



Early View

Review

ERS International Congress 2020 virtual: highlights from the Respiratory Intensive Care Assembly

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**ERS International Congress 2020 virtual: highlights from the Respiratory Intensive Care
Assembly**

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Abstract:

During the virtual European Respiratory Society (ERS) Congress 2020, early career members summarised the sessions organised by the Respiratory Intensive Care Assembly. The topics covered included diagnostic strategies in patients admitted to the Intensive Care unit (ICU) with acute respiratory failure, with a focus on patients with interstitial lung disease and for obvious reasons, SARS-CoV2 infection. These sessions are summarised in this paper, with take-home messages highlighted.

A. State-of-the-art session: respiratory critical care

1. The role of chest computed tomography scanning in the differential diagnosis of acute respiratory failure in ventilated patients. M. Prokop (Nijmegen, Netherlands)

According to M. Prokop, chest computed tomography (CT) scans can be used as a 'problem solver' to identify causes of acute respiratory failure (ARF) when a diagnosis cannot be established from the initial chest X-ray or ultrasound. M. Prokop was supportive of the use of inspiratory chest CT scans with a narrow slice width of 0.5 to 1mm, multiplane reconstructions and intravenous contrast injection; intravenous contrast differentiates pleural disease from intrapulmonary disease e.g. atelectasis from consolidation. He also suggested the use of expiratory chest CT scans in cases where air trapping is suspected. The speaker reminded us that except in low-dose CT, all contemporary chest CT scan now create high-resolution images and use radiation doses lower than natural background radiation (except for larger patients).

Different clinical situations were then described by M Prokop: on a chest X-ray, the absence of air bronchogram does not indicate extrapulmonary disease, and the volume of a pleural effusion must be greater than 200 ml to be visualized. In pleural effusions, ultrasound predicts drainage success better than CT [1, 2]. However, chest scans can differentiate empyema (obtuse angle with the pleura) from subpleural abscesses (acute angle); broncho-pleural fistula is a complication of pneumonia and may be suspected if air is observed within areas of in the consolidation. Chest CT scans are also the diagnostic gold standard for pulmonary embolism, and can be useful in estimating the size and extent of pneumothoraces or barotrauma, including in acute respiratory distress syndrome (ARDS), and in visualising interstitial and mediastinal emphysema..

However, chest CT scans are of modest utility in discriminating between other clinical conditions e.g. between edema, pneumonia, ARDS and organizing pneumonia, and in such cases additional clinical information is required. Ventral consolidations are however more typical for pneumonia, and symmetrical or moving consolidations are more suggestive of edema.

Take-home messages

- Chest CT scans are excellent in distinguishing atelectasis from other causes of consolidation, distinguishing consolidations, pleural from parenchymal disease, and in the diagnosis of pulmonary embolism and infarct
- Chest CT scans are of modest utility in identifying the cause of consolidation (e.g. organizing pneumonia, pulmonary hemorrhage, ARDS, edema)
- Chest CT scans are less useful in the differential diagnosis of pneumonia and predicting treatment outcomes of complex pleural effusions.

2. Role of broncho-alveolar lavage and lung biopsy in the diagnosis of acute respiratory failure in intensive care unit patients. V. Poletti (Forlì, Italy)

According to V. Poletti the most common causes of acute respiratory failure are ARDS and infections, but the two entities can overlap as infections can cause ARDS. There are histopathological differences between ARDS and others entities that may mimic it, such as idiopathic acute interstitial pneumonia, acute exacerbation of interstitial lung disease, organizing pneumonia, anti-GBM antibody disease, granulomatosis with polyangiitis, diffuse alveolar haemorrhage, acute hypersensitivity pneumonitis and acute hypereosinophilic pneumonia [3].

In patients with hypoxic respiratory failure, a bronchoscopy with broncho-alveolar lavage (BAL) can be considered only after clinical, biological, and radiological assessments have been performed. The most frequent side effects are fever and hypoxia, especially with large volume instillations. Rarely complications occurs such as hypoxemia, hypotension, bradycardia, haemoptysis, pneumothorax, bronchospasm, bleeding, takotsubo cardiomyopathy and acute exacerbations of interstitial lung disease (ILD) [4].

Diffuse alveolar damage (DAD) is the histological term used to describe specific changes that occur during acute lung injury. The pathological alterations observed in DAD vary depending on the interval between the onset of symptoms and the time of cyto-histological lung sampling. The early exudative stage of DAD is characterized by oedema and hyaline membrane formation. This progresses to a proliferative stage in which there is hyperplasia of type II pneumocytes, desquamation of alveolar lining cells, and thickening of the interstitium with fibroblast proliferation. Fibrosis may then occur with architectural destruction and honeycombing.

The cytopathological profile for DAD consists of more than 85% neutrophils, atypical type II pneumocytes and amorphous extracellular material (hyaline membranes). Cytological profiles that differ from this neutrophilic-predominant pattern should suggest ARDS mimics, for example diffuse alveolar haemorrhage (blood-stained BAL fluid, hemosiderin loaded macrophages) or disseminated malignancy [5]. Microbiological testing may identify a treatable cause of ARDS and has a high diagnostic yield in the immunocompromised patient [6].

Lung biopsy should be considered when BAL does not lead to a diagnosis. The decision to perform a lung biopsy should be made based on lung compliance and oxygenation, and consider, whether specific procedures and therapies are available, and the anticipated time to biopsy. DAD is present in the majority of patients with **acute respiratory failure secondary to** nonresolving ARDS and the frequency of DAD does not differ across the three stages of ARDS [7]. Although some studies show that open lung biopsy (OLB) provides a specific diagnosis in up to 84% of patients and leads to a change in management in up to 73% of cases, the risk of complications is unclear with only two

studies reporting complication rates: these varied from 0 to 56% [8, 9] and the wide range may be explained by differences in patient characteristics and definitions of complications. Studies investigating ARF studies did not compare differences in mortality between those who had an OLB from those who had not. Alternatives to OLB are transbronchial (cryo)biopsy. Limited data suggest that they have a high diagnostic yield with fewer complications than OLB [10] [11].

