

Respiratory physiology of COVID-19 and Influenza associated Acute Respiratory Distress Syndrome

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Research

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Abstract

Background: There is ongoing debate whether lung physiology of COVID-19 associated acute respiratory distress syndrome (ARDS) differs from ARDS of other origin.

Objective: The aim of this study was to analyze and compare how critically ill patients with COVID-19 and Influenza A or B were ventilated in our tertiary care center with or without extracorporeal membrane oxygenation (ECMO). We ask if acute lung failure due to COVID-19 requires different intensive care management compared to conventional ARDS.

Methods: 25 patients with COVID-19 associated ARDS were matched to a cohort of 25 Influenza patients treated in our center from 2011 to 2021. Subgroup analysis addressed whether patients on ECMO received different mechanical ventilation than patients without extracorporeal support.

Results: Compared to Influenza-associated ARDS, COVID-19 patients had higher ventilatory system compliance (40.7 ml/mbar [31.8 – 46.7 ml/mbar] vs. 31.4 ml/mbar [13.7 – 42.8 ml/mbar], $p = 0.198$), higher ventilatory ratio (1.57 [1.31 – 1.84] vs. 0.91 [0.44 – 1.38], $p = 0.006$) and higher minute ventilation at the time of intubation (mean minute ventilation 10.7 l/min [7.2 – 12.2 l/min] for COVID-19 vs. 6.0 l/min [2.5 – 10.1 l/min] for Influenza, $p = 0.013$). There were no measurable differences in P/F ratio, positive end-expiratory pressure (PEEP) and driving pressures (ΔP). Respiratory system compliance deteriorated considerably in COVID-19 patients on ECMO during 2 weeks of mechanical ventilation (C_{rs} , mean decrease over 2 weeks $-23.87 \text{ ml/mbar} \pm 32.94 \text{ ml/mbar}$, $p = 0.037$), but not in ventilated Influenza patients on ECMO and less so in ventilated COVID-19 patients without ECMO. For COVID-19 patients, low driving pressures on ECMO were strongly correlated to a decline in compliance after 2 weeks (Pearson's R 0.80, $p = 0.058$). Overall mortality was insignificantly lower for COVID-19 patients compared to Influenza patients (40% vs. 48%, $p = 0.31$). Outcome was insignificantly worse for patients requiring veno-venous ECMO in both groups (50% mortality for COVID-19 on ECMO vs. 27% without ECMO, $p = 0.30$ / 56% vs 34% mortality for Influenza A/B with and without ECMO, $p = 0.31$)

Conclusion: The pathophysiology of early COVID-19-associated ARDS differs from Influenza-associated acute lung failure by sustained respiratory mechanics during the early phase of ventilation. We question whether intubated COVID-19 patients on ECMO benefit from extremely low driving pressures, as this appears to accelerate derecruitment and consecutive loss of ventilatory system compliance.

Background

The emergence of COVID-19 in December 2019 brought a surge of patients with viral pneumonia and associated acute respiratory distress syndrome (ARDS)^{1,2} Shortly into the first pandemic wave, observations arose that respiratory failure due to COVID-19 could have unusual physiologic features³⁻⁶. In Mai 2020, Marini et al. described that patients with COVID-19 often present with severe hypoxemia yet only mildly impaired respiratory mechanics⁷. Since ARDS was historically accompanied by a loss of

functional residual capacity and compliance⁸, this observation sparked a debate whether respiratory failure due to COVID-19 might be a different disease. To this day, this question has not yet been sufficiently answered. Another recurrent observation in COVID-19 is a disproportionate increase in respiratory effort⁹, visible through elevated minute ventilation, leading to only marginal clearance of carbon dioxide (CO₂). Several mechanisms have been suggested to account for this discrepancy, e.g. ventilation-perfusion mismatch with increased functional right-to-left shunting¹⁰, increased CO₂ production¹¹ or a combination of both. The question if COVID-19 associated ARDS is different from 'classic' ARDS is not only of academic purpose, as it might imply a need for different intensive care management¹²⁻¹⁴.

Early in the first COVID-19 pandemic wave, centers started to employ extracorporeal membrane oxygenation for most severe COVID-19 cases, drawing on existing cohort data on ECMO for ARDS of different origin^{15,16}. While ECMO has proven to reduce mortality of patients with very severe COVID-19¹⁷, the mortality benefit for these patients was unsatisfyingly low¹⁸. It is well accepted that high driving pressures may increase mortality of ARDS patients through ventilator-associated lung injury (VILI)¹⁹. Consequently, these patients usually receive high positive end-expiratory pressure (PEEP) and low driving pressures at the cost of CO₂ elimination to limit mechanical power on the lungs²⁰. The strength of ECMO for ARDS patients is considered not only providing oxygenation but allowing protective ventilation through effective extracorporeal CO₂ clearance. If ventilator-induced lung injury contributes to disease progression in COVID-19, then protective ventilation during ECMO should decelerate the deterioration of lung mechanics.

We therefore aimed to check whether COVID-19 patients on ECMO benefitted from low driving pressures. Hence, we analyze in detail how intubated COVID-19 patients were ventilated on our ward in comparison to patients with Influenza-associated ARDS. Furthermore, we checked whether ventilation with high PEEP and low driving pressures during ECMO for COVID-19 had a positive impact on respiratory system physiology and compliance and compared these data to the Influenza cohort.

