



Role of high definition colonoscopy in colorectal adenomatous polyp detection

Tolga Erim, John M Rivas, Evelio Velis, Fernando Castro

Tolga Erim, Department of Gastroenterology, Cleveland Clinic Florida, Weston, FL 33331, United States

John M Rivas, Department of Internal Medicine, Cleveland Clinic Florida, Weston, FL 33331, United States

Evelio Velis, Health Services Administration Master Program, Barry University, Miami Shores, Florida 33161, United States

Fernando Castro, Department of Gastroenterology, Cleveland Clinic Florida, Weston, FL 33331, United States

Author contributions: Erim T, Rivas JM and Castro F contributed equally to this work; Erim T and Castro F designed the research; Erim T and Rivas J performed the research; Velis E contributed new reagents/analytic tools; Erim T, Rivas J, Velis E and Castro F analyzed the data; Erim T, Rivas JM and Castro F wrote the paper.

Supported by Cleveland Clinic Florida Institution Review Committee

Correspondence to: Dr. Tolga Erim, Department of Gastroenterology, Cleveland Clinic Florida, 2950 Cleveland Clinic Blvd, Weston, FL 33331, United States. erimt@ccf.org

Telephone: +1-954-6595000 Fax: +1-954-6595480

Received: December 17, 2010 Revised: March 11, 2011

Accepted: March 18, 2011

Published online: September 21, 2011

Abstract

AIM: To investigate the rates of polyp detection in a mixed risk population using standard definition (SDC) vs high definition colonoscopes (HDC).

METHODS: This was a retrospective cohort comparative study of 3 colonoscopists who each consecutively performed 150 SDC (307, 200 pixel) and 150 HDC (792, 576 pixels) in a community teaching hospital.

RESULTS: A total of 900 colonoscopies were evaluated (mean age 56, 46.8% men), 450 with each resolution. Polyps of any type were detected in 46.0% of patients using SDC and 43.3% with HDC ($P = 0.42$). There was no significant difference between the overall number of polyps, HDC (397) and SDC (410), detected among

all patients examined, ($P = 0.73$). One or more adenomatous polyps were detected in 24.2% of patients with HDC and 24.9% of patients with SDC colonoscopy ($P = 0.82$). There was no significant difference between HDC ($M = 0.41$) and SDC ($M = 0.42$) regarding adenomatous polyp ($P = 0.88$) or advanced adenoma ($P = 0.56$) detection rate among all patients examined.

CONCLUSION: HDC did not improve yield of adenomatous polyp, advanced adenoma or overall polyp detection in a population of individuals with mixed risk for colorectal cancer.

© 2011 Baishideng. All rights reserved.

Key words: High definition colonoscopy; Colon cancer screening; Adenomatous polyps

Peer reviewer: Dr. Shinji Tanaka, Director, Department of Endoscopy, Hiroshima University Hospital, 1-2-3 Kasumi, Minami-ku, Hiroshima 734-8551, Japan.

Erim T, Rivas JM, Velis E, Castro F. Role of high definition colonoscopy in colorectal adenomatous polyp detection. *World J Gastroenterol* 2011; 17(35): 4001-4006 Available from: URL: <http://www.wjgnet.com/1007-9327/full/v17/i35/4001.htm> DOI: <http://dx.doi.org/10.3748/wjg.v17.i35.4001>

INTRODUCTION

It is estimated that up to 15 million colonoscopies are performed annually in the United States^[1,2]. Colonoscopy and polypectomy have been estimated to prevent 50%-80% of colorectal cancers^[3-5]. However, recent trials have implied a lower level of protection and even a failure of colonoscopy to prevent right sided colon cancer^[6,7]. With adenoma miss rates of up to 20% demonstrated for moderate sized polyps, the potential of improved polyp detection in preventing colon cancer

deaths could be substantial^[8-10].

