



Electronic chromo-endoscopy: technical details and a clinical perspective

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Abstract: Precise endoscopic assessment is necessary to detect neoplastic changes in an early stage. Electronic or virtual chromo-endoscopy (ECE) is an alternative to conventional dye-based chromo-endoscopy which markedly improves capillary pattern and hence can detect micro-vessel morphological changes of early neoplasia to target biopsies and aid in diagnosis. The clinical significance increased after the advent of endoscopic treatment modalities like ESD/EMR which requires precise delineation of extent and depth of lesion. Most of the studies have used narrow-band imaging (NBI) (Olympus Medical Systems Tokyo, Japan), although data from i-SCAN (PENTAX Endoscopy, Tokyo, Japan) and flexible spectral imaging color enhancement (FICE) (Fujinon, Fujifilm Medical Co, Saitama, Japan) are emerging. Electronic chromo-endoscopy is convenient compared to dye-based chromo-endoscopy in the sense that it is available at the push of a button in endoscope and reduces procedure time substantially with comparable efficacy. Scope of this review is to discuss available electronic chromo-endoscopy modalities and their role in the diagnosis, surveillance, and management of early GI neoplasia.

Keywords: Electronic chromoendoscopy; narrow-band imaging (NBI); I-SCAN; flexible spectral imaging color enhancement (FICE); Blue laser imaging (BLI); Linked colour imaging (LCI)

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Introduction

White light endoscopy was an important milestone in gastroenterology enabling not only detection of lesion, assessment of length, extent and severity but also allowing biopsy and endoscopic management of diseases. However standard white light endoscopy (WLI) may miss the subtle lesions of superficial neoplasia specially the early cancer lesions which are usually flat and may appear isochromatic to surrounding mucosa on WLI (1).

Dye based chromo-endoscopy was the first step towards image enhancement. This involved endoscopy with spraying of various dyes on the mucosa in order to facilitate visualization of benign versus cancerous mucosal changes. Several staining agents broadly categorized as

absorptive (vital) stains, contrast stains, and reactive stains were used. Absorptive stains (e.g., Lugol's iodine solution and methylene blue) diffuse or are preferentially absorbed by specific epithelial cell membranes; contrast stains (e.g., indigo carmine) highlight surface topography and mucosal irregularities by permeating mucosal crevices & reactive stains (e.g., Congo red and phenol red) undergo chemical reactions with specific cellular constituents, resulting in a color change. This improved the contrast and was a useful adjunct to routine WLI (2).

Dye based Chromo-endoscopy has been widely used for various indications throughout the gastrointestinal tract (*Table 1*). However, the technique is often operator dependent and requires additional procedure time including

Table 1 Electronic chromoendoscopy and clinical applications

Scope of electronic chromoendoscopy	Clinical applications
Oesophagus	Barrett's oesophagus (<i>Figure 1A, 1B</i>) and Barrett associated CA Minimal change oesophagitis (<i>Figure 1C</i>) Early oesophageal carcinoma (<i>Figure 1D-1F</i>)
Stomach	Intestinal metaplasia (<i>Figure 2</i>) Helicobacter pylori infection Early gastric cancer (diagnosis) Gastric polyps
Duodenum	Celiac disease Ampullary dysplasia or duodenal carcinoma Familial adenomatous polyposis
Small intestine	Angiodysplasia
Biliary tract	Early detection of cholangiocarcinoma Differentiation between benign & malignant biliary strictures
Colorectum	Detection & characterization of colonic polyps Inflammatory bowel disease (IBD): dysplasia detection, mucosal healing

technical expertise in preparing and using the dye (3). The standardization on contrast dilution, dilution methods, concentration and the depth of the color hue, remains unclear. It requires additional instruments like the spray catheter and the dye may not spread evenly across the mucosal surface.

Additionally, switching between white light view and chromo-endoscopic view is not possible. The vascular microstructure is often difficult to assess after spraying dye and can be distorted by the use of dyes like acetic acid which causes vascular congestion. Methylene blue can precipitate met-hemoglobinemia in patients with Glucose-6-Phosphatase dehydrogenase deficiency (4). Moreover, the risks of aspiration pneumonitis particularly in sedated elderly patients limits its usage.

Cost effectiveness of dye spray chromo-endoscopy in clinical practice has not been established especially with the advent of high definition endoscopes (5). There is also concern regarding accelerated carcinogenesis with the use of methylene blue due to photo activation in Barrett's oesophagus (BE) screening (6).

Digital chromo-endoscopy in the form of narrow band imaging (NBI), Fuji intelligent chromo-endoscopy (FICE) and can overcome most of these limitations and is now preferred in clinical practice particularly for the early detection of cancers.

Definition of electronic chromoendoscopy

Electronic chromo-endoscopy or digital chromoendoscopy refers to advanced endoscopic imaging technologies that provide detailed contrast enhancement of the mucosal surface and blood vessels based on the principle of observing light transmittance at selected wavelength given interaction of particular tissue structures with light is wavelength dependent. The advantages of electronic chromo-endoscopy are ease of use, short learning curve, no need of special assembly or dye and shorter procedure time (7).

Electronic chromoendoscopy: technical details

ECE uses the principle of selective light transmittance. This is achieved by either optical filtering of white light (Narrow Band Imaging) or software driven post image processing (FICE, I-scan). These are hardware based first generation image enhanced endoscopy (IEE) systems. The major drawbacks of these systems are low resolution and dark images at distant view. This is overcome by blue laser imaging (BLI), which was released by Fujifilm corporation in 2013. BLI provides bright images at distant view which can improve colorectal polyp detection rate (8). Further improvement in image enhancement is achieved by novel techniques such as linked colour imaging (LCI) developed by Fujifilm corporation (Tokyo, Japan) (9).

