Brand interchangeability of pepsinogen tests in the real-world setting after eradication of *Helicobacter pylori*: a community-based study

Tsung-Hsien Chiang1,2, Yen-Nien Chen3, Yi-Ru Chen4, Yu-Hua Tseng5, Chun-Fu Shieh6, Cheng-Ying Liu7, Han-Mo Chiu1,8, Hung Chiang9, Chia-Tung Shun10, Ming-Shiang Wu1, Jaw-Town Lin1,11 and Yi-Chia Lee1,8,12*

Abstract

**Background:** Serum pepsinogen (PG) is recommended as a screening test for premalignant gastric lesions. However, real-world evidence demonstrating its applicability and equivalence between different test brands is limited.

**Methods:** Mass screening began in 2018 in a high-risk Taiwanese population after eradication of *Helicobacter pylori*, with the first stage of two PG tests (GastroPanel®️, Helsinki, Finland and LZ-Test®, Tokyo, Japan) and the second stage of endoscopy. A positive test was defined as PG-I < 30 ng/mL or PG-I/II ratio < 3 for GastroPanel®️ and PG-I ≤ 70 ng/mL and PG-I/II ratio ≤ 3 for LZ-Test®️. Index lesions included atrophic gastritis and intestinal metaplasia. Test performance was evaluated based on the participation rate, positivity rate, referral rate, positive predictive value (PPV), and the detection rate.

**Results:** Among 7616 eligible participants, 5117 (67.2%) received PG tests and 284 (5.6%) tested positive. Of those who tested positive, 105 (37.0%) underwent endoscopy. Overall PPVs for atrophic gastritis and intestinal metaplasia were 12.4% and 18.9%, respectively, with detection rates of 2.5 and 3.9 per 1000, respectively. Correlations of numerical measures between tests were high and the agreements of test results were substantial. The PPVs (16.3% vs. 16.3% and 23.8% vs. 21.3%, *P* = 1.00 and 0.71, respectively), detection rates (2.5 vs. 2.5 and 3.7 vs. 3.3 per 1000, *P* = 1.00 and 0.27, respectively), and the stage distributions of gastritis were all comparable, which were confirmed by multiple regression analyses.

**Conclusions:** PG testing is effective for mass screening after eradication of *H. pylori*. Tests from different manufacturers, even using different analytical methods and cutoff criteria, can perform equivalently.

**Keywords:** *Helicobacter pylori*, Pepsinogen, Screening, Positive predictive value, Detection rate

Introduction

Gastric cancer is the sixth most common cancer and the third most common cause of cancer deaths worldwide [1]. Screening and prevention strategies are urgently needed to reduce the significant burden of gastric cancer. *Helicobacter pylori* infection is the initiator for gastric cancer carcinogenesis [2], so active screening and eradication of this bacterium is considered one of the best strategies [3]. However, the benefit of such an approach depends on the magnitude of pre-existing molecular damages in the gastric mucosa [4]. For patients who already harbor premalignant gastric lesions after long-term inflammation, such as atrophic gastritis and intestinal metaplasia, they can retain irreversible changes after the eradication treatment [5, 6]. Therefore, when *H.
**Methods**

**Study population**

Our eligible population comprised 7616 inhabitants of an offshore archipelago (Matsu Islands, Taiwan), who were aged 30 years or more and registered in the population registry. Because of the high gastric cancer incidence in this population, a mass screening and eradication program of *H. pylori* has been implemented since 2004 [3]. All eligible inhabitants were invited by mail, telephone, or announcement in social media and newspapers to attend the screening program. Participants’ demographic data, social habits, and medical histories were recorded in a structured questionnaire. *H. pylori* infection was determined using the $^{13}$C-urea breath test, those testing positive received eradication treatment, and those who failed initial treatment were retreated [15, 16]. By the end of 2018, six rounds of mass eradication have been conducted. As a result, the prevalence rate of *H. pylori* infection in the Matsu Islands declined from 64.2% before the mass eradication program to about 10% recently, accompanied by a 53% reduction in gastric cancer cases [14].

**Study design**

Since scattered cases of gastric cancer occurred during the mass eradication program, a two-stage endoscopic screening program was added to the chemoprevention program in 2015, with the main purpose to identify participants who were still at high risk for gastric cancer after *H. pylori* eradication and provide them with endoscopic screening. This program was done using the serological test with PG, in which the test positives would be referred for endoscopic examination and histological evaluation. Preliminary results were reported previously [17].

In 2018, because various brands of PG were available in the market, we tested whether or not the performance of two brands, the GastroPanel® (Biohit HealthCare, Helsinki, Finland) and the LZ-Test® (Eiken Chemical Co., Ltd, Tokyo, Japan), was equivalent. We applied a parallel test design [18], in which participants with positive PG results for either one of the two tests would be referred for endoscopic examination. We assumed the possibility of false negatives could be lower than those based on a single test.

These tests also included measurements of the anti-*H. pylori* IgG or gastrin-17; however, since our initial experience indicated that the values of these two measures had been reduced after *H. pylori* eradication [17], we only analyzed the results of the PG-I, PG-II, and PG-I/II ratio in this study. The study protocol was approved by the Institutional Review Board (IRB) of National Taiwan University Hospital (IRB No: 201406021RINA), and written informed consent was obtained from all participants before they entered the study.

