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High-frequency oscillatory ventilation and pediatric cardiac surgery: Yes, we can!

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See related research by Bojan et al., http://ccforum.com/content/15/5/R259

**Abstract**

In the previous issue of *Critical Care*, Bojan and colleagues reported their experiences with high-frequency oscillatory ventilation (HFOV) after pediatric cardiac surgery. A total of 120 patients were treated with HFOV on the day of surgery, thus excluding rescue HFOV use. The main finding of the authors was that the duration of mechanical ventilation was significantly shorter in patients in whom HFOV was initiated on the day of surgery. Especially interesting about this work is that the authors used HFOV when there was evidence of pulmonary hypertension or right ventricular (RV) failure in their patients. This is an interesting approach as it is often assumed that high intra-thoracic pressures increase RV afterload and thus may enhance RV dysfunction. The findings of Bojan and colleagues may be explained by the fact that they were able to decrease the pulmonary vascular resistance by finding a proper balance between atelectasis and overdistension of the lung. It can be argued that it is possible to do so by applying positive end-expiratory pressure. But, at the same time, this may coincide with the delivery of high inspiratory pressures (>30 cm H₂O). As HFOV is, in fact, a continuous positive airway pressure system, its advantage is that it is possible to maintain sufficient lung volume without large injurious pressure swings. Although the observations by Bojan and colleagues need to be confirmed in a prospective randomized trial, they have provided arguments not to rule out the early use of HFOV in pediatric cardiac surgery patients.

High-frequency oscillatory ventilation (HFOV), as used by Bojan and colleagues, is, at least in theory, an ideal tool for lung-protective ventilation as it allows effective pulmonary gas exchange with the delivery of a very small tidal volume (Vᵢ) below dead space and diminished risk of atelectrauma [1,2]. Numerous animal and clinical studies have shown clearly that mechanical ventilation (MV) itself can initiate or exacerbate lung injury, termed ventilator-induced lung injury. The application of a Vᵢ and the repetitive opening and closure of alveoli have been identified as important pathophysiological mechanisms [3].

Until now, only two randomized controlled trials investigating the effect of HFOV on patient outcome have been performed [4,5] and the larger of these included only 58 children [4]. The main finding of this study was that HFOV did not significantly improve survival (66% with HFOV versus 59% with conventional MV) or total ventilator days (20 ± 22 days with HFOV versus 22 ± 17 days with conventional MV) [4]. As a consequence, HFOV is not universally employed in pediatric critical care. Also, it seems very unrealistic that any new pediatric HFOV trial will be initiated within the next few years. This means that well-designed observational studies are needed to shed light on some of the many other aspects of pediatric HFOV which remain to be explored, including the identification of patients who are most likely to benefit from HFOV, the timing of HFOV (early versus rescue), optimal oscillator settings, and monitoring during HFOV.

In the previous issue of *Critical Care*, Bojan and colleagues [1] reported their experiences with HFOV after pediatric cardiac surgery. A total of 120 patients were treated with HFOV on the day of surgery, thus excluding rescue HFOV use. Patients were transitioned to HFOV when hypoxemia and acidosis occurred despite increasing alveolar ventilation on conventional MV (if Vᵢ exceeded 10 mL/kg) or when there was evidence of pulmonary hypertension and right ventricular (RV) failure. The main finding was that the duration of MV was significantly shorter in patients in whom HFOV was
initiated on the day of surgery. This is a very positive finding in a relatively large sample size.

However, there are some concerns related to the methodology of the study and the practical use of HFOV in the authors’ institution. Some of the limitations inherent to their work have been properly addressed. But their study was designed as a retrospective one introducing confounding by indication. Hence, the authors have calculated a propensity score to minimize this, although the variables chosen for calculating it raise some concern. Included were all demographic and post-operative variables that yielded a P value of less than 0.10 in the univariate analysis between patients with HFOV and those without HFOV. However, it seems more rational to define, beforehand, which variables should be used for calculating the propensity score. As a consequence of this approach, hemodynamic and ventilator parameters have not been used. If the authors had taken these into account when calculating their propensity score, the results of the study might have been even more firm. Also, it is not customary in their institution to routinely use an open study might have been even firmer. Also, it is not customary in their institution to routinely use an open thoracic approach, hemodynamic and ventilator parameters have not been used. If the authors had taken these into account when calculating their propensity score, the results of the study might have been even more firm.

It is important to stress that the findings of Bojan and colleagues warrant prospective confirmation in a well-designed randomized controlled trial. In the mean time, the authors are to be congratulated for their work as their study has provided arguments not to rule out the early use of HFOV in pediatric cardiac surgery patients.

Abbreviations
HFOV, high-frequency oscillatory ventilation; MV, mechanical ventilation; PaO₂, arterial partial pressure of oxygen; PEEP, positive end-expiratory pressure; PVR, pulmonary vascular resistance; RV, right ventricular; V₁, tidal volume.

Competing interests
The author declares that he has no competing interests.

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