Introduction

Veno-venous extracorporeal membrane oxygenation (VV ECMO) is a rescue therapy for the most severe cases of respiratory failure. Criteria to start a patient on VV ECMO are published by ELSO (Extracorporeal Life Support Organization, Ann Arbor, MI, USA. www.elso.org) (1) and have been adopted with only slight variations by most ECMO centers. Once a patient is on VV ECMO for respiratory failure the purpose of the device is to sustain gas exchange and allow the injured lung to recover by reducing the need for applied pressures of mechanical ventilation. Ideally the process to wean a patient off from VV ECMO therapy starts once the lungs start to improve, which can be seen by improving compliance and better gas exchange (Figure 1).

The weaning process from the two modalities of ECMO, veno-arterial (VA) ECMO and VV ECMO, respectively,
are rather different (2-4). In the former both cardiac and pulmonary recovery have to be assessed before decision to decannulate. For VV ECMO the procedure is more straight-forward. Nevertheless, there are great differences in how the weaning process is managed from center to center. These differences may also be illustrated by published guidelines available on the internet (ELSO/www.elso.org) (1), with a bias towards adults and, of course, the more complicated VA ECMO.

The aim of the current overview is to describe experiences from different approaches to how weaning off from VV ECMO could be undertaken in safe and at the same time pragmatic ways.

Three experienced ECMO centers describe their approach to VV ECMO weaning in acute lung failure.

Settings

ECMO Centre Karolinska, Stockholm, became the second ECMO unit in Europe established in 1987 at a pediatric intensive care unit (PICU), and has been an ELSO Center of Excellence since 2007. The unit is dedicated to ECMO treatments, and all ages and diagnoses except post-cardiomyectomy are treated. ECMO is performed by ECMO specialists (PICU/ICU nurse) and ECMO physicians (anesthetists and intensivists). Both categories have passed an ELSO accredited course. Catheterization are performed by a cannulating surgeon. The basal staffing is one ECMO-specialist (ICU nurse), ½ ECMO Physician and ½ Nurse's aid per patient. The patient is never left alone. Approximately 90 annual ECMO runs are performed, and 55% are commenced on VA ECMO. A total of 1,200 patients have been treated up today. Integrated in the clinic is also an organization for mobile ECMO that have performed close to 900 national and international primary and secondary transports on ECMO.

After commencement of ECMO the ventilator pressures and fraction in spired oxygen over the ventilator (FiO₂) are reduced to “lung rest settings”, aiming to keep the lung open. This is also a matter of patient safety since the majority (80%) of the patients have a transfer ahead. After arrival to the ECMO ICU, FiO₂ may be in the range of 35–50%, and most patients supported with pressure control or pressure support mode depending on course of illness and patient comfort. In children, positive end expiratory pressure (PEEP) usually is set to 3–5 cmH₂O, and the driving pressure to 10–20 cmH₂O. For adults, the corresponding pressures would be 6–12 cmH₂O and 15–25 cmH₂O, respectively. Blood gases are taken every hour and the aim for oxygenation is oxygen saturation (SaO₂) >85%, and partial pressure for dissolved carbon dioxide in plasma (pCO₂) corresponding to a normal pH. Tracheostomy is performed in the adult at day two or three after which sedation is reduced to awake the patient. Pediatric and neonatal patients are not tracheostomized but sedation is reduced. Fluid balance is often an issue and most of the patients are submitted to continuous renal replacement therapy (CRRT) with fluid balance as the main indication.

The University Hospital Regensburg provides ECMO service since about 20 years and was the first German ECMO centre to receive the ELSO award of Excellence in Life support in 2014. Within the University Hospital
Regensburg ECMO is performed on different specialized ICUs: medical, cardiac surgery, and general surgical. Most cases of severe respiratory failure receiving VV ECMO are treated on the medical ICU, however. Approximately 140 annual ECMO runs are performed at the ECMO Center Regensburg, and 60% are commenced on VV ECMO. Staffing on the ICU for ECMO patients is ½ ICU nurse with experience in ECMO patient care, approximately 30 min/day of a physiotherapist and a minimum of one physician on the ward (max. 15 beds). After implementation of the ECMO, invasiveness of ventilation is reduced to diminish further ventilator-induced lung injury, however PEEP is not reduced to avoid atelectasis due to small TV. The aim for oxygenation is a partial pressure of oxygen (PaO₂) of >8 kPa [>60 mmHg] and partial pressure of carbon dioxide (PaCO₂) is adjusted to achieve a normal pH level. Arterial blood gases are routinely taken every 4 hours, and blood gases from both sides of the membrane lung (ML) are obtained at least once daily.

