Burden of cancer attributable to modifiable factors in Japan in 2015

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Abstract: This study estimated the cancer burden attributable to modifiable factors in Japan in 2015 using the best available epidemiological evidence and a standard methodology. We selected the following factors for inclusion in the estimates, namely tobacco smoking (active smoking and secondhand smoking), alcohol drinking, excess bodyweight, physical inactivity, infectious agents (*Helicobacter pylori*, hepatitis C virus, hepatitis B virus, human papilloma virus, Epstein-Barr virus, and human T-cell leukemia virus type 1), dietary intake (highly salted food, fruit, vegetables, dietary fiber, red meat, processed meat), exogenous hormone use, never breastfeeding and air pollution, given that these were considered modifiable, in theory at least. We first estimated the population attributable fraction (PAF) of each cancer attributable to these factors using representative relative risks of Japanese and the prevalence of exposures in Japanese around 2005, in consideration of the 10-year interval between exposure and cancer outcomes. Using nationwide cancer incidence and mortality statistics, we then estimated the attributable cancer incidence and mortality in 2015. We finally obtained the PAF for site-specific and total cancers attributable to all modifiable risk factors using this formula, with statistical consideration of the effect of overlap between risk factors. The results showed that 35.9% of all cancer incidence (43.4% in men and 25.3% in women) and 41.0% of all cancer mortality (49.7% in men and 26.8% in women) would be considered preventable by avoidance of these exposures. Infections and active smoking followed by alcohol drinking were the greatest contributing factors to cancer in Japan in 2015.

Keywords: cancer, modifiable factor, population attributable fraction, Japan

Introduction

The structure of disease in Japan has changed drastically over the past decades, largely due to changes in economics, demographics and lifestyle. Together with rapid aging, the transition from communicable diseases such as tuberculosis and pneumonia to non-communicable diseases such as cancer has challenged domestic health systems and public health.

Cancer has been the leading causes of death in Japan since 1981 (1). Recent statistics show that cancer

accounted for 27.3% of all deaths ($n = 1,381,093$) (31.1% of men and 23.2% of women) in 2019 (1). The national cancer registry documented 980,856 newly diagnosed cases of cancer (558,874 men and 421,964 women) in 2018, suggesting that 65.0% of men and 50.2% of women will be diagnosed with cancer at least once in their lifetime (2). For 2035, it is estimated that over 1,172,000 Japanese will be diagnosed with cancer, and 382,000 will die from it (3). Thus, effective cancer prevention and control is crucial to reducing the burden of cancer in Japan.

It has long been known that many cancers are caused by lifestyle and environmental factors that could potentially be prevented (4), suggesting the possibility of reducing the cancer burden caused by such exposures. However, the pattern of such modifiable factors and pattern of cancer differ among countries and regions, and cancer control policies should accordingly be tailored to reflect the local burden of cancer.

The proportion of preventable cancers is often referred to as the population attributable fraction (PAF). The first quantitative assessment of attributable causes of cancer mortality was reported in 1981 for the US by Doll and Peto (5). More recently, country-specific attributable causes of cancer using PAF have been reported in the US (5-7), Nordic countries (8,9), France (10-12), China (13), South Korea (14), UK (15,16), Australia (17), Canada (18), Germany (19), Brazil (20), Chile (21) and Vietnam (22) as well as Japan (23). In Japan, the burden of cancer attributable to known preventable risk factors was first assessed for the year 2005 (23), in which 55% of cancer among men and nearly 30% among women was attributed to preventable risk factors. That study also showed that tobacco smoking and infections were the major causes of cancer in Japan. Of note, the study applied to the prevalence of exposures occurring 15 years before, *i.e.* around 1990, based on a 15-year time lag between exposure and cancer outcome. This process was hampered by the limited availability of data sources for many exposures around 1990, which obliged the researchers to use less representative data or to exclude such exposures from analysis. Likewise, the number of Japanese epidemiological studies of risk assessment at that time was also limited, which led to insufficient evidence for PAF estimates. The subsequent decade saw the accumulation of additional epidemiological evidence and data on exposures of interest, and national representative data became available for many factors. In addition, the prevalence of many modifiable risk factors in Japan was shown to have dramatically changed by birth cohort. Accordingly, analysis to determine the burden of cancer attributed to such factors should be done with consideration to prevalence among birth cohorts at the time of exposure.

