

## Prospective Study

## Application of the Prague C and M criteria for endoscopic description of columnar-lined esophagus in South Korea

Jung Wan Choe, Young Choon Kim, Moon Kyung Joo, Hyo Jung Kim, Beom Jae Lee, Ji Hoon Kim, Jong Eun Yeon, Jong-Jae Park, Jae Seon Kim, Kwan Soo Byun, Young-Tae Bak

Jung Wan Choe, Young Choon Kim, Moon Kyung Joo, Hyo Jung Kim, Beom Jae Lee, Ji Hoon Kim, Jong Eun Yeon, Jong-Jae Park, Jae Seon Kim, Kwan Soo Byun, Young-Tae Bak, Department of Gastroenterology, Korea University Guro Hospital, Seoul 08308, South Korea

**Author contributions:** Choe JW and Kim YC worked in data interpretation, and writing this manuscript; Bak YT worked in data acquisition, data analysis, data interpretation and in writing of this manuscript; all authors read and approved the final form of the manuscript.

**Institutional review board statement:** The study was reviewed and approved by the institutional review boards of Korea University Guro Hospital in South Korea.

**Clinical trial registration statement:** Although this research is a prospective study, there is no need to register clinical trial. This study only analyzed the results of endoscopic features, not evaluated or compare the clinical outcome. So, we are so sorry not to provide the trial's registry.

**Informed consent statement:** All study participants, or their legal guardian, provided written consent prior to study enrollment.

**Conflict-of-interest statement:** The authors of this manuscript having no conflicts of interest to disclose.

**Data sharing statement:** No additional data are available.

**Open-Access:** This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

**Correspondence to:** Young-Tae Bak, MD, Department of Gastroenterology, Korea University Guro Hospital, 148 Guro-

dong-ro, Guro-gu, Seoul 08308, South Korea. [drbakyt@korea.ac.kr](mailto:drbakyt@korea.ac.kr)  
Telephone: +82-2-26261778  
Fax: +82-504-3666381

Received: July 7, 2015

Peer-review started: July 8, 2015

First decision: September 8, 2015

Revised: September 30, 2015

Accepted: December 1, 2015

Article in press: December 2, 2015

Published online: April 25, 2016

### Abstract

**AIM:** To ascertain whether the Prague circumferential (C) length and maximal (M) length criteria for grading the extent of Barrett's esophagus can be applied prior to its widespread application in South Korea.

**METHODS:** Two hundred and thirteen consecutive cases with endoscopic columnar-lined esophagus (CLE) were included and classified according to the Prague C and M criteria.

**RESULTS:** Of 213 cases with CLE, the distribution of maximum CLE lengths was: 0.5-0.9 cm in 99 cases (46.5%); 1.0-1.4 cm in 63 cases (29.6%); 1.5-1.9 cm in 15 cases (7.0%); 2.0-2.4 cm in 14 cases (6.6%); 2.5-2.9 cm in 1 case (0.5%); and 7.0 cm in 1 case (0.5%). Twenty cases (9.4%) had columnar islands alone. Two hundred and eight cases (97.7%) lacked the circumferential CLE component (COMx). Columnar islands were found in 70 cases (32.9%), of which 20 cases (9.4%) had columnar islands alone.

**CONCLUSION:** In regions where most CLE patients display short or ultrashort tongue-like appearance, more detailed descriptions of CLE's in < 1.0 cm lengths and

columnar islands, as well as avoidance of repeating the prefix "C0" need to be considered in parallel with the widespread application of the Prague system in South Korea.

**Key words:** Barrett's esophagus; Endoscopy; Columnar-lined esophagus; Prague criteria

© **The Author(s) 2016.** Published by Baishideng Publishing Group Inc. All rights reserved.

**Core tip:** This was a prospective study to assess the feasibility of the Prague circumferential length and maximal length criteria for the endoscopic description of columnar-lined esophagus in South Korea. In regions like South Korea where the prevalence and endoscopic features of this condition are quite different from the West, we suggest possible modifications that may fit the characteristics of the South Korean source population more properly.

