Colonoscopy quality assessment


Department of Digestive Diseases. University Hospital Santa María del Rosell. Cartagena, Murcia.
1 University Hospital Morales Meseguer. Murcia. 2 University Hospital Virgen de la Arrixaca. Murcia.
3 University Hospital Reina Sofía. Murcia. 4 Dirección General de Calidad Asistencial. Consejería de Sanidad de la Región de Murcia. 5 Dirección General de Salud Pública. Consejería de Sanidad de la Región de Murcia. Spain

ABSTRACT

Aim: colonoscopy has become accepted as the most effective method for colon exploration. Some application problems have been detected in the setting of normal clinical care due to its wide range of uses in recent years, and therefore there is a need to measure colonoscopy quality. For that purpose valid quality indicators are necessary to be defined. The application process of some quality indicators is presented in this study. The proposed indicators in this study are: quality of bowel preparation, cecal intubation rate, withdrawal time, adenoma detection rate, and adenoma removal rate.

Material and method: this is a prospective 12-month study where colonoscopies performed in the VI health area of Murcia Region were evaluated. From February 2006 to February 2007 a total of 609 subjects were eligible for colonoscopy after a positive fecal blood test in the setting of a colorectal cancer screening program. A sample of thirty patients (n: 30) was considered representative to assess the reliability of quality indicators and for a preliminary analysis of results.

Results: indicators results are: quality of bowel preparation (87%), kappa 0.74 (95% CI: 0.48-0.99); cecal intubation rate (90%), kappa 0.74 (95% CI: 0.49-0.99); adenoma detection and removal rate (96%), kappa 0.78 (95% CI: 0.53-0.99); withdrawal time: 13.36 min (95% CI: 10.48-16.11). Kappa: 0.78 (95% CI: 0.49-0.99).

Conclusions: quality indicators definition and application in colonoscopy performance is possible. More studies are necessary to define the role of these indicators in the setting of clinical practice.

Key words: Quality. Quality indicators. Colonoscopy. Endoscopy.

INTRODUCTION

Colonoscopy is presently the gold standard method for colonic examination. Not only is it widely accepted as a diagnostic technique but also as a procedure which enables treatment for a wide variety of conditions involving the intestinal tract. This unique feature explains why its practice is done in most hospitals of present-day healthcare systems.

Quality measurements for any process require firstly a definition of valid and reliable indicators to enable assessment (1). In recent years many attempts have been tried in order to define useful criteria for this purpose; however, its use has not been completely accepted to date by the different medical societies (2-5).

There is strong evidence that colonoscopy performance varies among different centers and endoscopists (6-12). The issue we presently face is the unavailability of adequate tools to measure these parameters.

The use of colonoscopy has proven the most effective method to reduce colorectal cancer (CRC) mortality (13-15). A wide variety of screening modalities have been developed to date, and a majority proved effective to reduce colorectal cancer incidence and mortality rates, with colonoscopy being the last resort in the screening sequence regardless of the initial screening method used (16-19).

In this study we present the application and adaptation of several parameters to a specific context in order to evaluate the quality of a screening program. This is a
complicated process which requires a staff used to specific quality evaluation methods (20,21).

The following recommendations should be followed when criteria or valid indicators are elaborated as quality measurement tools (Table I).

In this study we present four indicators elaborated for the assessment of colonoscopy quality in the setting of a screening program.

**Table I. Recommendations for criteria setup**

<table>
<thead>
<tr>
<th>Recommendation</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Use simple criteria</td>
<td>Limit the maximum number to 10</td>
</tr>
<tr>
<td>Include only essential parameters</td>
<td>Adapt to sources of healthcare professionals and their setting</td>
</tr>
<tr>
<td>Ensure consensus and acceptability by the involved staff</td>
<td>Make sure the content is updated and valid</td>
</tr>
<tr>
<td>Pay special attention to exceptions and explanations of used terminology</td>
<td>Ensure reliability</td>
</tr>
</tbody>
</table>

(Adapted from: Saturno PJ. Evaluación y mejora de la calidad en Servicios de Salud, 2000).