Hence, in this report, we estimated the cancer burden attributable to modifiable factors in Japan in 2015, using the most recent epidemiological evidence.

Methods

PAF in this project is defined as the fraction of cancer incidence or mortality that is attributable to a particular exposure and that could be avoided if the exposure were eliminated or reduced to an alternative scenario that would result in the lowest risk; or in other words, the theoretical minimum risk exposure distribution. We took several steps to accomplish this goal (Figure S1, <https://www.globalhealthmedicine.com/site/supplementaldata.html?ID=43>), namely I. Selection of risk factors with definitions of their theoretical minimum risk exposures and target cancers; II. Acquisition of essential data, namely *i*) cancer incidence and mortality statistics in Japan, *ii*) prevalence of risk factors in Japan, and *iii*) representative relative risks for each risk factors; and III. Estimation of PAF of cancer attributable to modifiable factors in Japan in 2015. Estimation of PAF was conducted first by risk factor, by specific cancer site. By adding up the obtained PAF of each factor related to each site, with statistical control of the effect of overlap between risk factors, we finally obtained the PAF for total cancers attributable to modifiable factors.

I. Selection of risk factors with definitions of their theoretical minimum risk exposures and target cancers

Selection of risk factors and related cancer sites

Risk factors included in the present PAF estimation were basically selected based on agents classified by the International Agency for Research on Cancer (IARC) as group 1 (carcinogens in humans) (24) and risk factors that were judged as "convincing", with the exception of "convincing" or "probable" for dietary factors, from the Third Expert Report, Diet, Nutrition, Physical Activity and Cancer: a Global Perspective (25). Related cancer sites were also selected from these expert reports, as well as from a report of the US Surgeon General in 2014 (26) for tobacco smoking. Additionally, we referred to domestic comprehensive evaluations, including a report by the Japanese Committee on Smoking and Health in 2016 (27), and a risk assessment by the Research Group for the Development and Evaluation of Cancer Prevention Strategies in Japan (28).

Risk factors meeting the following criteria, as applied in a previous estimate (23), were adopted into the present estimate: 1) there was evidence of a causal association between the factor and at least one type of cancer; 2) prevalence was available from national representative surveys; and 3) there were achievable alternative exposure levels that would reduce the risk.

Some established carcinogens, including some types of infectious agents such as *Schistosoma hematobium*, *Opisthorchis viverrini*, human immunodeficiency virus (HIV), and aflatoxin, were not included due to their extremely low prevalence in Japan. Furthermore, we did not include occupational exposures, or ultraviolet or radiation exposure, due to a lack of reliable prevalence data in Japan.

Theoretical minimum risk exposure level

We defined a theoretical minimum risk exposure level for each risk factor included in this estimate. Details are described in the following companion reports, with a focus on the PAF of cancer by each risk factor.

Risk factors, theoretical minimum risk exposure levels, and cancer sites considered in this estimate are

described in Table 1.

II. Acquisition of essential data

i) Cancer incidence and mortality statistics in Japan in 2015

Cancer incidence data in 2015 were estimated using the annual estimate of cancer incidence in 2013 by the Monitoring of Cancer Incidence in Japan Project (MCIJ) on the basis of data collected from population-based cancer registries in Japan (29). Estimation was done using an age and period spline model, a type which is used for short-term projections of cancer incidence in Japan (30). For some of the cancer sites, subsites, or histological subtypes which were not included in this published database, we further asked the MCIJ project to provide such detailed data, with permission for this purpose.

The sex- and age-specific incidence data for target cancers were coded by the International Statistical Classification of Diseases and Related Health Problems, 10th edition (ICD-10) (31), with the morphology code of

the International Classification of Disease for Oncology, 3rd edition (ICD-O-3) (32).