Choe JW, Kim YC, Joo MK, Kim HJ, Lee BJ, Kim JH, Yeon JE, Park JJ, Kim JS, Byun KS, Bak YT. Application of the Prague C and M criteria for endoscopic description of columnar-lined esophagus in South Korea. *World J Gastrointest Endosc* 2016; 8(8): 357-361 Available from: URL: <http://www.wjgnet.com/1948-5190/full/v8/i8/357.htm> DOI: <http://dx.doi.org/10.4253/wjge.v8.i8.357>

## INTRODUCTION

Barrett's esophagus (BE) is defined as a histological change of the distal tubular esophagus, from squamous to columnar epithelium, which displays an intestinal metaplasia containing goblet cells<sup>[1,2]</sup>. Because BE is characterized by an upward shift of the squamocolumnar junction (SCJ) proximal to the gastroesophageal junction (GEJ), the resulting columnar-lined mucosa of the distal esophagus can be identified by its salmon-pink color during endoscopic examination<sup>[3,4]</sup>. Moreover, multiple endoscopic biopsies at the extended columnar-lined epithelium are needed to confirm BE diagnosis.

BE is associated with gastroesophageal reflux disease (GERD) and is considered a premalignant lesion for esophageal adenocarcinoma<sup>[5,6]</sup>, the incidence of which is steadily rising in the United States and Europe<sup>[7,8]</sup>. Increasing GERD incidence in South Korea is considered to result from more consumption of westernized foods<sup>[9,10]</sup>. As patients with chronic GERD are at a higher risk of developing BE<sup>[11,12]</sup>, the expected increase in BE and esophageal cancer incidence rates in the future is a matter of potential concern in South Korea.

Various studies have examined BE length as a risk factor for esophageal adenocarcinoma<sup>[13-15]</sup>. Results from a study showed that a doubling in BE length resulted in a 1.7-fold increase in the risk of developing esophageal adenocarcinoma<sup>[15]</sup>, and others revealed that a significantly increased risk of dysplasia or adeno-

carcinoma was related to greater lengths of BE<sup>[13,14]</sup>. Therefore, accurate measuring of columnar-lined esophagus (CLE) lengths and describing in well-defined clinical terms are important in appropriate risk assessment and surveillance. Although previous diagnostic criteria for BE were based on the 3-cm length threshold of columnar-lined esophagus (CLE), by which BE was divided into 2 types, long ( $\geq 3$  cm) and short ( $< 3$  cm), this simple classification of variable endoscopic findings of CLE was a rather crude approach in describing BE. Furthermore, as considerable inter- and intra-observer variability in detecting and describing the CLE are common, the establishment of an accurate BE diagnosis and surveillance may be tricky<sup>[16-18]</sup>.

Therefore, the Prague classification system that measures the circumferential (C) and maximal (M) extents for endoscopic standardization of BE lengths was developed and finally introduced by the International Working Group for the Classification of Oesophagitis (IWGCO) in 2004<sup>[19]</sup>. However, the overall reliability and validity of the Prague C and M criteria for BE diagnosis continues to be challenged<sup>[20-22]</sup>. Moreover, its performance in South Korea where the incidence of BE is low and the short-segment BE is the predominant type remains unclear.

In the present study, we aimed to assess the feasibility of the Prague C and M criteria for the endoscopic description of CLE in South Korea where the prevalence and endoscopic features of this condition are quite different from the West and to suggest possible modifications that may fit the characteristics of the South Korean source population more properly.

## MATERIALS AND METHODS

This prospective study was conducted from the endoscopy data of consecutive CLE patients who underwent esophagogastroduodenoscopy (EGD) at Endoscopy Center of the Korea University Guro Hospital, Seoul, South Korea. Exclusion criteria included the presence of esophageal varices, acute upper gastrointestinal bleeding, malignancy near GEJ, and history of gastric surgery. Before each EGD, written informed consent was obtained. All endoscopic procedures were performed by an experienced endoscopist.

