Review Article



Status of Endoscopic Screening Strategies for Upper Gastrointestinal Tract Cancer



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Abstract

Upper gastrointestinal tract cancer (UGIC), which includes both gastric and esophageal cancer, is a major threat to human health. Patients in the early stage have a significant chance to obtain a better prognosis, when compared to patients in the advanced stage. Improving the detection rate of early UGIC is important to improve the survival rate and prognosis. The endoscopic screening of UGIC includes opportunistic and population-based screening, and this has been carried out in a few regions. Compared to these two gastroscopy screening strategies, the early detection ability of opportunistic screening is no less than that for population-based screening, and the compliance of population-based participation is better. Considering economic factors, bundled opportunistic gastroscopy screening is cost-effective. Overall, the screening strategy for UGIC is limited by economic, medical and geographical factors, and the prospect of opportunistic screening is considerable.

Introduction

Upper gastrointestinal cancer (UGIC) mainly includes gastric cancer and esophageal cancer. In the 2018 global cancer statistics, the incidence of gastric cancer (7.2%) and esophageal cancer (4.2%) accounted for 11.4% of globally new cancer cases, with a mortality rate of 16.1% (9.5% for gastric cancer and 6.6% for esophageal cancer).¹ The epidemiology of gastric cancer suggests that East Asia accounts for more than 60% of worldwide new cases,² while another phenomenon is that the age-standardized morbidity and mortality of gastric cancer are declining worldwide. This may be correlated to the reduction in *Helicobacter pylori* (*Hp*) infection.^{3,4} However, the incidence and burden of cases of gastric cancer continue to rise due to the aging population.⁵ Therefore, an effective screening strategy for UGIC is urgently needed.

The development of gastric and esophageal cancer may take decades before the emergence of clinical symptoms.^{6,7} This window makes it possible for the detection and treatment of early UGIC. Screening is an important method to improve its early diagnosis. For example, upper gastrointestinal barium radiography, which is an anti-cancer screening item, is acceptable and simple. Furthermore, serological screening, such as the ABCD serum screening strategy, can identify a part of the high-risk population of gastric cancer.^{8–10} At the same time, it is also necessary to understand that the serum antibody of Helicobacter pylori (Hp) cannot reflect the damage to the gastric mucosa caused by Hp, even if the result is negative. Moreover, various screening strategies, such as swallowpull balloon cytological examination,11 cytology combined with biomarker screening,¹² and expiratory markers,¹³ have been used in clinic to detect early esophageal cancer, since these improves the method of screening. At present, gastroscopy is the most important part of the screening and treatment of early UGIC. The present study reviews the present strategies for gastroscopy screening.

Development of gastroscopy screening strategies

Gastroscopy screening for UGIC was derived from the cancer screening program of high-risk regions, which aims to identify high-risk populations through different screening methods, including radiography and serological testing, in order to increase the detection rate, and improve the curative dissection of UGIC.

Past screening strategies

In the 1960s, a population-based gastric cancer screening program

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Keywords: Cancer screening; Gastroscopy; Upper Gastrointestinal tract cancer. Abbreviations: AFE, auto-fluorescence endoscopy; aOR, adjusted odds ratio; ASR, age-standardized rate; BLI-bright, blue laser imaging-bright; CAG, chronic atrophic gastritis; CI, confidence interval; EGC, early gastric cancer; ESCC, esophageal squamous cell carcinomas; GC, gastric cancer; GIM, gastric intestinal metaplasia; H&N, head and neck; HGIN, high-grade intraepithelial neoplasia; *Hp, Helicobacter pylori*; HR, hazard ratio; ICER, cost-effectiveness ratios; NBI, narrow-band imaging; NCSP, national cancer screening program; OR, odds ratio; QALY, quality-adjusted life-year; RR, relative risk; UGIC, upper gastrointestinal tract cancer; UGIS, upper gastrointestinal series; WLI, white light imaging.

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was initiated in Japan, and barium radiography was initially used as the main method. Then, gastroscopy was used in 1983, and the participation increased annually. This became the primary screening item in 2016, with a participation rate of nearly 90%.⁸ In South Korea, X-ray and gastroscopy examinations have been performed in the National Cancer Screening Program (NCSP) for the population-based screening of UGIC biennially, since 1999.¹⁴ The 10 years of screening data revealed that the number of people who underwent gastroscopy screening increased, accounting for 84.5% in 2016.¹⁵ From 2005 to 2014, China successively carried out gastric cancer population-based screening programs in rural areas (263 project sites), the Huaihe River Basin (32 project sites), and cities (66 cities).¹⁶ Opportunistic gastroscopy screening is a supplementary part of the population-based gastroscopy screening program, and this works in regions where population-based screening is not available.

