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Advancements in biliopancreatic endoscopy - A comprehensive review of artificial intelligence in EUS and ERCP

Authors:

Belén Agudo Castillo, Miguel Mascarenhas, Miguel Martins, Francisco Mendes, Daniel de la Iglesia, Antonio Miguel Martins Pinto da Costa, Carlos Esteban Fernández-Zarza, Mariano González-Haba Ruiz

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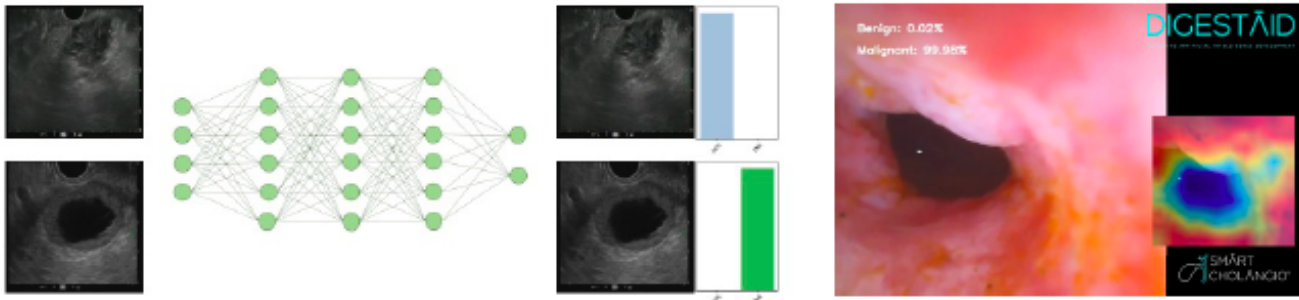
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A COMPREHENSIVE REVIEW OF ARTIFICIAL INTELLIGENCE IN EUS AND ERCP



Example of an output generated by artificial intelligence with differentiation between different types of solid pancreatic lesions in an endoscopic ultrasound exam. The bars represent the probability estimated by the network ADC - pancreatic adenocarcinoma or TNE - pancreatic neuroendocrine tumor. In the right case in a biliary stricture the software discriminates in real time between benign and malignant origin, while a heatmap highlights the areas with highest probability of malignancy (Source: Digestaid).

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Belén Agudo Castillo¹, Miguel Mascarenhas², Miguel Martins², Francisco Mendes², Daniel de la Iglesia¹, Antonio Miguel Martins Pinto da Costa¹, Carlos Esteban Fernández-Zarza¹, Mariano González-Haba Ruiz¹

¹*Department of Gastroenterology. Hospital Universitario Puerta de Hierro Majadahonda. Majadahonda, Madrid. Spain.* ²*Department of Gastroenterology. Centro Hospitalar Universitário de São João. Porto, Portugal.*

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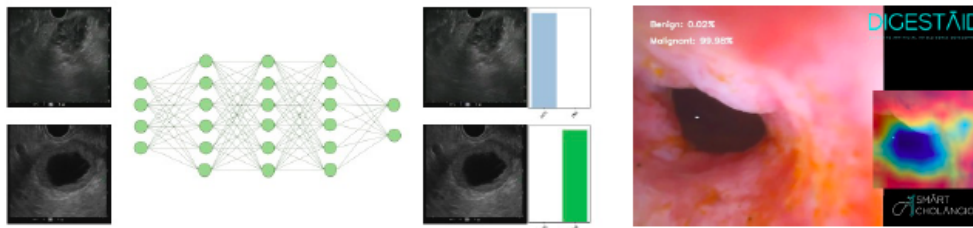
Correspondence: Belén Agudo Castillo. Department of Gastroenterology. Hospital Universitario Puerta de Hierro Majadahonda. C/ Joaquín Rodrigo, 1. 28222 Majadahonda, Madrid. Spain

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ABSTRACT

The development and implementation of artificial intelligence (AI), particularly deep learning (DL) models, has generated significant interest across various fields of gastroenterology. While research in luminal endoscopy has seen a rapid translation into clinical practice with approved AI devices, its potential extends far beyond, offering promising benefits for biliopancreatic endoscopy, including optical characterization of strictures during cholangioscopy and detection and classification of pancreatic lesions during diagnostic endoscopic ultrasound (EUS). This narrative review provides an update on the latest literature and available studies in this field, serving as a comprehensive guide to the current landscape of AI in biliopancreatic endoscopy, emphasizing technological advancements, main applications, ethical considerations, and future directions for research and clinical implementation.