The statistical data on cancer mortality from 2015 were obtained from the vital statistics of Japan (33). We obtained sex- and age-specific mortality data by cause of death from available data sources from the Health, Labour, and Welfare Statistics Association (34). As was done for the cancer incidence data, 4-digit ICD-10 codes were used to classify the cause of death.

Table 2 shows the summary of incidence and mortality of cancer data used in this estimate in Japan in 2015.

ii) Prevalence of risk factors in Japan

The current incidence or mortality of cancer reflects the cumulative effect of past exposures. For most cancers and risk factors, the latent period between "exposure" to the risk factor and the increase in risk of the relevant cancer has not been well established. We assumed that this would be 10 years on average, and thus examined the effects of cancers occurring in 2015 from exposure to risk factors in 2005. We collected prevalence data

Table 1. Risk factors considered, theoretical minimum risk exposure level, and cancer sites considered

Risk factors	Theoretical minimum risk exposure level		Cancer site
<i>Tobacco smoking</i>	Never		
1. Active smoking			lung; oral cavity and pharynx; nasal; esophagus; stomach; colon; rectum; liver; pancreas; bladder; cervix; acute myeloid leukaemia
2. Secondhand smoking			lung
<i>Alcohol drinking</i>	None		oral cavity, pharynx, esophagus, stomach, colorectum, liver, larynx, and breast
<i>Excess body weight</i>	BMI < 23 kg/m ²		esophageal adenocarcinoma, stomach (cardia), colorectum, liver, gallbladder, pancreas, breast* (pre- and post- menopausal), ovary*, endometrium*, advanced prostate**, kidney
<i>Physical activity</i>	Regular exercise		colorectum, breast*, endometrium*
<i>Infection</i>	No infection		
1. <i>Helicobacter pylori</i>			stomach (non-cardia); MALT lymphoma
2. Hepatitis C virus (HCV)			liver
3. Hepatitis B virus (HBV)			liver
4. Human papilloma virus (HPV)			oral cavity; oropharynx; anus; penis; vulva; vagina; cervix uteri
5. Epstein-Barr virus (EBV)			nasopharynx; Burkitt lymphoma; Hodgkin lymphoma
6. Human T-cell Leukemia Virus			adult T-cell lymphoma/leukemia (ATL)
Type 1 (HTLV-1)			
<i>Dietary intake</i>			
1. Highly salted food	None		stomach
2. Fruit			lung
3. Vegetables			stomach (distal)
4. Dietary fiber	Age (yrs)	Men/Women	colon
	5-9	≥ 11.5 g/≥ 11 g	
	10-14	≥ 15 g/≥ 14.5 g	
	15-19	≥ 19.5 g/≥ 17.5 g	
	20-69	≥ 20 g/≥ 18 g	
	70+	≥ 19 g/≥ 17 g	
5. Red and processed meat	< 500 g/week		colon; rectum
<i>Breast feeding</i>	Ever		breast*; ovary*; endometrium*
<i>Exogenous hormone use</i>	Never		
1. Menopausal hormone therapy			breast*; ovary*
2. Oral contraceptive use			breast*; ovary*; endometrium*
<i>Air pollution</i>	Ambient PM 2.5 ≤ 10 µg/cm ³		lung

*Calculated for women only. **Calculated for men only.

Table 2. Incidence^a and mortality^b of all and selected sites of cancer in Japan (2015)