Methods

Study design

Our ICU is a tertiary care unit with 24 beds dedicated to the treatment of COVID-19 patients. Most COVID-19 patients included in the study were intubated in primary or secondary care centers nearby and then transferred for evaluation of ECMO. Several patients were cannulated externally by our mobile ECMO teams and then transferred to Homburg on running ECMO. We first analyzed basic characteristics of 25 patients with COVID-19, treated in our center from March 2020 to March 2021. All included patients were mechanically ventilated and devoid of severe comorbidities. Patients aged between 18 and 70 with laboratory-confirmed SARS-CoV-2 infection, defined as a positive result on real-time RT-PCR assay from nasal or pharyngeal swabs or respiratory tract aspirates, with hypoxemia and a high respiratory drive

meeting the criteria for moderate ARDS according to the Berlin definition of ARDS were included in this cohort.

Most COVID-19 patients treated in our center could not be recruited for the study cohort due to one of the following exclusion criteria: age- or comorbidity-related contraindication for extracorporeal life support, additional circulatory failure, defined as requiring intravenous noradrenaline of more than 0.3 µg/kg/min, cardiopulmonary resuscitation prior to intubation for COVID-19-associated ARDS, mechanical ventilation for COVID-19 for more than 8 days prior to study inclusion.

14 patients from the COVID-19 cohort later received veno-venous extracorporeal membrane oxygenation due to respiratory failure.

A matched control cohort of 25 patients with ARDS due to Influenza A or B, treated in our center from 2011 to 2021, was then recruited for comparison. As severe COVID-19 affects predominantly men, matching both cohorts for sex was not possible. Instead, both groups were matched according to age and BMI as well as possible. The same exclusion criteria were applied for the recruitment of Influenza patients.

We compared the following parameters taken from day 1, 2, 4, 8 and 16 of invasive ventilation: positive end-expiratory pressure (PEEP), plateau pressure (PP), driving pressure (ΔP) and Respiratory Rate (RR).

Ventilation was in pressure-controlled mode in all patients. Generally, we targeted a protective ventilation approach, aiming at driving pressures below 15 cmH₂O and tidal volumes (VT) of approximately 6 ml per kg of predicted bodyweight (PBW). The mean arterial pressure target was 60 – 65 mmHg if physiological aims were reached. These included capillary refill time (i.e. warm periphery), sufficient urinary output (≥ 0.5 ml/kg/h) and normal lactate levels (≤ 2.0 mmol/l). Norepinephrine was the vasopressor of choice in patients with pH ≥ 7.25 . Inotrope use was initiated if ScvO₂ was below 65% despite adequate hemoglobin levels. Nutrition in both groups was done according to the same standards.

If patients were not deeply sedated and breathed spontaneously, the actual rate of assisted spontaneous breathing was counted whenever intended and actual respiratory rates diverged by more than 2 per minute.

Derived from the above-mentioned parameters as well as results from blood gas samples, we calculated the following parameters: P_aO_2/F_iO_2 ratio (arterial oxygen partial pressure divided by inspiratory oxygen fraction), ventilatory system compliance, both absolute and in relation to idealized body weight (tidal volume in relation to predicted body weight divided by driving pressure), ventilatory ratio (VR) as a measure of ventilator efficiency, calculated as $(VE_{real} \times PaCO_{2real})$ divided by $(100 \text{ mL/min} \times \text{predicted body weight (PBW; kg)} \times 40 \text{ mmHg (expected } PaCO_2))$.

We defined the primary endpoint 'death from any cause' to calculate 28-day and 60-day mortality for both cohorts and subgroups as well as the secondary endpoint 'free from ventilator support after 15 days of

ventilation' to compare weaning success in both groups.

Statistical analysis

Continuous variables are displayed as mean, standard deviation and 95% confidence intervals. For ordinal variables and non-normally distributed continuous variables, we calculated median and interquartile range.

Horizontal comparisons between groups were done by two-sided t-testing for normally distributed continuous variables and Man-Whitney-U-test for non-continuous parameters. Changes in parameters over time were analyzed by paired two-sided t-testing at a significance level of 0.05. Kaplan-Meier-curves were plotted using the corresponding SPSS tool with 'death from any cause' as counted event. We performed log rank tests to compare survival between groups. Analysis and plotting were performed with SPSS Statistics v. 2.6.0.0.

Results

I) Basic patient characteristics

Basic patient characteristics are displayed in **Table I**. As severe COVID-19 affects predominantly men, the COVID-19 group has an excess of male patients (20 of 25 (80%) vs. 16 of 25 (64%). Apart from a non-significant age difference (61,7 years [53,1 – 68,1 years] for COVID-19 vs. 55,7 years [45,5 – 65,4 years] for Influenza, $p = 0.08$), both groups were well comparable regarding height, weight, body-mass index (BMI) and oxygenation impairment at the beginning of invasive ventilation.

Most frequent comorbidities were adiposity, arterial hypertension, Diabetes mellitus and coronary heart disease.

Both groups showed severe hypoxemia at the time of intubation. Mean arterial oxygen partial pressure before intubation was slightly higher in the influenza group (COVID19: paO_2 84.4 mmHg [76.8 – 92.1 mmHg] vs. 93.3 mmHg [71.8 – 114.8 mmHg] for Influenza, $p = 0.31$). Prior to intubation, patients received either oxygen supplementation via high-flow nasal cannula or non-invasive ventilation. Mean inspiratory oxygen fraction during non-invasive oxygenation was slightly lower in the COVID-19 group (0.6 [0.49 – 0.70], $n = 14$) compared to control (0.75 [0.53 – 0.84], $n = 9$).

