

Gastric atrophy and oesophageal squamous cell carcinoma: possible interaction with dental health and oral hygiene habit

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BACKGROUND: Gastric fundal atrophy has been hypothesised to increase the risk of oesophageal squamous cell carcinoma (OSCC), but studies have shown inconsistent results.

METHODS: We measured serum pepsinogen I (PGI) and pepsinogen II (PGII) among 293 incident cases and 524 matched neighbourhood controls in a high-risk area of Northern Iran. Conditional logistic regression model was used to estimate odds ratios (ORs) and their 95% confidence intervals (CIs).

RESULTS: After controlling for age, sex, residence area and other potential confounders, gastric atrophy (defined by a validated criterion, PGI < 55 $\mu\text{g dl}^{-1}$) was associated with a two-fold increased risk (OR = 2.01, 95% CI: 1.18, 3.45) of OSCC in the absence of nonatrophic pangastritis (defined as PGII < 11.8 $\mu\text{g dl}^{-1}$). Stratification by PGII decreased the misclassification errors due to cancer-induced gastritis. Presence of both poor dental health, indicated by higher than median sum of decayed, missing, and filled teeth (DMFT score), and gastric atrophy further increased the risk of OSCC (OR = 4.15, 95% CI: 2.04, 8.42) with relative excess risk due to interaction (RERI) of 1.47 (95% CI: -1.15, 4.1). Coexistence of poor oral hygiene habit with gastric atrophy elevated OSCC risk eight times (OR = 8.65, 95% CI: 3.65, 20.46) and the additive interaction index was marginally statistically significant (RERI = 4.34, 95% CI: -1.07, 9.76).

CONCLUSION: Gastric atrophy is a risk factor for OSCC, and poor dental health and oral hygiene habit may act synergistically in increasing the risk.

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Oesophageal cancer is the eighth most common cause of cancer death worldwide (Jemal *et al*, 2011). Gastric atrophy as a consequence of chronic *Helicobacter pylori* infection was linked to an increased risk of oesophageal squamous cell carcinoma (OSCC) in a Swedish population-based case-control study (Ye *et al*, 2004). In the atrophic stomach, hypochlorhydria allows bacterial overgrowth (Viani *et al*, 2000) and may produce potential carcinogenic substances, such as nitrosamines, which are also related to alcohol drinking and tobacco smoking (Millonig *et al*, 2011). Histology is the gold standard for diagnosing gastric atrophy. However, because of its inconvenience in large epidemiological studies, serum level of pepsinogens has been applied as a replacement method. Pepsinogen I (PGI) is produced in the fundic glands and decreases proportionally with progression of fundic atrophy. Pepsinogen II (PGII) is synthesised in most

parts of the gastric mucosa and part of the duodenum and some studies have shown its importance in detecting inflammation (He *et al*, 2011) or gastric cancer screening (Abnet *et al*, 2011). Magnitude of the reported associations between serological gastric atrophy and OSCC varied from null to eightfold (Iijima *et al*, 2007; de Vries *et al*, 2009; Ren *et al*, 2009; Cook *et al*, 2010; Venerito *et al*, 2011).

Poor oral and dental hygiene are also linked to nitrosamine (Meurman and Uittamo, 2008) and acetaldehyde production (National Toxicology Program, 2010). In addition, poor oral hygiene and dental health have been shown to be associated with an increased risk of OSCC in high-risk areas (Abnet *et al*, 2001, 2008). Previous studies have not tested possible synergism between these two risk factors. Golestan province in Iran is located in the Asian Oesophageal Cancer Belt and has some of the highest incidence rates for OSCC (Mahboubi *et al*, 1973; Saidi *et al*, 2000). Unlike low-risk areas, tobacco and alcohol use are not major risk factors for OSCC in this belt (Nasrollahzadeh *et al*, 2008), from which 90% of the OSCC cases arise worldwide. We evaluated the association between gastric atrophy (measured by serum pepsinogen level) and OSCC in a population-based case-control

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study in Northern Iran and tested joint effect on OSCC risk in the presence of both gastric atrophy and poor oral hygiene.