A considerable effort is being spent on optimizing the yield of colonoscopy with respect to polyp detection. Several technologies such as wide-angle, cap-fitted, retroflexion colonoscopy and Third Eye Retroscope colonoscopy have been used in an attempt to increase mucosa exposure. Various optical enhancing technologies such as chromoendoscopy, narrow-band (NBI) and multi-band imaging, high definition, and autofluorescence have been studied as well. While some have been found to be effective in expert hands in tertiary care centers, many techniques suffer from issues of practicality. The rising demand of colon cancer screening and the advent of several different modalities for this purpose, such as computerized tomography colonoscopy, have stressed the importance of improved efficiency in colonoscopy.

At present, the only technical developments that are readily available and in use in routine practice settings in the United States are wide angle, high definition and NBI/multi-band imaging. High-definition endoscopes have been touted by manufacturers to show markedly clearer images in hopes that this would translate into higher polyp detection rates. In the current study, we present a comparison of polyp detection rate of endoscopists using standard definition and high definition endoscopy systems.

MATERIALS AND METHODS

Patients

Nine hundred consecutive patients who had colonoscopy between May 2007 and May 2008 by three experienced endoscopists (> 6000 colonoscopies each) were selected for analysis retrospectively. Patients were mixed risk and all colonoscopies were performed at the same endoscopy center of a community teaching hospital in Florida, United States. Colonoscopies performed by gastroenterology fellows were excluded from the study. The study was approved by the Institutional Review Board at Cleveland Clinic Florida.

Endoscopy equipment

The standard definition colonoscopies (SDC) were performed with an EPK-1000 processor (Pentax), EC-3430LK, EC-3830LK, EC-3470LK, and EC-3870LK model colonoscopes (Pentax), a 19-inch CRT monitor at a resolution of 640 × 480 producing a 307, 200 pixel image at distance of approximately 2.8 m from the endoscopists. The high definition colonoscopies (HDC) were performed with an EPX-4400 digital processor (Fujinon), EC-450HL5 and EC-450LS5 model colonoscopes (Fujinon), a 32-inch LCD monitor at a resolution of 1032 × 768 producing a 792, 576 pixel image at a distance of approximately 2.8 m from the endoscopists. Both standard definition and high definition colonoscopes had a 140° field of view.

The Fujinon system has the capability of multi-band imaging that produces images similar to the NBI endo-

scopes, commercially termed Fuji Intelligent Chromo Endoscopy (FICE). The difference lies in that the Fujinon system uses software to construct images based on preset RGB wavelength combinations. The NBI systems use optical filters that restrict the bandwidth of a transmitted light signal. Currently available NBI systems utilize 2 narrow-band filters that provide tissue illumination in the blue (415 nm) and green (540 nm) spectrums of light^[11]. The Fujinon equipment has ten factory-determined wavelength preset combinations.

Endoscopic procedures

Data from one hundred fifty consecutive patients who had colonoscopy with standard definition (SD) equipment were collected for three endoscopists from May - October 2007. Following the installation of the HDC system, all endoscopic procedures in our unit were performed exclusively using the high definition (HD) scopes and data was collected from 150 consecutive patients who had colonoscopy by the same three endoscopists from October 2007 - March 2008. The endoscopists were not aware of the study. Bowel preparation agents used were predominantly sodium phosphate and polyethylene glycol based regimens. The procedures were performed under a nurse administered standard sedation with Meperidine and Midazolam or anesthesiologist administered Propofol. Colonoscopy withdrawal times were recorded by the nursing staff.

Endoscopists were free to use the multi-band feature on the HDC system as needed. The system was initially set on the factory default preset of 0, which produced an image restricted to the following wavelengths: R 500 nm, G 445 nm, and B 415 nm. A push-button on the handle of the colonoscope was programmed to enable switching between conventional white-light image and the preset multi-band image. Endoscopists were also free to change to a different factory preset according to their preferences. The study was designed prior to arrival of the high definition system; however, data collection was started afterwards.

