Extracorporeal Membrane Oxygenation in the Treatment of Severe Pulmonary and Cardiac Compromise in Coronavirus Disease 2019: Experience with 32 Patients

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As coronavirus disease 2019 (COVID-19) cases surge worldwide, an urgent need exists to enhance our understanding of the role of extracorporeal membrane oxygenation (ECMO) in the management of severely ill patients with COVID-19 who develop acute respiratory and cardiac compromise refractory to conventional therapy. The purpose of this manuscript is to review our initial clinical experience in 32 patients with confirmed COVID-19 treated with ECMO. A multi-institutional registry and database was created and utilized to assess all patients who were supported with ECMO provided by SpecialtyCare. Data captured included patient characteristics, pre-COVID-19 risk factors and comorbidities, confirmation of COVID-19 diagnosis, features of ECMO support, specific medications utilized to treat COVID-19, and short-term outcomes through hospital discharge. This analysis includes all of our patients with COVID-19 supported with ECMO, with an analytic window starting March 17, 2020, when our first COVID-19 patient was placed on ECMO, and ending April 9, 2020. During the 24 days of this study, 32 consecutive patients with COVID-19 were placed on ECMO at nine different hospitals. As of the time of analysis, 17 remain on ECMO, 10 died before or shortly after decannulation, and five are alive and extubated after removal from ECMO, with one of these five discharged from the hospital. Adjunctive medication in the surviving patients while on ECMO was as follows: four of five survivors received intravenous steroids, three of five

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survivors received antiviral medications (Remdesivir), two of five survivors were treated with anti-interleukin-6-receptor monoclonal antibodies (Tocilizumab or Sarilumab), and one of five survivors received hydroxychloroquine. An analysis of 32 COVID-19 patients with severe pulmonary compromise supported with ECMO suggests that ECMO may play a useful role in salvaging select critically ill patients with COVID-19. Additional patient experience and associated clinical and laboratory data must be obtained to further define the optimal role of ECMO in patients with COVID-19 and acute respiratory distress syndrome (ARDS). These initial data may provide useful information to help define the best strategies to care for these challenging patients and may also provide a framework for much-needed future research about the use of ECMO to treat patients with COVID-19. ASAIO Journal XXX; XX:00-00.

Key Words: extracorporeal membrane oxygenation, coronavirus, coronavirus disease 2019, pulmonary failure, acute respiratory distress syndrome, heart failure, outcomes, quality

As of April 9, 2020, 1,579,690 patients around the world have been diagnosed with confirmed coronavirus disease 2019 (COVID-19), with 94,807 associated deaths (6.0% mortality worldwide).¹ Meanwhile, in the United States, as of April 9, 2020, 452,582 patients have been diagnosed with confirmed COVID-19, with 16,129 associated deaths to date (3.6% mortality in the United States).¹ Most deaths in patients with COVID-19 are due to severe respiratory failure, with a small group succumbing to combined pulmonary and cardiac failure.^{2,3}

Extracorporeal membrane oxygenation (ECMO) is an advanced life support modality that was initially used to treat severe neonatal respiratory failure.^{4,5} Over time, the use of ECMO has expanded, with ECMO presently utilized widely to treat multiple forms of severe acute respiratory or cardiac failure in neonates, infants, children, and adults.^{5,6} In the 2009 CESAR trial, adults with severe acute respiratory failure were randomized to treatment with ECMO *versus* maximal conventional ventilatory support and management (*e.g.*, steroids, prone positioning, bronchoscopy, and inhaled nitric oxide). In that study, 63% of patients (57/90) allocated to consideration for treatment using ECMO survived to 6 months without disability compared with 47% (41/87) of those allocated to conventional management.

In 2009 and 2010, the "swine-flu" H1N1 pandemic caused thousands of deaths in the United States. According to the

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Centers for Disease Control and Prevention (CDC) of the United States, from April 12, 2009, to April 10, 2010, there were "60.8 million cases (range: 43.3–89.3 million), 274,304 hospitalizations (range: 195,086–402,719), and 12,469 deaths (range: 8,868–18,306) in the United States due to the (H1N1) pdm09 virus."⁷ During the H1N1 pandemic, ECMO was used successfully to salvage patients with severe respiratory failure, with an associated 79% survival.⁸

As COVID-19 cases surge worldwide, an urgent need exists to enhance our understanding of the role of ECMO in the management of severely ill patients with COVID-19. The purpose of this report is to review our early clinical experience with the use of ECMO in 32 patients with confirmed COVID-19 and severe pulmonary compromise, some of whom also developed severe cardiac compromise.

