



SPECIAL ARTICLE

## Recommendations for interventional pulmonology during COVID-19 outbreak: a consensus statement from the Portuguese Pulmonology Society

Q4 F. Guedes<sup>a,b,c,\*</sup>, J.P. Boléo-Tomé<sup>d</sup>, L.V. Rodrigues<sup>e,f</sup>, H.N. Bastos<sup>g,h,i</sup>, S. Campinha<sup>j</sup>, M. de Santis<sup>k</sup>, L. Mota<sup>l</sup>, A. Bugalho<sup>m,n</sup>

<sup>a</sup> Centro Hospitalar do Porto (CHP), Hospital Geral de Santo António (HGSA), Unidade de Broncologia, Serviço de Pneumologia, Porto, Portugal

<sup>b</sup> Departamento de Clínicas Veterinárias, Instituto de Ciências Biomédicas de Abel Salazar (ICBAS), Universidade do Porto (UP), Porto, Portugal

<sup>c</sup> Centro de Estudos de Ciência Animal (CECA), Instituto de Ciências, Tecnologias e Agroambiente (ICETA) da Universidade do Porto, Praça Gomes Teixeira, Apartado 55142, 4051-401, Porto, Portugal

<sup>d</sup> Pulmonology Department, Hospital Prof. Doutor Fernando Fonseca, Amadora, Portugal

<sup>e</sup> Pulmonology Department, Hospital Sousa Martins, Unidade Local de Saúde da Guarda, Guarda, Portugal

<sup>f</sup> Faculty of Health Sciences, University of Beira Interior, Covilhã, Portugal

<sup>g</sup> Department of Pneumology, Centro Hospitalar São João, Porto, Portugal

<sup>h</sup> Faculty of Medicine, University of Porto, Porto, Portugal

<sup>i</sup> IBMC/i3S - Instituto de Biologia Molecular e Celular / Instituto de Investigação e Inovação em Saúde, University of Porto, Portugal

<sup>j</sup> Pulmonology Department, Vila Nova de Gaia-Espinho Hospital Center, Vila Nova de Gaia, Portugal

<sup>k</sup> Pulmonology Department, Instituto Português de Oncologia (IPO), Coimbra, Portugal

<sup>l</sup> Pulmonology Department, Hospital Pulido Valente, Centro Hospitalar Lisboa Norte, Lisboa, Portugal

<sup>m</sup> Pulmonology Department, CUF Infante Santo Hospital and CUF Descobertas Hospital, Lisbon, Portugal

<sup>n</sup> Comprehensive Health Research Centre, Chronic Diseases Research Center (CEDOC), NOVA Medical School, Lisbon, Portugal

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### KEYWORDS

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Consensus statement

**Abstract** Coronavirus disease 2019 (COVID-19) is an emerging infectious disease caused by a novel SARS-CoV-2 pathogen. Its capacity for human-to-human transmission through respiratory droplets, coupled with a high-level of population mobility, has resulted in a rapid dissemination worldwide. Healthcare workers have been particularly exposed to the risk of infection and represent a significant proportion of COVID-19 cases in the worst affected regions of Europe.

Like other open airway procedures or aerosol-generating procedures, bronchoscopy poses a significant risk of spreading contaminated droplets, and medical workers must adapt the

\* Corresponding author.

E-mail address: fernando.t.guedes@gmail.com (F. Guedes).

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procedures to ensure safety of both patients and staff. Several recommendation documents were published at the beginning of the pandemic, but as the situation evolves, our thoughts should not only focus on the present, but should also reflect on how we are going to deal with the presence of the virus in the community until there is a vaccine or specific treatment available. It is in this sense that this document aims to guide interventional pulmonology throughout this period, providing a set of recommendations on how to perform bronchoscopy or pleural procedures safely and efficiently.

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## Introduction

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46 Coronavirus Disease 2019 (COVID-19), a new infectious  
47 **Q6** disease that emerged in early December 2019 in Wuhan  
48 (China),<sup>1</sup> is triggered by a novel pathogen with phylogenetic  
49 similarity to what caused the severe acute respiratory syn-  
50 drome (SARS) outbreak in 2003, and was called SARS-CoV-2.<sup>2</sup>  
51 Its capacity for human-to-human transmission and interna-  
52 tional air travel facilitated the rapid dissemination on an  
53 unprecedented scale to the rest of the world.<sup>3,4</sup>

54 In Italy, the latest figures reported that 9% of COVID-19  
55 cases were health care workers (HCW), while in Spain the  
56 rate of medical staff infected reached 26%, the highest in  
57 Europe.<sup>5</sup> In Portugal, by 12th May 2020, 11.3% of infections  
58 occurred in HCW.<sup>6</sup> There are at least two explanations for  
59 such a high number of infected personnel. First, the lack of  
60 proper personal protective equipment (PPE) at the begin-  
61 ning of the epidemic, when assisting both confirmed and  
62 suspected patients with COVID-19. Second, the duration of  
63 exposure to infected patients undergoing aerosol-generating  
64 procedures, such as non-invasive ventilation (NIV) and bron-  
65 choscopy, directly resulting in a significant increase in the  
66 risk of transmission to HCW.

67 The Portuguese Society of Pulmonology recently issued  
68 a set of recommendations for bronchoscopic procedures,<sup>7</sup>  
69 shortly after the diagnosis of the first cases in Portugal.  
70 The document aimed to guarantee the protection of both  
71 patients and medical practitioners, and to ensure that the  
72 healthcare workforce would be conserved to fulfill their  
73 mission throughout the period. Since then, a significant  
74 amount of scientific evidence has been accumulated; so,  
75 the present document gives an update of the available lit-  
76 erature, providing practical suggestions for pulmonologists  
77 undergoing bronchoscopy or pleural procedures in the set-  
78 ting of the current and post-pandemic phases.