Keywords: Artificial intelligence. Endoscopic ultrasound. ERCP. Biliopancreatic endoscopy.

INTRODUCTION

Artificial intelligence (AI) involves the use of computers and algorithms to simulate human-like decision-making and problem-solving processes (1). It is widely considered

a revolutionary tool with the potential to transform medicine (2). Imaging-based specialties have emerged as leaders in AI model development (3). Gastroenterology, in particular, stands to benefit significantly from AI advancements, potentially leading to disruptive changes in clinical practice. Deep learning (DL), specifically convolutional neural networks (CNNs), are architectures inspired by the human visual cortex, excelling at image analysis (4). Due to reduced pre-processing requirements and less reliance on prior knowledge, CNNs often outperform other DL models in lesion detection and differentiation (5). Increased computational power and the development of sophisticated CNNs have driven an exponential growth in AI-related research within gastroenterology, as shown in recent reviews, highlighting the evolving applicability of AI in gastrointestinal endoscopy alongside relevant regulatory and ethical considerations for its general implementation in clinical practice (6).

Upper endoscopy was one of the first areas to test the development of AI models (7). In a recent meta-analysis, AI had comparable diagnostic accuracy to expert endoscopists in diagnosing Barrett's esophagus and revealed a non-significant increase in the diagnostic accuracy of esophageal squamous cell carcinoma (8). The role of AI in the diagnosis of early gastric cancer was also studied, showing a non-significant increase in diagnostic accuracy (9) (Fig. 1).

AI-aided colonoscopy is an important study field with focus on the diagnosis of dysplastic adenomatous lesions. A recent systematic review and meta-analysis showed that AI-aided colonoscopy had a significant increase in adenoma detection rate (ADR) and adenomas detected per colonoscopy (10). Interestingly, AI-aided colonoscopy was found to be more useful for endoscopists with lower ADR and shorter inspection time, while in experts it revealed a similar ADR. An important consideration is whether this increase in ADR involves detection of advanced adenomas, as most increased detection rates found in individual studies involved small adenomas and hyperplastic polyps (11,12). Indeed, recent research focuses on the potential impact of AI on ADR within colorectal cancer screening programs, where it could play a significant role (13). In capsule endoscopy (CE), the development of DL models holds disruptive potential in addressing its inherent challenges. The process is not only time-consuming but also susceptible to fatigue and errors (missed frames may lead to missed lesions) (14).

Complex CNNs are available that can automatically detect multiple clinically relevant lesions with a high diagnostic yield, and some are even capable of predicting their bleeding potential (15-18). The scientific progress in this field has advanced from detecting lesions in the small bowel to encompassing both the small bowel and the colon, while the ultimate goal is to turn this assessment into a comprehensive endoscopic approach (19). While further prospective and multicentric studies are necessary, AI-enhanced CE may have a transformative impact in medical practice. It may improve cost-effectiveness, broaden current CE indications (e.g., CE after negative upper endoscopy, in patients presenting with upper bleeding), and even change the current paradigm to include endoscopic oncological screening for GI cancers (20).

Beyond significantly reducing procedure interpretation time, the use of AI algorithms can play a crucial role in other gastroenterology procedures that involve a steep learning curve and suboptimal diagnostic accuracy with high inter-observer variability. This applies to specialized areas within gastroenterology such as the evaluation of biliary stenosis in cholangioscopy, the identification/differentiation of dysplastic precursor lesions in anoscopy, and the assessment of EUS-found pancreatic lesions (21-24). In these cases, the use of DL in the real-world clinical setting would not only help physicians in their decision process but also indicate the most likely location of the lesion, guiding the biopsy/treatment process. Although the current treatment paradigm regarding these lesions still demands histopathological assessment and confirmation, a prospective validation of these technological tools could lead to remarkable changes, perhaps even to considering the omission of biopsies, particularly as multimodal AI technologies progress and encompass both imaging data and other personal health records, assuming that computational processing power will keep pace with this technological advancement. This review offers a state-of-the-art examination of the current research and advancements in AI-assisted biliopancreatic endoscopy.

Initial studies exploring its use in ERCP, EUS and cholangioscopy have shown promising results in identifying key anatomical landmarks during these procedures, as well as in differentiating conditions such as pancreatic cancer, autoimmune pancreatitis, pancreatic cystic lesions, and biliary strictures (Figs. 2 and 3).