Materials and Methods

A real time cohort study was conducted of all patients with confirmed COVID-19 who were supported with ECMO therapy provided by SpecialtyCare; a multi-institutional registry and database was created and utilized to assess these patients. (SpecialtyCare is a United States provider of allied health services: predominantly perfusion services and intraoperative neuromonitoring, based in Brentwood, TN [https:// specialtycareus.com/]) Data captured included patient characteristics, pre-COVID-19 risk factors and comorbidities, confirmation of COVID-19 diagnosis, features of ECMO support, specific medications utilized to treat COVID-19, and shortterm outcomes through hospital discharge. This database is prospectively maintained on all patients and has been used for data collection and analysis. The database used is a component of the SCOPE registry; SpecialtyCare, Nashville, TN (https://specialtycareus.com/).

This analysis includes all of our patients with documented COVID-19 infection who were supported with ECMO, with an analytic window starting March 17, 2020, when our first COVID-19 patient was placed on ECMO, and ending April 9, 2020. Entry criteria for placement on ECMO was determined by the individual patient care team at the each of the nine hospitals submitting data; all patients were placed on ECMO with severe respiratory failure felt to be refractory to conventional management. The decision to initiate ECMO, the mode of therapy (*i.e.*, veno-veno, veno-arterial), and the cannulation strategy were all determined by the individual ECMO-Teams, as determined by their individual institutional protocols and guidelines.

Descriptive analysis of the entire cohort was performed using mean, standard deviation, median, and interquartile range, as appropriate. The primary outcome of interest was mortality during the index hospitalization. Potential differences in categorical variables by mortality group were assessed using χ^2 and Fisher exact tests, while possible differences in continuous variables by mortality group were assessed using Kruskal–Wallis rank-sum tests and Welch ANOVA.

Institutional Review Board (IRB) approval and waiver of the need for consent were obtained. The human subjects' research protocol for this study was reviewed and approved by an independent IRB. Institutional ethics review board approval was obtained for the use of data from the SCOPE registry (Protocol No. 012017, ADVARRA Center for IRB Intelligence, 6940 Columbia Gateway Drive, Suite 110, Columbia, MD 21046).

Results

During the 24 days of this study, 32 consecutive patients with COVID-19 were placed on ECMO at nine different hospitals. **Table 1** depicts the number of patients with COVID-19 placed on ECMO at each hospital and the geographic location of each hospital, as well as data about the number of patients in each State diagnosed with COVID-19, hospitalized for COVID-19 and dead. These regional data contextualize the data from each hospital.

As of the time of analysis, 17 out of 32 patients remain on ECMO, 10 died before or shortly after decannulation, and five are alive following discontinuation of ECMO. All of the five survivors have been separated from mechanical ventilation, with one having been discharged from the hospital to date. **Table 2** provides detailed data about all 32 patients with COVID-19 treated with ECMO. Of note, 14 of 32 patients (43.8%) had obesity, 11 of 32 patients (34.4%) had diabetes, four of 32 patients (12.5%) had heart disease, three of 32 patients (9.4%) had cancer, and three of 32 patients (9.4%) had asthma.

Table 3 provides detailed data about 15 patients with COVID-19 treated with ECMO and no longer on ECMO and compares the characteristics of the five survivors to the 10 nonsurvivors. Adjunctive medication in the surviving patients while on ECMO was as follows: four of five survivors received intravenous steroids, three of five survivors received antiviral medications (Remdesivir), two of five survivors were treated with anti-interleukin-6 (IL-6) receptor monoclonal antibodies (Tocilizumab or Sarilumab), and one of five survivors received hydroxychloroquine. In the 10 patients who died, documented causes of death were as follows: respiratory failure (6/10), disseminated intravascular coagulation (DIC, 2/10), multisystem organ failure (MSOF) including acute kidney injury (1/10), and cerebral bleeding while on ECMO (1/10).

None of these 32 patients were placed on ECMO during cardiopulmonary resuscitation (CPR) (*i.e.*, extracorporeal CPR [ECPR]) was not utilized in this cohort). All five survivors were supported only with veno-venous ECMO. Furthermore, no patients receiving partial or complete veno-arterial ECMO have survived decannulation (five patients were supported with partial or complete veno-arterial ECMO: three have died and two remain on ECMO). Zero patients were converted from veno-venous ECMO to veno-arterial ECMO. Zero patients were converted from veno-venous ECMO to V-AV ECMO (ECMO with systemic venous inflow with dual systemic venous and systemic arterial outflow [combined veno-venous and veno-arterial ECMO]).

Figure 1 depicts the current status of all 32 COVID-19 ECMO patients. **Figure 2** depicts the number of patients cannulated each week. **Figure 3** depicts the distribution of hours on ECMO, comparing the survivors with the nonsurvivors. **Figure 4** depicts the distribution of the age of the patients, comparing the survivors with the nonsurvivors.