MATERIALS AND METHODS

Case and control selection

Details of subject selection have been reported earlier (Nasrollahzadeh *et al*, 2008). Briefly, case subjects were recruited at Atrak clinic, the only specialised clinic for oesophageal cancer diagnosis and treatment in eastern Golestan, from December 2003 to June 2007. Included cases were those of histopathologically confirmed OSCC patients who underwent oesophago-gastro-duodenal videoscopy at Atrak clinic and agreed to participate in the study. The reports of the local cancer registry demonstrated that ~70% of all incident cases during the study period were referred to Atrak clinic (unpublished data). Attempts were made to select two population-based control subjects, individually matched to the cases by age (± 2 years), sex, and from the same neighbourhood or village using annually updated family health census. List of all potentially eligible controls for rural cases was ordered randomly in each village and for urban cases was ordered by geographic proximity to the case's residence. In all, 77% of the enrolled controls were the first selected neighbours, and 11% and 3%, respectively, were the second and third choices from the lists. Absence of the eligible control at the time of invitation was almost always the reason for not participating in the study. In total, we recruited 300 cases and 571 matched controls.

To investigate whether OSCC cases have higher risk of developing non-atrophic gastritis in histology, we used another set of controls. Controls were consisted of endoscopy room patients who were referred to Atrak clinic with a suspicion of upper gastrointestinal malignancies during same period of case enrolment and were matched for age and sex. Three biopsy tissues from lesser-curvature antrum, lesser-curvature body and cardia were available for 261 cases (88%) and 274 controls.

Interview and examination

After obtaining written informed consent from each subject, a nurse and a physician administered a structured questionnaire. A nutritionist administered a validated food frequency questionnaire (Malekshah *et al*, 2006). All interviews were conducted face to face at Atrak clinic for case subjects and at health centres for controls except those chosen from Gonbad city who were interviewed at Golestan Cohort Centre. No proxies were used.

The trained physicians examined each patient's oral cavity and teeth. Number of decayed, missing, and filled teeth (DMFT score) in addition to frequency of tooth brushing was recorded. Detailed information has been published elsewhere (Abnet *et al*, 2008). Reliability of measuring tooth count using κ statistic was 0.86 (Pourshams *et al*, 2005). Data was collected on demographic variables, ethnicity, alcohol consumption, lifelong history of tobacco or opium use, medication, and several potential confounders.

Serological data

Prior to endoscopy, an experienced nurse collected 12 ml venous blood sample from each case subject. A trained technician immediately separated serum and stored it at -80°C . Collected blood samples from matched controls were transferred on ice in a cooler box ($\sim 4^{\circ}\text{C}$). Time between collection and processing of neighbourhood control samples was <12 h. For those controls who were interviewed at the Golestan Cohort Centre, sample processing and storage were performed immediately. Samples were transported on dry ice to Karolinska Institutet, Sweden, and stored at -20°C prior to analysis. Serum pepsinogen I and II were measured blinded to case-control status using enzyme-linked

immunosorbent assays (Biohit, Helsinki, Finland). Aliquots from a pool of serum samples of healthy individuals were distributed among nine assay plates (two samples per plate). The coefficients of variation were 8% and 14% for PGI and PGII, respectively. *H. pylori* serology was evaluated qualitatively with western blot assay (Helico Blot 2.1; MP Biomedicals Asia Pacific Ltd, Singapore, Singapore). *H. pylori* infection was considered positive if (1) both 19.5- and 30-kDa bands were present or (2) any of the 35-, 37-, or 89-kDa band was present. CagA was positive if 116-kDa band was present. We have carried out a validation study in the same population with similar ethnic structure among 309 endoscopy clinic patients and compared pepsinogen serology with histology using modified Sydney classification as the gold standard. We reported $\text{PGI} < 55 \mu\text{g l}^{-1}$ (sensitivity: 61.9%, specificity: 94.8%) as the optimal cutoff point for serological diagnosis of gastric fundal atrophy in the study population. At cutoff concentration of $11.8 \mu\text{g l}^{-1}$, PGII demonstrated 84.2% sensitivity and 45.4% specificity to distinguish nonatrophic pangastritis (Nasrollahzadeh *et al*, 2011).

