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Age-specific risk factor profiles of adenocarcinomas of the esophagus: a pooled analysis from the international BEACON consortium

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Abstract

Esophageal (EA) and esophagogastric junction (EGJA) adenocarcinoma have been steadily increasing in frequency in younger people, however the etiology of these cancers is poorly understood. We therefore investigated associations of body-mass index (BMI), cigarette smoking, alcohol consumption, gastroesophageal reflux, and use of non-steroidal anti-inflammatory drugs (NSAIDs) in relation to age-specific risks of EA and EGJA. We pooled individual participant data from eight population-based, case-control studies within the international Barrett's and Esophageal Adenocarcinoma Consortium (BEACON). The analysis included 1,363 EA patients, 1,472 EGJA patients, and 5,728 control participants. Multivariable logistic regression was used to estimate odds ratios (ORs) and 95% confidence intervals (CIs) for age-specific (<50, 50–59, 60–69, 70 years) cancer outcomes, as well as interactions by age. BMI, smoking status and pack-years, recurrent gastroesophageal reflux, and frequency of gastroesophageal reflux were positively associated with EA and EGJA in each age group. Early-onset EA (<50 years) had stronger

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associations with recurrent gastroesophageal reflux (OR=8.06, 95%CI: 4.52, 14.37; $P_{\text{effect modification}}=0.01$) and BMI (OR_{BMI 30 vs. <25}=4.19, 95%CI: 2.23, 7.87; $P_{\text{effect modification}}=0.04$), relative to older age groups. In contrast, inverse associations of NSAID use were strongest in the oldest age group (>70 years), although this apparent difference was not statistically significant. Age-specific associations with EGJA showed similar, but slightly weaker patterns and no statistically significant differences by age were observed. Our study provides evidence that associations between obesity and gastroesophageal reflux are stronger among earlier onset EA cancers.

Keywords (MeSH terms)

risk factors; esophageal cancer; case-control studies; obesity; age of onset

Introduction

Incidence of esophageal adenocarcinoma (EA) and esophagogastric junction adenocarcinoma (EGJA) has increased sharply in Western populations^{1, 2} among all age groups^{3–5}. Survival has remained particularly poor, with five-year survival less than 20%⁶. A particularly disconcerting aspect is the growing proportion of advanced-stage tumors that occur at early-onset (<50 years), as opposed to later-onset^{7–9}, with most recent estimates from SEER indicating approximately 8.2% of EAs/EGJAs are diagnosed in US individuals less than 50 years of age. Understanding whether and how risk factor profiles vary by age could provide evidence-based information that may inform clinical practice, as well as provide etiologic insight.

A number of risk factors for EA and EGJA have been identified—irrespective of age at diagnosis. Increased risk is associated with male sex, white race, hiatal hernia^{10–12}, gastroesophageal reflux^{10, 13}, obesity¹⁴, and cigarette smoking^{15, 16}. Inverse associations with usage of nonsteroidal anti-inflammatory drugs (NSAIDs)¹⁷ and moderate alcohol consumption¹⁸ have also been observed. Few studies have conducted age-specific risk factor analyses for EA and EGJA. One study suggested that obesity is more strongly associated with risk of EA among younger patients (<50 years)¹⁹ while another reported that obesity among patients was associated with younger median age at diagnosis²⁰. Given the relative rarity of these malignancies prior studies have been limited in size, which has precluded detailed investigations across multiple age groups. Therefore, we leveraged the large sample size in the international Barrett's and Esophageal Adenocarcinoma Consortium (BEACON) and comprehensively evaluated risk factors by age at diagnosis, particularly focusing on early-onset (<50 years) disease.

Methods

Study Population

The BEACON consortium was formed in 2005 by an international group of investigators^{21–33}. Consortium data consist of population-based case-control and cohort

studies of EA, EGJA and Barrett's esophagus. The large total amount of consortium data enable the conduct of etiological studies of these conditions with excellent statistical power.

For this analysis, we included eight case-control studies that provided data from both EA and EGJA patients: the nationwide Australian Cancer Study (esophageal cancer component)³², Factors Influencing the Barrett's Adenocarcinoma Relationship (FINBAR) Study²¹, nationwide Swedish Esophageal and Cardia Center Study (SECC)²⁸, Larynx/Esophagus/Oral cavity (LEO) Study²⁹, Los Angeles County Multi-ethnic Study³³, Nebraska Health Study II³¹, Population Health Study²², and the United States (US) Multi-Center Study²⁷. Details about source populations, case definitions, recruitment procedures, participants, and study designs have been reported previously²¹⁻³⁴.

The main outcomes of interest were age-specific groups of EA and EGJA patients; particularly early-onset disease (<50 years) given that risk factor profiles for this rarer group are poorly understood. Subjects were limited to those of white non-Hispanic ancestry, because of relatively small numbers of non-white or Hispanic patients (26 black, 89 Hispanic, 42 other race or ethnic groups). The eight studies together provided 2,835 patients (1,363 EA, 1,472 EGJA) and 5,728 controls for analysis. Data acquisition and data pooling were approved by the Institutional Review Board or Research Ethics Committee of each participating institution included in the study.

Study Variables

Included studies provided information on age at diagnosis for case patients, age at interview for controls, sex, education, usual adult body mass index (BMI: weight divided by square of height [kg/m²]), alcohol consumption, cigarette smoking, and study center (for multi-center studies). Studies or their subsets also included information on reported heartburn and regurgitation^{21, 27, 28, 32, 33}, and NSAID usage^{21, 27, 32, 33}. Detailed information on exposure harmonization and detailed analyses of these risk factors has been reported previously for BMI¹⁴, alcohol consumption¹⁸, cigarette smoking^{16, 34}, heartburn/regurgitation¹⁵, and NSAIDs use¹⁷. The categories for age at diagnosis were selected *a priori* based on convenient cut-points (<50, 50-59, 60-69, 70 years) for ease of interpretation.

Statistical Analysis

Primary exposure variables of interest included BMI, alcohol consumption, cigarette smoking, heartburn and regurgitation, and NSAID usage. Because of the small number of early-onset esophageal cancer diagnoses in each study (Table 1), we pooled data for our analysis, rather than using a 2-step meta-analytic approach as in most prior BEACON studies that have focused on a single exposure in relation to cancer diagnosed at any age and include larger case groups, such as BMI¹⁴, alcohol¹⁸, smoking³⁴, or gastroesophageal reflux¹⁵. Multivariable logistic regression models were used to estimate odds ratios (ORs) and their 95% confidence intervals (CIs) for the association between each primary exposure variable and each outcome (EA or EGJA) stratified by age (diagnosis for cases, interview for controls; <50, 50-59, 60-69, 70 years). Covariates included in the multivariable logistic regression models (if not assessed as primary exposure variables) included age (continuous), sex, study, study-center (for multicenter studies), education (< or high

school), BMI (continuous), ever regular consumption of alcohol, and smoking status (never, current, former), as these variables are either of primary interest or known to be associated with these malignancies and thus potential confounders³⁵.