NBI

NBI is an endoscopic optical image enhancement technology developed by Olympus Medical Systems. It's based on the penetration properties of the light which is directly proportional to the wavelength (10). NBI filter in front of the xenon arch lamp produces 2 narrow bands centered at 415 and 540 nm. These 2 wavelengths correspond to the primary and secondary light absorption peaks of hemoglobin, respectively (11). Capillaries in the superficial mucosa are highlighted by the 415-nm wavelength and appear brown. The longer 540-nm penetrates more deeply and make the deeper veins appear blue-green (cyan) thus creating contrast with the

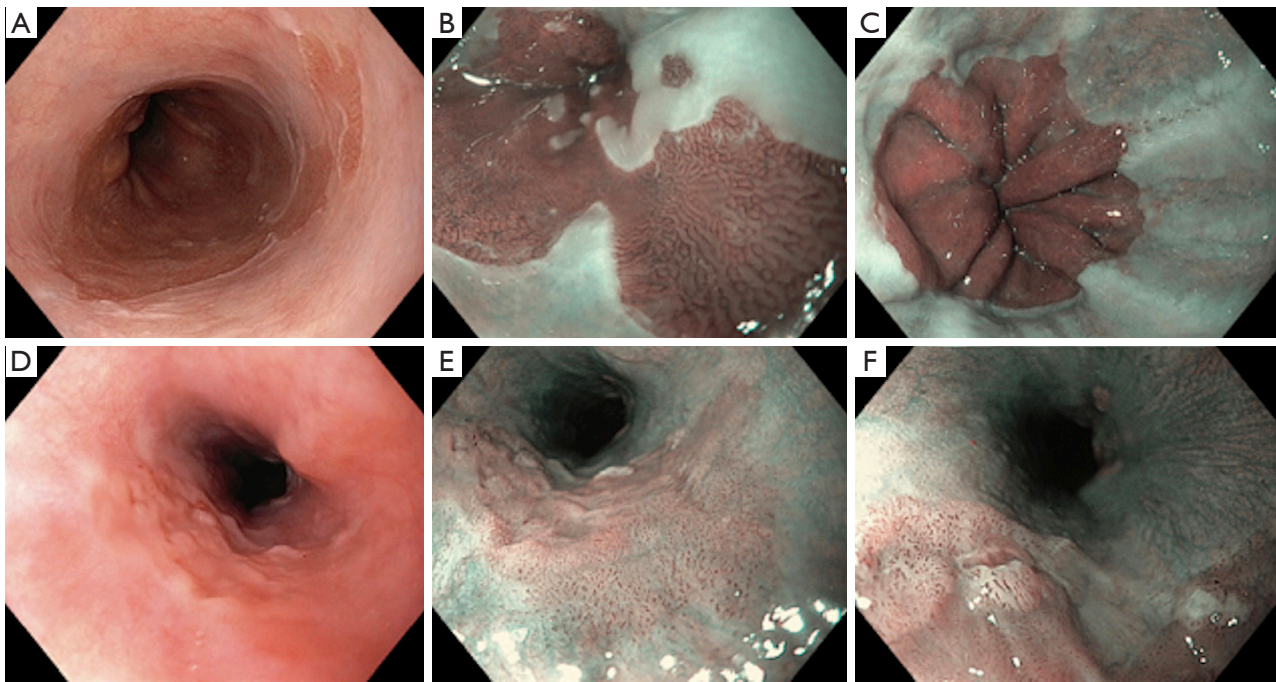


Figure 1 Role of narrow band imaging (NBI) in esophagus (Barrett's oesophagus, minimal change esophagitis and early oesophageal cancer). (A) Barrett's esophagus on white light imaging (WLI); (B) Barrett oesophagus on NBI: regular ridged pit pattern with normal micro-vasculature with no evidence of dysplasia; (C) minimal change esophagitis: dilated intra-papillary capillary loop pattern (IPCLs) type II. These are enlarged but arranged in a linear regular fashion; (D) early oesophageal cancer on WLI: mild nodularity on careful inspection; (E) early oesophageal cancer: NBI image, Brownish discoloration with irregular dilated IPCLs; (F) early oesophageal cancer: NBI with magnification showing small ulcerated area with Type IV IPCLs.

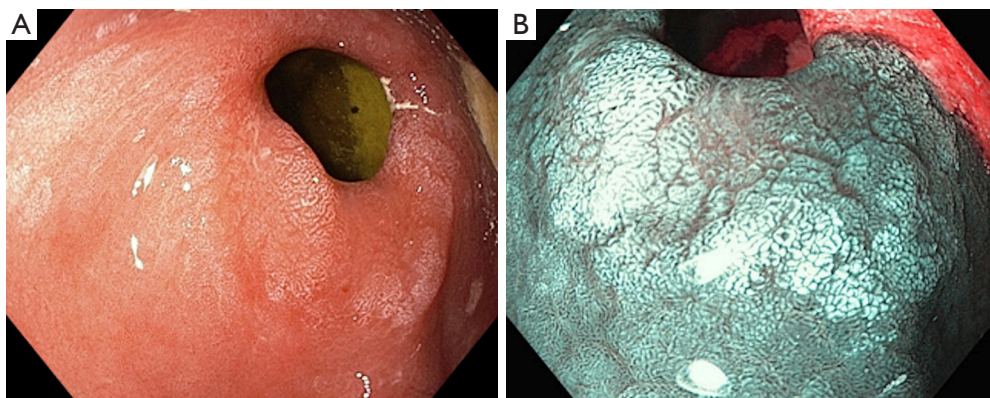


Figure 2 Gastric intestinal metaplasia. (A) White light imaging and (B) narrow band imaging.

nonvascular structures in the mucosa (*Figure 3*) (12,13).