GEJ and SCJ were carefully assessed during the insertion of the endoscope. The distal margin of the palisade blood vessels of the lower esophagus was used as a marker of GEJ<sup>[23]</sup>. If the palisade vessels could not be seen adequately, the proximal margins of the gastric folds were used to identify GEJ. SCJ was used as a marker for upper border of CLE. The length of CLE, that is the distance from GEJ to SCJ, was measured by the insertion depths with the centimeter markings on the endoscope. CLE's shorter than 0.5 cm in length were ignored to avoid possible observation errors that may lead to overdiagnosis. Careful observation was done to look for any presence of islands of columnar mucosa.

The C and M extents of CLE were recorded accord-

**Table 1** Application of Prague circumferential and maximal criteria in cases with ultrashort, short, and long columnar-lined esophagus (*n* = 213)

Lengths of CLE (cm)	<i>n</i> (%)	COMx cases (%)
0 (islands only)	20 (9.4)	20 (100)
0.5-0.9	99 (46.5)	99 (100)
1.0-1.5	63 (29.6)	61 (96.8)
1.5-1.9	15 (7.0)	14 (93.3)
2.0-2.4	14 (6.6)	12 (85.7)
2.5-2.9	1 (0.5)	1 (100)
≥ 3.0	1 (0.5)	1 (100)
Total	213 (100)	208 (97.7) <sup>1</sup>

<sup>1</sup>Exceptions: 2 cases with C1M1 and 3 cases with either C1M1.5, C1M2, or C1.5M2. CLE: Columnar-lined esophagus.

**Table 2** Application of Prague circumferential and maximal criteria in cases with short and long columnar-lined esophagus (*n* = 139)

Lengths of CLE (cm)	<i>n</i> (%)	COMx cases (%)
0 (islands only)	45 (32.4)	45 (100)
1.0-1.4	63 (45.3)	61 (96.8)
1.5-1.9	15 (10.8)	14 (93.3)
2.0-2.4	14 (10.1)	12 (85.7)
2.5-2.9	1 (0.7)	1 (100)
≥ 3.0	1 (0.7)	1 (100)
Total	139 (100)	134 (96.4) <sup>1</sup>

<sup>1</sup>Exceptions: 2 patients with C1M1 and 3 patients with either C1M1.5, C1M2, or C1.5M2. CLE: Columnar-lined esophagus.

ing to the Prague C and M criteria proposed by the IWGCO<sup>[19]</sup>. M lengths were divided into long (≥ 3 cm), short (1-2.9 cm), and ultrashort (< 1 cm) segments.

## RESULTS

### Patient demographic characteristics

A total of 213 CLE patients consisting of 154 men and 59 women, with 53.8 ± 12.3 years in age (mean ± SD) were enrolled.

### Distribution of CLE lengths and application of the Prague C and M criteria

#### Analysis of cases with CLE's including ultrashort CLE's:

Distribution of CLE's according to their M values, including those with ultrashort CLE's, is shown in Table 1. Among the total 213 cases, 99 (46.5%), 63 (29.6%), 15 (7.0%), 14 (6.6%), 1 (0.5%), and 1 (0.5%) had CLE's of 0.5-0.9 cm, 1.0-1.4 cm, 1.5-1.9 cm, 2.0-2.4 cm, 2.5-2.9 cm, and ≥ 3.0 cm in lengths, respectively. The remaining 20 cases (9.4%) had columnar islands alone. Therefore, 99 cases (46.5%) had ultrashort CLE's (CLE < 1.0 cm), 113 (53.1%) had short CLE's (1-2 cm) and only one (0.5%) had a long CLE (≥ 3 cm), showing a CLE of 7.0 cm in length.

When the cases were classified by the Prague criteria, 208 (97.7%) had no C component (COMx). Two cases had C1M1 and the remaining three cases had

either, C1M1.5, C1M1, or C1.5M2. Columnar islands were observed in 70 (32.9%) cases, of which 20 (9.4%) had columnar islands alone.