Data collection

The data was collected from electronic medical records, procedure nursing notes, procedure reports, and pathology reports. The numbers of detected polyps recorded on the procedure reports were corroborated with the pathology reports and the nursing notes. The main outcome parameter was the polyp detection rate in the two groups. Secondary outcome measures included: detection rates of adenomatous polyps, advanced adenomas, and cancer. Advanced adenoma was defined as adenomatous polyps having one or more features of: > 1 cm in diameter, high-grade dysplasia, and villous histology. Additional data was collected with regards to patient age, gender, race, indication for colonoscopy, polyp location, procedure time, withdrawal time, type of sedation, and prep quality.

Table 1 Patient Characteristics *n* (%)

Parameters	HD group (<i>n</i> = 450)	SD ² group (<i>n</i> = 450)	<i>P</i> value
Patients			
Mean age, years (SD ¹)	55 (± 12.5)	56 (± 11.4)	0.21
Men	213 (47.3)	208 (46)	0.86
Race			
White	233 (51.8)	281 (62.4)	0.10
African American	49 (10.9)	53 (11.8)	0.75
Hispanic	139 (30.9)	95 (21.1)	0.01
Others	29 (6.4)	21 (4.7)	0.31
Indication			
Screening	216 (48.0)	173 (38.4)	0.07
Non-screening	234 (52.0)	277 (61.6)	0.13
Cecal intubation	433 (96.2)	438 (97.3)	0.92
Poor prepare	14 (3.1)	17 (3.8)	0.72
Withdrawal all procedures, min (SD ¹)	11.3 (± 6.1)	10.8 (± 5.6)	0.20
Withdrawal non-polypectomy, min (SD ¹)	10.0 (± 5.9)	9.2 (± 4.2)	0.02

¹Standard deviation; ²Standard definition; HD: high definition; min: Minutes.

Table 2 Detection of all polyps, adenomas, and cancer *n* (%)

Parameters	HD group (<i>n</i> = 450)	SD ² group (<i>n</i> = 450)	<i>P</i> value
Total polyps detected	397	410	0.73
Non-adenomas	196 (51.3)	209 (52.0)	0.81
Non-advanced adenomas	150 (39.3)	150 (37.3)	0.84
Advanced adenomas	34 (8.9)	40 (10.0)	0.60
Cancer	2 (0.5)	3 (0.7)	1.00
Pathology not identified	15 (3.8)	8 (2.0)	0.14
< 6 mm	325 (81.9)	340 (82.9)	0.96
6-10 mm	50 (12.6)	44 (10.7)	0.45
> 10 mm	22 (5.5)	24 (5.9)	1.00
Size not specified	0	2	0.50
All patients			
With non-adenomas	84 (18.7)	92 (20.4)	0.56
With non-advanced adenomas	84 (18.7)	83 (18.4)	1.00
With advanced adenomas	25 (5.6)	29 (6.4)	0.67
With cancer	2 (0.4)	3 (0.7)	1.00
Polyps per pt, mean (SD ¹)	0.88 (± 1.63)	0.91 (± 1.41)	0.77
Adenomas per pt, mean (SD ¹)	0.41 (± 1.04)	0.42 (± 0.94)	0.88
Advanced adenoma per pt, mean (SD ¹)	0.076 (± 0.35)	0.089 (± 0.38)	0.56
Adenocarcinoma per pt (mean)	0.004	0.006	
Screening patients, <i>n</i>	216	173	0.07
With non-adenomas	49 (22.7)	36 (20.8)	0.81
With non-advanced adenomas	41 (19.0)	32 (18.5)	1.00
With advanced adenomas	13 (6.0)	14 (8.1)	0.55
With cancer	1 (0.5)	0 (0.0)	1.00
Non-screening patients, <i>n</i>	234	277	0.13
With non-adenomas	35 (15.0)	56 (20.2)	0.59
With non-advanced adenomas	43 (18.4)	51 (18.4)	0.21
With advanced adenomas	12 (5.1)	15 (5.4)	1.00
With cancer	1 (0.4)	3 (1.1)	0.63

¹Standard deviation; ²Standard definition; HD: high definition; Per pt: Per patient.