ARTIFICIAL INTELLIGENCE-ASSISTED EUS

EUS is a valuable diagnostic tool that is used in a variety of clinical applications, including differentiating benign from malignant pancreaticobiliary disorders, staging gastrointestinal (GI) tract tumors, evaluating subepithelial lesions (SELs), and obtaining diagnostic tissue samples (25). Compared to other endoscopic imaging modalities, fewer studies have investigated the use of AI for EUS image analysis. This can be attributed to several key challenges. Firstly, obtaining EUS images of targets with confirmed histological diagnoses poses greater difficulty when compared to luminal endoscopic techniques, where biopsies are more readily acquired. Secondly, the lower prevalence of pancreatic diseases compared to upper GI or colonic lesions results in a smaller pool of available data for AI model development. Finally, EUS images possess an inherently lower resolution and are susceptible to quality degradation from external factors such as movement artifacts because of patient breathing and heartbeat, which require real-time corrections and recordings by the AI models to compensate for image jitter and shifting.

The application of AI to enhance the differential diagnosis of pancreatic lesions using EUS images represents a pivotal frontier in current research. Despite its established role in pancreatic lesion diagnosis, EUS has some drawbacks including low specificity and operator dependence. AI-assisted EUS has the potential to address these limitations, with studies demonstrating improved diagnostic accuracy and reduced interobserver variability. Multiple studies utilizing support vector machines (SVM), principal component analysis, and neural networks have shown that AI algorithms achieve significantly higher sensitivity and specificity when compared to traditional EUS (26-28).

Pancreatic cancer

Pancreatic cancer (PC) has a poor overall five-year survival rate (12 %) and early diagnosis is crucial to significantly improve survival rate (29). While traditional imaging techniques have limitations, EUS offers a superior sensitivity for detecting small pancreatic lesions (30,31). Recognizing this advantage, researchers are actively exploring the application of AI in conjunction with EUS (EUS-AI) for PC detection

(32-34).

In a recent meta-analysis DL models demonstrated superior performance compared to conventional EUS diagnosis for PC detection, with 95 % sensitivity and 90 % specificity, suggesting a strong potential to improve early detection (33,35). Of particular interest is the accurate differentiation between PC and benign conditions like chronic pancreatitis (CP) and autoimmune pancreatitis (AIP) (36). Tonozuka et al. developed a DL-based computer-assisted diagnosis system to detect PC. Using control images from patients with CP or a normal pancreas, their system achieved exceptional performance with an area under the curve (AUC) of 0.92 and 0.94 (validation and testing, respectively) (26). Similarly, Zhu et al. employed an SVM predictive model built from EUS image parameters to differentiate between PC and CP. Their model demonstrated an average accuracy of 94.2 %, with sensitivity and specificity at 96.3 % and 93.4 %, respectively. While promising, these studies highlight the need for external validation to confirm generalizability (37). Marya et al. successfully employed an EUS-CNN model capable of differentiating AIP from other pancreatic conditions with promising results. Their model demonstrated a sensitivity of 90 % and a specificity of 78 % in distinguishing AIP from all other conditions. Specificity increased to 87 % when considering only AIP versus PC (38). These findings are significant, especially in light of a recent meta-analysis highlighting the limitations of EUS-tissue acquisition in accurately diagnosing AIP (39).

Pancreatic cystic lesions

AI holds promise for the endosonographic diagnosis and characterization of pancreatic cystic lesions (PCLs), an area where traditional EUS faces challenges with low interobserver agreement, especially in distinguishing neoplastic from non-neoplastic PCLs, with an accuracy ranging from 48 % to 94 % (40). Several studies have investigated the application of AI, particularly CNNs, to classify PCLs. Nguon and colleagues highlighted the potential of AI for distinguishing between mucinous and serous cystic neoplasms, achieving an accuracy of approximately 83 %, comparable to visual assessment by endoscopists. While this result is promising, the study's focus was limited to two specific types of cystic lesions (41). One notable retrospective study

using a dataset of 5,505 EUS images demonstrated that a high-precision CNN algorithm could distinguish mucinous from non-mucinous cysts with remarkable accuracy (98.5 %), sensitivity (98.3 %) and specificity (98.9 %), as well as an AUC of 1 (42). AI has also shown potential in predicting malignancy, such as in patients with pathologically confirmed intraductal papillary mucinous neoplasms (IPMNs). Here, the AI model achieved an impressive AUC of 0.91 for diagnosing malignant IPMNs, surpassing the diagnostic accuracy of human experts using pre-operative EUS (43). Machicado et al. explored the potential of AI in combination with EUS-guided needle-based confocal laser endomicroscopy (EUS-nCLE) for advanced IPMN diagnosis and risk stratification. Compared to established guidelines, their AI-assisted approach demonstrated higher sensitivity and accuracy with comparable specificity (44). These findings suggest that AI-assisted EUS holds significant potential to revolutionize PCL risk stratification, aiding clinical decision-making and guiding follow-up strategies.