Take-home messages

- BAL can identify a specific diagnosis in a minority of cases but the greatest yield is in alveolar filling processes. It is a relatively low risk technique in acutely ill patients
- When no diagnosis is made after BAL, lung biopsy may be indicated but it is mandatory to discuss this on a case-by-case basis, taking into account the risk-benefit ratio
- Limited data suggest fewer complications in transbronchial cryobiopsy compared to open lung biopsy

3. The future of pulmonary diagnostics in intensive care unit patients, L. Bos (Amsterdam, Netherlands)

L. Bos first presented the diagnostic challenges in intensive care *i.e.*, identifying the cause of respiratory failure, understanding why a patient's lung injury is not resolving, and knowing if the patient is developing pneumonia.

Lung ultrasound (LUS), currently used as a dichotomous diagnostic test, should be used more quantitatively, for example to describe the size of pleural effusions or the amount of pulmonary oedema. Quantification of B-lines, either by manual scoring or by a computerized pre-processing algorithm, correlates very well with measurements of extravascular lung water [12]. The effect of an intervention can also be monitored by dynamic LUS. LUS Scoring for lung re-aeration by chest CT strongly correlates with scoring by LUS, suggesting that the latter could be used as a bedside tool to monitor response to changes of ventilatory settings [13]. Another promising technique is the use of contrast-enhanced LUS to examine peripheral consolidations: hypoechoic areas suggest the absence of perfusion and the possibility of infarction [14].

Confocal lung endomicroscopy (CLE) is an imaging technique using a laser-based bronchoscopic intervention that assesses the alveolar morphology *in situ* by detection of elastin in the extracellular matrix. As it is difficult to differentiate between oedema, inflammation and underlying fibrosis on CT, CLE can help by showing distortion of normal architecture, which is particularly prominent in fibrosis [15].

Another promising tool is biological assessment. Breath contains hundreds of volatile metabolites that can be reflective of pulmonary infection, pulmonary inflammation, and/or oxidative stress

response. A rapid bedside breath analysis by gas chromatography and mass spectrometry shows potential as a biomarker in the diagnosis of ARDS and the exclusion of bacterial causes for nosocomial pneumonia [16] [17].

Several “omics” techniques have been used to better phenotype patients and to provide rapid results. The broad use of molecular techniques could improve diagnostic yield. Real-time metagenomic sequencing of BAL fluid seems to be a more sensitive and faster way to identify the causative pathogen in pneumonia compared to standard cultures [18] [19]. It can also show the diversity of the lung microbiome, which predicts intensive care outcomes [20] [21]. The use of transcriptomic profiles has shown that inflammatory markers in the blood are not very well correlated with markers in the lung. This is an additional argument to focus our diagnostic sampling on the lung [22]. These techniques can also give us more data about the host response to inflammation.

Take-home messages

- Lung ultrasound have the advantage of instant clinical feedback and are easily repeatable
- Confocal lung endoscopy detects in vivo pathology and is easily repeatable. However, it only provides a focal assessment and requires bronchoscopy
- Breath analysis provides rapid results but lacks specificity with sampling errors
- Comprehensive “omics” techniques are likely to be more accurate in providing a comprehensive snapshot of disease but are time/labour sensitive and require sampling of the lower airway

B. Hot topics. Interstitial lung disease patients in the intensive care unit

1. Diagnostic procedures in ICU: bronchoscopy and bronchoalveolar lavage or tissue? C. Ravaglia (Forlì, Italy)

Hypoxic respiratory failure is a frequent cause of admission to ICU and contributes significantly to morbidity and mortality. Diagnosis of lung infiltrates of unclear aetiology is a major challenge in ICU [23] [24]. BAL may be performed when it is not possible to retain a definite diagnosis or identify a specific cause after taking a complete history, physical examination, global clinical assessment, chest CT scan, laboratory tests (including immunologic and serologic testing), echocardiography and/or pulmonary artery catheterization). BAL is a relatively low risk technique, which is well tolerated, easily performed with flexible bronchoscopy, and usually safely performed in acutely ill patients [25].

Rapid on-site cytologic examination of BAL fluid may have a role in determining specific diagnoses [26]. In patients with long-standing ILDs with new onset lung infiltrates admitted to ICU with acute respiratory failure, BAL may be diagnostic only in a minority of cases (13%), and hospital mortality and therapeutic management appear to be unchanged regardless of BAL findings [27]. The greatest yield for BAL is obtained in patients with pathological alveolar filling processes (such as some

opportunistic infections, peripheral malignancies, alveolar proteinosis, alveolar haemorrhage, lipoid pneumonia, etc.)

Cytologic analysis on BAL fluid is most useful for lung malignancies (particularly for adenocarcinoma or tumours with lymphangitic growth patterns) and haematologic malignancies. Morphological analysis may be implemented by immunocytochemical or molecular tests to identify the cell lineage and the presence of monoclonality [5]. Total white cell count, differential cell counts, lymphocyte subpopulations and immunohistochemistry are mostly used to identify ILDs and lymphomatous or leukemic infiltrates.

As there are no established guidelines, the decision to perform a lung biopsy should be discussed during multidisciplinary meetings and answer the following questions: 1) Could the biopsy provide a specific diagnosis which is of benefit to the patient? 2) Could the patient tolerate further insult to lung compliance and oxygenation? The timing and procedure used to perform the biopsy should also be discussed [28].