Benefits of gastroscopy screening

Endoscopy has become the first choice for UGIC screening in Japan and South Korea. The UGIC screening program has achieved remarkable results in high-risk areas, leading to a 40% decrease in mortality.^{17,18} A meta-analysis of 342,013 subjects revealed a reduction in gastric cancer mortality in Asia.¹⁹ For the screening process in Korea, gastroscopy exhibited a stronger ability to identify (adjusted odds ratio [aOR] = 2.10, 95% confidence interval [CI] = 1.90–2.33) local gastric cancer, when compared to the indirect upper gastrointestinal series (UGIS),²⁰ and this reduced the mortality for esophageal cancer (hazard ratio [HR] = 0.497, 95% CI = 0.464-0.531)²¹ and gastric cancer (OR = 0.53, 95% CI = 0.51-0.56, in South Korea; OR = 0.695, 95% CI = 0.489-0.986, in Japan).^{22,23} The regional screening results in China also revealed that endoscopic screening reduced the mortality for esophageal cancer by 37%, and gastric cancer by 33%.²⁴ In addition, compared to upper gastrointestinal barium radiography, endoscopy has no ray exposure, and the cost-effectiveness of gastroscopy is seven times higher.²⁵

Accuracy of gastroscopy screening

Gastroscopy allows for the direct observation of the esophagus and gastric mucosa, as well as the morphology of vessels, and this can also make depression lesions more prominent using simple staining and biopsy techniques, which are conducive for detecting early cancer. A study reported that the sensitivity of gastroscopy to upper gastrointestinal malignancies ranges within 69.0-95.4%, and that the specificity for endoscopy screening and UGIS was 96.0% and 96.1%, respectively.²⁶ In China, the strategy of spraying Lugo iodine under gastroscopy to identify the unstained areas of precancerous lesions can reduce the morbidity, and increase the detection rate of early esophageal cancer.²⁷ In another study, the sensitivity of endoscopic screening for localized gastric cancer was 65.7%, which was significantly higher than that for UGIS.²⁸ Therefore, both opportunistic and population-based gastric cancer screening should include esophageal cancer screening, in order to provide additional benefits.

Developed techniques based on gastroscopy, such as narrowband imaging (NBI), have led to new progress in the detection of early UGIC.²⁹ For the detection of esophageal squamous cell carcinomas (ESCCs) in the head and neck (H&N) region, NBI has a sensitivity of 100% and 97.2%, respectively, and an accuracy was 86.7% and 88.9%, respectively. These results were significantly better, when compared to white light imaging (WLI) (p < 0.001).³⁰ For the detection of early gastric cancer, previous studies and meta-analyses have revealed that NBI is superior to WLI, and that the sensitivity, specificity, positive predictive value, negative predictive value, and accuracy for diagnosing early gastric cancer was 87.2%, 98.6%, 82.1%, 99.0% and 97.8%, respectively,³¹ with a diagnostic odds ratio of 102.75 (95% CI = 48.14-219.32).³²

For auto-fluorescence endoscopy (AFE) and blue laser imaging-bright (BLI-bright). Prospective double-blinded studies have revealed that the clinical value of AFE remains limited. Furthermore, although this has an advantage in detecting elevated gastric musical neoplasias, the sensitivity (74% vs. 64%, p > 0.05) and specificity (83% vs. 40%, p = 0.0003) are inferior to those for WLI.³³ Furthermore, for blue laser imaging-bright (BLI-bright) is superior to WLI, in terms of the real-time detection of gastric cancer (93.1% vs. 50.0%, p = 0.001), especially for lesions in the lower third of the stomach, depressed-type lesions, and small lesions of <10 mm.³⁴

Other gastroscopy screening methods

Magnetic-controlled capsule endoscopy screening is an emerging endoscopic technology, which can be used as a supplement to the intolerance of traditional endoscopy. Its accuracy has been verified in some studies, but the examination cost is high, and its promotion value needs further economic research.^{35,36}

Comparison of population-based and opportunistic screening

The purpose of the screening is to expand the population-accepted screening and improve the detection rate of early UGIC. However, a full-coverage gastroscopy screening would bring huge medical pressure. Hamashima *et al.* estimated the potential of endoscopic screening population was replaced with gastroscopy, the amount of gastroscopy increased by 8.6% of the current.³⁷ Furthermore, recent economic assessments have revealed that population-based endoscopic screening in high-risk areas is cost-effective.³⁸ However, this remains difficult to apply in low- and medium-risk areas, because the incremental cost-effectiveness ratio of screening remains high.³⁹

The population-based gastroscopy screening system has not been estimated in most regions in the world. Thus, opportunistic screening occupies the main position. The present study compared the object, effect, compliance and cost-effectiveness of two screening methods.

Screening object

Population-based screening is primarily differentiated from opportunistic screening, in which invitations to target populations are issued from population registers. In population-based gastroscopy screening, the objects are included in national or local cancer screening programs. The policymaker decides the screening intervals and target population, constructs the re-examination system, and provides the budget, such as South Korea's National Cancer Screening Program (NCSP).

Meanwhile, the objects for the opportunistic gastroscopy screening were individuals who were advised to undergo gastroscopy in the course of other medical services, and those who underwent an endoscopy that was paid by themselves, excluding populations supported by national programs. The screening interval was inconstant, and was based on the awareness of cancer prevention of patients, and the whole examination cost was shouldered by themselves. According to the budget, there are two types of opportunistic screening: individual and collective opportunistic screening. These are applied in Japan.