Currently available video processors and light sources are next generation Evis Exera III released in 2012 which produces brighter image compared to initial commercially available NBI systems such as Evis Exera II 180 system

(color CCD system) and the Evis Lucera 260 spectrum series (RGB sequential system). Second generation NBI systems have corrected the issue of darker image in earlier generation by improvements in the light source, system's lenses and mirrors. These modifications have increased

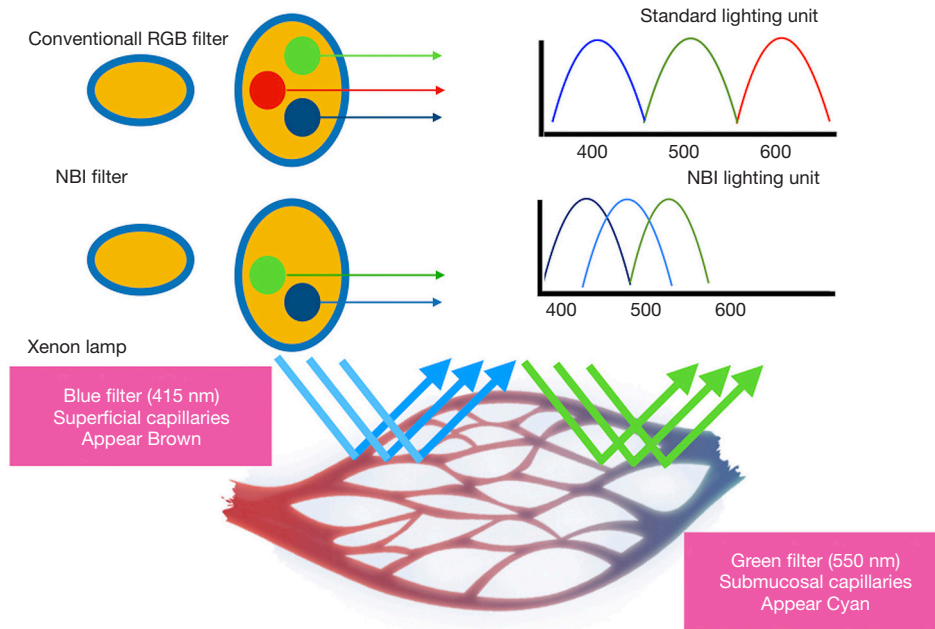


Figure 3 Principles of narrow band imaging.

the brightness of the image by minimising lamp light permeating from the glass fibre within the endoscope and 3D noise reduction capability (12).

FICE

FICE is a proprietary digital post-processing system of Fujinon (13). FICE is based on a Spectral Estimation Technology invented by Professor Yoichi Miyake from Tokyo. The Captured image is sent to the Spectral Estimation Matrix processing circuit in the EPX-4400 where various pixilated spectrums of the image are estimated. Since the spectrums by pixels are known, it is possible to implement imaging on a single wavelength. Such single wavelength images are randomly selected, and assigned to R (Red,) G (Green,) and B (Blue) respectively to build and display a F.I.C.E. enhanced color image (8,13).

Ten factory-determined presets are available. Each preset can be button activated from the computer keyboard and can be changed manually to possible permutations of the available wavelengths (from 400–695 nm) that can be manipulated in 5-nm increments. The endoscopy push button can be programmed to enable switching between the conventional white-light image and up to 3 FICE presets. However, the major limitation of the FICE system is that the optimal settings have not been clearly established (13).

i-SCAN

i-SCAN is a digital, post processing image enhancement technology from PENTAX Endoscopy similar to FICE that provides digital contrast to endoscopic images. There are 3 i-SCAN modes -1/2/3. They comprise of surface enhancement (SE-enhanced mucosal surface texture), Contrast enhancement (CE-sharpened views of surface vessels), Tone enhancement (TE-increases the contrast between the mucosa and blood vessels improving visibility of blood vessels) and Tone enhancement mode g (improves visualisation of grimly lit far-field regions) which functions by performing per pixel modifications of the white light image. i-SCAN 1 has CE and SE mode. i-SCAN 2 has CE, SE and TE mode whereas i-SCAN 3 has CE, SE, TE g mode (Table 2). i-SCAN 3 differs from i-SCAN 2 primarily in its ability to illuminate more distant regions better. Unlike NBI, red remains the predominant blood vessel color in all i-SCAN modes. Switch from WLE to i-SCAN occurs almost instantly. PENTAX processors with i-SCAN capability are EPK-i5010 with LH-150PC as the light source (8,14).

BLI

BLI uses monochromatic lasers (410 and 450 nm) in place of xenon light (NBI). The 410 nm laser acts like NBI with shorter wavelength which highlights the micro-vasculature

Table 2 Different I-SCAN modes and their functions

Mode	Enhancement	Functionality
i-SCAN (Modes off)	SE/CE/TE -off	High definition white light endoscopy
i-SCAN 1	SE/TE	Sharpens surface vessels and enhances surface texture of mucosa
i-SCAN 2	SE, CE, TEc	Increases contrast between mucosa and blood vessels, enhances surface texture of mucosa
i-SCAN 3	SE, CE, TEG	Increases contrast between mucosa and blood vessel including dimly illuminated far-field regions, enhances surface texture of mucosa

whereas 450 nm laser acts by exciting phosphors to create white light which provides white light illumination. Ratio of the intensity of the two lasers can be changed to provide a illumination for both normal and narrow band observation. Furthermore, brighter images can be derived by using optical filters (15,16).

BLI has 3 modes: (I) BLI mode- for detailed magnified observation; (II) BLI bright mode- both vascular and surface pattern analysis even from distant view; (III) white light mode- uses laser (450 nm) rather than conventional white light which better enhances the micro-vasculature. Most colorectal polyps can be diagnosed with good accuracy by BLI magnification. Hiroshima classification can be used to delineate polyps by BLI as it uses both surface and vascular pattern. According to this classification, A depicts non-adenomatous polyps, B depicts adenomatous polyps, C1 depicts intra-mucosal carcinoma, C2 depicts slight submucosal invasion (<1,000 µm) and C3 depicts deep submucosal invasion (>1,000 µm). According to a study BLI was accurate for all the subtypes except for C2 which requires conventional chromo-endoscopy with pit pattern analysis for better characterisation (15,16). Blue Light Imaging using LED light source (Eluxeo system) that is now available in Europe, US and Asia/Australia. The original BLI using Laser light (LASEREO) is commercially available in Japan and Asia-Pacific (17).

