



Editorial

What is the most appropriate spontaneous breathing trial before extubation in ICU ventilated patients?



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Mechanical ventilation, which is the most used organ replacement therapy in intensive care unit (ICU), is a life-saving intervention. In the ICU, the timing of liberation from invasive mechanical ventilation is an important issue for clinicians treating critically ill intubated patients receiving invasive mechanical ventilation [1–3]. After resolution of acute illness, or in other words when the resolution or improvement of the condition leading to intubation are done, mechanically ventilated patients are liberated from the ventilator, a process termed “weaning” from mechanical ventilation [4–6]. Weaning and extubation, though following each other in clinical practice, are two separate processes that pose distinct problems. Separating the ventilator from the patient (weaning) and removing the tracheal tube (extubation) are challenging issues [1]. Indeed, if the patient remains intubated too long, complications associated with prolonged invasive mechanical ventilation may appear [7]. On the other hand, if the patient is extubated too early, reintubation, which could occur in 5 to 25% of cases, is associated with higher morbidity and mortality [1–8].

Before extubation, it is recommended to perform a “weaning test”, which consists of a “spontaneous breathing trial” (SBT) [5,9]. The SBT aims to evaluate how the critically ill patient is able to breath “alone” with no ventilatory assistance. In the standard of care, two main weaning tests are suggested for non-selected adult patients: a T-piece trial (oxygen supply without positive pressure) and low pressure support ventilation (PSV), with a low level of PSV without positive end expiratory pressure (PEEP), from 5 to 7 cmH₂O, to compensate for the imposed workload due to the endotracheal tube and ventilator circuit [5]. In case of presence of a heated and moisture exchanger, the low level of PSV set should

take into account the imposed workload due to the additional dead space, which is evaluated from additional 2 to 5 cmH₂O PSV level.

Although these two weaning tests are not equivalent in terms of work of breathing (WOB) [10], no study has yet assessed the risk of extubation failure. Therefore, both weaning trials are recommended to assess whether a patient is ready to be extubated [5,11]. Moreover, the duration of the weaning test is also a subject of controversy ranging from a “short” 30-minute to a “long” 120-minute period.

In a recent Spanish clinical trial, Subira et al. [12] compared two different SBT techniques in 1153 mechanically ventilated patients in 18 ICUs. One was a “long” 120 minutes T-piece SBT (“highly demanding”) and the other was a “short” 30 minutes low-PSV SBT (“less demanding”). In both groups, patients who successfully completed their SBT were extubated. The authors reported that patients in the 30 minutes low-PSV SBT group were more likely to be extubated compared to patients in the 120 minutes T-piece SBT group. Successful extubation occurred in 82% of patients in the low-PSV group and in 74% of patients in the T-piece group ($P = 0.001$). Worthy of note is that patients in the PSV SBT group did not experience a significant higher reintubation rate in the 72 hours after extubation (11% vs. 12%, $P = 0.63$). No significant difference was shown for ICU length of stay (median 9 days vs. 10 days; $P = 0.69$) or hospital length of stay (median 24 days vs. 24 days ($P = 0.45$)). In comparison with patients in the T-piece group, those in the low-PSV group had significantly lower hospital mortality (10% vs. 15% [$P = 0.02$]) and 90-day mortality (13% vs. 17% [$P = 0.04$]).

The authors [12] should be congratulated for performing the largest randomised clinical trial ever conducted comparing the two common SBT techniques (T-piece vs. low PSV). They add a new piece to the puzzle of the weaning process in the overall non-selected patients. Their findings suggest that most patients can be tested for a short 30-minute period using a so-called “low-PSV” level of 8-cmH₂O and zero PEEP. Moreover, using the low-PSV SBT add a safety process as the patient is continuously monitored for the main physiological respiratory variables (respiratory rate, tidal volume, minute ventilation. . .).

Future research should evaluate which additional prophylactic strategies could be associated to the optimal SBT, with the aim to increase the success rate of extubation [13]. It could be proposed to evaluate:

- the optimal analgesia-sedation protocol as the weaning process of both analgesia-sedation and ventilator weaning are always linked [7];
- the use of reconnection to ventilator before extubation that was associated with reduced reintubation rate at 48 hours in a recent study [14] (reconnection to ventilator was made in 25% of included patients in the Subira study [12]);
- prophylactic non-invasive ventilation after extubation [15,16] (prophylactic non-invasive ventilation was performed in 9% of the low-PSV group and in 6% of the T-piece group in the Subira study [12]);
- the use of prophylactic high-flow nasal oxygen cannula after extubation [17–19] (which was made in 16% of the low-PSV group and in 13% of the T-piece group in the Subira study [12]).

In summary, the Subira study [12] suggests that for the majority of “easy to wean patients”, a short 30-minute period using a so-called “low-PSV” level of 8-cmH₂O and zero PEEP could be proposed as first line approach before extubation as SBT test.

Disclosure of interest

Pr. Jaber reports receiving consulting fees from Dräger, Fisher & Paykel and Fresenius. The other authors declare that they have no competing interest.

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