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New endoscopes and add-on devices to improve colonoscopy performance

Paraskevas Gkolfakis, Georgios Tziatzios, George D Dimitriadis, Konstantinos Triantafyllou

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Abstract

Colonoscopy is the gold standard for colorectal cancer prevention; however, it is still an imperfect modality. Precancerous lesions can be lost during screening examinations, thus increasing the risk of interval cancer. A variety of factors either patient-, or endoscopist dependent or even the procedure itself may contribute to loss of lesions. Sophisticated modalities including advanced technology endoscopes and add-on devices have been developed in an effort to eliminate colonoscopy's drawbacks and maximize its ability to detect potentially culprit polyps. Novel colonoscopes aim to widen the field of view. They incorporate more than one cameras enabling simultaneous image transmission. In that way the field of view can expand up to 330°. On the other hand a plethora of add-on devices attachable on the standard colonoscope promise to detect lesions in the proximal aspect of colonic folds either by offering a retrograde view of the lumen or by straightening the haustral folds during withdrawal. In this minireview we discuss how these recent advances affect colonoscopy performance by improving its quality indicators (cecal intubation rate, adenoma detection rate) and other metrics (polyp detection rate, adenomas per colonoscopy, polyp/adenoma miss rate) associated with examination's outcomes.

Key words: Colonoscopy; Quality indicators; Wide-angle view colonoscopes; Add-on devices

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Core tip: Accomplishing high intra-procedural colonoscopy quality indicators has been associated with better patients' outcomes. Recently, a number of novel wide-angle view endoscopes as well as different add-on devices have been developed aiming to further improve

these metrics. They promise detailed inspection of otherwise difficult to examine parts of the colonic mucosa. Herein, we present the current evidence regarding the efficacy of these scopes and devices.

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INTRODUCTION

Colorectal cancer (CRC) ranks second regarding cancer-related mortality^[1]. Colonoscopy interrupts the carcinogenesis by detecting and removing precancerous lesions, namely adenomas, thus providing the opportunity for neoplasia screening^[2,3]. Despite its efficacy and widespread use, it is an imperfect examination. Almost a quarter of existing colonic adenomas remain undetected during a screening colonoscopy, while more recent studies raise that percentage up to 40%^[4-7]. The so-called missed adenomas are considered independent risk factor for interval CRC^[8], defined as CRC rising within the surveillance intervals. Missed adenomas are of particular significance in the right colon (RC), where more than half of the interval CRC incidents occur^[9]. Furthermore, the usually flat serrated sessile adenomas (SSA) of the RC represent premalignant lesions of a distinct group of CRCs that also develop predominantly in the proximal colon^[10,11].

Missed adenomas are a consequence of multiple factors; poor bowel preparation^[12], inability to complete the colonoscopy by visualizing the cecum^[13], inadequate withdrawal times^[14], lack of expertise^[15] and poor inspection of the proximal side of the colonic folds, as well as of the region around the anatomic flexures and the ileocecal valve^[16,17].

Recent studies highlighted the importance of accurate adenoma detection during screening colonoscopies. Corley *et al.*^[18] evaluated more than 300000 examinations and proved that patients of both genders undergoing screening colonoscopy by an endoscopist with high adenoma detection rate (ADR) are protected against interval CRC both in the proximal and the distal colon in comparison with individuals undergoing colonoscopy by a physician with lower ADR. Similarly, a mathematical model showed that 1% increase of the ADR leads to 3% decrease of colon cancer incidence^[8].

Aiming to provide patients the highest level of health services, scientific endoscopy Societies have recommended specific quality indicators to measure colonoscopy outcomes^[19]. Similarly, endoscopy

industries make continuous efforts to develop novel endoscopes and several devices to improve colonoscopy's intrinsic technical imperfectness (Table 1). Almost a decade ago, a simple transparent plastic cap was one of the first devices introduced to increase endoscopists' performance. Since then several studies have been conducted that led to two meta-analyses^[20,21]. Their results indicate marginal efficacy of cap-assisted colonoscopy (CAC) to increase detection of patients with polyps. Due to the lack of further remarkable evolvement, cap-assisted colonoscopy will not be discussed in this paper. Marginal improvement of colonoscopy performance was also associated with the advent of high - definition endoscopy^[22]. Due to this marginal positive effect and high costs of the investment, this technology is not the standard of care worldwide yet, and its detailed presentation is beyond the scope of this minireview.

In this minireview we focus on the intra-procedural quality indicators: cecal intubation rate (CIR), polyp detection rate (PDR), adenoma detection rate (ADR), adenomas per colonoscopy (APC), as well as the polyp- and adenoma miss rates (PMR/AMR) (Table 2) in studies evaluating wide angle view (> 170°) colonoscopes and new add-on colonoscopy accessories. The term ADR -patients with at least one adenoma- will be used to describe not only the adenoma detection rate in screening/surveillance populations, but also in symptomatic individuals. To facilitate readers' comprehension the exact composition of each study population regarding its indication will be presented, whenever needed.

We conducted a comprehensive review of English literature published in MEDLINE electronic database from January 2008 until January 2017. The following key words were used: "wide-angle view colonoscopes", "Third-Eye Retroscope", "Full-Spectrum Endoscopy", "balloon assisted colonoscope", "Endocuff" and "Endorings". Moreover, data from abstracts presented during the Digestive Diseases Week and the United European Gastroenterology Week from 2010 to 2016 were retrieved and manually searched. First author name, year of publication, study design, number of participants, their age and indications, CIR, PDR, ADR APC, PMR and AMR were extracted either as reported by the authors or after appropriate calculation.