Cancer site	ICD-10	Men		Women		Both sexes	
		Incidence	Death	Incidence	Death	Incidence	Death
Base of Tongue	C01	654	26	38	2	692	28
Oral and mouth	C02-C06	6,409	1,844	4,544	1,467	10,954	3,311
Tonsil and Oropharynx	C09-C10	2,033	798	475	167	2,509	965
Nasopharynx	C11	581	224	277	75	858	299
Other oral pharynx	C00, C07-C08, C12-C14	4,802	2,366	1,150	411	5,952	2,777
Esophagus	C15	19,390	9,774	3,540	1,965	22,930	11,739
Stomach	C16	91,883	30,809	42,203	15,870	134,087	46,679
Colon	C18	50,394	17,063	44,310	17,275	94,703	34,338
Rectum	C19-C20	30,433	9,755	16,401	5,606	46,835	15,361
Anus	C21	521	209	472	202	993	411
Liver	C22	28,222	19,008	15,087	9,881	43,308	28,889
Gallbladder	C23	3,721	2,532	5,190	3,716	8,911	6,248
Pancreas	C25	19,523	16,186	18,524	15,680	38,046	31,866
Sino-nasal	C30-31	1,333	392	705	245	2,038	637
Larynx	C32	4,630	899	360	72	4,990	971
Lung	C33-C34	83,169	53,208	40,025	21,170	123,194	74,378
Breast	C50			84,709	13,584	84,709	13,584
Vulva	C51			867	262	867	262
Vagina	C52			363	151	363	151
Cervix	C53			11,253	2,813	11,253	2,813
Endometrium	C54			15,372	2,322	15,372	2,322
Ovary	C56			10,166	4,676	10,166	4,676
Penis	C60	412	141			412	141
Prostate	C61	82,896	11,326			82,896	11,326
Kidney	C64	13,471	3,182	5,954	1,584	19,426	4,766
Renal pelvis & ureter	C65-66	5,292	2,353	3,008	1,558	8,300	3,911
Bladder	C67	15,280	5,582	5,168	2,548	20,448	8,130
Hodgkin disease	C81	805	102	446	58	1,251	160
Non-Hodgkin Lymphoma	C82-C85, C96	15,195	6,568	12,581	5,140	27,776	11,708
Burkitt's lymphoma	C837	170	36	138	20	308	56
Adult T-cell Lymphoma/Leukemia	C915	912	446	663	506	1,575	952
Acute Myeloid leukemia	C920, C924, C925	2,970	2,968	1,983	1,808	4,953	4,776
All sites	C00-C96, C97 ^c	549,241	219,508	408,572	150,838	957,813	370,346

^aDerived from Monitoring of Cancer Incidence in Japan project. ^bDerived from the vital statistics of Japan. ^cC97 is included in death only.

of exposure to each risk factor from different sources, prioritizing Japanese national representative surveys. For exogenous hormone use (hormone replacement therapy and oral contraceptive use), the latency period was not considered since cancer risk decreases rapidly after cessation (35).

iii) Representative relative risk for each risk factor

Relative risk (RR) data were obtained from epidemiological studies identified from different sources, including PubMed, *Ichushi-web*, and other literature searches, either in English or Japanese. We used studies that reported RR and corresponding 95% confidence intervals (CIs). We employed priority ranking for the inclusion and selection of representative RRs. Among these studies, meta-analyses that included pooled analyses of Japanese populations were the most preferred source of RR. When such meta-analyses were not available, RR was derived from the most comprehensive Japanese studies. Cohort studies were prioritized over case-control studies. If no RR derived from Japanese population sources were available, the

data from other Asian populations were used, followed by non-Asian populations. Within meta- and pooled analyses where multiple RRs were available, RRs were selected based on characteristics that were most relevant to the evidence.

III. Estimation of PAF of cancer attributable to modifiable factors in Japan in 2015

Cancer site-specific PAF estimation by each risk factor

Cancer site-specific PAFs of each risk factor were calculated separately by sex and age groups and the age-specific attributed cases or deaths were aggregated to obtain the total number of attributed cases or deaths. In this study, we estimated PAF by a couple of methods. When exposure to a risk factor was dichotomous, a standard PAF formula proposed by Levin (36) was applied:

$$PAF = \frac{P(RR - 1)}{P(RR - 1) + 1}$$

where P is prevalence and RR is relative risk.

When there are multiple categories to be considered in the PAF calculation, the modified formula for κ multiple categories by Hanley was used (37):

$$PAF = \frac{\sum_{\kappa=1}^K P_{\kappa}(RR_{\kappa}-1)}{\sum_{\kappa=1}^K P_{\kappa}(RR_{\kappa}-1)+1}$$

$$\kappa = 1, 2, \dots, K$$

where P is prevalence, R is relative risk, and K is the number of categories for the risk factor.