	Risk level	Strategy	GC / Total	EGC / GC
Kim B ⁴³	High-risk	Population-based gastroscopy screening	100/34,416 (0.3%)	74/100 (74%)
		Opportunistic screening	26/11,238 (0.2%)	14/26 (53.8%)
Lau J ⁴⁴	Middle-risk	Opportunistic screening	5/1,414 (0.35%)	3/5 (60%)
Zhou Q ⁴⁵	High-risk	Opportunistic screening	190/6,701 (2.84%)	88/190 (46.3%)

Table 1. Detection ability of gastric cancer between the two strategies

EGC: Early Gastric Cancer, GC, Gastric Cancer.

Screening effect

There is uncertainty in the ability to prevent and decrease the mortality of UGIC between these two different endoscopic screening strategies. The effect of the population-based gastroscopy screening strategy for detecting UGIC has been verified in various countries. At present, the detection rate by gastroscopy for early gastric cancer in Japan is 75.1%, and the 5-year survival rate of gastric cancer is 64.6%.40 Furthermore, the gastroscopy detection rate for early gastric cancer in Korea is 63.9%, and the 5-year survival rate is 74.6%.⁴¹ The population-based gastroscopy strategy aims to improve the early diagnosis and treatment of UGIC, which has been carried out in 194 high-risk areas for gastric cancer in China since 2005. According to a 10-year statistical analysis of six highrisk areas, it was found that compared to areas without gastroscopy screening, the incidence and mortality of UGIC decreased by 23% (relative risk [RR] = 0.77, 95% CI = 0.74–0.81) and 57% (RR = 0.43, 95% CI = 0.40-0.47) in the gastroscopy screening group, and decreased by 14% (RR = 0.86, 95% CI = 0.84-0.89) and 31% (RR = 0.69, 95% CI = 0.66-0.72) for all participants.^{24,42}

Kim *et al.*⁴³ compared the population-based endoscopic screening group (n = 34,416) and opportunistic screening group (n = 11,238), and revealed that the detection rate for gastric cancer using these two screening strategies was 0.3% and 0.2%, respectively, but there was no significant difference (p = 0.299). Another study revealed that in middle-risk areas for gastric cancer, the detection rate for UGIC in asymptomatic patients by opportunistic endoscopy is 0.35%.⁴⁴ However, the proportion for early gastric cancer was significantly higher when the population-based gastroscopy screening strategy was applied, when compared to opportunistic gastroscopy screening (74.5% vs. 53.8%, p = 0.046).⁴³ This means that population-based endoscopy screening is superior to opportunistic screening, in terms of detecting early gastric cancer. The detection rate of cancer in some of the reviewed studies are presented in Table 1.⁴³⁻⁴⁵

Screening compliance

Compliance is the basis of screening surveillance. The NCSP data revealed that 67% of participants chose gastroscopy for screening in the future two years, and 47.8% of participants who accepted UGIS chose gastroscopy in the future. Furthermore, participants with a family history of gastric cancer had better compliance with gastroscopy (aOR = 2.05, 95% CI = 1.17-3.60).⁴⁶ A latest research in China revealed that the compliance rate for gastroscopy screening in urban and rural areas is 45.2% and 48.4%, respectively, and the demographic characteristics differed, or even reversed, in urban and rural populations with good compliance.⁴⁷

Among these two different screening strategies, people who underwent opportunistic gastroscopy screening had better compliance. A large sample study conducted in South Korea revealed that under the definition of regular gastroscopy for at least once every two years, people who underwent opportunistic gastroscopy had better compliance (27.1% vs. 19.1%, p < 0.001). The study also suggested that compared to screening methods, the compliance of the screening population is more important.⁴³ Combined with the study conducted by Jun,²² it was found that the death toll for gastric cancer obviously decreased with the increase in frequency of gastroscopy. Furthermore, it was observed that opportunistic gastroscopy has more advantages in screening populations that require multiple gastroscopy examinations, such as patients with chronic gastric mucosal atrophy or post-endoscopic treatment of early UGIC.

Screening cost-effectiveness

Zhou *et al.* investigated the cost of gastric cancer screening in high-risk populations (n = 27,970). They found that the total cost for gastric cancer in a screened population and in an unscreened group was \$4,041.10 and \$4,228.00, respectively. If 68.9% (20/29) of advanced gastric cancer patients are diagnosed at the early stage by screening, merely an extra of \$1,020.00 per screening would needed, providing economic and social benefits.⁴⁸ Furthermore, a study conducted for opportunistic gastroscopy screening in middle-risk areas (gastric cancer age-standardized rate [ASR] of 8.2/100,000.0) revealed that the detection rate for UGIC or precancerous lesions was 12.7%, and the cost of each case was \$3,960.00, and when the detection rate for UGIC was 0.35%, the cost of each case was \$141,400.00.⁴⁴

For people who underwent opportunistic gastroscopy, the bundled endoscopy strategy was found to be more cost-effective. Gupta et al. conducted a Markov model analysis, and the result suggested that gastroscopy bundled with colonoscopy can reduce the number of gastrointestinal cancer-related deaths by 61.1/100,000.0, which is one-third more than that for single gastroscopy or colonoscopy, and the incremental cost-effectiveness ratio (ICER) per qualityadjusted life-year (QALY) was \$95,559.00, which is lower than that for a single endoscopy examination (\$115,664.00).³⁹ However, there is a prerequisite, according to the study conducted by Areia.49 In middle-risk areas for gastric cancer (ASR = 13.1/100,000.0), the combined screening strategy for detecting UGIC would be cost-effective only when patients plan to undergo colonoscopy, and an additional gastroscopy would only cost €60.00. The study also suggested that a single gastroscopy would cost €137.00, and that it would be cost-effective to conduct independent gastroscopy screenings every five years, in which only the ASR for gastric cancer would be over 25/100,000. In summary, the average detection cost of UGIC is the lowest in opportunistic gastroscopy bundled with a planned colonoscopy, and performing an opportunistic gastroscopy every five years in high-risk areas would be cost-effective.

