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Perspective

Extracorporeal membrane oxygenation (ECMO): does it have a role in the treatment of severe COVID-19?

Xiaoyang Hong\textsuperscript{a,1}, Jing Xiong\textsuperscript{b,c,d,e,f,1}, Zhicheng Feng\textsuperscript{a,*,1}, Yuan Shi\textsuperscript{b,c,d,e,f,**}

\textsuperscript{a} Bayi Children’s Hospital, The Seventh Medical Center, PLA General Hospital, Beijing, China
\textsuperscript{b} Department of Neonatology, Ministry of Education Key Laboratory of Child Development and Disorders, PR China
\textsuperscript{c} National Clinical Research Center for Child Health and Disorders, PR China
\textsuperscript{d} China International Science and Technology Cooperation base of Child development and Critical Disorders, PR China
\textsuperscript{e} Children’s Hospital of Chongqing Medical University, PR China
\textsuperscript{f} Chongqing Key Laboratory of Pediatrics, Chongqing, 400014, PR China

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\textbf{A B S T R A C T}

The severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has emerged since December 2019 in Wuhan city, and has quickly spread throughout China and other countries. To date, no specific treatment has been proven to be effective for SARS-CoV-2 infection. According to World Health Organization (WHO), management of coronavirus disease 19 (COVID-19) has mainly focused on infection prevention, case detection and monitoring, and supportive care. Given to the previous experience, extracorporeal membrane oxygenation (ECMO) has been proven to be an effective therapy in the treatment of respiratory failure or acute respiratory distress syndrome (ARDS). On the basis of similar principle, ECMO may be also an effective therapy in the treatment of severe COVID-19. In this study, we described and discussed the clinical outcomes of ECMO for ARDS patients, ECMO use for severe COVID-19 in China, the indications of ECMO use, and some important issues associated with ECMO.

Since the end of 2019, an outbreak of pneumonia caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has occurred in China. In a recent report published in JAMA, 26.1% of 138 coronavirus disease 2019 (COVID-19) patients needed to be admitted to intensive care unit (ICU), of which 61.1% were suffering from acute respiratory distress syndrome (ARDS). The case fatality rate of COVID-19 has been reported about 4.3% (Wang et al., 2020). Until now, no specific treatment has been recommended for COVID-19.

Historical perspective

Extracorporeal membrane oxygenation (ECMO), which can provide effective respiratory or cardiac support, has been regarded as a rescue therapy for severe ARDS. ECMO therapy during the influenza A (H1N1) pandemic in 2009 appeared to benefit, with ECMO-treated patients with H1N1-related ARDS achieving a mortality of 21%, which greatly increased interest in its use (The Australia and New Zealand Extracorporeal Membrane Oxygenation, 2009). Another cohort study by using ECMO database of patients with H1N1-related ARDS showed that hospital mortality rate was 23.7% for ECMO-referred patients vs 52.5% for non-ECMO-referred patients (RR, 0.45 [95% CI, 0.26-0.79]; P = 0.006) when individual matching was used; 24.0% vs 46.7%, respectively (RR, 0.51 [95% CI, 0.31-0.81]; P = 0.008) when propensity score matching was used; and 24.0% vs 50.7%, respectively (RR, 0.47 [95% CI, 0.31-0.72]; P = 0.001) when GenMatch matching was used. These suggested that for patients with H1N1-related ARDS, ECMO-referred patients were associated with significantly lower hospital mortality compared with matched non-ECMO-referred patients (Noah et al., 2011). A clinical trial, named as CESAR, was encouraging as well (Peek et al., 2009). However, EOLIA Clinical Trial showed that 60day mortality with very severe ARDS patients was not significantly lower, yet was largely reduced in the ECMO group compared with the conventional mechanical ventilation group (35% vs 46%; RR, 0.76 [95% CI, 0.55-1.04]; P = 0.09), but there was a 28% crossover to ECMO for failure of conventional mechanical ventilation, suggesting a lack of clinical equipoise (Combes et al., 2018). Otherwise, a post hoc Bayesian

\footnote{\textsuperscript{*} Corresponding author at: Bayi Children’ Hospital, The Seventh Medical Center of Chinese PLA General Hospital, Beijing, 100000, China.}
\footnote{\textsuperscript{**} Corresponding author at: Department of Neonatology, Children’s Hospital of Chongqing Medical University, Chongqing, 400014, China.}
\footnote{\textsuperscript{1} Dr Hong and Dr Xiong contributed equally to this study.}