Statistical methods

Serum PGI and PGII were analysed as dichotomous variables using validated cutoff points. There was no *a priori* DMFT cutoff value, and cubic spline curve did not highlight any specific pattern, therefore the median DMFT was considered as a cutoff value for dichotomising it. Conditional logistic regression models were used to calculate odds ratios (ORs) and 95% confidence intervals (CIs). In design, case and control subjects were individually matched for age, sex, and place of residence. ORs were further adjusted for education, ethnicity (Turkmen *vs* non-Turkmen), tobacco and opium use, alcohol drinking, and fruit/vegetable consumption. Interaction on additive scale between dichotomous variables of gastric atrophy and poor dental health or oral hygiene habit was evaluated and the relative excess risk due to interaction (RERI), attributable proportion and synergy index were calculated. Delta method was applied to calculate 95% CI for measures of interaction. To address the effect of bias due to measurement error in serology assay, the ORs adjusted for misclassification as a function of different sensitivity and specificity for cases and controls were calculated using external validation data. Two-sided *P*-values ≤ 0.05 were considered as statistically significant. All analyses were done using Stata version 11.1 (StataCorp., College Station, TX, USA).

Ethical approval

This study was approved by the ethical committee of the Digestive Disease Research Centre of Tehran University of Medical Sciences, Iran, the Institutional Review Board of National Cancer Institute, USA, and the Stockholm Regional Ethics Vetting Board, Sweden.

RESULTS

The study included 300 incident case patients and 571 matched control subjects from whom serum samples were available for 293 (98%) cases and 524 (92%) controls. All cases had at least one matched control. Demographic characteristics of the study participants and potential confounders are summarised in Table 1. No substantial differences were observed between those who provided serum and those whose serum samples were not available.

Table 2 demonstrates proportion of subjects with serological gastric atrophy and ORs for association of PGI and PGII with OSCC risk after further controlling for education level and ethnicity (as socioeconomic index), opium and tobacco use, fruit, vegetable, and alcohol consumption. A $\text{PGI} < 55 \mu\text{g l}^{-1}$ was

Table 1 Characteristics of OSCC cases and matched controls, Golestan province, Iran, 2003–2007

Characteristics	Cases (n = 293)	Controls (n = 524)	P-value	Controls without serum samples (n = 47)	P-value ^a
Mean age (s.d.), years	64.4 (11.1)	65.5 (10.4)	0.95	62.1 (9.9)	0.07
Sex (%)					
Men	147 (50.2)	256 (48.8)	0.71	22 (46.8)	0.8
Women	146 (49.8)	268 (51.2)		25 (53.2)	
Residence area (%)					
Urban	80 (27.3)	133 (25.4)	0.54	17 (36.2)	0.1
Rural	213 (72.7)	391 (74.6)		30 (63.8)	
Ethnicity (%)					
Turkmen	166 (56.7)	282 (53.8)	0.43	30 (63.8)	0.18
Non-Turkmen	127 (43.3)	242 (46.2)		17 (36.2)	
Education (%)					
Illiterate	261 (89.1)	437 (83.4)	0.02	37 (87.7)	0.41
Primary school and higher	32 (10.9)	87 (16.6)		10 (21.3)	
Alcohol ever use (%)					
No	287 (98)	511 (97.5)	0.69	46 (97.9)	0.88
Yes	6 (2)	13 (2.5)		1 (2.1)	
Tobacco and opium use ^b (%)					
Neither	162 (55.5)	366 (70.0)	0.0002	32 (68.1)	0.58
Ever use tobacco	43 (14.7)	59 (11.3)		7 (14.9)	
Ever use opium	29 (10.0)	33 (6.3)		1 (2.1)	
Ever use tobacco and opium	58 (19.8)	65 (12.4)		7 (14.9)	
Fruit and vegetable consumption (%)					
≤ Median ^c	139 (47.4)	235 (44.8)	0.47	20 (42.5)	0.76
> Median	154 (52.6)	289 (55.1)		27 (57.5)	
DMFT (%) ^b					
≤ Median (28)	117 (40.3)	266 (51.0)	0.004	28 (59.6)	0.25
> Median	173 (59.7)	256 (49.0)		19 (40.4)	
Tooth brushing habit ^b					
Ever	59 (20.3)	221 (42.4)	<0.0001	18 (39.1)	0.66
Never	232 (79.7)	300 (57.6)		28 (60.9)	
PGI ($\mu\text{g l}^{-1}$), median (IQR)	109.3 (67.2–147.5)	94.9 (66.6–123.3)		NA	
PGII ($\mu\text{g l}^{-1}$), median (IQR)	14.2 (9.6–21.5)	12.3 (8.8–18.2)		NA	
PGI/PGII ratio, median (IQR)	7.0 (4.8–9.7)	7.2 (5.3–9.9)		NA	