Open lung biopsy is rarely performed in ICU patients because of poor physiological conditions of most patients and because DAD has traditionally been considered the pathological signature of ARDS thus making the biopsy appear less necessary [29]. However, in reality the sensitivity and specificity of DAD for ARDS is low [7]. The most frequent therapeutic change implemented after biopsy is the decision to initiate, continue, adjust the dose or stop the administration of steroids, but management of antimicrobials and end-of-life decision-making are also important [8, 9].

There is a need for a prospective multicentred, randomised controlled trial evaluating the survival benefit of lung biopsy for patients with ILF, acute respiratory failure and lung infiltrates.

Take-home messages

- Bronchoscopy is a relatively low-risk technique. It may have an important role in the ICU, particularly in the diagnosis of alveolar filling processes.
- When no definite diagnosis is obtained after BAL, lung biopsy may be indicated, however, there are no widely accepted guidelines and there is urgent need of a prospective randomized controlled trial evaluating the real-world benefit of lung biopsy for these patients
- Surgical lung biopsy may provide a specific diagnosis and lead to a change in management in the majority of patients

2. Radiological patterns in ICU: does it matter? N. Sverzellati (Parma (PR), Italy)

Patients admitted with acute respiratory failure should be assessed as to whether the condition could be related to undiagnosed interstitial lung disease (ILD). Assessment should take into account clinical risk factors for progression of interstitial lung abnormalities (ILA) e.g., cigarette smoking, inhalational exposure, medications (e.g., chemotherapy, immune checkpoint inhibitors), radiation therapy, and

thoracic surgery. When analysing the chest CT, physicians should assess patterns associated with progressive disease, i.e., the presence of non-fibrotic ILA with basal and peripheral predominance, and fibrotic ILA with basal and peripheral predominance with or without honeycombing.

Previously undiagnosed usual interstitial pneumonia (UIP) is identified in 50% of patients dying from acute interstitial pneumonia [30]. 59% of patients with acute respiratory failure complicating ILD do not have a pre-morbid diagnosis of ILD prior to ICU admission [31]. Signs of fibrosis on CT are associated with increased in-hospital mortality, 1-year mortality and failure to respond to high dose corticosteroids.

In acute respiratory failure, honeycombing is a better diagnostic marker than bronchiectasis for underlying UIP, because of difficulty discriminating irreversible traction bronchiectasis from free standing bronchiectasis during acute lung injury. Analysis of prior radiological examinations (even chest X-rays) is particularly helpful, if available.

Similar CT findings may be caused by different aetiologies. Interlobular septal thickening, tree-in-bud, cavitation, and features of organizing pneumonia are good diagnostic markers of organizing pneumonia, while ground glass opacification is less specific (Figure 1).

Extension and distribution of ground glass opacities are associated with poorer prognosis [32]. Asymmetric distribution of acute abnormalities are associated with better prognosis; 180-day mortality was lower in patients with asymmetrical distribution compared to those in which acute abnormalities were symmetrical [33].

Take home message:

- UIP is often not diagnosed before acute exacerbation
- Prior chest imaging e.g., CT scans are pivotal for interpretation of acute abnormalities

3. Selection of ILD patients for ICU and ECMO, C. Agerstrand (New York, United States of America)

The CESAR trial [34] recruited adults with severe ARF and randomized them to conventional care at their local hospital or referral and transfer to an ECMO centre. The authors met their primary outcome of an improvement in survival without severe disability at 6 months in the ECMO-referred group: 63% vs. 47% (RR 0.69 [CI 0.05-0.97] p=0.03). However, not all patients in the ECMO group actually received ECMO, as some improved with conventional management in the tertiary care centre to which they were referred., In addition, not all patients randomized to the control group received lung protective ventilation strategies that are known to improve outcomes in ARDS.

The EOLIA trial [35] sought to address the limitations of the pragmatically designed CESAR trial, and recruited 249 adults with severe ARDS and randomized them to a protocolized approach to mechanical ventilation or ECMO with an lung ultra protective ventilatory approach. Using ECMO in conjunction with ultra lung protective ventilation, 60-day mortality was not significantly lower with

ECMO (35%) than with a strategy of conventional mechanical ventilation (46%) (relative risk= 0.76 [95% confident interval: 0.55 to 1.04], p=0.09) however, some patients in the conventional mechanical ventilation also received ECMO as rescue therapy.

A retrospective analysis included 40 patients with interstitial lung disease (ILD) referred to ICU for acute respiratory failure treated with or without ECMO. Data suggest that ECMO is a lifesaving option for patients with interstitial lung disease and acute respiratory failure provided that they are candidates for lung transplantation. Indeed, ECMO was not able to improve the poor prognosis in patients that did not qualify for lung transplantation [36].

Various ECMO configurations can provide varying levels of support by making adjustments to oxygen delivery to the brain/coronary circulation and the level of hemodynamic support [37]. Venovenous ECMO can be used in the absence of concomitant pulmonary hypertension; venoarterial ECMO is better for patients with pulmonary hypertension.

In order to decide whether or not to start ECMO, several questions need to be answered such as the rehabilitation potential of the patient pre- and post-ECMO, as well as factors that affect the time to potential lung transplantation (e.g. extremes of height and weight, or rare blood type that may indicate the wait will be longer). Upper body venoarterial ECMO via the subclavian or innominate artery cannulation is advised in patients on a bridge to transplant ECMO [38] as it allows ambulatory ECMO that supports physical reconditioning.

Take home messages:

- ECMO may be beneficial in ILD exacerbations as a bridge to transplantation
- Early referral to a transplant centre is essential for potentially eligible patients

4. Management of ILD patients in ICU. M. Kokosi (London, United Kingdom)

Patients with idiopathic pulmonary fibrosis (IPF) have a significantly worse prognosis at 90 days compared to patients with non-IPF idiopathic interstitial pneumonia (IIP) or connective tissue disease (CTD) related ILD (which have similar outcomes to each other) [39].

