</td>
<td>16.7 (20.6)</td>
<td>38.7</td>
</tr>
<tr>
<td>Germany</td>
<td>2.0</td>
<td>2.6</td>
<td>95.3</td>
<td>18.1 (22.3)</td>
<td>43.5</td>
</tr>
<tr>
<td>Greece</td>
<td>21.0</td>
<td>19.0</td>
<td>60.0</td>
<td>14.4 (17.9)</td>
<td>23.4</td>
</tr>
<tr>
<td>Italy</td>
<td>19.4</td>
<td>6.0</td>
<td>74.6</td>
<td>12.4 (15.6)</td>
<td>25.8</td>
</tr>
<tr>
<td>Netherlands</td>
<td>16.7</td>
<td>19.1</td>
<td>64.1</td>
<td>15.4 (18.1)</td>
<td>27.1</td>
</tr>
<tr>
<td>Spain</td>
<td>24.7</td>
<td>31.6</td>
<td>43.8</td>
<td>13.4 (16.7)</td>
<td>16.1</td>
</tr>
<tr>
<td>UK</td>
<td>15.2</td>
<td>2.9</td>
<td>81.9</td>
<td>17.6 (20.6)</td>
<td>37.7</td>
</tr>
<tr>
<td>Total‡</td>
<td>11.7</td>
<td>9.0</td>
<td>79.3</td>
<td>15.9 (19.7)</td>
<td>33.2</td>
</tr>
</tbody>
</table>

*Mean alcohol consumption computed among lifetime consumers.
†>24 g/day in men; >12 g/day in women.
‡Weighted average by using population size data of population from each country.

We used 10 000 Monte-Carlo simulated alcohol attributable fractions, considering the uncertainty of the hazard rate ratios, to compute the variances, standard errors, and corresponding 95% confidence intervals of the alcohol attributable fractions. We estimated the absolute number of alcohol attributable cancer cases by multiplying the alcohol attributable fractions with the total number of incidental cancer cases from 2008 derived from the GLOBOCAN 2008 project.13

The analyses were performed with SAS, version 9.2, and R, version 2.9.1.

RESULTS

Across the countries investigated, alcohol consumption followed a north south gradient with Greece and Spain having the highest proportions of never and former consumers, and Denmark and Germany having the highest proportion of lifetime consumers (table 1). This gradient was also seen for the proportions of alcohol consumption higher than the recommended upper limit with Greece and Spain showing the lowest and Germany and Denmark the highest proportions.

Among male and female lifetime consumers, the risk for all the cancers we included increased with each additional drink a day (table 2). Former consumption compared with never was associated with a considerably higher risk for total and alcohol related cancer in men. We could not compute the risk for former consumers of alcohol and upper aerodigestive tract and liver cancer in men because of low number of cases in those who had never consumed alcohol. Hence, we...
Table 2 | Adjusted hazard ratios (HRRs)* (95% confidence intervals) per 12 g/day increment for lifetime consumers and for former versus never consumers (reference category) of alcohol

<table>
<thead>
<tr>
<th>Cancer site</th>
<th>Continuous (per 12 g/day)†</th>
<th>Former consumers‡</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No of cases</td>
<td>HRR (95% CI)</td>
</tr>
<tr>
<td><strong>Men</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total cancer</td>
<td>5726</td>
<td>1.03 (1.02 to 1.04)</td>
</tr>
<tr>
<td>Alcohol related</td>
<td>1235</td>
<td>1.10 (1.07 to 1.12)</td>
</tr>
<tr>
<td>Upper aerodigestive tract</td>
<td>272</td>
<td>1.17 (1.12 to 1.23)</td>
</tr>
<tr>
<td>Colonrectum</td>
<td>859</td>
<td>1.05 (1.02 to 1.09)</td>
</tr>
<tr>
<td>Liver</td>
<td>104</td>
<td>1.13 (1.04 to 1.22)</td>
</tr>
<tr>
<td><strong>Women</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total cancer</td>
<td>12 467</td>
<td>1.03 (1.01 to 1.05)</td>
</tr>
<tr>
<td>Alcohol related</td>
<td>6671</td>
<td>1.05 (1.03 to 1.07)</td>
</tr>
<tr>
<td>Upper aerodigestive tract</td>
<td>113</td>
<td>1.25 (1.10 to 1.42)</td>
</tr>
<tr>
<td>Colonrectum</td>
<td>1245</td>
<td>1.04 (0.99 to 1.09)</td>
</tr>
<tr>
<td>Liver</td>
<td>54</td>
<td>1.09 (0.89 to 1.33)</td>
</tr>
<tr>
<td>Breast</td>
<td>5259</td>
<td>1.05 (1.02 to 1.07)</td>
</tr>
</tbody>
</table>