LCI

LCI is a novel IEE technique which increases image contrast. Although novel modalities like BLI produces brighter images, it is still criticized for dark image contributing to low diagnostic rate. BLI uses green and blue light for red colour adjustment like NBI. Hence LCI has been introduced. It uses information of all three colours for signal processing. LCI differentiates the red colour spectrum more effectively than WLI. The increased colour contrast improves detection of inflammation and results in more accurate delineation. By

unique image processing, it makes red regions more vivid and white region clear white (9).

There is substantial evidence to suggest role of LCI in detecting early gastrointestinal neoplasms by enhancing color contrast between neoplasm and surrounding mucosa (18). This distinction is made by specific color allocation based mainly on the distribution of capillaries. It helps in early detection of pale and superficial neoplasms providing sufficient brightness to illuminate wide lumen by novel image processing which is a distinct advantage over NBI and BLI. Early gastric cancers and intestinal metaplasia appear orange-red and purple on LCI respectively (19). In colon, LCI improves the adenoma detection rate and decreases the polyp miss rate (20).

Role of NBI in BE

NBI sharpens the visual fields while inspecting squamo-columnar junction and can help detect Barrett's and dysplasia (Figure 1A,1B). Various studies have compared HD-WLE with HD-NBI for detection of Barrett's and associated dysplasia (21,22). A Randomized crossover study by Kara *et al.* showed that HD-WLE and HD-NBI are comparable in detecting high grade dysplasia (HGD) with a sensitivity of 79% and 86% respectively (21). But a multi-center, randomised study by Sharma *et al.* compared the two and have found that NBI-HD significantly detects more number of HGD (30% *vs.* 21%, P=0.01) and require less number of biopsies (3.6% *vs.* 7.3%, P<0.0001) compared to HD-WLE (22). In a meta-analysis, sensitivity, and specificity of HD-NBI in diagnosing, HGD is 96% and 94% respectively and, specialized intestinal metaplasia (SIM) is 95% and 65% respectively (23). This is supported by other studies and meta-analysis showed better detection of oesophageal dysplasia and cancer with NBI compared to WLE (24). There are also studies on reproducibility of NBI findings. However, the results showed moderate inter observer agreement and that the inter-observer agreement

was not improved by NBI over HD-WLE (25).

Using NBI, different classifications have been proposed. Classifications used for BE with magnification narrow band imaging or electronic chromo-endoscopy are listed (Table 3) (25-28).

Role of I-SCAN in BE

The role of I-SCAN in detecting dysplasia in Barrett's esophagus is evolving with influx of new data. I-SCAN post processing endoscopic imaging has been shown to be superior to white-light endoscopy in the detection of intestinal metaplasia in BE in previous studies (29,30). In a recent study by Lipman *et al.*, addition of acetic acid and Magnification imaging (PENTAX EG 2990 Zi magniview endoscope) to I-SCAN significantly increased the accuracy of the I-SCAN classification system (69% to 79%, $P=0.01$). BE associated dysplasia detection rate was significantly improved compared to HD-WLE and I-SCAN alone (83% vs. 76%; $P=0.047$). Inter-observer agreement of the classification system was moderate, although it improved to substantial when incorporated into a clinical decision-making protocol ($\kappa=0.69$). Small subgroup analysis of LGD lesions showed a comparable accuracy of 80% (28). In spite of that, it did not meet the ASGE PIVI threshold for adopting this instead of current practice of random biopsies given limitations of small sample size, low prevalence of dysplasia in examined cohort and use of videos rather than still images (30,31).

A further modification of the iSCAN system by optical enhancement (OE) uses both pre and post processing images which can improve dysplasia detection rate without the addition of ACA, thus reducing procedure time (32). as shown in study by Everson *et al.*, which also showed that prediction of histology was better with iSCAN OE compared to HD-WLE by both expert and trainee endoscopists (33).

Role of FICE in BE

There is very limited data on the use of FICE in BE, A small pilot study including 18 patients with acetic acid application, HGD was identified in 100% patients as compared to 14% using HD-WLE (34). FICE channel 4 images were significantly better than the conventional images (35). In a study consisting of 40 patients comparing FICE and WLE, visualization of palisade vessels and demarcation of BE and gastric mucosa was better made by FICE compared to

WLE (36). Pohl *et al.* compared chromoendoscopy with acetic acid application (CAA) with FICE for detection of HGD in BE in a randomised cross over trial and found that sensitivity of both the modalities were similar (87%) and found out sensitivity at around 87%. Sensitivity of directed biopsies for detection of neoplasia was 83% and 92% for CAA and FICE respectively, although the difference was not statistically significant (37).

Electronic chromo-endoscopy in esophageal squamous cell carcinoma (ESCC)

Electronic chromoendoscopy has enabled early detection of ESCC by predicting of depth and extent of the lesion, as submucosal invasion increases probability of lymph node metastasis. Esophageal stratified squamous epithelium is featureless with no pit pattern and regular palisading capillary network (Figure 1D,1E). Esophageal microvascular structure including intrapapillary capillary loop pattern (IPCL) can be visualized with clarity using NBI. IPCL is a smooth running small diameter (10 μm) capillary vessel positioned upright from a branching vessel. Branching vessels appear green while the IPCLs are observed as dark brown loops/dots on NBI. Minimal mucosal changes are easily missed by WLE because of flat and isochromatic appearance of superficial ESCC (38). Chai *et al.* demonstrated usefulness of HD-NBI compared to HD-WLE in detecting ESCC (adenoma detection rate 70.2% vs. 35.7% respectively, $P<0.01$). Most of the missed lesions were high grade intraepithelial lesions (39). In a recent systematic review and meta-analysis including 1,911 patients by Morita *et al.*, NBI fared better than conventional chromoendoscopy with Lugol's iodine to diagnose high grade intraepithelial neoplasia and ESCC (40).