ILD patients admitted to the ICU can be divided in three main categories: 1) ILD with a clear autoimmune background or drug-induced ILD; 2) ILD with no autoimmune features; and 3) patients with signs of autoimmune dysregulation but no clear diagnosis. The most common ILDs seen in ICU are IIPs, CTD-ILD, vasculitis, drug-induced ILD and acute eosinophilic pneumonia. Patients with IPF and overt fibrotic disease have poor outcomes [40] [41].

Patients with ILD admitted to ICU can also be divided into three phenotypes: phenotype A - progressive with potential for full reversibility; phenotype B - progressive with potential for partial reversibility; phenotype C - rapidly progressive with unpredictable outcomes. Reversibility in an acute exacerbation of ILD is more likely with *de novo* ILD, features of organising pneumonia on CT, a shorter time from diagnosis, antibody positivity (apart from melanoma-differentiation-associated

protein 5 (MDA5)), increased inflammatory markers and response to previous treatment. Organising pneumonia (OP) with no radiological evidence of fibrosis could be treated with IV methylprednisolone monotherapy (phenotype A). In contrast, a patient with Jo1 positive inflammatory myopathy or OP with established fibrosis would need dual therapy with IV methylprednisolone and cyclophosphamide or rituximab (phenotype B). Patients with MDA5 myositis (phenotype C) would need triple therapy with IV methylprednisolone, cyclophosphamide and rituximab. In these cases, the addition of Tacrolimus, Cyclosporin, or even plasmapheresis could be considered

Take home points

- Prognosis of ILD in ICU is poor, but most case series consist of mixed ILD including IPF
- Careful selection of ILD patients for ICU admission is crucial with involvement of ILD specialists in multidisciplinary discussions.
- The evaluation of reversibility is the cornerstone in the management of these patients

C. Management COVID-19

These sessions explained the most up-to-date advice at the time of the ERS congress for the clinical management of patients with COVID-19, including respiratory support and pharmacological management. Some knowledge may have now evolved.

1. Respiratory support in the management of SARS-CoV2: pre-ventilation, ventilation and ECMO. P. Navalesi (Padova, Italy)

The SARS-CoV2 pandemic resulted in unprecedented pressure on intensive care resources and bed capacity. In this context, the use of non-invasive respiratory support outside ICU has been demonstrated to be safe and effective [42] [42] although it may contribute to viral aerosolization [42] that needs to be reduced (Figure 3). Successful treatment with high flow nasal cannula (HFNC), continuous positive airway pressure (CPAP) or non-invasive ventilation (NIV) was more frequent in younger, less hypoxic patients with fewer comorbidities [42] [42]. No significant differences in negative outcomes including mortality and endotracheal intubation (ETI) rate have been identified based on the non-invasive modality used [42]. However, disease severity and ceilings of care may independently affect modality selection and outcomes.

Once a patient is placed on a ventilator, the key challenge is to avoid complications. Mechanical ventilation (in and of itself) does not produce lung healing - it merely keeps patients alive until their own biological mechanisms are able to recover. The objective is to avoid further lung inflammation [43].

Where ETI is required, individualisation of ventilatory support with the aim of preventing further complications is advised [43]. Lung-protective ventilation using conservative tidal volumes and

driving pressures has been recommended [44]. Determining patients' optimal ventilatory pressures is complicated by heterogeneity in lung recruitability [45] and patient factors including comorbidities such as obesity [46] [47] (Figure 2). P. Navalesi suggested the following three stages: 1) the use of lung protective ventilation, 2) early prone positioning *i.e.* positioning: within 72h of ETI with a first cycle of 16 hours then 12 hours and 3) if there is a lack of improvement, to contact an ECMO referral centre. Further investigation is required to inform timing of weaning invasive ventilation and extubation.

In SARS-CoV2-induced ARDS, ventilation/perfusion (V/Q) mismatching is typically exacerbated by hypercoagulability and development of pulmonary microthrombosis [48] [49]. Prone positioning may increase oxygenation by reducing V/Q mismatching [50], but patient response is variable due to differences in disease phenotype and pathophysiology.

WHO guidelines recommend the consideration of extracorporeal membrane oxygenation (ECMO) in patients with refractory hypoxaemia despite lung protective ventilation. Outcomes of ECMO-rescued patients appear similar in SARS-CoV2 infection compared to other causes of ARDS [51]. It is also important to consider the practical and ethical implications for healthcare services, and the wider patient population [52].

Take home messages

- Use of non-invasive respiratory support to manage hypoxaemia in selected patients with SARS-CoV2 infection may prevent requirement for intubation
- Pathophysiological differences exist between patients with acute respiratory failure and ARDS due to SARS-CoV2 infection and it is essential to tailor supportive respiratory management to the individual

2. Drugs: anti-virals. J. Beigel (Baltimore, United States of America)

J. Beigel reported **acute respiratory failure secondary to acute respiratory failure secondary to** the preliminary data from two trials: 1) RECOVERY (Randomised Evaluation of COVid-19 thERapY) and 2) ACTT-1 (Adaptive Covid-19 Treatment Trial)

Remdesivir

Remdesivir was one of the first antiviral treatments assessed in SARS-CoV2 infection. Results from the ACTT-1 study were presented [53]. The study was a double-blind, randomized, placebo-controlled trial of remdesivir versus placebo for up to 10 days. Preliminary results came from 538 patients randomized to remdesivir and 521 to placebo. In patients with mild to moderate disease, time to recovery was 5 days for both remdesivir and placebo. In patients with severe disease, time to recovery was 12 days in the remdesivir group and 18 days in the placebo group. The odds ratio (OR) for improvement was higher in the remdesivir group compared to the placebo group (OR for improvement, 1.50; 95% CI, 1.18 to 1.91; P=0.001; 844 patients) at the 15th day. Mortality was not

significantly different (hazard ratio - 0.70 (95% CI, 0.47 to 1.04). The data for patients not receiving oxygen or receiving high-flow oxygen, as well as those receiving extracorporeal membrane oxygenation (ECMO) or mechanical ventilation should be interpreted with caution due to the low number of patients recruited and followed up.

