

# Restarting care-related activities in the context of COVID-19: Prioritization, protocols, and procedures. Experience of a gastroenterology outpatient unit in Bogotá, Colombia

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## Abstract

The Colombian Government ordered a mandatory nationwide quarantine in March 2020 due to the spread of the SARS-CoV-2 virus. Since then, outpatient endoscopy units were closed and only urgent procedures were performed in the hospital setting, resulting in a repression of sensitive, priority and elective outpatient endoscopic procedures. The rate of spread of the virus was contained and it did not progress exponentially as in other countries; in the meantime, gastroenterology services were provided in the form of teleconsultation. The mitigation measures and the containment of the virus allowed the Mayor's Office of Bogotá and the National Government to issue notices with recommendations for the provision of some regulated outpatient services in May 2020, thus creating a window of opportunity to care for patients with sensitive diseases. Under this legal and epidemiological framework, the provision of digestive endoscopy services was restarted at EMDIAGNOSTICA S.A.S. This study presents the strategies for scheduling and performing endoscopic procedures in an outpatient gastroenterology unit during COVID-19 pandemic in Colombia and describes a system for prioritizing procedures according to medical criteria, ranging from care by teleconsultation and/ or an application of a telephone survey and the use of a medically necessary, time-sensitive (MENTS) scale adapted for digestive endoscopy. It also describes changes in infrastructure, methodology implemented for protection of human talent and patients, and post procedure follow-up for feedback, safety and satisfaction degree evaluation in care.

## Keywords

SARS-CoV-2; COVID-19; Endoscopy; Quality; Safety; Sedation.

## INTRODUCTION

The Coronavirus Disease 2019 (COVID-19), declared a pandemic by the World Health Organization (WHO) on March 11, 2020 (1), substantially changed the way how health care services are delivered to the community and by health care workers. In Colombia, a compulsory lockdown was decreed in March 2020, with the aim of delaying the peak of infections, allowing the hospital network to

make the necessary adaptations so as not to collapse (2,3). Isolation protocols were established on March 6 (4) after the first case was reported in the country, and they have been adjusted to allow the movement of people and return to work activities, but always considering the security of citizens as a priority (5, 6).

However, government guidelines specifically restricted the provision of some non-essential medical services (2,3). Specifically, services provided to diagnose time-sensitive

conditions (i.e., in which the time of diagnosis or the start of treatment have a direct impact on prognosis) had to comply with the lockdown and the preventive isolation measures established by the Government while proposing options to continue performing diagnostic procedures that are essential for the prognosis of some diseases without posing a greater risk to health.

Considering that gastrointestinal endoscopies are (or may be) aerosol generating procedures, strategies were developed to reduce the risk of transmission while being performed. After being closed for 7 weeks, the outpatient units were able to restart their services without knowing if they would be suspended again. Consequently, it was necessary to define how to restart these activities and who would undergo these procedures in endoscopy units.

In our unit, once the problem was identified, a strategy began to be planned to deal with the situation. Thus, a multidisciplinary group was created (gastroenterologists, anesthesiologists, lawyers, nurses, and financial advisors) who, through online meetings, established a protocol with recommendations to reactivate the provision of services based on the prioritization scheme described below.

## METHODOLOGY

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Before reopening the unit, a non-systematic search was conducted in the medical literature (Medline-Pubmed, Scielo, Embase, and Google-Scholar) on protocols, clinical practice guidelines, and reports in countries that had already experienced this situation. Moreover, we participated in several virtual conferences and reviewed the decrees issued by government agencies (2,3). It was possible to learn about the measures that had been adopted in this regard worldwide, reproduce the actions that proved to be effective, and modify or adapt those that could be improved.

Multiple international scientific organizations (7-16) designed and socialized guidelines that were reviewed and adapted to our context, since their application had to be adjusted to the needs of an outpatient endoscopy unit that does not provide emergency services. All these contingency measures were planned in accordance with the behavior of COVID-19 in Bogotá.

## LEGAL FRAMEWORK

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The Ministry of Health adopted the action plan for the provision of health care services during the pandemic containment and mitigation phases through Resolution 536 of 2020, and, based on it, care plans were structured in the context of the crisis. Thus, in April and May, guidelines were issued for the gradual reopening of services during the health emergency mitigation and control phases. Emphasis

was placed on outpatient surgery, surgery with low-risk intensive care unit (ICU) requirement, short hospital stay surgeries, and non-deferrable surgeries.

In update made on June, recommendations regarding outpatient services, surgery, and diagnostic support procedures were eliminated, stating that “care services may be provided, and undifferentiated procedures may be performed in such a way as to guarantee the protection and safety of users and the human talent associated with the provision of such care services.”