For some risk factors, risk of cancer per unit increase in exposure and average RR for the whole population based on the average exposure, with the assumption of a log-linear relationship between exposure and risk, was calculated using the following formula (11):

$$PAF = \frac{Risk - 1}{Risk}$$

$$Risk = \exp[\ln(\text{risk per unit}) \times \text{average exposure level}]$$

For some infectious agents, PAF was calculated using the prevalence of the infectious agent in cases only and the RR associated with that infection (37):

$$PAF = Pc \times \frac{RR - 1}{RR}$$

where Pc is prevalence among cases.

Total cancer PAF estimation by individual risk factors

Cancer site-specific PAF was multiplied by the number of incidence or mortality of site-specific cancer to obtain the site-specific number of attributable cancer incidence and mortality. By summing these site-specific attributed numbers of cancer incidence and mortality, we obtained the attributed number of total cancer incidence and mortality. Total cancer PAF was then obtained by dividing the number of attributed total cancer incidence and mortality by the number of observed total cancer incidence and mortality.

Site-specific and total cancer PAF estimation considering all modifiable risk factors

Risk factor-specific PAF of each site of cancer and total cancers was further aggregated to estimate total PAF with consideration of all modifiable risk factors. Most cancers are caused by multiple risk factors, and PAF for individual risk factors for the same cancer sites can therefore overlap and add up to over 100%. To obtain the PAF of total modifiable risk factors, we took into account the degree of overlap of PAF under the assumption of independence of exposures and risk (38). Combined PAF for the two risk factors A and B can be

calculated by the following formula (38):

$$PAF_{AB} = 1 - (1 - PAF_A) \times (1 - PAF_B)$$

This formula can be generalized to more than two risk factors.

We finally obtained the PAF for site-specific and total cancers attributable to all modifiable risk factors using this formula with statistical consideration of the effect of overlap between risk factors. These PAFs were then used to estimate the number of attributable incident cases and deaths of site-specific and total cancer.

Sensitivity analysis of PAFs

Calculation of the PAF is determined by the choice of estimated exposure prevalence at the population level and the RR associated with the risk factor. Use of a point estimate of RR involves uncertainty. To account for differences in PAF due to uncertainty in the RR estimation, we performed sensitivity analyses on the PAFs by using the lower and upper bounds of the 95% CIs for the RR estimates.

Results

The estimated PAFs of total and associated cancer sites for each modifiable factor were reported in separate articles in detail (39-48), and are summarized in Table 3. Table S1 (cancer incidence) and S2 (cancer mortality) (online data, <https://www.globalhealthmedicine.com/site/supplementaldata.html?ID=42>) describe the PAF estimates for relevant cancer sites and types and total cancers for each modifiable factor. Overall, 35.9% of all cancers diagnosed in Japan in 2015 (43.4% in men and 25.3% in women) were attributable to modifiable risk factors of cancer. The corresponding figure for cancer mortality was 41.0% for both sexes (49.7% in men and 26.8% in women). Infections and active smoking are the most important modifiable factors for cancer in Japan, followed by alcohol drinking, given that 16.6% of cancer incidence is explained by infections, 15.2% by active smoking, and 6.2% by alcohol drinking. Cancer mortality showed similar figures, among which 19.6% of cancer deaths were explained by active smoking, 17.7% by infections, and 6.5% by alcohol drinking. There was a difference in the pattern of PAF by sex (Figure 1), with the major attributable factors for men being active smoking (23.6% for incidence and 29.8% for mortality), infections (18.1% for incidence and 18.5% for mortality) and alcohol drinking (8.3% for incidence and 8.8% for mortality), versus infections (14.7% for incidence and 16.5% for mortality), alcohol drinking (3.5% for incidence and 3.0% for mortality) and active smoking (4.0% for incidence and 4.7% for mortality) for women. In both sexes, other factors such as secondhand smoking, excess body weight, physical

Table 3. Number and population attributable fraction (PAF, %) of cancer incidence and mortality attributable to selected modifiable risk factors in Japan in 2015

