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Dietary fiber intake and head and neck cancer risk: a pooled analysis in the International Head and Neck Cancer Epidemiology consortium¹

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Abstract

The possible role of dietary fiber in the etiology of head and neck cancers (HNCs) is unclear. We used individual-level pooled data from 10 case-control studies (5959 cases and 12,248 controls) participating in the International Head and Neck Cancer Epidemiology (INHANCE) consortium, to examine the association between fiber intake and cancer of the oral cavity/pharynx and larynx. Odds Ratios (ORs) and their 95% Confidence Intervals (CIs) were estimated using unconditional multiple logistic regression applied to quintile categories of non-alcohol energy-adjusted fiber intake and adjusted for tobacco and alcohol use and other known or putative confounders.

Fiber intake was inversely associated with oral and pharyngeal cancer combined (OR for 5th vs. 1st quintile category = 0.49, 95% CI: 0.40-0.59; *p* for trend <0.001) and with laryngeal cancer (OR = 0.66, 95% CI: 0.54-0.82, *p* for trend <0.001). There was, however, appreciable heterogeneity of the estimated effect across studies for oral and pharyngeal cancer combined. Nonetheless, inverse associations were consistently observed for the subsites of oral and pharyngeal cancers and within most strata of the considered covariates, for both cancer sites.

Our findings from a multicenter large-scale pooled analysis suggest that, although in the presence of between-study heterogeneity, a greater intake of fiber may lower HNC risk.

Keywords

dietary fiber intake; INHANCE; head and neck cancer; laryngeal cancer; oral cavity and pharyngeal cancer

Introduction

There are more than 500,000 new cases of head and neck cancers (HNCs) diagnosed worldwide every year¹. Approximately 60% of patients are diagnosed with advanced

disease, for which prognosis is poor even with a multimodal treatment approach^{2, 3}. This situation emphasizes the importance of primary prevention of HNCs.

Although the combination of tobacco smoking and alcohol drinking accounts for approximately 80% of HNC risk^{4, 5}, a role of dietary factors in HNCs has been reported^{6–8}. Higher intakes of non-starchy vegetables, foods containing carotenoids, and fruit in general are thought to probably protect against HNC⁸.

Fruit and vegetables are rich sources of compounds that have anti-carcinogenic properties, including vitamins, minerals, fiber, and phytochemicals^{9–12}. Among them, dietary fiber could protect against cancer^{13–17}. However, the association of dietary fiber with HNC has been sparingly assessed and, to date, the evidence is still limited^{18–28}, although most^{18–23}, ^{25–28}, but not all studies²⁴ indicated an inverse association with HNC risk.

The International Head and Neck Cancer Epidemiology (INHANCE) consortium was established in 2004 to elucidate the aetiology of HNCs through pooled analyses of individual-level data on HNCs on a large scale^{29, 30}. To date, it includes 35 case-control studies, for a total of 25,478 cases and 37,111 controls³¹. Selected aspects of diet have been investigated within the consortium. Among relevant foods and food groups, an inverse association with HNC risk was found for higher intakes of fruit and vegetables, while no association was observed for some cereal and grain products³². In addition, higher intakes of selected micronutrients and food components from natural sources, like vitamin E, vitamin C, folate and carotenoids, have been previously found to reduce HNCs risk^{33–36}.

The main objective of the current paper is to assess the association between fiber intake and the risk of 2 HNC outcomes - oral and pharyngeal cancer combined and laryngeal cancer – adjusting for several confounders, including tobacco and alcohol use. Moreover, we evaluate whether the effect estimates differ by tumor subsite or in strata of selected factors, and explore the potential interaction of fiber intake with smoking and alcohol on the 2 HNC outcomes of interest.

Materials and methods

Design and subjects

Within version 1.5 of the INHANCE consortium pooled data set, 10 case-control studies collected information on total dietary fiber at the individual level^{6, 37–46}. Details on the individual studies, harmonization of data and data pooling methods have been previously described for the consortium³⁰ and are summarized in Supplemental Table 1 (Online Resource). Informed consent was obtained from study subjects. The investigations were approved by the relevant institutional review boards, according to the rules adopted in each country at the time of data collection.

Selection of subjects

Cases were included if their cancer had been originally classified as an invasive cancer of oral cavity, oropharynx, hypopharynx, oral cavity or pharynx not otherwise specified, larynx, or HNC unspecified. Cases with cancers of the salivary glands or of the nasal cavity/ear/

paranasal sinuses were excluded. The International Classification of Diseases coding used for the classification into subsites was previously specified²⁹.

We removed from our analysis: 1. cases with missing information on the site of origin of their cancer; 2. subjects with missing information on dietary fiber intake; and 3. subjects with implausible (<500 or >5500 kcal) or missing values on daily non-alcohol energy intake. Thus, the present analysis was based on a total of 18,207 subjects, with 5959 HNC cases and 12,248 controls. There was a total of 1385 oral cavity cancer cases, 1653 oropharyngeal and 571 hypopharyngeal cancer cases (2224 pharyngeal cancer cases), 805 unspecified oral cavity/pharynx cases (giving a total of 4414 oral and pharyngeal cancer cases combined), and 1545 laryngeal cancer cases.