*Adjusted for smoking (dose and duration); education; physical activity; BMI; consumption of meat, fish, fruit and vegetables; fibre, and non-alcoholic energy intake (kJ/day); and, for women additionally, menopausal status, age at menarche, breast feeding, oral contraceptive use, hormone replacement therapy.
†Among lifetime consumers (102 648 men, 216 149 women). Log linear estimates used to establish risk functions.
‡Reference category=never consumers of alcohol.
§When there were no or a limited number of cases in reference category of never consumers, HRR for total cancer was used.

computed alcohol attributable fractions for upper aerodigestive tract and liver cancer in men based on the hazard rate ratio for former alcohol consumption and total cancer.

If we assume causality, these estimates would translate into 10% (95% confidence interval 7% to 13%) of total cancer in men (table 3) and 3.0% (1% to 5%) of total cancer in women (table 4) being attributable to alcohol consumption in these selected European countries. In both sexes the alcohol attributable fraction was highest for cancer of the upper aerodigestive tract (44% (31% to 56%) in men; 25% (5% to 46%) in women), followed by liver cancer (33% (11% to 54%) and 18% (−3% to 38%), respectively). Alcohol consumption was associated with 17% (10% to 25%) of cases of colorectal cancer in men and 4% (−1% to 10%) in women. Also, 5% (2% to 8%) of cases of breast cancer in women could be associated with total alcohol consumption. The alcohol attributable fractions varied across countries because of the differences in alcohol exposure, with relatively high alcohol attributable fractions for Spanish men compared with men in other countries. Confidence intervals of the alcohol attributable fractions, however, overlapped for all countries in both men and women.

Partial attributable fractions for alcohol consumption higher than two drinks a day in men accounted for 10% of colorectal cancer, 27% of liver cancer, and 38% of upper aerodigestive tract cancer (fig 2), which accounted for 57% to 87% of the total alcohol attributable fractions. The proportion of cancer associated with alcohol consumption higher than the recommended upper limit did not vary much by country in men, except for Greece and Spain, where partial alcohol attributable fractions were somewhat lower because of the lower proportions of men consuming more than two drinks a day. In women, partial alcohol attributable fractions accounted for 3% of colorectal cancer, 4% of breast cancer, 7% of liver cancer, and 25% of upper aerodigestive tract cancer (fig 3), which accounted for 40% to 98% of the total alcohol attributable fractions. For all cancers investigated in women, the partial alcohol attributable fraction was lowest in Spain, Greece, and Italy and highest in Germany, Denmark, and the UK. When we compared total with partial alcohol attributable fractions, a substantial part (40-98%) of the incidence of alcohol attributable cancer occurred because of alcohol consumption higher than the recommended upper limit in both men and women. The remaining part of the total alcohol attributable fraction (2-60%) was associated with consumption of less than the recommended upper limit and former consumption. In men, about three in 100 alcohol related cancer cases were associated with alcohol consumption of ≤24 g/day and more than 18 in 100 were associated with alcohol consumption >24 g/day. In women one in 100 alcohol attributable cancer cases was associated with alcohol consumption of ≤12 g/day and about four in 100 associated with alcohol consumption >12 g/day.