13 of 25 COVID-19 patients (42%) and 16 of 25 Influenza (64%) patients received veno-venous extracorporeal membrane oxygenation for respiratory failure few days after intubation.

II) Horizontal comparison of ventilation parameters

On the first day of invasive ventilation, there were no significant differences in P/F ratio (127.9 mmHg [112.8 – 161.3 mmHg] for COVID-19 vs. 135.4 mmHg [100.3 – 180.4] mmHg, $p = 0.56$), positive end-expiratory pressure (12.0 mbar [9.6 – 13.5 mbar] for COVID-19 vs. 13.0 mbar [9.0 – 15.5 mbar] for

Influenza, $p = 0.37$), plateau pressure (23.5 mbar [$20.8 - 29.0 \text{ mbar}$] for COVID-19 vs. 25.0 mbar [$24.0 - 28.5 \text{ mbar}$] for *Influenza*, $p = 0.56$) or driving pressure (13.5 mbar [$10.8 - 16.0 \text{ mbar}$] for COVID-19 vs. 12.0 mbar [$11.0 - 14.5 \text{ mbar}$] for *Influenza*, $p = 0.64$) between both groups.

Similar ventilation pressures produced higher tidal volumes in relation to predicted body weight in COVID-19 patients (7.69 ml/kgBW [$7.12 - 8.12 \text{ ml/kgBW}$]) compared to control (5.12 ml/kgBW [$3.14 - 8.40 \text{ ml/kgBW}$], $p = 0.06$), indicating less impaired ventilatory system compliance in the COVID-19 group. COVID-19 patients required significantly higher ventilation frequencies (20.0 min^{-1} [$15.5 - 21.3 \text{ min}^{-1}$] vs. 14.0 min^{-1} [$12.5 - 17.5 \text{ min}^{-1}$], $p = 0.011$) and minute ventilation (10.7 l/min [$7.2 - 12.2 \text{ l/min}$] vs. 6.0 l/min [$2.5 - 10.1 \text{ l/min}$], $p = 0.013$) for sufficient CO₂ elimination. Indeed, calculated static compliance of the respiratory system was higher in COVID-19 patients than *Influenza* patients (40.7 ml/mbar [$31.8 - 46.7 \text{ ml/mbar}$] vs. 31.4 ml/mbar [$13.7 - 42.8 \text{ ml/mbar}$], $p = 0.198$) throughout the observation period. We calculated ventilatory ratio (VR) for both groups to assess ventilation efficacy. VR was significantly higher in the COVID-19 cohort (1.57 [$1.31 - 1.84$] vs. 0.91 [$0.44 - 1.38$], $p = 0.006$). For a comprehensive overview of ventilation parameters, s. **Tables II, III** and **Figure I**.

III) Longitudinal comparison of ventilation parameters

Immediately after intubation, most patients in both groups received pressure-controlled ventilation (PCV) during deep sedation or relaxation. The proportion of patients with strictly controlled ventilation on the first day was 79% in the COVID-19 group (11/14) and 100% in the *Influenza* group (9/9). All patients for whom ventilation data from the first day of mechanical ventilation were missing could not be included into the analysis.

As ventilation progressed, an increasing number of patients could be weaned to augmented spontaneous breathing (ASB) with continuous positive airway pressure (CPAP). On day 8, 41% of COVID-19 patients and 38% of *Influenza* patients breathed spontaneously with pressure support. On day 16, the rate of patients still on controlled ventilation had dropped to 9.5% in the COVID-19 group (2/21) but stayed relatively unchanged at 36% (4/11) in the control group.

Except for few *Influenza* patients developing prolonged disease, we observed that COVID-19 patients were ventilated much longer than most *Influenza* patients. Odds Ratio in the overall cohort for breathing free from ventilator 15 days after intubation was 3.5 for *Influenza* compared to COVID-19.

To assess how ventilatory system compliance developed over time, we performed longitudinal pair-wise t-test comparisons of the initial compliance from the day of intubation to day 8 and day 16 of invasive ventilation.

We detected a continuous decrease of compliance in the COVID-19 cohort, with a significant reduction after 2 weeks of ventilation ($47.99 \text{ ml/mbar} \pm 32.80 \text{ ml/mbar}$ vs. $24.13 \text{ ml/mbar} \pm 10.70 \text{ ml/mbar}$, mean difference after 15 days being $-23.87 \text{ ml/mbar} \pm 32.94 \text{ ml/mbar}$, $p = 0.037$) (**Figure II**). In contrast, ventilatory system compliance improved in the *Influenza* cohort after 7 days ($28.32 \text{ ml/mbar} \pm 21.98$

ml/mbar vs. 34.85 ml/mbar ± 29.59 ml/mbar, p = 0.10). Since the course of ARDS tended to be much shorter for Influenza patients, we cannot provide a comparison in this cohort over 15 days, as only 2 patients would have qualified for pair-wise comparison between day 1 and 16.

IV) Subgroup analysis – ECMO vs. Non-ECMO

Gradual decline in compliance was particularly present in the subgroup of COVID-19 patients receiving extracorporeal support (*mean loss of compliance after 7 days 19.5 ml/mbar ± 42.6 ml/mbar, p = 0.31, mean loss after 15 days: -36.19 ml/mbar ± 41.88 ml/mbar, p = 0.09*). COVID-19 patients not on ECMO also deteriorated in compliance, however, the loss of compliance was not as strong (*-9.08 ml/mbar ± 4.43 ml/mbar after 15 days, p = 0.010*).