Definition of the exposure variable

Intakes of total energy, several nutrients and food components, including fiber, were derived by combining information from study-specific food-frequency questionnaires (FFQs) assessing subject's usual diet during a reference period preceding cancer diagnosis for cases or interview for controls – with that from country-specific food composition databases^{47–50}. In detail, the current analysis considered total dietary fiber, which is fiber from foods only, and expressed its intake in grams/day. Comparability of total dietary fiber intakes was also improved by selecting intakes of fiber obtained with the enzymatic gravimetric methods [AOAC (Association of Official Analytical Chemist) 1980 or equivalents), rather than with the Englyst or Southgate ones⁵¹, when more estimates of total fiber intakes were simultaneously available in the single studies.

Finally, to adjust for the (study-specific) effect of daily energy intake excluding alcohol, we computed 'non-alcohol energy-adjusted' fiber intake within each study, on both cases and controls, based on the residual method⁵².

Statistical analysis

We estimated the odds ratios (ORs) and the corresponding 95% confidence intervals (CIs) of oral and pharyngeal cancer combined (including oral cavity, oropharyngeal, hypopharyngeal, unspecified oral cavity/pharyngeal cancer) and laryngeal cancer for quintile categories of 'non-alcohol energy-adjusted' fiber intake (calculated on both cases and controls from all studies combined) using unconditional multiple logistic regression models⁵³. Models included adjustment for age, sex, race, study center, education, cigarette smoking status, cigarette smoking intensity (number of cigarettes per day), cigarette smoking duration, cigar smoking status, pipe smoking status, alcohol drinking intensity (number of drinks per day), and the product (interaction) term for cigarette intensity and alcohol drinking (see Table 1 for a complete list of the covariate categories used). Tests for linear trend were computed referring to the median values of 'non-alcohol energy-adjusted' fiber intake within the selected quintile categories.

For oral and pharyngeal cancer combined, separate analyses were conducted by anatomical subsite (oral cavity, oro-/hypo-pharynx, and oral cavity or pharynx not otherwise specified). For both cancer sites, we carried out stratified analyses by age, sex, education, geographic region, body mass index (BMI), tobacco (cigarettes, cigars, pipes, snuff or chewing

products) smoking status, alcohol consumption (see Tables 3 and 4 for categories used), and heterogeneity between/among strata (possible effect modification) was tested using likelihood ratio tests⁵³.

In addition, we investigated potential confounding by other dietary factors, including some nutrients [total carotenoids, vitamin C, vitamin E, and iron (for the last nutrient, information was not available in the Buffalo study)], total fruits, and total vegetables.

We also carried out a sensitivity analysis, excluding each study one at a time to ensure that the magnitude of the overall estimates were not dependent on any specific study⁵⁴. In all the analyses described, when the *p*-value for heterogeneity between studies was less than 0.1, we estimated the corresponding ORs and CIs specifying a random intercept-random slope generalized linear mixed model with a logit link function and binomial family⁵⁵. We also adopted a complete-case approach to the analysis, where subjects with no missing information on the final database (including information on cancer sites, exposure and confounding variables) were considered for the analysis. However, as the Japan study did not provide information on education (3495 subjects), we defined an extra category of education with all missing values, to avoid the exclusion of the study from the analysis.

All statistical tests were two-sided. Calculations were carried out using the open-source statistical computing environment R^{56} , with its libraries "Ime4"⁵⁷ and "nnet"⁵⁸.

Results

Table 1 shows selected characteristics of HNC cases (separately for oral and pharyngeal cancer combined and for laryngeal cancer) and controls under investigation. Over 70% of the included subjects were white. The US Multicenter study provided cases of oral and pharyngeal cancer only. Cases were less educated, more often smokers and alcohol drinkers than controls.

Supplemental Table 2 describes the distribution of raw values of fiber intake across studies and in all the studies combined. Study-specific distributions were skewed to the right. The reported summary statistics showed different values across studies, with the Buffalo, Italy Multicenter, and Milan studies showing the highest values and the Los Angeles study the lowest values of fiber intake.

Table 2 provides the ORs, and the corresponding 95% CIs, for oral and pharyngeal cancer combined and for laryngeal cancer by quintile categories of fiber intake. For oral and pharyngeal cancer combined, we reported mixed-effects estimates with appreciable heterogeneity of the effect estimates across studies (*p*-value for heterogeneity <0.001); for laryngeal cancer, however, there was less heterogeneity across studies (*p*-value = 0.633), so fixed effects are reported. Fiber intake was inversely associated with oral and pharyngeal cancer risk: the OR for the highest quintile category of fiber intake compared to the lowest one was 0.49 (95% CI: 0.40–0.59) (*p*-value for linear trend <0.001). For laryngeal cancer, the OR for the highest quintile category was 0.66 (95% CI: 0.54–0.82; *p*-value for linear trend <0.001). In the influence analyses, the point estimates of the ORs of oral and pharyngeal cancer combined did not materially change after the exclusion of any study; the

detected heterogeneity between studies was similar to that of the main analysis when excluding any study at a time from the analysis. However, after excluding the Italy Multicenter study from the main analysis, the OR of laryngeal cancer was closer to unity (OR=0.96, 95% CI: 0.74–1.24 for the last quintile category) and the corresponding heterogeneity between studies was reduced.