A brief case history of one of these patients is enlightening: A 51 white female with no past medical history sustained an orthopedic injury while on vacation in Vail, Colorado. On March 7, 2020, she underwent elective repair of her ankle injury. During extubation from the orthopedic procedure, frothy

Hospital	No. of Patients witl COVID-19 Placed on ECMO		Patients in State Diagnosed with COVID-19 As of April 13, 2020	Patients in State Hospitalized As of April 13, 2020	Patients in State in Intensive Care Unit As of April 13, 2020	Deaths in State As of April 13, 2020	
А	10	California	23,338*	3,124*	1,177*	758*	
В	7	Texas	15,492†	1,538†		364†	
С	3	Colorado	8,280‡	1,636‡		357‡	
D	3	Washington, DC	2,197§			72§	
E	3	Pennsylvania	26,490¶	2,440¶		647¶	
F	2	Florida	21,865	3,249		614	
G	2	Pennsylvania	26,490¶	2,440¶		647¶	
Н	1	Hawaii	530#			9#	
I	1	Indiana	8,955**	737**		436**	
Total: 32 patients, nine centers							

Table 1. Number of Patients with COVID-19 Placed on ECMO at Each Hospital and the Geographic Location of Each Hospital, Along with Data About the Number of Patients in Each State Diagnosed With COVID-19, Hospitalized for COVID-19 and Dead

*https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/Immunization/ncov2019.aspx.

thttps://txdshs.maps.arcgis.com/apps/opsdashboard/index.html#/ed483ecd702b4298ab01e8b9cafc8b83.

thttps://covid19.colorado.gov/case-data.

Shttps://coronavirus.dc.gov/release/coronavirus-data-april-14-2020.

¶https://www.health.pa.gov/topics/disease/coronavirus/Pages/Cases.aspx.

||https://floridahealthcovid19.gov/.

#https://health.hawaii.gov/coronavirusdisease2019/what-you-should-know/current-situation-in-hawaii/.
**https://www.coronavirus.in.gov/.

COVID-19, coronavirus disease 2019; ECMO, extracorporeal membrane oxygenation.

pink sputum was noted. The patient was isolated, tested for COVID-19, and found to be positive. She was soon reintubated and was transferred to a tertiary care center near Denver, Colorado on March 13, 2020. She developed acute respiratory distress syndrome (ARDS) and was proned. On March 19, 2020, she was placed on veno-venous ECMO. On March 22, 2020, she received her first of nine doses of compassionate use Remdesivir. On March 27, 2020, her respiratory status improved and her ECMO flow and ventilatory settings were weaned. Two days later, on March 29, 2020, after 10 days on ECMO. On April 1, 2020, she was extubated. On April 8, 2020, she was discharged from the hospital on room air and went to a rehabilitation facility for her orthopedic injury.

Discussion

Clinical guidelines for the management of patients with COVID-19 have been released by the World Health Organization (WHO)⁹ and the CDC.¹⁰ The Extracorporeal Life Support Organization (ELSO)¹¹ and The American Society for Artificial Internal Organs (ASAIO)¹² both recently published guidelines about the role of ECMO in treating patients with COVID-19. Nevertheless, the role of ECMO in the management of these challenging patients remains unclear. Here, we report on our recent initial experience in 32 severely ill COVID-19 patients with severe pulmonary compromise, some of whom also developed severe cardiac compromise. Although readily deployed, our initial experience demonstrated that 22 of 32 patients are alive (68%), with 17 of 32 (53.1%) alive on ECMO, although with only five of 15 (33.3%) surviving to date post-ECMO removal. Although our early limited experience does not allow for subgroup analysis, it is important to note that all five survivors were supported with only venoveno ECMO. Thus, the survival of patients treated with only veno-venous ECMO and separated from veno-venous ECMO is five out of 12 (41.7%)-outcomes that are reasonable in the context of contemporary use of ECMO for ARDS in adults.⁶ Our hope is that this early experience will provide information about the real-world results of ECMO in patients with COVID-19 pneumonia and facilitate decision-making at the bedside. Further, this analysis will inform and drive future research to improve outcomes. Out of this analysis and subsequent experience with ECMO, insight will arise as to the appropriate role, timing, and utility of ECMO in patients with severe COVID-19.

A recently published retrospective, single-center study included all 99 patients with confirmed COVID-19 pneumonia in Wuhan Jinyintan Hospital between January 1, 2020, and January 20, 2020.13 Patients had clinical manifestations of fever (82 patients [83%]), cough (81 patients [82%]), shortness of breath (31 patients [31%]), muscle ache (11 patients [11%]), confusion (nine patients [9%]), headache (eight patients [8%]), sore throat (five patients [5%]), rhinorrhea (four patients [4%]), chest pain (two patients [2%]), diarrhea (two patients [2%]), and nausea and vomiting (one patient [1%]). Radiographic evaluation documented that 74 patients (75%) showed bilateral pneumonia, 14 patients (14%) showed multiple mottling and ground-glass opacity, and one patient (1%) had pneumothorax. Seventeen patients (17%) developed ARDS. Oxygen therapy was used in 75 (76%), noninvasive (i.e., face mask) mechanical ventilation was used in 13 (13%), invasive mechanical ventilation was used in four (4%), continuous renal replacement therapy (CRRT) was used in nine (9%), and ECMO was used in three (3%). Eleven patients died of multiple organ failure (11%).