WIDE-ANGLE VIEWING ENDOSCOPES

One of the factors potentially accountable for missed lesions during colonoscopy is the relatively narrow field of view (140°-170°) of standard forward viewing (SFV) colonoscopes. In an effort to eliminate this limitation, novel wider field of view endoscopes have been manufactured, allowing meticulous inspection of the proximal aspect of the haustral folds. Table 3 summarizes data from studies regarding wide-angle view colonoscopy platforms.

Table 1 Available endoscopes and add-on devices for improving colonoscopy outcomes

Wide-angle view colonoscopes		Add-on devices	
Brand	Manufacturer	Brand	Manufacturer
Full-spectrum endoscopy platform (Fuse)	EndoChoice, GA, United States	Third-Eye Retroscope (TER)	Avantis Medical Systems, Inc, Sunnyvale, CA, United States
Extra-wide angle view colonoscope	Olympus Co., Tokyo, Japan	Third-Eye Panoramic	Avantis Medical Systems, Inc, Sunnyvale, CA, United States
Self-propelled disposable colonoscopy system (Aer-O-Scope)	GI View Ltd, Ramat Gan, Israel	Endocuff	Arc Medical Design, Leeds, England
		Endocuff-Vision	Arc Medical Design, Leeds, England
		EndoRings	EndoAid Ltd, Caesarea, Israel
		NaviAid G-EYE	SMART Medical Systems Ltd, Ra'anana, Israel

Table 2 Intra-procedural quality indicators

Metric	Definition	Suggested target (references)
Cecal intubation rate	The frequency of completed colonoscopies (cecum is visualized)	Overall: $\geq 90\%$ Screening: $\geq 95\%$ ^[19]
Polyp detection rate	The proportion of patients with at least one polyp	N/A
Adenoma detection rate	The proportion of patients with at least one adenoma	Men: $\geq 30\%$ Women: $\geq 20\%$ ^[19]
Adenoma per colonoscopy	The mean number of adenomas detected per colonoscopy	N/A
Polyp miss rate (PMR)	The proportion of polyps missed during a first pass and detected by a second one. It is used in back-to-back studies.	N/A
Adenoma miss rate	The proportion of adenomas missed during a first pass and detected by a second one. It is used in back-to-back studies.	N/A

N/A: Recommendation not available.

Full-spectrum endoscopy (Fuse) system

The full-spectrum endoscopy platform (Fuse, EndoChoice, GA, United States) consists of a video colonoscope and a processor. The colonoscope is a normal adult (168 cm working length, 12.8 mm outer diameter) flexible and reusable scope that allows both diagnostic and therapeutic procedures. It provides high-resolution, 330° field of view, achieved by three imagers and LED groups positioned one at the front and two at each side of the scope's distal tip. The images in the three monitors (Figure 1) reflect transmission from the respective lenses (right image for the right-sided lens, center image for the central positioned lens and left image for the left-sided lens). The endoscopist is allowed to perform all potential maneuvers, such as complete tip deflection (180° up/down direction and 160° left/right direction).

This novel platform has been proven to be safe and feasible with CIR almost 100% in two non-randomized studies^[23,24]. Gralnek *et al.*^[25] conducted an international, multicenter, randomized back-to-back study to investigate whether Fuse detects more missed adenomas in comparison to SFV colonoscopy. Participants ($n = 197$, mixed indications) were randomly assigned to undergo same day tandem colonoscopies (either Fuse colonoscopy first followed by SFV colonoscopy or vice versa). The Fuse system had significantly lower miss rates compared to SFV endoscopy for adenomas (7% vs 41%, $P < 0.0001$)

and polyps (10% vs 43%, $P < 0.0001$). The majority of the 20 adenomas that were missed during SFV examination and detected by the Fuse were sessile (90%), diminutive (70%) and RC sited (70%). In a similar design cross over study, Papanikolaou *et al.*^[26] showed that Fuse outperformed SFV complemented by examination of the right colon with scope retroflexion regarding adenoma (10.9% vs 33.7%, $P < 0.001$) and polyp (3.0% vs 33.5%, $P < 0.001$) miss rates. The same study showed that the incremental benefit of full-spectrum colonoscopy when performed, as a second examination, was 39% higher compared to that of conventional colonoscopy with retroflexion in the cecum, regarding-adenoma miss-rate overall (Figure 2). Moreover, an even higher incremental benefit was shown in favor of FC in the proximal colon. This benefit might be ameliorated by the fact that the majority of missed lesions measured less than 1 cm, in both study arms.