Within the field of AI-Assisted EUS, studies have explored the integration of AI with various techniques, including elastography, contrast-enhanced harmonic EUS (CH-EUS), and the assessment of cytology and histology samples obtained via fine-needle aspiration (FNA) or biopsy (FNB) (45-48).

An early prospective trial investigated the use of EUS elastography images, converted to vector data and analysed with simple neural networks. Despite a small sample size (necessitating 10-fold cross-validation), this achieved an AUC of 0.93 in classifying malignant tumors (49). In 2012 a larger multicentre blinded study (258 patients) validated the approach, outperforming hue histogram analysis (45).

Systems like CH-EUS MASTER use deep CNNs and random forest algorithms for real-time pancreatic mass diagnosis and biopsy guidance. It provides real-time mass identification/tracking, differentiates PC from CP using perfusion analysis and, by utilizing real-time feedback provided by the endoscopists throughout the procedure, the system aids in the selection of the most suitable type and size of puncture needle, offers guidance on optimal location and evaluates the quality of the obtained sample. Consequently, this integration of AI technology has the potential to decrease the number of punctures necessary to acquire an adequate sample, to enhance puncture precision, and to mitigate the likelihood of complications (50). Udristoiu et al.

advanced machine learning in this field by integrating a more complex approach, enabling the model to consider temporal data from contrast-enhanced imaging alongside other image types. Five image sets were extracted per EUS exam (greyscale, color Doppler, CH, elastography) and the results demonstrated 96.4 % specificity and 98.6 % sensitivity overall (51).

Advancements in rapid on-site evaluation (ROSE) aim to improve diagnostic yield and accuracy. Lin et al. developed an AI-ROSE model as potential substitute for manual ROSE during EUS-FNA. While the model demonstrates promise, its current sensitivity (under 80 %) indicates a need for refinement (52). An ideal AI-ROSE system should not only identify malignancy but also accurately assess sample adequacy. To maximize practicality, potential solutions include smartphone-based algorithms for rapid analysis or telepathology options for remote cytopathologist expertise (Table 1).

Table 1. Summary of studies on DL-assisted endoscopic ultrasonography in pancreatic disease

Field	Study	Patient population (n)	Objective	AI model	Outcomes
Pancreatic cancer	Saftoiu et al. 2012 (45)	PC (211); CP (47)	Differentiate cancer from benign masses	Multi-layered perceptron	Sensitivity = 87.59 %. Specificity = 82.94 %. AUC = 0.94
Pancreatic cancer	Udristou et al. 2021 (51)	PC (30); CP (20); pNET (15)	Diagnose focal pancreatic mass	Convolutional neural network and long short-term memory	Sensitivity = 98.6 %. Specificity = 97.4 %. AUC = 0.98
Pancreatic cancer	Tonozuka et al. 2021 (26)	PC (76); CP (34); Control (29)	Differentiate pancreatic cancer from chronic pancreatitis and normal pancreas	Convolutional neural network and pseudo-colored heatmap	Sensitivity = 92.4 %. Specificity = 84.1 %. AUC = 0.94

Pancreatic cancer	Kuwahara et al. 2023 (36)	PC (524). Non-cancer patients (170)	Differentiate pancreatic cancer from non-cancer pancreatic lesions	Deep convolutional generative adversarial network	Sensitivity = 94 %. Specificity = 82 %. AUC = 0.90
Pancreatic cystic lesions	Kurita et al. 2019 (53)	Mucinous cystic neoplasms (23); serous cystic neoplasms (15); IPMN (30); other cyst types (17)	Differentiate benign from malignant cyst	Multi-layered perceptron	Sensitivity = 95 %. Specificity = 91.9 %. AUC = 0.96
Pancreatic cystic lesions	Kuwahara et al. 2019 (43)	Benign IPMN (27). Malignant IPMN (23)	Predict malignancy of IPMN	Convolutional neural network	Sensitivity = 95.7 %. Specificity = 92.6 %. AUC = 0.98
Pancreatic cystic lesions	Vilas-Boas et al. 2022 (42)	Mucinous PCL (17); non-mucinous PCL (11)	Differentiate mucinous from non-mucinous PCLs	Convolutional neural network	Sensitivity = 98.3 %. Specificity = 98.9 %. AUC = 1
<p><i>PC: pancreatic cancer; CP: chronic pancreatitis; pNET: pancreatic neuroendocrine tumor; IPMN: intraductal papillary mucinous neoplasm; PCLs: pancreatic cystic lesion.</i></p>					

