HHS Public Access

Author manuscript

Int J Cancer. Author manuscript; available in PMC 2018 November 01.

Published in final edited form as:

Int J Cancer. 2017 November 01; 141(9): 1811–1821. doi:10.1002/ijc.30886.

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

Daisuke Kawakita^{1,2,3}, Yuan-Chin Amy Lee⁴, Federica Turati¹, Maria Parpinel⁵, Adriano Decarli^{1,6}, Diego Serraino⁷, Keitaro Matsuo³, Andrew F. Olshan⁸, Jose P. Zevallos⁹, Deborah M. Winn¹⁰, Kirsten Moysich¹¹, Zuo-Feng Zhang¹², Hal Morgenstern¹³, Fabio Levi¹⁴, Karl Kelsey¹⁵, Michael McClean¹⁶, Cristina Bosetti¹⁷, Werner Garavello¹⁸, Stimson Schantz¹⁹, Guo-Pei Yu²⁰, Paolo Boffetta²¹, Shu-Chun Chuang²², Mia Hashibe⁴, Monica Ferraroni¹, Carlo La Vecchia¹, and Valeria Edefonti¹

¹Branch of Medical Statistics, Biometry and Epidemiology "G. A. Maccacaro", Department of Clinical Sciences and Community Health, Università degli Studi di Milano, via G. Venezian, 1, 20133 Milano, Italy ²epartment of Otorhinolaryngology, Head and Neck Surgery, Nagoya City University Graduate School of Medical Sciences, 1 Kawasumi, Mizuho-cho, Mizuho-ku, Nagoya, 467-0001, Japan ³Division of Molecular Medicine, Aichi Cancer Center Research Institute, 1-1 Kanokoden, Chikusa-ku, Nagoya, 464-8681, Japan ⁴Division of Public Health, Department of Family & Preventive Medicine and Huntsman Cancer Institute, University of Utah School of Medicine, 375 Chipeta Way, Salt Lake City, UT 84108, USA ⁵Department of Medical and Biological Sciences, University of Udine, Piazzale M. Kolbe, 4, 33100 Udine, Italy ⁶Branch of Medical Statistics, Biometry and Bioinformatics, Fondazione IRCSS Istituto Nazionale Tumori di Milano, via G. Venezian, 1, 20133, Milano, Italy ⁷Epidemiology and Biostatistics Unit, CRO Aviano National Cancer Institute, IRCCS, via F. Gallini, 2, 33081 Aviano (PN), Italy 8University of North Carolina School of Public Health, Chapel Hill, NC, USA 9Department of Otolaryngology/Head and Neck Surgery, University of North Carolina at Chapel Hill, 170 Manning Drive, Campus Box # 7070, Chapel Hill, NC 27599-7070, USA 10 Division of Cancer Control and Population Sciences, National Cancer Institute, 9609 Medical Center Drive, Bethesda, MD 20892-9764, USA 11 Roswell

Corresponding Author: Monica Ferraroni, Ph.D., Branch of Medical Statistics, Biometry and Epidemiology "G. A. Maccacaro", Department of Clinical Sciences and Community Health, Università degli Studi di Milano, via Venezian 1, 20133 Milano, Italy; monica.ferraroni@unimi.it; telephone: 0039 02-50320859; fax: 0039 02-50320866.

Conflict of interest: All the authors declare that they have no conflict of interest on the topic of the current paper.

¹Funding: The INHANCE Pooled Data Project was supported by grants from the National Institutes of Health (NIH), National Cancer Institute, (NCI) R03CA113157 and NIDCR R03DE016611. Individual studies were funded by the following grants: 1. Italy Multicenter study: Italian Association for Research on Cancer (AIRC), Italian League Against Cancer, and Italian Ministry of Research; 2. Swiss study: the Swiss Research against Cancer/Oncosuisse [KFS-700 and OCS-1633]; 3. Los Angeles study: NIH [P50CA090388, R01DA011386, R03CA077954, T32CA009142, U01CA096134, R21ES011667] and the Alper Research Program for Environmental Genomics of the UCLA Jonsson Comprehensive Cancer Center; 4. Boston study: NIH [R01CA078609, R01CA100679]; 5. US multicenter study: The Intramural Program of the NCI, NIH, United States; 6. MSKCC study: NIH [R01CA051845]; 7. Japan study (2001–2005): Scientific Research grant from the Ministry of Education, Science, Sports, Culture and Technology of Japan (17015052) and grant for the Third-Term Comprehensive 10-Year Strategy for Cancer Control from the Ministry of Health, Labor and Welfare of Japan (H20-002); 8. North Carolina (2002–2006) study: NCI R01CA90731-01 and NIEHS P30ES010126; 9. Buffalo study: -; 10. Milan study (2006–2009): Italian Association for Research on Cancer (AIRC Grant Number 10068), Italian Foundation for Cancer Research (FIRC), and Italian Ministry of Education (PRIN 2009 X8YCBN) DK was supported by JSPS Grant-in-Aid for Young Scientists (B) (No.15K21283).