Lopinavir-ritonavir

Lopinavir-ritonavir, usually used to treat HIV infection, has been repurposed as a treatment for SARS-CoV2 infection. In a randomised trial, no significant difference was identified in the 28-day mortality between 1596 subjects with lopinavir-ritonavir and 3376 patients in usual care (22.1% lopinavir-ritonavir vs. 21.3% usual care). No benefit in the risk of progression to mechanical ventilation or length of hospital stay was observed [54]. The small number of patients on invasive mechanical ventilation included does not allow clear conclusions to be made about the effect of lopinavir-ritonavir in mechanically ventilated patients.

Hydroxychloroquine

Hydroxychloroquine has been suggested as a treatment option based on its in vitro activity against SARS-CoV2. No significant difference in 28-day mortality was identified between 1542 patients randomized to hydroxychloroquine and 3132 patients randomized to usual care (25.7% hydroxychloroquine vs. 23.5% usual care) [55]. No supportive evidence of beneficial effects on hospital stay or other outcomes was observed. These conclusions have been confirmed by a large systematic review and meta-analysis [56] [.

Take home message

- Remdesevir may improve the clinical outcome in severe COVID-19 patients.
- The effect of Remdesevir as a stand-alone treatment has not been established
- Combination Lopinavir-ritonavir did not demonstrate clinical benefit
- Hydroxychloroquine did not demonstrate clinical benefit

2. Drugs: anti-inflammatories. R. Haynes (Oxford, United Kingdom)

R. Haynes presented data from the RECOVERY trial (Randomised Evaluation of COVID-19 therapy), a randomised controlled trial of medications (dexamethasone, lopinavir-ritonavir tocilizumab, azithromycin, hydroxychloroquine, convalescent plasma), enrolling more than 15% of all UK hospitalized patients with COVID-19. Data from the COVACTA trial were also discussed.

Dexamethasone:

The dexamethasone study arm was inspired by the long-standing contradictory outcomes of corticosteroid use in severe viral respiratory infections [57]. 2104 patients were randomised to dexamethasone 6 mg once per day for 10 days and 4321 patients to usual care. In the

dexamethasone group, the incidence of death was lower than in the usual care group among patients receiving invasive mechanical ventilation (29.3% vs. 41.4%; rate ratio, 0.64; 95% CI, 0.51 to 0.81) and among those receiving oxygen without invasive mechanical ventilation (23.3% vs. 26.2%; rate ratio, 0.82; 95% CI, 0.72 to 0.94) but not among those who were receiving no respiratory support at randomization (17.8% vs. 14.0%; rate ratio, 1.19; 95% CI, 0.92 to 1.55). Benefit was also clear in patients who were treated more than 7 days after symptom onset, when inflammatory lung damage is likely to be more common. Patients in the dexamethasone group had a shorter duration of hospital stay than the usual care group (12 days vs. 13 days) and a greater probability of being discharged alive within 28 days (rate ratio, 1.10; 95% CI, 1.03 to 1.17).

Tocilizumab

The COVACTA trial evaluated the efficacy, safety, pharmacodynamics, and pharmacokinetics of tocilizumab, an interleukin 6 (IL-6) receptor antagonist, in 450 hospitalized patients with severe COVID-19 pneumonia [58]. Despite the fact that tocilizumab-treated patients spent a week less in hospital compared those in the placebo arm, treatment did not reduce mortality. The drawbacks of the study included broad eligibility criteria, heterogeneity of the study population, the timing of drug delivery and the timing of outcome analysis, which all limit interpretation of the results. Tocilizumab was not effective in preventing intubation or death in 243 moderately ill hospitalized patients with COVID-19. Some benefit or harm cannot be ruled out as the confidence intervals for efficacy comparisons were wide [59]. However, data published after the conference showed a survival benefit in critically ill patients treated with tocilizumab [60].

Take home message

- Dexamethasone 6 mg once daily for up to 10 days reduces 28-day mortality in patients with COVID-19 who are receiving respiratory support *i.e.*, invasive mechanical ventilation or oxygen (if oxygen saturations on room air are 92-94%).
- Results of ongoing clinical trials are awaited to determine the clinical efficacy of tocilizumab in hospitalized severe COVID-19 patients

3. Other drugs and convalescent plasma. M. Patout (Paris, France)

M. Patout discussed three topics regarding COVID-19 pathology and therapeutic approaches: 1) convalescent plasma; 2) thrombotic events and anticoagulation; 3) bacterial co-infection and antibiotics

Convalescent plasma

Convalescent plasma to treat COVID-19 disease was granted authorization for emergency use by the FDA in August 2020 . M. Patout presented data from the largest clinical study of convalescent plasma therapy: plasma transfusion in severe COVID-19 disease led to clinical improvement in 9 patients (36%) by day 7 and 12 patients (48%) by day 14 and was not associated with any clinical complications [61]. These findings are consistent with several case studies

investigating the use of convalescent plasma in severe cases [62] [62]. Despite promising results among patients with severe and/or life-threatening COVID-19 disease, several clinical questions regarding the efficacy and applicability of plasma transfusion remain, including the timing of transfusion, the volume of plasma and adjustment for body mass index, the lowest efficient titre of donor plasma, the optimal number of transfusions, the effect in early disease, and the impact on mortality.

