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**Usefulness of artificial intelligence in gastric neoplasms**

Kim JH *et al*. AI in gastric neoplasms

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**Abstract**

Recently, studies in many medical fields have reported that image analysis based on artificial intelligence (AI) can be used to analyze structures or features that are difficult to identify with human eyes. To diagnose early gastric cancer, related efforts such as narrow-band imaging technology are on-going. However, diagnosis is often difficult. Therefore, a diagnostic method based on AI for endoscopic imaging was developed and its effectiveness was confirmed in many studies. The gastric cancer diagnostic program based on AI showed relatively high diagnostic accuracy and could differentially diagnose non-neoplastic lesions including benign gastric ulcers and dysplasia. An AI system has also been developed that helps to predict the invasion depth of gastric cancer through endoscopic images and observe the stomach during endoscopy without blind spots. Therefore, if AI is used in the field of endoscopy, it is expected to aid in the diagnosis of gastric neoplasms and determine the application of endoscopic therapy by predicting the invasion depth.

**Key Words:** Artificial intelligence; Convolutional neural network; Gastric neoplasm; Esophagogastroduodenoscopy; Diagnosis; Invasion depth

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**Core Tip:** Recently, image analysis based on artificial intelligence (AI) has been applied in the field of diagnostic endoscopy in gastroenterology, and active research is also being conducted on gastric neoplasms. Several studies reported that AI-based early gastric cancer diagnosis and the prediction of invasion depth showed excellent performance and that the differential diagnosis from non-neoplastic lesions including benign gastric ulcers was possible. Therefore, if AI is used in clinical practice, it can be expected to help diagnose gastric neoplasms and determine treatment methods.

**INTRODUCTION**

Gastric cancer is the fifth most common malignant neoplasm in the world and the third most common cause of cancer-related death[1,2]. Although advanced gastric cancer (AGC) is associated with poor outcomes, the detection of early gastric cancer (EGC) can improve survival up to 90%[1,3]. Endoscopy is the most important tool for detecting and diagnosing gastric cancer. However, the accuracy of detection relies upon the expertise and experience of the endoscopist and complex factors of the gastrointestinal (GI) tract. Accordingly, endoscopy techniques and related fields such as image-enhanced endoscopy (IEE) have been developed to improve the diagnosis of EGC. Since its introduction in the 1950s, artificial intelligence (AI) such as deep learning (DL) has experienced remarkable progress in the last decade, and many researchers have studied the application of AI not only in the field of medical imaging but also in predicting patient prognosis based on medical records[4,5]. Many studies have utilized AI in endoscopic diagnosis. The application of AI in colonoscopy has significantly improved the adenoma detection rate (29.1% *vs* 20.3%, *P* < 0.001), and can even differentiate whether a detected polyp is non-neoplastic or neoplastic[6,7]. Based on such advancements, companies have already adapted AI for use in colonoscopy. Medtronic developed the GI Genius™ Intelligent Endoscopy Module that utilizes AI for the detection of colon polyps in real-time colonoscopy, while Olympus developed the EndoBRAIN-EYE[8]. In addition, Pentax and Fuji released the PENTAX medical Discovery™ and computer-assisted diagnosis (CAD)-EYE, respectively. Many studies have also been conducted in the field of AI in esophagogastroduodenoscopy (EGD). Thus, this article aimed to review recent developments and the use of AI in gastric neoplasms focusing on EGC, which has its unique characteristics among various GI diseases.

**AI TECHNOLOGY**

AI refers to machines that can do complex tasks like humans by imitating the cognitive functions of human intelligence such as learning and problem-solving (Figure 1). It was first introduced in 1955 and has been rapidly integrated into modern technologies and medicine[9]. Five subfields are included in AI, machine learning (ML), artificial neural network (ANN), natural language processing, DL, and computer vision[10]. ML is a field of AI where large amounts of data and algorithms are incorporated into the machine, and the machine automatically learns the input data by analyzing its patterns. Although the machine is capable of learning data patterns, the process still requires a certain amount of human instruction. DL is an important technique among many methods of ML, which is a process where the machine collects, analyzes, and processes data without receiving human instructions. Using massive amounts of data, the machine creates a learning model by extracting the key features of the given data. ANNs are the core technology of DL, and just as the human brain structure is formed by groups of neurons, the learning model of ML connects several computational nodes into several layers composed of an input layer, an output layer, and one or more hidden layers between them (Figure 2). The simplest type of neural network is called a perceptron, which consists of one input layer and one output node. The weight is a concept that gives a certain amount of importance to each input. The perceptron creates an output using inputs and weights. When an input is received, a weighted sum is calculated according to the weight, and when the value satisfies a specific criterion (activation function), the result is returned as 1 or 0. Convolutional neural network (CNN) is a kind of ANN, an algorithm that automatically learns features from the data, used mainly for image recognition[11]. It is an advanced ML model designed to think similarly to the human brain using large image datasets to learn patterns in correlating images. CNN is typically composed of three types of layers that extract features of the image and those that classify the data[12]. The convolution and pooling layers extract features of the image, while the fully connected layer is responsible for mapping them into output. The convolutional layer is a key in CNN, typically composed of a filter and an activation function. Using the image as input data, the filter extracts features of the image, and the activation function converts the value to a non-linear value. The CNN has multiple network layers of consecutive convolutional layers after pooling layers, and many filters are used as the input image is processed into consecutive convolutional layers. The extracted features are accumulated and become more complex to determine the characteristics of the input image. Subsequently, classification is performed through the fully connected layers, which are the last layers of CNN (Figure 3). As terms appearing in CNNs, one epoch refers to one forward and backward passes of the entire dataset to update the weight. The batch size is the number of training examples processed at one forward and backward pass, and iteration refers to the number of batches to complete one epoch.

Most examinations and diagnosis of GI tract diseases are performed through endoscopy and endoscopic imaging is one of the most effective applications of AI-based analytics in the field of medicine[9]. The use of CNN is ideal for endoscopic image recognition to detect and localize GI neoplasms. An AI algorithm learns what a neoplasm looks like in an endoscopic image using an image labeled by an endoscopist. After training, the CNN is tested on non-labeled new images to which it has not been previously exposed to and it is validated that the model can correctly identify previously unseen neoplasms. As a result, the algorithm can identify what it believes is a neoplasm in a real-time endoscopic video feed.

**AI IN GASTRIC NEOPLASMS**

Detection of gastric neoplasms. The detection of early-stage stomach cancer and precancerous lesions is essential to improving survival. Endoscopy is the most important and widely used detection tool for gastric cancer screening but since it is a manual procedure performed by an endoscopist, it is prone to technical and cognitive errors depending upon the endoscopist. EGC lesions usually show subtle changes of mucosa, such as elevation, depression, and redness. Moreover, they are surrounded by chronic inflammation or intestinal metaplasia. Therefore, there is a possibility of missing the subtle changes seen in the early forms of gastric cancer, especially in countries where the incidence of gastric cancer is low and where training is limited. Previous studies reported false-negative rates for detecting gastric cancer ranging between 4.6% and 25.8%[13-17]. A method to improve diagnostic accuracy involves the use of IEE such as narrow-band imaging (NBI) and blue laser imaging, which are more effective than conventional white light imaging alone[18,19]. However, such an optical diagnosis requires substantial expertise and experience[20], hindering its general use in gastroscopy. The 5-year survival rate of gastric cancer patients is highly correlated with the stage of gastric cancer at the time of diagnosis. Thus, it is paramount to improve the detection rates of EGC. Many groups have already started integrating AI into their routine practice to improve the overall detection rates of gastric cancer. AI-assisted evaluation can provide a better objective approach to improving diagnostic accuracy and avoiding unnecessary biopsies[10]. A list of studies using AI in gastric neoplasms is summarized in Table 1.