**Serological tests**

After an overnight fast, participants received blood sampling for both GastroPanel® and LZ-Test® at the same time. Participants were requested to stop taking acid suppressing drugs, particularly proton-pump inhibitors, at least one week before testing. After the blood sample was
The blood sample was then added to the wells, creating antibody-antigen complexes based on color intensity. The PG-specific antibodies were coated onto the surface of a microplate. The blood sample was then added to the wells, creating the antibody-antigen reaction. A secondary antibody linked to an enzyme was used to bind the antigen-antibody complexes. Finally, color-producing substances were added to react with the enzyme and color was produced. The PG concentration in the serum sample could be quantified according to the intensity of color. The cutoff concentrations recommended by the manufacturer were set at PG-I ≤ 30 ng/mL or PG-I/II ratio < 3 [19]. The sensitivity, specificity, and positive likelihood ratio in the prediction of premalignant gastric lesions had been reported as about 40%, 93%, and 6.0, respectively [20].

The LZ-Test® applied the latex-enhanced turbidimetric immunoassay (L-TIA) that measured the concentration of antigen-antibody complexes based on the optical turbidity. The latex reagent was prepared through the binding between the PG-specific antibodies and the latex particles. After the latex particles reacted with the PG presented in the blood samples, the agglutination reaction would change the turbidity, which could be therefore quantified. The cutoff concentrations recommended by the manufacturer were set at PG-I ≤ 70 ng/mL and PG-I/II ratio ≤ 3 [21]. The sensitivity, specificity, and positive likelihood ratio in the prediction of premalignant gastric lesions had been reported as about 59%, 89%, and 5.5, respectively [22].

Endoscopic and histological evaluations
During the endoscopic examination, biopsy specimens were taken from gastric mucosa in the antrum (from the greater and lesser curvatures 2–3 cm from the pylorus) and corpus (one each from the lesser and greater curvature at the middle corpus) according to the modified Sydney protocol [23]. Sampling was done at the same locations in all participants to maintain consistency. All specimens were fixed in formalin for histological evaluation. A senior histopathologist (Dr. H Chiang), blinded to participants’ clinical status and test results, performed all histological assessments. Specimens were graded, using the Updated Sydney Classification, as acute inflammation (polymorphonuclear infiltrates), chronic inflammation (lymphoplasmacytic infiltrates), atrophic gastritis (loss of glandular tissue and fibrous replacement), or intestinal metaplasia (presence of goblet cells and absorptive cells) [23]; the severity of each category was rated as none, mild, moderate, or marked. And the severity of premalignant lesions were classified using the criteria of Operative Link for Gastritis Assessment of Atrophic Gastritis (OLGA) and Operative Link for Gastritis Assessment of Intestinal Metaplasia (OLGIM) [24, 25].

Statistical analysis
For the baseline characteristics, categorical data were expressed as a percentage and continuous data were expressed as a mean, with standard deviation. The baseline characteristics of participants were compared using the Student t-test or chi-square test. The performance of the whole program was evaluated according to the consecutive steps of an organized screening program, including the invitation, participation, testing, referral, and the final diagnosis. The indicators included the test positivity rates (the number of positive cases divided by the number of participants), referral rate for endoscopic examinations (the number of endoscopies divided by the number of positive cases), positive predictive value (PPV: the number of participants diagnosed with premalignant gastric lesions divided by the number of diagnostic endoscopies), and the detection rate (the number participants diagnosed with premalignant gastric lesions divided by the number of participants).

Two brands of PG tests were then compared. First, the Pearson’s correlation coefficient (a level of 0.7 or more indicated a strong correlation) was applied for the numerical measures of PG-I, PG-II, and PG-I/II ratio, and the Kappa statistics (a level of 0.6 or more indicated substantial agreement) was applied for the positivity rates. Second, the positivity rate, referral rate, PPV, and detection rate were compared between two tests using the two-sample proportional test. To adjust for differences in baseline characteristics, logistic regression analyses were performed, with the results expressed as the crude and adjusted odds ratios and the corresponding 95% confidence intervals (CIs). Third, the positivity rates between two tests in patients with premalignant lesions were compared using the two-sample proportion test according to the OLGA and OLGIM stages.

All statistical analyses were performed using Stata 14 software (StataCorp LLC, College Station, TX, USA). All P-values were two-sided, with P<0.05 considered statistically significant.

Results
Baseline characteristics
The study flowchart is shown in Fig. 1. Among 7616 eligible participants, 5117 received the PG tests, leading to a coverage rate of 67.2%. Among these participants, 284
(5.6%) had positive results. Among participants tested positive, 51 participants tested positive for GastroPanel®, only, 92 tested positive for LZ-Test® only, and 141 tested positive for both tests. A total of 105 (37%) participants tested positive and underwent endoscopic examinations: 13 were diagnosed with atrophic gastritis, 19 diagnosed with intestinal metaplasia, and 7 diagnosed with both atrophic gastritis and intestinal metaplasia.

Baseline characteristics of participants are shown in Table 1. Participants who tested positive for either GastroPanel® or LZ-Test® were older than those who tested negative (58.1 vs. 54.8 years, \(P<0.001\)). A history of hypertension was more common in those who tested positive than those who tested negative (38.7% vs. 29.9%, \(P=0.002\)), which was probably because the subjects who tested positive were older than those who tested negative. The mean value of body mass index was similar between those who tested positive and negative. Also, the proportion of male sex, cigarette smokers, alcohol drinkers, betel nut chewers, and other medical conditions, including diabetes mellitus, hyperlipidemia, cardiovascular disease, stroke, chronic hepatitis B, chronic hepatitis C, and chronic kidney disease, were all comparable between participants with positive and negative results.

The comparison of baseline data between endoscopic receivers and refusers is shown in Additional file 1: Table S1. No significant differences were noted except that the proportion of hyperlipidemia and cardiovascular disease were higher in the endoscopic receivers.

