E-learning system to improve the endoscopic diagnosis of early gastric cancer

Kenshi Y ao, Takashi Y ao, Noriya Uedo, Hisashi Doyama, H ideki Ishikawa, Satoshi Nimura, Yuichi Takahashi

1 Department of Endoscopy, Fukuoka University Chikushi Hospital, Fukuoka; 2 Department of Human Pathology, Juntendo University Graduate School of Medicine, Tokyo; 3 Department of Gastrointestinal Oncology, Osaka International Cancer Institute, Osaka; 4 Department of Gastroenterology, Ishikawa Prefectural Central Hospital, Kanazawa; 5 Department of Molecular-Targeting Cancer Prevention, Graduate School of Medical Science, Kyoto Prefectural University of Medicine, Osaka; 6 Department of Pathology, Fukuoka University Chikushi Hospital, Fukuoka; 7 Department of Development, FAST Inc., Tokyo, Japan

We developed three e-learning systems for endoscopists to acquire the necessary skills to improve the diagnosis of early gastric cancer (EGC) and demonstrated their usefulness using randomized controlled trials. The subjects of the three e-learning systems were “detection,” “characterization,” and “preoperative assessment.” The contents of each e-learning system included “technique,” “knowledge,” and “obtaining experience.” All e-learning systems proved useful for endoscopists to learn how to diagnose EGC. Lecture videos describing “the technique” and “the knowledge” can be beneficial. In addition, repeating 100 self-study cases allows learners to gain “experience” and improve their diagnostic skills further. Web-based e-learning systems have more advantages than other teaching methods because the number of participants is unlimited. Histopathological diagnosis is the gold standard for the diagnosis of gastric cancer. Therefore, we developed a comprehensive diagnostic algorithm to standardize the histopathological diagnosis of gastric cancer. Once we have successfully shown that this algorithm is helpful for the accurate histopathological diagnosis of cancer, we will complete a series of e-learning systems designed to assess EGC accurately.

Keywords: Endoscopy; Learning; Stomach neoplasms

INTRODUCTION

Gastric cancer is reported to have the fourth highest mortality rate in the world.1 The stage of gastric cancer at diagnosis is often advanced, except in Japan and Korea. Therefore, the overall 5-year survival rate is <10%.1 However, the prognosis improves in patients diagnosed at an early stage.5 In addition, endoscopic treatment of early gastric cancer (EGC) ensures a patient's quality of life.3 Hence, early diagnosis of gastric cancer is imperative. Although esophagogastroduodenoscopy (EGD) is useful for detecting EGC,1 endoscopists have difficulty learning the “technique,” gaining “knowledge,” and “obtaining experience.” These three topics are essential to enable endoscopists to detect EGC using EGD.1 To address these difficulties, we developed three types of web-based e-learning systems, tested each through randomized controlled trials (RCTs), and demonstrated their usefulness in ensuring correct and precise endoscopic diagnosis of EGC.5,7 In this review, we introduce the basic diagnostic systems employed in e-learning, the outcomes of RCTs testing the usefulness of these three e-learning systems, and our future projects.
THE PRINCIPLES FOR DEVELOPING E-LEARNING SYSTEMS FOR THE ENDOSCOPIC ASSESSMENT OF EGC

The core principles of the e-learning system are that endoscopists should acquire “technique,” “knowledge,” and “experience” for the endoscopic diagnosis of EGC. Therefore, we constructed the contents for all e-learning systems imparting “technique,” “knowledge,” and “experience.”

To teach “technique” and “knowledge,” we employed video lectures. The alternative, “acquiring experience” by e-learning, is challenging. Therefore, we speculated that repeated self-study with 100 consecutive cases (100 cases of repeated self-study) could be useful for accumulating experience similar to that employed in machine learning (Fig. 1).

BASIC DIAGNOSTIC SYSTEMS AND THEIR PERFORMANCE IN “DETECTION,” “CHARACTERIZATION,” AND “PREOPERATIVE ASSESSMENT” EMPLOYED IN THE E-LEARNING SYSTEMS

Detection using conventional white-light imaging

We constructed a color plus surface classification system (CSCS) to characterize mucosal lesions detected using conventional white-light imaging (C-WLI) alone. The criteria according to the CSCS were as follows: (1) presence of a well-demarcated lesion with irregularity in color, and (2) presence of a well-demarcated lesion with an irregular surface. If the target met either criterion, the diagnosis was “cancerous.” When both were absent, the diagnosis was “noncancerous” (Fig. 2).

We reported that when we applied these criteria to clinical practice, the sensitivity was 81.0%, and the specificity was 88.1%. This diagnostic system can help endoscopists improve their skills in the early detection of EGC.