Likewise, the Ministry of Health stated that the actions to be followed and the possibility of reactivating the provision of health care services depended on the analysis, in each territory, of factors such as the trend of new cases, the availability of human resources and personal protection equipment (PPE), the use of the installed capacity, and the behavior in the provision of the service. Therefore, the implementation of any reactivation plan must necessarily go through the assessment of the conditions in the territories and the guidelines provided in this regard by the competent authority.

The Health Department of Bogotá issued Notice 029 of April 29, 2020, and Notice 036 of May 12, 2020, which contained recommendations for the operational organization of outpatient services in the context of the COVID-19 pandemic emergency. In said notices, it was concluded that the provision of the health care services in the framework of the health emergency is associated with a dynamism conditioned by the evolution of the pandemic at the territorial level and that, as circumstances evolve, these guidelines will be modified. This process requires constant updating and coordination among the actors involved.

## OPERATIONAL PHASE

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### Preparation of administrative and assistance personnel

Individual risk factors were estimated when classifying administrative and healthcare personnel available for on-site care during the pandemic. In addition, the staff was divided into two groups, so that endoscopy services could be provided rotating the staff every two weeks. Agreements and online workshops were made for the adaptation, use and disposal of PPE. Also, PPE necessary to guarantee at least 4 months of operation was acquired.

Furthermore, some documents and procedures necessary to ensure compliance and safe health care services delivery for patients, staff, and the community were adapted, modified, and created (**Table 1**). Finally, it was agreed that, at the beginning of each day, all staff would make a statement of their health status and risk of COVID-19 (**Table 2**). All

**Table 1.** Documents adapted, modified, or created to reactivate activities during the COVID-19 pandemic\*

COVID-19 Information Booklet for Patients
Informed consent for endoscopic procedures (endoscopy, colonoscopy, polypectomy)
Informed consent for face-to-face consultation
Informed consent for teleconsultation
Informed consent for sedation
Informed consent for patient companion
COVID-19 health and risk statement for patients, companions, providers, clients, administrative and care staff
Occupational health and safety management system
Process roadmap and verification
COVID-19 biosafety manual, checklist, and contingency measures
COVID-19 Sedation note
Note for face-to-face consultation and procedures
Process and checklist for scheduling endoscopic procedures for patients with semi-urgent and priority indications
Procedure and checklist for scheduling a face-to-face consultation during the pandemic
COVID-19 Sedation procedure

\*List of institutional documents from Emdiagnóstica S.A.S., which were adapted, modified, or edited to fit the COVID-19 pandemic scenario. It details the information provided to patients, informed consent forms, organizational documents related to safety and occupational health, protocols created to guide medical care provision, the form with the basic information recorded in the gastroenterology and anesthesiology clinical records, checklists, and roadmaps. COVID-19: Coronavirus Disease 2019

**Table 2.** Statement of health status and risk of COVID-19\*

Questions	Answers	
1. Do you suffer from severe chronic lung disease (asthma, COPD) or high blood pressure, chronic kidney failure, or serious heart disease (including heart attacks, heart valve disease, or replacements), or do you have an implantable cardioverter-defibrillator? Do you have a pacemaker? Do you suffer from known severe immune system disorders?	Yes	No
2. In the past 2 weeks, have you, or anyone you live with, had a fever of 101°F (38°C) or higher associated with headache, dry cough, breathing difficulty, sore throat, loss or change to your sense of smell or taste (ability to perceive flavors), or had recent diarrhea?	Yes	No
3. Do you, or someone you live with, have a diagnosis of COVID-19 infection or have been in contact with someone who has a suspected or confirmed diagnosis of this disease?	Yes	No
4. Have you, or someone you live with, recently been out of the country or in contact with travelers from abroad?	Yes	No
5. If you are part of the healthcare workforce, do you suspect that you have had unprotected contact with a COVID-19 patient?	Yes	No
6. Are you willing to fully and properly use all the personal protection equipment required, which will be provided during your stay at Emdiagnóstica S.A.S.?	Yes	No
7. Can you say that, to the best of your knowledge, you have not been diagnosed with COVID-19 infection to date?	Yes	No
8. Did you receive the PPE in good condition?	Yes	No

\*Form delivered daily, at the beginning of the day, to all the Emdiagnóstica S.A.S. staff to confirm their current state of health and to estimate the risk of COVID-19. Its completion is mandatory. Once the questionnaire has been completed in its entirety, the employee's body temperature has been taken, the risk of COVID-19 has been confirmed to be low, and the health condition is found to be optimal, the employee is authorized to enter the unit facilities. All forms are stored in the physical archive of Emdiagnóstica S.A.S. COVID-19: Coronavirus Disease 2019; COPD: chronic obstructive pulmonary disease; PPE: personal protective equipment.

patients would be considered suspected cases and, consequently, protective measures would be the same in all cases.