To evaluate the diagnostic accuracy of AI in the detection of gastric cancer, Hirasawa *et al*[21] used a CNN-based algorithm called the Single Shot MultiBox Detector to train using 13584 endoscopic images of gastric cancer, then tested using 2296 images (714 with confirmed gastric cancer) from 69 patients[21]. The overall sensitivity was 92.2% in the detection of gastric cancer, and the process took 47 s to analyze 2296 test images. The CNN accurately detected all invasive gastric cancer images. The detection rate for lesions larger than 6 mm was 98.6% while invasive cancers were all identified by AI. However, in the case of minute cancers that are less than 5 mm, 1 out of 6 (16.7 %) was detected, and 161 non-neoplastic lesions were included in the total 232 Lesions that were machine-identified as gastric cancer, which produced a lower positive predictive value (PPV) of 30.6%[21]. The most common cause of false-positive lesions was gastritis with a change in color tone or irregular mucosal surface, which are sometimes difficult to distinguish even by endoscopists, and the next most common cause was normal structures such as cardia, pylorus, and angle. Ishioka *et al*[22] applied the same algorithm to video images collected from 62 patients who underwent endoscopic submucosal dissection (ESD) for EGC[22]. When applied to the live video images, the diagnostic accuracy was 94.1%, and the median time for lesion detection was one second. Although the accuracy was low in minute cancers, AI showed great performance in lesions larger than 6 mm which looked very promising. In another study, Sakai *et al*[23] trained a CNN-based system with 348943 images (with data augmentation) obtained from 58 patients and tested 9650 images[23]. The accuracy of detecting gastric cancer by AI was 82.8%, and the image processing time was 4 ms *per* image.

Gastric cancer has many visual features that are challenging for endoscopists to describe. To improve diagnostic accuracy during endoscopy, several techniques have been developed to assist the gastroenterologist. Magnified NBI (M-NBI) has been shown to have higher detection rates for EGC, however, many endoscopists are not trained to confidently use M-NBI. To facilitate detection using M-NBI, Kanesaka *et al*[24] developed a CAD system to help diagnose EGC using only M-NBI images[24]. They used support vector machine to train with 66 EGC images and 60 non-cancer images, then tested detection and delineation of gastric cancer with 61 EGC and 20 non-cancer images. They reported an accuracy of 96.3%, a PPV of 98.3%, a sensitivity of 96.7%, and a specificity of 95%. Their CAD processed each image in 0.41 s[24]. In a related study, Li *et al*[25] used 386 non-cancerous M-NBI images and 1702 M-NBI images of EGC to train the Inception-v3 CNN model and tested 341 endoscopic images[25]. The sensitivity, specificity, and accuracy for the detection of EGC were 91.2%, 90.6%, and 90.9% respectively[25]. In another study by Horiuchi *et al*[26], the 22-layer GoogLeNet CNN model was trained using 1492 M-NBI images of EGC and 1078 M-NBI images of gastritis, then tested on 258 images (151 images of EGC)[26]. Further, the authors tried to determine if the differentiation between gastritis and cancer was possible. The reported accuracy for the detection of cancer was 85.3%. The sensitivity was 95.4%, the specificity was 71.0%, the PPV was 82.3%, and the negative predictive value (NPV) was 91.7%. The CNN falsely diagnosed 31 gastritis images as cancers, which were reported to have localized atrophy, atrophy of the fundic gland, and intestinal metaplasia[26]. The diagnostic performance of the same model was evaluated using 174 endoscopic videos (87 cancers and 87 non-cancers)[27]. The area under the curve (AUC) was 0.8684 and the accuracy, sensitivity, specificity, PPV, and NPV were 85.1%, 87.4%, 82.8%, 83.5%, and 86.7%, respectively. When compared to 11 experts, CAD was significantly more accurate than two experts, and not significantly different from eight experts[27].

As other studies were single-center results, or limited in the number of included cases, Luo *et al*[28] conducted a multi-center, case-controlled study of real-world endoscopic imaging to evaluate the accurate diagnosis of upper GI cancer with a CNN[28]. Using 157207 images obtained from 18765 participants from Sun Yat-sen University Cancer Center, the authors developed and validated the Gastrointestinal AI Diagnostic System (GRAIDS) algorithm through training, intrinsic verification, and internal validation. Then, they tested the performance of GRAIDS using a prospective validation dataset and additional external validation datasets obtained from five other hospitals, which included 879289 images from 65659 participants. The AUC in the external validation of the five participating hospitals ranged from 0.966 (95%CI: 0.965-0.967) to 0.990 (95%CI: 0.990-0.991)[28]. When compared to the diagnostic accuracy of the endoscopists, the diagnostic accuracy of the GRAIDS was 0.928 (95%CI: 0.919-0.937), which was significantly lower than the diagnostic accuracy of 0.967 (95%CI: 0.961-0.973; *P* < 0.0001) of the expert endoscopist (professor with more than 10 years of endoscopic experience) and 0.956 (95%CI: 0.949-0.963; *P* < 0.0001) of the competent endoscopist (attending doctor with more than five years of endoscopic experience), but significantly higher than the diagnostic accuracy of 0.886 (95%CI: 0.875-0.897; *P* < 0.0001) of the trainee endoscopist (resident with two years of endoscopic experience). The sensitivity of the GRAIDS was not significantly different from the expert [0.942 (95%CI: 0.924-0.957) *vs* 0.945 (95%CI: 0.927-0.959); *P* = 0.692]. When compared to the competent expert [0.858 (95%CI: 0.832-0.880), *P* < 0.0001] and the trainee endoscopist [0.722 (95%CI: 0.691-0.752), *P* < 0.0001], the sensitivity of the GRAIDS was confirmed to be superior. The PPV of the GRAIDS was 0.814 (95%CI: 0.788-0.838), the expert endoscopist was 0.932 (95%CI: 0.913-0.948), the competent endoscopist was 0.974 (95%CI: 0.960-0.984), and the trainee endoscopist was 0.824 (95%CI: 0.795-0.850). The PPV of the GRAIDS was lower than that of the expert and the competent endoscopist but was similar to that of the trainee. These problems are mainly because the GRAIDS misinterprets normal structures (the pylorus, angle, mucus, gastric wall elevation during peristalsis, *etc.*) as lesions, and validation was conducted with data that had a low prevalence (3.8%-9.5%) in upper GI cancer. However, it seems that normal structures can be easily distinguished by the endoscopist and confirmed as false positives. This was a notable study that used more than one million images obtained from more than 80000 patients from different centers in China. A study by Ikenoyama *et al*[29] also compared the diagnostic accuracy of AI to that of endoscopists[29]. The AI model from the previous study by Hirasawa *et al*[21]. was tested on images obtained from 75 patients with gastric cancer [66 with mucosal cancer (T1a), and nine with submucosal cancer], and the diagnostic accuracy was compared to that of 67 endoscopists (33 board-certified endoscopists with more than 18 years of experience, and 34 uncertified endoscopists with about eight years of experience). The sensitivity, specificity, PPV, and NPV of the CNN were 58.4%, 87.3%, 26.0%, and 96.5% respectively. Compared to the CNN, the endoscopists showed a sensitivity, specificity, PPV, and NPV of 31.9%, 97.2%, 46.2%, and 94.9%, respectively, which showed that the CNN had significantly higher sensitivity than the endoscopists. Also, the average time it took for the CNN to evaluate an image was 45.5 ± 1.8 s, which was much faster than the 173.0 ± 66.0 min taken by an endoscopist and suggested that AI accurately diagnosed EGC at a much higher speed[29].