Statistical analysis

The Statistical Package for Social Sciences (SPSS 16.0) was used in order to organize, validate and analyze the collected data. Quantitative data were summarized using mean values (M) and standard deviation; Student's *t* test were performed in order to detect significant differences between colonoscope types; equality of variances was inspected using Levene's tests. We examined associations between categorical variables, performing χ^2 tests or Fisher's exact test when appropriate.

RESULTS

A total of 900 colonoscopies were evaluated, comparing 450 patients each who had colonoscopy with SDC equipment and HDC equipment. Each endoscopist performed 300 colonoscopies equally divided between standard and high definition procedures. The mean age of the study population was 56, and 46.8% were men. There were no statistically significant differences in patient characteristics between the two groups with the exception of a higher number of Hispanic patients and those that had screening colonoscopy in the HDC group (Table 1). However, there was no overall difference in adenomatous polyp detection rate in Hispanics (23.9%) *vs* Non-Hispanics (24.6%) ($P = 0.86$) and the screening (25.7%) and non-screening (21.7%) groups ($P = 0.18$).

Cecal intubation, bowel prep quality and withdrawal times were also not statistically significantly different between the HDC and SDC groups. The cecum was reached in 96.7% of all cases. Average withdrawal time was 11.1 min, which included polypectomy time.

Polyps of any type were detected in 46.0% of patients with SDC and 43.3% of those patients who had HDC ($P = 0.42$). There was no significant difference between the overall number of polyps, HDC (397) and SDC (410), detected among all patients examined ($P = 0.73$). One or more adenomatous polyps were detected in 24.2% of patients with HDC and 24.9% of patients with SDC ($P = 0.82$). There was no significant difference between HDC (M = 0.41) and SDC (M = 0.42) regarding adenomatous polyp detection rate among all patients examined ($P = 0.88$). In addition, there was no significant difference between the study groups regarding advanced adenoma polyp detection rates ($P = 0.60$) or cancer detection rate among all patients examined ($P > 0.05$) (Table 2). There was no difference in polyp detection rates when each individual endoscopist's HDC and SDC detection rates were compared (data not shown).

Polyps detected during the procedures were also analyzed according to size. There was no significant difference between the detected number of polyps of sizes < 6 mm, 6-10 mm, and >10 mm in the HDC and SDC groups.

Gender was shown to be a significant variable as men in this study were found to have a higher incidence of all polyps ($P < 0.01$), adenomatous polyps ($P < 0.01$) and

Table 3 Detection of all polyps, adenomas and cancer, with respect to gender *n* (%)

Parameters	HD group (<i>n</i> = 450)	SD group (<i>n</i> = 450)	<i>P</i> value
Male			
Total polyps detected	264	245	0.74
Patients	213	208	0.86
With non-adenomas	42 (19.7)	46 (22.1)	0.48
With non-advanced adenomas	52 (24.4)	43 (20.7)	0.50
With advanced adenomas	17 (8.0)	21 (10.1)	0.50
With cancer	1 (0.5)	2 (1.0)	0.62
Female			
Total polyps detected	133	165	0.21
Patients	237	242	0.87
With non-adenomas	42 (17.7)	46 (19.0)	0.82
With non-advanced adenomas	32 (13.5)	40 (16.5)	0.45
With advanced adenomas	8 (3.4)	8 (3.3)	1.00
With cancer	1 (0.4)	1 (0.4)	1.00

SD: Standard definition; HD: High definition.