San Raffaele Hospital has started its ECMO activity on 2009 and the workload has been steadily growing up to 125 cases in 2016; 15% of them are in the VV mode. All patients are treated in a Cardiothoracic ICU which is staffed 24/7 with intensivists competent in extracorporeal supports. All cannulations are performed at bedsides by intensivists with transesophageal echocardiography (TEE) guidance. Dual lumen cannulation is the routine strategy since 2012. Staffing on the ICU for ECMO patients is ½ ICU nurse with experience in ECMO patient care, approximately 30 min/day of a physiotherapist and a minimum of one physician on the ward. A perfusionist is available 24/7; pre and post ML blood gases are taken at least once daily.

Results

MO Centre Karolinska

The weaning process from VV ECMO starts as soon as the tidal volumes start to recover. The ECMO blood flow has usually been reduced over the preceding days as well. The marker for progression is more to carbon dioxide clearance than oxygenation. The process is based on respiratory physiology. For sweep gas, oxygen (FiO₂ 1.0), or oxygen with a mix of carbon dioxide (0–5%) is used. The ECMO specialist adjusts the sweep gas flow accordingly to blood gas assessments and daily pre-set goals for pH and pCO₂. During the ECMO run hourly blood gases are sampled from the patient’s arterial line and/or the pre ML limb of the ECMO circuit. As the lung recovers and the clearance of CO₂ by the native lung increases, the sweep gas flow will successively be reduced. When a ML size-related minimal sweep gas flow is accomplished (adult: 2 L/min; pediatric/ neonatal: 1 L/min), incremental pCO₂ will be added to the sweep gas. The maximum fraction delivered CO₂ (FDCO₂) to the ML would correspond to 5 kPa [37.5 torr] ambient pressure.

The efficiency of the native lung to clear CO₂ is assessed by simultaneous samplings of blood gases before and after the ML. When the pCO₂ difference (pre- and post-ML) is less than 0.2–0.4 kPa (1.5–3.0 torr), the patient is regarded as “equilibrated”. It means that the ML neither clears or adds CO₂ to the patient’s blood. All CO₂ produced by the patient is thus cleared by the patient’s native lung. If the arterial SaO₂ is adequate, a trial off is performed: FiO₂ is set to 35–55%, and the sweep gas discontinued. If the procedure turns out well the sweep may be turned off for hours or overnight if necessary (till decannulation). Otherwise, the sweep gases are turned on again a new assessment will follow the next day.

University Hospital Regensburg

The weaning process starts after successful treatment of the underlying disease and improvement of lung function [FiO₂ <0.45, PEEP <10 cmH₂O, peak inspiratory pressure (PIP) <27 cmH₂O]. When the ventilator settings still allow a protective ventilation and CO₂ clearance through the native lung starts to increase the ECMO gas flow can be tapered to adjust pH values. Whenever ECMO settings are adjusted a blood gas control will be performed within 30 to 60 mins of adjustment. The extracorporeal blood flow is stepwise reduced to 1.5 L/min. Gas flow is tapered mostly in parallel to the blood flow and finally shut off for 30–60 minutes. If blood gases remain stable and the patient does not develop dyspnoea or tachypnoea the decision to remove the system is made. Depending on the current state of anticoagulation of the patient heparin treatment is stopped and activated partial thromboplastin time (aPTT), platelet count and INR are tested before the ECMO system is removed and decannulation with manual compression is carried out.

San Raffaele Hospital

After VV ECMO is implanted, the patient is kept sedated over the next 24 h in order to re-perfuse and re-establish metabolic homeostasis; respiratory rate is decreased to
10/min and TV kept at 6 mL/kg, FiO₂ is set at the lowest value in order to obtain oxygen saturation by pulse oximetry (SpO₂) >90%. Thereafter, if the patient is stable without cardiovascular impairment, sedation is stopped and the ventilator switched off, with the patient connected to a high flow continuous positive airway pressure (CPAP) with pressure set at similar values as PEEP on the ventilator. The patient can be apnoeic for a certain period of time. Once awake, dyspnoea is assessed and, if absent, we wait for full neurological recovery and eventually proceed to extubation. VV ECMO is kept at full support. Once there is clinical improvement on chest X-ray, ECMO flow is progressively reduced to 2.5–3 L/min and then sweep gas and FiO₂ tapered. After a successful ‘trial off’, anticoagulation is stopped and cannulas removed with manual compression.

In cases with septic shock and/or multiple organs dysfunction failure (MOF), the process of weaning is strongly dictated by the pattern of recovery of end organs, namely central and peripheral nervous system, and care is decided accordingly. Early tracheostomy is strongly advocated in this scenario.