***Classification of gastric neoplasms***

While many studies have tested the diagnostic accuracy of AI in differentiating cancerous lesions from normal mucosa, several attempts at classifying other non-cancerous lesions have been made (Table 1). Sun *et al*[30] created a network-based model that could classify ulcers into different types (benign ulcers or malignant ulcers) with a performance comparable to that of endoscopists[30]. The study reported that the DL model was able to identify and classify ulcers with a total accuracy of 86.6%, which was comparable to that of the endoscopist with the highest accuracy (86.3%) and higher than that of the endoscopist with the lowest accuracy (62.5%)[30]. Lee *et al*[31] developed a model that could distinguish gastric ulcers and malignancy[31]. Using the Inception-v3 network, ResNet50 and the Visual Geometry Group (VGG) Net to classify normal *vs* cancer, normal *vs* benign ulcers, and cancer *vs* benign ulcers, 180 normal images, 200 ulcer images, and 337 cancer images were used for training. When tested on 20 normal, 20 ulcers, and 30 cancer images, the best performance was observed in ResNet50, with a diagnostic accuracy of 0.9649 for differentiating between normal *vs* cancer, 0.9262 for differentiating between normal *vs* ulcers, and 0.7712 for differentiating between cancer *vs* ulcers. Based on such findings, AI was proposed as an efficient means for the classification of endoscopic images[23]. Cho *et al*[32] made a novel attempt at developing a DL model that could automatically classify gastric neoplasms using conventional endoscopic images[32]. Using 5017 images from 1269 participants, three CNN architectures (Inception-v4, ResNet-152, and Inception-ResNet-v2) were used to train and validate the classification of conventional endoscopic images. The images were classified into two categories from two perspectives, which were cancer *vs* non-cancer, and neoplasm *vs* non-neoplasm. All images were grouped into five categories, AGC, EGC, high-grade dysplasia (HGD), low-grade dysplasia (LGD), and non-neoplasm. To compare the diagnostic accuracy, six endoscopists with experience with more than 6000 endoscopies also viewed and classified the endoscopic images. The Inception-ResNet-v2 model was reported to have the best performance at classifying the images into the five categories, with an accuracy of 84.6% (95%CI: 83.69-85.5) and a mean classification time of 0.0264 s. The AUC was highest for the detection of AGC (range: 0.802-0.855) and the lowest for HGD (range: 0.491-0.522). In prospective validation, the performance of Inception-ResNet-v2 was not significantly inferior to that of the endoscopist with the worst performance. However, the endoscopist with the highest performance showed significantly better performance with a diagnostic accuracy of 87.6% (95%CI: 84.3-90.9) compared to 76.4% (95%CI: 72.1-80.7) for Inception-ResNet-v2. This suggested that AI could have the potential for classifying endoscopic lesions into several categories[32]. A recent study by Kim *et al*[33] assessed the ability of the CNN model to classify gastric mesenchymal tumors using endoscopic ultrasonography (EUS) images[33]. Using 905 EUS images from gastric mesenchymal tumors that were histologically confirmed by either resection or EUS-guided fine-needle biopsy, the CNN-CAD system was developed and validation was performed with 212 EUS images. The reported accuracy for detecting gastrointestinal stromal tumors was 79.2% using the CNN-CAD system, with a sensitivity and specificity of 83.0% and 75.5%, respectively. The performance was compared to that of six endoscopists (three experienced endoscopists who performed more than 500 EUS examinations, and three junior endoscopists who performed less than 200 EUS examinations). When compared to the diagnostic accuracy of the endoscopists, the sensitivity of CNN-CAD system was not significantly different from that of any of the endoscopists. The specificity and diagnostic accuracy of CNN-CAD system were significantly higher than that of two experienced endoscopists and one junior endoscopist[33], which suggested the potential application of AI in the classification of EUS images as well.

***Prediction of invasion depth***

The prediction of the invasion depth of gastric cancer (T-staging) is very important as it is an essential factor in determining the treatment method and prognosis of EGC. Tumors in the early stages that do not involve lymphovascular invasion and have an invasion depth no deeper than 500 μm of submucosa can be treated by endoscopic resection alone[34]. The gross findings of the tumor seen on endoscopy or EUS are used to determine the invasion depth of EGC. Some studies have reported that conventional endoscopy was comparable to EUS in predicting the invasion depth of EGC[35,36]. The reported overall accuracy of invasion depth using conventional endoscopy ranged between 69% and 79%[35,37]. In a study of depth prediction scores for differentiated EGCs, tumor sizes more than 30 mm, marked redness, an uneven surface, and marginal elevation were associated with deeper submucosal cancers[38]. However, gastric cancer depth can be difficult to determine by endoscopy alone and some patients may undergo surgery when endoscopic resection could have been an effective method of treatment. To overcome such problems, the utilization of AI to determine the depth of invasion has been studied (Table 1). Kubota *et al*[39] used retrospectively collected 902 conventional endoscopic gastric cancer images from 344 patients who underwent surgery or endoscopic resection to train and validate with a backpropagation algorithm for determining the depth of invasion[39]. The overall accuracy for detecting the depth of invasion was 64.7%, with 77.2% at the T1 stage (68.9% for T1a and 63.6% for T1b), 49.1% at the T2 stage, 51.0% at the T3 stage, and 55.3% at the T4 stage. This computer-aided system suggested a novel approach of using AI to determine cancer invasion depth by endoscopy[39]. Zhu *et al*[40] used 790 images from gastric cancer patients to train and another 203 images to validate ResNet50. The overall accuracy was reported to be 89.2%, which was significantly higher than the overall accuracy of 77.5% of the experienced endoscopists. The AUC for AI was 0.94 (95%CI: 0.90-0.97), and the sensitivity, specificity, PPV, and NPV were 76.5%, 95.6%, 89.7%, and 89.0%, respectively[40]. To test AI in the diagnostic accuracy for EGC stages, Yoon *et al*[41,42] included 800 patients, 428 patients with T1a and 372 patients with T1b histology-proven EGC, and selected 11539 images (896 T1a images, 809 T1b-EGC images, and 9834 non-cancer images) to train and validate the lesion-based VGG16-network and gradient-weighted class activation mapping (Grad-CAM)[41,42]. The overall AUC for EGC detection and invasion depth prediction was 0.981 and 0.851, respectively. Interestingly, the study also analyzed the factors affecting the AI prediction of invasion depth. The images of undifferentiated-type histology were associated with inaccurate predictions of invasion depth, especially in T1b cases[41]. As previous studies used already diagnosed gastric cancer images for training and testing, Cho *et al*[43] used Inception-ResNet-v2 and DenseNet-161 models to test the diagnostic accuracy of gastric neoplasms and invasion depth[43]. The authors used 2899 conventional endoscopic images obtained from 846 patients with confirmed pathology including LGD, HGD, EGC, and AGC. The AUC and diagnostic accuracy for determining the invasion depth were 0.887 and 77.3%, respectively, in the external validation set for the DenseNet-161 model. When applied to clinical simulation, the AI misdiagnosed only two cases that had submucosa invasion (misdiagnosed as mucosal lesions), which were also misdiagnosed by the endoscopists. In 89 patients who underwent surgery, 11 cases were actually mucosal-confined lesions, among which AI correctly classified six cases as mucosal lesions. The authors developed an algorithm with substantial performance in predicting invasion depth from the endoscopic images of neoplasms[43]. As other studies used images obtained from conventional endoscopy, Nagao *et al*[44] retrospectively collected 16557 gastric cancer images from 1084 cases to train and validate ResNet50 for predicting invasion depth by conventional white light, non-magnifying NBI, and indigo-carmine stained images[44]. The AUC using white light imaging, NBI, and indigo-carmine stain imaging were reported to be 0.9590, 0.9048, and 0.9481 respectively, and the lesion-based accuracy for predicting invasion depth using white light imaging, NBI, and indigo-carmine were 94.5%, 94.3%, and 95.5%, respectively[44].