The ability of Fuse system to improve colonoscopy outcomes has further been evaluated in parallel design non-randomized^[27,28] and randomized studies^[29-31]. Manes *et al.*^[27] conducted a non-randomized study ($n = 529$) comparing Fuse and standard HD colonoscope. The authors reported increased PDR (56.6% vs 44.3%, $P < 0.01$) and ADR (35.5% vs 29.9%) in the Fuse arm. In Denmark Roepstorff *et al.*^[28] recruited 205 consecutive individuals undergoing screening colonoscopy either with Fuse system or with the

Table 3 New endoscopes and colonoscopy performance improvement

Ref.	Study design	Technology	Comparator	N	Indication	Age (yr), range	CIR (%)	PDR (%)	ADR (%)	APC	PMR (%)	AMR (%)
Gralnek <i>et al</i> ^[25] , 2013	Single-center prospective,	FUSE	None	50	Mixed	18-70	100%	-	-	N/A	N/A	N/A
Gralnek <i>et al</i> ^[25] , 2014	Multicenter, prospective, randomized, tandem	FUSE	SFV	101 vs 96	Mixed	18-70	98.0% vs 98.9%	-	¹ 34.0% vs 28.0%	¹ 0.64 vs 0.33	10% vs 43%	7% vs 41%
Papanikolaou <i>et al</i> ^[26] , 2017	Multicenter, prospective randomized, tandem	FUSE	SFV+R	107 vs 108	Mixed	41-80	-	-	-	¹ 0.61 vs 0.50	13.0% vs 33.5%	10.9% vs 33.7%
Hassan <i>et al</i> ^[29] , 2016	Multicenter, prospective, randomized parallel	FUSE	SFV	328 vs 330	Screening after (+) FIT	50-69	92.1% vs 93.3%	-	43.6% vs 45.5%	0.81 vs 0.85	N/A	N/A
Song <i>et al</i> ^[24] , 2016	Singe-center retrospective,	FUSE	None	262	Mixed	22-80	100%	54.20%	36.3%	0.66	N/A	N/A
Rath <i>et al</i> ^[31] , 2015	Multicenter, prospective, parallel	FUSE	SFV	90	-	-	-	36% vs 24.0%	-	-	N/A	N/A
Manes <i>et al</i> ^[27] , 2016	Single-center prospective, parallel	FUSE	SFV	264 vs 265	Mixed	18-85	-	56.6% vs 44.3%	35.5% vs 29.9%	-	N/A	N/A
Roepstorff <i>et al</i> ^[28] , 2016	Single-center prospective, parallel	FUSE	SFV	109 vs 106	Screening	-	83.4% vs 93.4%	N/A	67.0% vs 59.6%	1.8 vs 1.4	N/A	N/A
Leong <i>et al</i> ^[30] , 2016	Single-center, prospective, randomized tandem	FUSE	SFV	25 vs 27	IBD	-	-	-	-	-	² 25.0% vs 71.4%	-
Uraoka <i>et al</i> ^[33] , 2015	Multicenter, feasibility	EWAVC	None	47	Mixed	-	100%	-	-	0.64	N/A	N/A
Uraoka <i>et al</i> ^[34] , 2013	Multicenter, prospective, randomized parallel	EWAVC	SFV	316	Mixed	-	-	-	50.6% vs 45.6%	1.1 vs 1.0	N/A	N/A
Gluck <i>et al</i> ^[35] , 2016	Single-center, prospective, tandem	Aer-O-Scope	SFV	56	Screening	27-72	98.2% vs 98.2%	-	21.4% vs 25.0%	-	12.5% for Aer-O-Scope	-

¹Refers to the first of the tandem examinations; ²Dysplasia miss-rate. N/A: Non-applicable; -: Data not provided; CIR: Cecal intubation rate; PDR: Polyp detection rate; ADR: Adenoma detection rate; APC: Adenoma per colonoscopy; PMR: Polyp miss rate; AMR: Adenoma miss rate; FUSE: Full-spectrum endoscopy platform; SFV: Standard forward view colonoscopy; SFV + R: Standard forward view colonoscopy + retroflexion in cecum; EWAC: Extra-wide-angle view colonoscopy; IBD: Inflammatory bowel disease.

conventional endoscope. Completion rate was lower with Fuse (83.4% vs 93.4%, $P = 0.04$) but Fuse showed numerically higher ADR (67% vs 59.6%, $P = 0.36$) and APC (1.8 vs 1.4, $P = 0.09$).

Hassan *et al*^[29] compared the ADR of Fuse and SFV study arms in 658 individuals undergoing colonoscopy after positive FIT test in the context of a population-based massive screening program. Of interest, both ADR and APC were similar (43.6 vs 45.5 and 0.81 vs 0.85, respectively) between Fuse and conventional colonoscopy. Statistical significant difference was neither shown for advanced adenomas, sessile serrated adenomas and proximal adenomas. Authors acknowledged that the high ADR in the control group, potentially related to the disease enriched (FIT+) population of the study might have hindered detection of difference. Apart from that, sample size issues also

rise, since randomized control trials of parallel design would normally require significantly more participants in order to achieve sufficient statistical power^[32].

Another small ($n = 90$), randomized, prospective study^[31] that assigned patients to undergo either Fuse or conventional colonoscopy showed higher PDR (36% vs 24%) associated with Fuse use.

Finally, the Fuse system has also been evaluated in patients with inflammatory bowel diseases (IBD). In a randomized back-to-back study from Australia^[30], 52 IBD patients underwent tandem colonoscopies with Fuse system and conventional colonoscopy in order to evaluate dysplasia miss rate of the first examination (25 patients had Fuse index colonoscopy and 27 started with the conventional examination). Fuse was associated with a significant lower dysplasia miss rate (25% vs 71.4%, $P = 0.0001$).



Figure 1 Fuse platform (EndoChoice, GA, United States) consists of the processor, the endoscope with one forward and two lateral cameras and a wide screen where you can appreciate the simultaneous monitor presentation from the three cameras (left, center, right) of the full-spectrum endoscopy system (Image courtesy of Endochoice, GA, United States).