In terms of total numbers of cases of alcohol related cancer, and if we accept that there is a causal association between alcohol consumption and occurrence of cancer, in 2008, 33 037 of 178 578 alcohol related cancer cases in men and 17 470 of 397 043 alcohol related cancer cases in women were associated with alcohol consumption.
consumption of more than two (one for women) drinks a day. Cancer of the upper aerodigestive tract accounted for the highest number of alcohol attributable cancer cases in men (22,022 cases), with Germany showing most cases (table 5). In women, breast cancer contributed most to the number of alcohol attributable cancer cases with 12,589 cases (fig 2, table 5). The numbers of total alcohol attributable cancer cases varied considerably by country, mainly because of different population sizes in the investigated countries but also because of varying alcohol attributable fractions across the countries.

The sensitivity analysis using the alcohol consumption data in the cancer cases only had similar results to those in tables 3 and 4 (data not shown). The maximum deviation was 3 percentage points in men for alcohol related cancers (29% vs 32%) and 2 percentage points in women for upper aerodigestive tract cancer (23% vs 25%).

Given there is a causal association between alcohol consumption and risk of cancer in people who have never smoked, sensitivity analyses with the hazard rate ratios of never smokers indicated noticeable differences compared with alcohol attributable fractions that were based on hazard rate ratios adjusted for smoking from the total cohort, particularly for liver cancer, for which the alcohol attributable fraction in men who had never smoked (AAF\textsubscript{Sens}) was 78% compared with 33% in the total population, and for upper aerodigestive tract cancer, for which the AAF\textsubscript{Sens} was 14% compared with 44%. The alcohol attributable fractions for colorectal cancer in women differed by 3 percentage points with AAF\textsubscript{Sens} 1% vs 4%, which was, however, within the confidence interval computed for the alcohol attributable fraction based on estimates of the total cohort.

### DISCUSSION

If we assume causality, our analysis shows that about 10% of total cancer in men and 3% in women could be attributed to current and former alcohol consumption in the European countries included in this study. In relative terms, the alcohol attributable fraction of cancer incidence was highest for cancer of the upper aerodigestive tract, followed by liver cancer. The highest absolute number of alcohol attributable cancer cases in men was found for upper aerodigestive tract and in women for breast cancer. Furthermore, a substantial part of the alcohol attributable cancer cases were associated with consumption of more than two or one standard drinks per day for men and women, respectively.

### Comparison with other studies

Few previous studies have reported on alcohol attributable mortality\textsuperscript{37-40} or incidence\textsuperscript{41-44} of cancer. Published estimates for alcohol attributable incidence of cancer in Europe\textsuperscript{46} and France\textsuperscript{44} were of similar magnitude to our estimates. For women both higher\textsuperscript{46} and lower\textsuperscript{44} alcohol attributable fractions for single cancer sites were reported. Differences could emerge because we considered also the risk of cancer associated with former consumption of alcohol, or because of the risk functions used by one study,\textsuperscript{44} which were not derived for men and women separately,\textsuperscript{46} or because of the different application of alcohol exposure data. One British study used population means of alcohol consumption,\textsuperscript{5} while we applied gamma distributions, which better represent the right skewness of the data on alcohol consumption. Because of the various methods used to compute the alcohol attributable fractions in previous studies and because of limited data, no comparable estimates across the European countries on the alcohol attributable burden of cancer have been available until now.

Besides the total burden, we also quantified the burden of cancer incidence associated with exceeding the recommended maximal daily limit of alcohol. We found that a substantial part of this incidence was associated with consumption above the recommended upper limit, indicating the potential for cancer prevention merely by adhering to the current recommendations. For cancer sites with markedly higher risks for former compared with never consumers—such as liver cancer—a noticeable part of the total alcohol attributable fractions was associated with former consumption. That also explains why for those cancers the partial alcohol attributable fractions associated with consumption above the recommended upper limit were lower than for cancers with less strong risk estimates in former consumers, such as breast cancer.