Abbreviations: DMFT = Sum of decayed, missed, and filled teeth; IQR = interquartile range; NA = not applicable; OSCC = oesophageal squamous cell carcinoma. ^aP-value for significance of difference between controls with and without serum samples. ^bNumbers of cases and controls were less than total because of missed data. ^cMedian fruit and vegetable consumption = 521.8 gr/day.

associated with a 1.39-fold increased risk of OSCC (95% CI: 0.93, 2.09). Histological diagnosis of non-atrophic gastritis was found in 30% of OSCC cases with 2-fold risk comparing clinic controls (OR = 1.97, 95% CI: 1.13, 3.45) (Supplementary Table). Because gastritis might be caused secondary to OSCC, we evaluated the association of gastric fundal atrophy with OSCC in combination with severe nonatrophic pangastritis. Combination of PGI < 55 $\mu\text{g l}^{-1}$ and PGII < 11.8 $\mu\text{g l}^{-1}$ was associated with a two-fold increased risk for OSCC. Further adjustment for tea-drinking habit as a suggested strong risk factor in this population (Islami et al, 2009) did not alter the observed estimates. *H. pylori* serology, measured as antibodies against either whole *H. pylori* or CagA, was not associated with OSCC risk.

We applied other published thresholds for atrophy, including PGI < 30 $\mu\text{g l}^{-1}$, PGI/PGII ratio < 3 and PGI < 70 $\mu\text{g l}^{-1}$, PGI/PGII ratio < 3. The measures of association did not change materially. Combination with PGII also showed similar results (Table 3).

Table 4 presents the results of joint effect of gastric atrophy, poor dental health, and oral hygiene habit on OSCC risk after breaking the matching factors. Adjusted OR for association of DMFT with OSCC was 1.60 (95% CI: 1.14, 2.24). Although based

on modest numbers of cases, presence of poor dental hygiene, indicated as higher than median DMFT, and gastric atrophy rendered a more than four-fold excess risk of OSCC compared with the group with neither poor dental hygiene nor atrophy. The RERI between low DMFT and atrophy was 1.47 (95% CI: -1.15, 4.1), synergy index (95% CI) 1.90 (0.6, 6.1), and the attributable proportion (95% CI) 0.36 (-0.14, 0.85). We used frequency of tooth brushing as another measure of dental and oral hygiene in combination with gastric atrophy, which resulted in an eight-fold increase in OSCC risk (OR: 8.65, 95% CI: 3.65, 20.46) compared with non-atrophic group who occasionally or regularly brush their teeth. The RERI between habit of never brushing teeth and atrophy was 4.34 (95% CI: -1.07, 9.76), synergy index 2.24 (95% CI: 0.97, 5.15), and attributable proportion 0.49 (95% CI: 0.14, 0.83). Although based on fewer sets, conditional logistic regression model showed similar results (data not shown).

To quantify the magnitude of misclassification bias that occurs in presence of nonatrophic gastritis, we estimated the classification probabilities using external validation study (Nasrollahzadeh et al, 2011). Effects of the range of PGII levels on diagnostic accuracy of atrophy and corrected ORs for the association with OSCC are

Table 2 Levels of pepsinogen I, pepsinogen II, and *H. pylori* serology and their associations with OSCC risk, Golestan province, Iran, 2003–2007

	Controls, N (%)	Cases, N (%)	Crude OR (95% CI)	Adjusted OR (95% CI) ^a
PgI ($\mu\text{g l}^{-1}$)				
≥ 55	449 (85.7)	238 (81.2)	Referent	Referent
< 55	75 (14.3)	55 (18.8)	1.33 (0.89–1.97)	1.39 (0.93–2.09)
PgII ^b ($\mu\text{g l}^{-1}$)				
< 11.8	246 (46.9)	113 (38.7)	Referent	Referent
≥ 11.8	278 (53.1)	179 (63.1)	1.37 (1.01–1.85)	1.40 (1.01–1.92)
PgI and PgII ^b ($\mu\text{g l}^{-1}$)				
PgI ≥ 55 and PgII < 11.8	189 (36.1)	71 (24.2)	Referent	Referent
PgI < 55 and PgII < 11.8	57 (10.9)	42 (14.3)	1.92 (1.15–3.22)	2.01 (1.18–3.45)
PgI ≥ 55 and PgII ≥ 11.8	260 (49.6)	167 (57)	Referent	Referent
PgI < 55 and PgII ≥ 11.8	18 (3.4)	12 (4.4)	1.02 (0.47–2.23)	1.20 (0.54–2.66)
<i>H. pylori</i> status				
<i>H. pylori</i> – cagA –	52 (10)	35 (12)	Referent	Referent
<i>H. pylori</i> + cagA –	34 (6.5)	14 (4.8)	0.64 (0.30–1.35)	0.70 (0.32–1.53)
<i>H. pylori</i> + cagA +	351 (67.2)	179 (61.3)	0.73 (0.46–1.17)	0.79 (0.48–1.29)
<i>H. pylori</i> – cagA +	85 (16.3)	64 (21.9)	1.05 (0.60–1.85)	1.15 (0.63–2.08)