Park Cancer Institute, Buffalo, NY, USA ¹²Department of Epidemiology, UCLA School of Public Health, 71-225 CHS, Box 951772, Los Angeles, CA 90095-1772, USA ¹³Departments of Epidemiology and Environmental Health Sciences, School of Public Health and Comprehensive Cancer Center, University of Michigan, Ann Arbor, MI, USA 14 Institute of Social and Preventive Medicine (IUMSP), Lausanne University Hospital (CHUV), Route de la Corniche 10, 1010 Lausanne, Switzerland ¹⁵Department of Epidemiology and Pathology and Laboratory Medicine, Brown University, 70 Ship Street, G-E3, Providence, RI 02912, USA 16 Department of Environmental Health, Boston University School of Public Health, 715 Albany Street, Talbot 4W Boston, MA 02118, USA ¹⁷Department of Epidemiology, IRCCS - Istituto di Ricerche Farmacologiche 'Mario Negri', via G. La Masa, 19, 20156 Milano, Italy 18 Department of Otorhinolaryngology, School of Medicine and Surgery, University of Milano - Bicocca, via Pergolesi, 20052 Monza, Italy ¹⁹Department of Otolaryngology, New York Eye and Ear Infirmary, 310 E. 14th Street, New York, NY 10003, USA ²⁰Medical Informatics Center, Peking University, Peking, China ²¹The Tisch Cancer Institute and Institute of Translational Epidemiology, Icahn School of Medicine at Mount Sinai, 1425 Madison Avenue, New York, NY 10029, USA 22 Institute of Population Health Sciences, National Health Research Institutes, Miaoli, Taiwan

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 R⁵⁶, with its libraries "lme4"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 vs lowest 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 consuming a lower than 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 combined (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 anti-inflammatory role, via the production of short-chain fatty acids by gut bacteria with anti-proliferative 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 laryngeal 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.

Acknowledgments

Acknowledgements: MH, MF, CLV, PB, AD, and VE designed research; KM, DS, CLV, AO, JPZ, DMW, VJ, KM, ZFZ, HM, FL, VE, CB, WG, KK, MM, SS, and GPY conducted research and provided single-study databases; SCC and YAL prepared the pooled dataset for the analysis; MP provided advice on nutritional issues; VE performed all statistical analyses; FT collected and discussed the existing literature on this topic; VE and DK wrote the paper; VE had primary responsibility for final content. All authors read and approved the final manuscript.

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

M 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

References

- Ferlay J, Shin HR, Bray F, Forman D, Mathers C, Parkin DM. Estimates of worldwide burden of cancer in 2008: GLOBOCAN 2008. International journal of cancer. 2010; 127:2893–917. [PubMed: 21351269]
- Seiwert TY, Cohen EE. State-of-the-art management of locally advanced head and neck cancer. British journal of cancer. 2005; 92:1341–8. [PubMed: 15846296]
- 3. Garavello W, Bertuccio P, Levi F, Lucchini F, Bosetti C, Malvezzi M, Negri E, La Vecchia C. The oral cancer epidemic in central and eastern Europe. International journal of cancer. 2010; 127:160–71. [PubMed: 19882710]
- Zeka A, Gore R, Kriebel D. Effects of alcohol and tobacco on aerodigestive cancer risks: a metaregression analysis. Cancer causes & control: CCC. 2003; 14:897–906. [PubMed: 14682447]

5. Hashibe M, Brennan P, Chuang SC, Boccia S, Castellsague X, Chen C, Curado MP, Dal Maso L, Daudt AW, Fabianova E, Fernandez L, Wunsch-Filho V, et al. Interaction between tobacco and alcohol use and the risk of head and neck cancer: pooled analysis in the International Head and Neck Cancer Epidemiology Consortium. Cancer epidemiology, biomarkers & prevention: a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology. 2009; 18:541–50.