After ECMO was initiated in most patients, we checked whether ventilation strategies changed following ECMO initiation. Indeed, for the COVID-19 group, patients on ECMO received significantly higher PEEP on day 8 and 16 compared to patients without ECMO (Day 8: 12.5 mbar ± 2.5 mbar vs. 10.3 mbar ± 2.0 mbar, $p = 0.025$, Day 16: 12.3 mbar ± 2.3 vs. 7.8 mbar ± 1.8 mbar, $p < 0.001$). Attempting to limit VILI, patients on ECMO were ventilated with lower driving pressures. This difference was small for day 8, as not all patients had been cannulated up to that day but was significant for day 16 (Day 8: 13.8 mbar ± 4.6 vs. 14.2 mbar ± 2.6 mbar, $p = 0.83$, Day 16: 11.3 mbar ± 2.3 mbar vs. 16.6 mbar ± 4.5 mbar, $p = 0.002$).

In contrast, there were no differences between PEEP and driving pressures between patients with and without ECMO for the Influenza group (not shown).

Since patients on ECMO tended to receive lower driving pressures but also seemed to deteriorate in ventilatory system compliance over time, we checked for correlations between a reduction in driving pressure and decreases in compliance over 15 days. We found that decreased driving pressures as a strategy of protective ventilation on ECMO was positively correlated to a loss of compliance after 15 days in bivariate correlation analysis (Pearson's $R = 0.80$, $p = 0.058$) in the COVID-19 group. In contrast, COVID-19 patients not receiving ECMO showed an inverse correlation between change in driving pressure and compliance on day 16 (Pearson's $R = -0.92$, $p = 0.025$). Compared to the COVID-19 group, this correlation was weak for Influenza patients on ECMO between the first and eighth day of ventilation (Pearson-coefficient -0.10 , $p = 0.88$).

IV) Outcome

We determined 'death from any cause' as an endpoint to assess survival rates after 28 and 60 days from the beginning of invasive ventilation.

28-day mortality was 16% for the COVID-19 group compared to 36% in the control group. As Kaplan-Meier-plots demonstrate (s. **Figure III A to C**), overall survival was slightly better for the COVID-19 group compared to control (40% vs. 48% for Influenza, $p = 0.31$). Requiring veno-venous ECMO was a negative predictive factor for survival in both groups (50% mortality for COVID-19 on ECMO vs. 27% without ECMO, $p = 0.30$ / 56% vs 34% mortality for Influenza A/B with and without ECMO, $p = 0.31$).

Discussion

Our study provides evidence that, during the early phase of mechanical ventilation, COVID-19 associated ARDS differs from Influenza-associated ARDS, mainly in lung mechanics. The ventilator ratio in patients with COVID-19 associated ARDS is statistically significant higher than in patients with Influenza-associated ARDS. Thus, the efficacy of CO₂ elimination in COVID-19 associated ARDS is highly impaired.

The finding that COVID-19 patients present with higher minute ventilation to achieve similar CO₂ clearance is in line with results from other groups^{6,21}. Recently, we were able to demonstrate that pulmonary shunt fraction is not elevated in COVID-19 patients compared to an ARDS cohort¹¹. Increased ventilatory effort is thus rather the result of heavily increased CO₂ production rather than right-to-left shunting. Almost all COVID-19 patients had high fevers, some as high as 40°C, which might have contributed to CO₂ accumulation. A recent retrospective analysis has connected fever of COVID-19 patients on admission to worse outcome²². Aggressive medical treatment of hyperthermia could limit respiratory drive and improve CO₂ clearance in both awake and intubated patients.

Although simplifying a complex disease, the concept of L- and H-type COVID-19 introduced by Gattinoni et al²³ reflects the observation that patients with COVID-19 tend to transition from sustained to impaired lung mechanics.

Early into the pandemic, intubation of COVID-19 patients was recommended permissively, mainly to prevent emergency intubation during acute deterioration and limit aerosol generation during non-invasive ventilation or high-flow oxygen supplementation²⁴. This approach has been questioned by emerging insight into COVID-19 lung physiology²⁵. The main advantages of invasive ventilation are maintenance of high positive end-expiratory pressure and facilitating lung recruitment. If patients with early COVID-19 associated respiratory failure possess sustained lung mechanics, it appears questionable whether they benefit from early intubation, as derecruitment is probably not the leading cause of hypoxemia. This insight has raised the dilemma which COVID-19 patients should be mechanically ventilated, provided they do not require emergency intubation, and if so, when intubation should be performed.

Awake patients with COVID-19 pneumonia typically produce high tidal volumes with high negative intrathoracic pressures during non-invasive ventilation or high-flow nasal cannula²⁶. The resulting mechanical strain puts them at risk of developing patient self-inflicted lung injury (pSILI), a concept which has already been described for other conditions with respiratory failure²⁷. On the other hand, once patients are intubated, the need for CO₂ control will most likely limit the possibilities of ventilating patients with tidal volumes low enough to limit ventilator-associated lung injury (VILI). Indeed, those COVID-19 patients from our cohort who did not receive veno-venous ECMO required higher tidal volumes and minute ventilation than what most physicians would consider 'protective' for ARDS. Furthermore, invasive ventilation increases the risk of ventilator-associated infections, possibly accelerating the transition from L- to H-type COVID-19²⁵. This assumption is in line with our data which show that

intubated COVID-19 patients without ECMO had required relatively high driving pressures throughout the course of disease. Interestingly, it is the ECMO group of COVID-19 patients in our cohort who had the heaviest losses in respiratory system compliance, possibly and paradoxically due to low driving pressures.