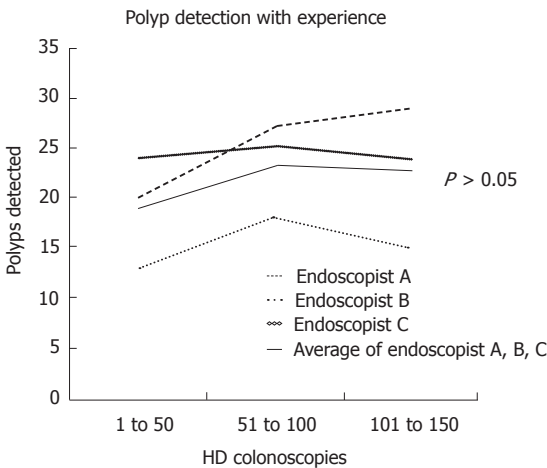


Figure 1 Comparison of polyp detection per endoscopist with gained high definition experience. HD: High definition.

Table 4 Comparison of standard *vs* high definition adenomatous polyp detection studies to date

Study	Date	Method	<i>n</i>	Colonoscope (resolution) (Angle) (NBI capable)		<i>P</i> value for adenoma detection rates
				Group 1	Group 2	
East <i>et al</i> ^[13]	2008	Prospective nonrandomized	130	Olympus SD 140	Olympus HD 140	0.200
Pellise <i>et al</i> ^[12]	2008	Prospective randomized	620	Olympus SD 140	Olympus HD 170	0.850
Tribonias <i>et al</i> ^[14]	2010	Prospective randomized	390	Olympus SD 140	Olympus HD 170	0.160
Burke <i>et al</i> ^[16]	2010	Retrospective	852	Olympus SD 140	Olympus HD 170	0.360
Buchner <i>et al</i> ^[15]	2010	Retrospective	2430	Olympus SD 140	Olympus HD 170 NBI	0.012

NBI: Narrow-band imaging; SD: Standard definition; HD: High definition.

advanced adenomas ($P < 0.01$). However, this disparity was consistent in both cohorts with no statistically significant difference between the HDC and SDC groups (Table 3).

We found that the overall polyp and adenoma detection rates did not change significantly between the first, second or third 50 HDC performed by our endoscopists when the three were coned ($P > 0.05$). Therefore, there does not seem to be a learning effect associated with use of HDC with on demand multi-band imaging capability by endoscopists who had not used them before (Figure 1).

DISCUSSION

Our goal in this study was to assess the performance of HDC with regards to polyp detection. Five prior studies have compared adenoma detection rate between standard and high definition white light colonoscopy with conflicting results. There have been methodological and technical differences between the studies (Table 4) with all using Olympus colonoscopes, whereas our study is the first performed using Fujinon HDC with FICE. Among the three prospective studies, Pellisé *et al*^[12] had the largest patient population. It was a prospective randomized controlled trial of 620 patients conducted in

Spain involving seven colonoscopists. Patients were randomized to either HDC with wide angle (170°) field or SDC with 140° view, with the investigators finding no difference in adenoma detection rate between the study groups (HDC 26% *vs* SDC 25%, $P = 0.85$). The study by East *et al*^[13] was not randomized, consisting of 130 patients who underwent either HDC with 140° view or SC 140° view by a single colonoscopist. Although HDC did not improve the yield of adenomatous polyp detection, there was a trend in this direction (71% *vs* 60%). The Tribonias *et al*^[14] study randomized 390 patients prospectively into HDC with wide angle *vs* SDC groups and, although there was a significant difference between the two groups with regards to overall rate of polyp detection, (HDC 63% *vs* SDC 53%, $P = 0.03$), there was no significant difference demonstrated in the detection of rate of adenomas (HDC 58% *vs* SDC 50%, $P = 0.16$).