**Performance of the whole program**

The number of positive tests, positivity rates, number of endoscopies, and the referral rate of endoscopic examinations, which are stratified by age, sex, and testing brand, are shown in Table 2. The positivity rates were 3.8%, 4.6%, and 5.6% for GastroPanel®, LZ-Test®, and the parallel combination (i.e. either one was positive), respectively. The positivity rate of the parallel combination was higher than that of GastroPanel® (5.6% vs. 3.8%, \(P<0.001\)) and LZ-Test® (5.6% vs. 4.6%, \(P=0.021\)). The positivity rates of the parallel combination were slightly higher for males (5.8% vs. 5.3%, \(P=0.43\)), but were significantly higher in the older-age group \((\geq 50\) years) than the younger-age group \((30–49\) years) (6.1% vs. 4.5%, \(P=0.016\)).

Results of the PPV and detection rate are shown in Table 3. In addition to the overall results, the results were also stratified by age, sex, and test brand. The overall PPV (i.e., two-test combination) for atrophic gastritis
was slightly lower than that of the individual test with either GastroPanel® or LZ-Test® (12.4% vs. 16.3% and 16.3%, both \( P = 0.45 \)). Findings for intestinal metaplasia were similar (18.9% vs. 23.8% and 21.3%, \( P = 0.42 \) and 0.69, respectively). Taking into consideration the referral rates, the detection rate for atrophic gastritis was the same for overall and the individual tests (2.5 per 1000). The detection rate for intestinal metaplasia was also comparable (3.9 vs. 3.7 and 3.3 per 1000, \( P = 0.60 \) and 0.10, respectively).

The PPVs for either atrophic gastritis or intestinal metaplasia was slightly higher in males than females, and was significantly higher in the older-age than the younger-age group. The detection rates for atrophic gastritis and intestinal metaplasia were both higher in males than females, and were both higher in the older-age than the younger-age group.

### Comparisons of the measures between two tests

As shown in Fig. 2, the quantitative measures of PG-I, PG-II, and PG-I/II ratio for GastroPanel® were about 40% higher, 26% lower, and 2.3-fold higher than those for LZ-Test®, respectively. However, the correlation coefficients between these two tests were 0.96 (95% CI 0.95–0.96), 0.91 (95% CI 0.90–0.91), and 0.76 (95% CI 0.75–0.77) for the PG-I, PG-II and the PG-I/II ratio, respectively, indicating a strong correlation (all \( P < 0.001 \)). For the positivity rate, the kappa value for agreement was 0.65 (95% CI 0.60–0.70), also indicating a substantial agreement (\( P < 0.001 \)).

### Comparisons of the screening indicators between two tests

In the comparison between two tests, the positivity rate was significantly lower in GastroPanel® than that in LZ-Test® (3.8% vs. 4.6%, \( P = 0.044 \)), particularly in the younger-age group (2.2% vs. 3.5%, \( P = 0.018 \)). The referral rate for diagnostic examination was slightly higher in GastroPanel® than LZ-Test® but the difference was not statistically significant (41.1% vs. 33.9%, \( P = 0.12 \)). The PPVs for atrophic gastritis were the same at 16.3% (\( P = 1.00 \)) and the results for intestinal metaplasia were 23.8% and 21.3% (\( P = 0.71 \)) for GastroPanel® and LZ-Test®, respectively. Two tests had the same detection rate for atrophic gastritis, 2.5 per 1000, and their detection rates as for intestinal metaplasia were also comparable (3.7 vs. 3.3 per 1000, \( P = 0.27 \)).

### Multiple regression analyses

Results for the logistic regression analyses are shown in Table 4. Univariate analyses showed non-significant differences between the two test brands regarding the PPV for atrophic gastritis (OR 1.00, 95% CI 0.42–2.38, \( P = 1.00 \)) or intestinal metaplasia (OR 1.16, 95% CI 0.54–2.48, \( P = 0.70 \)). Taking into consideration the referral rate for endoscopy, the results for the detection rate were still similar between two tests for either atrophic gastritis (OR

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### Table 1  Baseline characteristics of the screening participants stratified by the results of the PG tests

| Screening participants | Participants (n = 5117) | Negative result (n = 4833) | Positive result* (n = 284) | P value† |
|------------------------|-------------------------|----------------------------|---------------------------|---------|
| Mean age, years (SD; range) | 55.0 (12.9; 30–100) | 54.8 (12.9; 30–100) | 58.1 (13.5; 30–96) | <0.001 |
| Male sex, no. (%) | 2510 (49.1) | 2364 (48.4) | 146 (51.4) | 0.33 |
| Body mass index, kg/m² (SD; range) | 25.1 (4.0; 14.6–52.1) | 25.1 (4.0; 14.6–52.1) | 25.1 (3.9; 15.8–37.5) | 0.93 |
| Social habits, no. (%) | | | | |
| Current smoker | 752 (14.7) | 708 (14.6) | 44 (15.5) | 0.68 |
| Regular alcohol drinking | 320 (6.3) | 307 (6.3) | 13 (4.6) | 0.25 |
| Betel nut chewing | 221 (4.3) | 208 (4.3) | 13 (4.6) | 0.81 |
| Medical history, no. (%) | | | | |
| Hypertension | 1553 (30.3) | 1443 (29.9) | 110 (38.7) | 0.002 |
| Diabetes mellitus | 464 (9.1) | 441 (9.1) | 23 (8.1) | 0.57 |
| Hyperlipidemia | 375 (7.3) | 346 (7.2) | 29 (10.2) | 0.06 |
| Cardiovascular disease | 269 (5.3) | 253 (5.2) | 16 (5.6) | 0.77 |
| Stroke | 30 (0.6) | 29 (0.6) | 1 (0.4) | 0.67 |
| Chronic hepatitis B | 626 (12.2) | 586 (12.1) | 40 (14.1) | 0.32 |
| Chronic hepatitis C | 27 (0.5) | 24 (0.5) | 3 (1.1) | 0.18 |
| Chronic kidney disease | 56 (1.1) | 54 (1.1) | 2 (0.7) | 0.53 |