## Risk of transmission

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80 Respiratory droplets comprise the main route of SARS-  
81 CoV-2 transmission, although airborne transmission is also  
82 possible through aerosol-generating procedures, such as  
83 bronchoscopy.<sup>8</sup> One study during the H1N1 pandemic pro-  
84 vided experimental evidence that bronchoscopic procedures  
85 increases more than 4 times the viral copy number per litre  
86 in positive air samples.<sup>9</sup> While the heavy droplets rapidly  
87 settle, aerosol particles are much smaller (<5–10 μm) and

are dispersed in the air over extensive distances, posing a  
considerable risk of infection in enclosed spaces, specially  
if poorly ventilated.<sup>10</sup>

The contribution of asymptomatic carriers has also been  
subject of debate.<sup>11,12</sup> A significant proportion of them  
have lung abnormalities on chest CT scans<sup>13,14</sup> and a high  
level of viral shedding may be detected in presymptomatic  
patients,<sup>14</sup> so it is likely that transmission occurs in the early  
stages of infection when patients are either minimally symp-  
tomatic or asymptomatic. Unrecognized patients pose a real  
challenge to infection control and, when not promptly han-  
dled with appropriate airborne precautions, are one of the  
most critical factors for SARS-CoV-2 infection spread in the  
healthcare setting.

## Methods

The Portuguese Pulmonology Society appointed FG to chair  
this consensus group. Seven national IP specialists were  
selected based on their clinical expertise and different  
settings (university vs. non-university hospitals; state vs.  
private hospitals; pulmonologists vs. critical care special-  
ists; ...). At the first online consensus meeting, attended  
by all members, a primary draft with several sections was  
created. This was shared online and further improved by  
written comments and suggestions. Then, each IP specialist  
was assigned a specific section presented in this document  
and was responsible for reviewing and evaluating the rel-  
evant available literature related to the topic. Electronic  
databases (Pubmed, OVID Medline and Embase, Web of Sci-  
ence, Cochrane Central Register of Controlled Trials) were  
used to search for the terms "COVID-19" OR "SARS-CoV-2"  
AND ("bronchoscopy" OR "interventional pulmology" OR  
"thoracocentesis" OR "thoracocentesis" OR "pleural effu-  
sion" OR "pneumothorax" OR "rigid bronchoscopy" OR  
"thoracoscopy" OR "chest drain"). Position papers from  
major health organizations (US Centers for Disease Con-  
trol and Prevention, European Centre for Disease Prevention  
and Control and World Health Organization) and important  
scientific societies (European Respiratory Society, European  
Association for Bronchology and Interventional Pulmonology,  
American Association for Bronchology and Interventional  
Pulmonology, World Association for Bronchology and Inter-  
ventional Pulmonology and British Thoracic Society) were  
also reviewed.

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In a second online conference the complete draft was evaluated by all team members and two working groups were created. They were responsible for discussing and revising different sections, and editing the text for consistency. Afterwards, the final manuscript was distributed to the consensus group members and assessed for final approval.

## Adaptations of the interventional pulmonology (IP) department

Although there is still some heterogeneity in the definition and scope of “interventional pulmonology” (IP), it has become the most widely accepted term to describe the use of techniques for the diagnosis and treatment of a growing number of thoracic disorders.<sup>15</sup>

In the context of this document, the term IP is used to encompass the concepts of bronchoscopy (diagnostic or therapeutic), advanced bronchoscopy (flexible or rigid bronchoscopy and all its associated techniques), pleuroscopy (rigid or semi-flexible) and other simpler pleural techniques (such as thoracentesis, placement of thoracic drainage systems and indwelling pleural catheters). Though we acknowledge this wider definition of IP may be controversial, it covers all technical domains that most Portuguese pulmonologists need to address, and for the purpose of this document, it positions us to issue general recommendations. In the following subsections, specific scenarios of different technical specializations will be addressed in order to overcome this broader definition and to apply it better to individual settings.

The IP department is a high-risk area, given the type of procedures that are performed with airway manipulation and with multiple staff involved. Although this setting is generally designed to deal with occasional airborne infectious diseases, such as tuberculosis, it is not prepared to systematically assess high-risk cases that need additional resources, diminish productivity and effectiveness and generate a huge workload.

Thus, each IP unit must rethink their administrative and logistic circuits in different areas, as well as the type and timing of performed procedures, to protect both HCW and patients. Moreover, as international health associations advocate, an infection-control program in healthcare settings should be implemented, consisting of a three-level hierarchy, including administrative, environmental and engineering controls, and personal protection equipment (PPE).<sup>16</sup> In the following subsections, each of the above listed hierarchic levels are briefly presented.

### Administrative and organizational issues

Administrative and logistic measures are crucial to ensuring safety while still maintaining IP activity.<sup>17</sup> Some general precautions include:

- All referrals and requests to the IP unit must preferably be made by telephone or digital means.
- Upon schedule and 24–48 h prior to arrival at the IP Unit, patients should be contacted by telephone and submitted to a pre-screening checklist that includes questions about 1) recent symptoms suggestive of COVID-19 (e.g. fever,

cough, chills, muscle pain, shortness of breath/difficulty breathing, headache, sore throat, loss of taste or smell); 2) contact with suspicious/confirmed SARS-CoV-2 cases; and 3) occupational exposure.