Other scenarios

Initial applications of AI in EUS focused on pancreatic disorders but recent studies explore its potential for gastrointestinal SEL diagnosis, particularly regarding gastrointestinal stromal tumors (GISTs). A recent meta-analysis of seven studies (2,431 patients) demonstrated that the EUS-AI model employing CNNs achieved superior sensitivity (0.92) and specificity (0.82) in detecting GISTs when compared to conventional endoscopy. Additionally, the model exhibited potential for assessing the malignant risk of GISTs (54).

AI presents practical solutions for optimizing EUS training. A major challenge for novice endoscopists is accurately identifying anatomical structures. Deep-learning systems like BP MASTER tackle this challenge with station classification, segmentation, and

real-time EUS guidance. These tools substantially improve trainee accuracy in recognizing stations and interpreting images, potentially shortening the learning curve (55,56).

ARTIFICIAL INTELLIGENCE IN ERCP

One of the most challenging diagnoses in GI diseases is biliary strictures (BSs). BSs are defined as undetermined when cross-sectional imaging, as well as tissue sampling, are inconclusive or negative (57), and this represents a challenging clinical scenario. Almost 20 % of BSs are of indeterminate etiology at presentation (58).

Conventional sampling techniques such as ERCP-guided brush cytology or forceps biopsies are limited by low sensitivity (45 % and 48.1 %, respectively) and the combination of the two techniques can barely increase said sensitivity (59.4 %) (59). EUS-guided tissue acquisition can significantly improve the diagnosis and sampling of BSs, particularly those located in the distal bile duct. It should be strongly considered as part of any comprehensive BS workup. However, EUS has limitations when strictures are caused by intraductal vegetative lesions, when they are located in the biliary hilum, or when biliary stents are already present.

Digital single operator cholangioscopy (D-SOC) has gained popularity due to recent technological advances, its availability and its advantages, such as allowing a direct visualization of BSs and their surrounding mucosa, and the collection of targeted biopsies. D-SOC is a safe procedure and may be cost effective at initial ERCP in certain situations (60). While D-SOC demonstrates a high success rate in identifying BSs through visual assessment, with sensitivity and specificity rates of 94 % and 95 %, respectively, the accuracy of D-SOC-guided biopsies is lower (61). Sensitivity in this context ranges from 74 % to 80 %, while specificity remains high at 98 % (62,63). This highlights that tissue sampling might not be as reliable for diagnosis as endoscopic direct visualization. Some visual findings have been statistically associated with malignancy, like the presence of neovascularization or dilated tortuous vessels, an irregular or nodular biliary mucosa, tumors or masses, and an irregular surface with ulcerated, infiltrative or friable appearance (64,65). But to date there is suboptimal interobserver agreement among experts for interpreting the visual impression of a BS.