Our data suggest that reducing ventilation effort to a more protective scheme (e.g. 3-5 ml/kg predicted body weight) in patients on ECMO might give way to a subsequent loss of respiratory system compliance. We assume that reduced alveolar gas exchange might lead to further alveolar derecruitment in these patients. Subsequent alveolar edema might finally lead to the condition Gattinoni and others have described as 'H-type' COVID-19⁷. This effect was present even though, as outlined above, COVID-19 patients on ECMO had higher average PEEP than those without ECMO, suggesting that increase in PEEP alone is not sufficient to prevent derecruitment in these patients. While it is true that the danger of fatal hypoxia will probably be averted by ECMO, it is unknown whether this advantage can outweigh the long-term complications associated with derecruitment.

We did not detect a loss of compliance in our Influenza patients on ECMO, which is probably because respiratory system compliance was already severely impaired in these patients when they were intubated, hence protective ventilation on ECMO did not further jeopardize recruitment. As our mortality data shows, many COVID-19 patients do not recover from this state, questioning the role of highly protective ventilation on ECMO for COVID-19. A possible escape from this dilemma might be the initiation of awake ECMO in patients with L-type COVID-19. That way, vigorous respiratory effort could be limited easily through increased sweep gas flow, while avoiding invasive ventilation could reduce the risk of ventilator-associated pneumonia. Obviously, the number of patients qualifying for this strategy will be limited, as cognition and compliance should be unimpaired to avoid accidental decannulation. Cohort data is required to assess the possible benefits of awake ECMO in patients with early COVID-19.

Limitations

Our study has several limitations that need to be addressed. First, as we present real-life data, matching of COVID-19 and control cohorts was not ideal, since COVID-19 patients were noticeably, yet not significantly, older compared to control. Secondly, being a tertiary care center, many patients were intubated in external hospitals and later transferred to our center for ECMO evaluation. Hence, data on the first days of ventilation is missing for some patients. As this is the case for both COVID-19 and Influenza patients, systematic error will probably be limited. Finally, given that COVID-19 is a relatively new disease and Influenza patients were recruited from the last ten years, we cannot exclude that changes in internal clinical standards and improved management of ARDS patients might account for systemic bias between groups.

Conclusions

The pathophysiology of early COVID-19-associated ARDS differs from Influenza-associated acute lung failure by sustained respiratory mechanics during the early phase of ventilation. Patients with early COVID-19-associated severe respiratory failure frequently exhibit sustained lung mechanics. In these patients, the benefit from early intubation seems questionable, as derecruitment is not the leading cause of hypoxemia. Replacing intubation by awake ECMO might be a way out of this situation.

Abbreviations

| | |
|------------|-------------------------------------|
| ARDS | Acute respiratory distress syndrome |
| COVID-19 | Coronavirus disease 2019 |
| ΔP | driving pressure |
| ECMO | Extracorporeal membrane oxygenation |
| PEEP | positive end-expiratory pressure |
| PP | plateau pressure |
| RR | Respiratory Rate |
| VILI | Ventilator induced lung injury |
| VR | Ventilatory ratio |

Declarations

Ethics approval and consent to participate

Informed consent for the analysis of data is waived by the institutional review board due to the anonymous and retrospective analysis of data.

Consent for Publication

Not applicable.

Availability of data and materials

Data can be provided on request addressed to the corresponding author. All data sharing statements are subject to conformity with German data protection legislation and rules (Datenschutzgrundverordnung - DSGVO).

Conflict of interest disclosure

Robert Bals declares funding from AstraZeneca, Boehringer Ingelheim,

GlaxoSmithKline, Grifols, Novartis, CLS Behring, the German Federal Ministry of Education and Research (BMBF) Competence Network Asthma and COPD (ASCONET), Sander-Stiftung, Schwiete-Stiftung, Krebshilfe and Mukoviszidose eV. Conflicts that the editors consider relevant to the content of the manuscript have been disclosed. All other authors: No potential conflicts of interest

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Authorship Contribution

P.L. and S.M. drafted the study. N.K. oversaw data collection and analysis. S.M. aided with statistical analysis and drafted the manuscript. F.S., G.D., R.M., C.R., A.B., M.K., T.R., C.M., R.B. and P.L. revised the manuscript for important intellectual content. All authors have seen and approved the final version of the manuscript.

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Tables

Table I - Basic patient data

| | COVID (n = 25) | Influenza (n = 25) | P |
|---|-------------------------------|-------------------------------|------------|
| Age on admission [years] | 61,7 [53,1 – 68,1] | 55,7 [45,5 – 65,4] | p = 0.082* |
| Male / female | 20 / 5 | 16 / 9 | |
| Height [m] | 1,77 [1,73 – 1,81] | 1,73 [1,70 – 1,76] | p = 0.105 |
| Weight [kg] | 100,4 [91,6 – 109,2] | 95,9 [81,3 – 110,5] | p = 0.573 |
| Body-Mass Index (BMI) [kg/m²] | 32,3 [29,0 – 35,5] | 31,5 [27,0 – 36,0] | p = 0.781 |
| Comorbidities: | Adiposity (19) | Adiposity (10) | |
| | Arterial Hypertension (13) | Arterial Hypertension (10) | |
| | Diabetes mellitus Type II (3) | Diabetes mellitus Type II (9) | |
| | Coronary Heart Disease (3) | Coronary Heart Disease (6) | |
| | Congestive Heart Failure (3) | COPD (4) | |
| | Past pulmonary embolism (3) | Chronic kidney disease (3) | |
| | Past stroke (4) | Epilepsy (2) | |
| | Myasthenia gravis (1) | Atrial fibrillation | |
| | | Cystic fibrosis | |
| | | Asthma | |