#### Analysis of cases with CLE's excluding ultrashort CLE's:

Distribution of CLE's according to their M values among those excluding ultrashort CLE's is shown in Table 2. Among 139 cases, 63 (45.3%), 15 (10.8%), 14 (10.1%), 1 (0.7%), and 1 (0.7%) had CLE's of 1.0-1.4 cm, 1.5-1.9 cm, 2.0-2.4 cm, 2.5-2.9 cm and ≥ 3.0 cm in lengths, respectively. Therefore, 138 (99.3%) out of all 139 cases had short CLE's, and only one showed an exceptionally long CLE.

When 139 cases were classified by the Prague criteria, 134 (96.4%) had CLE's without C component (COMx). Two cases had C1M1 and the remaining three patients had either C1M1.5, C1M1, or C1.5M2. Columnar islands were found in 70 (50.4%) cases, of which 45 (32.4%) showing columnar islands alone.

## DISCUSSION

BE is a very well known risk factor for the development of dysplasia and esophageal adenocarcinoma<sup>[24-26]</sup>. The risk of dysplasia and adenocarcinoma in metaplastic epithelium reportedly increases in parallel to the lengths of BE<sup>[13-15]</sup>. A recent multicenter study conducted by Gaddam *et al.*<sup>[13]</sup> revealed that for every 1-cm extension in BE length, the risk of high-grade dysplasia and esophageal adenocarcinoma increased by 21%. The study demonstrated that the increase in BE lengths significantly widens the area of metaplasia, which is associated with the progression to high-grade dysplasia/esophageal adenocarcinoma<sup>[13]</sup>. Although a novel technique using a computer software program to create a two-dimensional image map of the esophagus has been introduced to accurately and reproducibly measure the extent of CLE<sup>[27]</sup>, such a complicated approach is not suitable for a daily clinical practice. Therefore, assessment of BE extent by simple measurement of the height of metaplastic CLE remains as the most commonly used procedure to distinguish short- from long-segment BE<sup>[13-15]</sup>. However, the study of the clinical course and therapeutic response of BE has been limited because this classic method only provides gross estimates of the area. This system does not measure the surface areas of metaplastic mucosa, which may be more important than the endoscopic lengths<sup>[19]</sup>. The presence of an irregular border of columnar tissue or interspersed metaplastic mucosal islands can hamper the precise measurement of the extent of CLE<sup>[20]</sup>.

The Prague C and M criteria, suggested by IWGCO, not only allows a more detailed description of the length of the endoscopically recognized CLE, using "C" and "M" values above the GEJ, but can also assist the objective calculation of the actual surface area, which may be more important in the risk assessment of the neoplastic transformation<sup>[19-21]</sup>. These advances in CLE description have facilitated the depiction and reporting of various

circumferential and tongue-like longitudinal CLE lengths by using a method that can be understood easily and comprehensively. Importantly, high inter-observer reliability in the grading of endoscopically suspected CLE was demonstrated among gastroenterology experts and trainees<sup>[22]</sup>.

In recent years, accelerated life style changes have increased the prevalence of GERD in Asian populations, including South Koreans<sup>[9,10,28,29]</sup>, and BE incidence is also expected to increase<sup>[12]</sup>. BE prevalence in South Korea was 0.2%-3.6% in the year 2000<sup>[11,12,30]</sup>, lower than in Western countries. Lengths and shapes of CLE's as well as their prevalence in South Korea are quite different from those of the Western countries. Long-segment BE is more common in Western countries, wherein 14%-31% of BE patients show this type<sup>[31,32]</sup>. However, most cases of BE are short-segment type in South Korea, where long-segment type BE's are extremely rare<sup>[11]</sup>. In our study, with the exception of the only one case, 212 (99.5%) out of 213 CLE cases were short-segment type (< 3 cm); and from these, 99 cases (46.5%) had ultrashort CLE (< 1 cm). Lee *et al.*<sup>[33]</sup> reported that the reliability coefficients of the C and M values in the endoscopic recognition of short-segment type CLE were 0.90 (95%CI: 0.80%-1.00%) and 0.92 (95%CI: 0.87%-0.98%), respectively. However, the reliability of such coefficients for the recognition of the ultrashort (< 1 cm) CLE extent type was very low, with C and M coefficients of 0.18 (95%CI: 0.03%-0.32%) and 0.21 (95%CI: 0.00%-0.51%), respectively<sup>[33]</sup>.