Moreover, some high-risk features can be present in certain benign instances, such as primary sclerosing cholangitis, which can result in false-positive malignant diagnoses. Given the aforementioned limitations in the diagnostic approach to BSs, there has been increasing interest in exploring the potential of AI to overcome them. AI can potentially impact BS diagnosis by providing categorization (i.e., discriminating malignant BSs from non-malignant BSs) as well as by improving a morphologic classification that has scarcely been assessed. A handful of important studies have been published in the last two years evaluating the accuracy of CNN in BSs. In 2022 Saraiva et al. evaluated the performance of their CNN in distinguishing between benign and malignant BSs. With a total of 11,855 images from 85 patients (9,695 malignant strictures and 2,160 benign findings), the model had an overall accuracy of 94.9 %, a sensitivity of 94.7 %, a specificity of 92.1 %, and an AUC of 0.988 in the cross-validation analysis (66). Several important publications emerged in this setting during 2023. One such study by Marya et al. evaluated the accuracy of their CNN for classifying BSs versus traditional ERCP-based sampling techniques. By analyzing 2,388,439 still images from 154 patients, their CNN demonstrated an overall accuracy of 0.906 for CNN-based video analysis, which is significantly higher than that of brush cytology (0.625, $p = 0.04$) or forceps biopsy sampling (0.609, $p = 0.03$). Their occlusion block heatmap analysis demonstrated that the most frequent image feature for a malignant BS was the presence of frond-like mucosa/papillary projections (67). Later, Carlos Robles-Medranda (CRM) et al. developed a new cholangioscopy-based CNN for recognizing neoplasia in indeterminate BSs in pre-recorded videos and real-time D-SOC procedures, and compared the model with cholangioscopy experts and nonexperts using the CRM and Mendoza classifications. This model achieved significant accuracy values for neoplasm diagnosis, with 90.5 % sensitivity, 68.2 % specificity, and 74.0 % and 87.8 % positive and negative predictive values, respectively, and also outperformed the two non-expert and one of two expert endoscopists (21). Simultaneously, Zhang et al. proposed a different model (MBSDeiT) capable of automatically selecting qualified DSOC images with high accuracy (AUC of 0.963-0.973 across internal and external testing data sets) and subsequently identifying 92.3 % of malignant BSs in prospective videos. MBSDeiT outperformed both expert and novice

endoscopists (23).

Finally, Saraiva et al. evaluated their CNN with 84,994 images from 129 D-SOC exams in two centers (Portugal and Spain). The model achieved 82.9 % overall accuracy, 83.5 % sensitivity, and 82.4 % specificity with an AUC and AUPRC of 0.92 and 0.93, respectively. This model additionally showed outstanding performance in detecting tumor vessels and papillary projections, with AUC values of 0.98 and 0.96, respectively (24). AI based on clinical biomarkers like alkaline phosphatase, intrahepatic bile duct diameter, and total bile duct diameter could also serve as an adjuvant for diagnosing malignant bile duct obstruction (68).

AI has also been used to evaluate different aspects of ERCP. A computer-assisted system using CasNet, a segmentation architecture of DL trained on 1,381 cholangiogram images, showed an effective assessment and classification of the degree of technical difficulty in endoscopic stone removal during ERCP (69). Additionally, CNN-based models have been developed to predict the location and cannulation difficulty of the ampulla (70). Machine learning models have also been proposed to predict the likelihood of post-ERCP pancreatitis and to identify new clinical features relevant for this risk (71) (Table 2).

Table 2. Main studies evaluating CNN-based DL for biliary stricture diagnosis

Author (year)	Patient population (n)	Objective	Study characteristics	Main outcomes
Saraiva et al. (2022) (66)	85	Automatic detection of malignant BSs in DSOC images	Pilot validation study	Overall accuracy 94.9 %, sensitivity 94.7 %, specificity 92.1 %, AUC 0.988 for differentiating malignant from benign BSs
Marya et al. (2023) (67)	154	Analyzing DSCO images in real-time to accurately classify biliary strictures	Multicenter validation study	CNN had greater accuracy for biliary stricture classification (0.906) compared to brush cytology (0.625, $p = 0.04$) or forceps biopsy sampling (0.609, $p = 0.03$).
Robles-Medranda et al., (2023) (21)	<i>Phase 1</i> 48 <i>Phase 2</i> 116	Validation of a CNN model for identifying malignancy in indeterminate BSs	International multicenter, two-stage validation study	90.5 % sensitivity, 68.2 % specificity, and 74.0 %, 87.8 % PPV and NPV, respectively, in distinguishing neoplastic lesions.
Zhang et al. (2023) (23)	150	Validation of a novel AI model to identify and predict malignant BSs	Multicenter diagnostic study	MBSDeiT accurately identified 92.3 % of malignant BSs in prospective testing videos
Saraiva et al. (2023)	129	Distinguishing benign from	International multicenter	Sensitivity 83.5 %, specificity 82.4 %,

LIMITATIONS OF ARTIFICIAL INTELLIGENCE AND ETHICAL ISSUES

Employing AI-assisted advancements in gastroenterological techniques, which enable endoscopists to see more and beyond in order to make better decisions, requires careful consideration of precautions to ensure trustworthiness. Technological development should follow the FAIR data principles (72). To maximize AI's potential in healthcare, it must be findable (with clear data labelling and unique patient identifiers), accessible (ensuring transparent data sources for verifying algorithm robustness), interoperable (compatible with various devices for wider use), and reusable (promoting the use of open-source frameworks and allowing datasets to be reused whenever they prove useful in addressing clinical challenges). In addition to complex data acquisition and standardization, privacy concerns arise as data is collected, requiring robust protection. Healthcare blockchain innovations may address this, offering decentralized and secure data frameworks (73,74). Addressing inherent selection biases is also critical to ensure a transparent and transferable AI, which is achievable only through high-quality training data (75).