Decreasing ORs with higher fiber intakes were also observed across oral cavity and pharyngeal cancer subsites: the OR for the highest vs lowest quintile category was 0.39 (95% CI: 0.29–0.52) for oral cavity, 0.54 (95% CI: 0.45–0.64) for oropharynx and hypopharynx combined, and 0.46 (95% CI: 0.33–0.65) for oral cavity or pharynx not otherwise specified (Supplemental Table 3 – Online Resource). In addition, the ORs for the oropharynx were similar to those of the hypopharynx site (e.g. for the highest vs lowest quintile category, OR=0.58, 95% CI: 0.48–0.70 and OR=0.55, 95% CI: 0.41–0.74, respectively) (data not shown) but, given the limited number of hypopharyngeal cancer cases, we combined the results of these subsites.

Tables 3 and 4 show the ORs of oral and pharyngeal cancer combined and laryngeal cancer by strata of selected variables. An inverse association between fiber intake and risk of either cancer site was present in most of the strata, in accordance with the main findings from Table 2. However, there was appreciable heterogeneity in risk for laryngeal cancer across strata of geographic region, with only the 3 European studies showing a moderate inverse association (OR for the 5th vs. 1st quintile category = 0.45; 95% CI: 0.34–0.60; p for heterogeneity = 0.015). An appreciable heterogeneity between studies was present for several strata in Table 3, but not for most strata in Table 4.

Figure 1 shows the combined effect of fiber intake and alcohol or tobacco consumption on oral and pharyngeal cancer combined and laryngeal cancer. No deviation from multiplicative interaction was found for either cancer site (*p*-value for interaction - Panel A: 0.123; Panel B: 0.084; Panel C: 0.612; Panel D: 0.430). In addition, when comparing never/light drinkers with a higher than median consumption of fiber, the ORs for subjects with a lower than median consumption and 5 drinks/day of alcohol were 11.45 (95% CI 9.61–13.65) for oral and pharyngeal cancer combined (Panel A), and 5.36 (95% CI 3.30–8.71) for laryngeal cancer (Panel B). Compared to never smokers with a higher than median consumption of fiber, the ORs for subjects consumption of fiber, the ORs for subjects consumption of median amount of fiber and being smokers of >20 cigarettes/day were 6.33 (95% CI 5.30–7.55) for oral and pharyngeal cancer (Panel C), and 26.34 (95% CI 18.28–37.95) for laryngeal cancer (Panel D).

In the sensitivity analyses considering additional adjustment for one extra nutrient at a time, the point estimates for quintiles of fiber intake were generally in line with the ones from the main analysis, although the ORs were higher for both cancer sites. The only exception was the adjustment by iron intake, which modified the OR of laryngeal cancer for the highest quintile category of fiber intake to 0.52 (95% CI: 0.44–0.62). With the additional adjustment by total fruit or total vegetable intake, the inverse association between dietary fiber intake and either tumor site was consistent, although the associations were less strong than in the main analysis for oral cavity and pharyngeal cancer combined.

Discussion

In this pooled analysis of 10 case–control studies providing information on dietary fiber within the INHANCE consortium – the largest dataset to date on the issue - we found inverse associations between dietary fiber intake and the risk of oral and pharyngeal cancer, as well as of laryngeal cancer. Similar results were observed across oral and pharyngeal cancer subsites and in most of the strata considered.

Several plausible mechanisms have been reported for such a favorable effect of dietary fiber^{13–16}. First, dietary fiber may reduce glycaemic load⁵⁹ and improve insulin sensitivity, favourably influencing insulin-like growth factor I (IGF-1), which is a promoter of the process of carcinogenesis at various sites⁶⁰. Second, dietary fiber appears to have an antiinflammatory role, via the production of short-chain fatty acids by gut bacteria with antiproliferative and pro-apoptosis properties^{14, 16}. Third, dietary fiber may also bind carcinogens and thereby limit their contact with upper digestive tract epithelia^{14, 17}. Finally, fiber-rich foods generally tend to have a higher content of antioxidants¹⁷. However, a higher fiber intake may simply be an indicator of a diet rich in fruit, vegetables and pulses, and whole gains, and, on this way, poorer in refined cereals, meat and animal fats, which have been positively associated with a higher HNC risk^{11, 61, 62}. Still, adjustment by fruit and vegetables, as well as selected micronutrients, only marginally affected the main results on fiber intake. Similarly, dietary fiber may simply be an indicator of a better general life-style pattern.