The largest patient population study on this topic, by Buchner *et al*^[15], was retrospective involving 2430 patients in two arms: HDC and SDC. The HRC were 170° wide-angle and NBI was used as needed. The SDC in the study had a 140° view and did not have NBI. The study found that the HDC were able to detect a statistically significant higher number of adenomatous polyps compared with SDC (28.8% *vs* 24.3%, $P = 0.012$). The most recently

published study was by Burke *et al.*^[16] and consisted of 426 individuals in each group and found no advantage of HDC in overall polyp detection rate, adenomas or advanced adenomas.

In our study, we found no difference in detection rates of overall polyps, adenomas, advanced adenomas, and cancer between the HD and SD groups. There was no difference in polyp detection rates when each individual endoscopist's HD and SD detection rates were compared despite having used 32-inch LCD high-resolution monitors with the Fujinon system whereas 19-inch CRT monitors were used with the standard definition colonoscopes. Although there are considerable methodological differences between the Pellisé *et al.*^[12] and our study, both studies show very similar results and conclusions. In fact, their adenomatous polyp detection rates are nearly identical to ours in the SD and HD arms: 0.45 ± 1.07 *vs* our 0.41 ± 1.04 adenomas per patient in SD and 0.43 ± 0.87 *vs* our 0.42 ± 0.94 adenomas per patient in HD. Our polyp detection rates are well in line with several other studies of white light colonoscopy with regards to prevalence of adenomas, advanced adenomas, cancer, and gender differences^[17-19]. We were also able to demonstrate that polyp detection rate did not improve as the endoscopist experience with HDC increased by comparing adenoma detection in consecutive groups of 50 colonoscopies ($P > 0.05$). This lack of learning effect was also demonstrated by Adler *et al.*^[20] in 2008 in a prospective randomized study of NBI *vs* conventional colonoscopy for adenoma detection. Although prior studies were meant to compare HD and SD, the HD equipment used in these reports also had a wide angle field of view and the study by East is the only one that used 140° scopes in both arms, but it was underpowered for detecting small differences in polyp detection rate. Similar to East, our study design eliminates the confounding factor of the wide angle field of view by using 140° scopes in the HD and SD groups.

There is significant variability amongst endoscopists in adenoma detection rates, making the endoscopist probably the most important variable in adenoma detection rate^[21]. We tried to minimize the impact factor of the endoscopist by assigning an equal number of overall cases per endoscopist (300) and dividing these equally amongst the study groups. Our study has a significant advantage in this.

It can be argued that our study's retrospective design was a limitation, but it may have also served to reduce endoscopist bias. Endoscopist bias is an inherent limitation of nearly all prospective colonoscopy study designs since the equipment cannot be hidden from the performer of the examination. A second limitation is that the population was a mixed-risk sample and there were slight differences with respect to Hispanics and screening patients. However, there was no statistically significant difference in the detection of adenoma, advanced adenoma or cancer between the Hispanic *vs* Non-Hispanic and screening *vs* non-screening groups. In fact, there was a slightly higher prevalence of adenomas in the populations that were overrepresented in the HD group. This would have

worked to bias the results in favor of HDC had it been a significant difference. In summary, the results of our study are relevant to most practices as the majority of the new colonoscopy equipment purchased in the future will have HD and NBI or multi-band imaging capabilities. Until recently, evidence regarding the potential of this new technology in improving yield of polyp detection was lacking. Complementing the results of Pellisé, and Burke, our study concludes that HDC with multi-band imaging capability does not detect more total polyps, adenomas, advanced adenomas or cancer. For now at least, the endoscopist and not the equipment used, continues to be the major factor in polyp detection.

COMMENTS

Background

It is estimated that up to 15 million colonoscopies are performed annually in the United States. Colonoscopy and polypectomy have been estimated to prevent 50%-80% of colorectal cancers. A considerable effort is being spent on optimizing the yield of colonoscopy with respect to polyp detection. At present, high definition colonoscopy (HDC) is being widely adapted but whether or not it impacts detection of colon polyps is debatable.