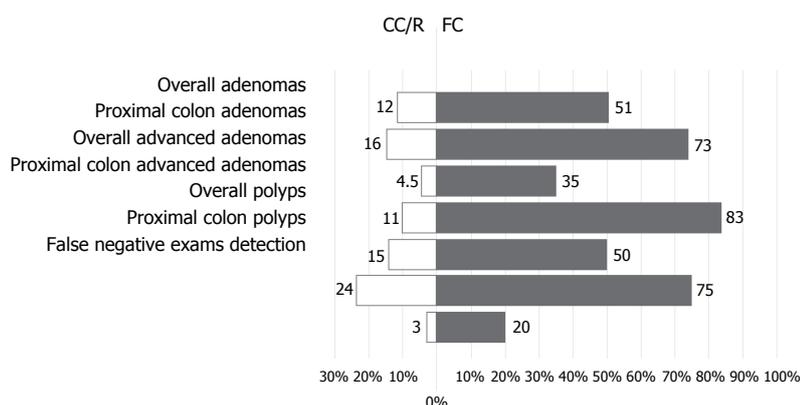


Figure 2 Incremental benefit from full-spectrum colonoscopy compared to conventional colonoscopy complemented with proximal colon examination with scope retroflexion^[26].

Extra-wide angle view colonoscope

This prototype colonoscope introduced by Olympus Co., Tokyo, Japan is composed of two lens' systems: a standard 140°-angle forward-viewing lens and a 144-232°-angle lateral-backward viewing lens. A video monitor puts together the images of both lens and presents them simultaneously as a single image. Following an initial feasibility study^[33] showing CIR

of 100%, Uraoka *et al.*^[34] compared this prototype scope to SFV in a randomized parallel design study regarding APC and ADR. The sample consisted of 316 individuals undergoing colonoscopy for various indications. The extra-wide angle view colonoscope (EWAVC) had similar to the SFV system APC (1.1 vs 1.0, $P = 0.36$) and ADR (50.6% vs 45.6%, $P = 0.43$). However, this novel system may be proven of special

importance in angulated and narrow regions of the colon (*i.e.*, sigmoid), as per segment analysis showed a statistically significant higher sigmoid-APC in favor of EAWVC (0.4 vs 0.2, $P = 0.04$).

Aer-O-Scope

The efficacy and safety of this self-propelled disposable colonoscopy (SPDC) system (Aer-O-Scope; GI View Ltd, Ramat Gan, Israel) has been evaluated in one non-randomized prospective study of 56 patients undergoing tandem screening colonoscopies^[35]. Its optical system consists of white-light LEDs and a CMOS high-definition digital camera; it allows a simultaneous 57° field of forward view an omni 360° view of a cylindrical band of the colon. Participants underwent colonoscopy with Aer-O-Scope first while SFV colonoscopy followed. SPDC was proven to be safe and effective. CIR were similar between SPDC and SFV endoscopy but SPDC failed to detect 5/40 of the polyps identified by the SFV (PMR 12.5%), leading to a lower ADR (21.4% vs 25%) in comparison to SFV.

“ADD-ON” COLONOSCOPY DEVICES

With the term “add-on” device we describe all those accessories appended on the distal end of a standard colonoscope to facilitate meticulous inspection of the colonic mucosa. These devices provide either a retrograde view of the lumen (Third-Eye Retroscope, Avantis Medical Systems, Inc, Sunnyvale, CA, United States), or wider field of view (Third-Eye Panoramic, Avantis Medical Systems, Inc, Sunnyvale, CA, United States) or flattening of colonic folds during withdrawal to allow visualization of their proximal side of the colonic folds (Endocuff, Arc Medical Design, Leeds, England; Endocuff-Vision, Arc Medical Design, Leeds, England; EndoRings, EndoAid Ltd., Caesarea, Israel and balloon assisted-colonoscopy using the G-EYE, SMART Medical Systems Ltd, Ra’anana, Israel). Table 4 summarizes data originating from the available studies that evaluated their safety, feasibility and efficacy in improving colonoscopy performance.

Third-Eye Retroscope and the Third-Eye panoramic

The Third-Eye Retroscope (TER) (Avantis Medical Systems, Inc, Sunnyvale, CA, United States) is one of the first auxiliary imaging devices that tried to extend the field of view of the standard forward viewing colonoscope^[36]. The TER is inserted through the working channel, it extends beyond the distal tip of the SFV scope to bend 180° in a J-shape form, looking opposite of the scope main lens; thus, it provides a complement retrograde view of the colonic lumen during scope withdrawal. Three open-label, one-arm prospective studies implementing the device on the SFV colonoscopy showed that in the absence of Third Eye the examinations’ polyp and adenoma miss rates would be 4.4%-12.9% and 7.8%-13.8%,

respectively^[37-39]. In accordance with these findings, Leufkens *et al*^[5] presented the results of a randomized tandem clinical study comparing PMR and AMR of SFV colonoscopy with SFV colonoscopy plus TER. In the per protocol analysis the TER arm was associated with significantly lower miss rates (PMR: 15.9% vs 32.8% and AMR: 18.4% vs 31.4%). However, the above studies underlined some limitations related to TER use. The device narrows almost 50% the diameter of the working channel making the suction of residues compulsory prior to withdrawal. Moreover, each time a polyp is detected by the retrograde view of TER the device must be removed to allow lesion removal, leading to significant prolongation of the procedure. Finally, the device procurement bears an additional financial burden. For these reasons the device has been abandoned.