Most widely used ME-NBI classification is by Inoue *et al.* (Figure 4) but due to its complexity it's not widely popular among the endoscopists (41).

A simplified NBI with magnifying endoscopy (NBI-ME) criteria (dyad criteria) by Dobashi *et al.* defined the lesion endoscopically as superficial ESCC based on 6 NBI-ME findings: inter-vascular background coloration, proliferation of IPCLs, dilatation/tortuosity/change in calibre/various shapes of IPCL (42). It was validated in a prospective comparative trial of 170 patients at high risk of ESCC, 77 were detected to have superficial ESCC with better sensitivity, by DF-NBI than WLI (91% vs. 51%, $P<0.001$) with high inter (0.77)/intra (0.82) observer agreement (43).

JES (Japanese Esophageal Society) has developed

Table 3 NBI Endoscopy—Barrett’s oesophagus classification systems

Morphological classifications and their performance	Kansas (25)	Nottingham (26)	Amsterdam (27)	Barrett’s International NBI group (BING) (28)
Morphology				
1) Normal	Mucosal pattern (MP): circular Vascular pattern (VP): normal	Type A: round/oval pits with regular microvasculature (MW)	MP: regular VP: regular Abnormal blood vessels (ABV): absent	MP: circular, ridged/villous, or tubular VP: blood vessels situated regularly along or between mucosa ridges and/or those showing normal, long, branching patterns
2) Intestinal metaplasia	MP: ridged/villous VP: normal	Type B: villous/ridge/linear pits with regular MW Type C: absent pits with regular MW	MP-regular VP-regular (villous/gyrus) ABV: absent	Not tested
3) Dysplasia	MP: irregular/distorted VP: abnormal	Type D: distorted pits with irregular MW	MP-irregular VP: irregular ABV: present	MP: absent or irregular patterns VP: focally or diffusely distributed vessels not following normal architecture of the mucosa
Performance of various classifications	Sensitivity, specificity, & Positive predictive value (PPV) of ridge/villous pattern for diagnosis of intestinal metaplasia (IM) without high grade dysplasia (HGD) were 93.5%, 86.7%, and 94.7%, respectively	87.9% in magnification endoscopy corresponded to histological diagnosis	The magnified NBI images had a sensitivity of 94%, a specificity of 76%, a PPV of 64%, and a NPV of 98% for HGIN	High confidence
	The sensitivity, specificity, and PPV of irregular/distorted pattern for HGD were 100%, 98.7%, and 95.3%, respectively	PPV & negative predictive value (NPV) for type A (Columnar mucosa without IM) was 100% & 97% respectively For Type B & C (IM) was 88% and 91% respectively For Type D (high grade dysplasia) was 81 & 99% Inter and Intraobserver agreement was Kappa of 0.71–0.87 (Nonexperts) and 0.78–0.91 (Experts)	No data on observer agreement	predictions by the experts yielded high accuracy sensitivity, and specificity (92%, 91%, and 93% respectively) Inter-observer agreement was substantial (k=0.681)

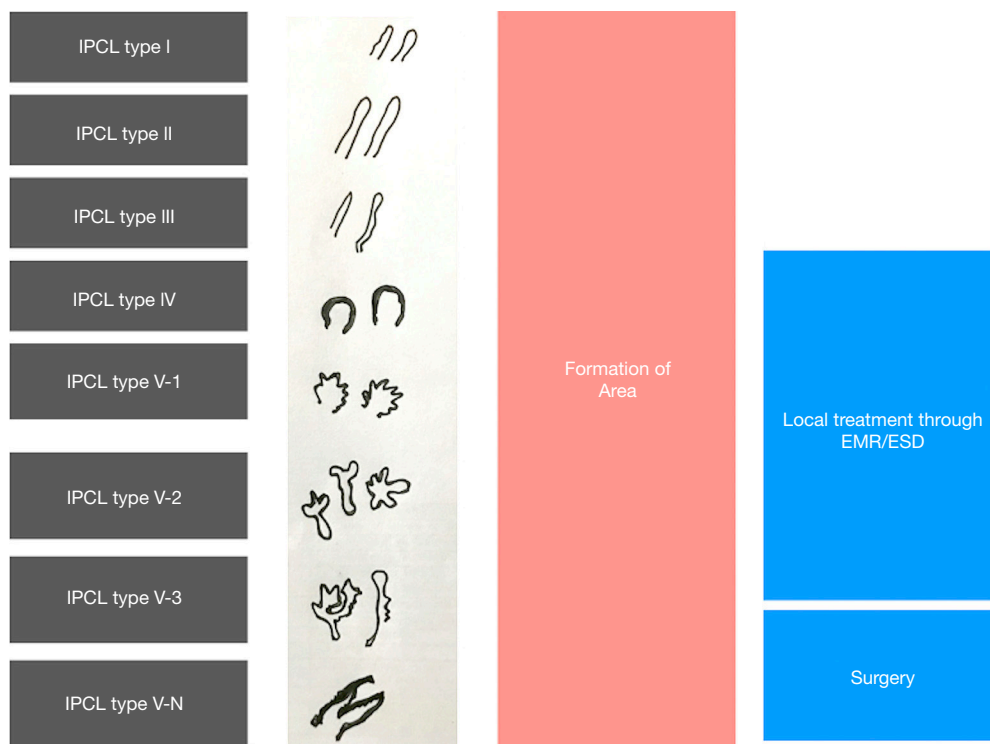


Figure 4 Original intra-papillary capillary loop pattern (IPCL) classification. Adopted from Inoue *et al.* (37).