**MATERIAL AND METHOD**

This study was performed in the setting of a CRC screening program. The aim of this study was to measure a number of quality parameters for this diagnosis confirmation stage by using some indicators (22).

The study population consisted of subjects participating in the screening program with a quantitatively positive immunochemical occult blood test (≥ 100 ng/ml) who accepted to undergo colonoscopy (609 participants) between February 2006 and February 2007. The unit of the study consisted of each participant enrolled in the program.

A number of indicators were settled upon based on the best scientific evidence available to date for quality assessment and measurement (2,3,5). The aforementioned indicators measure the magnitude of the scientific-technical quality of colonoscopy.

All participants were given detailed information about the screening process, which is registered in the clinical records of each participant. All colonoscopies performed were completely recorded and stored in a digital system. Each participant was assigned an individual number, which was used for case identification. A written informed consent was obtained from each participant to be included in an anonymous way.

The screening program was performed by two endoscopists who were specially trained to use these indicators. A pilot program was carried out in order to confirm the reliability of the proposed indicators. For this purpose a randomized sample of 30 patients was obtained to calculate a specific consistency index for each indicator. The analysis of such indicators was carried out by three endoscopists who did not participate in the screening program.

The data source for the indicators evaluating colonoscopy was the recorded examination, subsequently visualized to draw conclusions.

Indicators analyzed in this study were as follows:
- Bowel preparation quality.
- Cecal intubation rate.
- Polyp detection and resection rate.
- Colonoscope withdrawal time.

**Definition of criteria**

**Adequate bowel preparation for colonoscopy**

It is an indicator of the process and evaluates the quality of bowel cleanliness prior to colonoscopy. The colonoscope explores the tract from the anus to the cecum and the small bowel; for an optimal evaluation of the colonic mucosa, the patient must be prepared for the procedure by removing feces from the digestive tract (23,24).

To evaluate the quality of bowel preparation during colonoscopy the following classification was used (Table II):

<table>
<thead>
<tr>
<th>Grade</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Very good</td>
<td>Solid feces are not observed, or liquid easily to be suctioned (transparent or like urine)</td>
</tr>
<tr>
<td>2. Good</td>
<td>Semi-liquid feces are observed, not adhered to the colonic mucosa, easily suctioned; do not block visualization of the whole colonic mucosa</td>
</tr>
<tr>
<td>3. Fair</td>
<td>Solid feces are observed, adhered to the colonic mucosa, not allowing an adequate vision of the whole mucosa. The examination can be completed but it is not satisfactory to obtain information</td>
</tr>
<tr>
<td>4. Poor</td>
<td>Solid feces are observed impossible to suction, not allowing an adequate progression of the endoscope and block visualization of the whole colonic mucosa</td>
</tr>
</tbody>
</table>

**Complete examination of the colon (cecal intubation)**

An appropriate examination of the proximal colon or cecum with colonoscopy is essential. In daily practice a
complete colonoscopy implies an intubation of the entire colon until the cecum is visible. Only the visualization of the ileocecal valve and/or intubation of the terminal ileum provides reassurance on the procedure’s completeness (6,25-27). Thus, it is necessary to document the procedure in a digital source once the cecum is reached in order to confirm the fulfillment of this criterion. In procedures where the endoscope found a stenosis, this indicator was not evaluated regardless of stenosis nature.

**Resected polyps and flat lesions**

Every lesion detected during colonoscopy must be resected. Colonoscopy in a screening program is performed under the most suitable conditions regarding tolerance as well as preparation. Every detected lesion which can be endoscopically resected is removed. There is a strong relationship between polyp size and histological type so that any lesion with a size larger than 10 mm is considered of high risk (28-31). In recent years the neoplastic potential of lesions smaller in size has been analyzed, specially regarding flat lesions, making the resection of every visible polyp a necessary condition regardless of size (10,11). To achieve fulfillment for this indicator all visualized lesions should be endoscopically resected except certain lesions. Those lesions which endoscopically suggest a hyperplastic polyp (pale, sessile lesions that disappear after insufflation) and measure less than 3 mm in size can be biopsied with no necessity to resect all lesions to meet indicator criteria.