As previously noted, a subset of studies also included information on the following risk factors: heartburn, regurgitation or both as recurrent symptoms (weekly frequency: yes/no) and frequency of heartburn, regurgitation or both (never, <monthly, monthly to <weekly, weekly to <daily, daily) and regular NSAID use (ever/never; including aspirin usage and non-aspirin NSAID usage).

To assess effect modification by age group, we used the likelihood-ratio statistic to compare nested models of the age-adjusted main-effect data (i.e., not stratified by age) with a model that also included main-effect exposure-age category interaction terms (<50, 50–59, 60–69, 70 years; 3df). All analyses were performed using Stata software version 13. All statistical tests were two-sided and P-values less than 0.05 were considered to be statistically significant.

Results

In total, data from 1,363 EA and 1,472 EGJA case patients were available for the analysis (Table 1). Among these subjects, 125 EA and 174 EGJA patients had been diagnosed at <50 years of age (10.5%), 314 EA and 334 EGJA diagnosed at 50–59 years of age (22.9%), 473 EA and 526 EGJA diagnosed at 60–69 years of age (35.2%), and 451 EA and 438 EGJA patients diagnosed at 70 years of age (31.4%). In total, 5,728 population controls—including 790 aged <50 years, 1,286 aged 50–59 years, 1,936 aged 60–69 years, and 1,716 aged 70 years—were available for comparison (Table 1). Study participant characteristics by age at diagnosis/interview are shown in Supplementary Table 1.

The overall and age-specific risk factor profiles of EA and EGJA are provided in Tables 2 and 3. Overall we observed an increased risk of EA and EGJA with increasing BMI, whereas alcohol consumption categories up to 3–5 drinks per day, but not more, were inversely associated, each compared with the 0 drinks per day category. Current or former cigarette smoking status were each positively associated with increased risk of both EA and EGJA, and we observed a trend of increasing risks with increasing cumulative exposure to tobacco smoke (pack-years). Recurrent heartburn and/or recurrent regurgitation, and greater frequency of symptoms were associated with increased risk of both cancers. Lastly, we observed inverse associations with use of NSAIDs and aspirin, and a modest inverse association with non-aspirin NSAID usage with EA. In contrast, associations between NSAID use and EGJA were mostly null. These results are similar to individual exposures investigated in prior BEACON studies^{14, 15, 17, 18, 34}.

We next evaluated age-specific associations of each exposure in relation to EA. Obesity (BMI ≥30) was most strongly associated with early-onset EA (OR=4.19, 95% CI: 2.23, 7.87), with significant differences across age groups ($P_{\text{effect modification}}=0.042$). The magnitude of the association was higher in early-onset EA than in later-onset patients. ORs for the other age categories ranged between 2.6–2.8. When we limited our analysis to studies

including reflux variables and further adjusted models by recurrent reflux, little difference was seen in the association between BMI and EA in the youngest age group (Supplemental Table 3: OR: 2.1 vs OR: 2.3) and age remained an effect modifier ($P_{\text{effect modification}}=0.002$).

Alcohol consumption and number of drinks per day showed little to no associations with risk of EA among age groups <50, 50–59, and 60–69 years. However, for the age group ≥70 years the inverse association for regular drinking was stronger and gained nominal statistical significance (OR 0.62, 95%CI: 0.45, 0.85). Similar findings were found for frequency of alcohol consumption, up to and including the 5–<7 drinks per day category. However, tests for effect modification by age were not statistically significant for either ever regularly consuming alcohol ($P_{\text{effect modification}}=0.17$) or for frequency of alcohol consumption ($P_{\text{effect modification}}=0.33$).

Current and former cigarette smokers were at increased risk of EA in each age group assessed compared with never-smokers. ORs ranged between 1.81 and 3.75 for former and current smokers. Effect modification by age was observed ($P=0.028$), yet no obvious linear pattern across age groups was seen. Associations were strongest for 50–59 year olds and weakest for 60–69 year olds, and these differences persisted when pack-years of cigarette smoking was included in the model, albeit the P value for effect modification was slightly attenuated and not statistically significant ($P=0.073$).

We observed statistically significant positive associations between recurrent gastroesophageal reflux variables (≥ weekly symptoms) and EA risk for all age groups. The strongest associations were observed for early-onset EA, with ORs ranging from 6.62 to 8.06 and age was consistently a significant effect modifier ($P_{\text{effect modification}}=0.02–0.04$). In contrast, associations between EA and frequency of heartburn and/or regurgitation symptoms were similar across each age groups ($P_{\text{effect modification}}=0.16–0.57$).

Regular use of NSAIDs and aspirin were inversely associated with EA for the age group ≥70 years. For all other age groups assessed, weaker inverse associations of these exposures with EA risk were seen, although most of them were not statistically significant and neither were their tests effect modifications by age ($P=0.09–0.12$). No substantial differences for regular use of non-aspirin NSAIDs were seen by age group and were not associated.

Age-specific analyses of EGJA (Table 3) provided generally similar results to those observed for EA (Table 2). Age-specific associations of BMI, alcohol, and gastroesophageal reflux exposures in relation to EGJA were slightly attenuated when compared with those for EA, but were similar in patterns and direction. However, none of these apparent differences were statistically significant. Inverse associations were seen between users of non-aspirin NSAIDs and EGJA in older age groups (60–69 and ≥70) compared with non-users. However, unlike the EA analysis, no associations were observed with regular usage of aspirin.

Discussion

In this large pooled analysis of eight population-based case-control studies we found that recurrent heartburn/regurgitation and obesity were appreciably stronger risk factors for

early-onset EA relative to older age categories. Conversely, low/moderate alcohol consumption and use of aspirin and NSAIDs were inversely associated with late-onset EA, although effect modifications of these associations by age were not significant. Age-specific associations with EGJA showed similar, but slightly weaker patterns and no statistically significant differences by age were observed.