- Bravi F, Bosetti C, Filomeno M, Levi F, Garavello W, Galimberti S, Negri E, La Vecchia C. Foods, nutrients and the risk of oral and pharyngeal cancer. British journal of cancer. 2013; 109:2904–10.
 [PubMed: 24149181]
- 7. Edefonti V, Bravi F, La Vecchia C, Randi G, Ferraroni M, Garavello W, Franceschi S, Talamini R, Boffetta P, Decarli A. Nutrient-based dietary patterns and the risk of oral and pharyngeal cancer. Oral oncology. 2010; 46:343–8. [PubMed: 20226721]
- World Cancer Research Fund/American Institute for Cancer Research. Food, Nutrition, Physical Activity and the Prevention of Cancer: A Global Perspective. AICR; Washington, DC: 2007.
- 9. Pavia M, Pileggi C, Nobile CG, Angelillo IF. Association between fruit and vegetable consumption and oral cancer: a meta-analysis of observational studies. The American journal of clinical nutrition. 2006; 83:1126–34. [PubMed: 16685056]
- 10. Riboli E, Norat T. Epidemiologic evidence of the protective effect of fruit and vegetables on cancer risk. The American journal of clinical nutrition. 2003; 78:559S–69S. [PubMed: 12936950]
- 11. Lucenteforte E, Garavello W, Bosetti C, La Vecchia C. Dietary factors and oral and pharyngeal cancer risk. Oral oncology. 2009; 45:461–7. [PubMed: 18990606]
- Freedman ND, Park Y, Subar AF, Hollenbeck AR, Leitzmann MF, Schatzkin A, Abnet CC. Fruit and vegetable intake and head and neck cancer risk in a large United States prospective cohort study. International journal of cancer. 2008; 122:2330–6. [PubMed: 18092323]
- Ferguson LR, Chavan RR, Harris PJ. Changing concepts of dietary fiber: implications for carcinogenesis. Nutrition and cancer. 2001; 39:155–69. [PubMed: 11759275]
- 14. Kaczmarczyk MM, Miller MJ, Freund GG. The health benefits of dietary fiber: beyond the usual suspects of type 2 diabetes mellitus, cardiovascular disease and colon cancer. Metabolism: clinical and experimental. 2012; 61:1058–66. [PubMed: 22401879]
- 15. Latino-Martel P, Cottet V, Druesne-Pecollo N, Pierre FH, Touillaud M, Touvier M, Vasson MP, Deschasaux M, Le Merdy J, Barrandon E, Ancellin R. Alcoholic beverages, obesity, physical activity and other nutritional factors, and cancer risk: A review of the evidence. Critical reviews in oncology/hematology. 2016; 99:308–23. [PubMed: 26811140]
- 16. Moore MA, Park CB, Tsuda H. Soluble and insoluble fiber influences on cancer development. Critical reviews in oncology/hematology. 1998; 27:229–42. [PubMed: 9649935]
- Slavin J, Jacobs D, Marquart L. Whole-grain consumption and chronic disease: protective mechanisms. Nutrition and cancer. 1997; 27:14–21. [PubMed: 8970176]
- McLaughlin JK, Gridley G, Block G, Winn DM, Preston-Martin S, Schoenberg JB, Greenberg RS, Stemhagen A, Austin DF, Ershow AG, et al. Dietary factors in oral and pharyngeal cancer. Journal of the National Cancer Institute. 1988; 80:1237–43. [PubMed: 3418729]
- Gridley G, McLaughlin JK, Block G, Blot WJ, Winn DM, Greenberg RS, Schoenberg JB, Preston-Martin S, Austin DF, Fraumeni JF Jr. Diet and oral and pharyngeal cancer among blacks. Nutrition and cancer. 1990; 14:219–25. [PubMed: 2084618]
- Freudenheim JL, Graham S, Byers TE, Marshall JR, Haughey BP, Swanson MK, Wilkinson G. Diet, smoking, and alcohol in cancer of the larynx: a case-control study. Nutrition and cancer. 1992; 17:33–45. [PubMed: 1574443]
- 21. Marshall JR, Graham S, Haughey BP, Shedd D, O'Shea R, Brasure J, Wilkinson GS, West D. Smoking, alcohol, dentition and diet in the epidemiology of oral cancer. European journal of cancer Part B, Oral oncology. 1992; 28B:9–15.
- 22. Kune GA, Kune S, Field B, Watson LF, Cleland H, Merenstein D, Vitetta L. Oral and pharyngeal cancer, diet, smoking, alcohol, and serum vitamin A and beta-carotene levels: a case-control study in men. Nutrition and cancer. 1993; 20:61–70. [PubMed: 8415131]

23. Zheng T, Boyle P, Willett WC, Hu H, Dan J, Evstifeeva TV, Niu S, MacMahon B. A case-control study of oral cancer in Beijing, People's Republic of China. Associations with nutrient intakes, foods and food groups. European journal of cancer Part B, Oral oncology. 1993; 29B:45–55.