Alcohol consumption below the recommended upper limit accounted for a modest part of the total alcohol attributable fraction of alcohol related cancers, with at least three in 100 cases of cancer in men and one in 100 cases in women. This shows that following the current recommendation would not eliminate alcohol attributable cancer incidence completely. In contrast, for all cause mortality alcohol consumption is often shown to be associated with a lower risk for up to four drinks a day in men and two drinks a day in women.\textsuperscript{47} This lower risk is probably because of the lower risk of death from cardiovascular disease.
especially coronary heart disease and ischaemic stroke.\textsuperscript{48,50} Heavy alcohol consumption above the recommended upper limit, however, was shown to be not related to\textsuperscript{48} or detrimental for\textsuperscript{16} cardiovascular diseases, whereas for cancer, as shown by many studies\textsuperscript{46-51} including ours, there is no sensible limit below which the risk of cancer is decreased. Therefore, even though light to moderate alcohol consumption might decrease the risk for cardiovascular disease and mortality, the net effect of alcohol is harmful.\textsuperscript{1} Thus, alcohol consumption should not be recommended to prevent cardiovascular disease or all cause mortality.

Sensitivity analysis

Smoking, known to be closely related to alcohol consumption, could be a potential synergistic risk factor, particularly for cancer of the upper aerodigestive tract. A possible synergistic effect modification of smoking on the risk of alcohol and cancer could lead to an over-estimation of the alcohol attributable fraction.\textsuperscript{52} We observed a substantially higher alcohol attributable fraction for liver cancer and a considerably lower alcohol attributable fraction for upper aerodigestive tract cancer in men when we applied the hazard rate ratios for alcohol consumption among never smokers. Potential effect modification by smoking was also indicated for colorectal cancer in women, for which the alcohol attributable fraction computed by using hazard rate ratios among never smokers was lower than the overall alcohol attributable fraction, but within the 95% confidence interval. In the groups of never smokers the number of cases of cancer was limited in the EPIC study, which led to a limited power to assess the association between the consumption of alcohol and risk of cancer in this subgroup. This could be one explanation why the alcohol attributable fractions differed from the originally computed overall alcohol attributable fractions. We could also have overestimated the alcohol attributable fraction for upper aerodigestive tract cancer in men because of the effect of smoking. However, a recent pooled analysis of 17 European and American case-control studies investigating people who had ever consumed alcohol compared with those who had never consumed alcohol in relation to head and neck cancer in people who had never smoked,\textsuperscript{43,52} estimated a population attributable fraction of 29.5% in men and 31.5% women. These estimates are of similar magnitude to our estimates based on results from the total EPIC study population, suggesting the estimates of our overall alcohol attributable fractions to be valid. Regarding liver cancer, smoking is a recognised causal risk factor.\textsuperscript{3} Thus, there is no plausible explanation for the substantially higher alcohol attributable fraction for liver cancer based on risk estimates computed in those who had never smoked. Therefore, the limited power because of low numbers of cases and the resulting imprecise point estimates of the hazard rate ratios in never smokers is the most plausible explanation for the considerably higher alcohol attributable fraction of liver cancer.

### Table 4: Proportion of cancer cases attributable to alcohol use in women aged ≥15 years. Figures are percentages (95% confidence interval)