### Adaptation of the endoscopy unit facilities

Based on the studies by Repici et al. (17), Cennamo et al. (18) and the American Gastroenterological Association (AGA) (10), areas were demarcated according to the degree of exposure to droplets and aerosols, and the unit was equipped with a negative pressure air exchange system (15 cycles every hour) with high efficiency particulate air (HEPA) filters.

### Patient selection and procedure scheduling

The provision of the teleconsultation service was established when the preventive lockdown was decreed; in this case, the Government guidelines regarding remote appointments were accepted. This service was used to prioritize the performance of endoscopic procedures. Patient scheduling in this phase was based on national and international recommendations on prioritization, according to the relevance of the study, the health condition of the patient, and the local behavior of the COVID-19 infection curve. Moreover, the recommendations of the Asian-Pacific Society for Digestive Endoscopy (A-PSDE) for an outpatient endoscopy unit (15) were adapted to the local context (Table 3), as well as

the suggestions of the AGA (10) regarding the 8-week limit for performing time-sensitive procedures.

Thus, the procedures were scheduled based on the criteria of the gastroenterologist, on the diagnostic probability of having cancer and the presence of alarming symptoms, and on the risk/benefit balance during the pandemic. Three groups were created: priority 1 (procedures that, according to the gastroenterologist, had to be performed within the following two weeks), priority 2 (to be performed within weeks 3 and 4) and priority 3 (to be performed between weeks 5 and 8) (Figure 1). All patients were given recommendations on alarm symptoms that required performing the procedure earlier.

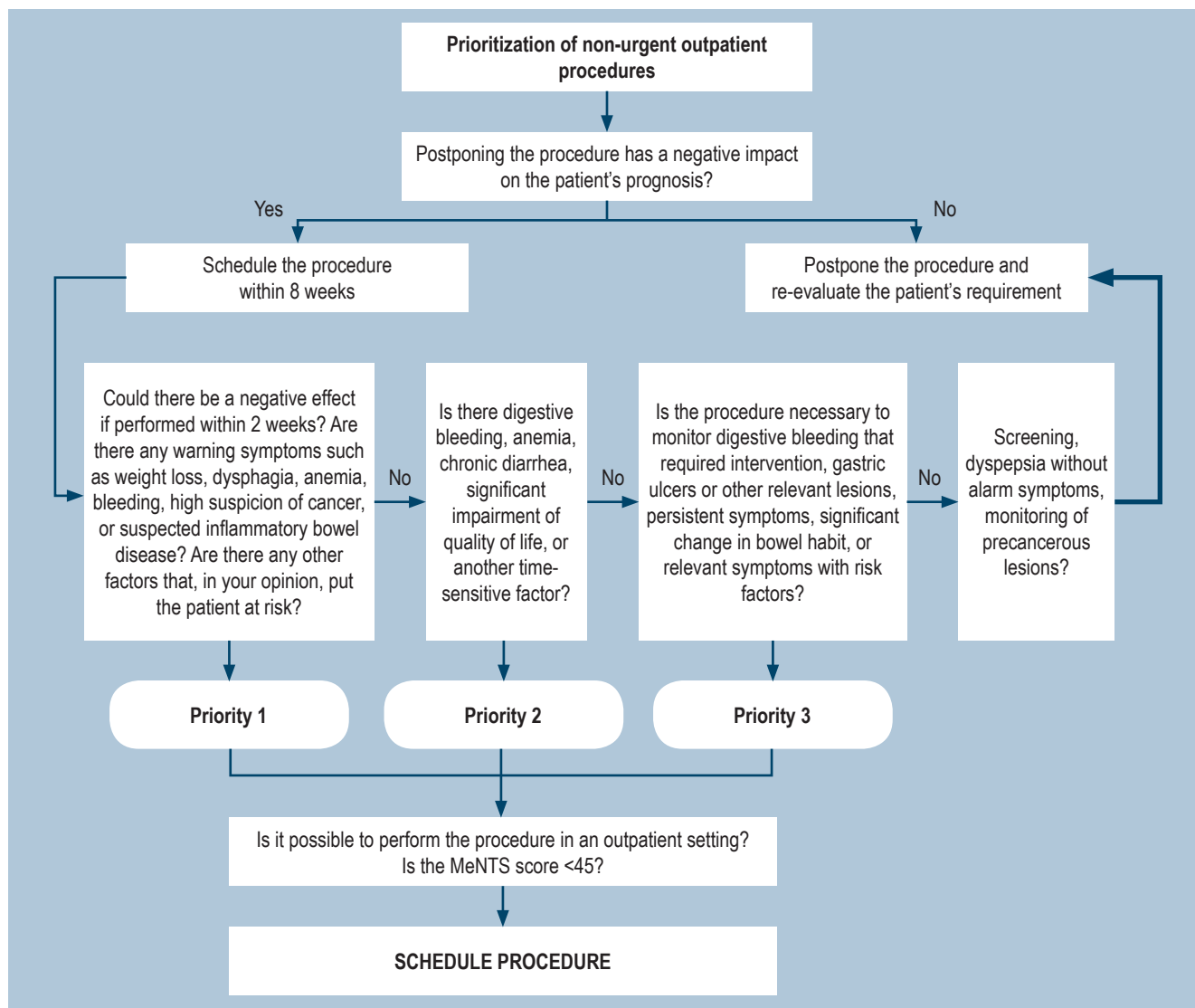
The day before assigning the appointment for the procedure and confirming it, the scheduling staff checked again for COVID-19 risk. Meanwhile, in case of referred patients, the procedure was performed, and a checklist was created to schedule prioritized patients (Table 4). When any of the patient's answers coincided with a warning signal, the patient was scheduled for teleconsultation, during which, besides defining the indication and the time of the procedure, the level of care in the context of COVID-19 was established.