In cases with more than 20 polyps detected the removal of all polyps can be performed in a second examination without invalidating the established criteria. In such patients genetic susceptibility to colorectal adenomatous lesions may be suspected.

In situations where the lesion cannot be removed at initial colonoscopy due to its large size, resection can be postponed. The minimal size to evaluate whether endoscopic resection is necessary in a second colonoscopy is 3 cm for polypoid lesions and 2 cm for flat lesions.

**Measuring examination time during colonoscopy withdrawal**

Examination time during colonoscope withdrawal is measured from the cecum to the anus. Endoscopic examination consists of the insertion of an optical system in the colon with the purpose of identifying treatable lesions (32).

Scientific evidence demonstrates that withdrawal is most suitable for the detection of lesions (33,34). Examinations where complete colonoscopy is achieved by a clear visualization of the cecum, as previously mentioned, will be evaluated.

It has been recently demonstrated that a withdrawal time superior to 6 minutes is associated with good endoscopic practice since it enables higher detection rates for colonic lesions (34,35).

If, during the examination, a specific technique different from lesion identification is performed, this time will not be considered to evaluate the indicator, and therefore will be taken from total examination time.

**Statistical analysis**

The statistical analysis was performed with the Epidat system (v 3.1 January 2006). Sample tests, inference over statistical parameters, and definition of confidence intervals were all performed with this program.

**RESULTS**

**Bowel preparation quality**

A total of 30 examinations were evaluated and bowel cleanliness was classified as follows: 3 (10%) “very good”, 23 (77%) “good”, 4 (13%) “fair”, and 0 “poor”. An analysis of indicator reliability was performed, and the global kappa index (K) for all three observers was: 0.47 (95% CI: 0.24-0.70). These consistent results were related to the assessment of bowel preparation using the proposed classification. Since the kappa value was in the limit of desirable, corrective modifications were performed. When a concordance analysis was done by gathering indicators as follows: “very good” or “good” as adequate examination, and “fair” or “poor” as inadequate examination, the value of kappa was modified to 0.74 (95% CI: 0.48-0.99) (Fig. 1).

**Fig. 1. Rate of patients according to grade of bowel cleanliness.**

Porcentaje de pacientes según nivel de limpieza.
Cecal intubation

Among the evaluated sample of 30 patients, the cecum was identified in 27 patients (90%) according to the previously mentioned criteria. Of the 3 patients where the cecum could not be reached, intolerance in spite of medication was the reason in two (6%); in one, cecum was not achieved due to an anatomical feature (dolichocolon). The kappa index obtained for indicator evaluation by all three observers was 0.74 (95% CI: 0.49-0.99).

Polyp resection

Of the 30 patients evaluated, this indicator was met in 25 patients (96%). Among the remaining 5, 4 cases were considered exceptions; 3 had hyperplastic polyposis, and one case required a second colonoscopy for polypectomy of a large-size polyp (greater than 3 cm). Only one case did not meet indicator criteria. A kappa index of 0.78 (95% CI: 0.53-0.99) was obtained in the consistency analysis (Fig. 2).

Withdrawal time

The following results were obtained: nearly 97% of patients met the established criteria. Mean withdrawal time was 13.36 min (95% CI: 10.48-16.11). A reliability analysis of minimal withdrawal time measurement was performed, which obtained a kappa index of 0.78 (95% CI: 0.49-0.99) (Fig. 3).

DISCUSSION

Quality is an essential paradigm in daily clinical activity. In recent years remarkable efforts have been made to devise tools able to establish optimal quality standards in the different processes of medical practice (2,5,18).

Gastrointestinal endoscopy has become a diagnostic-therapeutic tool essential for daily clinical practice, presently its practice being done in most hospitals of present-day healthcare systems.

This globalization has determined the present technological development, which enables us to assure there is no site in the gastrointestinal tract impossible to be explored. However, this same globalization has contributed to a certain level of heterogeneity, still to be defined, which does not allow comparisons between the different settings where it is performed (36). For this purpose it is necessary to establish resources capable of objectively assessing an activity regardless of the healthcare setting considered.