*Mann-Whitney-U

Table II – Comparison of ventilation parameters on the first day of invasive ventilation

| | COVID-19 | Influenza A / B | significance |
|---|---------------------------------|--------------------------------|------------------|
| Arterial oxygen partial pressure [mmHg] | 84.4 [76.8 – 92.1] n = 14 | 93.3 [71.8 – 114.8] n = 9 | p = 0.313 |
| Inspiratory oxygen fraction during non-invasive ventilation / HFNC | 0.6 [0.49 – 0.70] n = 14] | 0.75 [0.53 – 0.84] N = 9 | p = 0.227 |
| Arterial carbon dioxide partial pressure [mmHg] | 42.1 [38.1 – 46.0] n = 14 | 45.0 [38.1 – 51.9] n = 9 | p = 0.384 |
| pH | 7.39 [7.37 – 7.41] n = 14 | 7.34 [7.30 – 7.39]] N = 8 | p = 0.03 |
| Horovitz-Index | 127.9 [112.8 – 161.3] N = 14 | 135.4 [100.3 – 180.4] N = 9 | p = 0.557 |
| Positive end-expiratory pressure (PEEP) [cmH ₂ O / mbar] | 12.0 [9.6 – 13.5] N = 14 | 13.0 [9.0 – 15.5] N = 9 | p = 0.369 |
| Plateau pressure [cmH ₂ O / mbar] | 23.5 [20.8 – 29.0] N = 14 | 25.0 [24.0 – 28.5] N = 9 | p = 0.561 |
| Delta-P [cmH ₂ O / mbar] | 13.5 [10.8 – 16.0] N = 14 | 12.0 [11.0 – 14.5] N = 9 | p = 0.643 |
| Tidal volume [ml] | 551 [434.5 – 593.8] N = 14 | 431 [210 - 572] N = 9 | p = 0.124 |
| Tidal volume per kg predicted body weight [ml / kg] | 7.69 [7.12 – 8.12] N = 14 | 5.12 [3.14 – 8.40] N = 8 | p = 0.059 |
| Respiratory Rate* [min ⁻¹] | 20.0 [15.5 – | 14.0 [12.5 – | p = 0.011 |

| | | | |
|-----------------------------------|-------------------|------------------|------------------|
| | 21.3] | 17.5] | |
| | N = 14 | N = 9 | |
| Minute ventilation [l/min] | 10.7 [7.2 – 12.2] | 6.0 [2.5 – 10.1] | p = 0.013 |
| | N = 14 | N = 9 | |

* the real frequency of assisted spontaneous breathing was counted whenever intended and actual rates diverged by more than 2 per minute

Table III – Comparison of ventilatory system compliance and respiratory ratio on the first day of ventilation

| | COVID-19 | Influenza A / B | significance |
|--|----------------------------|---------------------------|---------------------|
| Ventilatory system compliance [ml/mbar] | 40.7 [31.8 – 46.7]; N = 14 | 31.4 [13.7 – 42.8]; N = 9 | P = 0.198 |
| Ventilatory system compliance / kg idealized body weight [ml/mbar/kg] | 0.57 [0.48 – 0.70]; N = 14 | 0.41 [0.20 – 0.59]; N = 8 | P = 0.150 |
| Ventilatory Ratio | 1.57 [1.31 – 1.84]; N = 14 | 0.91 [0.44 – 1.38]; N = 7 | P = 0.006 |