Therefore, the routine applicability of the Prague C and M criteria as a standardized validated method for the detailed endoscopic description of ultrashort BE and short-segment BE, the most dominant BE types in South Korea, requires further analysis. As our study showed, all ultrashort CLE and almost all short-segment CLE cases lacked the C component and were classified as C0Mx. Therefore, it appears appropriate for us to propose to omit of the prefix "C0" from all C0Mx cases in order to avoid needless repetitions when describing most cases in regions like South Korea. Because the presence of columnar islands is a frequent finding as we have observed in this study and they also may change to dysplasia<sup>[34]</sup>, we propose to add this to the Prague system, which currently does not include this category. Resultant examples following our proposals are: C2M5, if 2.0 cm of C component with 5.0 cm of M component; M2, if 2.0 cm of M component without C component; C2M5i or M2i, if columnar island(s) is/are found in addition to C2M5 or M2 CLE; and M0i, if only columnar island(s) is/are found.

In summary, the Prague C and M system is simple and useful in daily description of endoscopic feature of CLE's. However, in regions like South Korea where most cases with CLE display only short or ultrashort types without C component, we propose to omit the needless repetition of "C0" prefix from C0Mx and to add i component to describe the presence of columnar islands which also may have a potential to be dysplastic.

## COMMENTS

### Background

The Prague circumferential (C) length and maximal (M) length criteria have been adopted widely for grading the extent of Barrett's esophagus (BE). However, its validity in regions with low prevalence of BE, remains unclear. This study was designed to ascertain whether these criteria can be applied prior to its widespread application in South Korea.

### Research frontiers

The Prague C and M system is simple and useful in daily description of endoscopic feature of BE's. But, the overall reliability and validity of the Prague C and M criteria for BE diagnosis continues to be challenged. In this study, there are some suggestions of possible modifications that may fit the characteristics of the South Korean source population more properly.

### Innovations and breakthroughs

In regions like South Korea where most cases with columnar-lined esophagus display only short or ultrashort types without C component, the authors propose to omit the needless repetition of "C0" prefix from C0Mx and to add "i" component to describe the presence of columnar islands which also may have a potential to be dysplastic.

### Applications

This study serves as additional evidence supporting the investigation in parallel with the widespread application of the Prague system in South Korea.

### Terminology

Barrett's esophagus: A histological change of the distal tubular esophagus, from squamous to columnar epithelium, which displays an intestinal metaplasia containing goblet cells; The Prague classification criteria: A system to measure the C and M extents for endoscopic standardization of BE lengths.

### Peer-review

The study is has clear defined inclusion and exclusion criteria and is well conducted despite the lack of a control group. This study is innovative and would be interesting to see if the findings are reproducible in other countries where BE is not as common as in the West.

