POINT OF VIEW

The American-European Consensus Conference definition of the acute respiratory distress syndrome is dead, long live positive end-expiratory pressure!

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KEYWORDS
Acute respiratory distress syndrome; Positive end-expiratory pressure; Standard ventilatory settings; American-European Consensus Conference; PaO2/FiO2 ratio

Abstract In 1994, an American-European Consensus Conference (AECC) formalized the criteria for the diagnosis of the acute respiratory distress syndrome (ARDS). Although that definition is simple to apply in the clinical setting, it has been challenged over the years in several studies since the assessment of the oxygenation defect does not require standardized ventilatory support. We were the first to propose new guidelines, based on a specific, standard method of evaluating oxygenation status, a proposal that was later advocated by others. To address the limitations of the AECC definition, a modified ARDS definition has been proposed by a task force panel of experts, referred to as the Berlin Definition, using a terminology similar to that we previously proposed. However, that proposal has several methodological flaws. Since all ARDS patients start off with terrible oxygenation, the Berlin Definition offers no room for stratifying and identifying true ARDS patients since there is no further re-evaluation of the hypoxemia under standard ventilator setting in a specific time period. In this Point of View, we review the history of the definition of ARDS and discussed the methodological concerns regarding adopting this new, revised ARDS definition.

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Introduction and historical remarks

In August 1967, Ashbaugh et al. described for the first time a syndrome that they termed the ARDS. They studied a cohort of 272 patients who were receiving respiratory support, and from this cohort they identified 12 patients with a syndrome that was similar to the Infant Respiratory Distress Syndrome. The respiratory distress was defined as sudden, catastrophic, and often associated with a multi-organ system insult which led to tachypnea, hypoxemia, decreased respiratory system compliance, and bilateral pulmonary infiltrates on chest X-ray due to non-cardiogenic pulmonary edema. The mortality rate was 58% and on pathological examination the lungs of the non-survivors were heavy, atelectatic, with interstitial and alveolar edema, and hyaline membranes. Since that time, the hallmark of this syndrome has included: (i) a risk factor for the development of ARDS, (ii) severe hypoxemia with a relatively high FiO2, (iii) bilateral pulmonary infiltrates, and (iv) no clinical evidence of cardiogenic pulmonary edema, although acute lung injury (ALI) resulting in ARDS can also occur in the setting of left ventricular failure.

ARDS is caused by an inflammatory insult to the alveolar-capillary membrane that results in increased permeability and subsequent interstitial and alveolar edema. Unlike other forms of acute respiratory failure and like all forms of tissue inflammation, ALI during ARDS represents a complex process in which multiple cellular signaling pathways propagate or attenuate lung damage. Since it is difficult to measure changes in capillary and alveolar permeability at the bedside, diagnosis of ARDS is based on a combination of clinical, oxygenation, hemodynamic and radiographic criteria. These criteria allow the inclusion of a heterogeneous group of critically ill patients since various types of injury can lead to a similar pulmonary response. The original description of ARDS was incapable of identifying a uniform group of patients. Several patients from the original cohort would not be classified as ARDS today, since fluid overload was an important etiological factor. Thus, a precise definition is important for accurate identification and quantification of various aspects of the underlying pathophysiology and to select the best therapeutic approach in selected subgroups of patients.

Given that severe hypoxemia is the hallmark of ARDS, hypoxemia is crucial to the assessment of the severity of ARDS, for predicting the evolution in any given patient, and for assessing the response to treatment. In order to better characterize the severity of lung damage, the AECC defined ALI and ARDS as follows: (i) acute and sudden onset of severe respiratory distress; (ii) bilateral infiltrates on frontal chest radiograph; (iii) the absence of left atrial hypertension (a pulmonary capillary wedge pressure <18 mmHg or no clinical signs of left ventricular failure); and (iv) severe hypoxemia (assessed by the PaO2/FiO2 ratio). According to these guidelines, ALI exists when the PaO2/FiO2 ratio is ≤300 and >200 mmHg regardless of the PEEP and FiO2, and ARDS when the PaO2/FiO2 ratio is ≤200 mmHg again regardless of PEEP and FiO2. Although this definition formalized the criteria for the diagnosis of ARDS and is simple to apply in the clinical setting, it has been challenged over the years in several studies. All patients start off with terrible oxygenation and there is little room for stratifying the patients if there is no further re-evaluation of the hypoxemia. Also, the physiological thresholds of the AECC definition do not
require standardized ventilatory support. The use of PEEP can improve oxygenation indices sufficiently to change the physiology of the lung converting patients meeting the definition of ARDS to patients not meeting the ARDS definition. Therefore, a patient could fit the ARDS criteria when the PaO\textsubscript{2} is measured with zero PEEP but not when measured at a PEEP of 5 or 10 cmH\textsubscript{2}O, or when measured on FiO\textsubscript{2} > 0.35 but not when measured on FiO\textsubscript{2} = 0.5. \textsuperscript{2,3,5,11} These findings illustrate the major problems experienced when trying to compare the results of various clinical trials evaluating ventilatory strategies, since patients with very different levels of lung dysfunction and disease may have been enrolled.