Characterization using magnifying endoscopy with narrow-band imaging

We established a vessel plus surface classification system (VSCS) which has already proven helpful in the accurate assessment of EGC and in delineating EGCs for curative endoscopic resection. The VSCS has been approved through a multi-society consensus as a standardized magnifying endoscopy diagnostic system. The criteria were as follows: (1) presence of an irregular microvascular (MV) pattern with a demarcation line, and (2) presence of an irregular microsurface (MS) pattern with a demarcation line. If the target lesion met either or both criteria, the diagnosis of the lesion was “cancerous.” If both criteria were absent, the diagnosis was “noncancerous.”

We report the diagnostic performance of magnifying endoscopy with narrow-band imaging (M-NBI) with a high-confidence prediction according to the VSCS. The sensitivity was 85.7%, and the specificity was 99.4%. The case determined to be a false-negative was a signet-ring cell carcinoma that was pale in color and did not represent either an irregular MV pattern or an irregular MS pattern without a demarcation line. When we excluded the false-negative case that showed a pale superficial mucosal lesion, the sensitivity increased from 85.7% to 99.4%.

Fig. 1. An example of the self-study cases for the diagnosis of 100 cases. (A) One case comprises a set of three slides. The first slide shows one endoscopic photo where one lesion is present. First, the participant should click to choose whether the lesion is cancerous or noncancerous. (B) Immediately after clicking on their choice, an illustration indicating whether the answer is correct or incorrect appears on the second slide. (C) The third slide indicates brief instructions on characterizing the endoscopic findings to diagnose correctly and shows the original endoscopic image again. Adapted from Yao et al. EBioMedicine 2016;9:140–147, according to the Creative Commons license.
Fig. 2. Example of endoscopic images showing the differences between noncancerous and cancerous lesions according to the color plus surface classification system (CSCS) by conventional white-light imaging. (A) Endoscopic findings of a focal atrophic pale mucosal lesion. According to the CSCS, the lesion is not well demarcated without irregularity in color or surface. Therefore, this lesion is diagnosed as noncancerous. (B) Endoscopic findings of a pale early mucosal lesion. There is a well-demarcated lesion; the distribution of the color is irregular, and the surface pattern is irregular. Accordingly, this lesion is diagnosed as cancerous. Adapted from Yao et al. Gastric Cancer 2017;20(Suppl 1):28–38, according to the Creative Commons license.

Fig. 3. Vessel plus surface classification system (VSCS) by magnifying narrow-band imaging. Microvascular and microsurface patterns are classified as regular/irregular/absent. If the findings fulfill the following criteria, a cancer diagnosis is made (arrow, demarcation line): (1) presence of an irregular microvascular (MV) pattern with a demarcation line, and (2) presence of an irregular microsurface (MS) pattern with a demarcation line. Otherwise, the diagnosis is noncancerous. Adapted from Muto et al. Dig Endosc 2016;28:379–393, according to the Creative Commons license.
to 100%. Hence, when the VSCS is used for diagnosis using M-NBI, M-NBI is deemed an optical biopsy.

**Preoperative assessment to select the optimal therapeutic strategy for EGC (endoscopic vs. surgical resection) for predicting submucosal cancer**

To determine the indications for endoscopic resection, we need to assess the (1) histological findings (differentiated vs. undifferentiated type), (2) size, (3) depth of invasion (mucosal vs. submucosal invasion), and (4) presence or absence of an ulcer.

We focused on the endoscopic diagnosis of the depth of invasion by chromoendoscopy, referring to “the non-extension sign” (NES). The NES is a phenomenon characterized by focal thickness and rigidity of the submucosal layer caused by massive submucosal invasion of cancerous tissue. When endoscopic air is insufflated and the stomach is distended, the form of extension differs depending on the depth of the cancer. For example, T1a-T1b1 cancers and noncancerous mucosa are well extended and flattened. However, two distinct patterns are found in the case of T1b2 cancer. (1) The T1b2 submucosal invasive cancer is not flattened and is poorly extended, forming a trapezoid elevation (Fig. 4). (2) The tips of the mucosal folds, which converge onto the trapezoid elevation, become elevated (Fig. 4).

If the endoscopic findings show both or either of these findings, the lesion is determined to be NES-positive. If both findings are negative, it is assessed as NES-negative. The sensitivity of NES for predicting a T1b2 cancer was 92.0% (95% confidence interval [CI], 87.0%–97.0%), and the specificity was 97.7% (95% CI 96.7%–98.8%).

**OUTCOMES OF RCTS TESTING THE USEFULNESS OF THE E-LEARNING SYSTEMS**

**The first e-learning system: “detection”**

In the first step, mucosal lesions suspicious for EGC should be detected using C-WLI, which is commonly available worldwide. Accordingly, the first e-learning system was designed to enable endoscopists to increase the detection of EGC using C-WLI alone. The principles and details of this e-learning system are comprehensively reported elsewhere.