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analysis of EOLIA with various assumptions of prior belief and knowledge about ECMO efficacy in ARDS showed the posterior probability of a mortality reduction with ECMO in the EOLIA trial (Goldigger et al., 2018). In 2018, a retrospectively study on middle east respiratory syndrome (MERS) patients with refractory respiratory failure indicated that ECMO should be used as a rescue therapy, and ECMO group was associated with lower mortality in MERS patients with refractory hypoxemia compared with the conventional group (65 vs 100%, \( P = 0.02 \)) (Alshahrani et al., 2018) (Table 1).

**Application situation of COVID-19 in China**

According to the interim guidance formulated by the World Health Organization (WHO), ECMO should be considered as a rescue therapy for COVID-19 with refractory hypoxemia despite lung-protective ventilation (WHO, 2020). However, there is little experience with using ECMO to support SARS-CoV-2-infected patients (Wang et al., 2020; Yang et al., 2020; Guo et al., 2020; Guan et al., 2020; Huang et al., 2020). Most of studies didn’t report the clinical outcomes of ECMO use except for two studies. In the retrospective study conducted by Yang et al., 52 critically ill adult patients were identified with SARS-CoV-2 pneumonia and were admitted to intensive care unit (ICU), among them, 31 patients had died at 28 days, 6 patients were received ECMO, and 5 of them died and 1 patient was still on ECMO at the endpoint (Yang et al., 2020). Another retrospective study implemented by Guo et al. included 221 patients with laboratory confirmed SARS-CoV-2 pneumonia, 48 of severe patients developed ARDS, and 10 of them received invasive mechanical ventilation (IMV) and ECMO support. 2 patients had clinical benefits and had been discharged and 3 of them were non-survivors. The rest 5 patients were still on ECMO at the endpoint (Guo et al., 2020) (Table 2). Given lacking of clinical trial of ECMO on COVID-19, we could not conclude whether SARS-CoV-2-infected patients have benefited from ECMO at this time. But our concern may be settled by the ongoing trials in China (ChiCTR2000030744 and ChiCTR2000029804).

**Indications for the treatment of COVID-19 by ECMO**

Based on the entry criteria of EOLIA, ECMO should be considered when meeting one of the following three criteria despite optimization of mechanical ventilation for <7 days (\( \text{FiO}_2 > 0.80 \), tidal volume of 6 ml/kg predicted body weight, \( \text{PeePe}>10 \text{cmH}_2\text{O} \)) (Brodie et al., 2019): (1) \( \text{PaO}_2 / \text{FiO}_2 < 50 \text{ mmHg} \) for > 3 hours; (2) \( \text{PaO}_2 \); \( \text{FiO}_2 < 80 \text{ mmHg} \) for > 6 hours; (3) pH < 7.25 with \( \text{PaCO}_2 \geq 60 \text{ mmHg} \) for > 6 hours with a respiratory rate increased to 35 breaths per minute, adjusted for plateau pressure \(< 32 \text{ cmH}_2\text{O} \). Alternatively, after lung protective ventilation (tidal volume 6 ml/kg, \( \text{PeePe}>10 \text{cmH}_2\text{O} \)) was adopted and combined with lung recruitment maneuver, prone position ventilation, and high-frequency oscillation ventilation, patients are still under the condition of pure oxygen inhalation, in these situations, ECMO should be considered for ARDS as rescue therapy when meeting one of the following criteria: (1) \( \text{PaO}_2 / \text{FiO}_2<100 \text{ mmHg} \); (2) \( \text{P(A-a)O}_2>600 \text{ mmHg} \); (3) pH < 7.2 and plateau pressure >30 cmH\(_2\)O with respiratory rate more than 35 breaths per minute; (4) Age<65 years old; (5) Mechanical ventilation<7d; (6) Absence of contraindications (Critical Care Medicine Committee of the Chinese Association of Chest Physicians, 2019). Besides, for the patients with a harmful potential of ventilator-induced lung injury, lower ventilation and volumes and pressures may lead to hypercapnic acidosis, in this situation, extracorporeal carbon dioxide removal (ECCO2R) can be an important tool by providing direct removal of CO\(_2\) from blood (Brodie et al., 2019).