## REFERENCES

- 1 **Spechler SJ**, Goyal RK. Barrett's esophagus. *N Engl J Med* 1986; **315**: 362-371 [PMID: 2874485 DOI: 10.1056/NEJM198608073150605]
- 2 **Wang KK**, Sampliner RE. Updated guidelines 2008 for the diagnosis, surveillance and therapy of Barrett's esophagus. *Am J Gastroenterol* 2008; **103**: 788-797 [PMID: 18341497 DOI: 10.1111/j.1572-0241.2008.01835.x]
- 3 **Barrett NR**. Chronic peptic ulcer of the oesophagus and 'oesophagitis'. *Br J Surg* 1950; **38**: 175-182 [PMID: 14791960]
- 4 **Sharma P**, McQuaid K, Dent J, Fennerty MB, Sampliner R, Spechler S, Cameron A, Corley D, Falk G, Goldblum J, Hunter J, Jankowski J, Lundell L, Reid B, Shaheen NJ, Sonnenberg A, Wang K, Weinstein W. A critical review of the diagnosis and management of Barrett's esophagus: the AGA Chicago Workshop. *Gastroenterology* 2004; **127**: 310-330 [PMID: 15236196]
- 5 **Mann NS**, Tsai MF, Nair PK. Barrett's esophagus in patients with symptomatic reflux esophagitis. *Am J Gastroenterol* 1989; **84**: 1494-1496 [PMID: 2596449]
- 6 **Winters C**, Spurling TJ, Chobanian SJ, Curtis DJ, Esposito RL, Hacker JF, Johnson DA, Cruess DF, Cotelingam JD, Gurney MS. Barrett's esophagus. A prevalent, occult complication of gastroesophageal reflux disease. *Gastroenterology* 1987; **92**: 118-124 [PMID: 3781178]
- 7 **Botterweck AA**, Schouten LJ, Volovics A, Dorant E, van Den Brandt PA. Trends in incidence of adenocarcinoma of the oesophagus and gastric cardia in ten European countries. *Int J*