SD standard deviation  
† Test positives for either GastroPanel (Biohit HealthCare, Helsinki, Finland) or LZ-Test (Eiken Chemical Co., Ltd, Tokyo, Japan)  
‡ \( P < 0.05 \) in the comparison between the test positives and negatives
| Brands of PG test | Participants, no | Positive test, no | Positivity rate (%) | Endoscopic examination, no | Referral rate for endoscopic examination (%) |
|-------------------|------------------|------------------|---------------------|---------------------------|---------------------------------|
|                   | Test 1 | Test 2 | Test 1 + 2 | Test 1 | Test 2 | Test 1 + 2 | Test 1 | Test 2 | Test 1 + 2 | Test 1 | Test 2 | Test 1 + 2 | Test 1 | Test 2 | Test 1 + 2 |
| Male              |        |        |           |        |        |           |        |        |           |        |        |           |        |        |           |
| 30–49 years       | 842    | 17     | 29        | 37     | 2.0    | 3.4       | 4.4    | 10     | 9         | 15     |          |          | 58.8   | 31.0   | 405     |
| ≥ 50 years        | 1668   | 85     | 93        | 109    | 5.1    | 5.6       | 6.5    | 34     | 36        | 43     |          |          | 40.0   | 38.7   | 394     |
| Subtotal          | 2510   | 102    | 122       | 146    | 4.1    | 4.9       | 5.8    | 44     | 45        | 58     |          |          | 43.1   | 36.9   | 397     |
| Female            |        |        |           |        |        |           |        |        |           |        |        |           |        |        |           |
| 30–49 years       | 1006   | 24     | 36        | 46     | 2.4    | 3.6       | 4.6    | 8      | 8         | 12     |          |          | 33.3   | 22.2   | 261     |
| ≥ 50 years        | 1601   | 66     | 75        | 92     | 4.1    | 4.7       | 5.7    | 27     | 26        | 35     |          |          | 40.9   | 34.7   | 380     |
| Subtotal          | 2607   | 90     | 111       | 138    | 3.5    | 4.3       | 5.3    | 35     | 34        | 47     |          |          | 38.9   | 30.6   | 341     |
| Both sexes        |        |        |           |        |        |           |        |        |           |        |        |           |        |        |           |
| 30–49 years       | 1848   | 41     | 65        | 83     | 2.2†   | 3.5†      | 4.5    | 18     | 17        | 27     |          |          | 43.9   | 26.2   | 325     |
| ≥ 50 years        | 3269   | 151    | 168       | 201    | 4.6    | 5.1       | 6.1    | 61     | 62        | 78     |          |          | 40.4   | 36.9   | 388     |
| Total             | 5117   | 192    | 233       | 284    | 3.8†   | 4.6†      | 5.6    | 79     | 79        | 105    |          |          | 41.1   | 33.9   | 370     |

PG: pepsinogen; Test 1, GastroPanel (Biohit HealthCare, Helsinki, Finland); Test 2, LZ-Test (Eiken Chemical Co., Ltd, Tokyo, Japan); Test 1 + 2, two tests in parallel

† Test 1 + 2 indicated the parallel combination of two tests (either one of the two tests was positive)

† † † P < 0.05 in the comparison between test 1 and test 2
1.00, 95% CI 0.45–2.23, \( P = 1.00 \)) or intestinal metaplasia (OR 1.13, 95% CI 0.57–2.21, \( P = 0.73 \)).

When adjusted for age, sex, body mass index, social habits, and medical histories, the differences in PPVs for atrophic gastritis (adjusted OR 1.12, 95% CI 0.40–3.12, \( P = 0.83 \)) (Additional file 1: Figure S1) and intestinal metaplasia (adjusted OR: 1.75, 95% CI 0.65–4.67, \( P = 0.27 \)) (Additional file 1: Figure S2) between two tests were still non-significant. Results for the detection rates for atrophic gastritis (adjusted OR 1.00, 95% CI 0.45–2.24, \( P = 1.00 \)) (Additional file 1: Figure S3) and intestinal metaplasia (adjusted OR 1.20, 95% CI 0.60–2.40, \( P = 0.53 \)) (Additional file 1: Figure S4) were also non-significant.

Older age was borderline significantly associated with a higher PPV for atrophic gastritis (adjusted OR 3.68, 95% CI 0.67–20.16, \( P = 0.13 \)) while older age was significantly associated with a higher PPV for intestinal metaplasia (adjusted OR 5.97, 95% CI 1.12–31.77, \( P = 0.036 \)). Between sexes, differences in PPV were not statistically significant. The detection rate for atrophic gastritis was significantly higher in the older age group (adjusted OR 5.61; 95% CI 1.30–24.15, \( P = 0.021 \)) and slightly higher for