Risk factor	Theoretical minimum risk exposure level	Incidence		Mortality	
		PAF (%)	number	PAF (%)	number
Men					
Total Number			549,241		219,508
Active smoking	Never	23.6	129,502	29.8	65,417
Secondhand smoking	Never	0.2	1,095	0.3	735
Alcohol drinking	None	8.3	45,360	8.8	19,406
Excess body weight	BMI < 23 kg/m ²	1.0	5,539	1.0	2,197
Physical inactivity	Regular exercise	1.0	5,509	0.9	1,942
Infections	No infection	18.1	99,481	18.5	40,580
Highly salted food consumption	No intake	3.0	16,249	2.5	5,571
Vegetable consumption	≥ 350 g/day	0.3	1,667	1.0	2,197
Fruit consumption	≥ 100 g/day	0.1	570	0.1	253
Dietary fiber consumption	sufficient consumption*	1.2	6,837	1.0	2,124
Red and processed meat consumption	< 500 g/week	0.0	75	0.0	16
Air pollution	Ambient PM 2.5 ≤ 10 µg/cm ³	1.6	8,788	2.6	5,682
Above all risk factors (adjusted for overlaps)		43.4	238,497	49.7	109,151
Women					
Total Number			408,572		150,838
Active smoking	Never	4.0	16,263	4.7	7,105
Secondhand smoking	Never	0.9	3,483	1.3	1,932
Alcohol drinking	None	3.5	14,477	3.0	4,522
Excess body weight	BMI < 23 kg/m ²	0.3	1,228	0.3	429
Physical inactivity	Regular exercise	1.6	6,521	0.8	1,153
Infections	No infection	14.7	59,893	16.5	24,935
Highly salted food consumption	No intake	1.6	6,497	1.7	2,510
Vegetable consumption	≥ 350 g/day	0.1	564	0.3	429
Fruit consumption	≥ 100 g/day	0.0	69	0.0	18
Dietary fiber consumption	sufficient consumption*	0.8	3,143	0.9	1,370
Red and processed meat consumption	< 500 g/week	0.0	0	0.0	0
Breastfeeding	Ever	0.3	1,232	0.1	202
Exogenous hormone use	Never	0.4	1,454	0.2	279
Air pollution	Ambient PM 2.5 ≤ 10 µg/cm ³	0.7	3,014	1.0	1,582
Above all risk factors (adjusted for overlaps)		25.3	103,237	26.8	40,495
Both sexes					
Total Number			957,813		370,346
Active smoking	Never	15.2	145,765	19.6	72,521
Secondhand smoking	Never	0.5	4,579	0.7	2,667
Alcohol drinking	None	6.2	59,838	6.5	23,929
Excess body weight	BMI < 23 kg/m ²	0.7	6,767	0.7	2,625
Physical inactivity	Regular exercise	1.3	12,030	0.8	3,095
Infections	No infection	16.6	159,374	17.7	65,515
Highly salted food consumption	No intake	2.4	22,746	2.2	8,081
Vegetable consumption	≥ 350g/day	0.2	2,231	0.7	2,625
Fruit consumption	≥ 100g/day	0.1	640	0.1	271
Dietary fiber consumption	sufficient consumption*	1.0	9,979	0.9	3,494
Red and processed meat consumption	< 500 g/week	0.0	75	0.0	16
Breastfeeding	Ever	0.1	1,232	0.1	202
Exogenous hormone use	Never	0.2	1,454	0.1	279
Air pollution	Ambient PM 2.5 ≤ 10 µg/cm ³	1.2	11,922	2.0	7,264
Above all risk factors (adjusted for overlaps)		35.9	344,230	41.0	151,672

*Theoretical minimum risk exposure level for dietary fiber consumption by age group: Men 5-9: ≥ 11.5 g, 10-14: ≥ 15 g, 15-19: ≥ 19.5 g, 20-69: ≥ 20 g, 70+: ≥ 19 g; Women 5-9: ≥ 11 g, 10-14: ≥ 14.5 g, 15-19: ≥ 17.5 g, 20-69: ≥ 18 g, 70+: ≥ 17 g. The number of both sexes is not equal to the sum of men and women due to rounding.

inactivity, highly salted food intake, fruit intake, vegetable intake, dietary fiber intake, red and processed meat intake, breast feeding, exogenous hormone use, and air pollution accounted for a small share of both cancer incidence and mortality.