A few years later the same manufacturer developed another device called the Third-Eye® Panoramic. This plastic cap can be clipped on to the distal tip of all conventional colonoscopes and contains two side-viewing CMOS chips. The cap is connected to a thin plastic catheter that contains the transmission wires that runs along the scope’s shaft. The catheter ends to an external video processor connected to the conventional colonoscope’s video monitor, resulting in an extended field of view (more than 300°) through three - partially overlapping - images. This novel device has only been evaluated in a feasibility study^[40]. In this small study, the device was easy to use and the cecum was intubated in all cases.

Endocuff

Endocuff (Arc Medical Design, Leeds, United Kingdom) is a plastic, 2 cm long cuff that can be mounted onto the tip of the scope. Endocuff entails two rows of “finger”-like projections, which remain smooth during insertion and bend in the withdrawal phase to flatten the colonic folds and allow assessment of a greater, otherwise unsighted, mucosal area (Figure 3). Endocuff and its “descendant” Endocuff-Vision have been evaluated in numerous studies (Table 4). Feasibility studies^[41,42] showed that Endocuff was safe, since only minor insignificant mucosal lacerations were the adverse events related with its use. In these studies the rates of cecal intubation were higher than 98%. Three randomized parallel design studies^[43-45] have been published comparing Endocuff-assisted colonoscopy (EAC) to conventional colonoscopy in terms of polyp and adenoma detection. Two studies from Germany^[43,44], each recruiting almost 500 patients undergoing colonoscopy for various indications, favored Endocuff use to detect more patients with at least one polyp/adenoma compared to the conventional procedure (PDR: 55.4% vs 38.4% and 56% vs 42% and ADR: 35.4% vs 20.7% and 36% vs 28%, respectively). On the other hand, a similar multicenter Dutch study^[45] that randomized more than 1000 patients of various

Table 4 Add-on devices and colonoscopy performance improvement

Ref.	Study design	Device	Comparator	N	Indication	Age (yr)	CIR (%)	PDR (%)	ADR (%)	APC	PMR (%)	AMR (%)
Triadafilopoulos <i>et al</i> ^[36] , 2008	Single-center, prospective, pilot	TER	² SFV	24	Screening Surveillance	mean: 64					³ 10.5	³ 11.1
Waye <i>et al</i> ^[39] , 2010	Multicenter, prospective, open-label	TER	² SFV	249	Screening Surveillance	mean: 63				0.61 vs 0.55	³ 11.7%	³ 9.9%
DeMarco <i>et al</i> ^[37] , 2010	Multicenter, prospective, open-label	TER	² SFV	298	Mixed	mean: 57				0.39 vs 0.34	³ 12.9%	³ 13.8%
Leufkens <i>et al</i> ^[5] , 2011	Multicenter, prospective, randomized, tandem	TER	SFV	176 vs 173	Mixed	range: 23-83					15.9 % vs 32.8% (PP)	18.4% vs 31.4% (PP)
Mishkin <i>et al</i> ^[38] , 2012	Single-center, prospective	TER	² SFV	68	Mixed						³ 4.4%	³ 7.8%
Abstract Rubin <i>et al</i> ^[40] , 2015	Single center, Prospective, feasibility	TEP	² SFV	33	Mixed	mean: 60	100%		44% overall			
Gralnek <i>et al</i> ^[62] , 2014	Single-center, prospective, cohort	G-EYE	None	47	Mixed	mean: 59	100%	53.2	44.70%	0.76	N/A	N/A
Halpern <i>et al</i> ^[63] , 2014	Multicenter, prospective, randomized, tandem	G-EYE	SFV	54 vs 52	Mixed	mean: 55 vs 58	100% vs 100%	-	¹ 40.4% vs 25.9%	-	-	7.5% vs 44.7%
Halpern <i>et al</i> ^[63] , 2014	Multicenter, prospective, randomized, parallel	G-EYE	SFV	105 vs 117	Screening Surveillance	≥ 50	-	-	35.4% vs 23.5%	0.63 vs 0.36	N/A	N/A
Rey <i>et al</i> ^[64] , 2015	Multicenter, prospective, randomized, tandem	G-EYE	SFV	25 vs 24	Referral for colonoscopy	-	-	-	-	-	17 vs 41	-
Hendel <i>et al</i> ^[66] , 2015	Multicenter, prospective, randomized, parallel	G-EYE HD	SFV	54 vs 50	Mixed	≥ 50	-	76% vs 46%	59% vs 39%	1.15 vs 0.66	N/A	N/A
Shirin <i>et al</i> ^[67] , 2016	Multicenter, prospective, randomized, parallel	G-EYE HD	SFV	242 vs 238	Mixed	mean: 65	-	-	49.2% vs 33.8%	0.93 vs 0.57	N/A	N/A
Dik <i>et al</i> ^[61] , 2015	Multicenter, prospective, randomized, tandem	Endorings	SFV	57 vs 59	Mixed	mean: 59	100% vs 100%	¹ 68.4% vs 40.7%	¹ 49.% vs 28.8%	¹ 1.05 vs 0.51	9.1% vs 52.8%	10.4% vs 48.3%
Lenze <i>et al</i> ^[41] , 2014	Single-center, retrospective	Endocuff	None	50	Mixed	mean: 57	98%	-	34%	0.72	N/A	N/A
Floer <i>et al</i> ^[43] , 2014	Multicenter, prospective, randomized, parallel	Endocuff	SFV	249 vs 243	Mixed	median: 64	96% vs 94%	55.4% vs 38.4%	35.4% vs 20.7%	0.58 vs 0.36	N/A	N/A
Biecker <i>et al</i> ^[44] , 2015	Two-center, prospective, randomized, parallel	Endocuff	SFV	245 vs 253	Mixed	median: 67	98% vs 98%	56% vs 42%	36% vs 28%	-	N/A	N/A
Sawatzki <i>et al</i> ^[42] , 2015	Multicenter, prospective, feasibility	Endocuff	None	104	Screening Surveillance	mean: 59	99%	72%	47%	-	N/A	N/A
Van Doorn <i>et al</i> ^[45] , 2015	Two-center, prospective, randomized, parallel	Endocuff	SFV	1033 (ITT: 504 vs 529 PP: 486 vs 514)	Mixed	median: 65 vs 65	ITT: 98% vs 99%	-	ITT: 52% vs 52%	ITT: 1.36 vs 1.17	N/A	N/A
							PP: 94% vs 99%		PP: 54% vs 53%	PP: 1.44 vs 1.19		