To our knowledge this is the first pooled analysis aimed to evaluate and compare age-specific risk factor profiles for EGJA and one of only a few for EA. In a prior study of age-specific risk factors of EA (cases: n=356), the authors also found that obesity was associated with a younger age at cancer diagnosis²⁰. Our pooled analysis corroborates that finding and provides detailed investigations of the major risk factors across multiple age groups. Although the rise in EA incidence predates the rise in obesity³⁶, it is still possible that obesity is implicated in more recent increases in EA incidence; as evidenced by the strong associations with BE and EA observed in prior BEACON analyses^{37, 38} and other studies³⁹.

The predominant mechanism by which central adiposity is proposed to increase risk of esophageal cancer is by disrupting the integrity of the lower esophageal sphincter; leading to increased propensity for gastroesophageal reflux⁴⁰. Independent of the “mechanical effect” of central adiposity on cancer risk, proinflammatory effects of excess adipose tissue may increase risk of EA via systemic proinflammatory effects^{14, 40, 41}. In this study we also observed some evidence for an effect of obesity on risk of EA that is independent of reflux symptoms.

No prior study, independent of the studies contributing to this pooled analysis, has investigated age-specific associations between recurrent heartburn/regurgitation and risk of EA. Reflux increases secretion of numerous proinflammatory cytokines such as IL-6 and TNF- α and reactive oxygen species^{42–44}. As a recognized hallmark of cancer⁴⁵, inflammation induced by reflux is likely to contribute substantially to the development and progression of esophageal adenocarcinoma⁴⁶. Among a subset of the EA patients included in this study, four SNPs in three apoptosis genes (BCL2, CASP8, and TNFRSF10A) were previously identified as significantly associated with early-onset EA (≤ 55 vs >55 years)⁹. Taken together, one possible explanation for our results is that patients diagnosed with EA at younger ages may have increased likelihood of genetic susceptibility to the deleterious effect of reflux, perhaps having decreased ability to expunge damaged cells via apoptosis. This process could lead to the accumulation of oncogenic mutations in the esophageal cell population, heightening the risk of early-onset EA.

An important strength of our study was the availability of individual participant data from eight population-based, case-control studies, which provided sufficient case numbers required to identify risk factor profiles for early and late-onset disease and to assess effect modification by age. Our exposure variables are believed to have a high degree of reliability, since each was constructed using a unified approach across studies and each exposure in relation to EA and EGJA has been examined and published. Our overall results corroborate the findings of these prior studies^{14, 15, 17, 18, 34} and provide confidence in our statistical approach.

However, several limitations of our analysis should be considered. We have not assessed heritability or genetic factors. These factors could induce differential residual confounding by age at diagnosis, possibly contributing to the effect modification by age effects that we observed^{47–50}. The categories for age at diagnosis were selected *a priori* based on convenient cut-points for ease of interpretation, however, similar trends were observed when age-at-diagnosis categories were based on quartiles of age among controls (data not shown). It is important to note that our analysis includes self-reported symptoms of heartburn and regurgitation, yet reflux exposures can occur without apparent symptoms. Symptoms have been shown to correlate with greater severity of acid reflux exposure⁵¹; nonetheless our results may only apply to people with symptomatic heartburn or regurgitation.

Overall, this study confirms that smoking, obesity, recurrent or frequent heartburn/regurgitation are all associated with increased risk of EA and EGJA across all ages studied. In addition, we found evidence that recurrent heartburn/regurgitation and obesity are more strongly associated with risk of early-onset EA, compared with older age groups. Understanding the mechanisms through which obesity and reflux confer increased risks of esophageal cancer at younger ages might yield important insights for prevention and control of this cancer. Moreover, as the clinical community aims to incorporate more evidence-based health decisions, it is prudent that age-specific risk estimates be considered.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations

EA	esophageal adenocarcinoma
EGJA	esophagogastric junction adenocarcinoma
BMI	body mass index
NSAIDs	non-steroidal anti-inflammatory drugs
OR	odds ratio
CI	confidence interval.

References

1. Drahos J, Wu M, Anderson WF, Trivers KF, King J, Rosenberg PS, Ehemann C, Cook MB. Regional variations in esophageal cancer rates by census region in the United States, 1999–2008. *PloS one*. 2013; 8:e67913. [PubMed: 23861830]
2. Thrift AP, Whiteman DC. The incidence of esophageal adenocarcinoma continues to rise: analysis of period and birth cohort effects on recent trends. *Annals of oncology : official journal of the European Society for Medical Oncology / ESMO*. 2012; 23:3155–3162. [PubMed: 22847812]
3. Brown LM, Devesa SS, Chow WH. Incidence of adenocarcinoma of the esophagus among white Americans by sex, stage, and age. *Journal of the National Cancer Institute*. 2008; 100:1184–1187. [PubMed: 18695138]
4. Cook MB, Chow WH, Devesa SS. Oesophageal cancer incidence in the United States by race, sex, and histologic type, 1977–2005. *Br J Cancer*. 2009; 101:855–859. [PubMed: 19672254]
5. Devesa SS, Blot WJ, Fraumeni JF Jr. Changing patterns in the incidence of esophageal and gastric carcinoma in the United States. *Cancer*. 1998; 83:2049–2053. [PubMed: 9827707]
6. Dubecz A, Gall I, Solymosi N, Schweigert M, Peters JH, Feith M, Stein HJ. Temporal trends in long-term survival and cure rates in esophageal cancer: a SEER database analysis. *Journal of thoracic oncology : official publication of the International Association for the Study of Lung Cancer*. 2012; 7:443–447.
7. Hashemi N, Loren D, DiMarino AJ, Cohen S. Presentation and prognosis of esophageal adenocarcinoma in patients below age 50. *Digestive diseases and sciences*. 2009; 54:1708–1712. [PubMed: 19030991]
8. Portale G, Peters JH, Hsieh CC, Tamhankar AP, Almogy G, Hagen JA, Demeester SR, Bremner CG, Demeester TR. Esophageal adenocarcinoma in patients < or = 50 years old: delayed diagnosis and advanced disease at presentation. *The American surgeon*. 2004; 70:954–958. [PubMed: 15586504]
9. Wu IC, Zhao Y, Zhai R, Liu G, Ter-Minassian M, Asomaning K, Su L, Liu CY, Chen F, Kulke MH, Heist RS, Christiani DC. Association between polymorphisms in cancer-related genes and early onset of esophageal adenocarcinoma. *Neoplasia*. 2011; 13:386–392. [PubMed: 21472143]
10. Chow WH, Finkle WD, McLaughlin JK, Frankl H, Ziel HK, Fraumeni JF Jr. The relation of gastroesophageal reflux disease and its treatment to adenocarcinomas of the esophagus and gastric cardia. *JAMA : the journal of the American Medical Association*. 1995; 274:474–477. [PubMed: 7629956]
11. Wu AH, Tseng CC, Bernstein L. Hiatal hernia, reflux symptoms, body size, and risk of esophageal and gastric adenocarcinoma. *Cancer*. 2003; 98:940–948. [PubMed: 12942560]
12. Pohl H, Wrobel K, Bojarski C, Voderholzer W, Sonnenberg A, Rosch T, Baumgart DC. Risk factors in the development of esophageal adenocarcinoma. *The American journal of gastroenterology*. 2013; 108:200–207. [PubMed: 23247577]
13. Lagergren J, Bergstrom R, Lindgren A, Nyren O. Symptomatic gastroesophageal reflux as a risk factor for esophageal adenocarcinoma. *The New England journal of medicine*. 1999; 340:825–831. [PubMed: 10080844]
14. Hoyo C, Cook MB, Kamangar F, Freedman ND, Whiteman DC, Bernstein L, Brown LM, Risch HA, Ye W, Sharp L, Wu AH, Ward MH, et al. Body mass index in relation to oesophageal and oesophagogastric junction adenocarcinomas: a pooled analysis from the International BEACON Consortium. *International journal of epidemiology*. 2012; 41:1706–1718. [PubMed: 23148106]
15. Cook MB, Corley DA, Murray LJ, Liao LM, Kamangar F, Ye W, Gammon MD, Risch HA, Casson AG, Freedman ND, Chow WH, Wu AH, et al. Gastroesophageal Reflux in Relation to Adenocarcinomas of the Esophagus: A Pooled Analysis from the Barrett's and Esophageal Adenocarcinoma Consortium (BEACON). *PloS one*. 2014; 9:e103508. [PubMed: 25075959]
16. Lubin JH, Cook MB, Pandeya N, Vaughan TL, Abnet CC, Giffen C, Webb PM, Murray LJ, Casson AG, Risch HA, Ye W, Kamangar F, et al. The importance of exposure rate on odds ratios by cigarette smoking and alcohol consumption for esophageal adenocarcinoma and squamous cell carcinoma in the Barrett's Esophagus and Esophageal Adenocarcinoma Consortium. *Cancer epidemiology*. 2012; 36:306–316. [PubMed: 22504051]