An early PEEP/FiO\textsubscript{2} trial identifies different degrees of ARDS severity

In 1999, Villar et al.\textsuperscript{3} proposed the need for different guidelines, based on a specific, standard method of evaluating oxygenation status, a proposal that was later advocated by others. \textsuperscript{11} In order to determine the impact of various PEEP and FiO\textsubscript{2} levels on the stratification of patients meeting the AECC ARDS definition, Villar et al.\textsuperscript{3} evaluated the impact of standard ventilation settings applied on the day patients met the AECC ARDS criteria and 24 h later. They studied 170 patients and found that only 58% of them fulfilled ARDS criteria when evaluated on PEEP \geq 10 cmH\textsubscript{2}O and FiO\textsubscript{2} \geq 0.5 at 24 h after meeting the AECC ARDS definition. The ICU mortality of those patients was 46%. By contrast, 32% of patients were classified as having ALI (their mortality was 20%), and 10% of patients had a PaO\textsubscript{2}/FiO\textsubscript{2} > 300 mmHg and were simply categorized as having acute respiratory failure (their ICU mortality was 6%). This study demonstrated the large variability in the severity of lung damage in patients who initially meet the AECC definition of ARDS and the strong correlation between oxygenation impairment at 24 h after ARDS onset and ICU outcome. The major implication of these findings is that the use of the AECC ARDS definition to enroll patients into clinical trials may result in the inclusion of patients with highly variable severity of lung injury and mortalities. If the subjects in a trial have a very low risk of the condition that the intervention is hypothesized to prevent, the trial – regardless of sample size – will not verify the value of the intervention. \textsuperscript{12} For example, in the ARDSnet trial, \textsuperscript{13} 17% of enrolled patients did not have ARDS but were included in the overall analysis. Consequently, it can be argued that the ARDSnet trial failed to focus on the highest risk patients.

A PEEP and FiO\textsubscript{2} trial 24 h after ARDS onset is an easy and simple strategy to find or identify subpopulations of ARDS patients at highest risk. \textsuperscript{4} It is not difficult to explain why patients initially meeting AECC ARDS criteria would fail to meet the same criteria on standard ventilator setting. It is well established that changes in PEEP and FiO\textsubscript{2} alter the PaO\textsubscript{2}/FiO\textsubscript{2} values in lung-injured patients. \textsuperscript{14,15} PEEP can improve oxygenation sufficiently to change the physiology of the lung. \textsuperscript{15} If PEEP is inadequate, the lung collapses during expiration allowing alveolar derecruitment and causing ventilator-induced lung injury. The lack of a standard PEEP and FiO\textsubscript{2} setting in the AECC definitions may explain the negative results of published trials of various therapeutic interventions on patients with ALI/ARDS. \textsuperscript{16-18} Few of these negative trials used the AECC definition for ARDS, and none of the trials identified ARDS subtypes for test purposes.

The “Berlin Definition”

A proposal for an update of the AECC ARDS definition has been published recently\textsuperscript{6} by a task force panel of
experts using a similar terminology as we had previously reported. Using teleconferencing, in-person discussions and retrospective data, they proposed an ARDS classification with three severity categories (mild, moderate, and severe) for empirical evaluation. The term “mild” ARDS was used for defining those patients who are considered as ALI in the AECC definition (300 ≥ PaO₂/FIO₂ > 200 mmHg). The term “moderate” was used for patients with a PaO₂/FIO₂ ≥ 100 mmHg but <200, and the term “severe” for those with a PaO₂/FIO₂ < 100 mmHg. The panel used 7 datasets: 4 from multicenter studies (enrolling 4188 patients with a PaO₂/FIO₂ < 300 mmHg), and 3 from small, single-center studies (enrolling 269 patients). By considering only those patients from the multicenter studies who were managed with PEEP ≥ 5 cmH₂O at the time of study enrollment, the panel found that hospital mortality increased with every stage of severity (mild 27%, moderate 32%, severe 45%). In the database from the 3 small, single-center studies comprising 269 patients, the hospital mortality increased as well with every stage of ARDS (mild 20%, moderate 41%, severe 52%). Although the authors stated that the purpose of their empirical definition was not to develop a prognostic tool, this exercise should be cautiously generalized for the following methodological reasons.