- 24. De Stefani E, Ronco A, Mendilaharsu M, Deneo-Pellegrini H. Diet and risk of cancer of the upper aerodigestive tract-II. Nutrients Oral oncology. 1999; 35:22–6. [PubMed: 10211306]
- Soler M, Bosetti C, Franceschi S, Negri E, Zambon P, Talamini R, Conti E, La Vecchia C. Fiber intake and the risk of oral, pharyngeal and esophageal cancer. International journal of cancer. 2001; 91:283–7. [PubMed: 11169948]
- 26. Kasum CM, Jacobs DR Jr, Nicodemus K, Folsom AR. Dietary risk factors for upper aerodigestive tract cancers. International journal of cancer. 2002; 99:267–72. [PubMed: 11979443]
- 27. Pelucchi C, Talamini R, Levi F, Bosetti C, La Vecchia C, Negri E, Parpinel M, Franceschi S. Fibre intake and laryngeal cancer risk. Annals of oncology: official journal of the European Society for Medical Oncology/ESMO. 2003; 14:162–7.
- 28. Lam TK, Cross AJ, Freedman N, Park Y, Hollenbeck AR, Schatzkin A, Abnet C. Dietary fiber and grain consumption in relation to head and neck cancer in the NIH-AARP Diet and Health Study. Cancer causes & control: CCC. 2011; 22:1405–14. [PubMed: 21785948]
- 29. Hashibe M, Brennan P, Benhamou S, Castellsague X, Chen C, Curado MP, Dal Maso L, Daudt AW, Fabianova E, Fernandez L, Wunsch-Filho V, Franceschi S, et al. Alcohol drinking in never users of tobacco, cigarette smoking in never drinkers, and the risk of head and neck cancer: pooled analysis in the International Head and Neck Cancer Epidemiology Consortium. Journal of the National Cancer Institute. 2007; 99:777–89. [PubMed: 17505073]
- Conway DI, Hashibe M, Boffetta P, consortium I. Wunsch-Filho V, Muscat J, La Vecchia C, Winn DM. Enhancing epidemiologic research on head and neck cancer: INHANCE - The international head and neck cancer epidemiology consortium. Oral oncology. 2009; 45:743–6. [PubMed: 19442571]
- 31. Winn DM, Lee YC, Hashibe M, Boffetta P. The INHANCE consortium: toward a better understanding of the causes and mechanisms of head and neck cancer. Oral diseases. 2015; 21:685–93. [PubMed: 25809224]
- 32. Chuang SC, Jenab M, Heck JE, Bosetti C, Talamini R, Matsuo K, Castellsague X, Franceschi S, Herrero R, Winn DM, La Vecchia C, Morgenstern H, et al. Diet and the risk of head and neck cancer: a pooled analysis in the INHANCE consortium. Cancer causes & control: CCC. 2012; 23:69–88.
- 33. Edefonti V, Hashibe M, Parpinel M, Ferraroni M, Turati F, Serraino D, Matsuo K, Olshan AF, Zevallos JP, Winn DM, Moysich K, Zhang ZF, et al. Vitamin E intake from natural sources and head and neck cancer risk: a pooled analysis in the International Head and Neck Cancer Epidemiology consortium. British journal of cancer. 2015; 113:182–92. [PubMed: 25989276]
- 34. Edefonti V, Hashibe M, Parpinel M, Turati F, Serraino D, Matsuo K, Olshan AF, Zevallos JP, Winn DM, Moysich K, Zhang ZF, Morgenstern H, et al. Natural vitamin C intake and the risk of head and neck cancer: A pooled analysis in the International Head and Neck Cancer Epidemiology Consortium. International journal of cancer. 2015; 137:448–62. [PubMed: 25627906]
- 35. Galeone C, Edefonti V, Parpinel M, Leoncini E, Matsuo K, Talamini R, Olshan AF, Zevallos JP, Winn DM, Jayaprakash V, Moysich K, Zhang ZF, et al. Folate intake and the risk of oral cavity and pharyngeal cancer: a pooled analysis within the International Head and Neck Cancer Epidemiology Consortium. International journal of cancer. 2015; 136:904–14. [PubMed: 24974959]
- 36. Leoncini E, Edefonti V, Hashibe M, Parpinel M, Cadoni G, Ferraroni M, Serraino D, Matsuo K, Olshan AF, Zevallos JP, Winn DM, Moysich K, et al. Carotenoid intake and head and neck cancer: a pooled analysis in the International Head and Neck Cancer Epidemiology Consortium. European journal of epidemiology. 2015
- 37. Blot WJ, McLaughlin JK, Winn DM, Austin DF, Greenberg RS, Preston-Martin S, Bernstein L, Schoenberg JB, Stemhagen A, Fraumeni JF Jr. Smoking and drinking in relation to oral and pharyngeal cancer. Cancer research. 1988; 48:3282–7. [PubMed: 3365707]
- 38. Schantz SP, Zhang ZF, Spitz MS, Sun M, Hsu TC. Genetic susceptibility to head and neck cancer: interaction between nutrition and mutagen sensitivity. The Laryngoscope. 1997; 107:765–81. [PubMed: 9185733]