In this study, participants took a pretest and were randomly assigned to either the e-learning group (ELG) or non-e-learning group (NELG). Only the participants assigned to the ELG could perform e-learning. After 2 months, both groups underwent a posttest. Eligibility was determined by 515 endoscopists from 35 countries, and 332 endoscopists were enrolled in the trial. We then randomly assigned 166 participants to either the ELG or NELG. Finally, we analyzed the data obtained from 151 participants who completed e-learning (ELG) and 144 (NELG) who did not have access to e-learning (Fig. 5).

The mean improvement rate±standard deviation in the ELG was 1.24±0.26, which was significantly higher than that of the NELG (1.00±0.16) (p<0.001, unpaired t-test) (Table 1). Regarding the other analysis, the scores of the ELG were better than those of the NELG, which means that the e-learning system was beneficial regardless of the pretest rating, participants’ expertise, or geographical origin (Table 1).

**The second e-learning system: characterization**

After detecting a suspicious mucosal lesion by C-WLI, the next process should be “characterization.” M-NBI helps differentiate between cancerous and noncancerous lesions and reduces the number of biopsies.

Although VSCS is simple, endoscopists require substantial effort to acquire “technique,” “knowledge,” and “experience” to be accustomed to this system. Because no systematic learning
Fig. 5. Flowchart of the enrollment of participants, randomization, and analysis records.

Table 1. Degree of improvement in test scores between the e-learning and non-e-learning groups

<table>
<thead>
<tr>
<th>Group</th>
<th>E-learning group</th>
<th>Non-e-learning group</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>Mean±SD (%)</td>
<td>n</td>
</tr>
<tr>
<td>Overall</td>
<td>151</td>
<td>1.24±0.26</td>
</tr>
<tr>
<td>Lower score group</td>
<td>86</td>
<td>1.34±0.29</td>
</tr>
<tr>
<td>Higher score group</td>
<td>65</td>
<td>1.19±0.14</td>
</tr>
<tr>
<td>Less experienced group</td>
<td>84</td>
<td>1.28±0.26</td>
</tr>
<tr>
<td>More experienced group</td>
<td>67</td>
<td>1.19±0.26</td>
</tr>
<tr>
<td>Asia-Oceania</td>
<td>32</td>
<td>1.33±0.34</td>
</tr>
<tr>
<td>Europe</td>
<td>22</td>
<td>1.18±0.24</td>
</tr>
<tr>
<td>Latin America</td>
<td>97</td>
<td>1.23±0.23</td>
</tr>
</tbody>
</table>

<sup>a</sup>p<0.001 for the e-learning group versus the non-e-learning group.  
<sup>b</sup>p=0.002 for the e-learning group versus the non-e-learning group.

Adapted from Yao et al. EBioMedicine 2016;9:140–147, according to the Creative Commons license.<sup>5</sup>

A total of 395 endoscopists from 77 facilities who completed test 1 were randomly assigned to either the ELG or NELG. The ELG could learn the M-NBI diagnosis through the e-learning system. After the ELG finished e-learning, both the ELG and NELG took test 2.

This trial demonstrated the usefulness of this second e-learning system for improving the diagnosis of EGC by M-NBI.

The third e-learning system: “preoperative diagnosis”

After completing two e-learning systems for “detection” and
“characterization,” we were confident that acquiring “technique,” “knowledge,” and “experience” is beneficial to endoscopists to improve the diagnosis of EGC. Teaching “technique” and “knowledge” by giving lectures using recorded video clips in an e-learning context is not so difficult. However, the “accumulation of experience” is difficult for every endoscopist. The experience of endoscopists in hospitals with few cases is limited. We assumed that for predicting submucosal EGC according to endoscopic findings, “experience” is essential for acquiring the diagnostic ability in addition to learning “the technique” and gaining “the knowledge.”

Accordingly, in contrast to the two previous RCTs, the purpose of the third study was to investigate whether this “accumulation of experience” is independently affected in improving the ability of the endoscopist to predict submucosal EGC. We carried out an RCT that included 423 endoscopists from 93 institutions all over Japan. After the participants took a pretest, they learned the “technique” and “knowledge” by watching recorded videos and the received posttest 1. After posttest 1, we randomly assigned the participants to either the self-study group (SSG) or non-self-study group (NSSG). Only participants

Fig. 6. Flowchart of the enrollment of participants, randomization, and analysis records.

![Flowchart](image)

Fig. 7. The changes in accuracy between tests 1 and 2 in the e-learning and non-e-learning groups. The mean accuracy with 95% confidence intervals in tests 1 and 2 for both groups. The change in accuracy in tests 1 and 2 is significantly higher in the e-learning group than in the non-e-learning group (Δ7.4 vs. Δ0.14 points, respectively; p<0.001, unpaired t-test). Adapted from Nakanishi et al. Endoscopy 2017;49:957–967, permission from the publisher.
in the SSG could log on to the self-study training system, where they practiced 100 case quizzes. After completing the self-study, participants in the SSG and NSSG received posttest 2. The primary endpoint was the difference in the mean scores of posttest 2 between the SSG and NSSG. The maximum score was 100.