The endoscopic surveillance of the esophageal adenocarcinoma precancerous status, Barrett's esophagus, is cost-effective.⁵⁰ However, its large-scale endoscopic screening would not be costeffective.⁵¹ In high-risk regions, performing a gastroscopy every 1–3 years as a screening strategy for individuals at the age of 40 would be cost-effective.⁵² In the general population of over 50

	Strategy	Country	Risk level	Object and aim disease	Threshold to pay	Advise
Gupta N ³⁹	Opportunistic gastroscopy bundled with colonoscopy	Various	Various	50 years old, UGIC	\$50,000.00	For GC/ESCC/EAC, not cost-effective
Inadomi J ⁵⁰	Review	Various	Various	50 years old, BE	\$100,000.00	Cost-effective
Xia R ³⁸	Program screening	China	High-risk area	40–69 years old, UGIC	\$30,828.00	Gastroscopy every two years is cost-effective
Wu B ⁵²	Program screening	China	High-risk area	Over 40 years old, ESCC	\$1,151.00	Gastroscopy every 1–3 years is cost-effective
Shah SC ⁵⁴	Strategy 1: Opportunistic gastroscopy bundled with colonoscopy every three years, when GIM is identified; Strategy 2: Biennial opportunistic gastroscopy bundled with colonoscopy; Strategy 3: No gastroscopy	United States	Low-risk (Asian- American)	50 years old, GC	\$100,000.00	Strategy 1 is cost- effective (\$75,959.00– 74,329.00/QALY)
Saumoy M ⁵⁵	Strategy 1: Opportunistic gastroscopy bundled with colonoscopy every three years, when GIM is identified; Strategy 2: Biennial opportunistic gastroscopy bundled with colonoscopy; Strategy 3: No gastroscopy	United States	Low-risk area	50 years old, GC	\$100,000.00	Strategy 1 is cost- effective for non- Hispanic black (\$80,278.00/QALY), Hispanic (\$76,070.00/ QALY), and Asians (\$71,451.00/QALY), but not for non- Hispanic white (\$122,428.00/QALY)
Kowada A ⁵³	Biennial screening for mild-moderate CAG, annual for severe CAG	Japan	High-risk area	GC after <i>Hp</i> eradication	\$100,000.00	Cost-effective
Lau J ⁴⁴	Opportunistic screening	Singapore	Middle- risk area	UGIC+HGIN	Not available	\$3,950.00/lesion
Areia M ⁴⁹	Strategy 1: Upper endoscopy only every five years; Strategy 2: Opportunistic gastroscopy bundled with colonoscopy every 5–10 years; Strategy 3: Gastroscopy after biennial serology screening positive.	Portugal	Middle- risk area	GC	€37,000.00	Strategy 1 is cost- effective when the risk is over 25/100,000; Strategy 2 is cost- effective when the risk is over 10/100,000

CAG, Chronic Atrophic Gastritis; EAC, Esophageal adenocarcinoma; ESCC, Esophageal squamous cell carcinoma; GC, Gastric Cancer; GIM, Gastric Intestinal Metaplasia; HGIN, Highgrade Intraepithelial Neoplasia.; QALY, Quality-Adjusted Life Year; UGIC, Upper Gastrointestinal Cancer.

years old, the ICER for the screening and surveillance of Barrett's esophagus, combined with the screening strategy, was \$95,559.00/ QALY, indicating high cost.³⁹ For gastric mucosal atrophy after the eradication of *Hp*, biennial gastroscopy examination for patients with mild-to-moderate gastric mucosal atrophy and annual gastroscopy surveillance for patients with severe gastric mucosal atrophy are the most cost-effective, with a willingness-to-pay threshold of \$100,000.00/QALY.⁵³ The cost-effectiveness of this strategy is also correlated to race. A study conducted in the United States revealed that gastroscopy screening and surveillance are cost-effective for Asians elder than 50 years old, non-Hispanic blacks, and Hispanics, and the ICER per QALY was \$71,451.00/ QALY, \$80,278.00/QALY and \$76,070.00/QALY, respectively (willingness-to-pay level = \$100,000.00/QALY).^{54,55} The cost-effectiveness details of the lectures are presented in Table 2.^{38,39,44,49,50,52-55} It should be noted that the willingness-to-pay threshold was assumed to be the per capita gross domestic product, and that the indigenization of economic effect was necessary.^{56,57}

Summary

Compared to opportunistic gastroscopy, there was no significant difference in the detection rate for gastric cancer in the populationbased gastroscopy screening strategy, but the proportion of early gastric cancer detected using the population-based gastroscopy screening strategy was significantly higher. The advantage of opportunistic gastroscopy is that the compliance of participants is better, and asymptomatic people would benefit more from this strategy. Opportunistic gastroscopy bundled with planned colonoscopy can lead to higher benefits in UGIC screening. Similarly, if the screening for esophageal cancer is incidentally carried out during the gastric cancer screening, additional benefits can be obtained.