- Patients who have recent respiratory and infectious symptoms and/or chest imaging suggestive of COVID-19, should have their elective procedures postponed and rescheduled after all symptoms are resolved.
- On arrival at the IP Unit, all patients must be asked again for respiratory symptoms and have their temperature checked.
- If possible, all patients should have at least one negative RT-PCR for SARS-CoV-2 in the 24–48 h preceding the exam. In patients with a positive RT-PCR SARS-CoV-2, the decision to proceed with the intervention will be based on the urgency of the procedure (Chart 1 and Table 1).
- The IP unit should keep a record of deferred patients to reschedule their procedures according to the COVID-19 outbreak situation, as proposed in Table 2.

### Environmental and engineering control

#### Physical space preparation

The design of strategies to minimize risks and a protocol fitting the characteristics of each specific Unit are crucial.<sup>18</sup>

- Reception, administrative, clinical and waiting areas should separate confirmed/high-risk patients from negative/low-risk ones. In addition, inpatients should be segregated from outpatients, either by time or physical location, to prevent cross infection.
- Specific circuits and written workflow plans must be prepared, covering the pre-procedural area, procedural room, post-procedural area, decontamination and reprocessing. The implementation of a flowchart with different areas and walking paths using a visual colour zone system can be useful: 1) red zone for contaminated areas; 2) yellow zone for transition areas, and 3) green zone for non-COVID-19 safe areas<sup>19</sup> (Fig. 1A). These need to be formulated by internal elements from the IP unit with the cooperation of a multidisciplinary team of hospital members, including administration, engineers, and infection control board.
- A specific place to store and retrieve all items required for PPE should be defined inside the Unit.
- A designated area in the Unit should be selected, close to the procedural suite, for gowning and removal of all PPE, according to hospital protocol and standards, in order to reduce exposure to contaminated particles and droplets. When an anteroom is available, it may be used as an area for donning and doffing of PPE (Fig. 1B).
- Stations should be created to facilitate frequent hand hygiene and to distribute waste containers according to local infectious control recommendations. Posters and other visual aids should be placed at strategic locations around the intervention suite to act as reminders.
- Emergency procedures in COVID-19 positive patients should preferably be performed within the ICU environment, with controlled airway through cuffed endotracheal tube and assisted ventilation.

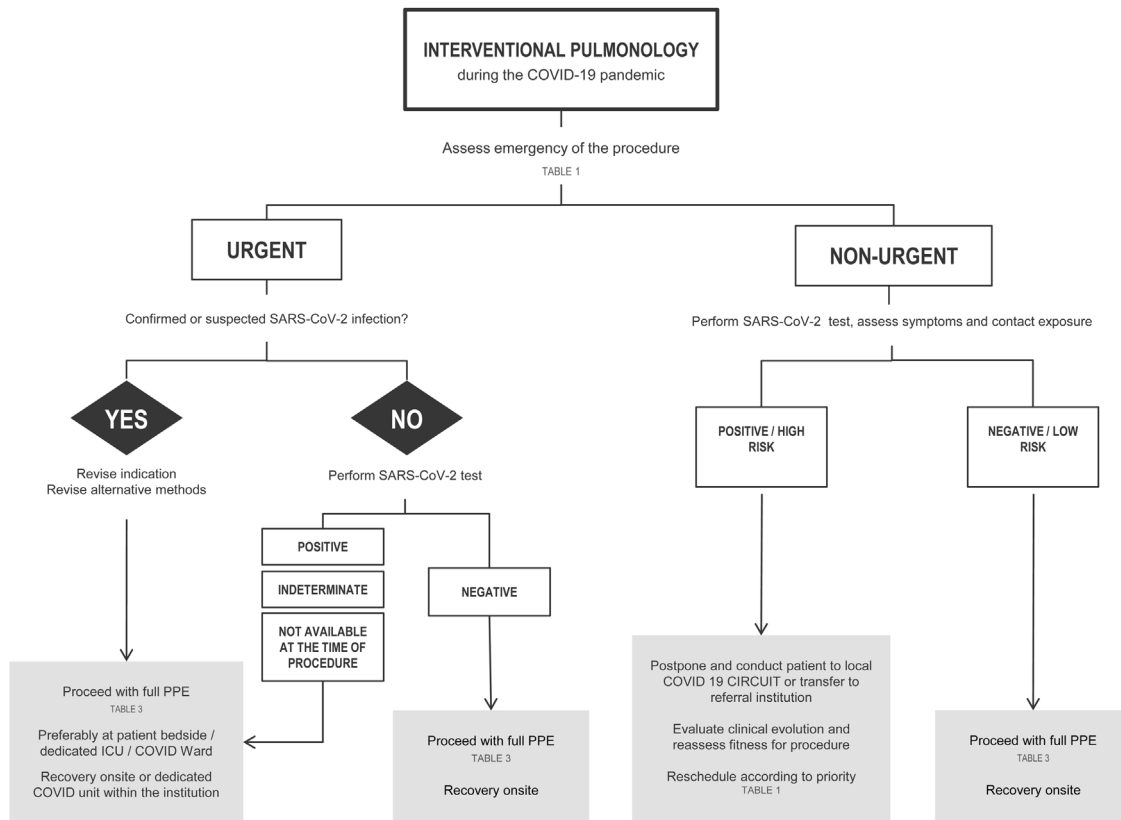


Chart 1 Proposed triage of IP procedures during the COVID-19 outbreak.

Table 1 Priorization of IP exams according to SARS-CoV-2 status and procedure urgency.