An additional publication from Wuhan describes 52 critically ill adult patients with COVID-19 pneumonia who were admitted to the intensive care unit (ICU) of Wuhan Jin Yin-tan hospital (Wuhan, China) between late December 2019, and January 26, 2020. Six of these patients were supported with ECMO with an alarmingly high mortality of 83% (5/6).^{14,15} Clearly, very limited published information exists about the role of ECMO in patients with COVID-19.

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	Overall
N Days from COVID diagnosis to intubation (mean [SD]) Days from COVID diagnosis to intubation (median [IQR]) Days from intubation to cannulation (mean [SD]) Days from intubation to cannulation (median [IQR]) Days on ECMO (mean [SD]) Days on ECMO (median [IQR]) Hours on ECMO (mean [SD]) Hours on ECMO (meain [IQR]) Age (median [IQR]) Age (median [IQR]) Gender (count [%])	32 2.47 (3.52) 1.00 (1.00–3.00) 4.26 (2.35) 4.00 (2.00–6.50) 7.33 (3.31) 6.00 (5.00–10.00) 166.53 (81.31) 143.00 (105.00–233.00) 52.41 (12.49) 52.41 (12.49)
Female Male	10 (31.2) 22 (68.8)
Cancer (count [%]) No Unknown Yes	28 (87.5) 1 (3.1) 3 (9.4)
Diabetes (count [%]) No Unknown Yes	20 (62.5) 1 (3.1) 11 (34.4)
Heart disease (count [%]) No Unknown Yes	27 (84.4) 1 (3.1) 4 (12.5)
Obesity (count [%]) No Unknown Yes	17 (53.1) 1 (3.1) 14 (43.8)
Asthma (count [%]) No Unknown Yes	27 (84.4) 2 (6.2) 3 (9.4)
Proned Before ECMO (count [%]) No Unknown Yes	7 (21.9) 5 (15.6) 20 (62.5)
CVVH or CRRT used (count [%]) No Unknown Yes	10 (31.2) 10 (31.2) 12 (37.5)
ECMO type (count [%]) Unknown V-A V-AV to V-V V-V V-V, VV-A V-V, VV-A V-V, VV-V Anticoagulation type (count [%])	1 (3.1) 3 (9.4) 1 (3.1) 25 (78.1) 1 (3.1) 1 (3.1)
Argatroban Heparin Unknown	2 (6.2) 28 (87.5) 2 (6.2)
Anti-interleukin-6 receptor monoclonal antibodies (count [%]) No Yes	26 (81.2) 6 (18.8)
Anti-viral medication (count [%]) No Yes	26 (81.2) 6 (18.8)
Hydroxychloroquine (count [%]) No Yes	31 (96.9) 1 (3.1)
Intravenous steroids (count [%]) Unknown Yes	27 (84.4) 5 (15.6)

COVID-19, coronavirus disease 2019; CRRT, continuous renal replacement therapy; CVVH, continuous venovenous hemofiltration; ECMO, extracorporeal membrane oxygenation; IQR, interquartile range; SD, standard deviation; V-A, veno-arterial extracorporeal membrane oxygenation; V-AV, extracorporeal membrane oxygenation with systemic venous inflow with dual systemic venous and systemic arterial outflow (*i.e.*, V-V and V-A combined); V-V, veno-venous extracorporeal membrane oxygenation; VV-A, V-A extracorporeal membrane oxygenation with dual systemic venous cannulation for inflow (typically bicaval systemic venous drainage); VV-V, V-V extracorporeal membrane oxygenation with dual systemic venous cannulation for inflow (typically bicaval systemic venous drainage).

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Table 3. Overview of Patients with COVID-19 Treated with ECMO and No Longer on ECMO