17. Liao LM, Vaughan TL, Corley DA, Cook MB, Casson AG, Kamangar F, Abnet CC, Risch HA, Giffen C, Freedman ND, Chow WH, Sadeghi S, et al. Nonsteroidal anti-inflammatory drug use reduces risk of adenocarcinomas of the esophagus and esophagogastric junction in a pooled analysis. *Gastroenterology*. 2012; 142:442–452. e5. quiz e22–3. [PubMed: 22108196]
18. Freedman ND, Murray LJ, Kamangar F, Abnet CC, Cook MB, Nyren O, Ye W, Wu AH, Bernstein L, Brown LM, Ward MH, Pandeya N, et al. Alcohol intake and risk of oesophageal adenocarcinoma: a pooled analysis from the BEACON Consortium. *Gut*. 2011; 60:1029–1037. [PubMed: 21406386]
19. Chow WH, Blot WJ, Vaughan TL, Risch HA, Gammon MD, Stanford JL, Dubrow R, Schoenberg JB, Mayne ST, Farrow DC, Ahsan H, West AB, et al. Body mass index and risk of adenocarcinomas of the esophagus and gastric cardia. *Journal of the National Cancer Institute*. 1998; 90:150–155. [PubMed: 9450576]
20. Chak A, Falk G, Grady WM, Kinnard M, Elston R, Mittal S, King JF, Willis JE, Kondru A, Brock W, Barnholtz-Sloan J. Assessment of familiarity, obesity, and other risk factors for early age of cancer diagnosis in adenocarcinomas of the esophagus and gastroesophageal junction. *The American journal of gastroenterology*. 2009; 104:1913–1921. [PubMed: 19491834]
21. Anderson LA, Watson RG, Murphy SJ, Johnston BT, Comber H, Mc Guigan J, Reynolds JV, Murray LJ. Risk factors for Barrett's oesophagus and oesophageal adenocarcinoma: results from the FINBAR study. *World journal of gastroenterology : WJG*. 2007; 13:1585–1594. [PubMed: 17461453]
22. Brown LM, Silverman DT, Pottern LM, Schoenberg JB, Greenberg RS, Swanson GM, Liff JM, Schwartz AG, Hayes RB, Blot WJ, et al. Adenocarcinoma of the esophagus and esophagogastric junction in white men in the United States: alcohol, tobacco, and socioeconomic factors. *Cancer causes & control : CCC*. 1994; 5:333–340. [PubMed: 8080945]
23. Chen H, Ward MH, Graubard BI, Heineman EF, Markin RM, Potischman NA, Russell RM, Weisenburger DD, Tucker KL. Dietary patterns and adenocarcinoma of the esophagus and distal stomach. *The American journal of clinical nutrition*. 2002; 75:137–144. [PubMed: 11756071]
24. Cheng KK, Sharp L, McKinney PA, Logan RF, Chilvers CE, Cook-Mozaffari P, Ahmed A, Day NE. A case-control study of oesophageal adenocarcinoma in women: a preventable disease. *British journal of cancer*. 2000; 83:127–132. [PubMed: 10883680]
25. Corley DA, Kubo A, Zhao W. Abdominal obesity and the risk of esophageal and gastric cardia carcinomas. *Cancer epidemiology, biomarkers & prevention : a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology*. 2008; 17:352–358.
26. Freedman ND, Abnet CC, Leitzmann MF, Mouw T, Subar AF, Hollenbeck AR, Schatzkin A. A prospective study of tobacco, alcohol, and the risk of esophageal and gastric cancer subtypes. *American journal of epidemiology*. 2007; 165:1424–1433. [PubMed: 17420181]
27. Gammon MD, Schoenberg JB, Ahsan H, Risch HA, Vaughan TL, Chow WH, Rotterdam H, West AB, Dubrow R, Stanford JL, Mayne ST, Farrow DC, et al. Tobacco, alcohol, and socioeconomic status and adenocarcinomas of the esophagus and gastric cardia. *Journal of the National Cancer Institute*. 1997; 89:1277–1284. [PubMed: 9293918]
28. Lagergren J, Bergstrom R, Lindgren A, Nyren O. The role of tobacco, snuff and alcohol use in the aetiology of cancer of the oesophagus and gastric cardia. *International journal of cancer Journal international du cancer*. 2000; 85:340–346. [PubMed: 10652424]
29. Vaughan TL, Davis S, Kristal A, Thomas DB. Obesity, alcohol, and tobacco as risk factors for cancers of the esophagus and gastric cardia: adenocarcinoma versus squamous cell carcinoma. *Cancer epidemiology, biomarkers & prevention : a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology*. 1995; 4:85–92.
30. Veugeliers PJ, Porter GA, Guernsey DL, Casson AG. Obesity and lifestyle risk factors for gastroesophageal reflux disease, Barrett esophagus and esophageal adenocarcinoma. *Diseases of the esophagus : official journal of the International Society for Diseases of the Esophagus / ISDE*. 2006; 19:321–328.
31. Ward MH, Sinha R, Heineman EF, Rothman N, Markin R, Weisenburger DD, Correa P, Zahm SH. Risk of adenocarcinoma of the stomach and esophagus with meat cooking method and doneness