### Table 3 The performance indicators of screening, stratified by the age, sex, and the brands of tests

| Brands of PG test ¹ | Positive predictive value (%)* | Detection rate (per 1000)* |
|---------------------|--------------------------------|---------------------------|
|                     | Atrophic gastritis | Intestinal metaplasia | Atrophic gastritis | Intestinal metaplasia |
|                     | Test 1  Test 2 Test 1 + 2 | Test 1  Test 2 Test 1 + 2 | Test 1  Test 2 Test 1 + 2 | Test 1  Test 2 Test 1 + 2 |
| **Male**            |                  |                          |                  |                          |
| 30–49 years         | 10.0  10.0  6.7   | 10.0  11.1  6.7         | 1.2  1.2  1.2     | 1.2  1.2  1.2          |
| ≥ 50 years          | 20.6  19.4  16.3  | 32.3  28.0  25.6        | 4.2  4.2  4.2     | 4.2  4.2  4.2          |
| **Subtotal**        | 18.2  17.4  13.8  | 27.3  26.7  20.7        | 3.2  3.2  3.2     | 4.8  4.8  4.8          |
| **Female**          |                  |                          |                  |                          |
| 30–49 years         | 11.1  12.5  8.3   | 11.1  11.1  7.7         | 1.0  1.0  1.0     | 1.0  1.0  1.0          |
| ≥ 50 years          | 14.8  15.4  11.4  | 22.2  15.4  20.0        | 2.5  2.5  2.5     | 3.7  2.5  4.4          |
| **Subtotal**        | 13.8  14.7  10.6  | 19.4  14.3  16.7        | 1.9  1.9  1.9     | 2.7  1.9  3.1          |
| **Both sexes**      |                  |                          |                  |                          |
| 30–49 years         | 10.5  11.1  7.4   | 10.5  11.1  7.1         | 1.1  1.1  1.1     | 1.1  1.1  1.1          |
| ≥ 50 years          | 18.0  17.7  14.1  | 27.9  24.2  23.1        | 3.4  3.4  3.4     | 5.2  4.6  5.5          |
| **Total**           | 16.3  16.3  12.4  | 23.8  21.3  18.9        | 2.5  2.5  2.5     | 3.7  3.3  3.9          |

PG = pepsinogen; Test 1 = GastroPanel (Biohit HealthCare, Helsinki, Finland); Test 2 = LZ-Test (Eiken Chemical Co., Ltd, Tokyo, Japan); Test 1 + 2 = two tests in parallel

*Positive predictive value was defined as the number of participants with premalignant lesions/ the number of participants positive to PG tests having attended an upper endoscopy. Detection rate was defined as the number of participants with premalignant lesions detected/ the number of participants having received the PG testing

¹ None reaches the significant \( P \) level of < 0.05 in the comparison between Test 1 and Test 2

### Fig. 2 Correlations of numerical measures between two tests using scatter plots. The \( R \) indicates the correlation coefficient (Y axis: GastroPanel and X axis: LZ-Test). The linear regression lines for the PG-I, PG-II, and PG-I/II ratio are \( Y = 1.40X + 7.78 \), \( Y = 0.74X - 0.99 \), and \( Y = 2.31X + 1.45 \), respectively
Regarding intestinal metaplasia, significantly higher detection rates were noted for either the older age group (adjusted OR 8.91; 95% CI 2.12–37.51, \( P = 0.003 \)) or the male sex group (adjusted OR 2.23; 95% CI 1.01–4.96, \( P = 0.049 \)).

**Comparison of positivity rates according to OLGA and OLGIM stages**

In subjects with histologically documented atrophic gastritis and intestinal metaplasia, 84.6% and 78.9% of them tested positive for both tests, respectively. The details are shown in Fig. 3; in patients with atrophic gastritis (all were graded with OLGA stage 1 diseases), both 92.3% yielded positive results for GastroPanel\textsuperscript{®} and LZ-Test\textsuperscript{®}. In patients with intestinal metaplasia, 94.7% and 84.2% yielded positive results for GastroPanel\textsuperscript{®} and LZ-Test\textsuperscript{®}, respectively; the difference was not significant (\( P = 0.29 \)). In details, the positivity rates were remarkably similar in patients with either the OLGIM stage 1 (100% vs. 75.0%, \( P = 0.13 \)), stage 2 (87.5% vs. 87.5%, \( P = 1.00 \)), or the stage 3 diseases (100% vs. 100%, \( P = 1.00 \)).

**Discussion**

This community-based study not only confirmed the applicability of PG testing for mass screening, but also made extensive comparisons between two different brands of PG testing. We found that although their absolute measures of PG-I, PG-II, and the PG-I/II ratio were significantly different, the correlations between them were high and the agreements in test results were substantial. We also found that, although they applied different immunoassay methods for measurement and defined different cutoff criteria, the performance of the two tests was comparable in the prediction of premalignant gastric lesions, particularly in terms of PPV and detection rate. The findings were replicated in multiple regression analysis that adjusted for differences in the baseline characteristics of participants. The histological severities of the identified atrophic gastritis and intestinal metaplasia were also similar based on unified scoring systems. Collectively, the results indicate that two different brands of PG tests perform equivalently in predicting premalignant gastric lesions in the real-world environment, supporting their interchangeability.
We evaluated the real-world performance of PG testing according to the step-by-step principles of screening [26]. First, our high population participation rate of about 67% supported PG testing as a tool for mass screening. Second, regarding the referral for endoscopic diagnosis, the endoscopic rate of about one third was relatively low, which was due to the invasive nature and the lower population acceptability of endoscopic screening. Third, regarding the PPV, the value of about one fifth has indicated the moderate predictability of PG testing in a high-risk population [27]. Finally, the detection rate of about 3 cases per 1000 participants, which took into account the positivity rate, referral rate, and the PPV, replicated the results of population-wide screening programs for other types of cancers [28].

Our study demonstrated a high correlation of quantitative measures between the two test brands, which is supported by two previous studies. First, one Japanese study, based on 304 blood samples, showed the correlation coefficients of 0.98, 0.98, and 0.92, respectively, for the PG-I, PG-II, and PG-I//II ratio between GastroPanel® and LZ-Test® [11]. Another Latvian study, based on 805 blood samples, also showed the high correlation coefficients of 0.89, 0.90, and 0.86 for the PG-I, PG-II, and PG-I//II ratio, respectively, between these two tests [13]. However, whether these results could be interpreted as equivalent in the real-world setting remained unclear. It was primarily because their absolute measures and cutoff definitions were so different. In our study, the prevalence rate of the premalignant conditions has declined after mass eradication of H. pylori [14] so the PPVs of the PG tests also decreased. As a result, the overlap of positive results between two tests may appear lower (141/284, 49.6%). Nonetheless, when we focused on the subjects with histologically documented premalignant conditions, a large proportion of them tested positive for both tests. Therefore, when we interpreted the findings with the concept of sensitivity (the number of test positives divided by the number of patients diagnosed with premalignant lesions), which was not affected by the lower prevalence rate of the premalignant lesions, the test consistency was actually high.