	Mortality on ECMO	Successful Wean from ECMO	р
N	10	5	
Days from COVID diagnosis to intubation (mean [SD])	0.00 (1.41)	3.00 (3.46)	0.308
Days from COVID diagnosis to intubation (median [IQR])	0.00 (-0.50 to 0.50)	1.00 (1.00–3.00)	0.135
Days from intubation to cannulation (mean [SD])	4.67 (2.08)	3.80 (2.39)	0.623
Days from intubation to cannulation (median [IQR])	4.00 (3.50-5.50)	4.00 (2.00-5.00)	0.651
Days on ECMO (mean [SD])	6.80 (3.08)	8.40 (3.85)	0.397
Days on ECMO (median [IQR])	6.00 (5.00–9.50)	8.00 (6.00–10.00)	0.497
Hours on ECMO (mean [SD])	153.40 (76.54)	192.80 (93.08)	0.396
Hours on ECMO (median [IQR])	135.50 (103.50–223.50)	170.00 (133.00–232.00)	0.54
Age (mean [SD])	56.70 (12.81)	52.80 (10.47)	0.567
Age (median [IQR])	56.70 (12.81)	52.80 (10.47)	0.567
Gender (count [%])	00110(12101)	02.00 (.0)	0.000
Female	2 (20.0)	3 (60.0)	0.333
Male	8 (80.0)	2 (40.0)	0.000
Cancer (count [%])	0 (00.0)	2 (40.0)	
No	9 (90.0)	4 (80.0)	1
Yes	1 (10.0)	1 (20.0)	1
Diabetes (count [%])	1 (10.0)	1 (20.0)	
No	5 (50.0)	3 (60.0)	1
Yes	5 (50.0)		I
	5 (50.0)	2 (40.0)	
Heart disease (count [%])	0 (00 0)	E (100 0)	4
No	9 (90.0)	5 (100.0)	1
Yes	1 (10.0)	0 (0.0)	
Obesity (count [%])		0 (10 0)	
No	5 (50.0)	2 (40.0)	1
Yes	5 (50.0)	3 (60.0)	
Asthma (count [%])		(00.0)	
No	10 (100.0)	4 (80.0)	0.714
Yes	0 (0.0)	1 (20.0)	
Proned before ECMO (count [%])			
No	3 (30.0)	1 (20.0)	0.962
Unknown	2 (20.0)	0 (0.0)	
Yes	5 (50.0)	4 (80.0)	
CVVH or CRRT used (count [%])			
No	2 (20.0)	3 (60.0)	0.782
Unknown	4 (40.0)	0 (0.0)	
Yes	4 (40.0)	2 (40.0)	
ECMO type (count [%])			
V-A	1 (10.0)	0 (0.0)	0.604
V-AV to V-V	1 (10.0)	0 (0.0)	
V-V	6 (60.0)	5 (100.0)	
V-V, VV-A	1 (10.0)	0 (0.0)	
V-V, VV-V	1 (10.0)	0 (0.0)	
Anticoagulation type (count [%])	()		
Argatroban	2 (20.0)	0 (0.0)	0.788
Heparin	8 (80.0)	5 (100.0)	
Anti-interleukin-6 receptor monoclonal antibodies (count [%l)	e (10010)	
No	9 (90.0)	3 (60.0)	0.494
Yes	1 (10.0)	2 (40.0)	0.404
Anti-viral medication (count [%])	1 (10.0)	2 (40.0)	
No	9 (90.0)	2 (40.0)	0.494
Yes	1 (10.0)	3 (60.0)	0.494
	1 (10.0)	3 (60.0)	
Hydroxychloroquine (count [%])	10 (100 0)	4 (80.0)	0 71 4
No	10 (100.0)	4 (80.0)	0.714
Yes	0 (0.0)	1 (20.0)	
ntravenous steroids (count [%])	0 (00 C)	1 (00 0)	N 1 A
Unknown	9 (90.0)	1 (20.0)	NA
Yes	1 (10.0)	4 (80.0)	

COVID-19, coronavirus disease 2019; CRRT, continuous renal replacement therapy; CVVH, continuous venovenous hemofiltration; ECMO, extracorporeal membrane oxygenation; IQR, interquartile range; NA, not available; SD, standard deviation; V-A, veno-arterial extracorporeal membrane oxygenation; V-AV, extracorporeal membrane oxygenation with systemic venous inflow with dual systemic venous and systemic arterial outflow (*i.e.*, V-V and V-A combined); V-V, veno-venous extracorporeal membrane oxygenation; VV-A, V-A extracorporeal membrane oxygenation with dual systemic venous cannulation for inflow (typically bicaval systemic venous drainage); VV-V, V-V extracorporeal membrane oxygenation with dual systemic venous cannulation for inflow (typically bicaval systemic venous drainage).

Our analysis of 32 patients from nine hospitals reveals that ECMO may play a meaningful role in salvaging select critically ill patients with COVID-19. Out of 32 consecutive patients with

COVID-19 were placed on ECMO at nine different hospitals: 17 remain on ECMO, 10 died before or shortly after decannulation, five are alive after separation from ECMO. All five

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Status of 32 COVID-19 ECMO Patients at Time of Publication



Figure 1. The current status of all 32 COVID-19 ECMO patients. COVID-19, coronavirus disease 2019; ECMO, extracorporeal membrane oxygenation.

of the survivors are extubated, and one has been discharged from the hospital on room air. Thus, five out of 15 patients (33.33%) who have been decannulated so far have survived. Our early experience seems to indicate that patients who require veno-arterial support have a poor prognosis in comparison to patients who require only veno-venous support. From our experience to date, further study will be necessary to tease out predictors of those COVID patients most likely to benefit from this therapy. Hopefully, as centers gain experience with this challenging and complex clinical problem, patient selection and outcomes will improve.