De Palma <i>et al</i> ^[46] , 2017	Single-center, prospective, crossover, tandem	Endocuff	SFV	137 vs 137	Mixed	mean: 55 vs 56	100% vs 100%	-	¹ 27.7% vs 28.5%	¹ 0.63 vs 0.52	-	1.1% vs 29.7%
Floer <i>et al</i> ^[48] , 2014 Abstract	Multicenter, prospective, randomized, parallel	Endocuff	SFV	652	Screening	mean: 64	98.5% vs 99.1%	55.4% vs 39.9%	-	0.9 vs 0.54	N/A	N/A
Marsano <i>et al</i> ^[50] , 2014 Abstract	Multicenter, retrospective	Endocuff	SFV	165 vs 153	Screening Surveillance	-	-	-	46.6% vs 30%	0.8 vs 0.38	N/A	N/A
Chin <i>et al</i> ^[53] , 2015 Abstract	Single-center, cohort	Endocuff	SFV	93 vs 143	Mixed	-	-	78.5% vs 57.3%	44.1% vs 27.3%	-	N/A	N/A
Patel <i>et al</i> ^[52] , 2016 Abstract	Single-center, cohort	Endocuff	SFV	452 vs 597	Mixed	-	-	79.0% vs 57.4%	51.8% vs 36.3%	1.59 vs 0.91	N/A	N/A
Higham-Kessler <i>et al</i> ^[56] , 2016 Abstract	Single-center, cohort	Endocuff	SFV	77 vs 153	Screening Surveillance	-	-	-	67% vs 62.7%	-	N/A	N/A
Garcia <i>et al</i> ^[51] , 2016 Abstract	Single-center, randomized, parallel	Endocuff	SFV	174 vs 163	Screening	mean: 61	-	29.9% vs 15.9%	22.4% vs 13.4%	0.31 vs 0.22	N/A	N/A
Wada <i>et al</i> ^[49] , 2016 Abstract	Two-center, randomized, parallel	Endocuff	SFV	239 vs 207	-	-	EAC: 98.8%	62% vs 50%	55% vs 40%	-	N/A	N/A
Bensuleiman <i>et al</i> ^[54] , 2016 Abstract	Single-center, prospective, randomized, parallel	Endocuff	CAC	84 vs 75	Screening	-	98% vs 99%	-	53% vs 59%	1.03 vs 1.00	N/A	N/A
Cavallaro <i>et al</i> ^[55] , 2016 Abstract	Single-center, cohort	Endocuff	SFV	605 vs 579	Screening Surveillance	mean: 60 vs 60	-	-	53% vs 48%	1.1 vs 0.88	N/A	N/A
Triantafyllou <i>et al</i> ^[47] , 2016 Abstract	Multicenter, prospective, randomized, tandem	Endocuff	SFV	100 vs 100	Mixed	mean: 61	-	-	-	¹ 0.93 vs 0.53	-	14.7% vs 37.6%
Tsiamoulos <i>et al</i> ^[58] , 2015 Abstract	Single-center, cohort	Endocuff-vision	SFV	133 vs 266	Screening	-	-	-	68.9% vs 58.4%	2.2 vs 1.4	N/A	N/A
Bhattacharyya <i>et al</i> ^[60] , 2016 Abstract	Single-center, prospective, randomized, parallel	Endocuff-vision	SFV	266 vs 265	Screening	-	-	70.3% vs 69.8%	60.9% vs 63%	1.26 vs 1.35	N/A	N/A
Ngu <i>et al</i> ^[59] , 2016 Abstract	Multicenter, prospective, randomized, parallel	Endocuff-vision	SFV	1772	Mixed	mean: 62	96.7% vs 96.4%	-	40.9% vs 36.2%	0.95 vs 0.75	N/A	N/A

¹Refers to the first of the tandem examinations; ²Use of TER/TEP on SFV; ³Miss rate if TER/TEP was not used. N/A: Non applicable; -: Data not provided; CIR: Cecal intubation rate; PDR: Polyp detection rate; ADR: Adenoma detection rate; APC: Adenoma per colonoscopy; PMR: Polyp miss rate; AMR: Adenoma miss rate; SFV: Standard forward view colonoscopy; CAC: Cap-assisted colonoscopy; TER: Third-Eye Retroscope; TEP: Third Eye Panoramic Cap; EAC: Endocuff-assisted colonoscopy; ITT: Intention to treat analysis; PP: Per protocol analysis.

indications failed to reveal any advantage of Endocuff use regarding the proportion of patients with at least one adenoma (ADR: 52% for both arms) and the mean number of adenomas per patient (APC: 1.36 vs 1.17).