Prospect of opportunistic screening

It remains difficult to carry out a broad population-based screening program for UGIC, which is limited by the number of endoscopic physicians, the risk of UGIC, and the cost-effectiveness of screening. Furthermore, the mortality rate of cancer is positively correlated with the proportion of advanced cancer.⁵⁸ Moreover, the medical cost of hospitalized patients with UGIC is rapidly increasing, and the average cost has reached ¥ 17,567.0 by 2016. In order to reduce the mortality for UGIC, there is a need to improve the detection rate of cancer in the early stage, regardless of the strategy used.

For areas without a population-based screening program, opportunistic gastroscopy would be an important channel for people to receive screening. Studies conducted in Japan and South Korea suggested that people over 40 years old should perform gastroscopy surveillance every two years. This is suggested for high-risk populations, such as first-degree relatives of UGIC, *Hp* infection, high salt diet, and smokers.^{59,60} Opportunistic gastroscopy screening for UGIC can be performed in low- and moderate-risk areas, and this can be combined with colorectal cancer screening by colonoscopy.

Conclusions

Opportunistic gastroscopy is important for the early detection and treatment of UGIC. Based on the collected data, it was considered that opportunistic gastroscopy screening can be bundled with colonoscopy as a strategy to optimize its cost-effectiveness in lowand moderate-risk areas and populations. In high-risk areas for UGIC, population-based gastroscopy screening can be used as the main approach. However, the evaluation of monitoring intervals and cost-effectiveness ratios should be localized.

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Conflict of interest

Prof. Bin Lyu has been an editorial board member of Cancer Screening and Prevention since February 2022. The authors have no other conflict of interests related to this publication.

Author contributions

BL contributed to study concept and design, XLJ performed the acquisition of data for the references, XLJ drafted the manuscript, BL performed the critical revision of the manuscript, and BL supervised the study.

References

[1] Bray F, Ferlay J, Soerjomataram I, Siegel RL, Torre LA, Jemal A. Global

cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. CA Cancer J Clin 2018;68(6):394–424. doi:10.3322/caac.21492.

- [2] Sekiguchi M, Oda I, Matsuda T, Saito Y. Epidemiological Trends and Future Perspectives of Gastric Cancer in Eastern Asia. Digestion 2022;103(1):22–28. doi:10.1159/000518483, PMID:34515086.
- [3] Choi Y, Gwack J, Kim Y, Bae J, Jun JK, Ko KP, et al. Long term trends and the future gastric cancer mortality in Korea: 19832013. Cancer Res Treat 2006;38(1):7–12. doi:10.4143/crt.2006.38.1.7, PMID:1977 1252.
- [4] 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(2):243–250. doi:10.1136/gutjnl-2020-322200, PMID:32792335.
- [5] GBD 2017 Stomach Cancer Collaborators. The global, regional, and national burden of stomach cancer in 195 countries, 1990-2017: a systematic analysis for the Global Burden of Disease study 2017. Lancet Gastroenterol Hepatol 2020;5(1):42–54. doi:10.1016/S2468-1253(19)30328-0, PMID:31648970.
- [6] Correa P. Gastric cancer: overview. Gastroenterol Clin North Am 2013;
 42(2):211–217. doi:10.1016/j.gtc.2013.01.002, PMID:23639637.
- [7] Wang JW, Guan CT, Wang LL, Chang LY, Hao CQ, Li BY, et al. Natural History Analysis of 101 Severe Dysplasia and Esophageal Carcinoma Cases by Endoscopy. Gastroenterol Res Pract 2017;2017:9612854. doi:10.1155/2017/9612854, PMID:28465681.
- [8] Hamashima C, Systematic Review Group and Guideline Development Group for Gastric Cancer Screening Guidelines. Update version of the Japanese Guidelines for Gastric Cancer Screening. Jpn J Clin Oncol 2018;48(7):673–683. doi:10.1093/jjco/hyy077, PMID:29889263.
- [9] Park CH, Kim EH, Jung DH, Chung H, Park JC, Shin SK, *et al.* The new modified ABCD method for gastric neoplasm screening. Gastric Cancer 2016;19(1):128–135. doi:10.1007/s10120-015-0473-4, PMID:256 63259.
- [10] Tu H, Sun L, Dong X, Gong Y, Xu Q, Jing J, et al. A Serological Biopsy Using Five Stomach-Specific Circulating Biomarkers for Gastric Cancer Risk Assessment: A Multi-Phase Study. Am J Gastroenterol 2017;112(5):704–715. doi:10.1038/ajg.2017.55, PMID:28323271.
- [11] Patel AA, Strome M, Blitzer A. Directed balloon cytology of the esophagus: A novel device for obtaining circumferential cytologic sampling. Laryngoscope 2017;127(5):1032–1035. doi:10.1002/lary. 26472, PMID:28092117.
- [12] Roshandel G, Merat S, Sotoudeh M, Khoshnia M, Poustchi H, Lao-Sirieix P, et al. Pilot study of cytological testing for oesophageal squamous cell dysplasia in a high-risk area in Northern Iran. Br J Cancer 2014; 111(12):2235–2241. doi:10.1038/bjc.2014.506, PMID:25247319.
- [13] Zou X, Zhou W, Lu Y, Shen C, Hu Z, Wang H, et al. Exhaled gases online measurements for esophageal cancer patients and healthy people by proton transfer reaction mass spectrometry. J Gastroenterol Hepatol 2016;31(11):1837–1843. doi:10.1111/jgh.13380, PMID:26996099.
- [14] Goto R, Hamashima C, Mun S, Lee WC. Why screening rates vary between Korea and Japan—differences between two national healthcare systems. Asian Pac J Cancer Prev 2015;16(2):395–400. doi:10.7314/apjcp.2015.16.2.395, PMID:25684461.
- [15] Ryu JE, Choi E, Lee K, Jun JK, Suh M, Jung KW, et al. Trends in the Performance of the Korean National Cancer Screening Program for Gastric Cancer from 2007 to 2016. Cancer Res Treat 2022;54(3):842– 849. doi:10.4143/crt.2021.482, PMID:34607395.
- [16] Zhu J, Wang SM, Chen R, Li XQ, Wei WW. [Progress on screening for gastric cancer]. Zhonghua Zhong Liu Za Zhi 2020;42(7):603–608. doi:10.3760/cma.j.cn112152-20191125-00759, PMID:32842452.
- [17] Rosero-Bixby L, Sierra R. X-ray screening seems to reduce gastric cancer mortality by half in a community-controlled trial in Costa Rica. Br J Cancer 2007;97(7):837–843. doi:10.1038/sj.bjc.6603729, PMID:17912238.
- [18] Lee KJ, Inoue M, Otani T, Iwasaki M, Sasazuki S, Tsugane S, et al. Gastric cancer screening and subsequent risk of gastric cancer: a largescale population-based cohort study, with a 13-year follow-up in Japan. Int J Cancer 2006;118(9):2315–2321. doi:10.1002/ijc.21664, PMID:16331632.
- [19] Zhang X, Li M, Chen S, Hu J, Guo Q, Liu R, et al. Endoscopic Screening