First, none of the patients included in the empirical analysis were prospectively enrolled for the purpose of revising the ARDS definition and/or evaluating risk stratification. Second, the categorization of patients was done based on the PaO₂/FIO₂ value at the time of inclusion into their respective observational study or randomized clinical trial. There is no information on whether those baseline values of PaO₂/FIO₂ were calculated at the time of ARDS onset or during any time of their disease process, or whether the PaO₂ was measured under a known FIO₂ and PEEP level. In our studies, PaO₂/FIO₂ was always calculated from the PaO₂ values measured 30 min after each standard ventilator setting under a specified FIO₂ and PEEP level. Third, 24% of patients included had a PaO₂/FIO₂ > 200 at the time of enrollment (classified as non-ARDS by the AECC definition). We did not include those patients in our studies because in many centers they are usually not treated with endotracheal intubation and invasive MV. However, we have reported that in about 40% of ARDS patients, PaO₂/FIO₂ increased above 200 mmHg when they were evaluated on PEEP > 10 and FIO₂ ≥ 0.5 at 24 h after being diagnosed as having ARDS according to the AECC definition. Fourth, the empirical definition did not consider the level of FIO₂ for PaO₂/FIO₂ categorization despite the fact that changes in the applied FIO₂ results in changes in PaO₂/FIO₂. In addition, since it is likely that a significant proportion of patients included in those multicenter studies were on FIO₂ ≤ 0.4 at the time of study enrollment, there is no information on how many patients could not meet ARDS criteria if they were evaluated at a minimum level of FIO₂ = 0.5. Fifth, 518 patients were eliminated from the empirical analysis because PEEP was missing or was <5 cmH₂O. In our prospective studies, we did not exclude any patient based on the baseline PEEP or FIO₂ at the time of ARDS diagnosis. Based on the wide range of FIO₂ and PEEP levels at the time of AECC ARDS diagnosis, we found that a standard ventilator setting can improve oxygenation sufficiently to convert patients meeting the AECC ARDS definition on PEEP < 5 cmH₂O and/or on FIO₂ ≤ 0.4 to non-ARDS patients. Sixth, since there was no standardization of ventilator settings for measuring PaO₂, and since more than 50% of patients were on PEEP < 10 cmH₂O at baseline, the experts’ panel only selected a unique level of PEEP (≥5 cmH₂O) as a requirement for the proposed empirical ARDS definition. We had found that when all patients are evaluated under the same standard ventilator settings, only the evaluation under a PEEP > 10 reached statistical significance for association of PaO₂/FIO₂ category with ICU mortality. Seventh, among other considerations (strict entry criteria for the trials), those studies do not include all consecutive ARDS patients admitted into the ICU since patients with pre-existent chronic diseases, patients over a certain age, and patients with established organ dysfunction were excluded. Eighth, the 4 multicenter studies were a case-mix of observational studies and clinical trials performed from 1996 to 2000, when patients were ventilated with tidal volumes ≥10 ml/kg predicted body weight (currently considered as an injurious tidal volume) and low levels of PEEP, and studies performed after the year 2000 (when patients were ventilated with lower tidal volumes). It has been postulated that the development of ARDS should have decreased because of advances in supportive care, particularly the application of protective mechanical ventilation. Ninth, there is a great discrepancy between the number of patients included in the database from the three-single centers and the number of patients reported in the original publications. We have re-examined those publications and have noted that they only studied 75 non-consecutive, selected ALI/ARDS instead of the 269 that are claimed by the experts’ panel. Tenth, the experts’ panel failed to evaluate their patients at 24 h after ARDS diagnosis. There is no information about the number of patients who did not meet ARDS criteria at 24 h of study entry under the same arbitrary condition (PEEP ≥ 5). Taking into consideration all the above, and based on our previous studies, we postulate that at least 50% of those patients would have a PaO₂/FIO₂ > 200 mmHg, and at least 25% would have a PaO₂/FIO₂ > 300 mmHg 24 h after ARDS identification on a PEEP ≥ 10 and FIO₂ ≥ 0.5.

In conclusion, we need more specific guidelines based on a standard method of evaluating oxygenation status (i.e., a specific level of PEEP and FIO₂) in order to properly classify the severity of patients with and as ARDS. As we have reported in the several studies discussed in this review, a large variability in the severity of lung damage exists in patients meeting the AECC definition of ARDS and a strong correlation exists between oxygenation impairment at 24 h after ARDS onset and ICU outcome. The judicious use of PEEP, FIO₂, and tidal volume in critically ill patients in the last decade has made ARDS a rare syndrome in today’s modern ICUs. However, no matter how infrequently we observe its presence we need to be able to properly classify its severity. The AECC ARDS definition is dead, long live PEEP!

**Funding**

Supported by Instituto de Salud Carlos III, Spain (PI 10/0393) and the Asociación Científica Pulmón y Ventilación Mecánica.
Conflict of interest

The authors have no conflict of interest to declare.

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