Two studies have evaluated Endocuff regarding adenoma miss rate^[46,47]. De Palma *et al*^[46] randomized 274 patients to undergo same day back-to-back colonoscopies (either with Endocuff use first and then without it or vice versa). In this study any lesion detected during the first procedure was left in situ in order to be redetected -or not- during the second

one. Adenoma miss rate was significantly lower when Endocuff was used (1.1% vs 29.7%, $P < 0.001$). Similarly, we recently presented the results of a multicenter tandem study^[47] showing that Endocuff use outperformed its comparator (standard colonoscopy) in terms of AMR, overall (14.7% vs 37.6%, $P = 0.0004$) and in the proximal colon (10.4% vs 39%, $P = 0.004$).

There is also certain amount of data from parallel design studies, reporting increased PDR/ADR^[48-53] in the Endocuff arms published in abstract form only. However, three other studies failed to reveal Endocuff

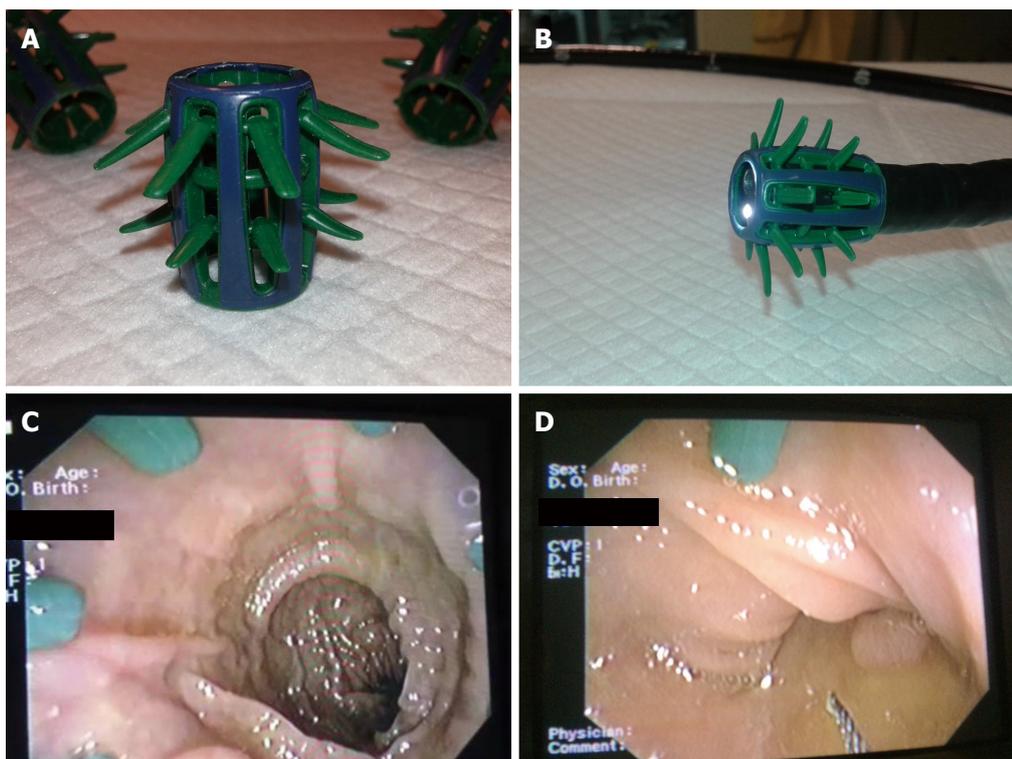


Figure 3 Endocuff (A) fitted onto the tip of the scope (B); device induced flattening of colonic folds during scope withdrawal (C) and assisting lesion reveal during polypectomy (D).



Figure 4 Endocuff-Vision with a single row of projections (Photo courtesy of Dr. Z. Tsiamoulos).

superiority^[54-56]. A recent meta-analysis tried to sum up these data^[57]. Taking into account data from 8 studies ($n = 4387$) the authors concluded that Endocuff use is associated with higher ADR compared to standard colonoscopy (50.4% vs 43.3%, OR = 1.49, 95%CI: 1.23-1.80, $I^2 = 50%$, $P < 0.01$).

Endocuff-Vision (Arc Medical Design, Leeds, England) - the evolution of the initial device, with a single row of projections (Figure 4) has also been evaluated in studies measuring colonoscopy outcomes. Tsiamoulos *et al*^[58] reported extremely high ADR for Endocuff-Vision assisted and conventional colonoscopy in a screening population. However, both ADR and APC were even higher in the Endocuff-Vision arms

(68.9% vs 58.5 and 2.2 vs 1.4, respectively). In a large study of more than 1700 patients^[59], Endocuff-Vision use was associated with a significant higher ADR (40.9% vs 36.2%) in patients of various indications for colonoscopy. Contrariwise, these findings were not confirmed in a single-center prospective parallel design study that involved screening population^[60]: similar ADR and APC between Endocuff-Vision-assisted and cap-assisted colonoscopy were noted.