URGENT PROCEDURES		NON-URGENT PROCEDURES	
COVID-19 STATUS	PROCEDURE	NON-DELAYABLE (<2-4 weeks)	DELAYABLE (≥4 weeks)
		COVID-19 NEGATIVE/POSITIVE	Massive hemoptysis with airway compromise Acute foreign body aspiration Severe symptomatic central airway obstruction Suspicion of alternative (non-COVID-19) acute and severe infectious disease Airway management in difficult and non-delayable endotracheal intubation or complicated percutaneous tracheotomy Large and symptomatic pleural occupation (air, fluid, blood, pus)
COVID-19 POSITIVE/HIGH RISK	Removal of copious secretions and mucus plugs Possibility of superinfection (community acquired or nosocomial) Severe suspicious cases of COVID-19 that need to be confirmed by bronchoscopy and minimal bronchoalveolar lavage, after at least 2 negative/inconclusive nasopharyngeal RT-PCR SARS-CoV-2 tests	COVID-19 NEGATIVE/LOW RISK	

**Table 2** Schedule of IP procedures according to the stage of COVID-19 pandemic.

COVID-19 in the community	IP Unit
Exponential increase of new cases	Urgent cases – only
Rapid increase of new cases	Urgent cases – only Elective but not delayable – evaluate case-by-case
Decrease in new cases	Urgent cases – full capacity Elective, but not delayable – full capacity Elective and delayable – resume partial capacity
Absence of new cases in the last 2 weeks	Resume all cases with full capacity

- Elective procedures should be reserved for COVID-19 negative patients (Chart 1 and Table 1). Nevertheless, these procedures should still be performed in a dedicated negative pressure room (see below, ventilation requirement) with strict isolation precautions and sufficient ventilation to avoid aerosol contamination.<sup>20</sup> If these requirements are not met in the bronchoscopy suite, then in a different venue, such as an operating theatre, isolation room or the ICU with negative pressure, if available.
- If negative pressure rooms are unavailable throughout the institution, a specific and dedicated room with adequate natural ventilation (see requirement below) may be an alternative, provided that appropriate intervals between procedures are reserved and that the suspected COVID-19 cases be programmed after all planned non-COVID daily activity, so that the unit can be carefully cleaned (following the disinfection policy) and ventilated.
- Keep the endoscopy room for procedures only (all other activities, such as planning, reporting and laboratory requisition should take place elsewhere).
- Suspected and confirmed cases of COVID-19 must be placed in an airborne infection isolation room with negative pressure before and after the procedure. Low-risk and negative patients can remain in the pre-procedural and recovery area, if there is adequate room ventilation,

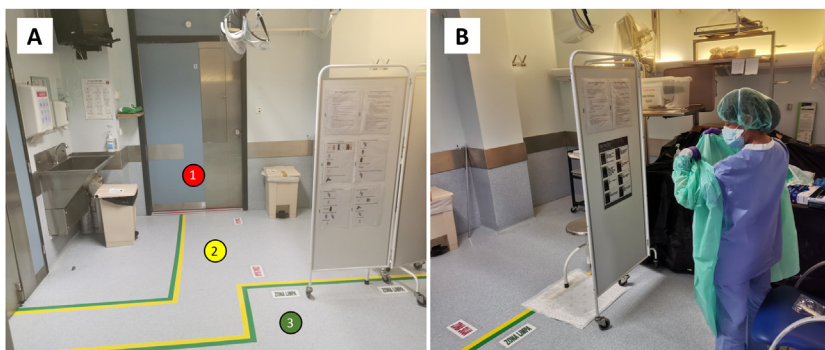
protective equipment (e.g. surgical mask) and physical distance (>2 m) from other negative patients.

**Ventilation**

- Patient source control strategies, such as wearing a mask should be encouraged.
- Whenever feasible, it is recommended procedures are performed in a room that meets the ventilation requirements for Airborne Infection Isolation (AII), ensuring the dilution and removal of contaminated air. The preferred system is a negative pressure room with at least 12 air changes per hour (ACH) with airflow direction control (single-pass or recirculation systems with HEPA filtration). Alternatively, natural ventilation with an airflow of at least 160 L/s is an option.<sup>19,21</sup>
- Enough time should be allowed to ensure that contaminated air is removed from the room before performing another procedure in the same room (depending on ACH and disinfection methods, but at least 30 min). Local adaptations must be considered according to the characteristics of the IP unit.

**Cleaning and disinfecting patient care equipment and rooms**

- Endoscopes are considered semi-critical medical instruments according to the Spaulding classification.<sup>22</sup> Recommendations from the Centers for Disease Control and Prevention (CDC) on reprocessing procedures should be followed. These include pre-cleaning, leak-testing, manual cleaning and visual inspection followed by disinfection/sterilization.
- A high-level manual disinfection or using an automated endoscope reprocessor is recommended.
  - Proper storage and documentation are also an integral part of the reprocessing workflow.
  - A pathway of contaminated equipment must be defined, as well as adequate packaging to minimize exposure (for example, a hermetic box).
  - If available, disposable bronchoscopes are recommended for confirmed COVID-19 patients with clear advantages in portability, post-procedural handling and cross contamination risk.<sup>23</sup>



**Figure 1** A. Implementation of specific circuits with colour visual zone system to distinguish contaminated (1, red zone), transition (2, yellow zone) and safe cleaned areas (3, green zone). B. Designated area for donning and doffing of PPE, where posters and other visual aids were placed strategically to act as reminders.