***Blind-spot monitoring***

Observing the whole stomach is a basic prerequisite for the diagnosis of gastric cancer at an early stage. To avoid blind spots, standardized procedures and guidelines have been made to map the entire stomach during gastroscopy. The European Society of Gastrointestinal Endoscopy published a protocol including 10 images of the stomach while the systematic screening protocol for the stomach published by Japanese researchers suggested 22 standard images of the stomach to avoid missing suspicious cancerous lesions[45,46]. However, insufficient supervision and the lack of practical tools make it difficult to follow protocols, which is related to the quality of endoscopic examinations[47]. To localize blind spots during EGD that may have been missed by an endoscopist, Wu *et al*[48] developed the WISENSE system, a real-time CNN to detect blind spots (Table 1)[48]. As the scope was inserted into the stomach, the deep CNN (DCNN) captured images and filled them into the corresponding part of the model, which enabled the endoscopist to identify the blind spots. These blind spots of the gastric mucosa, such as the lesser curvature of the antrum and the fundus, are areas that may hide lesions. If blind spots are not viewed during endoscopy, lesions could be missed. Trained on 34513 images of gastric locations agreed upon by at least four endoscopists, WISENSE was able to detect blind spots with an accuracy of 90.0% by identifying anatomic landmarks in EGD. In a single-center randomized control trial, 153 patients had their blind spots detected by WISENSE *vs* 150 in the control group without AI. The blind spot rate, defined as the proportion of the number of unobserved sites in 26 sites, was 5.9% in the WISENSE group which was significantly less than 22.5% in the control group (*P* < 0.001), suggesting that AI can also be used to improve the quality of EGD by identifying blind spots[48]. In another study by Wu *et al*[49], a DCNN was used for detecting gastric cancer and identifying blind spots. There were 24549 images used for training, and a grid model for the stomach was developed to generate a virtual stomach model[49]. The study reported a diagnostic accuracy of 92.5% for detecting malignancy, which was significantly higher than that of six experts. The reported sensitivity, specificity, PPV, and NPV were 94.0%, 91.0%, 91.3%, and 93.8%, respectively. The DCNN correctly identified the EGD images into 10 parts with an accuracy of 90.0% and into 26 parts with an accuracy of 65.9%, which was not significantly different from those of the endoscopists. When the model was tested on endoscopic videos, the DCNN accurately presented the covered parts synchronized with the process of EGD to verify that the entire stomach was mapped[49]. A related study by Chen *et al*[50] used ENDOANGEL (developed from WISENSE) to compare blind-spot monitoring in sedated conventional EGD, unsedated ultrathin endoscopy, and unsedated conventional EGD[50]. This prospective, 3-parallel-group, randomized study reported that the blind-spot rate using AI was significantly lower than that of unsedated endoscopy with an ultrathin endoscope and unsedated conventional EGD (3.4% *vs* 21.8% *vs* 31.2%, *P* < 0.05). Although the number of studies is limited, the application of AI in monitoring blind spots is very promising.

***Prediction of curative endoscopic resection***

Expanded indications for ESD in EGC include the undifferentiated type that is less than 2 cm and does not have ulcerations. However, observational studies have reported conflicting results. Thus, ESD in such groups has been considered an investigational treatment[34]. A meta-analysis of curative resection for EGC with undifferentiated type histology reported a rate of 61.4%, suggesting ESD as a feasible treatment for undifferentiated-type EGC[51]. To aid in the accurate prediction of curative resection in such cases, Bang *et al*[52] selected ML models that could predict curative resection in undifferentiated-type EGC. The XGBoost classifier presented the best performance with an accuracy of 81.5% in the first external validation and 89.8% in the second external validation[52]. The size of the lesion was the most important feature that could be explained by AI analysis. As such, AI could aid in decisions for therapeutic management.

**FUTURE PERSPECTIVES OF AI**

The real-time application of AI in the field of medicine is within reach. Endoscopic models that automatically detect colon polyps or gastric cancers during endoscopy sessions and highlight them using segmentation box have already received approval for use in Europe, Japan, and other countries, while many systems are currently under development[8]. Many software codes have been provided as open-source codes, which can be freely utilized in research or actual practice. Architectures can be modified by fine-tuning an already established pre-trained model by adjusting layers of the ANN, increasing the learning epoch, adjusting the batch size, adjusting the iteration, or modifying hyperparameters such as the optimizer. Aside from adjusting the complex algorithms to optimize the model, recent developments have enabled the automatic optimization of hyperparameters in ML (*i.e.*, AutoGluon) that makes AI more user-friendly and easier to use for clinicians unfamiliar with AI[52]. Most research on AI for gastroenterology has focused on developing algorithms for the detection of lesions, the classification of images to improve diagnostic accuracy, predicting prognosis, and to improve the quality of screening endoscopy. In the near future, AI will most likely be applied to therapeutic management. Recently, AI-based treatment methods have been developed using technologies such as microendoscopy, decision support system-based treatment modalities, robot-assisted treatment, application, and digital therapeutics[53]. However, such development comes with social issues other than technology, such as patient safety, ethics, legal responsibility, government approval, and cost-effectiveness, which need to be addressed as well. Although studies have shown that accuracy of detecting gastric cancer by AI is comparable to some doctors, experienced doctors with expertise have shown better performance than AI. This means that there is limitation to relying solely on AI alone. However, beneficial factors from application of AI, such as improved efficacy and time spent on repetitive task, must be acknowledged as well. Accordingly, the most applicable field of AI would be medical image data processing that could aid in improved diagnostic performance of trainees and non-expert doctors. The AI algorithm, especially DL, is comparable to a black box that learns from training data. Using the patterns learned from the training data, the output values can be predicted from newly input data. This means that efficacy and accuracy are highly dependent upon the quality and quantity of the training data. Like any other clinical research, the quality and quantity of the usable data are undeniably essential in proving the quality of the evidence and the outcome. It is important to gather high-quality clinical data, while developing a model that accurately tests the data is equally important. To effectively utilize such an AI algorithm in clinical practice, further studies and discussions on the usefulness, profitability, possible risks, medicolegal responsibility, and regulatory measures of AI are needed.