- preference. *International journal of cancer Journal international du cancer*. 1997; 71:14–19. [PubMed: 9096659]
32. Whiteman DC, Sadeghi S, Pandeya N, Smithers BM, Gotley DC, Bain CJ, Webb PM, Green AC. Combined effects of obesity, acid reflux and smoking on the risk of adenocarcinomas of the oesophagus. *Gut*. 2008; 57:173–180. [PubMed: 17932103]
 33. Wu AH, Wan P, Bernstein L. A multiethnic population-based study of smoking, alcohol and body size and risk of adenocarcinomas of the stomach and esophagus (United States). *Cancer causes & control : CCC*. 2001; 12:721–732. [PubMed: 11562112]
 34. Cook MB, Kamangar F, Whiteman DC, Freedman ND, Gammon MD, Bernstein L, Brown LM, Risch HA, Ye W, Sharp L, Pandeya N, Webb PM, et al. Cigarette smoking and adenocarcinomas of the esophagus and esophagogastric junction: a pooled analysis from the international BEACON consortium. *Journal of the National Cancer Institute*. 2010; 102:1344–1353. [PubMed: 20716718]
 35. Reid BJ, Li X, Galipeau PC, Vaughan TL. Barrett's oesophagus and oesophageal adenocarcinoma: time for a new synthesis. *Nature reviews Cancer*. 2010; 10:87–101. [PubMed: 20094044]
 36. Abrams JA, Sharaiha RZ, Gonsalves L, Lightdale CJ, Neugut AI. Dating the rise of esophageal adenocarcinoma: analysis of Connecticut Tumor Registry data, 1940–2007. *Cancer Epidemiol Biomarkers Prev*. 2011; 20:183–186. [PubMed: 21127287]
 37. Kubo A, Cook MB, Shaheen NJ, Vaughan TL, Whiteman DC, Murray L, Corley DA. Sex-specific associations between body mass index, waist circumference and the risk of Barrett's oesophagus: a pooled analysis from the international BEACON consortium. *Gut*. 2013; 62:1684–1691. [PubMed: 23355549]
 38. Hoyo C, Cook MB, Kamangar F, Freedman ND, Whiteman DC, Bernstein L, Brown LM, Risch HA, Ye W, Sharp L, Wu AH, Ward MH, et al. Body mass index in relation to oesophageal and oesophagogastric junction adenocarcinomas: a pooled analysis from the International BEACON Consortium. *Int J Epidemiol*. 2012
 39. Kong CY, Nattinger KJ, Hayeck TJ, Omer ZB, Wang YC, Spechler SJ, McMahon PM, Gazelle GS, Hur C. The impact of obesity on the rise in esophageal adenocarcinoma incidence: estimates from a disease simulation model. *Cancer Epidemiol Biomarkers Prev*. 2011; 20:2450–2456. [PubMed: 21930957]
 40. Ryan AM, Duong M, Healy L, Ryan SA, Parekh N, Reynolds JV, Power DG. Obesity, metabolic syndrome and esophageal adenocarcinoma: epidemiology, etiology and new targets. *Cancer epidemiology*. 2011; 35:309–319. [PubMed: 21470937]
 41. Drahos J, Ricker W, Parsons R, Pfeiffer RM, Warren JL, Cook MB. Metabolic Syndrome Increases Risk of Barrett Esophagus in the Absence of Gastroesophageal Reflux: An Analysis of SEER-Medicare Data. *Journal of clinical gastroenterology*. 2014
 42. Chiu HY, Chen CW, Lin HT, Hsieh CC, Lin SS, Cheng CM. Study of gastric fluid induced cytokine and chemokine expression in airway smooth muscle cells and airway remodeling. *Cytokine*. 2011; 56:726–731. [PubMed: 21996013]
 43. O'Riordan JM, Abdel-latif MM, Ravi N, McNamara D, Byrne PJ, McDonald GS, Keeling PW, Kelleher D, Reynolds JV. Proinflammatory cytokine and nuclear factor kappa-B expression along the inflammation-metaplasia-dysplasia-adenocarcinoma sequence in the esophagus. *The American journal of gastroenterology*. 2005; 100:1257–1264. [PubMed: 15929754]
 44. Souza RF, Huo X, Mittal V, Schuler CM, Carmack SW, Zhang HY, Zhang X, Yu C, Hormi-Carver K, Genta RM, Spechler SJ. Gastroesophageal reflux might cause esophagitis through a cytokine-mediated mechanism rather than caustic acid injury. *Gastroenterology*. 2009; 137:1776–1784. [PubMed: 19660463]
 45. Hanahan D, Weinberg RA. Hallmarks of cancer: the next generation. *Cell*. 2011; 144:646–674. [PubMed: 21376230]
 46. Kavanagh ME, O'Sullivan KE, O'Hanlon C, O'Sullivan JN, Lysaght J, Reynolds JV. The esophagitis to adenocarcinoma sequence; the role of inflammation. *Cancer letters*. 2014; 345:182–189. [PubMed: 23994342]
 47. Dhillon PK, Farrow DC, Vaughan TL, Chow WH, Risch HA, Gammon MD, Mayne ST, Stanford JL, Schoenberg JB, Ahsan H, Dubrow R, West AB, et al. Family history of cancer and risk of