Some of the studies contributing data to our pooled analysis have already published separate reports on dietary fiber and HNC cancer risk^{18, 19, 25, 27}. Besides them, at least 2 other cohorts^{26, 28} and 5 case-control studies^{20–24}, most of which conducted before the 2000s, provided results on the issue. Among the most recent and largest studies, the National Institutes of Health (NIH)-AARP Diet and Health Study, a US cohort of ~500,000 elderly participants including 1867 HNC cases developed during ~11 years of follow-up, found an inverse association with fiber intake among women (OR 0.61, 95% CI: 0.42-0.89 for the upper quintile), consistent across subsites, and a weaker one among men (OR 0.88, 95% CI: 0.73–1.05)²⁸. Similarly, in a cohort of over 34,000 postmenopausal women and 169 incident cases from the Iowa Women's Health Study, an inverse association for fiber intake was observed for cancers of the upper aerodigestive tract combined (OR 0.57, 95% CI: 0.36-0.92 for the upper tertile). In detail, a higher fiber intake decreased oral and pharyngeal cancer (53 cases, OR 0.49), but not laryngeal cancer risk (21 cases, OR 1.82)²⁶. Among the earlier investigations, a Chinese case-control study including 404 matched cases of oral cancer showed a strong inverse association for total dietary fiber (OR 0.38, 95% CI: 0.19-0.74 for the upper quartile), and for dietary fiber from vegetables and fruits, but no relation with fiber from other sources²³. Reduced risks of oral and pharyngeal cancer for higher intakes of fibers were also observed in an Australian case-control study on 41 male cases (OR 0.14, 95% CI: 0.1–0.4 for the upper tertile)²², and in a US case-control study on 290 matched cases (OR 0.6, 95% CI: 0.4-0.9 for 1-standard deviation increase in intake)²¹. In addition, an inverse association with larvngeal cancer was found for energy-adjusted fiber intake in a US case-control study including 250 male cases²⁰. The only study reporting little or no association with fiber intake was conducted in Uruguay, had a case-control design and

included 133 cases of cancer of the upper aerodigestive tract, among which 33 were oral/pharyngeal cancers and 34 were laryngeal cancers²⁴.

In interpreting our findings, there were some limitations. Concerning fiber intake, we were unable to assess the separate effect of different subtypes of fiber on HNC risk. Indeed, separate information on dietary fiber from vegetables, fruits or cereal grains was available in 4 studies and that on total soluble and insoluble fiber in 5 studies only. In addition, pooling dietary data is challenging³⁴ and the various populations included in the analysis differ in many respects, including exposure to alcohol and tobacco. For these and other reasons, some heterogeneity among studies is to be expected. In our analysis, we identified heterogeneity among studies for oral and pharyngeal cancer combined and in several strata of interest, including subsites of oral and pharyngeal cancer. Our inspection of study-specific distributions, stratified and influence analyses showed also heterogeneity between European and American studies in the case of laryngeal cancer. However, it is difficult to isolate the effect of control sources (hospital-versus population-based) from that of geographic region (the 3 studies from Europe were all hospital-based and 4/6 American studies were population-based). The identified heterogeneity cannot, therefore, be attributed to selection bias and to different types of controls. In addition, our results may be biased by a nondifferential misclassification of individual intakes (i.e. due to measurement error), and by differential misclassification derived from recall bias.

Our analysis also had several strengths. The large sample size provides the opportunity to consider cancer subsites and subgroups of interest with adequate statistical power. We could control for the potential confounding effect of tobacco smoking, alcohol drinking and their interaction, using information on status, duration and intensity for smoking and intensity for alcohol. Moreover, the inverse association with fiber intake was consistent across strata of tobacco smoking and alcohol drinking. We also assessed the presence of a potential bias related to the assumption of a single unknown education level in the Japan study, by comparing fiber effect estimates in the non-Japanese study population, adjusting versus not adjusting for education (but adjusting for other covariates). This sensitivity analysis provided reassuring results, with very similar ORs for both cancer sites (e.g. for the last quintile category, OR of oral cavity and pharyngeal cancer combined = 0.49, 95% CI: 0.40-0.59 and 0.49, 95% CI: 0.40–0.59, with and without the adjustment for education level, respectively; OR of laryngeal cancer = 0.57, 95% CI: 0.48–0.68 and 0.57, 95% CI: 0.48–0.69, with and without the adjustment for education level, respectively). In addition, we applied uniform criteria to define our exposure of interest. Finally, we found that effect estimates were similar in the different tumor subsites, suggesting that the action of dietary fiber represents a general mechanism, rather than a site-specific one.

In conclusion, findings from this large-scale pooled analysis indicated that a relatively high intake of fiber may play a protective role against HNC. Future studies that examine country-specific sources of dietary fiber - including fruits, vegetables, beans, nuts, brown rice, whole-grain breads, biscuits, and pasta - are warranted to further elucidate which foods are the main determinants of the inverse association observed between fiber intake and the incidence of HNCs.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations

BMI	Body Mass Index
CI	Confidence Interval
DMV	Department of Motor Vehicles
FFQ	Food-Frequency Questionnaire
HNC	head and neck cancer
INHANCE	International Head and Neck Cancer Epidemiology consortium
L	large
Μ	medium
MSKCC	Memorial Sloan Kettering Cancer Center
NCI	National Cancer Institute
NA	not available
NE	Not estimable
NIH	National Institutes of Health
OR	Odds Ratio
S	small

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Novelty and Impact

We investigated the association between dietary fiber intake and cancers of the oral cavity/pharynx and larynx, using individual-level pooled data from the INHANCE consortium. Significant inverse associations were observed overall, across subsites, and strata, although in the presence of heterogeneity among studies. These conclusions were derived from a re-analysis of studies, which used standardized definitions of outcomes, exposure and confounders, a standardized statistical approach, and had enough power to inspect disease-exposure associations within strata.