**Table 3** Specifications for personal protective equipment during IP procedures.

PPE	Characteristics/specifications/standards	Observations
Gloves	<input type="radio"/> Single-use <input type="radio"/> Waterproof <input type="radio"/> Standard EN ISO 374-2:2014, 374-3:2014 e 374.5:2016 <input type="radio"/> Double gloves: - first: long sleeved gloves - second: nitrile gloves	
Eye Protection	<input type="radio"/> Goggles with lateral protection <input type="radio"/> Face shield	If not for single use, perform disinfection with ethanol base solution or 0.1% sodium hypochlorite
Gowns	<input type="radio"/> Single-use  <input type="radio"/> Waterproof <input type="radio"/> Long sleeved <input type="radio"/> Standard EN 14605:2009	Consider biological risk protection EN 14126:2004, if confirmed positive patient
Cap	<input type="radio"/> Single-use	Consider hood cap, if confirmed positive patient
Shoe cover	<input type="radio"/> Single-use	
Respiratory	<input type="radio"/> FFP2/N95	Perform seal check before enter the endoscopy suite.
Protection	<input type="radio"/> Single-use	Consider FFP3, if confirmed positive patient

PPE, personal protective equipment.

- Floors and surfaces of the endoscopy suite must be disinfected after each procedure.
- Intermediate level disinfectants with proven activity against enveloped viruses include 0.1% sodium hypochlorite, 62–71% ethanol, 0.5% hydrogen peroxide and quaternary ammonium compounds.<sup>19,24,25</sup>

### Personal protective equipment

IP procedures are considered to be consistently subjected to the highest risk of exposure. In this setting, full precautions must be taken to cover all different possible types of transmission (contact, droplet and airborne).<sup>26</sup> Personnel involved in the reprocessing procedure must also wear protective equipment consisting of eye protection, respiratory mask FP2, long sleeved gown and double gloves.<sup>27,28</sup> The recommendations for the use of PPE are shown in [Table 3](#).

### Specimen transportation

Samples from the upper and lower respiratory tract, including pleural effusion, are deemed to be the most potentially infectious. Consequently, they should be handled as Category 3 pathogen and double-bagged (first the specimen must be bagged in the patient’s room and then taken out of the room and placed in a separate pre-labeled specimen bag). All specimens must be manually delivered.<sup>28,29</sup>

### Safety rules for staff and patients

It is also important to define proper new rules for both HCW and patients circulating in the IP unit, as listed below:

### Health professionals

- The IP Unit should reduce and prioritise the allocation of human resources according to the outbreak evolution and hospital needs. The minimum number of staff required to ensure a correct operation must be clearly defined.
- It is essential that all personnel follow, train and maintain competency in effective hand hygiene and every aspect of PPE (theoretical, training and simulation sessions) so that everyone is familiar with their role.
- All interactions with patients, including informed consent, should be done with appropriate PPE and frequent hand washing. The staff should not reduce the level of awareness and protection, and the idea that patients with suspected COVID-19 should be handled in the same manner as confirmed cases must be reinforced.
- A core team that includes only essential HCW should perform the procedure on SARS-CoV-2 positive patients. The most experienced staff should be responsible for the exam to reduce time and deal effectively with possible complications. Other healthcare personnel, such as residents, medical students and visitors should not be inside the unit and the examination room before, during or after the procedure.
- Of note, the scheduled exams must be done during normal working hours (avoiding an emergency basis or setting) and in an appropriate, designated room that fulfils all the standards required for care.

### Patients and other personnel

- Respiratory and contact isolation should be standard and mandatory for all patients. Outpatients and inpatients should always enter the IP Unit with a suitable face mask

and keep it on at all times (until the beginning and after the end of the procedure) to minimize the risk of transmission. No unnecessary personal items should be brought into the IP unit.

- Family members and caregivers should not stay in the IP waiting rooms. In case of children or patients in need of support, the Unit can allow a single relative to enter the preparation area to provide aid.
- The entry into the Unit of suppliers and medical devices sales representatives must be restricted.

## Prioritization of procedures

Scheduled elective procedures should be reviewed and cancelled if potentially delayable, until local control of the outbreak is achieved. After flattening the infectious curve, many elective IP procedures will have to be performed, as they are essential to provide a definitive diagnosis and effective treatment. At this time, it is advisable to evaluate the delayed requests and to optimize the procedure planning based on clinical needs and operational capability.

A suggested rational approach for stratification of procedures is provided in Table 1, but we recognize that, in certain cases, the indication may not be straightforward, and the risk-benefit must be weighted on an individual basis by the IP team.<sup>31</sup> Although rescheduling certain procedures is obvious in other cases it may not be desirable or ethical. It is important to note that these indications may change according to local epidemiological conditions and the response capabilities of the healthcare system. Several societies have recommended different levels of procedure stratification.<sup>26,30,31</sup> Briefly, what is recommended is a step-wise reopening of elective IP procedures according to the national and local COVID-19 outbreak situation, depending on the number of new confirmed cases, hospital admitted cases (ward and ICU), availability of equipment and healthcare staff, time elapse and number of postponed IP cases. Some authors<sup>34</sup> have proposed a summary of the elective endoscopic procedure by phases, as shown in Table 2. Anyway, it should be noted that the evolving procedural criteria should always be communicated to other physicians who refer patients for invasive respiratory procedures and to the hospital administration.

## Recommendations for bronchoscopy

### Bronchoscopy under spontaneous ventilation

The following recommendations are expert opinion-based and should be adapted to local regulations and guidelines. In an optimal scenario, it is safer to perform elective bronchoscopy under general anesthesia and orotracheal intubation, clinical conditions permitting. If this is not possible, bronchoscopy can be performed under spontaneous ventilation. Some recommendations are listed below:

- Operator should be standing behind the patient's head to reduce direct exposure. Oxygen supplementation should be done without the use of humidification, either through

a nasal cannula or preferably with an oxygen mask with an entrance to the bronchoscope (Fig. 2A).