Table 4 Modified from Oyama *et al.*—criteria of the Japanese Endoscopic Society (JES) magnifying endoscopic classification (44)

Type of vessels	Scheme	Definitions	Invasion depth	Histology
A		Normal intrapapillary capillary loop pattern (IPCL) or abnormal micro vessels (7–10 μm) without severe irregularity	No invasion	Normal epithelium, inflammation, and low-grade intraepithelial neoplasia
B1		Abnormal micro vessels with severe irregularity or dilated abnormal (20 μm) vessels, Type B vessels with loop like formation	T1a-Epithelium, T1a-Lamina propria mucosa	High grade intraepithelial neoplasm or invasive squamous cell carcinoma
B2		Type B vessels with out a loop like formation	T1a-MM (muscularis mucous) or T1b-SM1 (submucosa)	
B3		Highly dilated blood vessels (60 μm) with calibers >3 times B2 vessels	T1b-SM2 or deeper	

a simplified magnifying endoscopic classification for estimating invasion depth of superficial esophageal squamous cell carcinomas (Table 4) (44). Overall accuracy of type B1, B2, B3 microvessels to diagnose invasion is 90.5% which is fairly good.

Standardization of the ME-NBI reports is necessary to develop consensus among the endoscopists. NBI

should be used routinely for surveillance and diagnosis of superficial ESCC. FICE can be complementary to lugol chromoendoscopy and magnifying endoscopy for detecting ESCC due to better observation of IPCLs compared to conventional chromoendoscopy. I-scan usage in the management of squamous cell carcinoma of the oesophagus is scarce, routine use cannot be recommended at present.

Table 5 Gastric lesions classification with narrow band imaging (40)

Morphology and diagnosis	A	B	Hp+ (H-pylori infection)	C
Mucosal pattern	Regular circular	Regular ridge/tubulovillous/ Light blue crest*	Regular	Irregular or absent white opaque substance on the surface**
Vascular pattern	Regular thin/peripheral (gastric body) or thick/ central (gastric antrum) vessels	Regular	Regular with variable vascular density.	Irregular
Diagnosis	Normal	Intestinal metaplasia	H pylori infection	Dysplasia

*, "Light blue crest", defined as fine, blue line on the crest of the epithelial surfaces or gyri, being highly specific for the diagnosis of intestinal metaplasia; **, "White opaque substance" above the mucosa can be associated with dysplasia and cancer.

Electronic chromo-endoscopy in Stomach-Gastric Intestinal Metaplasia, dysplasia, and early gastric cancer

Electronic chromo-endoscopy has its primary utility in diagnosing early gastric cancer and certain premalignant conditions (e.g., gastric intestinal metaplasia) (*Figure 2, Table 5*) (45). Most of the studies to date are using NBI systems.

This simplified NBI classification diagnoses gastric intestinal metaplasia with 83% accuracy for normal histology (pattern A), 84% for intestinal metaplasia (pattern B) and 95% for dysplasia (pattern C) with high reproducibility. More than 90% of the individuals at risk for gastric adenocarcinoma could be identified by application of this simplified classification by better detection of gastric intra-epithelial metaplasia (GIM) (46). Endoscopic grading with score >5 for Intestinal metaplasia is identified as optimal cutoff for extensive metaplasia during surveillance but validation studies are lacking (47).

MESDA-C (Magnifying endoscopy simple diagnostic algorithm for early gastric cancer-EGC), has shown its utility in the diagnosis for the evaluation of a suspicious gastric lesion. In a suspicious lesion, the presence of a demarcation line with irregular microvascular/microsurface pattern diagnosed EGC with 97% accuracy (48).

Miscellaneous applications of electronic chromo-endoscopy

NBI can also show increased number, dilatation and tortuosity of IPCLs and greater presence of microerosions compared to controls in patients with GERD (49). i-SCAN also showed improved diagnosis of GERD compared to WLE (50).

Villous atrophy associated with celiac disease can be diagnosed with FICE with 100% accuracy and with NBI with sensitivity of 93% and 98% specificity (51). i-SCAN was also useful in complete atrophy (52). Role of NBI in FAP for duodenal adenoma and also to delineate ampullary dysplasia has also been described (53,54).

Electronic chromo-endoscopy in colonic polyps

Narrow band imaging has been studied most extensively in the management of the colonic polyps, but literature regarding other modalities like I-SCAN and FICE is also evolving.

A consensus-based classification NICE (NBI international colorectal endoscopic classification systems) based on color, vessels, and the surface pattern criteria for endoscopic diagnosis of small colorectal polyps has been developed (*Figure 5, Table 6*) (55,56).

Microvessel network and thickness of the vessel are ways of assessing histological grade and depth of invasion of colorectal tumors. On real time colonoscopy, endoscopists confidently made diagnosis in 75% of consecutive small colorectal polyps using NICE classification with high accuracy, sensitivity, and positive predictive values (PPV) (55). In a multicentric prospective study by Puig *et al.*, analysis of more than 2000 colonic lesions >10 mm, NICE classification identified lesions with deep invasion with 95% accuracy even by non-experts (57).

Japan Narrow-Band Imaging Expert Team (JNET) classification based on NBI-ME imaging divides NICE type 2 into type 2A (low grade adenomas) and type 2B (high grade adenomas/submucosally invasive cancer) (58). Retrospective analysis of over 2,900 colorectal lesions concluded that JNET classification very reliably predicts histology, except for

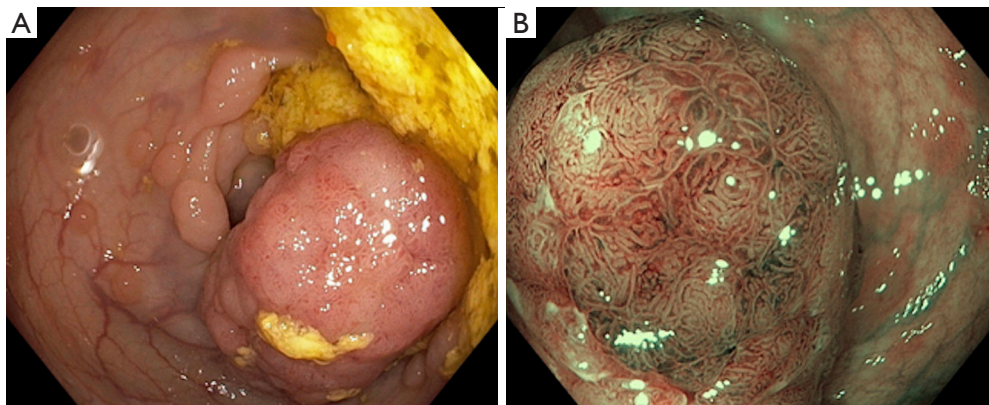


Figure 5 Adenomatous colonic polyp. (A) White light imaging, (B) narrow band imaging.