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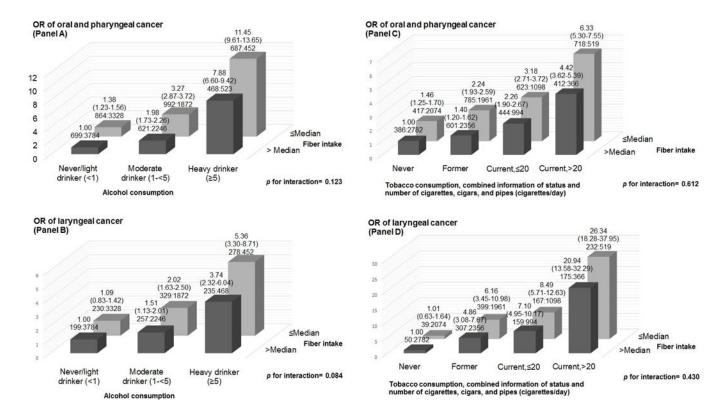


Fig 1.

Odds ratios (ORs) of oral and pharyngeal, and laryngeal cancer, and corresponding 95% confidence intervals (CIs), according to alcohol or tobacco consumption and "non-alcohol energy-adjusted" fiber intake. International Head and Neck Cancer Epidemiology (INHANCE) consortium.

(a) The odds ratios were derived from mixed-effects logistic regression models adjusted for age, sex, education, race/ethnicity, study center, combined smoking habits of cigarettes, cigars and pipes, and alcohol drinking, when appropriate.

(b) The number of cases and controls within each category was indicated below the corresponding OR as: "number of cases: number of controls."

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Distribution of cases of oral and pharyngeal cancer combined, laryngeal cancer and controls according to selected variables. International Head and Neck Cancer Epidemiology (INHANCE) consortium.

	Oral and pharyngeal cases	(%)	Laryngeal cases	(%)	Controls	(%)
Age (years)						
<40	208	4.7	26	1.7	681	5.6
>=40to<=44	194	4.4	45	2.9	563	4.6
>=45to<=49	446	10.1	123	8.0	949	7.7
>=50to<=54	645	14.6	188	12.2	1731	14.1
>=55to<=59	816	18.5	271	17.5	2079	17.0
>=60to<=64	713	16.2	290	18.8	2029	16.6
>=65to<=69	658	14.9	279	18.1	1931	15.8
>=70to<=74	474	10.7	227	14.7	1540	12.6
>=75	260	5.9	96	6.2	743	6.1
Missing	0	0.0	0	0.0	2	0.0
Sex						
Female	1187	26.9	244	15.8	3541	28.9
Male	3223	73.0	1300	84.1	8702	71.0
Missing	4	0.1	1	0.1	5	0.0
Race						
Black	387	8.8	116	7.5	535	4.4
Others (with Asians)	463	10.5	101	6.5	3089	25.2
White (with Hispanics)	3555	80.5	1324	85.7	8596	70.2
Missing	9	0.2	4	0.3	28	0.2
Study center						
Boston	313	7.1	71	4.6	611	5.0
Buffalo	396	9.0	168	10.9	1190	9.7
Italy Multicenter						
Milan	169	3.8	24	1.6	621	5.1
Pordenone	471	10.7	409	26.5	1528	12.5
Latina	95	2.2	0	0.0	425	3.5

	Oral and nharvnooal cases	(%)	I arvnoegl cases	(%)	Controls	(%)
[anan (2001–2005)	407	6.0	86	295	3002	245
Tos Angeles	246	i 9	90	0.0	808	89
Milan (2006-2000)	131	3.0	000	17.0	601	99
		0.0 F	007	<u>1.1</u>	120	0 0 0
INDACC	14	1.1	70	7.1	C71	1.0
North Carolina (2002–2006)	687	15.6	374	24.2	1120	9.1
Switzerland	367	8.3	121	7.8	877	7.2
US Multicenter						
Atlanta	129	2.9	0	0.0	134	1.1
New Jersey	467	10.6	0	0.0	459	3.7
Los Angeles	398	9.0	0	0.0	501	4.1
San Francisco	64	1.4	0	0.0	138	1.1
Education						
<= Junior high school	863	19.6	603	39.0	2723	22.2
Some high school	885	20.0	258	16.7	1240	10.1
High school graduate	588	13.3	237	15.3	1267	10.3
Technical school, some college	1174	26.6	214	13.9	2305	18.8
>= college graduate	491	11.1	145	9.4	1703	13.9
Missing	413	9.4	88	5.7	3010	24.6
Cigarette smoking status						
Never	806	18.3	91	5.8	4868	39.7
Former	1387	31.4	707	45.8	4330	35.4
Current	2210	50.1	735	47.6	2986	24.4
Missing	11	0.2	12	0.8	64	0.5
Cigarette smoking intensity (number of cigarettes/day)						
Never smoker	806	18.3	91	5.9	4868	39.7
>0to<=10	471	10.7	149	9.6	1949	15.9
>10to<=20	1466	33.2	628	40.6	3169	25.9
>20	1633	37.0	661	42.8	2137	17.4
Missing	38	0.9	16	1.0	125	1.0
Cigarette smoking duration (years)						
Never smoker	806	18.3	91	5.9	4868	39.7