in Asian Countries Is Associated With Reduced Gastric Cancer Mortality: A Meta-analysis and Systematic Review. Gastroenterology 2018;155(2):347–354.e9. doi:10.1053/j.gastro.2018.04.026, PMID: 29723507.

- [20] Choi KS, Jun JK, Suh M, Park B, Noh DK, Song SH, et al. Effect of endoscopy screening on stage at gastric cancer diagnosis: results of the National Cancer Screening Programme in Korea. Br J Cancer 2015;112(3):608–612. doi:10.1038/bjc.2014.608, PMID:25490528.
- [21] Kim JH, Han KD, Lee JK, Kim HS, Cha JM, Park S, et al. Association between the National Cancer Screening Programme (NSCP) for gastric cancer and oesophageal cancer mortality. Br J Cancer 2020;123(3):480–486. doi:10.1038/s41416-020-0883-x, PMID:3239 8860.
- [22] Jun JK, Choi KS, Lee HY, Suh M, Park B, Song SH, et al. Effectiveness of the Korean National Cancer Screening Program in Reducing Gastric Cancer Mortality. Gastroenterology 2017;152(6):1319–1328.e7. doi:10.1053/j.gastro.2017.01.029, PMID:28147224.
- [23] Hamashima C, Ogoshi K, Okamoto M, Shabana M, Kishimoto T, Fukao A. A community-based, case-control study evaluating mortality reduction from gastric cancer by endoscopic screening in Japan. PLoS One 2013;8(11):e79088. doi:10.1371/journal.pone.0079088, PMID:24236091.
- [24] Wang GQ, Wei WW. A new transition of the screening, early diagnosis and early treatment project of the upper gastrointestinal cancer: opportunistic screening. Zhonghua Yu Fang Yi Xue Za Zhi 2019;53(11):1084–1087.doi:10.3760/cma.j.issn.0253-9624.2019.11. 002, PMID:31683391.
- [25] Tashiro A, Sano M, Kinameri K, Fujita K, Takeuchi Y. Comparing mass screening techniques for gastric cancer in Japan. World J Gastroenterol 2006;12(30):4873–4874. doi:10.3748/wjg.v12.i30.4873, PMID:1693 7471.
- [26] Hamashima C, Okamoto M, Shabana M, Osaki Y, Kishimoto T. Sensitivity of endoscopic screening for gastric cancer by the incidence method. Int J Cancer 2013;133(3):653–659. doi:10.1002/ijc.28065, PMID:23364866.
- [27] Ding W, Tong S, Gou Y, Sun C, Wang H, Chen Z, et al. Human epidermal growth factor receptor 2: a significant indicator for predicting progression in non-muscle-invasive bladder cancer especially in high-risk groups. World J Urol 2015;33(12):1951–1957. doi:10.1007/ s00345-015-1557-9, PMID:25894367.
- [28] Choi KS, Jun JK, Park EC, Park S, Jung KW, Han MA, et al. Performance of different gastric cancer screening methods in Korea: a populationbased study. PLoS One 2012;7(11):e50041. doi:10.1371/journal. pone.0050041, PMID:23209638.
- [29] Gono K, Obi T, Yamaguchi M, Ohyama N, Machida H, Sano Y, et al. Appearance of enhanced tissue features in narrow-band endoscopic imaging. J Biomed Opt 2004;9(3):568–577. doi:10.1117/1.1695563, PMID:15189095.
- [30] Muto M, Minashi K, Yano T, Saito Y, Oda I, Nonaka S, et al. Early detection of superficial squamous cell carcinoma in the head and neck region and esophagus by narrow band imaging: a multicenter randomized controlled trial. J Clin Oncol 2010;28(9):1566–1572. doi:10.1200/JCO.2009.25.4680, PMID:20177025.
- [31] Yu H, Yang AM, Lu XH, Zhou WX, Yao F, Fei GJ, et al. Magnifying narrow-band imaging endoscopy is superior in diagnosis of early gastric cancer. World J Gastroenterol 2015;21(30):9156–9162. doi:10.3748/ wjg.v21.i30.9156, PMID:26290643.
- [32] Hu YY, Lian QW, Lin ZH, Zhong J, Xue M, Wang LJ. Diagnostic performance of magnifying narrow-band imaging for early gastric cancer: A meta-analysis. World J Gastroenterol 2015;21(25):7884–7894. doi:10.3748/wjg.v21.i25.7884, PMID:26167089.
- [33] Kato M, Kaise M, Yonezawa J, Yoshida Y, Tajiri H. Autofluorescence endoscopy versus conventional white light endoscopy for the detection of superficial gastric neoplasia: a prospective comparative study. Endoscopy 2007;39(11):937–941. doi:10.1055/s-2007-966857, PMID: 18008201.
- [34] Dohi O, Yagi N, Naito Y, Fukui A, Gen Y, Iwai N, *et al*. Blue laser imagingbright improves the real-time detection rate of early gastric cancer: a randomized controlled study. Gastrointest Endosc 2019;89(1):47–57. doi:10.1016/j.gie.2018.08.049, PMID:30189197.
- [35] Zou WB, Hou XH, Xin L, Liu J, Bo LM, Yu GY, et al. Magnetic-controlled