Figures

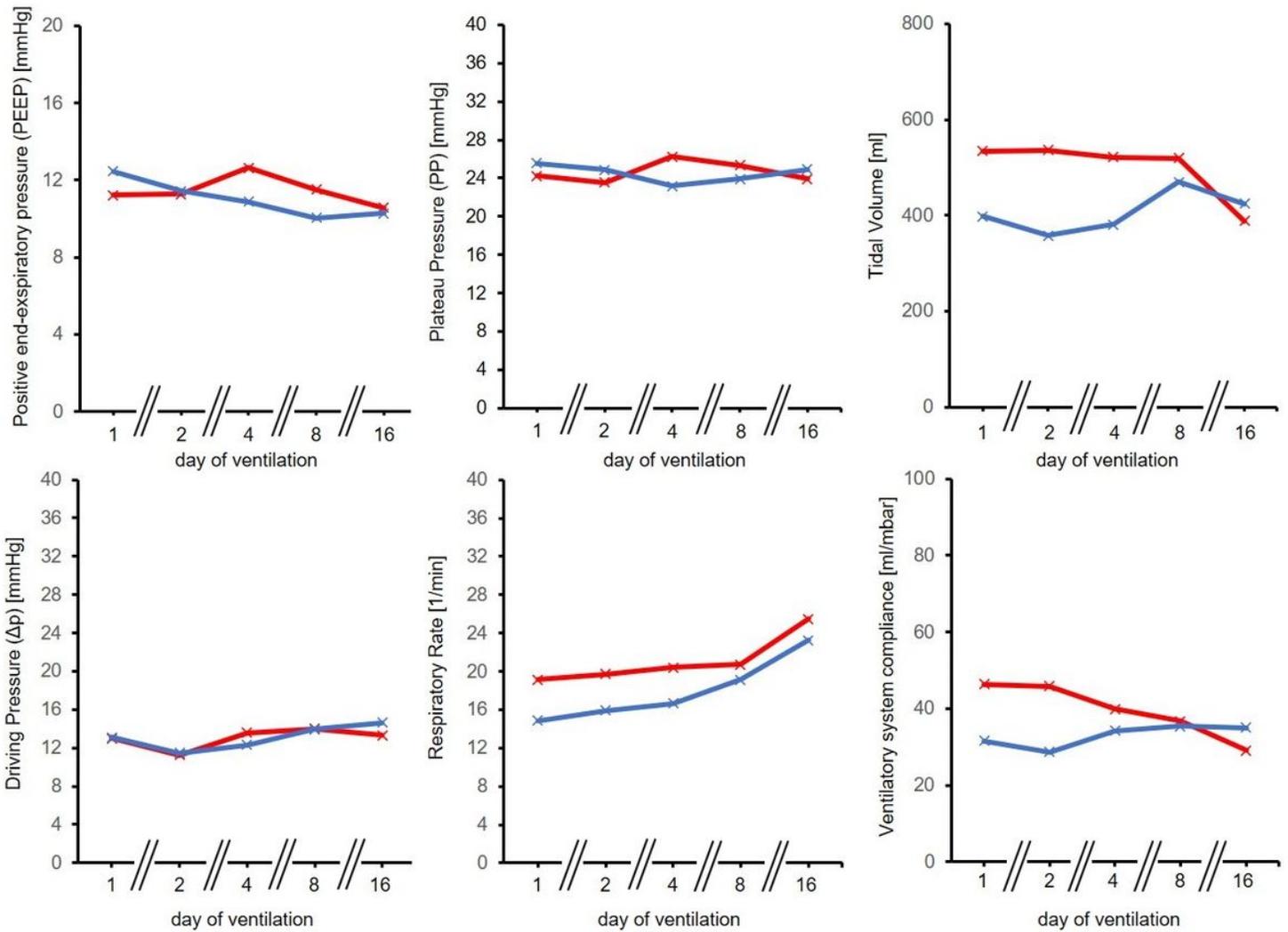


Figure 1

Development of ventilation parameters over time for COVID-19 and Influenza A / B Time course was similar in both groups for positive end-expiratory pressure (PEEP), Plateau Pressure (PP) and Driving Pressure (Δp). Respiratory Rate (RR), Tidal volumes and ventilatory system compliance were initially higher for the COVID-19 group.