The ELSO maintains an on-line worldwide registry of COVID-19 patients treated with ECMO.¹⁶ As of April 9, 2020, in the ELSO registry, the entire global experience of COVID-19 patients supported with ECMO is 216 suspected or confirmed COVID-19 patients and 212 confirmed COVID-19 patients, with nine out of 30 (30%) discharged alive. The North American experience of COVID-19 patients supported with ECMO is 150 confirmed COVID-19 patients, with 108 patients still on ECMO and 42 patients listed as completed ECMO. These data from ELSO provide an overall snapshot of the scope of ECMO use in patients with COVID-19, both worldwide and in North

America. Our multi-institutional analysis of 32 patients with COVID-19 treated at nine hospitals provides more detailed information about the early challenges and results.

To provide guidance about the use of ECMO in severely ill COVID-19 patients, the ASAIO has recently developed and published a recommendations statement as to emerging and best practices for ECMO in COVID-19, as a living document that will be updated periodically to help fine-tune ECMO selection and best practices.¹² To compliment this recent publication, ASAIO has developed a database specific to ECMO use in severe COVID-19 to aid in this effort. Merging and synergizing data between databases such as those obtained by Specialty-Care, ELSO, and ASAIO, as well as other sources, will begin to provide insight about the relevant exposure, demographics, comorbidities, and clinical and laboratory variables that may be predictive of outcome, inform selection of patients, guide timing of initiation of ECMO, or even suggest futility.

Of note, it is also interesting that four of the five ECMO survivors received intravenous steroids. Although the decision to use steroids was determined by the individual providers, the use of steroids was discouraged in the early Chinese reports, as well as in some of the current COVID-19 management



Figure 2. The number of patients cannulated each week. COVID-19, coronavirus disease 2019; ECMO, extracorporeal membrane oxygenation.

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Distribution of Hours on ECMO by Outcome for 15 Cases



Figure 3. The distribution of hours on ECMO, comparing the survivors with the nonsurvivors. ECMO, extracorporeal membrane oxygenation.

guidelines.¹⁷ This observation clearly illustrates that there is still much to be understood about the potential therapeutic options in this extremely heterogeneous population.

The concerning poor outcomes associated with veno-arterial ECMO in patients with COVID-19 suggest that the combined COVID-19 respiratory and cardiac failure might convey an inherently poor prognosis, regardless or treatment, or that alternative support therapies might be better suited for this complex pathophysiologic scenario. For example, anecdotal unpublished experiences suggest that veno-veno ECMO might be used to support isolated respiratory failure and if the patient has concomitant cardiac failure, then targeted right or left ventricular temporary percutaneous support might be considered (*i.e.*, right ventricular or left ventricular Impella [Abiomed, Inc., Danvers, MA]).

Our initial findings also illustrate the need for further data regarding the optimal cannulation strategy in these patients. Ultimately, the situation of each patient will need to be individualized by the local team with consideration of the physiologic needs, comfort in cannulation and cannula management (especially since some of these patients might be considered for prone positioning), as well as available resources. There may be some debate regarding cannula types and locations of vascular access in patients with severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). It is important to remember that the causative pathogen of COVID-19 is highly contagious and transmitted typically via respiratory droplets/fomites (although there are some concerns for other modes of viral transmission).7 Therefore, it is critical that at the time of cannulation, strict sterile technique along with respiratory droplet isolation precautions, including negative airflow isolation, be adhered to by the cannulating and management team. Cannulation in the context of COVID-19 is performed with full airborne and droplet precautions. The cannulation team is restricted to the surgeon, one assistant, and the perfusionist and is performed in a negative pressure room. All team members must wear appropriate personal protective equipment, beyond the sterile gowns, gloves, and hats used in the operating room, including appropriate N-95 masks and full protective eye-wear.¹⁸ Ultrasound-guided access of the right internal jugular vein and right femoral vein can minimize the duration of cannulation. Avoiding the use of dual lumen bicaval cannulas will decrease the need for either TEE or fluoroscopy, each of which may unnecessarily increase exposure and time. Another potential strategy



Distribution of Age by Outcome for 15 Cases

Figure 4. The distribution of age of the patients, comparing the survivors with the nonsurvivors.

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is to position the isolated patient with the ECMO console facing towards a window so that the ECMO specialist is able to view the control panel and parameters without having to stay in the room, thereby minimizing patient contact and potential pathogen exposure.