39. Levi F, Pasche C, La Vecchia C, Lucchini F, Franceschi S, Monnier P. Food groups and risk of oral and pharyngeal cancer. International journal of cancer. 1998; 77:705–9. [PubMed: 9688303]

- 40. Bosetti C, Gallus S, Trichopoulou A, Talamini R, Franceschi S, Negri E, La Vecchia C. Influence of the Mediterranean diet on the risk of cancers of the upper aerodigestive tract. Cancer epidemiology, biomarkers & prevention: a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology. 2003; 12:1091–4.
- 41. Peters ES, McClean MD, Liu M, Eisen EA, Mueller N, Kelsey KT. The ADH1C polymorphism modifies the risk of squamous cell carcinoma of the head and neck associated with alcohol and tobacco use. Cancer epidemiology, biomarkers & prevention: a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology. 2005; 14:476–82.
- 42. Cui Y, Morgenstern H, Greenland S, Tashkin DP, Mao J, Cao W, Cozen W, Mack TM, Zhang ZF. Polymorphism of Xeroderma Pigmentosum group G and the risk of lung cancer and squamous cell carcinomas of the oropharynx, larynx and esophagus. International journal of cancer. 2006; 118:714–20. [PubMed: 16094634]
- 43. Hashibe M, Morgenstern H, Cui Y, Tashkin DP, Zhang ZF, Cozen W, Mack TM, Greenland S. Marijuana use and the risk of lung and upper aerodigestive tract cancers: results of a population-based case-control study. Cancer epidemiology, biomarkers & prevention: a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology. 2006; 15:1829–34.
- 44. Jayaprakash V, Rigual NR, Moysich KB, Loree TR, Nasca MA, Menezes RJ, Reid ME. Chemoprevention of head and neck cancer with aspirin: a case-control study. Archives of otolaryngology-head & neck surgery. 2006; 132:1231–6. [PubMed: 17116820]
- 45. Suzuki T, Wakai K, Matsuo K, Hirose K, Ito H, Kuriki K, Sato S, Ueda R, Hasegawa Y, Tajima K. Effect of dietary antioxidants and risk of oral, pharyngeal and laryngeal squamous cell carcinoma according to smoking and drinking habits. Cancer science. 2006; 97:760–7. [PubMed: 16800818]
- 46. Divaris K, Olshan AF, Smith J, Bell ME, Weissler MC, Funkhouser WK, Bradshaw PT. Oral health and risk for head and neck squamous cell carcinoma: the Carolina Head and Neck Cancer Study. Cancer causes & control: CCC. 2010; 21:567–75. [PubMed: 20049634]
- 47. Dresser, CM. From nutrient data to a data base for a health and nutrition examination survey. Organization, coding and values-real or imputed. Proceeding of the 8th National Nutrient Data Base Conference Minneapolis; MN, USA. 1983; p. 92-104.
- 48. US Department of Agriculture (USDA), Agricultural Research Service. USDA National Nutrient Database for Standard Reference, Release 26 and previous versions. Nutrient Data Laboratory Home Page. http://www.ars.usda.gov/Services/docs.htm?docid=8964
- 49. Resource Council, Science and Technology Agency, the Government of Japan. Standard Tables of Food Composition in Japan, 5th Revised Version (in Japanese with English translation). Ministry of Finance Printing Bureau; Tokyo, Japan: 2000.
- 50. Gnagnarella, P., Salvini, S., Parpinel, M. Food Composition Database for Epidemiological Studies in Italy. Version 1.2015. Available online at: http://www.bda-ieo.it/
- Greenfield, H., Southgate, DAT. Food composition data Production, Management and Use. 2nd. Rome, Italy: FAO/WHO; 2003.
- 52. Willett W, Stampfer MJ. Total energy intake: implications for epidemiologic analyses. American journal of epidemiology. 1986; 124:17–27. [PubMed: 3521261]
- Hosmer, DW., Lemeshow, S. Applied Logistic Regression. 2nd. New York, NY: John Wiley & Sons, Inc; 2000.
- 54. Chapter 9: Analysing data and undertaking meta-analyses. In: Deeks, JJ.H, J., Altman, DG., editors; Higgins, JPT., Green, S., editors. Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0: The Cochrane Collaboration. 2011.
- 55. Pinheiro, JC., B, D. Mixed-Effects Models in S and S-PLUSed. New York, NY: Springer-Verlag; 2000
- 56. R Development Core Team. R: A Language and Environment for Statistical Computing 2016. Vienna, Austria: R Foundation for Statistical Computing; http://www.R-project.org (27 July 2016, date last accessed)