The MeNTS-ENDOGI scale was used for this purpose (Figure 2). The version used in the paper by Prachand et al. (19) was adapted, based on adjustments made on the assessed criteria, to apply it in the scenario of endoscopic procedures and achieve objectivity in the decision. The safety of

**Table 3.** Performance of endoscopic procedures during the COVID-19 pandemic\*

Endoscopy services during the COVID-19 pandemic with available PPE			
Behavior of COVID-19 in the community	Endoscopy services offered	Strategy	Average programming time per room
Exponential increase in new cases	None		
Rapid (non-exponential) increase in new cases	Performing time-sensitive procedures, postponing elective endoscopies	Prioritization 1, 2 and 3	Every hour
Downward trend in new cases	Performing time-sensitive procedures, while initiating elective procedures	Prioritization 1, 2 and 3. Diagnosis in patients with risk factors. High risk screening. Cancer monitoring. Procedures required by other specialties to start any management.	Every 40 min
No new cases diagnosed in the last 2 weeks	Normalization	Adding necessary diagnostic procedures without risk factors Injury follow-up. Average risk screening.	30 min

\*Strategies used to define the scheduling and performance of endoscopic diagnostic procedures during the COVID-19 pandemic, with available PPE, and considering the behavior of the local epidemiological curve, based on the recommendations of the AGA (10) and the A-PSDE (15). COVID-19: Coronavirus Disease 2019; PPE: personal protective equipment.



**Figure 1.** Flow chart for prioritizing endoscopic procedures during the mitigation phase and the rise of the COVID-19 curve. The decision to schedule or not a procedure is based on a judicious clinical evaluation and the determination of the appropriateness of the study during the pandemic period. In addition, the estimation of the MeNTS-ENDOGI score is taken into consideration. MeNTS: *Medically Necessary Time-Sensitive score*.

the outpatient environment for the patient was set in up to 45 points. If the score was higher, the recommendation was to refer the patient to a hospital to perform the procedure.

## OPERATIVE PHASE

### Personal protective equipment and biosecurity equipment

The guidelines issued by the *American College of Surgeons* (20), AGA (10), the Ministry of Health and the Colombian Association of Infectiology (11) were followed to establish

the use of PPE according to risk areas and activity, including patients (Table 5). Meanwhile, for safe use and disposal of PPE, online training and computer graphics were available in each room. Only PPE delivered by the institution was allowed for use.

### Sedation protocol

All endoscopic procedures are performed in the unit under sedation administered by anesthesiologists. Prior to the pandemic, it was customary to keep patients under moderate sedation according to the Practice Guidelines for

**Table 4.** Process and checklist for scheduling endoscopic procedures for priority patients\*

Questions	Answers	
1. Do you agree to schedule the procedure despite the COVID-19 pandemic? Leave a note in the medical history and schedule for teleconsultation	Yes	No
2. Have you experienced alarming weight loss, anemia, or significant changes in bowel habits (diarrhea/constipation)? Have you had difficulty swallowing, have you had digestive bleeding, or do you know if your doctor has a high suspicion of cancer in your case?	Yes	No
3. Are you on blood thinners? If yes, continue with question 4; otherwise, go to question 7	Yes	No
4. If you are on blood thinners, did you follow the medical recommendations for this procedure, such as suspension or bridging therapy?	Yes	No
5. Are you over 70 years of age?	Yes	
6. Do you live in a nursing home or senior citizen facility?	Yes	No
7. Do you have severe chronic lung disease (asthma, COPD), or suffer from inadequately controlled high blood pressure, chronic kidney failure or severe heart disease (including heart attacks, heart valve disease, or replacement), or have an implantable cardioverter defibrillator? Do you have a pacemaker? Do you suffer from known severe immune system disorders?	Yes	No
8. Are you obese or diabetic? Do you suffer from sleep apnea? If YES, schedule an online appointment	Yes	No
9. Are you pregnant?	Yes	No
10. Are you able to walk 3 blocks or climb two floors without being short of breath?	Yes	No
11. In the past 2 weeks, have you, or anyone you live with, had a fever of 101°F (38°C) or higher associated with headache, dry cough, breathing difficulty, sore throat, loss or change to your sense of smell or taste (ability to perceive flavors), or had recent diarrhea?	Yes	No
12. Do you, or someone you live with, have a diagnosis of COVID-19 infection or have been in contact with someone who has suspected or confirmed diagnosis of this disease?	Yes	No
13. Have you, or someone you live with, recently been out of the country or in contact with travelers from abroad in the past 2 weeks?	Yes	No