capsule endoscopy vs. gastroscopy for gastric diseases: a two-center self-controlled comparative trial. Endoscopy 2015;47(6):525–528. doi:10.1055/s-0034-1391123, PMID:25590177.

- [36] Xiao YF, Wu ZX, He S, Zhou YY, Zhao YB, He JL, et al. Fully automated magnetically controlled capsule endoscopy for examination of the stomach and small bowel: a prospective, feasibility, two-centre study. Lancet Gastroenterol Hepatol 2021;6(11):914–921. doi:10.1016/ S2468-1253(21)00274-0, PMID:34555347.
- [37] Hamashima C, Goto R. Potential capacity of endoscopic screening for gastric cancer in Japan. Cancer Sci 2017;108(1):101–107. doi:10.1111/cas.13100, PMID:27727490.
- [38] Xia R, Zeng H, Liu W, Xie L, Shen M, Li P, et al. Estimated Cost-effectiveness of Endoscopic Screening for Upper Gastrointestinal Tract Cancer in High-Risk Areas in China. JAMA Netw Open 2021;4(8):e2121403. doi:10.1001/jamanetworkopen.2021.21403, PMID:34402889.
- [39] Gupta N, Bansal A, Wani SB, Gaddam S, Rastogi A, Sharma P. Endoscopy for upper GI cancer screening in the general population: a cost-utility analysis. Gastrointest Endosc 2011;74(3):610–624.e2. doi:10.1016/j.gie.2011.05.001, PMID:21741639.
- [40] Matsuda T, Ajiki W, Marugame T, Ioka A, Tsukuma H, Sobue T, et al. Population-based survival of cancer patients diagnosed between 1993 and 1999 in Japan: a chronological and international comparative study. Jpn J Clin Oncol 2011;41(1):40–51. doi:10.1093/jjco/ hyq167, PMID:20819833.
- [41] Hong S, Won YJ, Park YR, Jung KW, Kong HJ, Lee ES, et al. Cancer Statistics in Korea: Incidence, Mortality, Survival, and Prevalence in 2017. Cancer Res Treat 2020;52(2):335–350. doi:10.4143/crt.2020.206, PMID:32178489.
- [42] Chen R, Liu Y, Song G, Li B, Zhao D, Hua Z, et al. Effectiveness of onetime endoscopic screening programme in prevention of upper gastrointestinal cancer in China: a multicentre population-based cohort study. Gut 2021;70(2):251–260. doi:10.1136/gutjnl-2019-320200, PMID:32241902.
- [43] Kim BJ, Heo C, Kim BK, Kim JY, Kim JG. Effectiveness of gastric cancer screening programs in South Korea: organized vs opportunistic models. World J Gastroenterol 2013;19(5):736–741. doi:10.3748/wjg. v19.i5.736, PMID:23430471.
- [44] Lau JWL, Khoo MJW, Leong XH, Lim TZ, Shabbir A, Yeoh KG, et al. Opportunistic upper endoscopy during colonoscopy as a screening strategy for countries with intermediate gastric cancer risk. J Gastroenterol Hepatol 2021;36(4):1081–1087. doi:10.1111/jgh.15290, PMID:33037826.
- [45] Zhou Q, Chen Y, Pan J, Zhou L, Lin J. Application of a novel scoring system for gastric cancer opportunistic screening in hospital visits. BMC Gastroenterol 2022;22(1):223. doi:10.1186/s12876-022-02315-9, PMID:35527297.
- [46] Choi KS, Kwak MS, Lee HY, Jun JK, Hahm MI, Park EC. Screening for gastric cancer in Korea: population-based preferences for endoscopy versus upper gastrointestinal series. Cancer Epidemiol Biomarkers Prev 2009;18(5):1390–1398. doi:10.1158/1055-9965.EPI-08-0940, PMID:19383892.
- [47] Li H, Cao MM, Sun DQ, He SY, Yan XX, Yang F, et al. [A comparative analysis of the distribution of the high-risk population of upper gastrointestinal cancer and endoscopic screening compliance in two urban areas and two rural areas in China]. Zhonghua Zhong Liu Za Zhi 2022;44(6):531–539. doi:10.3760/cma.j.cn112152-20210916-00707, PMID:35754227.
- [48] Zhou L, Guan P, Sun LP, He QC, Yuan Y, Zhou BS. Health economic assessment for screening of gastric cancer in a high risk population in northeastern china. Chin J Cancer Res 2011;23(1):21–24. doi:10.1007/s11670-011-0021-7, PMID:23467677.
- [49] Areia M, Spaander MC, Kuipers EJ, Dinis-Ribeiro M. Endoscopic screening for gastric cancer: A cost-utility analysis for countries with an intermediate gastric cancer risk. United European Gastroenterol J 2018;6(2):192–202. doi:10.1177/2050640617722902, PMID:2951 1549.
- [50] Inadomi JM, Saxena N. Screening and Surveillance for Barrett's Esophagus: Is It Cost-Effective? Dig Dis Sci 2018;63(8):2094–2104. doi:10.1007/s10620-018-5148-7, PMID:29948571.
- [51] di Pietro M, Canto MI, Fitzgerald RC. Endoscopic Management of Early Adenocarcinoma and Squamous Cell Carcinoma of the Esophagus:

Lyu B. et al: Screening strategies of UGIC

Screening, Diagnosis, and Therapy. Gastroenterology 2018;154(2):421–436. doi:10.1053/j.gastro.2017.07.041, PMID:28778650.

- [52] Wu B, Wang Z, Zhang Q. Age at Initiation and Frequency of Screening to Prevent Esophageal Squamous Cell Carcinoma in High-risk Regions: an Economic Evaluation. Cancer Prev Res (Phila) 2020;13(6):543– 550. doi:10.1158/1940-6207.CAPR-19-0477, PMID:32152149.
- [53] Kowada A. Endoscopy Is Cost-Effective for Gastric Cancer Screening After Successful Helicobacter pylori Eradication. Dig Dis Sci 2021; 66(12):4220–4226. doi:10.1007/s10620-020-06813-2, PMID:33417 196.
- [54] Shah SC, Canakis A, Peek RM Jr, Saumoy M. Endoscopy for Gastric Cancer Screening Is Cost Effective for Asian Americans in the United States. Clin Gastroenterol Hepatol 2020;18(13):3026–3039. doi:10.1016/j.cgh.2020.07.031, PMID:32707341.
- [55] Saumoy M, Schneider Y, Shen N, Kahaleh M, Sharaiha RZ, Shah SC. Cost Effectiveness of Gastric Cancer Screening According to Race and Ethnicity. Gastroenterology 2018;155(3):648–660. doi:10.1053/j. gastro.2018.05.026, PMID:29778607.
- [56] Cameron D, Ubels J, Norström F. On what basis are medical cost-

effectiveness thresholds set? Clashing opinions and an absence of data: a systematic review. Glob Health Action 2018;11(1):1447828. d oi:10.1080/16549716.2018.1447828, PMID:29564962.

- [57] Chi YL, Blecher M, Chalkidou K, Culyer A, Claxton K, Edoka I, et al. What next after GDP-based cost-effectiveness thresholds? Gates Open Res 2020;4:176. doi:10.12688/gatesopenres.13201.1, PMID:33575544.
- [58] Zeng H, Chen W, Zheng R, Zhang S, Ji JS, Zou X, et al. Changing cancer survival in China during 2003-15: a pooled analysis of 17 populationbased cancer registries. Lancet Glob Health 2018;6(5):e555–e567. doi:10.1016/S2214-109X(18)30127-X, PMID:29653628.
- [59] Praud D, Rota M, Pelucchi C, Bertuccio P, Rosso T, Galeone C, et al. Cigarette smoking and gastric cancer in the Stomach Cancer Pooling (StoP) Project. Eur J Cancer Prev 2018;27(2):124–133. doi:10.1097/ CEJ.00000000000290, PMID:27560662.
- [60] Wang XQ, Terry PD, Yan H. Review of salt consumption and stomach cancer risk: epidemiological and biological evidence. World J Gastroenterol 2009;15(18):2204–2213. doi:10.3748/wjg.15.2204, PMID: 19437559.