- 57. Bates D, M M, Bolker B. Ime4: linear mixed-effects models using S4 classes. 2011
- 58. Venables, WN., R, B. Modern Applied Statistics with S. 4th. New York, NY: Springer; 2002.
- 59. Augustin LS, Kendall CW, Jenkins DJ, Willett WC, Astrup A, Barclay AW, Bjorck I, Brand-Miller JC, Brighenti F, Buyken AE, Ceriello A, La Vecchia C, et al. Glycemic index, glycemic load and glycemic response: An International Scientific Consensus Summit from the International Carbohydrate Quality Consortium (ICQC). Nutr Metab Cardiovasc Dis. 2015; 25:795–815. [PubMed: 26160327]
- 60. Yu H, Rohan T. Role of the insulin-like growth factor family in cancer development and progression. Journal of the National Cancer Institute. 2000; 92:1472–89. [PubMed: 10995803]
- 61. Franceschi S, Favero A, Conti E, Talamini R, Volpe R, Negri E, Barzan L, La Vecchia C. Food groups, oils and butter, and cancer of the oral cavity and pharynx. British journal of cancer. 1999; 80:614–20. [PubMed: 10408875]
- 62. Bosetti C, La Vecchia C, Talamini R, Negri E, Levi F, Dal Maso L, Franceschi S. Food groups and laryngeal cancer risk: a case-control study from Italy and Switzerland. International journal of cancer. 2002; 100:355–60. [PubMed: 12115553]

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.

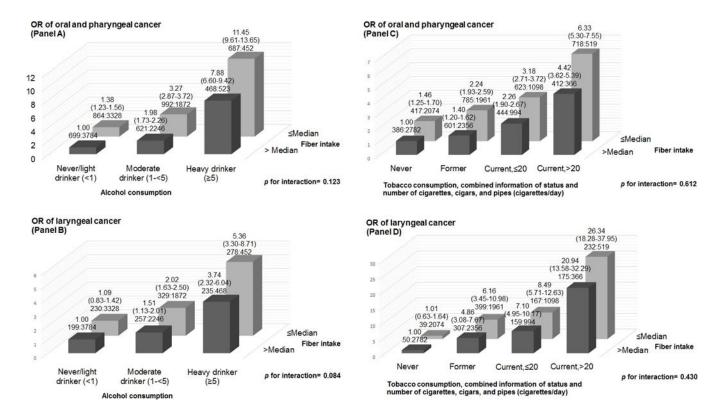


Fig 1.
Odds ratios (ORs) of oral and pharyngeal, and laryngeal cancer, and corresponding 95% confidence intervals (CIs), according to alcohol or tobacco consumption and "non-alcohol energy-adjusted" fiber intake. International Head and Neck Cancer Epidemiology (INHANCE) consortium.

- (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."

Table 1

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 1 | 12.2 | 1731 | 14.1 |
| >=55to<=59 | 816 | 18.5 | 271 17 | 17.5 | 2079 | 17.0 |
| >=60to<=64 | 713 | 16.2 | 290 | 18.8 | 2029 | 16.6 |
| >=65to<=69 | 859 | 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 1: | 15.8 | 3541 | 28.9 |
| Male | 3223 | 73.0 | 1300 8 | 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 8 | 85.7 | 8596 | 70.2 |
| Missing | 6 | 0.2 | 4 | 0.3 | 28 | 0.2 |
| Study center | | | | | | |
| Boston | 313 | 7.1 | 711 | 4.6 | 611 | 5.0 |
| Buffalo | 396 | 9.0 | 168 1 | 10.9 | 1190 | 9.7 |
| Italy Multicenter | | | | | | |
| Milan | 169 | 3.8 | 24 | 1.6 | 621 | 5.1 |
| Pordenone | 471 | 10.7 | 409 2 | 26.5 | 1528 | 12.5 |
| Latina | 95 | 2.2 | 0 | 0.0 | 425 | 3.5 |