**What should we do next?**

Indeed, many factors could affect the outcomes of ECMO treatment, including the duration of mechanical ventilation, the severity of underlying disease, the experience of trained medical staff, and ECMO equipment. Early evaluation, rapid assembly, and cannulation timely are important. Regardless of the efficacy of ECMO, under the special situation of the SARS-CoV-2 outbreak, we should also pay more attention to the safety of medical staff since they get infected easily when manipulating ECMO. Some approaches, such as intubation, ventilator venting, and sputum suction pose a high risk of infection to medical staff. Therefore, all related staff should be supplied with sufficient protection and be restricted in the independent area. As the pandemic spread, a shortage of ECMO consoles may be another problem to be solved due to a surge of critically ill patients worldwide. Furthermore, trained staff and isolation rooms should be in full preparedness to meet the coming challenges.

**Author contributions**

XH conceptualized the study and revised the final manuscript. JX drafted the initial manuscript. ZF reviewed the manuscript for important intellectual content. YS conceptualized and designed the study and critically reviewed the manuscript for important intellectual content. All authors reviewed the manuscript. XH and JX contributed equally to this study.

**Table 1**

| Application | Publication time | Study design | Outcomes of ECMO | Reference |
|-------------|-----------------|--------------|------------------|-----------|
| Influenza A (H1N1) | 2009 | Observational study | A mortality rate of 21% in the ECMO-treated patients | (The Australia and New Zealand Extracorporeal Membrane Oxygenation, 2009) |
| ARDS | 2009 | Multicenter RCT | 63% (57/90) of patients considered by ECMO survived to 6 months without disability compared with 47% (41/87) of those allocated to conventional management (RR, 0.69; 95% CI 0.05–0.97, \( P = 0.03 \)) | (Peek et al., 2009) |
| ARDS (CESAR) | 2011 | Cohort study | ECMO-referred patients was associated with lower mortality compared with non-ECMO-referred patients | (Noah et al., 2011) |
| Influenza A (H1N1) | 2018 | Multicenter RCT | 60-day mortality was not significantly lower with ECMO than with a strategy of conventional mechanical ventilation (35% vs 46%, \( P = 0.09 \)) | (Combes et al., 2018) |
| ARDS (EOLIA) | 2018 | Retrospective study | ECMO use was associated with lower mortality in MERS patients with refractory hypoxemia (65% vs 100%, \( P = 0.02 \)) | (Alshahrani et al., 2018) |

ARDS = acute respiratory distress syndrome; CI = confidence interval; ECMO = extracorporeal membrane oxygenation; MERS = middle east respiratory syndrome; RCT = randomized controlled trial; RR = relative risk.
Table 2
Current clinical uses of ECMO for COVID-19

| Application                                      | Study design   | Cases on ECMO (total cases) | Outcomes of ECMO                          | Reference |
|--------------------------------------------------|----------------|-----------------------------|-------------------------------------------|-----------|
| Critically ill patients with SARS-CoV-2 pneumonia | Single center, retrospective study | 6 (52)                     | Five patients died while one patient was still on ECMO at the endpoint | (Yang et al., 2020) |
| Patients with ARDS caused by SARS-CoV-2          | Single center, retrospective study | 10 (221)                    | Two patients were discharged, three patients died, and five patients were still on ECMO at the endpoint | (Guoqin et al., 2020) |
| Critically ill patients with SARS-CoV-2 pneumonia | Single center, retrospective study | 4 (138)                     | NA                                        | (Guan et al., 2020) |
| Critically ill patients with SARS-CoV-2 pneumonia | Multicenter retrospective study | 5 (1099)                    | NA                                        | (Huang et al., 2020) |
| Critically ill patients with SARS-CoV-2 pneumonia | Single center prospective study | 2 (41)                      | NA                                        | (Brodie et al., 2019) |

ARDS = acute respiratory distress syndrome; COVID-19 = coronavirus disease 2019; ECMO = extracorporeal membrane oxygenation; NA = not available; SARS-CoV-2 = severe acute respiratory syndrome coronavirus 2.

Conflict of Interest

The authors declare no conflicts of interest.

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Ethical Approval

Not applicable.

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