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**Retrospective Study**

**Capsule endoscopy and single-balloon enteroscopy in small bowel diseases: Competing or complementary?**

Ma JJ *et al***.** Evaluating small bowel disease by CE and/or SBE

Jing-Jing Ma, Ying Wang, Xiao-Min Xu, Jie-Wen Su, Wen-Yu Jiang, Jian-Xia Jiang, Lin Lin, Dao-Quan Zhang, Jing Ding, Li Chen, Ting Jiang, Ying-Hong Xu, Gui Tao, Hong-Jie Zhang

**Jing-Jing Ma, Xiao-Min Xu, Jie-Wen Su, Wen-Yu Jiang, Jian-Xia Jiang, Lin Lin, Dao-Quan Zhang, Jing Ding, Li Chen, Ting Jiang, Ying-Hong Xu, Gui Tao, Hong-Jie Zhang,** Department of Gastroenterology, The First Affiliated Hospital of Nanjing Medical University, Nanjing 210029, Jiangsu Province, China

**Ying Wang,** Department of Gastroenterology, Jiangsu Provincial Hospital, Nanjing 210024, Jiangsu Province, China

**Author contributions:** Ma JJ, Wang Y and Xu XM contributed equally to this work; Ma JJ and Wang Y performed this study; Ma JJ, Xu XM, Su JW and Jiang WY carried out data analysis and wrote the paper; Ma JJ, Wang Y, Jiang JX, Lin L, Zhang DQ, Ding J, Chen L, Jiang T, Xu YH, Tao G, Zhang HJ performed capsule endoscopy and single balloon enteroscopy; Ma JJ and Zhang HJ designed the study. Zhang HJ supervised the report; all authors have read and approved the final version to be published.

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**Correspondence to: Hong-Jie Zhang, PhD,** Department of Gastroenterology, The First Affiliated Hospital of Nanjing Medical University, 300 Guangzhou Road, Nanjing 210029, Jiangsu Province, China. hjzhang06@163.com

**Telephone:** +86-25-83718836-6920

**Fax:** +86-25-83674636

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**Abstract**

***AIM***

To evaluate diagnostic yields of capsule endoscopy (CE) and/or single-balloon enteroscopy (SBE) in patients with suspected small bowel diseases.

***METHODS***

We retrospectively analyzed 700 patients with suspected small bowel diseases from September 2010 to March 2016. CE, SBE, or SBE with prior CE was performed in 401, 353, and 47 patients, respectively. Data from clinical and endoscopy records were collected for analysis. Indications, procedure times, diagnostic yields, and complications were summarized and evaluated.

***RESULTS***

The overall diagnostic yield for the CE group was 57.6%. The diagnostic yield of CE in patients with obscure gastrointestinal bleeding (OGIB) was significantly greater than that in patients with no bleeding (70.5% *vs* 43.8%, *P* < 0.01). The overall diagnostic yield of SBE was 69.7%. There was no difference in the diagnostic yield of SBE between patients with OGIB and those with no bleeding (72.5% *vs* 68.9%, *P* = 0.534). Forty-seven patients underwent CE prior to SBE. Among them, the diagnostic yield of SBE with positive findings on prior CE was 93.3%. In addition, SBE detected two cases with superficial ulcer and erosive lesions in the small bowel, which were missed by CE. However, one case with lymphoma and two with Crohn’s disease were not confirmed by SBE. The rate of capsule retention was 2.0%. There were no significant complications during or after the SBE examinations.

***CONCLUSION***

SBE is a safe and effective technique for diagnosing small bowel diseases. SBE with prior CE seemed to improve diagnostic yield of small bowel diseases.

**Key words:** Capsule endoscopy; Single-balloon enteroscopy; Balloon-assisted enteroscopy; Small bowel diseases; Diagnosis

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**Core tip:** The aims of this study were to evaluate diagnostic yields associated with capsule endoscopy (CE), single-balloon enteroscopy (SBE), or their combined use in patients with suspected small bowel diseases, and to demonstrate the appropriate diagnostic algorithms for diagnosing different small bowel diseases. This study revealed the diagnostic yield of SBE with positive findings on prior CE was high (93.3%). CE followed by SBE represents an especially effective strategy for determining the cause of small bowel disease when findings from an initial CE examination are indeterminate.