Estimated PAFs of all modifiable factors aggregated

for cancer sites/types associated with one or more modifiable factors are shown in Table 4. Cancer of the uterine cervix and adult T-cell leukemia/lymphoma had the highest fractions of potentially modifiable factors (100%) for which the infectious agents were considered necessary causal factors. Other cancer sites/types with

Figure 1. Population attributable fraction (PAF, %) of cancer incidence and mortality in Japan in 2015 by exposure.

Table 4. Summary table of population attributable fraction of cancer incidence and mortality in Japan in 2015 by cancer site

Site	ICD-10	Population Attributable Fraction (%)					
		Men		Women		Both sexes	
		Incidence	Mortality	Incidence	Mortality	Incidence	Mortality
Base of Tongue	C01	72.3	71.8	58.4	56.0	68.9	68.3
Oral and mouth	C02-C06	59.8	58.6	27.6	22.9	50.8	49.4
Tonsil and Oropharynx	C09-C10	72.3	71.8	58.4	56.0	68.9	68.3
Nasopharynx	C11	82.4	82.6	83.5	83.1	83.6	83.8
Other oral pharynx	C00, C07-C08, C12-C14	58.6	57.2	24.5	19.5	48.9	47.4
Esophagus	C15	82.4	81.4	42.9	40.2	77.8	76.2
Stomach	C16	85.0	86.6	87.1	90.1	86.0	88.0
Colon	C18	42.0	40.7	15.1	15.1	31.3	33.1
Rectum	C19-C20	34.1	33.2	9.7	8.8	25.0	23.4
Anus	C21	88.0	88.0	88.0	88.0	88.0	88.0
Liver	C22	74.3	75.3	70.0	72.6	73.8	75.3
Gallbladder	C23	3.1	2.7	0.9	0.8	1.8	1.6
Pancreas	C25	26.8	26.7	7.7	7.2	17.5	17.1
Sinonasal	C30-31	48.9	48.4	19.7	17.1	38.8	36.4
Larynx	C32	77.0	75.4	13.5	11.8	73.0	71.3
Lung	C33-C34	66.3	65.6	33.2	31.2	55.2	55.4
Breast	C50			14.0	14.2	14.0	14.2
Vulva	C51			48.0	48.0	48.0	48.0
Vagina	C52			78.0	78.0	78.0	78.0
Cervix	C53			100.0	100.0	100.0	100.0
Endometrium	C54			16.1	18.5	16.1	18.5
Ovary	C56			1.0	1.1	1.0	1.1
Penis	C60	51.0	51.0			51.0	51.0
Prostate	C61	1.4	3.0			1.4	3.0
Kidney	C64	28.1	26.5	3.1	2.7	20.6	18.6
Renal pelvis & ureter	C65-C66	66.1	64.7	2.8	2.4	43.2	39.9
Bladder	C67	41.6	40.7	9.5	8.0	33.5	30.5
Hodgkin disease	C81	56.0	56.0	56.0	56.0	56.0	56.0
Non-Hodgkin Lymphoma	C82-C85, C96	3.7	3.7	3.7	3.7	3.7	3.7
Burkitt's lymphoma	C837	30.0	30.0	30.0	30.0	30.0	30.0
Adult T-cell Lymphoma/Leukemia	C915	100.0	100.0	100.0	100.0	100.0	100.0
Acute Myeloid leukemia	C920, C924, C925	20.6	21.2	5.3	4.4	14.5	14.8
All sites	ALL	43.4	49.7	25.3	26.8	35.9	41.0

Figure 2. Major attributable causes of cancer incidence-international comparison. Japan 2015 (the present estimate), Japan 2005 (23), China 2005 (13,52-55), South Korea 2009 (14,56-59), Vietnam 2018 (22), Australia 2010 (60), UK 2015 (16), US 2014 (7), Canada 2015 (18), France 2015 (12), Germany 2018 (19, 61-63), Brazil 2012 (20), Chile 2018 (21).

higher fractions tended to be associated with infectious agents and smoking.