- Epidemiol* 2000; **29**: 645-654 [PMID: 10922340]
- 8 **Powell J**, McConkey CC, Gillison EW, Spychal RT. Continuing rising trend in oesophageal adenocarcinoma. *Int J Cancer* 2002; **102**: 422-427 [PMID: 12402314 DOI: 10.1002/ijc.10721]
  - 9 **Lee SJ**, Song CW, Jeon YT, Chun HJ, Lee HS, Um SH, Lee SW, Choi JH, Kim CD, Ryu HS, Hyun JH. Prevalence of endoscopic reflux esophagitis among Koreans. *J Gastroenterol Hepatol* 2001; **16**: 373-376 [PMID: 11354273]
  - 10 **Yoo SS**, Lee WH, Ha J, Choi SP, Kim HJ, Kim TH, Lee OJ. [The prevalence of esophageal disorders in the subjects examined for health screening]. *Korean J Gastroenterol* 2007; **50**: 306-312 [PMID: 18159162]
  - 11 **Kim JY**, Kim YS, Jung MK, Park JJ, Kang DH, Kim JS, Song CW, Lee SW, Bak YT. Prevalence of Barrett's esophagus in Korea. *J Gastroenterol Hepatol* 2005; **20**: 633-636 [PMID: 15836715 DOI: 10.1111/j.1440-1746.2005.03749.x]
  - 12 **Park JJ**, Kim JW, Kim HJ, Chung MG, Park SM, Baik GH, Nah BK, Nam SY, Seo KS, Ko BS, Jang JY, Kim BG, Kim JW, Choi YS, Joo MK, Kim JI, Cho MY, Kim N, Park SH, Jung HC, Chung IS. The prevalence of and risk factors for Barrett's esophagus in a Korean population: A nationwide multicenter prospective study. *J Clin Gastroenterol* 2009; **43**: 907-914 [PMID: 19417682 DOI: 10.1097/MCG.0b013e318196bd11]
  - 13 **Gaddam S**, Young PE, Alsop BR, Gupta N, Gavini H, Higbee AD, Wani SB, Singh M, Rastogi A, Bansal A, Cash BD, Lieberman DA, Sampliner RE, Falk GW, Sharma P. Relationship Between Barrett's Esophagus (BE) Length and the Risk of High Grade Dysplasia (HGD) and Esophageal Adenocarcinoma (EAC) in Patients With Non Dysplastic Barrett's Esophagus Results From a Large Multicenter Cohort. *Gastroenterology* 2011; **140**: S81-S81 [DOI: 10.1016/S0016-5085(11)60329-6]
  - 14 **Iftikhar SY**, James PD, Steele RJ, Hardcastle JD, Atkinson M. Length of Barrett's oesophagus: an important factor in the development of dysplasia and adenocarcinoma. *Gut* 1992; **33**: 1155-1158 [PMID: 1427364]
  - 15 **Menke-Pluymers MB**, Hop WC, Dees J, van Blankenstein M, Tilanus HW. Risk factors for the development of an adenocarcinoma in columnar-lined (Barrett) esophagus. The Rotterdam Esophageal Tumor Study Group. *Cancer* 1993; **72**: 1155-1158 [PMID: 8339208]
  - 16 **Sharma P**, Morales TG, Sampliner RE. Short segment Barrett's esophagus--the need for standardization of the definition and of endoscopic criteria. *Am J Gastroenterol* 1998; **93**: 1033-1036 [PMID: 9672325 DOI: 10.1111/j.1572-0241.1998.00324.x]
  - 17 **Dekel R**, Wakelin DE, Wendel C, Green C, Sampliner RE, Garewal HS, Martinez P, Fass R. Progression or regression of Barrett's esophagus--is it all in the eye of the beholder? *Am J Gastroenterol* 2003; **98**: 2612-2615 [PMID: 14687805 DOI: 10.1111/j.1572-0241.2003.07680.x]
  - 18 **Kim SL**, Waring JP, Spechler SJ, Sampliner RE, Doos WG, Krol WF, Williford WO. Diagnostic inconsistencies in Barrett's esophagus. Department of Veterans Affairs Gastroesophageal Reflux Study Group. *Gastroenterology* 1994; **107**: 945-949 [PMID: 7926484]
  - 19 **Sharma P**, Dent J, Armstrong D, Bergman JJ, Gossner L, Hoshihara Y, Jankowski JA, Junghard O, Lundell L, Tytgat GN, Vieth M. The development and validation of an endoscopic grading system for Barrett's esophagus: the Prague C & M criteria. *Gastroenterology* 2006; **131**: 1392-1399 [PMID: 17101315 DOI: 10.1053/j.gastro.2006.08.032]
  - 20 **Anand O**, Wani S, Sharma P. When and how to grade Barrett's columnar metaplasia: the Prague system. *Best Pract Res Clin Gastroenterol* 2008; **22**: 661-669 [PMID: 18656823]
  - 21 **Chang CY**, Lee YC, Lee CT, Tu CH, Hwang JC, Chiang H, Tai CM, Chiang TH, Wu MS, Lin JT. The application of Prague C and M criteria in the diagnosis of Barrett's esophagus in an ethnic Chinese population. *Am J Gastroenterol* 2009; **104**: 13-20 [PMID: 19098843 DOI: 10.1038/ajg.2008.43]