<table>
<thead>
<tr>
<th>Cancer site</th>
<th>Denmark</th>
<th>France</th>
<th>Germany</th>
<th>Greece</th>
<th>Italy</th>
<th>Netherlands</th>
<th>Spain</th>
<th>UK</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total cancer</td>
<td>3 (1 to 5)</td>
<td>3 (1 to 5)</td>
<td>3 (1 to 5)</td>
<td>3 (2 to 4)</td>
<td>2 (1 to 3)</td>
<td>3 (2 to 5)</td>
<td>4 (3 to 5)</td>
<td>3 (1 to 5)</td>
<td>3 (1 to 5)</td>
</tr>
<tr>
<td>Alcohol related</td>
<td>7 (4 to 10)</td>
<td>6 (4 to 9)</td>
<td>7 (4 to 10)</td>
<td>4 (3 to 6)</td>
<td>4 (2 to 6)</td>
<td>5 (3 to 7)</td>
<td>4 (3 to 5)</td>
<td>6 (3 to 9)</td>
<td>5 (3 to 8)</td>
</tr>
<tr>
<td>Upper aerodigestive tract</td>
<td>32 (9 to 55)</td>
<td>30 (8 to 52)</td>
<td>35 (11 to 59)</td>
<td>15 (1 to 31)</td>
<td>18 (4 to 32)</td>
<td>17 (9 to 34)</td>
<td>5 (9 to 18)</td>
<td>30 (9 to 51)</td>
<td>25 (9 to 46)</td>
</tr>
<tr>
<td>Colorectum</td>
<td>5 (2 to 12)</td>
<td>5 (1 to 11)</td>
<td>6 (2 to 13)</td>
<td>4 (0 to 7)</td>
<td>3 (1 to 7)</td>
<td>4 (0 to 8)</td>
<td>3 (1 to 6)</td>
<td>5 (2 to 11)</td>
<td>4 (1 to 10)</td>
</tr>
<tr>
<td>Liver</td>
<td>18 (8 to 44)</td>
<td>17 (7 to 42)</td>
<td>15 (6 to 46)</td>
<td>24 (12 to 36)</td>
<td>13 (3 to 29)</td>
<td>24 (11 to 38)</td>
<td>31 (24 to 38)</td>
<td>13 (13 to 39)</td>
<td>18 (3 to 38)</td>
</tr>
<tr>
<td>Breast</td>
<td>6 (3 to 10)</td>
<td>6 (3 to 9)</td>
<td>7 (3 to 10)</td>
<td>4 (2 to 6)</td>
<td>4 (2 to 6)</td>
<td>4 (2 to 6)</td>
<td>3 (2 to 4)</td>
<td>5 (2 to 8)</td>
<td>5 (2 to 8)</td>
</tr>
</tbody>
</table>

### Table 5: Total number* of alcohol attributable cancer cases for general population in 2008 in selected countries

<table>
<thead>
<tr>
<th>Cancer site</th>
<th>Men</th>
<th>Women</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>1313</td>
<td>503</td>
</tr>
<tr>
<td>Alcohol related</td>
<td>996</td>
<td>4415</td>
</tr>
<tr>
<td>Upper aerodigestive tract</td>
<td>445</td>
<td>348</td>
</tr>
<tr>
<td>Colorectum</td>
<td>348</td>
<td>6305</td>
</tr>
<tr>
<td>Liver</td>
<td>64</td>
<td>1797</td>
</tr>
<tr>
<td>Breast</td>
<td>259</td>
<td>2395</td>
</tr>
</tbody>
</table>

*Numbers refer to total burden of incident cancer cases associated with current and former alcohol consumption in population. Sum of cases for single countries might not exactly add to total number of cases because of country specific estimation of numbers and separate estimation for all countries combined.
WHAT IS ALREADY KNOWN ON THIS TOPIC
Alcohol consumption has been causally related to cancers of the oral cavity, pharynx, larynx, oesophagus, liver, colorectum, and female breast
Current estimates of the alcohol attributable burden of these cancers refer mostly to current alcohol consumption and to Europe as a whole, and do not include the risk associated with previous alcohol consumption

WHAT THIS STUDY ADDS
If we assume causality between alcohol consumption and cancer, about 10% of all cancer cases in men and 3% of all cancer cases in women are attributable to current and former alcohol consumption in the investigated European countries
For cancers that are causally related to alcohol consumption, the proportions were 32% in men and 5% in women, with a substantial part (40-98%) being attributable to current alcohol consumption above the recommended upper limit of two drinks a day in men and one drink a day in women