Research frontiers

HDC is widely touted by manufacturers to improve polyp detection. Yet, several studies have compared detection of polyps with standard definition colonoscopy (SDC) *vs* HDC with variation in results.

Innovations and breakthroughs

A major confounding factor in previous studies on this subject is that the endoscopists are aware of the high definition equipment and this may have led to bias in polyp detection rates. This study uniquely eliminates the issue of endoscopist bias by using a retrospective model of consecutive colonoscopies.

Applications

By providing added evidence of HDC's role in polyp detection, this study may shift opinion further to the side of lack of benefit in improving yield.

Terminology

Standard colonoscopes typically use 640×480 resolution monitors producing a 307, 200 pixel image. The HDC with high resolution monitors can produce a 1032×768 resolution and a 792, 576 pixel image.

Peer review

The authors examined whether or not HDC resulted in detection of more polyps. The results show no significant difference in polyp detection between SDC and HDC. The results complement the conclusion of other recent studies in this field and suggest that high definition by itself may not improve yield of polyp detection.

REFERENCES

- 1 Seeff LC, Richards TB, Shapiro JA, Nadel MR, Manninen DL, Given LS, Dong FB, Wings LD, McKenna MT. How many endoscopies are performed for colorectal cancer screening? Results from CDC's survey of endoscopic capacity. *Gastroenterology* 2004; **127**: 1670-1677
- 2 Seeff LC, Manninen DL, Dong FB, Chattopadhyay SK, Nadel MR, Tangka FK, Molinari NA. Is there endoscopic capacity to provide colorectal cancer screening to the unscreened population in the United States? *Gastroenterology* 2004; **127**: 1661-1669
- 3 Winawer SJ, Zauber AG, Ho MN, O'Brien MJ, Gottlieb LS, Sternberg SS, Wayne JD, Schapiro M, Bond JH, Panish JF. Prevention of colorectal cancer by colonoscopic polypectomy. The National Polyp Study Workgroup. *N Engl J Med* 1993; **329**: 1977-1981
- 4 Citarda F, Tomaselli G, Capocaccia R, Barcherini S, Crespi M. Efficacy in standard clinical practice of colonoscopic pol-