EndoRings

EndoRings (EndoAid Ltd., Caesarea, Israel) is a silicone-rubber add-on device consisting of three circular rings. It fits onto the distal tip of the endoscope and allows not only the mechanical stretching of the haustral folds during withdrawal, but also maintains the lumen in the center of the inspection field. At the time of insertion the view of field is not affected since the device does not project beyond the distal end of the scope, allowing the unimpeded cecal intubation. This device has been evaluated only in one multicenter randomized tandem study^[61]. In the per protocol analysis of 116 patients of mixed indications, the use of EndoRings was associated with a statistically significant lower polyp (9.1% vs 52.8%, $P < 0.001$) and adenoma (10.4% vs 48.3%, $P < 0.001$) miss rate. The benefit of EndoRings use was higher for the detection of diminutive adenomas (AMR: 13.5% vs 54.2%, $P < 0.001$) and adenomas found both at the proximal and distal colon (AMR: 10.6% vs 58.1%, $P <$

0.001 and 10% vs 37%, $P < 0.001$, respectively)^[61].

Balloon-assisted colonoscopy-The G-EYE

The NaviAid G-EYE (SMART Medical Systems Ltd, Ra'anana, Israel) is a novel balloon-colonoscopy consisting of a standard adult colonoscope combined with an inflatable balloon at the bending part of the scope. The balloon is located 1-2 cm proximally to the distal tip of the colonoscope and it can be inflated up to 60mm diameter with unremarkable alteration in scope's outer caliber^[62]. A special inflation system - the SPARK²C - manipulated by the endoscopist *via* a foot-pedal, inflates the balloon once cecum intubation achieved and retains a constant pressure within the colon during withdrawal. With the balloon inflated during withdrawal, colonic folds and flexures are mechanically straightened revealing potential suspicious lesions located in their proximal aspect^[62]. Two randomized tandem studies^[63,64] evaluated G-EYE's lesions miss rates compared to SFV. Both studies examined individuals undergoing colonoscopy for various reasons. Halpern *et al*^[63] demonstrated a significant lower adenoma miss rate for G-EYE (7.5% vs 44.7%, $P = 0.0002$), while Rey *et al*^[64] showed a lower polyp miss rate in favor of the G-EYE (7% vs 41%). In terms of ADR and adenomas per colonoscopy this device has been evaluated in three randomized parallel design studies^[65,67]. Halpern *et al*^[65] randomized 222 individuals undergoing screening colonoscopy to receive either balloon-assisted colonoscopy or SFV examination. The G-EYE use was related to higher ADR and APC (35.4% vs 23.5% and 0.63 vs 0.36). The last two multicenter randomized trials^[66,67] used G-EYE in combination with a HD colonoscope. In both studies the reported rate of adenoma detection was higher in the G-EYE arm (59% vs 39%, and 49.2% vs 33.8%, respectively).

CRITICAL APPRAISAL AND CONCLUSION

The volume of presented data clearly illustrates the unmet need of optimizing technology to improve colonoscopy performance. The results of the aforementioned studies of novel wide-angle view endoscopes and add-on devices appear promising. Despite some contradictory results the majority of the data are in favor of the new endoscopes/devices regarding polyp and adenoma detection rates, as well as, polyp and adenomas miss rates. However, these data should be interpreted cautiously for a number of reasons:

Firstly, 50% of the reviewed studies have been published as abstracts only. The Extra-wide Angle View Colonoscope and the Third-Eye[®] Panoramic are still under development, while Aer-O-Scope and Third Eye have been abandoned. Moreover, several new colonoscopy add on devices appear in the endoscopy accessories market without having been adequately

evaluated, yet.

Secondly, heterogeneity characterizes the presented studies. Different target populations and lack of a solid integrated design do not allow safe generalization of the results. It should be noted that the plethora of parallel design studies has not enrolled adequate number of participants to detect differences in ADR with sound statistical power. Moreover, the comparator to the examined novelties comprises either standard or high definition endoscopes or both categories, thus adding more confounders to data interpretation. Of note, there are no direct comparisons between new wide angle view colonoscopes and add on devices regarding colonoscopy outcomes, yet and we can hardly expect any to come in the literature soon.

Thirdly, more attention should be paid to studies recruiting asymptomatic subjects at average risk for CRC. This is the particular population in which it is proven that improvement in colonoscopy outcomes (*e.g.*, increased ADR) is correlated to improved patients' outcomes (reduced risk for interval CRC).

Fourthly, it is still unknown if these novelties are of benefit for the low or the average performing endoscopist only or the benefit is also extended to the high detectors. Whether different levels of endoscopists' experience and performance or different endoscopic environment (*e.g.*, academic vs community or private practice) could lead to different acceptance of these technologies and to different levels of quality indicators improvement, pends to be answered.

Finally, cost is an important factor that could influence the widespread use of these novelties. It has been shown that in the era of financial recession expensive technologies used for patients' management are not favored^[68]. In this setting, attachable cuffs and rings present a relatively low cost investment.

Summing up, new wide-angle view endoscopes and add-on devices are promising technologies to improve colonoscopy and patients outcomes. More studies are definitely needed in order to provide answers to the aforementioned open questions. Until conclusive data are obtained, endoscopists should use these novelties in a personalized manner taking into account their availability and staff experience. At the same time, the fundamental principles of colonoscopy like adequate bowel preparation, meticulous inspection independently of endoscope and devices used and suitable withdrawal time should govern our practice.

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