Advantages and limitations
Our results are limited by the underlying data quality to generate the relative risk functions and by the data on alcohol exposure. The risk estimates were adjusted for several confounders—for example, dietary factors or extensive adjustment for smoking, the main confounder for the association between alcohol and risk of cancer. Different confounding in the various countries we investigated is unlikely to play a major role because the relative risk estimates were observed to be relatively homogenous across the countries. Also, the relative risk estimates are in line with results from meta-analyses and with the evaluation of the carcinogenicity of alcohol, which supports the validity of our effect estimates for alcohol consumption and risk of cancer. Loss to follow-up, though low, could have led to underestimation of the risk of cancer, as high risk or exposed people are more likely to be lost to follow-up. Recall bias is unlikely to have had an impact on the relative risk estimates as the exposure information was assessed before the cancer occurred. Also, under-reporting of alcohol consumption because of social desirability should not have affected our risk estimates, assuming that under-reporting was independent of later case and non-case status and that this did not change the ranking of the study participants. As the EPIC study population is a convenience sample, the transferability of study results to the general population could be questioned. Selective participation, however, should not impair aetiological conclusions expressed as relative risk estimates because these effect measures are both internally and externally valid.

The data on alcohol consumption were representative for the countries investigated, and they were quantified comparably across all countries. Thus, the exposure information is highly comprehensive and comparable, resulting in directly comparable alcohol attributable fractions across the selected countries. Comparison of previous country specific alcohol attributable fraction estimates was impeded as the studies used different methods and strategies to compute the alcohol attributable burden of cancer incidence. As the prevalence of alcohol exposure differs across the European countries, however, we would also expect the burden of alcohol attributable cancer incidence to differ across the countries, which is of potential interest for public health policy makers.

We have provided a systematic and comparable overview of the alcohol attributable cancer incidence for several European countries, and presented alcohol attributable fractions for causally related cancer sites based on empirical original data for both the total (current and former) alcohol consumption as well as for alcohol consumption higher than the recommended upper limit. We considered the risk from former alcohol consumption and could thus capture, in contrast with previous studies, the full burden of cancer incidence associated with alcohol consumption. The gamma distributions we used for current alcohol consumption overcome the limitation of using categorical risk estimates and categories of proportions of alcohol consumption. This is of particular value because the risk of cancer increases linearly and alcohol consumption follows a strongly right skewed distribution. The attributable burden of cancer incidence associated with consuming above the recommended upper limit illustrates the potential of avoidable cancer incidence, if the recommendations of the WCRF/AICR are followed. Until now, it was only speculated that reducing alcohol consumption to two drinks a day in men and one drink a day in women would be beneficial in terms of incidence of cancer. We have now computed quantitative measures, both relative (alcohol attributable fraction) and absolute (total number of cancer cases), for the burden associated with alcohol consumption above the recommended upper limit.

Conclusions and policy implications
In conclusion, if we assume causality between alcohol consumption and overall and specific cancer incidence, a considerable proportion of the most common and most lethal cancers is attributable to former and current alcohol consumption in the selected European countries, especially to consumption above the recommended upper limit. This strongly underlines the necessity to continue and to increase efforts to reduce alcohol consumption in Europe, both on the individual and the population level.

Contributors: All authors had full access to all of the data (including statistical reports and tables) and take responsibility for the integrity of the data and the accuracy of the data analysis. All authors MS, MMB, TP, HB, JR, TK, and ER were responsible for study concept and design. HB, TJK, K-TK, PP, AB, MD, M-DC, MS, LR, AA, PV, RT, FB, DP, AT, RK, FC-C, KO, AMT, AO, CD, M-CB-R, VB, DZ, SR, CVG, NW, MI, NS, DR, PAW, and ER acquired the data. MS, MMB, HB, TP, JR, TK, GG, PB, NEA, and PP analysed and interpreted the data. MS, MB, TP, HB, JR, TK, NEA, KO, PP, and PB drafted the manuscript, which was critically revised for important intellectual content by all the authors. MS, JR, TK, GG, TP, and MMB were responsible for the statistical analysis. MMB, JR, and HB supervised the study. MS is guarantor.

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Ethical approval: The EPIC study was approved by the IARC ethical committee and by the local ethics committees relevant for each study centre. All participants gave informed consent.

Data sharing: No additional data available.

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