Thrombotic events and anticoagulation

There is overwhelming evidence that SARS-CoV2 infection is associated with coagulopathy and increased thrombotic risk. The first European autopsy study established deep venous thrombosis in 58% of patients and pulmonary embolism in 30% [63]. Klok *et al.* reported thrombotic complications in 31% of 84 ICU subjects: Venous thromboembolism was confirmed by ultrasonography in 27%, arterial thrombosis in 3.7% and pulmonary embolism in 81% [64]. The mechanisms underlying the increased prothrombotic risk in SARS-CoV2 infection are diverse, including direct activation of coagulation by SARS-CoV2, endothelial dysfunction, increased levels of von-Willebrand factor, systemic inflammation, Toll-like receptor activation, and tissue factor pathway activation [65, 66].

Anticoagulation appeared to be effective in a large cohort of 4,389 hospitalized COVID-19 patients [67]. Patients were categorized into three groups: 1) 1,530 (34.9%) without anticoagulation; 2) 1,959 (44.6%) prophylactic anticoagulation and 3) 900 (20.5%) with therapeutic anticoagulation [63]. Those with prophylactic or therapeutic anticoagulation had a lower risk of in-hospital mortality with respectively adjusted hazard ratio (aHR) -0.50 [95%CI: 0.45-0.57] and aHR-0.53 [95% CI: 0.45-0.62] respectively) in comparison to no anticoagulation. Prophylactic and therapeutic dosages) were associated with a lower risk of intubation compared to no anticoagulation (aHR - 0.72 [95%CI: 0.58-0.89] and (aHR-0.69 [95%CI: 0.51-0.94] respectively). Major bleeding was highest among patients with therapeutic anticoagulation (3.0%) and similar for patients on prophylactic (1.7%) or no anticoagulation (1.9%). The authors concluded that anticoagulation in patients hospitalized with SARS-CoV2 is associated with lower mortality and risk of intubation without a statistically significant difference between prophylactic and therapeutic dosage regimens. However, these data have not been confirmed by randomized controlled trials.

Bacterial co-infections and antibiotics

Patients with SARS-CoV2 infection are commonly started empirically on antibacterial therapy when first hospitalized. The Michigan study reported that empiric antibacterial therapy was prescribed in 56.6% of 1705 hospitalized SARS-CoV2 patients within 2 days of hospital admission. However, microbiologically confirmed community-onset bacterial co-infection was established in only 3.5% (59/1705) of these patients [68]. These results are similar to other studies in UK [69] and New York [70]. Moreover, the cytokine storm, rather than bacterial sepsis, may mislead the clinician in a subset of patients, and result in clinical deterioration. The risks of unnecessary antibacterial therapy and the low rate of confirmed early bacterial co-infection in patients with SARS-CoV2 infection suggest against routine antibacterial therapy.

Take home message

- The administration of convalescent plasma is a safe treatment option in severe COVID-19; its contribution as a stand-alone treatment remains elusive
- There is a low frequency of early (up to 48h) bacterial coinfection in hospitalized COVID-19 patients and no evidence of concomitant fungal infection
- Hospitalised COVID-19 patients are at increased thrombotic risk. Anticoagulation is reasonable and is associated with lower rates of death and intubation. There is still doubt as to whether prophylactic or therapeutic doses of anticoagulation should be administered to prevent thrombosis in hospitalised COVID-19 patients

D. Hot topics in intensive care: COVID-19: Lessons learned

This session explained the most up-to-date advice at the time of the ERS congress for the clinical management of patients with COVID-19, including respiratory support and pharmacological management. Some knowledge may have now evolved.

1. Clinical characteristics and outcome, A. Demoule (Paris, France)

A. Demoule began by describing demographic data. The population admitted to ICU with COVID-19 was younger compared to those usually admitted to ICU: this was because of the comparatively high mortality of older people, and because such patients were less frequently admitted to ICU due to greater disease severity [71].

Ethnicity data from the USA cohort [72] show that minority ethnic backgrounds were over represented in ICU admissions (30 % African-American and 30% Hispanic), likely due to socioeconomic and professional factors.

Regarding comorbidities, 75-85% of the ICU population was obese or overweight, 50-60% had hypertension and 17-25% had type 2 diabetes. COPD was present in only 4-8% of patients.

Age, male gender, quantity of organ failure, severity of hypoxemia, morbid obesity and the number of beds in ICU departments has had an important impact on the prognosis of SARS-CoV2 pneumonia [71].

Take home message

- Risk factors for death from severe COVID-19 include metabolic syndrome, age, and gender
- Lung compliance is not dramatically reduced in all patients

2. Respiratory support and viral transmission, S. Nava (Bologna, Italy)

S. Nava started by explaining that most patients with acute respiratory failure due to SARS-CoV2 were managed outside the ICU [73].

In hypoxaemia unresponsive to conventional oxygen therapy, continuous positive airway pressure (CPAP) may improve intubation-free survival compared to oxygen therapy alone [74]. Response is variable as demonstrated by an observational study of patients managed with helmet CPAP, in which 21.7% (n = 34) required invasive mechanical ventilation (IMV) and 22.9% (n = 36) died [75]. After adjustment for potential confounders, the failure rate of CPAP is not significantly different to that of other non-invasive respiratory support (NRS) modalities *i.e.*, in the rate of mortality, endotracheal intubation and length of hospital stay. Mortality rate increased with age and comorbidity class progression [42]. Improvements in oxygenation observed on proning non-intubated patients was maintained after resupination in half of patients. Further studies are warranted to ascertain the potential benefit of this technique in improving respiratory and global outcomes [76].