- For flexible bronchoscopy, a transnasal approach should be preferred, and a surgical mask should be placed over the patient's mouth to minimize droplet emission (Fig. 2B).
- In hypoxemic patients, bronchoscopy can be performed under NIV, using a closed circuit ventilation (double circuit with viral filters in both arms) and non-ventilated masks with a dedicated bronchoscope entrance (Fig. 2C). High performance NIV ventilators with FiO<sub>2</sub> regulation are preferable. From the end of the procedure, NIV should be continued for 1–2 h, titrating the FiO<sub>2</sub> to obtain an SpO<sub>2</sub> of around 94–95%.
- Bronchoscopy under nasal high-flow oxygen therapy is not recommended and thus should be avoided.
- Nebulized medications should be avoided before or after the procedure.
- Proper sedation should be used to minimize cough reflex and to increase patient cooperation.
- An oral aspiration cannula should be available during the procedure (Fig. 2B).
- A transparent protective box may enhance safety by containing dispersal of droplet particles (Fig. 2D). The box is placed over the patient's head prior to bronchoscopy, with the anesthesia equipment already in place. The bronchoscope is inserted through the covered opening behind the patient (Fig. 2E).

### Bronchoscopy in the intubated patient

The following recommendations are directed for patients under mechanical ventilation in an ICU setting due to respiratory failure. As reported, 5% of COVID-19 patients can develop respiratory failure and will need ventilatory support<sup>32</sup>; moreover, associated bacterial, viral and fungal co-infection should not be neglected.<sup>33</sup> In critically ill patients under invasive ventilation, ventilator-associated pneumonia occurs in up to 30% and lobar collapse is frequent and multifactorial.<sup>34</sup> The same adaptations apply to elective procedures under general anesthesia, performed in the Bronchoscopy Unit or Operating Theatre.

- A cuffed endotracheal tube is preferred over supraglottic devices, such as a laryngeal mask; cuff pressure should be maintained between 25–30 cmH<sub>2</sub>O.<sup>35</sup>
- General anesthesia with muscle relaxant is recommended to reduce the aerosol production.
- FiO<sub>2</sub> should be adjusted to 100%.
- Volume control, pressure-limited mode is preferable and PEEP should be kept at the same level during the procedure. Adjustments can be made dynamically, with a prior assessment of the anticipated risks (e.g., lung derecruitment and desaturation, arrhythmias, pneumothorax).
- To avoid aerosol dispersion, a simple and appropriate maneuver consists of clamping the ventilation circuit just before introduction of bronchoscope, repeating the same step just before withdrawal.
- Bronchoscope removal and reinsertion should be avoided during the procedure.
- In hypoxemic patients, if bronchoalveolar lavage is needed for diagnostic purposes, the volume used should be



**Figure 2** Strategies to minimize droplets dispersal during bronchoscopy. A. The bronchoscope may be introduced through an opening made at the oxygen mask, in this case with an additional plastic sheet covering the patient's head. B. Transnasal approach, with oxygen supplementation through nasal cannula and a surgical mask placed over the patient's mouth and the oral aspiration canulla. C. Bronchoscopy can be performed under ventilatory support, using a closed circuit ventilation and non-ventilated masks with a dedicated bronchoscope entrance. D. Transparent protective box may contain droplet particles inside. E. Protective box placed over the patient's head during endobronchial ultrasound. F. Rigid bronchoscopy with rubber caps on the ports of the scope and a plastic covering.

477 reduced to a minimum. If a SARS-Cov-2 diagnosis is  
478 needed, a minimum of 2–3 mL of recovered lavage is  
479 enough.<sup>26</sup>

### 480 Rigid bronchoscopy

481 Rigid bronchoscopy is used for diagnostic and therapeutic  
482 purposes, in procedures where flexible bronchoscopy  
483 would be deemed difficult or even impossible, like obtain-  
484 ing larger samples of endobronchial lesions, foreign body  
485 removal, management of central airway obstruction (includ-  
486 ing ablative techniques, like electrocautery, argon plasma  
487 coagulation, laser, cryotherapy, among others, and place-  
488 ment of airway stents) or massive hemoptysis.<sup>36</sup>

489 There are different ventilation strategies used during  
490 rigid bronchoscopy, although manual jet ventilation and high  
491 frequency jet ventilation are much the most frequent.<sup>37</sup>  
492 Common to these two techniques is the fact that the prox-  
493 imal end of the bronchoscope is open to allow the passage

of instruments, thus ventilation is achieved providing 100%  
oxygen under high pressure (usually 50 psi) through an open  
system.<sup>39,40</sup> The use of these ventilation techniques means  
that aerosols are released into the room, making it a high-  
risk procedure.

In patients with suspected or confirmed COVID-19 diagnosis, rigid bronchoscopy should be avoided, except for urgent cases (Table 1). Clinical scenarios are mostly therapeutic, like acute foreign body aspiration, massive hemoptysis (when there is no place for embolization), severe symptomatic central airway obstruction (either benign or malignant) and migrated stents. In a clinically stable patient, upon suspicion of foreign body aspiration, one should consider non-contrast computerized tomography (CT) to confirm the presence of a foreign body before rigid bronchoscopy, to avoid unnecessary exams.<sup>38,39</sup>

In some centers, the rigid scope is used to perform other techniques, like Endobronchial Ultrasound-Transbronchial Needle Aspiration (EBUS-TBNA) or transbronchial cry-



513 obiopsy; this provides comfort to the operator and safety in  
514 case of major bleeding. The authors recommend performing  
515 these diagnostic procedures through cuffed endotracheal  
516 tube to minimize the risk of exposure. However, the operator  
517 must be ready to convert to rigid bronchoscopy, if necessary.