**CONCLUSION**

AI in the field of endoscopy was first applied for the detection of colon polyps. As described in this review article, many studies have already been published as stepping-stones toward the application of AI in detecting gastric neoplasms such as EGC. As there is a lack of such prospective studies in the detection of EGC, randomized controlled studies are needed to advance the technique. It is expected that the application of AI would not only provide guidelines for the endoscopic treatment of EGC or avoid unnecessary surgery by predicting the invasion depth but also help improve the overall prognosis of patients with EGC. There is no doubt that the development of AI-based endoscopy would also help to alleviate physical fatigue that can be a burden to endoscopists. Such achievements can only be done when the application of AI can improve the quality of imaging diagnosis beyond that of human capability, and optical biopsy is possible. This is possible by improving AI performance using the specific characteristics of different organs and diseases. AI is being studied and developed by scientists all over the world in various fields with hopes of providing accuracy and convenience. In the field of medicine, medical records and imaging are becoming digitalized and a new phase in the history of medicine is expected within five to 10 years. Accordingly, clinicians and researchers need to carefully approach and evaluate the results of further clinical studies using AI-based technology with great interest.

**REFERENCES**

1 **Sano T**, Coit DG, Kim HH, Roviello F, Kassab P, Wittekind C, Yamamoto Y, Ohashi Y. Proposal of a new stage grouping of gastric cancer for TNM classification: International Gastric Cancer Association staging project. *Gastric Cancer* 2017; **20**: 217-225 [PMID: 26897166 DOI: 10.1007/s10120-016-0601-9]

2 **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**: 394-424 [PMID: 30207593 DOI: 10.3322/caac.21492]

3 **Amin MB**, Greene FL, Edge SB, Compton CC, Gershenwald JE, Brookland RK, Meyer L, Gress DM, Byrd DR, Winchester DP. The Eighth Edition AJCC Cancer Staging Manual: Continuing to build a bridge from a population-based to a more "personalized" approach to cancer staging. *CA Cancer J Clin* 2017; **67**: 93-99 [PMID: 28094848 DOI: 10.3322/caac.21388]

4 **Erickson BJ**, Korfiatis P, Akkus Z, Kline TL. Machine Learning for Medical Imaging. *Radiographics* 2017; **37**: 505-515 [PMID: 28212054 DOI: 10.1148/rg.2017160130]

5 **Heo J**, Yoon JG, Park H, Kim YD, Nam HS, Heo JH. Machine Learning-Based Model for Prediction of Outcomes in Acute Stroke. *Stroke* 2019; **50**: 1263-1265 [PMID: 30890116 DOI: 10.1161/STROKEAHA.118.024293]

6 **Song EM**, Park B, Ha CA, Hwang SW, Park SH, Yang DH, Ye BD, Myung SJ, Yang SK, Kim N, Byeon JS. Endoscopic diagnosis and treatment planning for colorectal polyps using a deep-learning model. *Sci Rep* 2020; **10**: 30 [PMID: 31913337 DOI: 10.1038/s41598-019-56697-0]

7 **Wang P**, Berzin TM, Glissen Brown JR, Bharadwaj S, Becq A, Xiao X, Liu P, Li L, Song Y, Zhang D, Li Y, Xu G, Tu M, Liu X. Real-time automatic detection system increases colonoscopic polyp and adenoma detection rates: a prospective randomised controlled study. *Gut* 2019; **68**: 1813-1819 [PMID: 30814121 DOI: 10.1136/gutjnl-2018-317500]

8 **Attardo S**, Chandrasekar VT, Spadaccini M, Maselli R, Patel HK, Desai M, Capogreco A, Badalamenti M, Galtieri PA, Pellegatta G, Fugazza A, Carrara S, Anderloni A, Occhipinti P, Hassan C, Sharma P, Repici A. Artificial intelligence technologies for the detection of colorectal lesions: The future is now. *World J Gastroenterol* 2020; **26**: 5606-5616 [PMID: 33088155 DOI: 10.3748/wjg.v26.i37.5606]

9 **Abadir AP**, Ali MF, Karnes W, Samarasena JB. Artificial Intelligence in Gastrointestinal Endoscopy. *Clin Endosc* 2020; **53**: 132-141 [PMID: 32252506 DOI: 10.5946/ce.2020.038]

10 **Jin P**, Ji X, Kang W, Li Y, Liu H, Ma F, Ma S, Hu H, Li W, Tian Y. Artificial intelligence in gastric cancer: a systematic review. *J Cancer Res Clin Oncol* 2020; **146**: 2339-2350 [PMID: 32613386 DOI: 10.1007/s00432-020-03304-9]

11 **Ebigbo A**, Palm C, Probst A, Mendel R, Manzeneder J, Prinz F, de Souza LA, Papa JP, Siersema P, Messmann H. A technical review of artificial intelligence as applied to gastrointestinal endoscopy: clarifying the terminology. *Endosc Int Open* 2019; **7**: E1616-E1623 [PMID: 31788542 DOI: 10.1055/a-1010-5705]

12 **Yamashita R**, Nishio M, Do RKG, Togashi K. Convolutional neural networks: an overview and application in radiology. *Insights Imaging* 2018; **9**: 611-629 [PMID: 29934920 DOI: 10.1007/s13244-018-0639-9]

13 **Amin A**, Gilmour H, Graham L, Paterson-Brown S, Terrace J, Crofts TJ. Gastric adenocarcinoma missed at endoscopy. *J R Coll Surg Edinb* 2002; **47**: 681-684 [PMID: 12463707]

14 **Hosokawa O**, Hattori M, Douden K, Hayashi H, Ohta K, Kaizaki Y. Difference in accuracy between gastroscopy and colonoscopy for detection of cancer. *Hepatogastroenterology* 2007; **54**: 442-444 [PMID: 17523293]

15 **Menon S**, Trudgill N. How commonly is upper gastrointestinal cancer missed at endoscopy? A meta-analysis. *Endosc Int Open* 2014; **2**: E46-E50 [PMID: 26135259 DOI: 10.1055/s-0034-1365524]

16 **Voutilainen ME**, Juhola MT. Evaluation of the diagnostic accuracy of gastroscopy to detect gastric tumours: clinicopathological features and prognosis of patients with gastric cancer missed on endoscopy. *Eur J Gastroenterol Hepatol* 2005; **17**: 1345-1349 [PMID: 16292088 DOI: 10.1097/00042737-200512000-00013]

17 **Yalamarthi S**, Witherspoon P, McCole D, Auld CD. Missed diagnoses in patients with upper gastrointestinal cancers. *Endoscopy* 2004; **36**: 874-879 [PMID: 15452783 DOI: 10.1055/s-2004-825853]

18 **Diao W**, Huang X, Shen L, Zeng Z. Diagnostic ability of blue laser imaging combined with magnifying endoscopy for early esophageal cancer. *Dig Liver Dis* 2018; **50**: 1035-1040 [PMID: 29685806 DOI: 10.1016/j.dld.2018.03.027]