As experience matures, a better understanding of contraindications to ECMO in COVID-19 patients is necessary and will emerge. Although there are few absolute contraindications, given the concerns for limited resources, as protocols are developing, there are concerns that advanced relative age (*i.e.*, >65 years/old), multiple comorbidities, acute or chronic endorgan failure, and recent cardiopulmonary arrest are inherently associated with a poor prognosis in COVID-19 patients placed on ECMO. Some have advocated restricting mechanical support to veno-venous rather than veno-arterial ECMO. Each patient must be considered on a case-by-case basis, with great hesitation regarding candidacy in the context of advanced age, and those comorbidities that portend a poor prognosis, including diabetes, heart disease, obesity, and especially patients with underlying terminal disease, central nervous system hemorrhage, and evidence of MSOF. Finally, many centers have adopted a policy that COVID-19 patients are not candidates for ECPR, a policy related to both poor prognosis and protection of the healthcare team.

Our multicenter experiences suggest that there is some potential role for ECMO in appropriately selected patients with COVID-19. Although the risk factors and variables that contribute to optimal outcomes are inherently complex and probably reflect individual center experiences and available resources, it can be argued that it would be unethical to withhold ECMO—or consideration for referral to an experienced ECMO center—in patients who might potentially benefit from this therapy.¹⁹

Future Directions

Much remains to be learned about the role of ECMO in these patients. From our analysis to date, no specific demographic, clinical, or laboratory data, to date, is predictive of outcome with ECMO in patients with COVID-19. Similarly, the role of multiple medications in the treatment of COVID-19 remains unclear, including intravenous steroids while on ECMO, antiviral medications (Remdesivir), anti-IL-6 receptor monoclonal antibodies (Tocilizumab, Siltuximab, or Sarilumab), and hydroxychloroquine.

Accumulating evidence suggests that a subgroup of patients with severe COVID-19 have a cytokine storm syndrome in which a cascade of activated cytokines leads to harmful autoamplifying inflammatory cytokine production.^{20,,21,,22} Termed the "cytokine storm," this response often leads to organ damage and increases the risk of death. Among COVID-19 patients which have received ECMO, a strong positive correlation exists between mortality and high cytokine levels, most notably IL-6.15,22-24 Ruan et al24 documented that IL-6 concentrations differed significantly between survivors and nonsurvivors of COVID-19, with nonsurvivors having up to 1.7-times higher values.²² Multiple therapeutic strategies might mitigate "cytokine storm," including antibody therapies (e.g., Tocilizumab, Sarilumab, Siltuximab), therapeutic plasma exchange (TPE), and even direct removal of cytokines. TPE can reduce cytokine levels by separating and removing plasma from blood and replacing the removed plasma with fresh frozen plasma.²⁵ CytoSorb (https://cytosorbents.com/products/cyto-sorb/) is an extracorporeal cytokine adsorber that has been approved in the European Union to reduce toxic levels of cytokines; this technology might be combined with ECMO to treat cytokine storm associated with sever COVID-19 pneumonia. Initial experience with this approach has recently been reported.^{26,27} It is a fact that each of these theoretical treatment options merit additional investigation.

Limitations

This analysis reports very preliminary data. Additional follow-up is required on all surviving patients. Further patient accrual will enhance continued analysis of outcomes. Although our experience is the largest published analysis of COVID-19 patients treated with ECMO to date, we recognize that our dataset is relatively small, and we plan to continue gathering data to provide additional insight as to guideposts for patient selection and predictors of outcomes. It is our hope that by sharing our experience, other centers and patients may benefit.

Conclusions

Our early experience and analysis of 32 patients from nine hospitals reveals that ECMO plays a role in the stabilization and survival of select critically ill patients with COVID-19. During 24 days, 32 consecutive patients with COVID-19 placed on ECMO at nine different hospitals: 17 remain on ECMO, 10 died before or shortly after decannulation, five are alive and extubated after separation from ECMO, and one of these five has been discharged from the hospital. Additional gathering and analysis of data will inform appropriate selection of patients and provide guidance as to best use of ECMO in terms of timing, implementation, duration of support, and best criteria for discontinuation. A tremendous amount of information still needs to be learned about the role of ECMO in treating patients these sickest of patients with COVID-19.

References

- Coronavirus COVID-19 Global Cases by the Center for Systems Science and Engineering (CSSE). Available at: https://coronavirus.jhu.edu/map.html. Accessed April 7, 2020.
- Clerkin KJ, Fried JA, Raikhelkar J, et al: Coronavirus disease 2019 (COVID-19) and cardiovascular disease. *Circulation* 2020. [Epub ahead of print]
- 3. Madjid M, Safavi-Naeini P, Solomon SD, Vardeny O: Potential effects of coronaviruses on the cardiovascular system: A review. *JAMA Cardiol* 2020. [Epub ahead of print]
- UK collaborative randomised trial of neonatal extracorporeal membrane oxygenation: UK collaborative ECMO trail group. *Lancet* 348:75–82, 1996.
- Shah SM, Kays DW, Ghazarian SR, et al: Eighteen years of paediatric extracorporeal membrane oxygenation and ventricular assist devices: Insight regarding late outcomes. Cardiol Young 28: 1316–1322, 2018.
- Peek GJ, Mugford M, Tiruvoipati R, et al; CESAR trial collaboration: Efficacy and economic assessment of conventional ventilatory support versus extracorporeal membrane oxygenation for severe adult respiratory failure (CESAR): A multicentre randomised controlled trial. *Lancet* 374: 1351–1363, 2009.
- 2009 H1N1 Pandemic (H1N1pdm09 virus). Available at: https:// www.cdc.gov/flu/pandemic-resources/2009-h1n1-pandemic. html. Accessed April 7, 2020.
- 8. Davies A, Jones D, Bailey M, et al; Australia and New Zealand Extracorporeal Membrane Oxygenation (ANZ ECMO) Influenza