- esophageal and gastric cancers in the United States. *International journal of cancer Journal international du cancer*. 2001; 93:148–152. [PubMed: 11391635]
48. Ji J, Hemminki K. Familial risk for esophageal cancer: an updated epidemiologic study from Sweden. *Clinical gastroenterology and hepatology : the official clinical practice journal of the American Gastroenterological Association*. 2006; 4:840–845. [PubMed: 16678496]
49. Jiang X, Tseng CC, Bernstein L, Wu AH. Family history of cancer and gastroesophageal disorders and risk of esophageal and gastric adenocarcinomas: a case-control study. *BMC cancer*. 2014; 14:60. [PubMed: 24495377]
50. Zhang ZF, Kurtz RC, Sun M, Karpeh M Jr, Yu GP, Gargon N, Fein JS, Georgopoulos SK, Harlap S. Adenocarcinomas of the esophagus and gastric cardia: medical conditions, tobacco, alcohol, and socioeconomic factors. *Cancer epidemiology, biomarkers & prevention : a publication of the American Association for Cancer Research, cosponsored by the American Society of Preventive Oncology*. 1996; 5:761–768.
51. Bredenoord AJ, Weusten BL, Curvers WL, Timmer R, Smout AJ. Determinants of perception of heartburn and regurgitation. *Gut*. 2006; 55:313–318. [PubMed: 16120760]

Novelty & Impact

Incidence in esophageal adenocarcinoma has increased sharply among all age groups; however one disconcerting trend is the increasing proportion of advanced-stage tumors occurring at younger (< 50 years) as opposed to older ages. The relative rarity of these malignancies has precluded prior studies from assessing risk factors across age groups. Pooling data from 8 case-control studies we found that recurrent heartburn/regurgitation and obesity were appreciably stronger risk factors for early-onset EA relative to older age-categories.

Table 1

Characteristics of BEACON studies included in the analysis

Name	Country	Recruitment Period	Control Population						Esophageal adenocarcinoma Cases						Esophagogastric junction adenocarcinoma Cases								
			Age						Age						Age								
			Overall	≤50	50-59	60-69	70	Overall	≤50	50-59	60-69	70	Overall	≤50	50-59	60-69	70	Overall	≤50	50-59	60-69	70	
Australian Cancer Study ³²	Australia	2001-2005	1444	249	376	494	325	342	29	97	119	97	401	44	108	135	114						
Factors Influencing the Barrett's Adenocarcinoma Relationship Study ²²	Ireland	2002-2004	252	45	48	74	85	126	20	25	38	43	87	9	21	33	24						
Swedish Esophageal Cancer Study ²⁹	Sweden	1995-1997	816	48	160	245	363	189	7	31	61	90	262	26	49	84	103						
Larynx/Esophagus/Oral Cavity Study ³⁰	United States	1983-19990	687	98	174	303	112	119	8	29	62	20	140	24	29	63	24						
Los Angeles Country Multi-ethnic Study ³³	United States	1992-1997	810	84	163	309	254	165	16	43	72	34	200	20	52	79	49						
Nebraska Health Study II ³⁴	United States	1988-1993	421	85	43	78	215	96	9	19	24	44	34	5	3	7	19						
Population Health Study ²³	United States	1986-1989	702	116	198	211	177	58	5	19	20	14	113	13	28	45	27						
US Multi-Center Study ²⁸	United States	1993-1995	596	65	124	222	185	268	31	51	77	109	235	33	44	80	78						
Total			5728	790	1286	1936	1716	1363	125	314	473	451	1472	174	334	526	438						

Number of cases and controls listed are those with complete data (sex, age, BMI, smoking, education) after restricting to white non-Hispanic participants

Age at interview for controls and age at diagnosis for EA and EGJA cases

Table 2

Associations between risk factors and esophageal adenocarcinoma (EA) for all age groups (overall) and stratified by age

	Age					Age effect modification p-value
	Overall (all ages) OR (95%CI)	<50 yrs OR (95%CI)	50-59 yrs OR (95%CI)	60-69 yrs OR (95%CI)	70 yrs OR (95%CI)	
Body Mass Index (BMI)						
Normal (<25)	referent	referent	referent	referent	referent	0.042
Overweight (25-29)	1.58 (1.37, 1.82)	1.64 (0.99, 2.73)	1.61 (1.18, 2.20)	1.93 (1.51, 2.46)	1.19 (0.93, 1.52)	
Obese (30)	2.83 (2.36, 3.40)	4.19 (2.23, 7.87)	2.64 (1.81, 3.84)	2.73 (1.99, 3.74)	2.81 (2.00, 3.94)	
Alcohol Consumption						
Alcohol ever	0.78 (0.64, 0.96)	0.95 (0.42, 2.13)	0.73 (0.45, 1.17)	1.00 (0.70, 1.42)	0.62 (0.45, 0.85)	0.166
Alcoholic drinks per day						
None	referent	referent	referent	referent	referent	0.335
>0-<0.5	0.78 (0.62, 0.98)	1.05 (0.43, 2.58)	0.72 (0.42, 1.24)	0.99 (0.66, 1.48)	0.63 (0.43, 0.92)	
0.5-<1.0	0.55 (0.42, 0.72)	0.44 (0.14, 1.36)	0.47 (0.25, 0.88)	0.72 (0.45, 1.13)	0.53 (0.34, 0.82)	
1-<3	0.74 (0.58, 0.93)	0.85 (0.35, 2.07)	0.71 (0.42, 1.20)	0.84 (0.56, 1.26)	0.66 (0.45, 0.96)	
3-<5	0.69 (0.52, 0.91)	0.93 (0.34, 2.53)	0.51 (0.27, 0.94)	0.90 (0.56, 1.45)	0.55 (0.33, 0.89)	
7	0.93 (0.66, 1.30)	1.55 (0.47, 5.10)	0.85 (0.41, 1.74)	1.34 (0.76, 2.36)	0.48 (0.24, 0.97)	
Cigarette Smoking						
Smoking Status						
Never smoker	referent	referent	referent	referent	referent	0.028
Former smoker	2.36 (1.94, 2.88)	2.44 (0.88, 6.74)	2.95 (1.89, 4.62)	1.86 (1.33, 2.60)	2.51 (1.80, 3.49)	
Current smoker	2.65 (2.17, 3.24)	2.72 (1.11, 6.65)	3.75 (2.46, 5.69)	1.81 (1.30, 2.54)	2.80 (1.93, 4.07)	
Pack years						
Never	referent	referent	referent	referent	referent	0.073
<14 yrs	1.27 (1.05, 1.55)	2.02 (1.10, 3.68)	1.47 (0.96, 2.25)	0.98 (0.69, 1.38)	1.39 (0.99, 1.96)	
14->30 yrs	2.01 (1.66, 2.45)	3.62 (2.04, 6.42)	2.13 (1.41, 3.23)	1.79 (1.28, 2.50)	1.66 (1.14, 2.42)	
30->45 yrs	2.00 (1.63, 2.45)	2.88 (1.35, 6.14)	2.35 (1.54, 3.59)	1.42 (1.00, 2.01)	2.11 (1.45, 3.07)	
45 yrs	2.50 (2.09, 3.00)	2.68 (1.10, 6.53)	3.41 (2.29, 5.07)	1.84 (1.36, 2.48)	2.61 (1.92, 3.54)	