Ma JJ, Wang Y, Xu XM, Su JW, Jiang WY, Jiang JX, Lin L, Zhang DQ, Ding J, Chen L, Jiang T, Xu YH, Tao G, Zhang HJ. Capsule endoscopy and single-balloon enteroscopy in small bowel diseases: Competing or complementary? *World J Gastroenterol* 2016; In press

**INTRODUCTION**

The small bowel has long been considered a “black box” for gastroenterologists because of its length and complex anatomy. Before 2000, it was not possible to reach most of the small bowel using conventional endoscopic techniques. The diagnosis of small bowel diseases has been a challenge for gastroenterologists. The development of capsule endoscopy (CE) and balloon-assisted enteroscopy (BAE) represents a decisive breakthrough in the field. CE is painless and can be used to explore the entire small bowel in a single examination. It is considered the best choice for an initial diagnostic examination when a patient is suspected of small bowel disease[1-3]. However, CE has some technical limitations, including lack of therapeutic capability and risk of capsule retention.

BAE was introduced as a breakthrough technique for examining the deep small bowel, and comprises double-balloon endoscopy (DBE) and single-balloon enteroscopy (SBE). DBE was introduced in 2001 and is considered the standard technique of deep endoscopy to visualize the small bowel. DBE allows the endoscopist to visualize the small bowel and to perform therapeutic intervention. However, preparation and handling of DBE is complex. SBE was introduced in 2008, which has a simpler and easier-to-handle small bowel endoscopy system. Compared with DBE, SBE may be less efficient in terms of depth of insertion and complete visualization of the small bowel. However, some studies have shown that SBE is not inferior to DBE in relation to diagnostic yield[4]. Both CE and BAE are reported to have similarly high diagnostic yields for small bowel diseases[5,6]. DBE is considered an effective complementary technique, which can be used after initial diagnostic CE examination[7]. However, there are limited data on the role of CE in comparison and in combination with SBE for assessment of small bowel diseases[8].

We performed a retrospective study to (1) compare the diagnostic yields of CE, SBE, or their combined use; (2) determine their performance characteristics in patients with suspected small bowel diseases; and (3) demonstrate the appropriate diagnostic algorithms for different small bowel diseases.

**MATERIALS AND METHODS**

***Study design and patient selection***

We retrospectively analyzed the records of 700 patients suspected of small bowel diseases who underwent CE and/or SBE between September 2010 and March 2016 at the First Affiliated Hospital of Nanjing Medical University. All patients underwent routine clinical examinations and laboratory tests (including hemoglobin level and stool tests), abdominal ultrasound or computed tomography (CT), upper gastrointestinal endoscopy and colonoscopy. CE and SBE were performed after obtaining informed consent from the patients. Indications for the study included obscure gastrointestinal bleeding (OGIB), abdominal pain, diarrhea, or other symptoms. The characteristics of all patients and procedures were extracted from electronic medical records and the endoscopy reporting system. Final diagnosis was based on SBE findings, CE findings, surgical pathology, and/or clinical follow-up. The diagnostic yield was calculated by dividing the total number of patients who underwent the procedure by the number of cases with positive findings that could explain the patient’s symptoms. This study was approved by the Ethics Committee of the First Affiliated Hospital of Nanjing Medical University.

***CE procedure***

CE studies were performed using the OMOM CE system (Jinshan Science and Technology Company, Chongqing, China) or MiroCam TM system (IntroMedic, Seoul, South Korea). Each patient underwent bowel preparation with 3 L polyethylene glycol solution the day before the procedure, and fasted overnight. CE data collected included gastric transit time (GTT), small bowel transit time (SBTT), abnormal findings during the procedure, total recording time, quality of the bowel preparation, and complete visualization rate of the small bowel. The complete video of each CE examination was viewed by two independent and experienced gastroenterologists.

***SBE procedure***

SBE procedures were performed using the SBE endoscope system (SIF-Q260; Olympus, Tokyo, Japan). For anterograde SBE, patients generally needed an overnight fast. For a retrograde approach, patients underwent bowel preparation with 3 L polyethylene glycol solution the day before the procedure, and fasted overnight. The examination itself was carried out with conventional sedation with propofol and opioid. All procedures were performed by one of three experienced endoscopists; each of whom had previously conducted at least 50 SBE procedures. Procedures were carried out *via* the anterograde or retrograde approach, depending on whether the suspected pathology was in the proximal or distal small bowel.

***Statistical analysis***

Continuous data are expressed as mean ± standard deviation and range, and categorical data are showed as percentages. Student’s *t* test was used to compare age distributions between the CE and SBE groups. The χ2 test was used to compare positive-detection rates and sex distribution between the CE and SBE groups. *P* < 0.05 was considered statistically significant. All statistical analyses were conducted using SPSS version 20.0 (SPSS Inc., Chicago, IL, United States).