\*List of questions asked to each patient by the Em Diagnostica S.A.S. appointment scheduling center by telephone to define whether to schedule the procedure or, failing that (areas shaded in blue), refer the patient to a specialized gastroenterology assessment in the teleconsultation modality to individualize and establish the behavior. This form works as a screening test to classify patients according to their risk and optimize appointment scheduling. COVID-19: Coronavirus Disease 2019; COPD: chronic obstructive pulmonary disease.

Sedation and Analgesia by non-Anesthesiologist (21). Due to the requirement of depth of anesthesia to intubate safely and pass the cricopharyngeus muscle without cough reflex, it was decided to administer the medications conventionally used (remifentanyl and propofol) until this objective was achieved.

However, to reduce prolonged periods of apnea with desaturation and hypoxemia, it was established that all patients would receive routine preoxygenation for 3 min with oxygen by nasal cannula at 2-3 L/min (below the isolation barrier), with deep inspirations. In case the patient's apnea was accompanied by desaturation (oxygen saturation by pulse oximetry [SpO<sub>2</sub>] <90 %), the defined action plan was to increase oxygen flow up to 5 L/min and perform maneuvers for airway permeability.

If desaturation persisted, the indication was to suspend the endoscopic procedure and administer oxygen through a well-sealed mask using a two-handed E-C clamp technique to avoid endotracheal intubation or positive pressure ventilation as much as possible. A checklist of complications associated with sedation was adopted (22).

### Endoscopic procedures

Patients were scheduled alternately in each of the three available rooms, allowing for ventilation, cleaning, and disinfection of each area prior to the admission of the next patient. This way, despite having negative pressure, each room is used again after 1 hr., initially using 30% of the installed capacity. On the other hand, a plastic barrier is

### A. Score model

### B. The procedure can be scheduled in the outpatient environment

### C. The procedure cannot be scheduled in the outpatient environment

MeNTS-ENDOGI - Sedación	Puntaje	0
1. Duración del procedimiento menos de 5 <input type="checkbox"/> de 5 a 10 <input type="checkbox"/> de 10 a 20 <input type="checkbox"/> de 21 a 30 <input type="checkbox"/> más de 30 <input type="checkbox"/>		
2. Número de personas en la sala 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 o más <input type="checkbox"/>		
3. Probabilidad de intubación (porcentaje) menos de 1% <input type="checkbox"/> de 1 a 5 % <input type="checkbox"/> de 6 a 10% <input type="checkbox"/> de 11 a 25% <input type="checkbox"/> más de 25% <input type="checkbox"/>		
4. Tipo de procedimiento Endoscopia <input type="checkbox"/> Colonoscopia <input type="checkbox"/> Endoscopia y colonoscopia <input type="checkbox"/>		
5. Opciones no invasivas efectivas para el diagnóstico y tratamiento No disponibles <input type="checkbox"/> Efectividad menor a 40% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 41% y 65% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 66% y 80% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 81% y 95% en comparación con métodos invasivos <input type="checkbox"/> Disponibles, con efectividad equivalente al método invasivo <input type="checkbox"/>		
6. Disponibilidad de recursos / insumos para procedimiento no invasivo No aplicables <input type="checkbox"/> Difícil conseguir <input type="checkbox"/> Medicamento disponible <input type="checkbox"/> Fácil conseguir <input type="checkbox"/> Disponibles <input type="checkbox"/>		
7. Efecto en el progreso de la enfermedad a 2 semanas en caso de no realizar el procedimiento Significativamente peor <input type="checkbox"/> Peor <input type="checkbox"/> Moderadamente peor <input type="checkbox"/> Levemente peor <input type="checkbox"/> Equivalente <input type="checkbox"/>		
8. Edad (años) menos de 20 <input type="checkbox"/> de 21 a 40 <input type="checkbox"/> de 41 a 50 <input type="checkbox"/> de 51 a 65 <input type="checkbox"/> más de 65 <input type="checkbox"/>		
9. Enfermedad pulmonar (asma, EPOC, fibrosis quística) Ninguna <input type="checkbox"/> Uso ocasional de inhaladores <input type="checkbox"/> No controlada <input type="checkbox"/>		
10. Apnea obstructiva del sueño No <input type="checkbox"/> Leve / moderado <input type="checkbox"/> uso de CPAP / BPAP <input type="checkbox"/>		
11. Uso de medicamentos para enfermedad cardiovascular (HTA, falla cardiaca, enfermedad coronaria) No <input type="checkbox"/> 1 medicamento <input type="checkbox"/> 2 medicamentos <input type="checkbox"/> 3 medicamentos <input type="checkbox"/> 4 medicamentos o más <input type="checkbox"/>		
12. Diabetes No <input type="checkbox"/> En tratamiento con dieta <input type="checkbox"/> En tratamiento vía oral <input type="checkbox"/> En tratamiento con insulinas <input type="checkbox"/>		
13. Inmunosupresión No <input type="checkbox"/> Moderada <input type="checkbox"/> Severa <input type="checkbox"/>		
14. Síntomas de enfermedad similar a influenza (fiebre, tos, dolor de garganta, mialgias, diarrea) No <input type="checkbox"/> Sí <input type="checkbox"/>		
15. Contacto con una persona COVID-19 positiva o sospechosa en los últimos 14 días No <input type="checkbox"/> Posiblemente no <input type="checkbox"/> Con dudas <input type="checkbox"/> Posiblemente sí <input type="checkbox"/> Sí <input type="checkbox"/>		