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	Oral and pharyngeal cases	(%)	(%) Laryngeal cases	(%)	Controls	(%)
>0to<=20	443	10.0	102	6.6	2166	2166 17.7
>20	3132	71.0	1343	86.9	5123	41.8
Missing	33	0.7	6	0.6	91	0.7
Cigar smoking status						
Never cigar user	3583	81.2	1323	85.6	8545	69.8
Ever smoked >=100 cigars in a lifetime	394	8.9	118	7.6	636	5.2
Missing	437	9.6	104	6.7	3067	25.0
Pipe smoking status						
Never pipe user	3579	81.1	1325	85.8	8327	68.0
Ever smoked>=100 pipes in a lifetime	399	9.0	115	7.4	864	7.1
Missing	436	9.9	105	6.8	3057	25.0
Alcohol drinking intensity (number of drinks/day)						
Never drinker	548	12.4	187	12.1	3156	25.8
<1	1030	23.3	250	16.2	4022	32.8
>=lto3	973	22.0	344	22.3	2934	24.0
>=3to5	647	14.7	250	16.2	1215	9.6
>=2	1216	27.5	514	33.3	921	7.5

ABBREVIATIONS: MSKCC: Memorial Sloan Kettering Cancer Center.

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Table 2

Odds ratios (ORs)^a of oral and pharyngeal cancer combined, and laryngeal cancer and corresponding 95% confidence intervals (CIs) on fiber intake quintile categories. International Head and Neck Cancer Epidemiology (INHANCE) consortium.

	Oral and pharyngeal cases Controls OR $(95\% \text{ CI})^c$ $p_{studies}^{}d$ Laryngeal cases Controls OR $(95\% \text{ CI})^c$ $p_{studies}^{}d$	Controls	OR (95% CI) ^c	$P_{studies}^{}d$	Laryngeal cases	Controls	OR (95% CI) ^c	$p_{studies}^{d}$
I Quintile ^b	1062	1430	1430 1 (reference)		373	1430	1430 1 (reference)	
II Quintile ^b	793	1751	1751 0.71 (0.61–0.82)		262	1751	0.74 (0.60-0.90)	
III Quintile ^b	721	1889	1889 0.65 (0.57–0.75) <0.001	<0.001	252	1889	0.72 (0.58–0.88)	0.633
IV Quintile ^b	677	1986	0.55 (0.46–0.66)		251	1986	0.71 (0.57–0.87)	
V Quintile ^b	641	1940	1940 0.49 (0.40–0.59)		265	1940	1940 0.66 (0.54–0.82)	
Pfor linear trend			<0.001				<0.001	

Estimated from multiple logistic regression models adjusted for age, sex, race, study center, education, cigarette smoking status, cigarette smoking intensity (number of cigarettes per day), cigarette smoking duration, cigar smoking status, pipe smoking status, alcohol drinking intensity (number of drinks per day), and the product (interaction) term for cigarette intensity and alcohol drinking.

 b The cut-offs for the quintile categories of fiber intake were: -0.786, -0.322, 0.129, and 0.729, respectively.

^CFor the oral and pharyngeal cancer, heterogeneity between studies was appreciable and we reported the mixed-effects estimates derived from the corresponding generalized linear mixed model; for laryngeal cancer, there was less heterogeneity between studies and we reported the fixed-effects estimates.

 d P for heterogeneity between studies.

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Table 3

Odds ratios (ORs)^{a,b} of oral and pharyngeal cancer combined and corresponding 95% confidence intervals (CIs) on fiber intake quintile categories, in strata of selected covariates. International Head and Neck Cancer Epidemiology (INHANCE) consortium.