Investigators: Extracorporeal membrane oxygenation for 2009 influenza A(H1N1) acute respiratory distress syndrome. *JAMA* 302: 1888–1895, 2009.

- World Health Organization: Clinical Management of Severe Acute Respiratory Infection (SARI) When COVID-19 Disease Is Suspected. Interim guidance. March 3, 2020. Available at: https://www.who.int/docs/default-source/coronaviruse/clinicalmanagement-of-novel-cov.pdf. Accessed April 7, 2020.
- Interim Clinical Guidance for Management of Patients with Confirmed Coronavirus Disease (COVID-19). Available at: https://www.cdc.gov/coronavirus/2019-ncov/hcp/clinical-guidance-management-patients.html. Accessed April 7, 2020.
- Bartlett RH, Ogino MT, Brodie D, et al: Initial ELSO guidance document: ECMO for COVID-19 patients with severe cardiopulmonary failure. ASAIO J 2020. [Epub ahead of print]
- Rajagopal K, Keller S, Akhanti B, et al: Advanced pulmonary and cardiac support of COVID-19 patients: Emerging recommendations from ASAIO - A "Living Working Document." ASAIO J 2020.
- Chen N, Zhou M, Dong X, et al: Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: A descriptive study. *Lancet* 395: 507–513, 2020.
- Yang X, Yu Y, Xu J, et al: Clinical course and outcomes of critically ill patients with SARS-CoV-2 pneumonia in Wuhan, China: A single-centered, retrospective, observational study. Lancet Respir Med 2020. [Epub ahead of print]
- 15. Henry BM: COVID-19, ECMO, and lymphopenia: A word of caution. *Lancet Respir Med* 8: e24, 2020.
- The Extracorporeal Life Support Organization (ELSO): ECMO in COVID-19. Available at: https://www.elso.org/Registry/ FullCOVID19RegistryDashboard.aspx. Accessed April 9, 2020.
- Matthay MA, Aldrich JM, Gotts JE: Treatment for severe acute respiratory distress syndrome from COVID-19. *Lancet Respir Med* 2020. [Epub ahead of print]

- Firstenberg MS, Libby M, Ochs M, Hanna J, Mangino JE, Forrester J: Isolation protocol for a COVID-2019 patient requiring emergent surgical intervention: Case presentation. *Patient Saf Surg* 2020. [Epub ahead of print]
- Abrams D, Ferguson ND, Brochard L, et al: ECMO for ARDS: From salvage to standard of care? *Lancet Respir Med* 7: 108–110, 2019.
- Chousterman BG, Swirski FK, Weber GF: Cytokine storm and sepsis disease pathogenesis. *Semin Immunopathol* 39: 517– 528, 2017.
- Kellum JA, Kong L, Fink MP, et al; GenIMS Investigators: Understanding the inflammatory cytokine response in pneumonia and sepsis: Results of the Genetic and Inflammatory Markers of Sepsis (GenIMS) Study. Arch Intern Med 167: 1655– 1663, 2007.
- Mehta P, McAuley DF, Brown M, Sanchez E, Tattersall RS, Manson JJ; HLH Across Speciality Collaboration, UK: COVID-19: Consider cytokine storm syndromes and immunosuppression. *Lancet* 395: 1033–1034, 2020.
- Risnes I, Wagner K, Ueland T, Mollnes T, Aukrust P, Svennevig J: Interleukin-6 may predict survival in extracorporeal membrane oxygenation treatment. *Perfusion* 23: 173–178, 2008.
- Ruan Q, Yang K, Wang W, Jiang L, Song J: Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients from Wuhan, China. *Intensive Care Med* 2020. [Epub ahead of print]
- Fortenberry JD, Paden ML: Extracorporeal therapies in the treatment of sepsis: Experience and promise. Semin Pediatr Infect Dis 17: 72–79, 2006.
- Harm S, Schildböck C, Hartmann J: Cytokine removal in extracorporeal blood purification: An in vitro study. *Blood Purif* 49: 33–43, 2020.
- Hartman ME, Hernandez RA, Patel K, et al: COVID-19 respiratory failure: Targeting inflammation on VV-ECMO support. ASAIO J 2020. [Epub ahead of print]