Table 6 NBI International Colorectal Endoscopic (NICE) Classification (50,51)

	Type 1	Type 2	Type 3
Color	Same or lighter than background	Browner relative to background	Brown to dark brown relative to background; sometimes patchy whiter areas
Vessels	None, or isolated lacy vessels coursing across the lesion	Brown vessels surrounding white structures	Has areas of disrupted or missing vessels
Surface pattern	Dark or white spots of uniform size, or homogeneous absence of pattern	Oval, tubular or branched white structure surrounded by brown vessels.	Amorphous or absent surface pattern
Most likely pathology	Hyperplastic	Adenoma (<i>Figure 5</i>)	Deep submucosal invasive cancer

Type 2 consists of Vienna classification types 3, 4 & superficial 5 (all adenomas with either low or high grade dysplasia or with superficial submucosal carcinoma). The presence of HGD or superficial submucosal carcinoma may be suggested by an irregular vessel or surface pattern, and is often associated with atypical morphology.

type 2B lesions which is less specific and need additional pit pattern examination using dye-based chromo-endoscopy (59). Although small validation studies have been published, large scale validation is still lacking (60).

The “Workgroup Serrated Polyp and Polyposis” (WASP) classification incorporates premalignant lesions like sessile serrated polyps/adenomas (SSAs) and traditional serrated adenomas (TSA) unlike NICE classification. Based on WASP classification, ≥ 2 features out of clouded surface, indistinct borders, irregular shape and dark spots inside crypts are diagnostic of sessile serrated polyps optically (61).

NBI might be better than standard-WL colonoscopy and equal to HD-WLE for colorectal polyps (62). HD-NBI does not fare well in terms of increased ADR or flat adenomas, nor it improves the miss rates in screening colonoscopy (63). In a meta-analysis, HD-NBI improved detection of flat adenomas nearly two-fold at the cost of increased withdrawal time (64). In an RCT comparing new

generation (190-NBI) colonoscopy with HD-WLE showed higher colorectal adenoma/polyp detection rate without difference in adenoma miss rate (65). Further improvements in NBI technology like LUCERA ELITE improved polyp visibility and HD PHL (High definition Pentax Hiline) improve ADR by detecting flat adenomas with higher accuracy (66,67).

In a meta-analysis of 13 studies evaluating comparison of sensitivities of various modalities of IEE, NBI (60%) and magnification NBI (80%) were better than WLE for discriminating SSA/Ps from non-neoplastic lesions where’s modalities like FICE had poor sensitivity (47%) to in comparison with WLE, NBI (89% *vs.* 75%) & M-NBI (78% *vs.* 63%) demonstrated greater sensitivity (68).

A systematic review and meta-analysis by Backes *et al.* showed that both NBI and magnification chromo-endoscopy (MCE) are comparable in efficacy in optical diagnosis of T1 colorectal cancer (CRC) with deep

submucosal invasion when used in adjunct to gross morphological features (GMF). This can be useful to reduce piece-meal resection for T1 CRC or unnecessary surgical referral for endoscopically resectable lesion (69). Another systematic review by Zhang *et al.* concluded that M-NBI and MCE had comparable specificities but M-NBI has lower sensitivity in diagnosing deep submucosal CRC (70). ASGE & ESGE endorses use of advanced endoscopic imaging for evaluation & management of colorectal lesions.

Accurate optical diagnosis and resection of small adenomas (<10 mm) during colonoscopy can allow hyperplastic polyps to be left *in situ* and help determining surveillance intervals without need for tissue diagnosis. The sensitivity of NBI was 83.4% and hence not currently accurate enough to replace histology in determining surveillance intervals (71).

i-SCAN improves resolution of epithelial mucosal surface and vessels, and hence i-SCAN has better efficacy compared to HD-WLE in histological prediction of diminutive colorectal polyps. I-SCAN is comparable to NBI in predicting histology of diminutive colorectal polyps (72,73). Kidambi *et al.* in a recent randomized trial of 740 patients comparing i-SCAN 1 with HD-WLE colonoscopy, found 9.4% improvement in the ADR and 10.3% improvement in the neoplasia detection rate with the use of i-SCAN 1 which was mainly due to better detection of flat, diminutive, right sided adenomas (74). Hoffman in another study found i-SCAN with HD-colonoscopy found more adenomas and flat lesions as compared to SD-WLE (75). However conflicting results has been seen in few other studies. In a prospective cohort study of 84 patients, i-SCAN failed to show any significant improvements in the histology prediction over HD-WLE (76). It was not clear whether the increased sensitivity of I-SCAN with HD was due to HD or I-SCAN. This was answered in a study by Bowman in a prospective study of 1,936 average risk patients for screening colonoscopies, which compared i-SCAN to HD-WLE to detect adenomas and found out that i-SCAN significantly detected more adenomas (618 *vs.* 402, $P<0.01$) and advanced adenomas more than 1 cm (79 *vs.* 47, $P<0.021$) as compared to HD-WL colonoscopy (77). HD i-SCAN improved ADR when compared to SD-WL colonoscopy with i-SCAN and it also meets ASGE PIVI standards for optical diagnosis of diminutive polyps but only in expert hands and also needs training. Its performance is similar to standard i-SCAN, NBI and FICE (78).