MeNTS-ENDOGI - Sedación	Puntaje	24
1. Duración del procedimiento menos de 5 <input type="checkbox"/> de 5 a 10 <input type="checkbox"/> de 10 a 20 <input checked="" type="checkbox"/> de 21 a 30 <input type="checkbox"/> más de 30 <input type="checkbox"/>		
2. Número de personas en la sala 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input checked="" type="checkbox"/> 4 <input type="checkbox"/> 5 o más <input type="checkbox"/>		
3. Probabilidad de intubación (porcentaje) menos de 1% <input checked="" type="checkbox"/> de 1 a 5 % <input type="checkbox"/> de 6 a 10% <input type="checkbox"/> de 11 a 25% <input type="checkbox"/> más de 25% <input type="checkbox"/>		
4. Tipo de procedimiento Endoscopia <input type="checkbox"/> Colonoscopia <input checked="" type="checkbox"/> Endoscopia y colonoscopia <input type="checkbox"/>		
5. Opciones no invasivas efectivas para el diagnóstico y tratamiento No disponibles <input type="checkbox"/> Efectividad menor a 40% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 41% y 65% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 66% y 80% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 81% y 95% en comparación con métodos invasivos <input type="checkbox"/> Disponibles, con efectividad equivalente al método invasivo <input type="checkbox"/>		
6. Disponibilidad de recursos / insumos para procedimiento no invasivo No aplicables <input type="checkbox"/> Difícil conseguir <input type="checkbox"/> Medicamento disponible <input type="checkbox"/> Fácil conseguir <input type="checkbox"/> Disponibles <input type="checkbox"/>		
7. Efecto en el progreso de la enfermedad a 2 semanas en caso de no realizar el procedimiento Significativamente peor <input type="checkbox"/> Peor <input checked="" type="checkbox"/> Moderadamente peor <input type="checkbox"/> Levemente peor <input type="checkbox"/> Equivalente <input type="checkbox"/>		
8. Edad (años) menos de 20 <input type="checkbox"/> de 21 a 40 <input type="checkbox"/> de 41 a 50 <input checked="" type="checkbox"/> de 51 a 65 <input type="checkbox"/> más de 65 <input type="checkbox"/>		
9. Enfermedad pulmonar (asma, EPOC, fibrosis quística) Ninguna <input type="checkbox"/> Uso ocasional de inhaladores <input type="checkbox"/> No controlada <input type="checkbox"/>		
10. Apnea obstructiva del sueño No <input type="checkbox"/> Leve / moderado <input type="checkbox"/> uso de CPAP / BPAP <input type="checkbox"/>		
11. Uso de medicamentos para enfermedad cardiovascular (HTA, falla cardiaca, enfermedad coronaria) No <input type="checkbox"/> 1 medicamento <input type="checkbox"/> 2 medicamentos <input type="checkbox"/> 3 medicamentos <input type="checkbox"/> 4 medicamentos o más <input type="checkbox"/>		
12. Diabetes No <input type="checkbox"/> En tratamiento con dieta <input type="checkbox"/> En tratamiento vía oral <input type="checkbox"/> En tratamiento con insulinas <input type="checkbox"/>		
13. Inmunosupresión No <input type="checkbox"/> Moderada <input type="checkbox"/> Severa <input type="checkbox"/>		
14. Síntomas de enfermedad similar a influenza (fiebre, tos, dolor de garganta, mialgias, diarrea) No <input type="checkbox"/> Sí <input type="checkbox"/>		
15. Contacto con una persona COVID-19 positiva o sospechosa en los últimos 14 días No <input type="checkbox"/> Posiblemente no <input type="checkbox"/> Con dudas <input type="checkbox"/> Posiblemente sí <input type="checkbox"/> Sí <input type="checkbox"/>		