Reflux#	Age					Age effect modification p-value
	Overall (all ages) OR (95%CI)	<50 yrs OR (95%CI)	50-59 yrs OR (95%CI)	60-69 yrs OR (95%CI)	70 yrs OR (95%CI)	
Recurrent heartburn	4.52 (3.84, 5.33)	6.62 (3.66, 11.97)	3.17 (2.19, 4.60)	5.03 (3.82, 6.64)	5.21 (3.88, 6.99)	0.043
Recurrent regurgitation	4.45 (3.74, 5.31)	7.98 (4.26, 14.96)	3.31 (2.20, 4.98)	4.38 (3.27, 5.85)	5.24 (3.81, 7.19)	0.033
Recurrent heartburn or regurgitation	4.70 (4.03, 5.49)	8.06 (4.52, 14.37)	3.15 (2.22, 4.45)	5.33 (4.08, 6.95)	5.20 (3.95, 6.85)	0.011
Heartburn frequency						
Never	referent	referent	referent	referent	referent	0.348
< Monthly	0.88 (0.67, 1.16)	0.55 (0.21, 1.49)	0.65 (0.39, 1.11)	1.08 (0.68, 1.71)	0.86 (0.51, 1.44)	
Monthly to < weekly	2.74 (2.10, 3.57)	1.87 (0.61, 5.75)	1.93 (1.14, 3.27)	3.69 (2.35, 5.80)	2.63 (1.63, 4.26)	
Weekly to < daily	4.15 (3.35, 5.15)	5.21 (2.45, 11.06)	2.52 (1.51, 4.18)	5.27 (3.64, 7.61)	4.25 (2.93, 6.15)	
Daily	7.04 (5.47, 9.08)	6.84 (2.92, 16.02)	4.65 (2.72, 7.94)	7.75 (5.02, 11.97)	11.41 (7.04, 18.48)	
Regurgitation frequency						
Never	referent	referent	referent	referent	referent	0.573
< Monthly	0.70 (0.55, 0.91)	0.52 (0.21, 1.26)	0.56 (0.34, 0.91)	0.73 (0.47, 1.14)	0.75 (0.46, 1.22)	
Monthly to < weekly	2.82 (2.21, 3.59)	2.36 (0.88, 6.32)	2.21 (1.31, 3.74)	2.59 (1.70, 3.92)	3.69 (2.43, 5.62)	
Weekly to < daily	5.03 (4.02, 6.31)	7.03 (3.22, 15.33)	3.27 (1.95, 5.49)	5.65 (3.87, 8.23)	5.65 (3.77, 8.46)	
Daily	4.83 (3.64, 6.42)	7.00 (2.53, 19.40)	3.25 (1.65, 6.40)	4.33 (2.70, 6.94)	6.58 (3.99, 10.86)	
Heartburn or regurgitation frequency						
Never	referent	referent	referent	referent	referent	0.158
Monthly to < weekly	1.28 (1.04, 1.59)	1.23 (0.57, 2.65)	0.79 (0.51, 1.21)	1.63 (1.11, 2.39)	1.42 (0.97, 2.07)	
Weekly to < daily	4.40 (3.57, 5.44)	7.05 (3.23, 15.41)	2.25 (1.39, 3.64)	5.95 (4.09, 8.67)	4.56 (3.20, 6.51)	
Daily	6.86 (5.41, 8.70)	9.59 (3.97, 23.18)	4.01 (2.40, 6.73)	8.16 (5.39, 12.34)	9.55 (6.24, 14.62)	
NSAID Usage#						
NSAIDS	0.68 (0.57, 0.81)	0.75 (0.39, 1.44)	0.88 (0.60, 1.29)	0.72 (0.54, 0.96)	0.47 (0.34, 0.65)	0.090
Non-aspirin NSAIDS	0.82 (0.67, 0.99)	0.58 (0.29, 1.13)	0.99 (0.67, 1.46)	0.74 (0.53, 1.03)	0.88 (0.61, 1.25)	0.128
Aspirin	0.70 (0.59, 0.83)	0.81 (0.43, 1.54)	0.82 (0.57, 1.17)	0.76 (0.57, 1.00)	0.50 (0.37, 0.69)	0.116

Subset of case-control studies with exposure variable
 Adjusted for age, sex, BMI (<25, 25-29, 30), alcohol consumption (ever/never), education, smoking (never/former/current), and study-specific variables.

Abbreviations: OR, odds ratio; CI, confidence interval; yrs, years

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Table 3

Associations between risk factors and esophagogastric junction adenocarcinoma (EGJA) for all age groups (overall) and stratified by age