  - 22 **Vahabzadeh B**, Seetharam AB, Cook MB, Wani S, Rastogi A, Bansal A, Early DS, Sharma P. Validation of the Prague C & M criteria for the endoscopic grading of Barrett's esophagus by gastroenterology trainees: a multicenter study. *Gastrointest Endosc* 2012; **75**: 236-241 [PMID: 22248595 DOI: 10.1016/j.gie.2011.09.017]
  - 23 **Choi DW**, Oh SN, Baek SJ, Ahn SH, Chang YJ, Jeong WS, Kim HJ, Yeon JE, Park JJ, Kim JS, Byun KS, Bak YT, Lee CH. Endoscopically observed lower esophageal capillary patterns. *Korean J Intern Med* 2002; **17**: 245-248 [PMID: 12647639]
  - 24 **Cameron AJ**, Ott BJ, Payne WS. The incidence of adenocarcinoma in columnar-lined (Barrett's) esophagus. *N Engl J Med* 1985; **313**: 857-859 [PMID: 4033716 DOI: 10.1056/NEJM198510033131404]
  - 25 **Hameeteman W**, Tytgat GN, Houthoff HJ, van den Tweel JG. Barrett's esophagus: development of dysplasia and adenocarcinoma. *Gastroenterology* 1989; **96**: 1249-1256 [PMID: 2703113]
  - 26 **Van der Veen AH**, Dees J, Blankenstein JD, Van Blankenstein M. Adenocarcinoma in Barrett's oesophagus: an overrated risk. *Gut* 1989; **30**: 14-18 [PMID: 2920919]
  - 27 **Kim R**, Baggott BB, Rose S, Shar AO, Mallory DL, Lasky SS, Kressloff M, Faccenda LY, Reynolds JC. Quantitative endoscopy: precise computerized measurement of metaplastic epithelial surface area in Barrett's esophagus. *Gastroenterology* 1995; **108**: 360-366 [PMID: 7835577]
  - 28 **Rosaia MS**, Goh KL. Gastro-oesophageal reflux disease, reflux oesophagitis and non-erosive reflux disease in a multiracial Asian population: a prospective, endoscopy based study. *Eur J Gastroenterol Hepatol* 2004; **16**: 495-501 [PMID: 15097043]
  - 29 **Wong WM**, Lam SK, Hui WM, Lai KC, Chan CK, Hu WH, Xia HH, Hui CK, Yuen MF, Chan AO, Wong BC. Long-term prospective follow-up of endoscopic oesophagitis in southern Chinese--prevalence and spectrum of the disease. *Aliment Pharmacol Ther* 2002; **16**: 2037-2042 [PMID: 12452935 DOI: 10.1046/j.1365-2036.2002.01373.x]
  - 30 **Kim JH**, Rhee PL, Lee JH, Lee H, Choi YS, Son HJ, Kim JJ, Rhee JC. Prevalence and risk factors of Barrett's esophagus in Korea. *J Gastroenterol Hepatol* 2007; **22**: 908-912 [PMID: 17565647 DOI: 10.1111/j.1440-1746.2006.04448.x]
  - 31 **Ronkainen J**, Aro P, Storskrubb T, Johansson SE, Lind T, Bolling-Sternevald E, Vieth M, Stolte M, Talley NJ, Agr us L. Prevalence of Barrett's esophagus in the general population: an endoscopic study. *Gastroenterology* 2005; **129**: 1825-1831 [PMID: 16344051 DOI: 10.1053/j.gastro.2005.08.053]
  - 32 **Csendes A**, Smok G, Burdiles P, Korn O, Gradiz M, Rojas J, Recio M. Prevalence of intestinal metaplasia according to the length of the specialized columnar epithelium lining the distal esophagus in patients with gastroesophageal reflux. *Dis Esophagus* 2003; **16**: 24-28 [PMID: 12581250]
  - 33 **Lee YC**, Cook MB, Bhatia S, Chow WH, El-Omar EM, Goto H, Lin JT, Li YQ, Rhee PL, Sharma P, Sung JJ, Wong JY, Wu JC, Ho KY. Interobserver reliability in the endoscopic diagnosis and grading of Barrett's esophagus: an Asian multinational study. *Endoscopy* 2010; **42**: 699-704 [PMID: 20806154 DOI: 10.1055/s-0030-1255629]
  - 34 **Dunbar KB**, Okolo P, Montgomery E, Canto MI. Confocal laser endomicroscopy in Barrett's esophagus and endoscopically inapparent Barrett's neoplasia: a prospective, randomized, double-blind, controlled, crossover trial. *Gastrointest Endosc* 2009; **70**: 645-654 [PMID: 19559419 DOI: 10.1016/j.gie.2009.02.009]

P- Reviewer: Meshikhes AWN, Slomiany BL S- Editor: Ji FF

L- Editor: A E- Editor: Liu SQ





Published by **Baishideng Publishing Group Inc**

8226 Regency Drive, Pleasanton, CA 94588, USA

Telephone: +1-925-223-8242

Fax: +1-925-223-8243

E-mail: [bpgoffice@wjgnet.com](mailto:bpgoffice@wjgnet.com)

Help Desk: <http://www.wjgnet.com/esps/helpdesk.aspx>

<http://www.wjgnet.com>