		OR (95	OR (95% CI)		
	II Quintile	III Quintile	IV Quintile	V Quintile	P studies ^C
Age (years)					
<55	0.66 (0.50-0.87)	0.57 (0.45–0.72)	0.57 (0.44–0.76)	0.60 (0.48–0.77)	<0.001
55	0.72 (0.61–0.85)	0.70 (0.59–0.82)	0.70 (0.59–0.82) 0.54 (0.45–0.66)	0.46 (0.35–0.59)	0.007
p_{strata}^{d}		0.0	0.073		
Sex					
Female	0.88 (0.68–1.14)	$0.80\ (0.60{-}1.06)$	0.80 (0.60–1.06) 0.67 (0.52–0.87)	0.57 (0.43–0.75)	0.617
Male	0.67 (0.56-0.80)	0.63 (0.54–0.73)	0.63 (0.54–0.73) 0.53 (0.43–0.66)	0.48 (0.39–0.59)	<0.001
P_{strata}^{d}		0.6	0.635		
Education					
high school graduate	0.64 (0.54–0.76)	0.66 (0.54–0.79)	0.66 (0.54–0.79) 0.55 (0.45–0.68)	0.49 (0.39–0.61)	0.00
some college	0.80 (0.65–0.97)	0.66 (0.54–0.81)	0.66 (0.54–0.81) 0.59 (0.47–0.73)	0.54 (0.43–0.69)	0.174
$p_{strata}d$		0.6	0.640		
Geographic region c					
Europe	0.56 (0.44–0.71)	0.59 (0.47–0.74)	0.48 (0.38–0.61)	0.45 (0.34–0.61)	0.017
America	0.79 (0.67–0.93)	0.69 (0.57–0.83)	0.60 (0.47–0.77)	0.50 (0.39–0.66)	<0.001
Asia	0.94 (0.69–1.27)	0.62 (0.44–0.86)	0.42 (0.29–0.61)	0.38 (0.26-0.56)	NE
p_{strata}^{d}		0.1	0.178		
Body mass index					
<25 kg/m ²	0.75 (0.62–0.92)	0.63 (0.52–0.76)	0.55 (0.45–0.67)	0.45 (0.35–0.57)	0.264
25 kg/m^2	0.69 (0.54–0.88)	0.69 (0.57–0.83)	0.58 (0.46-0.73)	0.58 (0.46–0.71)	<0.001
p_{strata}^{d}		0.1	0.117		
Tobacco smoking status					
Never user	0.75 (0.55–1.03)	0.87 (0.65–1.16)	0.87 (0.65–1.16) 0.67 (0.48–0.93)	0.63 (0.44–0.89)	0.095
Former user	0.90 (0.69–1.17)	0.65 (0.48–0.88)	0.69 (0.49–0.98)	0.65 (0.49–0.86)	0.012

		OR (95	OR (95% CI)		
	II Quintile	III Quintile	IV Quintile	V Quintile	P studies ^C
Current user	0.66 (0.54–0.82)	0.57 (0.46-0.70)	0.66 (0.54-0.82) 0.57 (0.46-0.70) 0.53 (0.42-0.65) 0.51 (0.40-0.66) 0.169	0.51 (0.40–0.66)	0.169
$p_{strata}d$		0.1	0.188		
Alcohol drinking intensity f	ity ^f				
Never/light drinker	0.86 (0.71–1.05)	0.83 (0.68–1.01)	0.86 (0.71–1.05) 0.83 (0.68–1.01) 0.71 (0.56–0.88) 0.59 (0.44–0.78)	0.59 (0.44–0.78)	0.003

		0.085	0.0		p_{strata}^{d}
0.009	0.43 (0.31–0.60)	0.45 (0.32–0.63)	0.49 (0.33-0.72) 0.46 (0.33-0.66) 0.45 (0.32-0.63) 0.43 (0.31-0.60)	0.49 (0.33–0.72)	Heavy drinker
0.007	0.42 (0.34–0.54)	0.49 (0.39–0.62)	$0.67 \; (0.52 - 0.87) 0.56 \; (0.44 - 0.72) 0.49 \; (0.39 - 0.62) 0.42 \; (0.34 - 0.54)$	0.67 (0.52–0.87)	Moderate drinker
0.003	0.59 (0.44–0.78)	0.71 (0.56–0.88)	$0.86\ (0.71-1.05) 0.83\ (0.68-1.01) 0.71\ (0.56-0.88) 0.59\ (0.44-0.78)$	$0.86\ (0.71{-}1.05)$	Never/light drinker

ABBREVIATIONS: NE: Not estimable.

smoking duration, cigar smoking status, pipe smoking status, alcohol drinking intensity (number of drinks per day), and the product (interaction) term for cigarette intensity and alcohol drinking, when a Estimated from multiple logistic regression models adjusted for age, sex, race, study center, education, cigarette smoking status, cigarette smoking intensity (number of cigarettes per day), cigarette appropriate.

 $b_{\rm The \ I}$ Quintile category was considered as the reference one.

^c for heterogeneity between studies. When the p-value was less than 0.1 within strata, we reported mixed-effects estimates derived from the corresponding generalized linear mixed model.

 d^{d} for heterogeneity across strata. When fixed-and mixed-effects models were estimated for different categories of the same stratification variable, likelihood ratio tests for heterogeneity across strata had to be based on comparable mixed-effects models and therefore we re-fitted one or more mixed-effects models to replace the original fixed-effects ones. We consistently reported the corresponding stratumspecific mixed-effects models instead of the fixed-effects ones.

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Table 4

Odds ratios (ORs)^{a,b} of laryngeal cancer and corresponding 95% confidence intervals (CIs) on fiber intake quintile categories, in strata of selected covariates. International Head and Neck Cancer Epidemiology (INHANCE) consortium.