**RESULTS**

***Patient characteristics***

Seven hundred patients who underwent CE and/or SBE were reviewed in the present study. Of these, 401 individuals (248 male, 153 female; mean age, 49.4 ± 16.0 years) underwent 404 CE procedures; 353 individuals (235 male, 118 female; mean age, 42.1 ± 15.8 years) underwent 419 SBE procedures; 47 individuals (38 male, 9 female; mean age, 45.3 ± 15.1 years) underwent both CE and SBE (CE first) (Table 1). Main indications for CE and/or SBE were OGIB (37.1%, 243 with overt OGIB and 17 with occult OGIB), chronic abdominal pain (42.6%), chronic diarrhea (11.3%), and other complaints (9.0%). The demographic data of these patients are shown in Table 1. The mean age of these patients was 46.1 ± 16.5 years (range, 11–85 years). There was no significant difference in sex distribution between the CE and SBE group (*P* = 0.177). The average age of the CE group was older compared with the SBE group (*P*< 0.01).

***CE***

A total of 401 patients underwent 404 CE procedures. Complete visualization of the small bowel was achieved in 73.5% (297/404). The mean recording time was 555 ± 115 min (192–721 min). Mean GTT was 51 ± 62 min (range 1–565 min) and mean SBTT was 352 ± 157 min (range 33–715 min). The overall diagnostic yield for small bowel disease by CE was 57.6% (231/401). The main findings included: mucosal erosion and superficial ulcer in 98 patients (42.4%), angiodysplasia in 67 (29.0%), Crohn’s disease in 26 (11.3%), and masses (tumors and polyps) in 24 (10.4%). Other ﬁndings were parasites in 5.6% (13/231), diverticulum in 2.6% (6/231) and ongoing bleeding in 2.2% (5/231) (Table 2, Supplementary Figure 1). The diagnostic yield of CE in patients with OGIB was greater than that in those with no bleeding (70.5% *vs* 43.8%, *P* < 0.01) (Table 3). For eight patients, capsules were retained at the lesion sites, leading to a capsule-retention rate of 2.0%. Five of these patients were diagnosed with Crohn’s disease, two were diagnosed with lymphoma, and another patient had diverticulum with ulceration. Retained capsules were subsequently removed *via* surgery. In this study, we also compared with whether there is any difference between the OMOM system and MiroCam system. The complete visualization of the small bowel was achieved in 72.4% (197/272) with OMOM and 78.8% (104/132) with MiroCam (*P =* 0.169). The overall diagnostic yield for small bowel diseases was 57.2% (155/271) by OMOM CE and 60.8% (79/130) by MiroCam (*P =* 0.497). The results showed no significant difference regarding to the rates of complete small-bowel examination or diagnostic yields between MiroCam and OMOM capsule endoscopy (Supplementary Table 1).

***SBE***

A total of 419 SBE procedures were performed in 353 patients: 98 anterograde and 321 retrograde, and 24 combined anterograde and retrograde SBEs were conducted. No adverse events occurred during or after these procedures. The mean examination time was 65.5 ± 26.6 min (15–120 min). The overall diagnostic yield for small bowel disease by SBE was 69.7% (246/353). The main findings were as follows: mucosal erosion and superficial ulceration in 111 patients (45.1%), Crohn’s disease in 86 (35.0%), angiodysplasia in 21 (8.5%), and masses (tumors and polyps) in 26 (10.6%). Other ﬁndings were diverticulum (1.2%, 3/246) and parasites (0.4%, 1/246) (Table 2, Supplementary Figure 2). The diagnostic yield for small bowel diseases by SBE was greater than that by CE (69.7% *vs* 57.6%). There was no significant difference in the diagnostic yield of SBE between patients with OGIB and those with no bleeding (72.5% *vs* 68.9%, *P* = 0.534). In a subgroup analysis, the diagnostic yield for OGIB by SBE was similar to by CE (72.5% *vs* 70.5%). In addition, in patients with no bleeding, the diagnostic yield for small bowel diseases tended to be greater using SBE compared with CE (68.9% *vs* 43.8%, *P* < 0.01) (Table 3).

***CE combined with SBE***

Forty-seven patients underwent CE (including 30 with OGIB, 11 with abdominal pain, 1 with diarrhea, and 5 with weight loss) and were subsequently subjected to SBE. The small intestinal findings on SBE in patients with negative evaluation or definite findings on CE are shown in Table 4. Of 47 patients, 45 had positive findings by CE examination, which was followed by SBE, and 42 had positive findings by SBE examination. The diagnostic yield of SBE with findings on prior CE was 93.3% (42/45), which was a high diagnostic yield. Two cases of superficial ulcer and mucosal erosion that were missed by CE were found by SBE. However, CE also detected one mass and two cases of Crohn’s disease that lesions only in the small intestine were not detected by SBE.