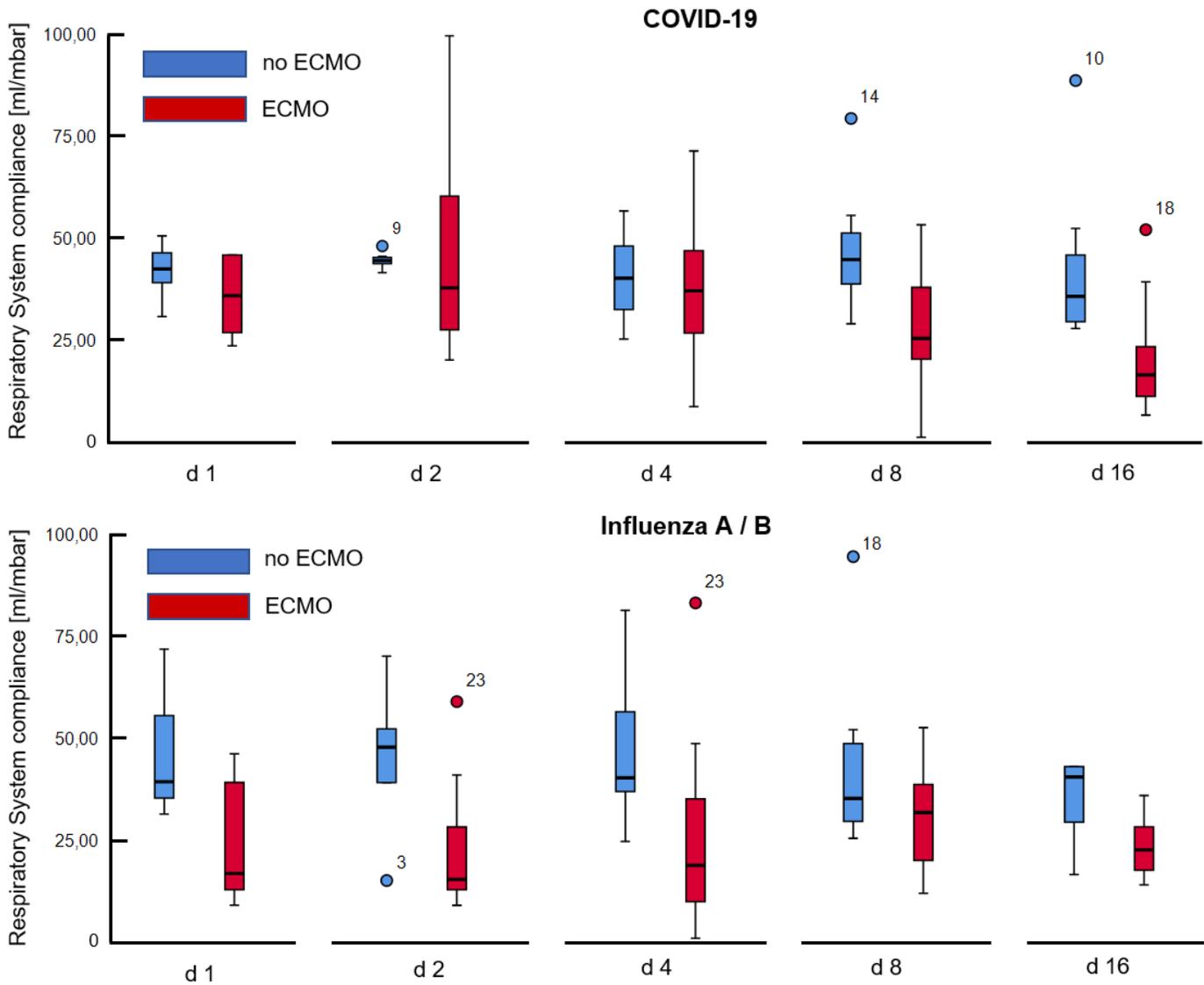


Figure 2

Development of Respiratory system compliance over time for COVID-19 and Influenza associated ARDS. Histograms depict that ventilatory system compliance showed gradual decline over 2 weeks for COVID-19 patients on veno-venous ECMO, while compliance was severely impaired in Influenza patients from the very beginning of invasive ventilation, improving insignificantly over time.

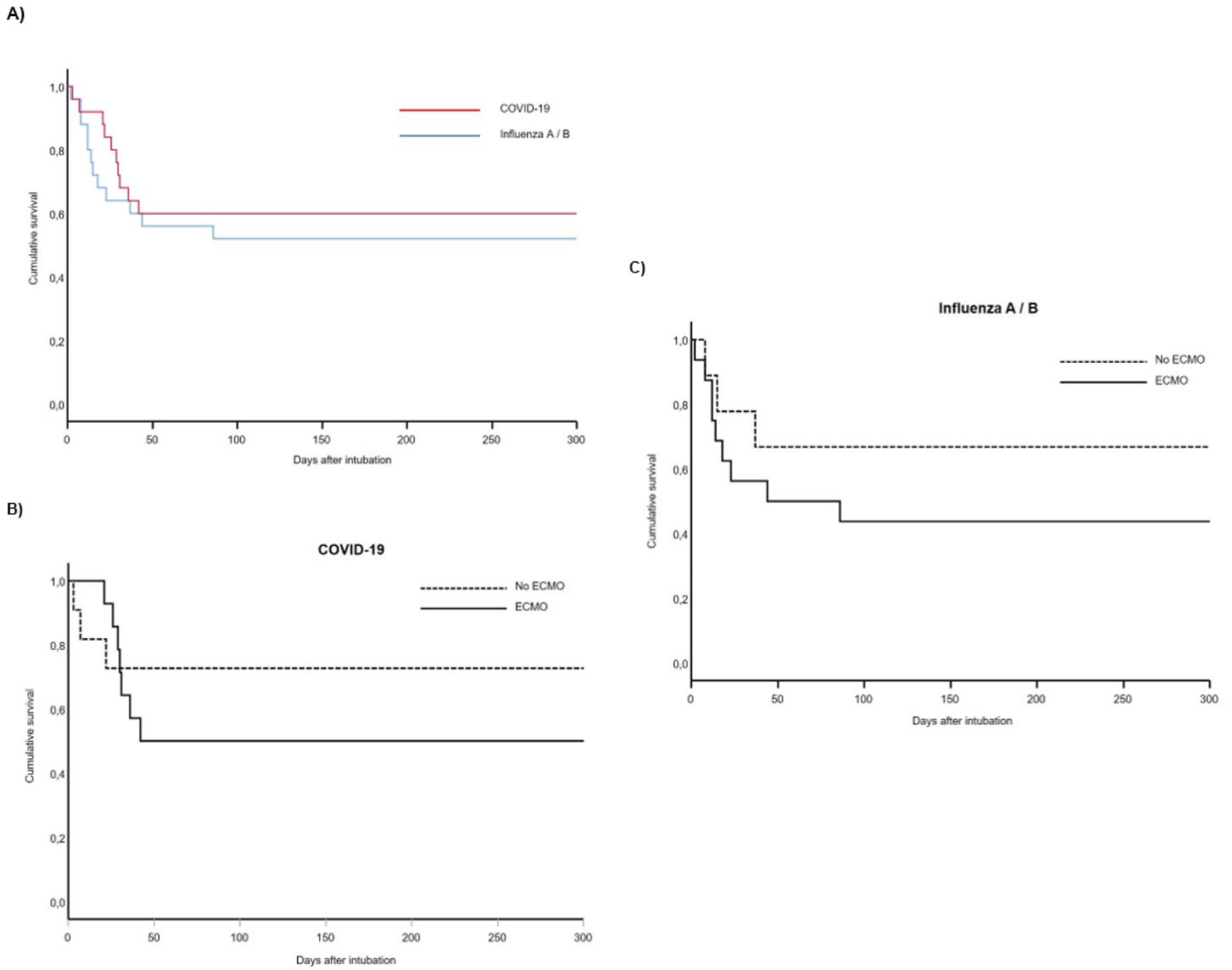


Figure 3

Kaplan-Meier-plots for COVID-19 and Influenza patients A) overall cumulative survival for COVID-19 and Influenza patients B) Subgroup analysis of cumulative survival for COVID-19 patients on ECMO and without ECMO C) Subgroup analysis of cumulative survival for Influenza patients on ECMO and without ECMO