There is growing interest in the elaboration of such models, and different scientific societies have introduced their own models, adapting them to their own practical settings (2,5,18). The observed tendency with these models is a progression to simplicity, since the more complex they are, the more problematic their application becomes, with a consequent loss in efficacy.

When a model is introduced the possibility of its modification must be clearly established since one of the main features should be continuous improvement.

One of the main problems emerging when a quality model is put to the test for a certain medical practice is framework; routine medical practice frequently has difficulties when all parts making up a process need to be controlled; consequently, a low rate of implementation...
for such models is mostly observed. Many attempts to consolidate a proposed quality model have been made in the setting of a CRC screening program (2,5). A screening program enables the organization of such activity by settling a healthcare process; the management of each process in the entire healthcare framework generated by the screening strategy allows controlling the whole activity performed in a more adequate fashion. Therefore, carrying out the study in the screening program enabled us to control colonoscopy practice, performed in this framework in a more simple way than in classic medical practice, and consequently allowing that the proposed model be put to the test. The proposed process for this study has a clearly established model defined by a number of indicators whose application will enable the settling of a certain quality level and its support by monitoring techniques. Measuring bowel cleanliness is somewhat difficult when levels have to be established. The possibility to establish levels according to the amount of clean mucosa measured in percentage seems unfeasible from a practical point of view due to the impossibility to define the different paths of the colon in absolute terms. One of the most relevant conclusions to reach from the assessment of this indicator was the necessity to group levels together in order to simplify interpretations on cleansing quality. Thus, when colonoscopy cleansing quality has to be evaluated a scale of two levels is most reliable: valid or not valid. The results presented in this study reveal that more than 90% of examinations were considered valid for bowel cleansing.

In published series of observational studies, a rate of fulfillment for this indicator above 70% represented an adequate level of cleanliness (2,3,5). Cecal intubation implies the end of the examination, and therefore a complete colon examination is assumed. Population-based series of cecal intubation not related to screening programs report rates of 90% as their highest levels (2,3,7). Thus, cecum intubation rates up to 97% have been reported in the setting of a screening program (2,3,6,25). The results observed in this study reveal that more than 90% of examinations were considered valid for bowel cleansing.

In published series of observational studies, a rate of fulfillment for this indicator above 70% represented an adequate level of cleanliness (2,3,5). Cecal intubation implies the end of the examination, and therefore a complete colon examination is assumed. Population-based series of cecal intubation not related to screening programs report rates of 90% as their highest levels (2,3,7). Thus, cecum intubation rates up to 97% have been reported in the setting of a screening program (2,3,6,25). The results observed in this study reveal that more than 90% of examinations were considered valid for bowel cleansing.

In the indicators assessed in our study, it is important to mention that to a certain extent they are indirect markers that do not directly establish whether colonoscopy was “high quality”, but determine the most suitable conditions in which colonoscopy can be performed. The only result indicator that directly reflects colonoscopy quality is the one related to polyp resection and removal. This indicator summarizes in a schematic fashion how the procedure should be performed – more polyps analyzed means better visualization, hence a more thorough bowel preparation.

One of the issues or criticisms raised regarding this study is the fact that endoscopists are aware of their being observed since the entire exploration is recorded and subsequently visualized. Such effect can theoretically determine a bias that may modify study data, even though the aim of the study is the systematization of how colonoscopy should be performed. In this first instance, the purpose is not an evaluation of the endoscopist’s skills but to assess whether we can define and put into practice some valid indicators to evaluate a certain manner of colonoscopy performance. This bias could become beneficial subsequently in a positive way. More studies are necessary to clarify which are the most reliable indicators to measure colonoscopy quality. Once the validity of these indicators is defined and demonstrated, they will be used in the setting of a screening program in order to evaluate the required quality parameters.

The efficacy of CRC screening programs depends on several factors; high-quality colonoscopic examinations are desirable to facilitate success for such programs (38).

REFERENCES