A total of 423 endoscopists who completed the pretest were included in the study. Of these, 415 completed posttest 1 and were randomly allocated to either the SSG ($n=208$) or NSSG ($n=207$). Finally, 204 participants from the SSG and 205 from the NSSG were analyzed (Fig. 8). After watching the lecture videos, the mean score increased from 72 points on the pretest to 77 points on posttest 1. The posttest 2 score of the SSG participants was 80 points, which was significantly higher than that (76 points) of the NSSG participants ($p<0.001$, unpaired $t$-test) (Fig. 9).

**DISCUSSION AND FUTURE PLANS**

In this article, we introduced three web-based e-learning systems to teach the endoscopic diagnosis of EGC, focusing on “detection,” “characterization,” and “preoperative diagnosis.” We clearly demonstrated that these three e-learning systems are useful as shown by RCTs. Before introducing an e-learning system, we also demonstrated a basic diagnostic system and evidence for each system because an e-learning system should be

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**Fig. 8.** Flowchart of participants’ enrollment, randomization, and analysis records. The difference with this e-learning study is that to investigate the effect of self-study (accumulating “experience” alone), participants are randomized and allocated to a self-study group or non-self-study group after both groups finish a video lecture showing “technique” and “knowledge.”
constructed with concrete diagnostic procedures proven useful in well-designed clinical trials.

The e-learning included three components: “technique,” “knowledge,” and “experience.” The endoscopist can learn “technique” and “knowledge” by viewing lecture videos; however, it is difficult for the endoscopist to acquire “experience.” Accordingly, we developed a 100-case self-training program. The efficacies of the first and second e-learning systems were assessed after participants, who were allocated to the ELG, studied “technique,” “knowledge,” and “experience” altogether. The third e-learning system trial was designed differently from the previous trials because we speculated that 100-case self-training could increase the learning effect. Accordingly, we randomized the participants after completing the video lectures to independently assess the efficacy of the 100-case self-training program. Interestingly, the video lectures significantly improved the test scores; however, this learning effect did not last long without experience. The participants who completed the 100-case self-training program showed significantly greater improvement in the test scores. These results indicated that acquiring “technique” and “knowledge” is not enough to achieve improved diagnostic ability and that accumulating “experience” is mandatory.

Conventional instruction, such as teaching in clinical practice by a senior endoscopist, hands-on training using models or patients, attending lectures, and reading papers or books, is useful for imparting “knowledge” and “technique.” However, the number of learners is limited, and these forms of instruction cannot provide an accumulation of “experience.” As described previously, e-learning offers several advantages over conventional instruction. It can provide “technique” and “knowledge” by watching video clips that are easy for endoscopists to understand, and repeated training programs can be helpful for endoscopists to accumulate “experience,” which is similar to machine learning. Moreover, the number of learners is not limited, and they can access e-learning whenever and wherever they prefer.

One of the limitations of these trials is that outcome measurement was carried out using test scores on the Internet. We
have not yet assessed the improvements in clinical practice after e-learning. We aimed to plan clinical trials to assess the improvement in endoscopic diagnosis in clinical practice after participants completed e-learning.

We will consider investigating the efficacy of an e-learning system to improve the histopathological diagnosis of EGC, as this is the gold standard for endoscopic diagnosis. As previously mentioned, a diagnostic system should first be constructed. To date, no standardized diagnostic systems have been established. We developed a comprehensive algorithm for making a histological diagnosis of gastric cancer (Fig. 10). We tested whether the algorithm can function as a standard diagnostic system for e-learning content. In the near future, we will initiate the RCT by inviting international pathologists (UMIN000044545). If this trial is successfully completed, our series of e-learning systems for improving the endoscopic and histopathological diagnosis of EGC can be finalized.

CONCLUSIONS

We developed an e-learning system to improve the endoscopic diagnosis of EGC and demonstrated its usefulness.

Conflicts of Interest

The authors have no potential conflicts of interest.

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Author Contributions

Conceptualization: KY, NU, HI; Methodology: TY, SN; Project administration: NU, HD, HI; Resources: TY, SN; Software: YT; Writing–original draft: KY; Writing–review & editing: all authors.

ORCID

Kenshi Yao https://orcid.org/0000-0003-0863-3649
Takashi Yao https://orcid.org/0000-0003-2287-7658
Noriya Uedo https://orcid.org/0000-0002-3029-9272
Hisashi Doyama https://orcid.org/0000-0002-7457-0167
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