In our study, we applied the two tests in parallel so the prevalence rates of premalignant gastric lesions were the same. The observed similarities in the PPVs (i.e., the prevalence rate of premalignant gastric lesions multiplied by the positive likelihood ratio of the screening test) would indicate that the positive likelihood ratios of two tests should be close to each other. Using histology as the reference standard, the positive likelihood ratio (i.e., the sensitivity divided by one minus specificity) was determined with the diagnostic accuracy study. Our findings supported the robustness of cutoff criteria selection from both companies because their criteria were defined based on the judicious tradeoffs between true positive and false positive results in the receiver operating characteristic curve analyses, which had been optimized according to the similar need for clinical practice [20, 22, 29].

Our previous study has estimated a prevalence rate of about 3% for premalignant gastric lesions in this population that had undergone six rounds of mass eradication [3, 14]. Given the PPVs of about 16% observed in this study, the positive likelihood ratio of PG testing could be estimated at about 5, which was consistent with the results from the previous studies investigating subjects without H. pylori eradication [20, 22, 29]. Given this test capability, we also evaluated how to improve the PPV on a real-world setting. First, we applied two tests in parallel, hoping to increase the test sensitivity so as to reduce the false-negative results. However, this hypothesis was rejected as the PPV and detection rate were comparable when the tests were either applied alone or in combination.

The strengths of our study include the invitation of all candidates in a high-risk community, the step-by-step evaluation following pre-established screening principles, and the head-to-head comparisons of two widely used PG tests in a real-world environment. Our findings can not only be applied in a high-risk population, but can also be generalized to other high-risk populations living in lower-risk countries, such as first-generation immigrants from high-prevalence areas. However, certain limitations should be noted. First, although our study confirmed the consistent performance of PG testing in predicting the presence of premalignant gastric lesions, detecting gastric cancer would have required a longer follow-up of this population, as our previous study has done [10]. Second, research has shown that, after H. pylori eradication, the histology can be improved.
but genetic damage may persist, which would lead to a decreased capacity of PG testing in the prediction of gastric cancer risk [17, 32]. Our study demonstrated a low prevalence rate of advanced-stage pre-malignancies (i.e., stage III–IV OLGA and OLGIM), particularly atrophic gastritis, which was likely related to the effect of *H. pylori* eradication on gastritis regression [14, 17]. Other serological markers or molecular testing based on histological samples could be possible solutions and directions for future research [33, 34]. Finally, regarding the generalizability, although we have demonstrated the equivalent performance between these two brands of PG tests for mass screening, it still need further investigate the interchangeability for other methods to measure the PG concentration except ELISA and L-TIA.

**Conclusions**

We demonstrate high correlations of either the PG-I, PG-II, or PG-I/II measures, good agreements in the test positivity, and the equivalent performance between two brands of PG tests for mass screening, which support the interchangeability of their test results. We also show that targeting older participants will increase the predictability of PG testing in a high-risk population when *H. pylori* screening and treatment has been implemented as common practice. These findings provide important implications in identifying those who will retain gastric cancer risk to properly identify who will benefit most from endoscopic screening.

**Abbreviations**

PG: Pepsinogen; CI: Confidence interval; *H. pylori*: *Helicobacter pylori*; OLGA: Operative Link for Gastritis Assessment of Atrophic Gastritis; OLGIM: Operative Link for Gastritis Assessment of Intestinal Metaplasia; OR: Odds ratio; PPV: Positive predictive value.

**Supplementary Information**

The online version contains supplementary material available at https://doi.org/10.1186/s12876-022-02155-7.

**Acknowledgements**

The authors would like to express their special thanks to the staff of the Eighth Core Lab, Department of Medical Research, National Taiwan University Hospital, for the laboratory testing, and the staff in the Health Bureau of Lienchiang County, for their administrative support in the Matsu islands.

**Authors’ contributions**

THC and YCL: conception and design of the study; statistical analysis; drafting, clinical reasoning and critical revision of the manuscript for important intellectual content. HC and CTS: pathological analysis. CYL: policy development and administrative support. YCL: program execution and evaluation. All authors: administrative, technical, or material support; approval of the final version of the manuscript. YCL: full responsibility for the decision to submit for publication. All authors read and approved the final manuscript.

**Funding**

This study was supported by the Ministry of Science and Technology, R.O.C. (105-2314-B-002-069-MY3 and 108-2314-B-002-140-MY3) and the National Taiwan University Hospital (110-P08). The funding sources had no role in study design, data collection, analysis, interpretation, report writing, or the decision to submit this paper for publication.

**Availability of data and materials**

Most data relevant to the study are included in the article. Additional data are available by contacting the corresponding author.

**Declarations**

**Ethics approval and consent to participate**

All study procedures were in accordance with the ethical standards of the responsible committee on human experimentation and with the Helsinki Declaration of 1964 and later versions. The study protocol was approved by the Institutional Review Board (IRB) of National Taiwan University Hospital (IRB No: 201406021RINA). Informed consent was obtained from all participants.

**Consent for publication**

Not applicable.

**Competing interests**

None of the authors have commercial affiliations that constitute any conflict of interest.