**DISCUSSION**

Current options for diagnosing small bowel diseases include push enteroscopy, CE, DBE, SBE, and intraoperative enteroscopy. Push enteroscopy has a limited depth of insertion. Intraoperative enteroscopy is the most invasive method and its use has diminished with the development of CE and BAE. CE is widely used to screen for various small bowel diseases, but this procedure is limited by a lack of therapeutic ability, and imprecise localization and collection of biopsy specimens. DBE is a deep enteroscopy technique that overcomes these shortcomings. However, there are also issues with the DBE technique such as complex preparation and handling procedure. SBE was recently introduced as an alternative deep enteroscopy technique. Some studies have demonstrated that SBE can also provide high diagnostic yield and enable many therapeutic interventions[9-12]. In the present study, the diagnostic yield of SBE for small bowel diseases was 69.7%, which suggests that SBE has a high diagnostic yield. However, previous studies showed the complete visualization rate of the small bowel using SBE was lower than that using DBE (the rate of complete enteroscopy using DBE was 40%–80%, while using SBE was 0%–25%)[13]. This indicated that CE, DBE and SBE all have advantages and limitations. Therefore, it is important to select the appropriate diagnostic algorithms when small bowel disease is suspected. This decision should be made on a case-by-case basis and depends on the clinical scenario, diagnostic yield, involved risks, availability, and patient preference.

In this study, OGIB was a common indication for small bowel endoscopy. Unless contraindicated, CE is recommended as the initial diagnostic test for patients with suspected OGIB[14], because CE is minimally invasive, easily tolerated, and can theoretically visualize the entire small bowel. Here, the diagnostic yield of CE for small bowel abnormalities in patients with OGIB was 70.5%. This result is supported by previous studies[15,16]. CE and BAE are also considered complementary procedures for the evaluation and treatment of OGIB[17-20]. Previous studies have supported using the non-invasive CE technique for patients with OGIB, with a subsequent DBE examination if necessary[21]. In the present study, CE found small bowel lesions in 30 patients with OGIB who were subsequently subjected to SBE. Twenty-eight patients had confirmed diagnosis by SBE examination. If false-negative rates were considered, our data suggested that both SBE and CE did miss some lesions. This study supported that CE evaluation remains the preferred initial strategy for patients with OGIB because of its relative non-invasiveness and acceptable diagnostic yield. However, SBE is useful in cases in which the CE result is ambiguous, and further examination or a biopsy is required. For patients with no bleeding, previous studies have not detected a difference between DBE and CE in identifying small bowel abnormalities[22]. However, we found that for identifying small bowel abnormalities in patients with no bleeding, the diagnostic yield of SBE was higher than that of CE (68.9% *vs* 43.8%).

SBE has the potential to become a useful technique for deep enteroscopy. SBE has a reasonable depth of insertion, can be administered using standard conscious sedation, and can be used with existing endoscopy systems. In addition, the SBE technique is easy to learn and can be rapidly incorporated into an endoscopy unit[23,24]. In our study, SBE generated a high diagnostic yield for small bowel diseases (overall diagnostic yield, 69.7%), for patients with OGIB and those without bleeding (72.5% *vs* 68.5%). A previous study recommended an initial CE examination, followed by DBE if necessary[7]. Here, we combined CE and SBE techniques to detect small bowel diseases and found that 45 patients had positive findings by CE examination, which was followed by SBE, and 42 patients had positive findings by SBE examination. The diagnostic yield of SBE with prior CE was 93.3% (42/45), which was a high diagnostic yield.

In summary, SBE appears to be a safe and effective method for diagnosing small bowel disease, especially for patients with OGIB. CE followed by SBE represents an especially effective strategy for determining the cause of small bowel disease when findings from initial CE examination are indeterminate.

**COMMENT**

***Background***

The diagnosis of small bowel diseases was difficult until the advent of capsule endoscopy (CE) and balloon assisted enteroscopy (BAE). Both CE and BAE were reported to have similarly high diagnostic yields of small bowel disorders. Single balloon enteroscopy (SBE), which is an alternative technique of Double balloon enteroscopy (DBE) for examining the deep small bowel, is simpler and easier-to-handle. There is limited data on the role of CE both in comparison and combination with SBE in the assessment of small bowel diseases.