Discussion

This report provides the latest estimates of the cancer burden attributable to modifiable factors in Japan in 2015. Compared with the first assessment for the year 2005, when data sources for the PAF estimates for many exposures occurring around 1990 were generally limited, more epidemiological evidence and data on exposures of interest have since been accumulated, and nationally representative data for the prevalence of many factors have become available. Further, the prevalence of many modifiable risk factors in Japan has dramatically changed with birth cohort; accordingly, use of appropriate prevalence by birth cohort at the time of exposure to these factors allows a more precise estimation of cancer burden.

Compared with the first PAF estimate for 2005 (23), we observed a nearly 5% reduction in the PAF of cancer incidence and mortality attributable to active smoking in men. This reduction is partly attributable to the decreased prevalence of active smoking in men over these decades (49). A similar reduction was also observed for the PAF attributable to infections. Infections such as *Helicobacter pylori* and hepatitis C and B viruses appear to be the

major contributors to cancer in Japan. Improvements in socioeconomic and hygiene status at the time of infection has definitely contributed to a reduction in prevalence, and prevalence rates estimated by birth cohort rather than age group might accordingly provide more precise estimates. In contrast, however, a notable increase in PAF was attributed to alcohol drinking in women.

International comparison clearly indicates that the PAF of infections is relatively high among Asian countries, namely Japan, China, South Korea and Vietnam; but relatively low among European countries and their migration countries, such as UK, US, France, Germany, Australia and Canada (Figure 2). Compared with the previous Japanese estimate for 2005, prevalence of major infectious agents in Japan has decreased, especially in younger birth cohorts (50,51). This decrease will in turn lead to drastic reductions in gastric and liver cancers - the major infection-related cancers in Japan - in the next couple of decades. PAF of other modifiable factors were generally similar among different countries. Of note, however, the larger difference in PAF by active smoking between men and women observed for Japan, China, and South Korea may reflect the difference in smoking rates by sex in East Asia.

Attribution of modifiable factors other than infections, active smoking, and alcohol drinking to

cancer in Japan was generally small (< 3%). Excess bodyweight, which is regarded as an important risk factor of cancers, was attributed a very small portion ($\leq 1\%$), in spite of the use of a more severe cut-off point (BMI < 23) as a theoretical minimal risk exposure level due to the average BMI levels being less than 25.

There are several limitations for these estimates. We did not include factors with insufficient exposure or relative risk data, such as solar ultraviolet radiation exposure and ionizing radiation. Also, we could not include occupational exposures due to a lack of adequate prevalence data in the Japanese general population. Given that occupational exposure in Japan is expected to be non-negligible, its inclusion would improve the PAF estimates. Furthermore, a multivariate approach is more realistic considering the multifactorial etiology of cancer in general. We took a statistical approach to account of the overlapping effect of exposures, due to the substantial lack of information for most interactions and the joint prevalence of multiple exposures.

Even though we used the best estimates of prevalence and relative risks available for Japanese and the most suitable methodology, the current estimates are focused on modifiable factors, and did not include unmodifiable factors such as genetic, socioeconomic disparity and reproductive factors which cannot be changed at the individual level. Inclusion of these factors could help provide more comprehensive estimates of attributable causes of cancer, both modifiable and non-modifiable.

In addition to the inclusion of attributable fractions of modifiable factors, national cancer control policy will be further strengthened by the inclusion of economic burden and burden of disability-adjusted life years (DALYs) under these modifiable factors. Inclusion of these variables in combination will aid understanding of the effective population intervention for each factor, and in turn lead to appropriate cancer control policy measures at the national level.

Conclusions

We estimated the cancer burden attributable to modifiable factors in Japan in 2015 using the best available epidemiological evidence. From our estimates, around 40% of cancer incidence and mortality may be preventable by avoiding exposure to known modifiable factors. Our findings clearly indicate that infections and active smoking followed by alcohol drinking are the greatest contributing factors and should be the most highly prioritized targets in current cancer control actions.

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