Clinical review: Scoring systems in the critically ill

Jean-Louis Vincent*1 and Rui Moreno2

Abstract
General illness severity scores are widely used in the ICU to predict outcome, characterize disease severity and degree of organ dysfunction, and assess resource use. In this article we review the most commonly used scoring systems in each of these three groups. We examine the history of the development of the initial major systems in each group, discuss the construction of subsequent versions, and, when available, provide recent comparative data regarding their performance. Importantly, the different types of scores should be seen as complementary, rather than competitive and mutually exclusive. It is possible that their combined use could provide a more accurate indication of disease severity and prognosis. All these scoring systems will need to be updated with time as ICU populations change and new diagnostic, therapeutic and prognostic techniques become available.

Introduction
Scoring systems used in critically ill patients can be broadly divided into those that are specific for an organ or disease (for example, the Glasgow Coma Scale (GCS)) and those that are generic for all ICU patients. In this article, we focus on the generic scores, which can broadly be divided into scores that assess disease severity on admission and use it to predict outcome (for example, Acute Physiology and Chronic Health Evaluation (APACHE), Simplified Acute Physiology Score (SAPS), Mortality Probability Model (MPM)), scores that assess the presence and severity of organ dysfunction (for example, Multiple Organ Dysfunction Score (MODS), Sequential Organ Failure Assessment (SOFA)), and scores that assess nursing workload use (for example, Therapeutic Intervention Scoring System (TISS), Nine Equivalents of Nursing Manpower Use Score (NEMS)).

The objective of this review is to give the intensivist without any particular knowledge or expertise in this area an overview of the current status of these instruments and their possible applications. For a more detailed explanation of the development, application and limitations of these models, the reader is referred to a recent review [1].

Outcome prediction scores
The original outcome prediction scores were developed more than 25 years ago to provide an indication of the risk of death of groups of ICU patients; they were not designed for individual prognostication. Patient demographics, disease prevalence, and intensive care practice have changed considerably since [2], and statistical and computational techniques have also progressed. As a result, all three of the major scores in this category have been recently updated to ensure their continued accuracy in today’s ICU (Table 1).

Acute Physiology and Chronic Health Evaluation
The original APACHE score was developed in 1981 to classify groups of patients according to severity of illness and was divided into two sections: a physiology score to assess the degree of acute illness; and a preadmission evaluation to determine the chronic health status of the patient [3]. In 1985, the original model was revised and simplified to create APACHE II [4], now the world’s most widely used severity of illness score. In APACHE II, there are just 12 physiological variables, compared to 34 in the original score. The effects of age and chronic health status are incorporated directly into the model, weighted according to their relative impact, to give a single score with a maximum of 71. The worst value recorded during the first 24 hours of a patient’s admission to the ICU is used for each physiological variable. The principal diagnosis leading to ICU admission is added as a category weight so that the predicted mortality is computed based on the patient’s APACHE II score and their principal diagnosis at admission. The reason for ICU admission is, therefore, an important variable in predicting mortality, even when previous health status and the degree of acute physiological dysfunction are similar.

APACHE III was developed in 1991 [5] and was validated and further updated in 1998 [6]. Equations for
predicting risk-adjusted ICU length of stay were also developed using the APACHE III model [7]. Most recently, APACHE IV was developed using a database of over 100,000 patients admitted to 104 ICUs in 45 hospitals in the USA in 2002/2003, and remodeling APACHE III with the same physiological variables and weights but different predictor variables and refined statistical methods [8]. APACHE IV again provides ICU length of stay prediction equations, which can provide benchmarks for the assessment and comparison of ICU efficiency and resource use [9].

Simplified Acute Physiology Score
SAPS, developed and validated in France in 1984, used 13 weighted physiological variables and age to predict risk of death in ICU patients [10]. Like the APACHE scores, SAPS was calculated from the worst values obtained during the first 24 hours of ICU admission. In 1993, Le Gall and colleagues [11] used logistic regression analysis to develop SAPS II, which includes 17 variables: 12 physiological variables, age, type of admission, and 3 variables related to underlying disease. The SAPS II score was validated using data from consecutive admissions to 137 ICUs in 12 countries [11].

In 2005, a completely new SAPS model, the SAPS 3, was created. Complex statistical techniques were used to select and weight variables using a database of 16,784 patients from 303 ICUs in 35 countries [12]. The SAPS 3 score includes 20 variables divided into three subscores related to patient characteristics prior to admission, the circumstance of the admission, and the degree of physiological derangement within 1 hour (in contrast to the 24-hour time window in the SAPS II model) before or after ICU admission. The total score can range from 0 to 217. Unlike the other scores, SAPS 3 includes customized equations for prediction of hospital mortality in seven geographical regions: Australasia; Central, South America; Central, Western Europe; Eastern Europe; North Europe; Southern Europe, Mediterranean; and North America. It should be noted that the sample size for development of some of these equations was relatively small, which may compromise their prognostic accuracy. The SAPS 3 score has been shown to exhibit good discrimination, calibration, and goodness of fit [12]. SAPS 3 has also been used to examine variability in resource use between ICUs using the standardized resource use parameter based on the length of stay in the ICU adjusted for severity of acute illness [13].