FICE (Fujinon) usage study of colonic polyps is scarce and comes from 2 prospective randomized multicentric

studies comparing it with WLE. In one study of 764 patients, there was no significant difference in ADR in FICE group compared to WLE and targeted indigo carmine spraying but sensitivities were comparable in predicting histology (79). Second large RCT by Ainalai *et al.* including 1,318 patients showed no advantage of the FICE technique over conventional high resolution endoscopy in terms of ADR (80). Chung *et al.* in a randomized study enrolled 1,650 subjects with 550 in each arm of NBI, FICE and WL, it showed that compared to WLE, NBI nor FICE did not increase ADR rate in the hand of non-experts (81). In a network meta-analysis by Li *et al.*, in recent study showed that ADR was better with virtual chromo-endoscopy compared to SD-WLE and HD-WLE (82). In a study by Teixeira *et al.*, FICE capillary classification accurately determined nature of colorectal lesions in 98.3% with high sensitivity, specificity, positive and negative predictive value (NPV) with high interobserver agreement (83). a prospective series from UK showed that FICE along with indigo carmine application significantly improves *in vivo* diagnosis of colonic polyps over WLI and can lead to significant cost savings avoiding unwanted biopsies (84).

Electronic chromoendoscopy in ulcerative colitis (Figure 6)

Chromoendoscopy is recommended standard of care for detecting dysplasia in inflammatory bowel disease (IBD) according to all international guidelines (85,86). Chromoendoscopy had an incremental yield of 6% (95% CI, 1–14%) for the diagnosis of dysplasia over NBI. NBI allowed better detection of *in vivo* angiogenesis (marker of colonic inflammation) compared to WLE in IBD patients (87). Study with i-SCAN including 78 consecutive patients showed that subtle vascular and mucosal abnormalities in patients with Mayo endoscopy subscore of 0 or 1 at WLE correlated closely with histological and clinical outcome (88). Randomised parallel group trial including 112 patients showed that there was no difference between NBI and HD-WLE with regard to dysplasia detection (89).

Bisschops *et al.* showed although chromo-endoscopy (CE) and NBI do not differ significantly for detection of dysplasia UC, given the easy applicability of NBI and longer withdrawal time for CE, NBI could replace CE (90).

Kudo *et al.* studied microvascular pattern for accurate healing in ulcerative colitis (91). In a randomized non-inferiority trial of 270 patients, Iacucci *et al.* showed that

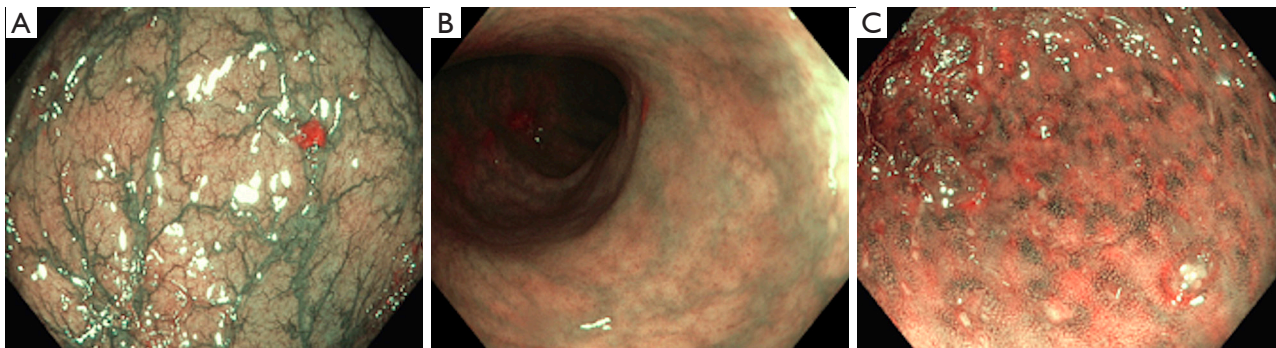


Figure 6 NBI in colorectum. (A) Normal colonic mucosa on NBI; (B) oedematous mucosa with loss of vascular pattern on NBI in UC; (C) UC-active inflammation. NBI, narrow-band imaging; UC, ulcerative colitis.

electronic chromoendoscopy and HD-WLE were non-inferior to dye spraying colonoscopy and neoplastic pit pattern (III–V) (OR 21.5) and location within the right colon (OR 6.52) were associated with neoplasia (92). Same group has developed and validated FACILE (Frankfurt Advanced Chromoendoscopic IBD Lesions) which showed that irregular surface, irregular vessel architecture, signs of inflammation and non-polypoidal morphology were the best predictors of dysplastic lesions on histology (93). Using kudo pit pattern, experts differentiated between neoplastic and non-neoplastic lesions with good sensitivity and specificity (94). Use of Kudo pit pattern to predict histology in the presence of inflammation in UC when assessed by standard colonoscopes in non-experts hands is still controversial (95).

Cassinotti *et al.* in first study of 205 patients using FICE and studying kudo pit pattern in differentiation of the lesions in ulcerative colitis and it accurately identified with sensitivity, specificity, PPV and NPV of 92%, 76%, 3.8 and 0.12 respectively. Fibrin cap as a marker of inflammation improves the identification further (96). To conclude electronic chromoendoscopy will be useful in surveillance and management of ulcerative colitis.

Conclusions

Electronic chromoendoscopy has helped in identifying, delineating and assessing the depth of the lesions in the routine endoscopies at the switch of a button, but validated simple classification systems are the need of the hour to make it more acceptable in clinical practice. NBI is the most popular advanced imaging technique but data regarding FICE, i-SCAN is also emerging. Narrow band imaging has

proved to be an useful adjunct in surveillance endoscopies and colonoscopies allowing targeted biopsy with improved detection. Prediction of nature of polyp together with a “Resect and discard” policy for diminutive polyps has become possible but further multicentric studies are needed before we put this into practice.

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