***Research frontiers***

In this study, the authors aimed to evaluate diagnostic yields associated with CE, SBE, or their combined use in patients with suspected small bowel diseases and demonstrate the appropriate selection for different small bowel diseases.

***Innovations and breakthroughs***

This study was a single-center experience in China involving 700 patients who underwent CE and/or SBE. The difference of diagnostic yield of detecting small bowel diseases between CE and SBE was evaluated. The diagnostic yields of different indications and findings of CE and/or SBE were analyzed in detail. At the same time, the advantage of SBE combined with prior CE was also evaluated.

***Applications***

Both CE and SBE have high diagnostic yields of small bowel disorders. SBE has a similar diagnostic yield for patients with obscure gastrointestinal bleeding and a higher diagnostic yield with non-bleeding compared with CE. CE followed by SBE represents an especially effective strategy for determining small bowel disease.

***Terminology***

CE refers to a miniature capsule shaped camera that takes multiple pictures as its passes through the small intestine. SBE is a method of enteroscopy that can lead to the observation of the small intestine *via* the mouth or anus with the help of one balloon which is attached to the distal end of a soft overtube.

***Peer-review***

An interesting study, showing the role of small bowel evaluation with CE and SBE. It deserves to be published, it will add to the literature on the subject. Very good language and good presentation of the data.

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Grade D (Fair): D

Grade E (Poor): 0

**Table 1 Patient characteristics and indications for single-balloon enteroscopy and capsule endoscopy**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | **CE** | **SBE** | **Both**  **(CE prior to SBE)** | ***P* value**  **(CE *vs* SBE)** |
| No. of patients | 401 | 353 | 47 |  |
| Age (yr)  mean (range) | 49.4 ± 16.0  (13–85) | 42.1 ± 15.8  (11–84) | 45.3 ± 15.1  ( 15–77) | < 0.01 |
| Male/female | 248/153 | 235/118 | 38/9 | 0.177 |
| Main indications, *n* (%) |  |  |  |  |
| OGIB | 207 (51.6) | 80 (22.7) | 30 (63.8) |  |
| Abdominal pain | 133 (31.2) | 184 (52.1) | 11 (23.4) |  |
| Diarrhea | 30 (7.5) | 52 (14.7) | 1 (2.1) |  |
| Other | 31 (7.7) | 37 (10.5) | 5 (10.6) |  |

CE: Capsule endoscopy; OGIB: Obscure gastrointestinal bleeding; SBE: Single-balloon enteroscopy.

**Table 2 Comparison of findings between single-balloon enteroscopy and capsule endoscopy *n* (%)**

|  |  |  |
| --- | --- | --- |
| **Findings** | **CE (*n* = 401)** | **SBE (*n* = 353)** |
| Overall detection rate | 231 (57.6) | 246 (69.7) |
| Superficial ulcer and erosion | 98 (42.4) | 111 (45.1) |
| Angiodysplasia | 67(29.0) | 21 (8.5) |
| Mass | 24 (10.4) | 26 (10.6) |
| Crohn’s disease | 26 (11.3) | 86 (35.0) |
| Parasites | 13 (5.6) | 1 (0.4) |
| Diverticulum | 6 (2.6) | 3 (1.2) |
| Bleeding | 5 (2.2) | 0 |

CE: Capsule endoscopy; SBE: Single-balloon enteroscopy.

**Table 3** **Subgroup analysis of diagnostic yield of capsule endoscopy or single-balloon enteroscopy in patients with obscure gastrointestinal bleeding or non-bleeding patients**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Diagnostic yield** | | ***P* value** |
| **OGIB** | **Non-bleeding** |  |
| CE | 70.5 | 43.8% | < 0.01 |
| SBE | 72.5 | 68.9% | 0.534 |

CE: Capsule endoscopy; OGIB: Obscure gastrointestinal bleeding; SBE: Single-balloon enteroscopy.

**Table 4** **Identification of positive findings on prior capsule endoscopy or by single-balloon enteroscopy**

|  |  |  |  |
| --- | --- | --- | --- |
| **Findings** | **CE: negative diagnosis** | **CE: definite diagnosis** | **CE: definite diagnosis** |
| **SBE: definite diagnosis** | **SBE: definite diagnosis** | **SBE: negative diagnosis** |
| Angiodysplasia |  | 9 |  |
| Erosion and superficial ulcer | 2 | 18 |  |
| Mass |  | 3 | 1 |
| Crohn’s disease | 0 | 11 | 2 |
| Parasites |  | 1 |  |

CE: Capsule endoscopy; SBE: Single-balloon enteroscopy.