Furthermore, there are two AI-related ethical challenges that should be considered. One concern is the “black-box” characteristic of these algorithms, implying that AI models can identify patterns (e.g., lesions) imperceptible by physicians (76). Even though understanding how some medical interventions work remains a challenge (for example, how certain drugs improve a patient's outlook without a fully known mechanism), there is a stronger resistance to AI making decisions in medicine without any human involvement (77). The second concern tends to arise as a consequence of the first one. If an AI model identifies a lesion that a physician disagrees with, and it turns out there was a lesion, should the doctor be held responsible? Conversely, if the machine fails to detect a lesion that later is found (e.g., false negative), who should be accountable: the doctor or the AI model development company? FDA is currently approving computer-aided detection and diagnosis (CADe and CADx, respectively) systems as “Software as a Medical Device” (SaMD). SaMD clearance implies that such technology aids in detecting clinically relevant lesions but does not make a diagnosis,

with the ultimate responsibility lying with the physician (78). Currently, there are three commercially approved technologies: Gi Genius® by Medtronic™, SKOUT® by Iterative Health™, and Veritai® by Satisfai Health™.

The development of AI models is crucial for enhancing the diagnostic capabilities of EUS. However, to ensure widespread clinical use, AI models must work accurately across different EUS devices. A significant concern in the development of DL models is the possibility of an imbalanced dataset that is not adapted to the population in which the technology will be used, thus limiting the external validity of the results.

The use of AI-based endoscopic imaging for the diagnosis of BSs has several potential clinical benefits that include reducing tissue sampling techniques, resulting in fewer procedures and their associated costs and adverse events, and also reducing the paradoxical gap between visual impression and histology. It can also have an academical benefit, providing expert and non-expert endoscopists a second opinion on lesions suggestive of neoplasia, helping to obtain a targeted sample, and reducing the current suboptimal level of interobserver agreement.

While promising, robust clinical adoption of AI necessitates further development and rigorous external validation. Currently, a significant limitation is reliance on relatively small datasets, hindering the ability of algorithms to generalize across diverse patient populations. Continued research utilizing larger, more comprehensive datasets is crucial for ensuring reliable performance. Understanding real-world benefits, addressing ethical considerations, adherence to FAIR data principles, and a focus on clinical validation hold the key to revolutionizing diagnostic accuracy, optimizing interventions, and ultimately improving patient outcomes. Despite these challenges, the potential of AI in this field remains undeniable.

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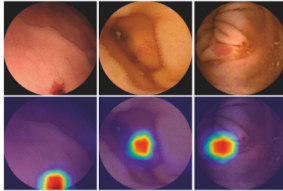
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Main types of AI used in digestive endoscopy

IMAGE RECOGNITION:

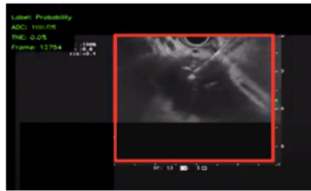
AI algorithms analyze endoscopic images to identify abnormalities.



e.g. AI-detects vascular lesion in stomach, small bowel or colon.

HISTOLOGY PREDICTION:

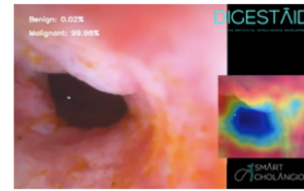
AI systems can predict histopathological features.



e.g. Probability of adenocarcinoma is shown in upper left quadrant.

REAL-TIME GUIDANCE:

AI can assist in real-time procedures and reduces exploration time.



e.g. AI-assisted cholangioscopy for guided biopsies.

Fig. 1. Main types of AI used in digestive endoscopy.