19 **Dutta AK**, Sajith KG, Pulimood AB, Chacko A. Narrow band imaging *vs* white light gastroscopy in detecting potentially premalignant gastric lesions: a randomized prospective crossover study. *Indian J Gastroenterol* 2013; **32**: 37-42 [PMID: 22983839 DOI: 10.1007/s12664-012-0246-5]

20 **Ishihara R**, Takeuchi Y, Chatani R, Kidu T, Inoue T, Hanaoka N, Yamamoto S, Higashino K, Uedo N, Iishi H, Tatsuta M, Tomita Y, Ishiguro S. Prospective evaluation of narrow-band imaging endoscopy for screening of esophageal squamous mucosal high-grade neoplasia in experienced and less experienced endoscopists. *Dis Esophagus* 2010; **23**: 480-486 [PMID: 20095991 DOI: 10.1111/j.1442-2050.2009.01039.x]

21 **Hirasawa T**, Aoyama K, Tanimoto T, Ishihara S, Shichijo S, Ozawa T, Ohnishi T, Fujishiro M, Matsuo K, Fujisaki J, Tada T. Application of artificial intelligence using a convolutional neural network for detecting gastric cancer in endoscopic images. *Gastric Cancer* 2018; **21**: 653-660 [PMID: 29335825 DOI: 10.1007/s10120-018-0793-2]

22 **Ishioka M**, Hirasawa T, Tada T. Detecting gastric cancer from video images using convolutional neural networks. *Dig Endosc* 2019; **31**: e34-e35 [PMID: 30449050 DOI: 10.1111/den.13306]

23 **Sakai Y**, Takemoto S, Hori K, Nishimura M, Ikematsu H, Yano T, Yokota H. Automatic detection of early gastric cancer in endoscopic images using a transferring convolutional neural network. *Annu Int Conf IEEE Eng Med Biol Soc* 2018; **2018**: 4138-4141 [PMID: 30441266 DOI: 10.1109/EMBC.2018.8513274]

24 **Kanesaka T**, Lee TC, Uedo N, Lin KP, Chen HZ, Lee JY, Wang HP, Chang HT. Computer-aided diagnosis for identifying and delineating early gastric cancers in magnifying narrow-band imaging. *Gastrointest Endosc* 2018; **87**: 1339-1344 [PMID: 29225083 DOI: 10.1016/j.gie.2017.11.029]

25 **Li L**, Chen Y, Shen Z, Zhang X, Sang J, Ding Y, Yang X, Li J, Chen M, Jin C, Chen C, Yu C. Convolutional neural network for the diagnosis of early gastric cancer based on magnifying narrow band imaging. *Gastric Cancer* 2020; **23**: 126-132 [PMID: 31332619 DOI: 10.1007/s10120-019-00992-2]

26 **Horiuchi Y**, Aoyama K, Tokai Y, Hirasawa T, Yoshimizu S, Ishiyama A, Yoshio T, Tsuchida T, Fujisaki J, Tada T. Convolutional Neural Network for Differentiating Gastric Cancer from Gastritis Using Magnified Endoscopy with Narrow Band Imaging. *Dig Dis Sci* 2020; **65**: 1355-1363 [PMID: 31584138 DOI: 10.1007/s10620-019-05862-6]

27 **Horiuchi Y**, Hirasawa T, Ishizuka N, Tokai Y, Namikawa K, Yoshimizu S, Ishiyama A, Yoshio T, Tsuchida T, Fujisaki J, Tada T. Performance of a computer-aided diagnosis system in diagnosing early gastric cancer using magnifying endoscopy videos with narrow-band imaging (with videos). *Gastrointest Endosc* 2020; **92**: 856-865.e1 [PMID: 32422155 DOI: 10.1016/j.gie.2020.04.079]

28 **Luo H**, Xu G, Li C, He L, Luo L, Wang Z, Jing B, Deng Y, Jin Y, Li Y, Li B, Tan W, He C, Seeruttun SR, Wu Q, Huang J, Huang DW, Chen B, Lin SB, Chen QM, Yuan CM, Chen HX, Pu HY, Zhou F, He Y, Xu RH. Real-time artificial intelligence for detection of upper gastrointestinal cancer by endoscopy: a multicentre, case-control, diagnostic study. *Lancet Oncol* 2019; **20**: 1645-1654 [PMID: 31591062 DOI: 10.1016/S1470-2045(19)30637-0]

29 **Ikenoyama Y**, Hirasawa T, Ishioka M, Namikawa K, Yoshimizu S, Horiuchi Y, Ishiyama A, Yoshio T, Tsuchida T, Takeuchi Y, Shichijo S, Katayama N, Fujisaki J, Tada T. Detecting early gastric cancer: Comparison between the diagnostic ability of convolutional neural networks and endoscopists. *Dig Endosc* 2021; **33**: 141-150 [PMID: 32282110 DOI: 10.1111/den.13688]

30 **Sun JY**, Sang WL, Kang MC, Kim SW, Ko SJ. A Novel Gastric Ulcer Differentiation System Using Convolutional Neural Networks. Proceedings of the 2018 IEEE 31st International Symposium on Computer-Based Medical Systems (CBMS); 2018 June 1; IEEE, 2018: 351-356

31 **Lee JH**, Kim YJ, Kim YW, Park S, Choi YI, Kim YJ, Park DK, Kim KG, Chung JW. Spotting malignancies from gastric endoscopic images using deep learning. *Surg Endosc* 2019; **33**: 3790-3797 [PMID: 30719560 DOI: 10.1007/s00464-019-06677-2]

32 **Cho BJ**, Bang CS, Park SW, Yang YJ, Seo SI, Lim H, Shin WG, Hong JT, Yoo YT, Hong SH, Choi JH, Lee JJ, Baik GH. Automated classification of gastric neoplasms in endoscopic images using a convolutional neural network. *Endoscopy* 2019; **51**: 1121-1129 [PMID: 31443108 DOI: 10.1055/a-0981-6133]

33 **Kim YH**, Kim GH, Kim KB, Lee MW, Lee BE, Baek DH, Kim DH, Park JC. Application of A Convolutional Neural Network in The Diagnosis of Gastric Mesenchymal Tumors on Endoscopic Ultrasonography Images. *J Clin Med* 2020; **9** [PMID: 33003602 DOI: 10.3390/jcm9103162]

34 **Japanese Gastric Cancer Association.**. Japanese gastric cancer treatment guidelines 2014 (ver. 4). *Gastric Cancer* 2017; **20**: 1-19 [PMID: 27342689 DOI: 10.1007/s10120-016-0622-4]

35 **Choi J**, Kim SG, Im JP, Kim JS, Jung HC, Song IS. Comparison of endoscopic ultrasonography and conventional endoscopy for prediction of depth of tumor invasion in early gastric cancer. *Endoscopy* 2010; **42**: 705-713 [PMID: 20652857 DOI: 10.1055/s-0030-1255617]

36 **Pei Q**, Wang L, Pan J, Ling T, Lv Y, Zou X. Endoscopic ultrasonography for staging depth of invasion in early gastric cancer: A meta-analysis. *J Gastroenterol Hepatol* 2015; **30**: 1566-1573 [PMID: 26094975 DOI: 10.1111/jgh.13014]