Abbreviations: cagA = cytotoxin-associated gene A; CI = confidence interval; *H. pylori* = *Helicobacter pylori*; OR = odds ratio; OSCC = oesophageal squamous cell carcinoma; PgI = pepsinogen I; PgII = pepsinogen II. ^aAdjusted for ethnicity (Non-Turkmen or Turkmen), alcohol consumption (never or ever), tobacco or opium use (none, only tobacco, only opium, or both), education level (illiterate, primary school or more), and vegetable/fruit consumption. ^bFor one case, PgII level was missing and just PgI level was available.

Table 3 Gastric atrophy defined by other serology criteria in literature and its association with OSCC risk

	Controls, N (%)	Cases, N (%)	Adjusted OR (95% CI) ^a
PgI			
PgI ≥ 30	500 (95.4)	274 (93.5)	Referent
PgI < 30	24 (4.6)	19 (6.5)	1.52 (0.79–2.93)
Stratifying by PgII level ^b			
PgI ≥ 30 and PgII < 11.8	226 (43.1)	97 (33.1)	Referent
PgI < 30 and PgII < 11.8	4 (0.8)	3 (1.0)	1.94 (0.91–4.13)
PgI ≥ 30 and PgII ≥ 11.8	274 (52.3)	176 (60.4)	Referent
PgI < 30 and PgII ≥ 11.8	20 (3.8)	16 (5.5)	1.10 (0.23–5.17)
PgI/PgII ratio ^b			
Ratio ≥ 3	490 (93.5)	265 (90.8)	Referent
Ratio < 3	34 (6.5)	27 (9.2)	1.50 (0.85–2.60)
Stratifying by PgII level ^b			
Ratio ≥ 3 and PgII < 11.8	233 (44.5)	100 (34.1)	Referent
Ratio < 3 and PgII < 11.8	21 (4.0)	14 (4.8)	2.32 (1.00–5.39)
Ratio ≥ 3 and PgII ≥ 11.8	257 (49.0)	165 (56.7)	Referent
Ratio < 30 and PgII ≥ 11.8	13 (2.5)	13 (4.4)	1.03 (0.49–2.20)
PgI and PgII/PgII ratio ^b			
PgI ≥ 70 or ratio ≥ 3	497 (94.9)	267 (91.5)	Referent
PgI < 70 and ratio < 3	27 (5.1)	25 (8.5)	1.69 (0.93–3.10)
Stratifying by PgII level ^b			
(PgI ≥ 70 or ratio ≥ 3) and PgII < 11.8	233 (44.5)	100 (34.19)	Referent
(PgI < 70 and ratio < 3) and PgII < 11.8	14 (2.6)	12 (4.1)	2.31 (0.99–5.38)
(PgI ≥ 70 or ratio ≥ 3) and PgII ≥ 11.8	264 (50.4)	167 (57.4)	Referent
(PgI < 70 and ratio < 3) and PgII ≥ 11.8	13 (2.5)	13 (4.4)	1.24 (0.52–2.94)

Abbreviations: CI = confidence interval; OR = odds ratio; OSCC = oesophageal squamous cell carcinoma; PgI = pepsinogen I; PgII = pepsinogen II. ^aAdjusted for ethnicity (Non-Turkmen or Turkmen), alcohol consumption (never or ever), tobacco or opium use (none, only tobacco, only opium, or both), education level (illiterate, primary school or more), and vegetable/fruit consumption. ^bFor one case, PgII level was missing and just PgI level was available.

presented in Table 5. Differential misclassification of gastric atrophy for cut points with low specificity (<80%) was far from the null. When exposure detection was better among controls, the resulted OR was extensively biased.

DISCUSSION

In this study, gastric atrophy, as indicated by low serum pepsinogen, was associated with an increased risk of OSCC.