518 In a patient undergoing rigid bronchoscopy, it is recom-  
519 mended that:

- 520 - Rigid bronchoscopy should always be performed in a neg-  
521 ative pressure room.
- 522 - Controlled ventilation is preferred, with the rigid bron-  
523 choscope used like an endotracheal tube.
- 524 - Air leaks should be reduced using rubber caps on the  
525 ports of the rigid scope, as well as using a plastic cover-  
526 ing (Fig. 2F) or filling the mouth with gauze.<sup>40</sup> While this  
527 strategy is more appealing to minimize aerosol spread,  
528 the operator may find it challenging to handle instruments  
529 through the working channel.

### 530 Recommendations for pleural techniques

531 Pleural effusion does not appear to be a prominent feature  
532 of COVID-19. It occurs in 5.3–5.8% of patients, according to  
533 two recent meta-analyses.<sup>41,42</sup> There have been occasional  
534 reports of bilateral effusion that resolved spontaneously.<sup>43</sup>  
535 As bacterial superinfection is common in severe patients,  
536 they can also develop complicated effusions or empyema,  
537 requiring targeted treatment. There have been a few  
538 anecdotal reports of spontaneous pneumothorax and pneu-  
539 momediastinum in severe COVID-19 pneumonia, requiring  
540 drainage.<sup>44,45</sup> This may be more frequent in critical patients  
541 on invasive ventilation, which can lead to bronchopleural  
542 fistulae.<sup>46</sup> It is, therefore, plausible that pleural drainage  
543 may be necessary in some COVID-19 patients, either in the  
544 ICU or in the ward, and indications for drainage do not  
545 differ from the standard clinical guidelines. However, as  
546 with any invasive procedure in confirmed COVID-19 patients,  
547 all precautions regarding the full use of protective equip-  
548 ment should be taken. The procedure must be performed  
549 by trained and dedicated staff to reduce its duration and  
550 to minimize the risk of complications. In other situations,  
551 the use of ultrasound may be very helpful with COVID-  
552 19 patients.<sup>47</sup> Besides its wide availability, safety and low  
553 cost, it is easy to use at the bedside and allows medical  
554 staff to detect small pleural effusions and to guide pleural  
555 fluid collection and drainage, if needed. On the other hand,  
556 even patients without suspected COVID-19 can have assymp-  
557 tomatic infection; so, any procedure should be considered as  
558 a possible COVID-19 case and precautions should be taken.  
559 Indeed, although some procedures may be postponed, in  
560 many situations they should not be deferred, especially in  
561 suspected or confirmed cancer patients. It is crucial that  
562 cancer patients do not experience delays in diagnostic or  
563 therapeutic procedures due to the present contingency.<sup>48</sup>

564 Few societies have published guidelines addressing pleu-  
565 ral procedures during COVID-19 pandemic. The British  
566 Thoracic Society has issued guidance on pleural ser-  
567 vices provision,<sup>49</sup> mainly to minimize hospital visits and  
568 admissions and to ensure both patient and staff safety.  
569 Nonetheless, although we recognize lack of published evi-

dence supporting these recommendations, this document  
will adopt some of them.

570 First of all, despite pleural procedures not being listed  
571 as Aerosol Generating Procedure (AGP) in the CDC updated  
572 recommendations,<sup>50</sup> they should be considered so and Level  
573 2 PPE should be worn, as described above. Other societies  
574 have considered potential AGP as any procedure “*likely  
575 to induce coughing, that should be performed cautiously  
576 and avoided if possible*”.<sup>51</sup> This is particularly relevant to  
577 open procedures, such as thoracoscopy and indwelling pleu-  
578 ral catheter insertion, and in case of pneumothorax with  
579 suspected bronchial-pleural fistula. Thus, we recommend  
580 taking the following precautions:  
581  
582

### 583 Pleural effusion

- 584 - On suspicion of malignant effusion, diagnostic pleural fluid  
585 aspiration should be performed, especially if the patient  
586 is a candidate for systemic therapy or if the effusion is  
587 symptomatic (Table 1).
- 588 - Consider placement of indwelling pleural catheters for  
589 recurrent malignant pleural effusions, avoiding repeated  
590 hospital drainage or admission.
- 591 - On suspicion of pleural infection, pleural fluid aspiration  
592 and analysis should be performed.
- 593 - In case of benign pleural effusions, the risk-benefit should  
594 be discussed with the attending physician; procedures can  
595 be postponed if the patient is not symptomatic.
- 596 - Thoracoscopy is not recommended as a principle; how-  
597 ever, in the absence of alternative options, its risk-benefit  
598 must be assessed.

### 599 Pneumothorax

- 600 - Spontaneous primary pneumothorax can be managed in  
601 outpatient care if the risk assessment allows and if there  
602 is local team experience; needle aspiration and discharge  
603 should be considered if the patient is minimally symp-  
604 tomatic.
- 605 - When pleural drainage placement is necessary, if there is  
606 local expertise and the patient is at low risk, the use of  
607 pleural vent systems should be considered, thus allowing  
608 the patient to be managed at home.