	Age					Age effect modification p-value
	Overall (all ages) OR (95%CI)	<50 yrs OR (95%CI)	50-59 yrs OR (95%CI)	60-69 yrs OR (95%CI)	70 yrs OR (95%CI)	
Body Mass Index (BMI)						
Normal (<25)	referent	referent	referent	referent	referent	0.472
Overweight (25-29)	1.33 (1.16, 1.52)	1.49 (0.97, 2.29)	1.56 (1.16, 2.11)	1.46 (1.16, 1.84)	1.03 (0.81, 1.32)	
Obese (30)	2.32 (1.94, 2.76)	3.33 (1.97, 5.62)	2.74 (1.91, 3.95)	2.14 (1.58, 2.89)	2.07 (1.47, 2.91)	
Alcohol Consumption						
Alcohol ever	0.73 (0.60, 0.88)	0.75 (0.41, 1.35)	0.65 (0.42, 0.99)	0.81 (0.57, 1.15)	0.76 (0.55, 1.07)	0.833
Alcoholic drinks per day						
None	referent	referent	referent	referent	referent	0.107
>0-<0.5	0.75 (0.60, 0.94)	1.06 (0.55, 2.06)	0.71 (0.43, 1.16)	0.66 (0.44, 0.99)	0.88 (0.60, 1.30)	
0.5-<1.0	0.68 (0.53, 0.87)	0.64 (0.30, 1.37)	0.50 (0.29, 0.88)	0.77 (0.50, 1.19)	0.85 (0.55, 1.30)	
1-<3	0.65 (0.52, 0.81)	0.57 (0.29, 1.14)	0.58 (0.36, 0.94)	0.83 (0.56, 1.21)	0.62 (0.41, 0.92)	
3-<5	0.76 (0.58, 0.98)	0.91 (0.42, 1.97)	0.56 (0.32, 0.98)	1.07 (0.69, 1.66)	0.56 (0.34, 0.92)	
7	0.86 (0.61, 1.19)	0.20 (0.04, 1.07)	0.93 (0.48, 1.83)	1.16 (0.67, 2.00)	0.74 (0.39, 1.41)	
Cigarette Smoking						
Smoking Status						
Never smoker	referent	referent	referent	referent	referent	0.730
Former smoker	2.48 (2.03, 3.02)	3.24 (1.41, 7.48)	2.64 (1.68, 4.14)	2.66 (1.89, 3.73)	2.31 (1.65, 3.23)	
Current smoker	3.24 (2.68, 3.92)	3.79 (1.85, 7.80)	3.81 (2.54, 5.73)	3.78 (2.74, 5.21)	2.30 (1.57, 3.36)	
Pack years						
Never	referent	referent	referent	referent	referent	0.615
<14 yrs	1.28 (1.06, 1.55)	1.04 (0.62, 1.72)	1.65 (1.09, 2.49)	1.26 (0.88, 1.79)	1.36 (0.97, 1.92)	
14->30 yrs	2.28 (1.89, 2.75)	2.12 (1.29, 3.48)	2.21 (1.46, 3.34)	2.65 (1.90, 3.69)	2.15 (1.52, 3.05)	
30->45 yrs	2.59 (2.14, 3.14)	3.18 (1.71, 5.94)	3.45 (2.33, 5.11)	2.51 (1.80, 3.52)	2.15 (1.49, 3.10)	
45 yrs	2.88 (2.41, 3.44)	3.71 (1.81, 7.60)	3.36 (2.27, 4.98)	3.25 (2.40, 4.40)	2.30 (1.69, 3.15)	

	Age						Age effect modification p-value
	Overall (all ages) OR (95%CI)	<50 yrs OR (95%CI)	50-59 yrs OR (95%CI)	60-69 yrs OR (95%CI)	70 yrs OR (95%CI)		
Reflux[#]							
Recurrent heartburn	2.42 (2.04, 2.87)	3.41 (1.99, 5.83)	2.36 (1.63, 3.44)	2.11 (1.58, 2.83)	2.63 (1.94, 3.57)	0.598	
Recurrent regurgitation	2.25 (1.86, 2.71)	2.99 (1.63, 5.47)	2.16 (1.41, 3.31)	2.16 (1.58, 2.94)	2.23 (1.58, 3.14)	0.627	
Recurrent heartburn or regurgitation	2.46 (2.10, 2.87)	3.35 (2.02, 5.55)	2.33 (1.65, 3.29)	2.49 (1.92, 3.24)	2.33 (1.76, 3.09)	0.590	
Heartburn frequency							
Never	referent	referent	referent	referent	referent	0.081	
< Monthly	0.53 (0.41, 0.68)	0.63 (0.30, 1.31)	0.49 (0.29, 0.83)	0.49 (0.31, 0.77)	0.51 (0.31, 0.83)		
Monthly to < weekly	1.59 (1.23, 2.06)	1.19 (0.46, 3.11)	1.70 (1.02, 2.85)	1.76 (1.14, 2.71)	1.30 (0.80, 2.13)		
Weekly to < daily	2.24 (1.81, 2.78)	3.17 (1.60, 6.30)	2.49 (1.56, 3.98)	2.11 (1.45, 3.05)	1.98 (1.35, 2.91)		
Daily	2.75 (2.09, 3.62)	2.54 (1.10, 5.86)	2.25 (1.23, 4.10)	2.04 (1.26, 3.30)	4.53 (2.76, 7.44)		
Regurgitation frequency							
Never	referent	referent	referent	referent	referent	0.603	
< Monthly	0.57 (0.46, 0.72)	0.52 (0.26, 1.01)	0.34 (0.21, 0.56)	0.71 (0.48, 1.04)	0.66 (0.43, 1.01)		
Monthly to < weekly	1.63 (1.28, 2.08)	1.32 (0.57, 3.08)	1.31 (0.78, 2.21)	1.89 (1.27, 2.81)	1.69 (1.08, 2.64)		
Weekly to < daily	1.99 (1.56, 2.53)	1.35 (0.58, 3.19)	1.60 (0.93, 2.75)	2.26 (1.51, 3.38)	2.16 (1.40, 3.32)		
Daily	2.44 (1.82, 3.27)	5.22 (2.12, 12.86)	2.25 (1.13, 4.47)	2.13 (1.30, 3.50)	2.22 (1.28, 3.84)		
Heartburn or regurgitation frequency							
Never	referent	referent	referent	referent	referent	0.833	
Monthly to < weekly	0.81 (0.67, 0.98)	0.62 (0.34, 1.15)	0.73 (0.49, 1.09)	0.94 (0.68, 1.31)	0.77 (0.54, 1.10)		
Weekly to < daily	2.12 (1.73, 2.60)	1.88 (0.93, 3.80)	2.21 (1.42, 3.43)	2.44 (1.72, 3.46)	1.84 (1.28, 2.65)		
Daily	2.72 (2.13, 3.47)	4.09 (1.91, 8.75)	2.23 (1.27, 3.90)	2.50 (1.64, 3.80)	3.03 (1.95, 4.71)		
NSAID Usage[#]							
NSAIDS	0.89 (0.75, 1.05)	0.64 (0.37, 1.12)	1.06 (0.74, 1.53)	1.02 (0.77, 1.35)	0.72 (0.52, 0.98)	0.313	
Non-aspirin NSAIDS	0.76 (0.63, 0.91)	0.81 (0.46, 1.44)	1.03 (0.71, 1.50)	0.67 (0.49, 0.93)	0.69 (0.48, 0.98)	0.376	
Aspirin	0.91 (0.77, 1.07)	0.55 (0.31, 0.97)	0.92 (0.65, 1.29)	1.05 (0.80, 1.37)	0.82 (0.61, 1.11)	0.288	

[#] Subset of case-control studies with exposure variable Adjusted for age, sex, BMI (<25, 25-29, 30), alcohol consumption (ever/never), education, smoking (never/former/current), and study-specific variables.

Abbreviations: OR, odds ratio; CI, confidence interval; yrs, years

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