Table 4 Combined effects of gastric atrophy and dental health or oral hygiene habit on OSCC risk after breaking the matching factors

	Gastric atrophy –		Gastric atrophy +	
	Case/control	OR ^a (95% CI)	Case/control	OR ^a (95% CI)
DMFT ≤28	29/104	Referent	12/26	1.65 (0.74–3.67) P = 0.20
DMFT >28	40/84	1.71 (0.98–2.98) P = 0.06	30/31	3.47 (1.81–6.64) P < 0.001
Ever tooth brushing	14/87	Referent	5/24	1.40 (0.44–4.39) P = 0.57
Never tooth brushing	57/100	4.12 (2.05–8.28) P < 0.001	37/33	8.77 (3.89–19.72) P < 0.001

Abbreviations: CI = confidence interval; DMFT = sum of decayed, missed, and filled teeth; OR = odds ratio; OSCC = oesophageal squamous cell carcinoma. Measure of interaction on additive scale: relative excess risk due to interaction with low DMFT (95% CI): 1.47 (–1.15 to 4.1) and with never tooth brushing habit (95% CI): 4.34 (–1.07 to 9.76). Measure of interaction on multiplicative scale: between atrophy and low DMFT: OR (95% CI): 1.44 (0.63–3.32); P = 0.39 and measure of interaction between atrophy and never tooth brushing on multiplicative scale: OR (95% CI): 1.78 (0.30–10.42); P = 0.52. ^aORs (95% CI) levels adjusted for age, sex, residence area, ethnicity (Non-Turkmen or Turkmen), alcohol consumption (never or ever), tobacco or opium use (none, only tobacco, only opium, or both), education level (illiterate, primary school or more), and vegetable/fruit consumption.

Table 5 Gastric atrophy defined by PGI <55 µg l^{–1} and its association with OSCC risk under assumption of various sensitivity (se) and specificity (sp) among cases and controls

			Controls						
			All PGII	Only PGII <5	Only PGII <11.8	Only PGII <15	Only PGII <19	Only PGII <30	Only PGII >11.8
			Se: 0.62	0.99	0.83	0.80	0.80	0.76	0.37
			Sp: 0.95	0.68	0.89	0.91	0.92	0.94	0.98
Cases	Se	Sp							
All PGII	0.62	0.95	1.63	a	6.6	3.9	3.3	2.4	0.59
Only PGII <5	0.99	0.68	a	0.79	a	a	a	a	a
Only PGII <11.8	0.83	0.89	0.6	a	2.5	1.5	1.3	0.9	0.2
Only PGII <15	0.80	0.91	0.8	a	3.3	1.97	1.7	1.2	0.3
Only PGII <19	0.80	0.92	0.9	a	3.6	2.2	1.83	1.3	0.3
Only PGII <30	0.76	0.94	1.1	a	4.6	2.8	2.3	1.65	1.66
Only PGII >11.8	0.37	0.98	4.7	a	19.1	11.4	9.6	6.8	1.69

Abbreviations: OSCC = oesophageal squamous cell carcinoma; PGI = pepsinogen I; PGII = pepsinogen II. ORs are adjusted for age, sex, and residence area in unconditional logistic regression model. ^aThese cells yielded negative adjusted counts, which were impossible values for true counts, thus corrected ORs could not be estimated.

Pepsinogens are markers of mixed conditions, including gastritis and atrophy (He *et al*, 2011), which could be of distinct etiologies in cancer patients. In spite of observed association between serological nonatrophic pangastritis with OSCC in our study, the probability of developing gastritis secondary to oesophageal cancer is not ignorable because of several evidences. Most importantly, studies with cohort design did not detect significant association between PGII and OSCC risk. In addition, cancer patients may develop gastritis due to non-steroidal anti-inflammatory drug (NSAID) use, opportunistic infections in stomach (cytomegalovirus, Epstein–Barr virus), malnutrition, anaemia, and consequent use of iron tablets (Abraham *et al*, 1999; Lauwers *et al*, 2010). Stage of tumours in most of our study population were III or IV at the time of diagnosis, and dysphagia was the presenting symptom, which led to high proportion of NSAID consumption among cases compared with controls (24.7% vs 15.8%, P = 0.001). On the other hand, development of marked atrophy needs long time period (Adamu *et al*, 2011) and its initiation during OSCC progression with short lead time is unlikely. Thus, in studying the relationship between gastric atrophy and OSCC when serum collection period is close to cancer diagnosis, PGI provides more valid estimate than PGI/PGII ratio.