Kawakita et al.

| | Oral and pharyngeal cases | (%) | Laryngeal cases | (%) | Controls | (%) |
|--|---------------------------|------|-----------------|------|----------|------|
| Japan (2001–2005) | 407 | 9.2 | 98 | 5.6 | 3002 | 24.5 |
| Los Angeles | 246 | 5.6 | 09 | 3.9 | 828 | 8.9 |
| Milan (2006–2009) | 131 | 3.0 | 200 | 12.9 | 691 | 5.6 |
| MSKCC | 74 | 1.7 | 32 | 2.1 | 123 | 1.0 |
| North Carolina (2002–2006) | 289 | 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 | 908 | 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 | 49 | 0.5 |
| Cigarette smoking intensity (number of cigarettes/day) | | | | | | |
| Never smoker | 908 | 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 | 808 | 18.3 | 91 | 5.9 | 4868 | 39.7 |
| | | | | | | |

| | Oral and pharyngeal cases | (%) | (%) Laryngeal cases | (%) | Controls | (%) |
|---|---------------------------|------|---------------------|------|----------|------|
| >0to<=20 | 443 | 10.0 | 102 | 9.9 | 2166 | 17.7 |
| >20 | 3132 | 71.0 | 1343 | 86.9 | 5123 | 41.8 |
| Missing | 33 | 0.7 | 6 | 9.0 | 91 | 0.7 |
| Cigar smoking status | | | | | | |
| Never cigar user | 3583 | 81.2 | 1323 | 85.6 | 8545 | 8.69 |
| Ever smoked >=100 cigars in a lifetime | 394 | 8.9 | 118 | 7.6 | 636 | 5.2 |
| Missing | 437 | 6.6 | 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 | 0.6 | 115 | 7.4 | 864 | 7.1 |
| Missing | 436 | 6.6 | 105 | 8.9 | 3057 | 25.0 |
| Alcohol drinking intensity (number of drinks/day) | | | | | | |
| Never drinker | 548 | 12.4 | 187 | 12.1 | 3156 | 25.8 |
| $\overline{\lor}$ | 1030 | 23.3 | 250 | 16.2 | 4022 | 32.8 |
| >=1to3 | 973 | 22.0 | 344 | 22.3 | 2934 | 24.0 |
| >=3to5 | 647 | 14.7 | 250 | 16.2 | 1215 | 9.9 |
| >= ? | 1216 | 27.5 | 514 | 33.3 | 921 | 7.5 |

ABBREVIATIONS: MSKCC: Memorial Sloan Kettering Cancer Center.

Author Manuscript

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% CI) $^{\mathcal{C}}$ $p_{studies}^{d}$ Laryngeal cases Controls OR (95% CI) $^{\mathcal{C}}$ $p_{studies}^{d}$ | Controls | OR $(95\% \text{ CI})^c$ | $p_{studies}^{d}$ | Laryngeal cases | Controls | OR $(95\% \text{ CI})^{\mathcal{C}}$ | $p_{studies}^{}$ |
|-------------------|---|----------|--------------------------|-------------------|-----------------|----------|--------------------------------------|------------------|
| I Quintile b | 1062 | 1430 | 1430 1 (reference) | | 373 | 1430 | 1430 1 (reference) | |
| II Quintile b | 793 | 1751 | 0.71 (0.61–0.82) | | 262 | 1751 | 0.74 (0.60–0.90) | |
| III Quintile b | 721 | 1889 | 0.65 (0.57-0.75) | <0.001 | 252 | 1889 | 0.72 (0.58-0.88) | 0.633 |
| IV Quintile b | <i>LL</i> 9 | 1986 | 0.55 (0.46–0.66) | | 251 | 1986 | 0.71 (0.57–0.87) | |
| V Quintile b | 641 | 1940 | 0.49 (0.40-0.59) | | 265 | 1940 | 0.66 (0.54–0.82) | |
| Ptor linear trend | | | <0.001 | | | | <0.001 | |

astimated 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.

Crothe 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. Page 19

 $^{\it d}_{\it P}$ for heterogeneity between studies.