MeNTS-ENDOGI - Sedación	Puntaje	50
1. Duración del procedimiento menos de 5 <input type="checkbox"/> de 5 a 10 <input type="checkbox"/> de 10 a 20 <input type="checkbox"/> de 21 a 30 <input checked="" type="checkbox"/> más de 30 <input type="checkbox"/>		
2. Número de personas en la sala 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input checked="" type="checkbox"/> 4 <input type="checkbox"/> 5 o más <input type="checkbox"/>		
3. Probabilidad de intubación (porcentaje) menos de 1% <input checked="" type="checkbox"/> de 1 a 5 % <input type="checkbox"/> de 6 a 10% <input type="checkbox"/> de 11 a 25% <input type="checkbox"/> más de 25% <input type="checkbox"/>		
4. Tipo de procedimiento Endoscopia <input type="checkbox"/> Colonoscopia <input type="checkbox"/> Endoscopia y colonoscopia <input checked="" type="checkbox"/>		
5. Opciones no invasivas efectivas para el diagnóstico y tratamiento No disponibles <input type="checkbox"/> Efectividad menor a 40% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 41% y 65% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 66% y 80% en comparación con métodos invasivos <input type="checkbox"/> Efectividad entre 81% y 95% en comparación con métodos invasivos <input type="checkbox"/> Disponibles, con efectividad equivalente al método invasivo <input type="checkbox"/>		
6. Disponibilidad de recursos / insumos para procedimiento no invasivo No aplicables <input type="checkbox"/> Difícil conseguir <input type="checkbox"/> Medicamento disponible <input type="checkbox"/> Fácil conseguir <input type="checkbox"/> Disponibles <input type="checkbox"/>		
7. Efecto en el progreso de la enfermedad a 2 semanas en caso de no realizar el procedimiento Significativamente peor <input type="checkbox"/> Peor <input type="checkbox"/> Moderadamente peor <input type="checkbox"/> Levemente peor <input type="checkbox"/> Equivalente <input type="checkbox"/>		
8. Edad (años) menos de 20 <input type="checkbox"/> de 21 a 40 <input type="checkbox"/> de 41 a 50 <input type="checkbox"/> de 51 a 65 <input type="checkbox"/> más de 65 <input checked="" type="checkbox"/>		
9. Enfermedad pulmonar (asma, EPOC, fibrosis quística) Ninguna <input type="checkbox"/> Uso ocasional de inhaladores <input type="checkbox"/> No controlada <input type="checkbox"/>		
10. Apnea obstructiva del sueño No <input type="checkbox"/> Leve / moderado <input type="checkbox"/> uso de CPAP / BPAP <input type="checkbox"/>		
11. Uso de medicamentos para enfermedad cardiovascular (HTA, falla cardiaca, enfermedad coronaria) No <input type="checkbox"/> 1 medicamento <input type="checkbox"/> 2 medicamentos <input type="checkbox"/> 3 medicamentos <input type="checkbox"/> 4 medicamentos o más <input type="checkbox"/>		
12. Diabetes No <input type="checkbox"/> En tratamiento con dieta <input type="checkbox"/> En tratamiento vía oral <input type="checkbox"/> En tratamiento con insulinas <input checked="" type="checkbox"/>		
13. Inmunosupresión No <input type="checkbox"/> Moderada <input type="checkbox"/> Severa <input type="checkbox"/>		
14. Síntomas de enfermedad similar a influenza (fiebre, tos, dolor de garganta, mialgias, diarrea) No <input type="checkbox"/> Sí <input checked="" type="checkbox"/>		
15. Contacto con una persona COVID-19 positiva o sospechosa en los últimos 14 días No <input type="checkbox"/> Posiblemente no <input type="checkbox"/> Con dudas <input type="checkbox"/> Posiblemente sí <input checked="" type="checkbox"/> Sí <input type="checkbox"/>		

**Figure 2.** MeNTS-ENDOGI score. These schemes represent the scale used by Emdiagnóstica S.A.S. The scale was adapted based on the original MeNTS model by Prachand et al. (19). Panel A shows the blank format before completion. Panel B shows the result after administering the MeNTS-ENDOGI scale to a patient with low risk of complications and dissemination of COVID-19, who can be scheduled for an outpatient procedure. Panel C presents the results of a patient who, due to risk factors (clinical factors and risk of dissemination of COVID-19 infection), cannot be scheduled for an outpatient procedure. Both cases are simulated scenarios and do not disclose information from real patients. Emdiagnóstica S.A.S. set 45 points as the cut-off point to define whether the procedure should be performed in an outpatient or inpatient environment.