The second part of the presentation focused on viral transmission: 11.4% (n = 42) of healthcare workers in contact with aerosol generating procedures (AGPs) had positive SARS-CoV2 PCR, 3 of whom required hospital admission [42]. However, the infection of healthcare workers remains inconclusive; one study showed the prevalence of SARS-CoV2 antibodies was 13.4% in a cohort of 40 329 health care personnel, including those without direct patient contact such as administrative staff [77]. Of doctors, General Practitioners were predominantly affected, likely due to close proximity to infected patients less frequent use of personal and protective equipment (PPE), particularly in early stages of the pandemic [42]. Although devastating for colleagues and healthcare organisations, the pandemic has reconnected healthcare workers to the communities they serve, with recognition and praise given in response to their dedication and sacrifice.

Take home messages

- Response to CPAP in patients with SARS-CoV2 is variable and requires further investigation to improve patient selection
- Further investigation is required to determine SARS-CoV2 transmission from aerosol generating procedures to inform safeguarding of healthcare workers.

3. Management of severe SARS-CoV2 in resource-poor settings, A. Dondorp (Bangkok, Thailand)

Nine Asian countries have been involved in a large ICU network with South American and African countries to capture important epidemiological data on ICU mortality and comorbidity. Indeed, 84% of the world's population lives in low or middle-income countries and there is no accurate data on the impact of SARS-CoV2 these areas.

The quality of care often depends on the healthcare setting, and many poorer countries depend on a variety of donated ventilators which complicates management. A quarter of hospitals lack sufficient oxygen supply. Therefore, awake proning in patients with non-invasive ventilation (NIV) has been particularly useful in these settings.

The suggested minimum level of facilities and staffing for mechanical ventilation in low-resource settings is continuous pulse oximetry while on the ventilator, 24-hour presence of a member of staff with airway training, and a nurse-to-patient ratio of at least 3:1 [78].

Lung ultrasound is a useful tool for lung imaging in resource-poor settings [79] because it is affordable, can be easily taught, and can help in diagnosis when testing is not available and also in assessing 'non-focal' vs 'focal' lesions, or 'non-compliant' or 'non-recrutable' lesions.

One of the main challenges of the management of SARS-CoV2 infection in these settings was the fact that PPE was in short supply; in addition, there was a lack of negative pressure rooms, isolation rooms, and clean water.

The emphasis of the management of severe SARS-CoV2 infection in resource-poor settings should focus on the availability and quality of non-invasive respiratory support, the safety of staff and the use of lung ultrasound for lung imaging.

Take home message

- In resource-poor settings, severe SARS-CoV-2 infection should be preferentially managed using non-invasive respiratory support
- In resource-poor settings, safety of staff is challenging because of the lack of PPE and equipment

4. How to organize research during a pandemic, M. Gong (Bronx, United States of America)

M. Gong presented an algorithm of the organization of clinical research used in Montefiore, Italy, during the pandemic: each trial was attributed to a senior clinical investigator, a non-clinical research fellow who coordinated all studies, and 2 research groups; 1) experienced critical care research coordinators that screened and enrolled patients, collected collect samples and performed hospital-based study procedures and 2) volunteer research coordinators (medical students) who performed data entry and query resolution. All other clinical research was suspended.

Clinical research during a pandemic is possible but challenging. It is important to adapt infrastructures to pandemic standards, including expediting regulatory approvals. Because of the urgency of trials, it also requires national leadership. Pandemic research became as urgent as pandemic clinical care: human subject review and approval was centralized and expedited; regulatory paperwork were reviewed and signed in rapid fashion; research pharmacy and clinical pharmacy collaborated and

were able to implement trials within days; and alternatives to written consent were accepted e.g. electronic signatures, or in the USA, verbal consent and electronic signature

Take homes messages

- Clinical research during a pandemic is possible by adapting infrastructures to pandemic standards and expediting regulatory approvals
- It is important to take into account the tension some physicians feel between clinical care and clinical trials during a pandemic

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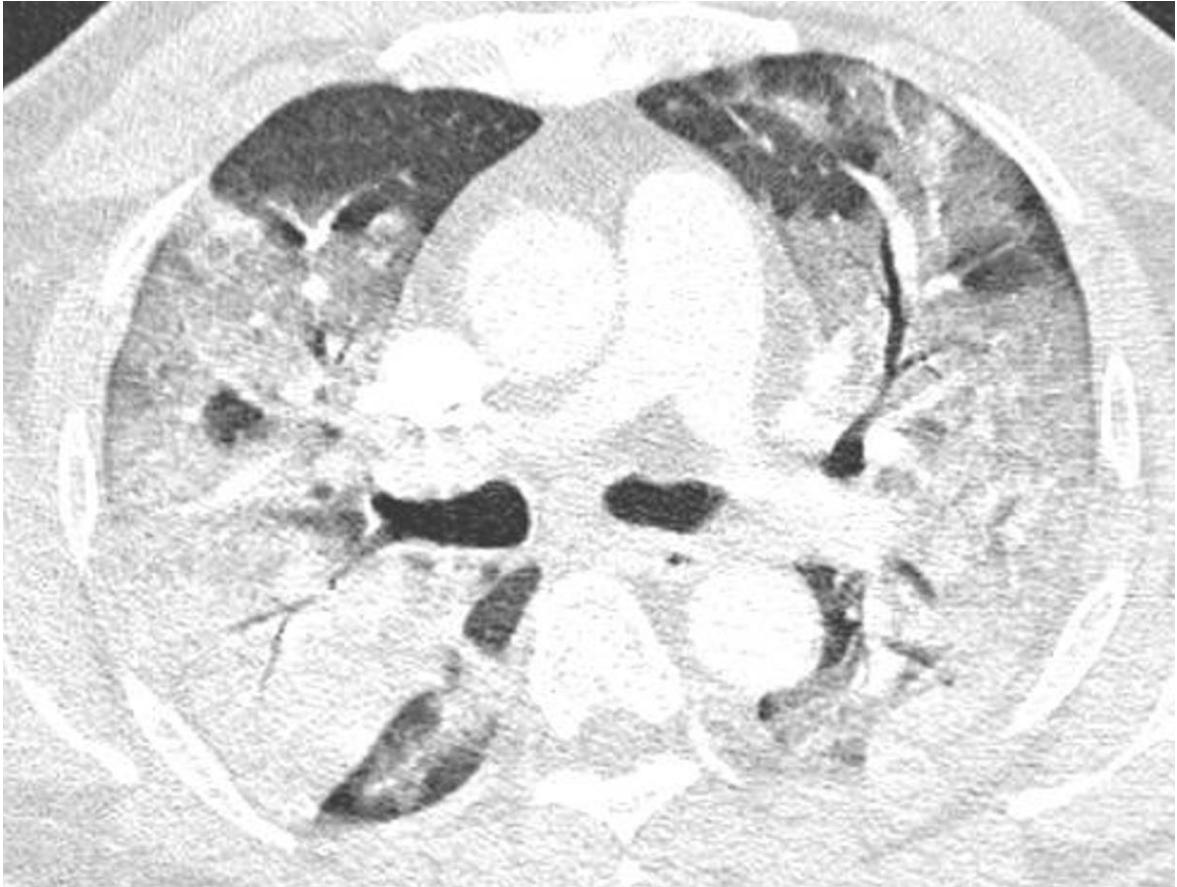