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 (9 | 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) | 0.54 (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) | 0.67 (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) | 0.53 (0.43–0.66) | 0.48 (0.39–0.59) | <0.001 |
| p_{strata}^{d} | | 0.0 | 0.635 | | |
| Education | | | | | |
| high school graduate | 0.64 (0.54–0.76) | 0.66 (0.54-0.79) | 0.55 (0.45–0.68) | 0.49 (0.39–0.61) | 0.009 |
| some college | 0.80 (0.65-0.97) | 0.66 (0.54-0.81) | 0.59 (0.47–0.73) | 0.54 (0.43–0.69) | 0.174 |
| ρ_{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 |
| ρ_{strata}^{d} | | 0.1 | 0.178 | | |
| Body mass index | | | | | |
| $<25 \text{ kg/m}^2$ | 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 |
| $25 \mathrm{kg/m^2}$ | 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) | 0.67 (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 |
| | | | | | |

Author Manuscript

| | | OR (99 | OR (95% CI) | | |
|---------------------------------|-------------------------|------------------|---|------------------|-----------|
| | II Quintile | III Quintile | IV Quintile | V Quintile | P studies |
| 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) | 0.51 (0.40–0.66) | 0.169 |
| $p_{strata}^{}^{}$ | | 0.1 | 0.188 | | |
| Alcohol drinking intensity f | sity^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 |
| Moderate drinker | 0.67 (0.52–0.87) | 0.56 (0.44–0.72) | $0.67 \ (0.52-0.87) \ 0.56 \ (0.44-0.72) \ 0.49 \ (0.39-0.62) \ 0.42 \ (0.34-0.54)$ | 0.42 (0.34–0.54) | 0.007 |
| Heavy drinker | 0.49 (0.33-0.72) | 0.46 (0.33–0.66) | 0.49 (0.33-0.72) 0.46 (0.33-0.66) 0.45 (0.32-0.63) 0.43 (0.31-0.60) | 0.43 (0.31–0.60) | 0.009 |
| P _{strata} d | | 0.0 | 0.085 | | |

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 astimated 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.

 $\stackrel{b}{h}$ The I Quintile category was considered as the reference one.

CP 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 Por 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.

e Europe included Italy Multicenter, Switzerland and Milan (2006–2009) studies. North America included Boston, Buffalo, Los Angeles, Memorial Sloan Kettering Cancer Center, North Carolina (2002– 2006), and US Multicenter studies. Asia included Japan study only. As Asia included Japan study only, there was no possibility to assess heterogeneity between studies in the Asia stratum. f The never/light drinker category included never drinkers and subjects who drink less than 1 drink per day; the moderate drinker category included subjects drinking between 1 (included) and 5 drinks per day; the heavy drinker category included subjects drinking 5 drinks per day or more.

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.

| | II Quintile | III Quintile | IV Quintile | V Quintile | Pstudies |
|----------------------------|------------------|------------------|------------------|------------------|----------|
| 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} | | 0.7 | 0.721 | | |
| 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 |
| $p_{strata}^{}$ | | 0.0 | 0.050 | | |
| ${f Geographic\ region}^c$ | | | | | |
| 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 | | | | | |
| $<25 \text{ kg/m}^2$ | 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 |
| $25 \mathrm{kg/m^2}$ | 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 |
| $P_{strata}^{}^{}$ | | 0.6 | 699.0 | | |
| Alcohol drinking intensity $^{\it f}$ | 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 |
| Moderate drin | 0.73 (0.53-1.00) | 0.74 (0.54–1.02) | 0.66 (0.43–1.01) 0.53 (0.38–0.74) | 0.53 (0.38–0.74) | 0.130 |
| Heavy drinker | 0.73 (0.37–1.41) | 0.53 (0.35-0.81) | $0.73\ (0.37-1.41) \qquad 0.53\ (0.35-0.81) \qquad 0.43\ (0.28-0.66) \qquad 0.49\ (0.33-0.73)$ | 0.49 (0.33–0.73) | 0.236 |
| $p_{strata}^{}$ | | 0.3 | 0.383 | | |

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 astimated 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 The I Quintile category was considered as the reference one.

CP 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 Por 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.

e Europe included Italy Multicenter, Switzerland and Milan (2006–2009) studies. North America included Boston, Buffalo, Los Angeles, Memorial Sloan Kettering Cancer Center, North Carolina (2002– 2006), and US Multicenter studies. Asia included Japan study only. As Asia included Japan study only, there was no possibility to assess heterogeneity between studies in the Asia stratum. f The never/light drinker category included never drinkers and subjects who drink less than 1 drink per day; the moderate drinker category included subjects drinking between 1 (included) and 5 drinks per day; the heavy drinker category included subjects drinking 5 drinks per day or more.