37 **Sano T**, Okuyama Y, Kobori O, Shimizu T, Morioka Y. Early gastric cancer. Endoscopic diagnosis of depth of invasion. *Dig Dis Sci* 1990; **35**: 1340-1344 [PMID: 2226095 DOI: 10.1007/BF01536738]

38 **Abe S**, Oda I, Shimazu T, Kinjo T, Tada K, Sakamoto T, Kusano C, Gotoda T. Depth-predicting score for differentiated early gastric cancer. *Gastric Cancer* 2011; **14**: 35-40 [PMID: 21327924 DOI: 10.1007/s10120-011-0002-z]

39 **Kubota K**, Kuroda J, Yoshida M, Ohta K, Kitajima M. Medical image analysis: computer-aided diagnosis of gastric cancer invasion on endoscopic images. *Surg Endosc* 2012; **26**: 1485-1489 [PMID: 22083334 DOI: 10.1007/s00464-011-2036-z]

40 **Zhu Y**, Wang QC, Xu MD, Zhang Z, Cheng J, Zhong YS, Zhang YQ, Chen WF, Yao LQ, Zhou PH, Li QL. Application of convolutional neural network in the diagnosis of the invasion depth of gastric cancer based on conventional endoscopy. *Gastrointest Endosc* 2019; **89**: 806-815.e1 [PMID: 30452913 DOI: 10.1016/j.gie.2018.11.011]

41 **Yoon HJ**, Kim S, Kim JH, Keum JS, Oh SI, Jo J, Chun J, Youn YH, Park H, Kwon IG, Choi SH, Noh SH. A Lesion-Based Convolutional Neural Network Improves Endoscopic Detection and Depth Prediction of Early Gastric Cancer. *J Clin Med* 2019; **8** [PMID: 31454949 DOI: 10.3390/jcm8091310]

42 **Yoon HJ**, Kim JH. Lesion-Based Convolutional Neural Network in Diagnosis of Early Gastric Cancer. *Clin Endosc* 2020; **53**: 127-131 [PMID: 32252505 DOI: 10.5946/ce.2020.046]

43 **Cho BJ**, Bang CS, Lee JJ, Seo CW, Kim JH. Prediction of Submucosal Invasion for Gastric Neoplasms in Endoscopic Images Using Deep-Learning. *J Clin Med* 2020; **9** [PMID: 32549190 DOI: 10.3390/jcm9061858]

44 **Nagao S**, Tsuji Y, Sakaguchi Y, Takahashi Y, Minatsuki C, Niimi K, Yamashita H, Yamamichi N, Seto Y, Tada T, Koike K. Highly accurate artificial intelligence systems to predict the invasion depth of gastric cancer: efficacy of conventional white-light imaging, nonmagnifying narrow-band imaging, and indigo-carmine dye contrast imaging. *Gastrointest Endosc* 2020; **92**: 866-873.e1 [PMID: 32592776 DOI: 10.1016/j.gie.2020.06.047]

45 **Bisschops R**, Areia M, Coron E, Dobru D, Kaskas B, Kuvaev R, Pech O, Ragunath K, Weusten B, Familiari P, Domagk D, Valori R, Kaminski MF, Spada C, Bretthauer M, Bennett C, Senore C, Dinis-Ribeiro M, Rutter MD. Performance measures for upper gastrointestinal endoscopy: a European Society of Gastrointestinal Endoscopy (ESGE) Quality Improvement Initiative. *Endoscopy* 2016; **48**: 843-864 [PMID: 27548885 DOI: 10.1055/s-0042-113128]

46 **Yao K**. The endoscopic diagnosis of early gastric cancer. *Ann Gastroenterol* 2013; **26**: 11-22 [PMID: 24714327]

47 **Rutter MD**, Rees CJ. Quality in gastrointestinal endoscopy. *Endoscopy* 2014; **46**: 526-528 [PMID: 24788539 DOI: 10.1055/s-0034-1365738]

48 **Wu L**, Zhang J, Zhou W, An P, Shen L, Liu J, Jiang X, Huang X, Mu G, Wan X, Lv X, Gao J, Cui N, Hu S, Chen Y, Hu X, Li J, Chen D, Gong D, He X, Ding Q, Zhu X, Li S, Wei X, Li X, Wang X, Zhou J, Zhang M, Yu HG. Randomised controlled trial of WISENSE, a real-time quality improving system for monitoring blind spots during esophagogastroduodenoscopy. *Gut* 2019; **68**: 2161-2169 [PMID: 30858305 DOI: 10.1136/gutjnl-2018-317366]

49 **Wu L**, Zhou W, Wan X, Zhang J, Shen L, Hu S, Ding Q, Mu G, Yin A, Huang X, Liu J, Jiang X, Wang Z, Deng Y, Liu M, Lin R, Ling T, Li P, Wu Q, Jin P, Chen J, Yu H. A deep neural network improves endoscopic detection of early gastric cancer without blind spots. *Endoscopy* 2019; **51**: 522-531 [PMID: 30861533 DOI: 10.1055/a-0855-3532]

50 **Chen D**, Wu L, Li Y, Zhang J, Liu J, Huang L, Jiang X, Huang X, Mu G, Hu S, Hu X, Gong D, He X, Yu H. Comparing blind spots of unsedated ultrafine, sedated, and unsedated conventional gastroscopy with and without artificial intelligence: a prospective, single-blind, 3-parallel-group, randomized, single-center trial. *Gastrointest Endosc* 2020; **91**: 332-339.e3 [PMID: 31541626 DOI: 10.1016/j.gie.2019.09.016]

51 **Bang CS**, Baik GH, Shin IS, Kim JB, Suk KT, Yoon JH, Kim YS, Kim DJ, Shin WG, Kim KH, Kim HY, Lim H, Kang HS, Kim JH, Kim JB, Jung SW, Kae SH, Jang HJ, Choi MH. Endoscopic submucosal dissection for early gastric cancer with undifferentiated-type histology: A meta-analysis. *World J Gastroenterol* 2015; **21**: 6032-6043 [PMID: 26019470 DOI: 10.3748/wjg.v21.i19.6032]

52 **Bang CS,** Ahn JY, Kim JH, Kim YI, Choi IJ, Shin WG. Establishing Machine Learning Models to Predict Curative Resection in Early Gastric Cancer with Undifferentiated Histology: Development and Usability Study. *J Med Internet Res* 2021; **23:** e25053 [PMID: 33856358 DOI: 10.2196/25053]

53 **Bang CS**. [Deep Learning in Upper Gastrointestinal Disorders: Status and Future Perspectives]. *Korean J Gastroenterol* 2020; **75**: 120-131 [PMID: 32209800 DOI: 10.4166/kjg.2020.75.3.120]

**Footnotes**

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**Figure Legends**



**Figure 1 Overview of artificial intelligence, machine learning, and deep learning.** Artificial intelligence refers to machines that can do complex tasks like humans by imitating human intelligence. One of the most important ways to achieve artificial intelligence is machine learning. Machine can learn by itself from the data provided to make accurate decisions. Deep learning is an important technique among many methods of machine learning. It is a kind of artificial neural network and learns data through an information input/output layer similar to neurons in the brain.



**Figure 2 Illustrative model of artificial neural network.** Once endoscopic image is selected as input layer, hidden layers are connected to next layer. Through this network, the input image is classified into output layer.