Low sensitivity for atrophy detection might lead to loss of precision and power by unnecessarily reducing the exposed sample size. We chose atrophy cutoff point at high specificity level (97%) however, false negative probability of 0.4 and low prevalence of gastric atrophy could potentially result in misclassification of exposure among cases and controls. Which we believe, if evolves, would be differential.

Histology is the gold standard for atrophy diagnosis, however, detection of atrophy is difficult because of the limitation in number of biopsies and patchy nature of atrophy. Furthermore, mucosal inflammation and separation of glands by inflammatory infiltrate would add more diagnostic difficulties in severe gastritis (Owen, 2003). Because inflammation among our cancer patients was more probable than healthy controls, the error probabilities for unexposed cases and controls might not be identical. To decrease the effect of mixed category and to reduce the problem of non-identical false-negative probabilities, we stratified the presence of nonatrophic pangastritis and observed a two-fold increased OSCC risk associated with atrophy.

Notably, reducing interpretations to a dichotomy could degrade the information. False-negative results are inherent in serological diagnosis of atrophy due to the low sensitivity of the diagnostic method. It is believed that resulted misclassification will drive the

association toward null if non-differential assumptions are met. However, atrophy is an uncommon exposure and its binary value is the result of categorising continuous PGI variable. Therefore, we performed a sensitivity analysis with the external validation data. Our results confirmed that the effect of misclassification due to ignoring nonatrophic pangastritis on association between atrophy and OSCC was substantial. This finding suggests that PGII as a marker of severe inflammation could be used as a stratifying factor.

In this study, we could not detect any association between infection with *cagA*-positive *H. pylori* strains and OSCC risk, which is in contrast to the results from the study among low-risk Swedish population (Ye *et al*, 2004), Japanese alcoholic males (Yokoyama *et al*, 2009), and in accordance to the reports from studies in Linxian of China (Kamangar *et al*, 2007), Magdeburg of Germany (Venerito *et al*, 2011), and among Finnish male smokers (Cook *et al*, 2010). Spontaneous disappearance of *H. pylori* over time or *H. pylori* eradication is possibly common in endemic areas. Acquired immunodeficiency among advanced malnourished cancer patients could be another explanation for reduced immune response to antigens. These possibilities could lead to heterogeneity of *H. pylori*-negative category as a reference. Hence, our risk estimates for *H. pylori* and *CagA* infection are inconclusive because of difficulties in achieving a clean reference group in an endemic area for *H. pylori* infection and among aged population in a case-control design.

Our study might indicate areas to search for the underlying mechanism. There was a suggestive evidence that the estimated joint effect of poor dental health or never tooth brushing and gastric atrophy together was larger than the sum of their effects alone, which is in favour of the hypothesis of the bacterial overgrowth in atrophic stomach and low-hygiene periodontal mucosa (Meurman and Bascones-Martinez, 2011). Endogenous formation of nitrosamines in the oral cavity of those with poor oral hygiene is eightfold higher than that in those with good oral hygiene (Nair *et al*, 1996). Upstream microbial community members from the oral mucosa and oesophagus have the potential to translocate to the stomach, and they might act as reservoirs for recolonisation. Gastric microbiota shows considerable overlap with oral microbiota (Bik *et al*, 2006), however, the mechanism of these two bacterial environment in OSCC carcinogenesis could be

different. Our statistical inference on the presence of interaction between poor oral hygiene and gastric atrophy has less precision than the main finding of the study because of the modest sample size. The observed interaction might be due to other mechanisms, as tooth loss may be an indicator of a less healthy life style. However, we did not detect association between education as a socioeconomic status indicator and DMFT in our data. Also, low DMFT could be a marker for distinct diet because of reduced masticatory ability, hence, the detected association might be due to residual confounding.

This study was a population-based study, matching improved statistical efficiency, and sample size was the largest among the studies so far published. Markers of the main exposure and confounders were validated and extensive information on cofounders was available.

In conclusion, our population-based case-control study confirms that gastric atrophy is a risk factor for OSCC in high-risk area of northern Iran. Excluding nonatrophic pangastritis through stratifying by PGII level decreases the possibility of PGI misclassification due to cancer-induced inflammation. Gastric atrophy and poor oral hygiene may act synergistically in increasing the risk of OSCC.

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Conflict of interest

The authors declare no conflict of interest.

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