- ypectomy in reducing colorectal cancer incidence. *Gut* 2001; **48**: 812-815
- 5 **Thiis-Evensen E**, Hoff GS, Sauar J, Langmark F, Majak BM, Vatn MH. Population-based surveillance by colonoscopy: effect on the incidence of colorectal cancer. Telemark Polyp Study I. *Scand J Gastroenterol* 1999; **34**: 414-420
- 6 **Robertson DJ**, Greenberg ER, Beach M, Sandler RS, Ahnen D, Haile RW, Burke CA, Snover DC, Bresalier RS, McKeown-Eyssen G, Mandel JS, Bond JH, Van Stolk RU, Summers RW, Rothstein R, Church TR, Cole BF, Byers T, Mott L, Baron JA. Colorectal cancer in patients under close colonoscopic surveillance. *Gastroenterology* 2005; **129**: 34-41
- 7 **Baxter NN**, Goldwasser MA, Paszat LF, Saskin R, Urbach DR, Rabeneck L. Association of colonoscopy and death from colorectal cancer. *Ann Intern Med* 2009; **150**: 1-8
- 8 **Rex DK**, Cutler CS, Lemmel GT, Rahmani EY, Clark DW, Helper DJ, Lehman GA, Mark DG. Colonoscopic miss rates of adenomas determined by back-to-back colonoscopies. *Gastroenterology* 1997; **112**: 24-28
- 9 **Hixson LJ**, Fennerty MB, Sampliner RE, Garewal HS. Prospective blinded trial of the colonoscopic miss-rate of large colorectal polyps. *Gastrointest Endosc* 1991; **37**: 125-127
- 10 **Heresbach D**, Barrioz T, Lapalus MG, Coumaros D, Bauret P, Potier P, Sautereau D, Boustière C, Grimaud JC, Barthélémy C, Sée J, Serraj I, D'Halluin PN, Branger B, Ponchon T. Miss rate for colorectal neoplastic polyps: a prospective multicenter study of back-to-back video colonoscopies. *Endoscopy* 2008; **40**: 284-290
- 11 **Song LM**, Adler DG, Conway JD, Diehl DL, Farraye FA, Kantsevov SV, Kwon R, Mamula P, Rodriguez B, Shah RJ, Tierney WM. Narrow band imaging and multiband imaging. *Gastrointest Endosc* 2008; **67**: 581-589
- 12 **Pellisé M**, Fernández-Esparrach G, Cárdenas A, Sendino O, Ricart E, Vaquero E, Gimeno-García AZ, de Miguel CR, Zabalza M, Ginès A, Piqué JM, Llach J, Castells A. Impact of wide-angle, high-definition endoscopy in the diagnosis of colorectal neoplasia: a randomized controlled trial. *Gastroenterology* 2008; **135**: 1062-1068
- 13 **East JE**, Stavrinidis M, Thomas-Gibson S, Guenther T, Tekkis PP, Saunders BP. A comparative study of standard vs. high definition colonoscopy for adenoma and hyperplastic polyp detection with optimized withdrawal technique. *Aliment Pharmacol Ther* 2008; **28**: 768-776
- 14 **Tribonias G**, Theodoropoulou A, Konstantinidis K, Vardas E, Karmiris K, Chroniaris N, Chlouverakis G, Paspatis GA. Comparison of standard vs high-definition, wide-angle colonoscopy for polyp detection: a randomized controlled trial. *Colorectal Dis* 2010; **12**: e260-e266
- 15 **Buchner AM**, Shahid MW, Heckman MG, McNeil RB, Cleveland P, Gill KR, Schore A, Ghabril M, Raimondo M, Gross SA, Wallace MB. High-definition colonoscopy detects colorectal polyps at a higher rate than standard white-light colonoscopy. *Clin Gastroenterol Hepatol* 2010; **8**: 364-370
- 16 **Burke CA**, Choure AG, Sanaka MR, Lopez R. A comparison of high-definition vs conventional colonoscopes for polyp detection. *Dig Dis Sci* 2010; **55**: 1716-1720
- 17 **Kanna B**, Schori M, Azeez S, Kumar S, Soni A. Colorectal tumors within an urban minority population in New York City. *J Gen Intern Med* 2007; **22**: 835-840
- 18 **Schoenfeld P**, Cash B, Flood A, Dobhan R, Eastone J, Coyle W, Kikendall JW, Kim HM, Weiss DG, Emory T, Schatzkin A, Lieberman D. Colonoscopic screening of average-risk women for colorectal neoplasia. *N Engl J Med* 2005; **352**: 2061-2068
- 19 **Adler A**, Aschenbeck J, Yenerim T, Mayr M, Aminimalai A, Drossel R, Schröder A, Scheel M, Wiedenmann B, Rösch T. Narrow-band vs white-light high definition television endoscopic imaging for screening colonoscopy: a prospective randomized trial. *Gastroenterology* 2009; **136**: 410-6.e1; quiz 715
- 20 **Adler A**, Pohl H, Papanikolaou IS, Abou-Rebyeh H, Schachschal G, Veltzke-Schlieker W, Khalifa AC, Setka E, Koch M, Wiedenmann B, Rösch T. A prospective randomised study on narrow-band imaging vs conventional colonoscopy for adenoma detection: does narrow-band imaging induce a learning effect? *Gut* 2008; **57**: 59-64
- 21 **Chen SC**, Rex DK. Endoscopist can be more powerful than age and male gender in predicting adenoma detection at colonoscopy. *Am J Gastroenterol* 2007; **102**: 856-861

S- Editor Sun H L- Editor Rutherford A E- Editor Xiong L