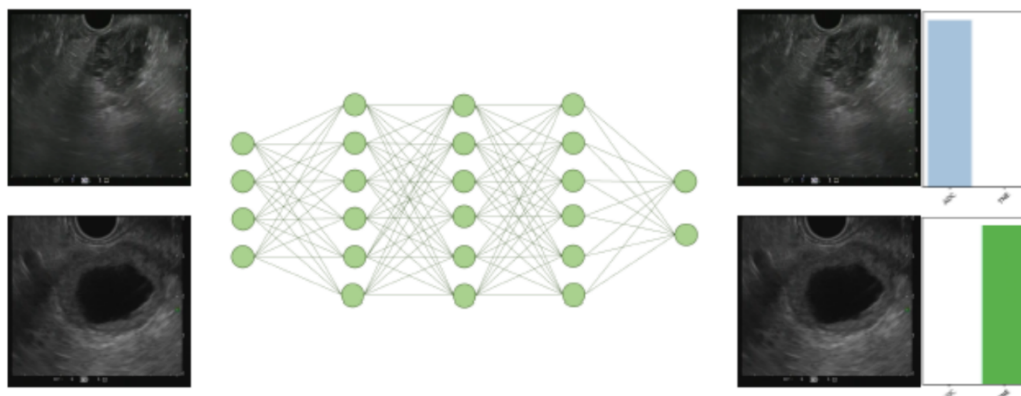


Figure 2. Output provided by artificial intelligence for distinguishing between various types of solid pancreatic lesions during an endoscopic ultrasound (EUS) examination. The bars depict the probabilities estimated by the network. ADC - pancreatic adenocarcinoma or TNE - pancreatic neuroendocrine tumor

Fig. 2. Output provided by artificial intelligence for distinguishing between various types of solid pancreatic lesions during an endoscopic ultrasound (EUS) examination. The bars depict the probabilities estimated by the network (ADC: pancreatic adenocarcinoma or TNE: pancreatic neuroendocrine tumor).

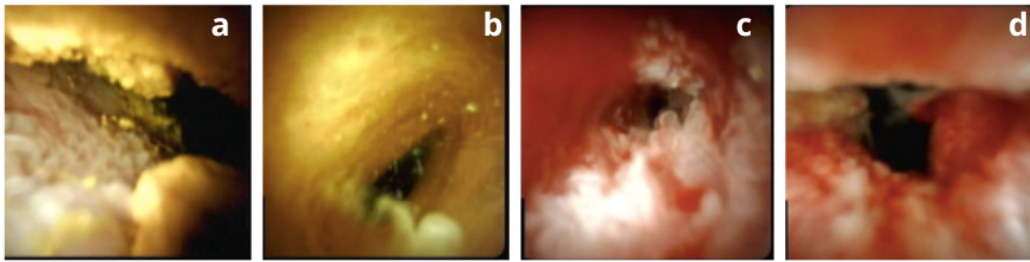


Figure 3.A. Images of benign (a,b) and malignant (c,d) biliary stenosis in a digital single-operator cholangioscopy.

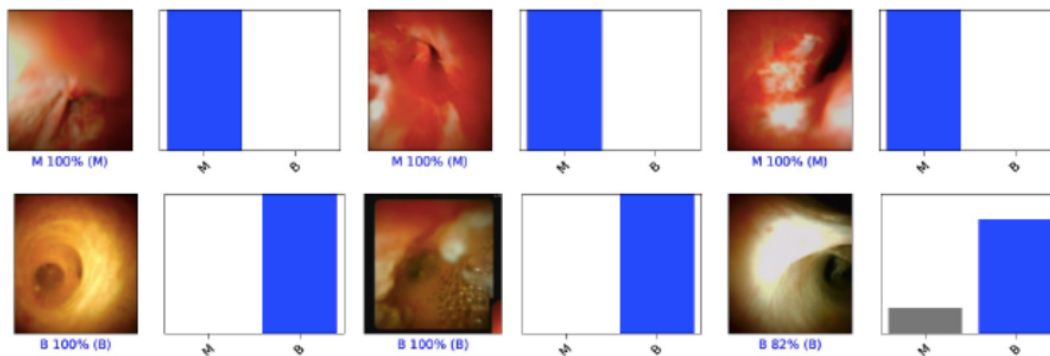


Figure 3.B. Output generated by the convolutional neural network for biliary stenosis in cholangioscopy. The bars represent the probability estimated by the model. M-Malignant, B-Benign.

Fig. 3. Images of biliary stenosis in digital single operator cholangioscopy (A). Output of a convolutional neural network for differentiation between malignant and benign biliary stenosis (B) (M: malignant; B: benign).