**Author details**

1. Department of Internal Medicine, College of Medicine, National Taiwan University, Taipei, Taiwan. 2. Department of Integrated Diagnostics and Therapeutics, National Taiwan University Hospital, Taipei, Taiwan. 3. Department of Medicine, National Taiwan University Hospital Cancer Center, Taipei, Taiwan. 4. Department of Health Care Management, National Taipei University of Nursing and Health Sciences, Taipei, Taiwan. 5. Lienchiang County Public Health Bureau, Nangan Township, Lienchiang County, Matsu, Taiwan. 6. Lienchiang County Hospital, Nangan Township, Lienchiang County, Matsu, Taiwan. 7. Lienchiang County Government, Nangan Township, Lienchiang County, Matsu, Taiwan. 8. Graduate Institute of Epidemiology and Preventive Medicine, College of Public Health, National Taiwan University, Taipei, Taiwan. 9. Taipei Institute of Pathology, Taipei, Taiwan. 10. Department and Graduate Institute of Forensic Medicine, College of Medicine, National Taiwan University, Taipei, Taiwan. 11. Division of Gastroenterology and Hepatology, Department of Internal Medicine, E-DA Hospital, Kaohsiung, Taiwan. 12. Department of Medical Research, National Taiwan University Hospital, Taipei, Taiwan.

**Received:** 13 November 2021 **Accepted:** 14 February 2022

**Published online:** 18 February 2022

**References**

1. Sung H, Ferlay J, Siegel RL, Laversanne M, Soerjomataram I, Jemal A, et al. Global cancer statistics 2020: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. CA Cancer J Clin. 2021;71:209–49.
2. Smyth EC, Nilsson M, Grabesch H, van Grieken NC, Lordick F. Gastric cancer. Lancet. 2020;396:635–48.
3. Lee YC, Chen TH, Chiu HM, Shun CT, Chiang H, Liu TY, et al. The benefit of mass eradication of *Helicobacter pylori* infection: a community-based study of gastric cancer prevention. Gut. 2013;62:676–82.
4. Lee YC, Chiang TH, Chou CK, Tu YK, Liao WC, Wu MS, et al. Association between *Helicobacter pylori* eradication and gastric cancer incidence: a systematic review and meta-analysis. Gastroenterology. 2016;150:1113–24.
5. de Vries AC, van Grieken NC, Looman CW, Casparie MK, de Vries E, Meijer GA, et al. Gastric cancer risk in patients with premalignant gastric
26. Rabeneck L, Lansdorp-Vogelaar I. Assessment of a cancer screening program. Lancet. 2018;392:2413–21.

25. Capelle LG, de Vries AC, Haringsma J, Ter Borg F, de Vries RA, Bruno MJ, et al. Screening and eradication of Helicobacter pylori for gastric cancer prevention: the Maastricht V/Florina Consensus Report. Gut. 2011;60:178–83.

24. Rugge M, Meggio A, Ihamaki T, Siurala M, Rottger J. Relationships among serum pepsinogen I, serum pepsinogen II, and gastric mucosal histology. A study in relatives of patients with pernicious anemia. Gastroenterology. 1982;83:204–9.

23. Bang CS, Lee JJ, Baik GH. Prediction of chronic atrophic gastritis and gastric cancer in a high-risk population. Gastroenterology. 2021;160:1086–98.

22. Chiang TH, Chuang SL, Chen SL, Chiu HM, Yen AM, Chiu SY, et al. Development and validation of a serum microRNA biomarker panel for detecting gastric cancer in a high-risk population. Gastroenterology. 2021;160:1082–9.

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lesions: a nationwide cohort study in the Netherlands. Gastroenterology. 2008;134:945–52.

6. Song H, Ekheden KG, Zheng Z, Ericsson J, Nyren O, Ye W. Incidence of gastric cancer among patients with gastric precancerous lesions: observational cohort study in a low risk Western population. BMJ. 2015;351:h3867.

5. Miki K, Urta Y. Using serum pepsinogens wisely in a clinical practice. J Dig Dis. 2007;8:8–14.

4. Samloff IM, Varis K, Ihamaki T, Siurala M, Rottger J. Relationships among serum pepsinogen I, serum pepsinogen II, and gastric mucosal histology. A study in relatives of patients with pernicious anemia. Gastroenterology. 1982;83:204–9.

2. Chiang TH, Chiu SY, Chen SL, Yen AM, Fann JC, Liu CY, et al. Serum pepsinogen as a predictor for gastric cancer death: a 16-year community-based cohort study. J Clin Gastroenterol. 2019;53:e186–93.

1. Miki K, Fujishiro M. Cautious comparison between East and West is necessary in terms of the serum pepsinogen test. Dig Endosc. 2009;21:134–5.

30. Malfertheiner P, Megraud F, O’Morain CA, Gisbert JP, Kuipers EJ, Axon AT, et al. Management of Helicobacter pylori infection-the Maastricht V/Florina Consensus Report. Gut. 2011;60:178–83.

29. Syrani K. A panel of serum biomarkers (GastroPanel®) in non-invasive diagnosis of atrophic gastritis. Systematic review and meta-analysis. Anticancer Res. 2016;36:1333–44.

28. Chiang TH, Chuang SL, Chen SL, Chiu HM, Yen AM, Chiu SY, et al. Different in performance of fecal immunochemical tests with the same hemoglobin cutoff concentration in a nationwide colorectal cancer screening program. Gastroenterology. 2014;147:1317–26.

27. Asada K, Nakajima T, Shimazu T, Yamamichi N, Maekita T, Yokoi C, et al. Demonstration of the usefulness of epigenetic cancer risk prediction by a multicentre prospective cohort study. Gut. 2015;64:388–96.

26. Rabeneck L, Lansdorp-Vogelaar I. Assessment of a cancer screening program. Best Pract Res Clin Gastroenterol. 2015;29:979–85.