**Table 5.** PPE according to the area of exposure and the task performed

Exposure area / task performed	PPE
Procedure area (care staff: gastroenterologists, anesthesiologists, nursing assistants)	N95 respirators Goggles Face shield Hat Double non-surgical gowns (the external gown is changed for each new visit) Two pairs of gloves Shoe cover
Administrative area (secretaries, receptionist, administrative assistant)	Mask and face shield
Patients in procedure area	Mask provided by the unit Plastic barrier Fenestrated drape

List of PPE classified according to exposure risk level. For risk assignment, the work area and the task performed by each employee were considered (to determine the degree of exposure to droplets and aerosols). The American College of Surgeons guidelines (20) were used to make this list. PPE: personal protective equipment.

used for endoscopy (23) and a fenestrated drape for colonoscopy. In all cases, the registration of quality indicators for endoscopy and colonoscopy are recorded in the roadmap and verification of processes form, as usual, while the lesions found are resected depending on the characterization of each lesion. In addition, the health care staff changes their external anti-fluid gown for receiving new patients.

## POSTOPERATIVE STAGE

### Follow-up and feedback

A satisfaction survey is administered to all patients for follow-up after the procedure and for monitoring complications after polypectomy between 12-48 hr. and at 7 and 14 days. They are also asked about their satisfaction level and their sense of safety while they were in the unit (Table 6).

## DISCUSSION

The mandatory lockdown managed to contain the pandemic's exponential growth in the country, providing

**Table 6.** Questions asked to patients to monitor their condition during the COVID-19 pandemic\*

1. Following your endoscopic procedure, have you experienced fever, cough, difficulty smelling or tasting, or difficulty breathing? If yes, notify it immediately to the medical management office
2. Following your endoscopic procedure, has your health deteriorated? If yes, notify it immediately to the medical management office.
3. Have any unusual symptoms appeared after your endoscopic procedure?
4. If yes, please state which symptom you have experienced and notify it immediately to the medical management office.
5. Do you think the safety measures were appropriate during the procedure? In your opinion, the endoscopic procedure was: A). Very unsafe B). Unsafe C). Safe D). Very safe
6. Do you have any recommendations or observations to make?

\*Questionnaire for telephone follow-up of patients who underwent an endoscopic diagnostic study at Emdiagnóstica S.A.S. Data obtained are used to assess the possible cases of COVID-19 that have been overlooked due to the window period (asymptomatic patients) and to estimate the degree of patient satisfaction with the service received. The level of safety perceived by the patient during their stay at Emdiagnóstica S.A.S. is also evaluated.

time to avoid the collapse of the ICUs. However, it also led to accumulating an exponential number of non-urgent but time-sensitive procedures that could not be performed due to the closure of outpatient endoscopy units.

Therefore, such procedures were scheduled in the current time window of care to optimize the available resources in the endoscopy units that were prepared to provide safe care services for an uncertain period, which mainly depended on the percentage of ICUs occupation.

The Colombian scenario was different from that of other countries with exponential curves, as they did not implement this containment period. The current pandemic has required leadership and teamwork from health services provision companies to create solutions and evaluate proposals

applicable to a new scenario that did not resemble any other country considering the epidemiological behavior.

The adaptations made to our unit have allowed us to provide our services again with 30% of the installed capacity before the epidemiological peak was reached, planning progressive increases according to the descent of the curve. Determining the criteria for the appropriate utilization of the installed capacity, human talent, and inputs, always in line with the local situation, was essential to answer the question: how to resume our activities?

We acknowledge that the COVID-19 pandemic context is dynamic and even unpredictable by presenting this document as a working experience. Based on the proposal of scheduling patients according to the priorities in this window period, we intend to solve the problem of whom to schedule first in order to optimize diagnostic resources and provide timely treatment to impact the prognosis of patients directly.

The results obtained by registering the procedures, monitoring patients, taking actions to reduce contagion risk, and the diagnoses made are subjects that must be considered in further studies. Resuming these activities has made evident the changes in the practice of the specialty, which imply significant challenges for the sustainable operation of endoscopy units.

Likewise, economic aspects such as higher costs and fewer procedures could threaten the stability of health care institutions and independent providers. Filtering, prioritizing, weighting cases based on clinical criteria, and planning expenses carefully has become even more important now, since the legal framework is constantly changing. Our challenge now is to adapt to continue providing sustainable and quality services.

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