**Figure 3 Overview of convolutional neural network.** It is composed of stacks of convolutional layers, pooling layers, and fully connected layers. Convolutional and pooling layer extract features of input images, while fully connected layers make output based on classification.

**Table 1 Recently published articles on application of artificial intelligence in gastric neoplasms**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Ref.** | **Purpose** | **AI type** | **Endoscopy type** | **Subjects** | **Outcomes** |
| **Detection of gastric neoplasms** |
| Hirasawa *et al*[[21](#_ENREF_21)], 2018 | Detect EGC | CNN (SSD) | Conventional endoscopy | Training: 13,584 images; Test: 2296 images from 69 patients. | Sensitivity 92.2%, PPV 30.6% |
| Ishioka *et al*[[22](#_ENREF_22)], 2019 | Real time detection of EGC | CNN (SSD) | Conventional endoscopy | Live video of 62 patients | Accuracy 94.1%, median time 1 s (range: 0-44 s) |
| Sakai *et al*[[23](#_ENREF_23)], 2018 | Detect EGC | CNN | Conventional endoscopy | Training: 348943 images; Test: 9650 images | Accuracy 82.8% |
| Kanesaka *et al*[[24](#_ENREF_24)], 2018 | Detect EGC | SVM | M-NBI | Training: 126 images; Test: 81 images | Accuracy 96.3%, sensitivity 96.7%, specificity 95% |
| Li *et al*[[25](#_ENREF_25)], 2020 | Detect EGC | CNN (Inception-v3) | M-NBI | Training: 2088 images; Test: 341 images | Accuracy 91.2%, sensitivity 90.6%, specificity 90.9% |
| Horiuchi *et al*[[26](#_ENREF_26)], 2020 | Classifying EGC from gastritis | CNN (GoogLeNet) | M-NBI | Training: 2570 images; Test: 258 images. | Accuracy 85.3%, sensitivity 95.4%, specificity 71.0%, test speed 51.83 images/s (0.02 s/image) |
| Horiuchi *et al*[[27](#_ENREF_27)], 2020 | Detect EGC | CNN (GoogLeNet) | M-NBI videos | 174 videos | Accuracy 85.1%, AUC 0.8684, sensitivity 87.4%, specificity 82.8%, PPV 83.5%, NPV 86.7% |
| Luo *et al*[[28](#_ENREF_28)], 2019 | Real time detection of EGC | GRAIDS | Conventional endoscopy | 1036496 images from 84424 patients | Sensitivity (0.942) similar to the expert (0.945), superior to the competent (0.858) and the trainee (0.722) endoscopist |
| Ikenoyama *et al*[[29](#_ENREF_29)], 2021 | Detect EGC | CNN (SSD) | WLI, NBI chromoendoscopy | Training: 13584 images; Test: 2940 images. | Sensitivity 58.4%, specificity 87.3%, PPV 26.0%, NPV 96.5% |
| **Classification of gastric neoplasms** |
| Sun *et al*[[30](#_ENREF_30)], 2018 | Classify ulcers | DCNN | Conventional endoscopy | 854 images | Accuracy 86.6%, sensitivity 90.8%, specificity 83.5%  |
| Lee *et al*[[31](#_ENREF_31)], 2019 | Detect EGC and benign ulcer | CNN (ResNet50, Inception-v3, VGG16) | Conventional endoscopy | Training: 717 images; Test: 70 images | AUC 0.95, 0.97, and 0.85 in Inception, ResNet50, and VGG16 |
| Cho *et al*[[32](#_ENREF_32)], 2019 | Detect AGC, EGC, dysplasia | CNN (Inception-v4, ResNet152, Inception-Resnet-v2) | Conventional endoscopy | 5217 images from 1469 patients | Gastric cancer: accuracy 81.9%, AUC 0.877; Gastric neoplasm: accuracy 85.5%, AUC 0.927  |
| Kim *et al*[[33](#_ENREF_33)], 2020 | Classify gastric mesenchymal tumors | CNN | Endoscopic ultrasonography | Training: 905 images; Test: 212 images. | Accuracy 79.2%, sensitivity 83.0%. specificity 75.5%  |
| **Prediction of invasion depth** |
| Kubota *et al*[[39](#_ENREF_39)], 2012 | Predict invasion depth | Back propagation | Conventional endoscopy | Training: 800 images; Test: 90 images | Accuracy 77.9%, 29.1%, 51.0% and 55.3% in T1, T2, T3, and T4 stage; Accuracy 68.9% and 63.6% in T1a and T1b stage |
| Zhu *et al*[[40](#_ENREF_40)], 2019 | Predict invasion depth | CNN (ResNet50) | Conventional endoscopy | Training: 790 images; Test: 203 images | AUC 0.94 (95% CI: 0.90-0.97), overall accuracy 89.2%, sensitivity 76.5%, specificity 95.6% |
| Yoon *et al*[[41](#_ENREF_41)], 2019 | Detect cancer, and predict invasion depth | CNN (VGG16, Grad-CAM) | Conventional endoscopy | 11539 images | Detection AUC 0.981, depth prediction AUC 0.851 (undifferentiated type histology with a lower accuracy) |
| Cho *et al*[[43](#_ENREF_43)], 2020 | Predict invasion depth | CNN (Inception-ResNet-v2, DenseNet-161) | Conventional endoscopy | Training: 2899 images, test: 206 images | Internal validation: accuracy 84.1%, AUC 0.887; External validation: accuracy 77.3%, AUC 0.887  |
| Nagao *et al*[[44](#_ENREF_44)], 2020 | Predict invasion depth | CNN (ResNet50) | WLI, NBI, indigo-carmine  | 16557 images from 1084 cases of gastric cancer | WLI: AUC 0.9590, sensitivity 89.2%, specificity 98.7%, accuracy 94.4%, PPV 98.3%, NPV 91.7%; NBI: AUC 0.9048; Indigo-carmine: AUC 0.9191 |
| **Blind-spot monitoring** |
| Wu *et al*[[48](#_ENREF_48)], 2019 | Detect blind spot | DCNN | Conventional endoscopy | 34513 images | Accuracy of detecting blind spot: 90.0%; Blind spot rate: 5.9% |
| Wu *et al*[[49](#_ENREF_49)], 2019 | Detect EGC and blind spot | DCNN | Conventional endoscopy | 24549 images | Accuracy 92.5%, sensitivity 94.0%, specificity 91.0%, PPV 91.3%, NPV 93.8% |
| Chen *et al*[[50](#_ENREF_50)], 2020 | Detect blind spot | DCNN | Live video | 437 patients | Blind spot rate: AI 3.4%, ultrathin endoscope 21.8%, conventional endoscope 31.2% |

AI: Artificial intelligence; EGC: Early gastric cancer; CNN: Convolutional neural network; SSD: Single Shot MultiBox Detector; PPV: Positive predict value; SVM: Support vector machine; M-NBI: Magnified narrow band imaging; AUC: Area under curve; GRAIDS: Gastrointestinal Artificial Intelligence Diagnostic System; NBI: Narrow band imaging; NPV: Negative predict value; DCNN: Deep convolutional neural network; VGG: Visual Geometry Group; AGC: Advanced gastric cancer; Grad-CAM: Gradient-weighted class activation mapping.