25. Capelle LG, de Vries AC, Haringsma J, Ter Borg F, de Vries RA, Bruno MJ, et al. The staging of gastritis with the OLGA system by using intestinal metaplasia as an accurate alternative for atrophic gastritis. Gastrointest Endosc. 2010;71:1150–8.

24. Rugge M, Meggio A, Piscalli F, Giacomelli L, De Pretis G, et al. Gastritis staging in clinical practice: the OLGA staging system. Gut. 2007;56:631–6.

23. Bang CS, Lee JJ, Baik GH. Prediction of chronic atrophic gastritis and gastric cancer in a high-risk population. Gastroenterology. 2021;160:1086–98.

22. Chiang TH, Chuang SL, Chen SL, Chiu HM, Yen AM, Chiu SY, et al. Different in performance of fecal immunochemical tests with the same hemoglobin cutoff concentration in a nationwide colorectal cancer screening program. Gastroenterology. 2014;147:1317–26.

21. Chiang TH, Maeda M, Yamada H, Chen CC, Chen SL, Chiu SY, et al. Risk stratification for gastric cancer after Helicobacter pylori eradication: a population-based study on Matsu Islands. J Gastroenterol Hepatol. 2021;36:671–9.

20. van Roon AH, Wiltschut JA, Hol L, van Ballegooijen M, Reijerink JC, van Manen J, et al. Diagnostic yield improves with collection of 2 samples in fecal immunochemical test screening without affecting attendance. Clin Gastroenterol Hepatol. 2011;9:333–9.

19. GastroPanel ELISA. https://www.biohithealthcare.com/en/products/diagnostic-tests/gastropanel/. Accessed 3 Nov 2021.

18. Chiang TH, Maeda M, Yamada H, Chen CC, Chen SL, Chiu SY, et al. Risk stratification for gastric cancer after Helicobacter pylori eradication: a population-based study on Matsu Islands. J Gastroenterol Hepatol. 2021;36:671–9.

17. Chiang TH, Chang WJ, Chen SL, Yen AM, Fann JC, Chiu SY, et al. Mass eradication of Helicobacter pylori to reduce gastric cancer incidence and mortality: a long-term cohort study on Matsu Islands. Gut. 2021;70:243–50.

16. Lee YC, Wu HM, Chen TH, Liu TY, Chiu HM, Chang CC, et al. A community-based study of Helicobacter pylori therapy using the strategy of test, treat, retest, and re-treat initial treatment failures. Helicobacter. 2006;11:418–24.

15. Lee YC, Wu HM, Chen TH, Liu TY, Chiu HM, Chang CC, et al. Sequential therapy for 10 days versus triple therapy for 14 days in the eradication of Helicobacter pylori in the community and hospital populations: a randomised trial. Gut. 2016;65:1784–92.

14. Chiang TH, Maeda M, Yamada H, Chen CC, Chen SL, Chiu SY, et al. Risk stratification for gastric cancer after Helicobacter pylori eradication: a population-based study on Matsu Islands. J Gastroenterol Hepatol. 2021;36:671–9.

13. Leja M, Camargo MC, Polaka L, Isejevs S, Liepniece-Karele I, Janciauskas D, et al. Detection of gastric atrophy by circulating pepsinogens: a comparison of three assays. Helicobacter. 2017;22:e12393.

12. Cho EJ, Kim HK, Jeong TD, Ko DH, Bae SE, Lee JS, et al. Method evaluation of pepsinogen I/II assay based on chemiluminescent immunoassays and comparison with other test methods. Clin Chim Acta. 2016;452:149–54.

11. Leja M, Camargo MC, Polaka L, Isejevs S, Liepniece-Karele I, Janciauskas D, et al. Detection of gastric atrophy by circulating pepsinogens: a comparison of three assays. Helicobacter. 2017;22:e12393.

10. Chiang TH, Chiu SY, Chen SL, Yen AM, Fann JC, Liu CY, et al. Serum pepsinogen as a predictor for gastric cancer death: a 16-year community-based cohort study. J Clin Gastroenterol. 2019;53:e186–93.

9. Samloff IM, Varis K, Ihamaki T, Siurala M, Rottger J. Relationships among serum pepsinogen I, serum pepsinogen II, and gastric mucosal histology. A study in relatives of patients with pernicious anemia. Gastroenterology. 1982;83:204–9.

8. Miki K, Urita Y. Using serum pepsinogens wisely in a clinical practice. J Dig Dis. 2007;8:8–14.

7. Miki K, Urta Y. Using serum pepsinogens wisely in a clinical practice. J Dig Dis. 2007;8:8–14.

6. Song H, Ekheden KG, Zheng Z, Ericsson J, Nyren O, Ye W. Incidence of gastric cancer among patients with gastric precancerous lesions: observational cohort study in a low risk Western population. BMJ. 2015;351:h3867.

5. Miki K, Urta Y. Using serum pepsinogens wisely in a clinical practice. J Dig Dis. 2007;8:8–14.

4. Samloff IM, Varis K, Ihamaki T, Siurala M, Rottger J. Relationships among serum pepsinogen I, serum pepsinogen II, and gastric mucosal histology. A study in relatives of patients with pernicious anemia. Gastroenterology. 1982;83:204–9.

3. Bang CS, Lee JJ, Baik GH. Prediction of chronic atrophic gastritis and gastric neoplasms by serum pepsinogen assay: a systematic review and meta-analysis of diagnostic test accuracy. J Clin Med. 2019;8:657.

2. Chiang TH, Maeda M, Yamada H, Chen CC, Chen SL, Chiu SY, et al. Risk stratification for gastric cancer after Helicobacter pylori eradication: a population-based study on Matsu Islands. J Gastroenterol Hepatol. 2021;36:671–9.