### 609 Chest drainage placement and care

- 610 - Extra care must be taken when placing the chest tubes, in  
611 order to avoid open communication with the pleural space  
612 and the potential emission of droplets and aerosols.
- 613 - When chest tubes are placed in ventilated patients, con-  
614 sideration should be given to clamping the ventilator  
615 circuit before assessing the pleural cavity, so that positive  
616 pressure spreading of pleural air or fluid can be prevented.
- 617 - Whenever possible, the use of non-wired pleural drainage  
618 should be considered; it can be connected to the drainage  
619 system before insertion into the pleural cavity (closed cir-  
620 cuit).
- 621 - When pleuroscopy is required, the use of one way valve  
622 trocars should be preferred to assess the pleural cavity  
623 and properly seal the entrance port of the pleuroscope.

Table 4 IP Unit checklist during COVID-19 outbreak.

PRE-PROCEDURE PLANNING	PROCEDURE EXECUTION	POST-PROCEDURE
<p><b>Adapt the IP unit to include:</b></p> <ul style="list-style-type: none"> <li>✓ Preprocedural area</li> <li>✓ Procedural room with negative pressure or adequate ventilation<sup>1</sup></li> <li>✓ Postprocedural area for decontamination, reprocessing and removal of PPE</li> </ul> <p><b>Revise prioritization of all procedures</b> (table x and x)</p> <p><b>Perform telephone pre-screening check-list</b> (upon schedule and 24-48h before the exam)<sup>2</sup></p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Symptoms</li> <li><input type="checkbox"/> Contact history</li> <li><input type="checkbox"/> Occupational exposure</li> </ul> <p><b>Test for SARS-CoV-2 in the 24-48h preceding the exam</b></p> <p><b>Proceed according to priority / test result / risk probability</b> (Flowchart 1, Table 1)</p>	<p><b>Gown adequate PPE</b> (Table 3) and perform procedure at adequate endoscopy suite<sup>1</sup></p> <p><b>Minimize direct exposure</b></p> <ul style="list-style-type: none"> <li>✓ Stand behind the patient's head</li> <li>✓ Place surgical mask over the patient's mouth</li> <li>✓ Adapt oral aspiration cannula</li> <li>✓ Avoid humidified O2 and nebulized drugs</li> <li>✓ Use appropriate sedation</li> </ul> <p><b>In ventilated patients:</b></p> <ul style="list-style-type: none"> <li>✓ Prefer cuffed endotracheal tubes over supraglottic devices</li> <li>✓ Clamp the ventilation circuit when removing or inserting the bronchoscope and immediately before assessing the pleural cavity when placing pleural drainages</li> </ul> <p><b>Avoid rigid bronchoscopy</b> (except if indicated for urgent purposes)</p> <p><b>Avoid thoracoscopy</b> (except if lack of alternative and after adequate risk/benefit stratification)</p> <p><b>Reduce teams and time of the procedure to minimal necessary</b></p>	<p><b>Collect respiratory samples</b> (including pleural fluid) in closed circuits and double-bag for transport</p> <p><b>If available, use disposable bronchoscopes for confirmed positive / highly suspicious COVID-19 patients</b></p> <p><b>Reprocessing of bronchoscopes must be considered an AGP</b></p> <ul style="list-style-type: none"> <li>✓ Gown adequate PPE (Table 3)</li> <li>✓ Perform procedure at adequate unit<sup>1</sup></li> </ul> <p><b>Disinfect floor and surfaces of the endoscopy suite after each procedure</b></p> <p><b>Allow adequate time (&gt;30min) to elapse before next procedure</b></p> <p><b>Perform sequential removal of PPE and adequate hand hygiene</b></p>

1. The preferred system is a negative pressure room with at least 12 air changes per hour with controlled of airflow direction (single pass or recirculation systems with HEPA filtration). Alternatively, natural ventilation with an airflow of at least 160 L/s is an option.  
2. Pre-screening checklist should include questions regarding (1) recent symptoms suggestive of COVID-19 (namely fever, cough, shortness of breath/difficulty breathing, chills, muscle pain, headache, sore throat, loss of taste or smell), (2) contact with suspicious or confirmed SARS-CoV-2 cases, (3) occupational exposure, and (4) recent travel history to a high-incidence region  
AGP: Aerosol generating procedures. PPE: personal protective equipment. IP: Interventional Pulmonology.

624 - In case of prolonged air-leaks, the use of wall suction  
625 should be weighted to create a closed system.

626 **Concluding remarks**

627 As with other societal consensus papers, this document  
628 was developed by a restricted panel of experts from  
629 the Portuguese Society of Pulmonology; individual clinical  
630 judgment and local resources may lead to alternative  
631 perspectives. This guidance was based on the current knowl-  
632 edge of COVID-19, but, as new data appears, this statement  
633 should be revised in the future to accommodate updated  
634 recommendations. At present, one of the controversial  
635 assumptions is that, every patient, even if asymptomatic,  
636 should be assumed as potentially infected with SARS-CoV-  
637 2. Therefore, it is mandatory that contact precautions and  
638 proper training on donning and doffing of PPE be provided  
639 to all HCWs involved in IP. Another key element is to plan  
640 in advance and keep each IP Unit well-organized (Table 4).  
641 Although the reduction in the number of elective procedures  
642 represents one of the central strategies to improve safety,  
643 it is crucial that patients do not suffer unnecessary delays in  
644 diagnostic or therapeutic procedures due to the current con-  
645 tingency. Taken together, our ultimate intention is to bring  
646 full attention to this and future outbreaks or other emerging  
647 medical situations.

648 **Conflicts of interest**

649  The authors have no conflicts of interest to declare.

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