Figure 1: chest computed tomography in a patient admitted in ICU for acute respiratory failure secondary to eosinophilic acute pneumonia



Figure 2: chest computed tomography in a patient admitted in ICU for acute respiratory failure secondary to SARS-CoV-2 infection

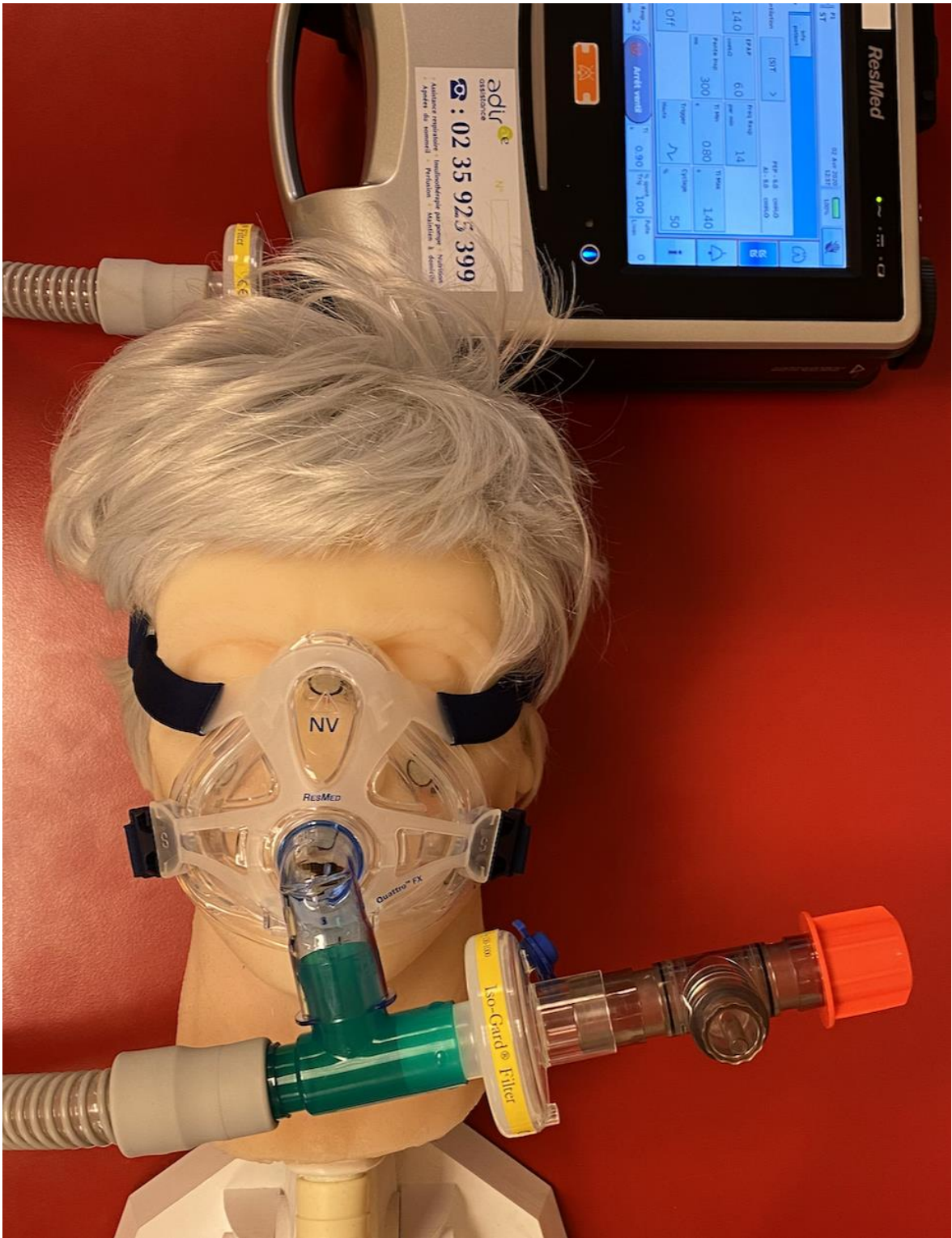


Figure 3: a non-invasive ventilation circuit used to minimize viral aerosolization (evaluated in [80])