		OR (95% CI)	(% CI)		
	II Quintile	III Quintile	IV Quintile	V Quintile	P studies ^C
Age (years)					
<55	0.98 (0.64–1.50)	0.76 (0.46–1.23)	0.65 (0.40–1.04)	0.66 (0.44–1.01)	0.131
55	0.70 (0.52–0.93)	0.67 (0.53–0.86)	0.69 (0.50-0.93)	0.60 (0.48–0.76)	0.075
p_{strata}^{d}		0.5	0.534		
Sex					
Female	0.79 (0.46–1.33)	0.81 (0.50–1.31)	0.63 (0.38–1.04)	$0.68 \ (0.41 - 1.13)$	0.243
Male	0.81 (0.59–1.11)	0.66 (0.51–0.87)	0.68 (0.48–0.96)	0.64 (0.48–0.86)	0.002
p_{strata}^{d}		0.721	21		
Education					
high school graduate	0.63 (0.48–0.82)	0.67 (0.53–0.85)	0.61 (0.45–0.85)	$0.65\ (0.48-0.88)$	0.050
some college	1.18 (0.79–1.77)	0.58 (0.32–1.04)	0.90 (0.60–1.34)	0.59 (0.39–0.89)	0.056
$p_{strata}d$		0.0	0.050		
Geographic region e					
Europe	0.55 (0.41–0.74)	0.54 (0.39–0.73)	0.52 (0.33-0.80)	$0.45\ (0.34-0.60)$	0.002
America	1.19 (0.76–1.87)	0.92 (0.63–1.36)	1.08 (0.80–1.48)	1.00 (0.74–1.37)	0.238
Asia	1.33 (0.68–2.61)	1.50 (0.78–2.91)	0.33 (0.12–0.85)	$0.85\ (0.40{-}1.83)$	NE
$p_{strata}d$		0.0	0.015		
Body mass index					
<25 kg/m ²	0.81 (0.59–1.12)	0.61 (0.44–0.84)	0.70 (0.45–1.07)	0.56 (0.37–0.85)	0.335
25 kg/m^2	0.78 (0.52–1.17)	0.70 (0.49–1.00)	0.67 (0.47–0.96)	0.66 (0.50–0.86)	0.004
p_{strata}^{d}		0.5	0.560		
Tobacco smoking status					
Never user	0.57 (0.24–1.38)	0.65 (0.27–1.56)	0.73 (0.33–1.63)	0.87 (0.39–1.94)	0.399
Former user	$0.83\ (0.58{-}1.19)$	0.73 (0.51–1.05)	0.64 (0.40–1.03)	0.81 (0.58–1.14)	0.375

		OR (95	OR (95% CI)		
	II Quintile	III Quintile	IV Quintile	V Quintile	$p_{studies}^{c}$
Current user	0.81 (0.54–1.20)	0.68 (0.47-0.99)	$0.81 \ (0.54 - 1.20) 0.68 \ (0.47 - 0.99) 0.57 \ (0.41 - 0.78) 0.60 \ (0.42 - 0.85) <0.001$	0.60 (0.42–0.85)	<0.001
$p_{strata}d$		0.6	0.669		
Alcohol drinking intensity f	ity ^f				
Never/light drinker	0.99 (0.64–1.53)	0.69 (0.39–1.23)	0.99 (0.64–1.53) 0.69 (0.39–1.23) 0.99 (0.70–1.40) 0.87 (0.60–1.25)	0.87 (0.60–1.25)	0.030

				٦
0.49 (0.33–0.73)	0.73 (0.37–1.41) 0.53 (0.35–0.81) 0.43 (0.28–0.66) 0.49 (0.33–0.73)	0.53 (0.35–0.81)	0.73 (0.37–1.41)	Heavy drinker
0.53 (0.38–0.74)	0.73 (0.53-1.00) 0.74 (0.54-1.02) 0.66 (0.43-1.01) 0.53 (0.38-0.74)	0.74 (0.54–1.02)	0.73 (0.53–1.00)	Moderate drin
0.87 (0.60–1.25)	0.99 (0.64–1.53) 0.69 (0.39–1.23) 0.99 (0.70–1.40) 0.87 (0.60–1.25)	0.69 (0.39–1.23)	0.99 (0.64–1.53)	Never/light drinker

0.130 0.236

ABBREVIATIONS: NE: Not estimable.

smoking duration, cigar smoking status, pipe smoking status, alcohol drinking intensity (number of drinks per day), and the product (interaction) term for cigarette intensity and alcohol drinking, when a Estimated from multiple logistic regression models adjusted for age, sex, race, study center, education, cigarette smoking status, cigarette smoking intensity (number of cigarettes per day), cigarette appropriate.

 $\boldsymbol{b}_{\mathrm{The}}$ I Quintile category was considered as the reference one.

^c for heterogeneity between studies. When the p-value was less than 0.1 within strata, we reported mixed-effects estimates derived from the corresponding generalized linear mixed model.

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