If the patient succeeds to extubation, non-invasive ventilation with face mask is systematically delivered. On top of that, oral feeding and mobilization can be easily achieved.

**Discussion**

Once ECMO has been established in severe acute respiratory distress syndrome (ARDS) patients, a still open question is when and how both extracorporeal and ventilation support could be discontinued safely. It has to acknowledge that several patients die after and despite being weaned. Thus, we could argue that, if not standardized, the withdrawal criteria applied might be inappropriate and imprudent. Moreover, the re-institution of ECMO and/or mechanical ventilation, albeit infrequent, is usually associated with poor outcome. This emphasizes the relevance of proper understanding how both lung disease and general clinical conditions are evolving and, consequently, what is the potential role of ECMO for each specific patient as bridge to recovery, to transplant, or to “nowhere”, in the attempt to establish appropriateness and timing for both extracorporeal and ventilatory support discontinuation (5-6).

Differences in weaning strategies for VV ECMO are based on expert opinions rather than evidence. This fact may show that weaning can be done in many ways. While some centres keep the blood flow of the ECMO device high (>3 L/min) to prevent any additional clotting within the systems there are also centres that reduce the blood flow to a minimum of 1 L/min (adults), following the idea of reduced shear stress for the circulated blood. The blood flow per se is of no concern in VV ECMO after the sweep gas has been turned off since it is isovolumic.

Using a physiologic Karolinska approach to wean is a pragmatic way of indirectly start to “wean” as soon as the tidal volume start to recover. The CO₂ clearance capacity of the patient’s lung is continuously assessed during the recovery phase. The risk with the CO₂ approach would be carbon dioxide narcosis if the partial pressure of CO₂ in the sweep gas increase to a load above the limit of total CO₂ clearance (sum of native and ML capacity). To our knowledge this has never occurred.

In the literature including published guidelines, different strategies have been described (1-4,7) where either a trial off is performed or not. The weaning process itself does not seem to be controversial.

It remains to be understood whether the preferable approach is whether to stop ECMO or not when the patient is still mechanically ventilated. Ventilatory strategies attempting to enable spontaneous breathing and, thus, encouraging earlier weaning from mechanical ventilation could prevent the complications of sedation as well as muscle deconditioning and fatigue, which may contribute to prolong the duration of ventilatory support itself. Indeed, early extubation is an alternative briefly suggested by the ELSO guidelines and it may turn beneficial allowing to avoid the risk of ventilation-induced lung injury and ventilator-associated pneumonia, both potentially leading to exacerbation of the initial lung tissue damage.

Recently, this topic has been gaining interest (8), but the evidence available so far is still limited or specific to subsets of patients, particularly end-stage respiratory disease awaiting lung transplantation (9-12). This makes it insufficient to definitively assess whether potential benefits could balance the risks arising from perhaps longer duration of ECMO support. The idea to reduce ECMO flow to 2.5–3 L/min stems from the assumption that at this configuration the mechanical performance of the circuit would be the best without risks for stagnation and clotting, nor blood trauma.

On top of that, several adjuvant strategies have been proposed to “speed up” time of recovery, i.e., time to wean or increase the margins to come off ECMO. Levosimendan may increase the power and strength of the diaphragm (13,14), and does also seem to improve recovery to wean.
after lung transplantation (15). In sheep submitted to ECMO, triiodothyronine (T3) decrease significantly: supplementation of T3 seems to restore lactate metabolism, improve metabolic flux, and enhance diaphragm contractile function during weaning (9,16). The conversion to T3 from T4 is dependent on the seleno-enzyme deiodinase-2 (DI2). Instead of hormonal treatment, micronutrients may suffice: in critical illness, it has been shown that serum selenium decrease with a concomitant decrease of glutathione peroxidase activity (GPX). High-dose substitution of selenium restores GPX activity in both the newborn, child and adult (17-19). The physiology may be similar for the DI2 activity. Kimmoun et al. (20) described that proning may shorten time for weaning. However, echocardiography does not seem to push weaning in VV ECMO (21).

In the light of these findings, further investigations are necessary in order to standardize the weaning process from VV ECMO. Now that VV ECMO treatment is consolidated and indications for implantation have been clarified (22), the efforts should be focused on the identification of the most appropriate strategy for weaning, and its timing.

Acknowledgements

None.

Footnote

Conflicts of Interest: LM Broman, MV Malfertheiner and F Pappalardo are members of the independent research consortium European ECMO Advisory Board, the EuroELSO Workgroup for Innovative Technologies, and the Medical Advisory Board of Eurosets Medical Srl, Medolla, Italy. The other author has no conflicts of interest to declare.

References