Mortality Probability Model
The first MPM, developed from data from patients in one ICU, consisted of an admission model using seven admission variables, and a 24-hour model using seven 24-hour variables [14]. A revised MPM, MPM II, was
developed in 1993 using logistic regression techniques on a large database of 12,610 ICU patients from 12 countries [15]. MPM II also consists of two scores: MPM₀, the admission model, which contains 15 variables; and MPM₄₂₄, the 24-hour model, which contains 5 of the admission variables and 8 additional variables and is designed for patients who stay in the ICU for more than 24 hours. Unlike the APACHE and SAPS systems where variables are weighted, in MPM II each variable (except age, which is entered as the actual age in years), is designated as present or absent and given a score of 1 or 0 accordingly. A logistic regression equation is then used to provide a probability of hospital mortality. The authors also developed a Weighted Hospital Days scale (WHD-94) by subjectively assigning weights to days in the ICU and to hospital days after ICU discharge from the first ICU stay, and an equation to predict an ICU’s mean WHD-94, thus providing an index of resource utilization [16].

MPM has recently been updated using a database of 124,885 patients from 135 ICUs in 98 hospitals (all in North America except for one in Brazil) collected in 2001 to 2004 [17]. MPM₀-III uses 16 variables, including 3 physiological parameters, obtained within 1 hour of ICU admission to estimate mortality probability at hospital discharge; the MPM₀ characterization is, therefore, based on patient condition largely before ICU care begins. The WHD-94 predictive equation has also been updated [18].

Discussion

Several studies have compared the different outcome prediction scoring systems. For example, in a study of 10,393 patients from Scottish ICUs, Livingston and colleagues [19] compared the APACHE II and III, an APACHE II using United Kingdom-derived coefficients (UK APACHE II), SAPS II, and MPM₀ and MPM₄. These authors reported that all models showed good discrimination, although observed mortality was significantly different from that predicted by all models. SAPS II had the best performance overall, but APACHE II had better calibration. In a retrospective study of 11,300 patients from 35 hospitals in California, Kuzniewicz and colleagues [20] recently used logistic regression to re-estimate the coefficients for the APACHE IV, MPM₀-III and SAPS II scores and applied the new equations to assess risk-adjusted mortality rates. These authors noted that discrimination and calibration were adequate for all models, with discrimination of APACHE IV slightly better than that of the other two scores (area under the receiver operating characteristic curve 0.892 for APACHE IV, 0.873 for SAPS II, and 0.809 for MPM₀ III, \( P < 0.001 \)).

In addition to using a more geographically heterogeneous database for development, the SAPS 3 model attempted to address any geographic variation by providing separate customized equations of different geographical regions. Nevertheless, local customization may still help improve the calibration of these scores in individual countries or regions as demonstrated for the APACHE III in Cleveland, Ohio [21], or more recently for the SAPS 3 score in Austria [22]. In a retrospective analysis of prospectively collected data from a surgical ICU, Sakr and colleagues [23] reported that the discriminative ability of SAPS 3 was similar to that of APACHE II and SAPS II (area under the receiver operating characteristic curve 0.80 for APACHE II, 0.83 for SAPS II, and 0.84 for SAPS 3). All three scores had poor calibration, which improved after customization to the local population. In the UK, investigators have developed a new scoring system specifically for use in UK ICU patients [24]. This score uses elements of the APACHE, SAPS, and MPM systems and was developed using the large Intensive Care National Audit and Research Centre (ICNARC) database and calibrated for adult critically ill patients admitted to ICUs in the UK. It performed better than SAPS II, APACHE II and III, and MPM II [24], but has not been compared to the latest versions of these scores.

When using these instruments, in addition to the issues related to local customization and regular updates discussed above, a few important limitations should be kept in mind. First, all general outcome prediction models can only at their best predict the behavior of a group of patients that exactly matches the patients in the development population. For example, the APACHE and MPM scores were largely based on North American populations and the SAPS score on European patients, while SAPS 3 developers used a database that included a geographically more heterogeneous group of patients [12]. In addition, in most of the scores, specific populations were excluded from the original databases (for example, patients with burns, patients aged less than 16 or 18 years, patients with a very short length of ICU stay, and so on).

Second, the accuracy of any scoring system is highly dependent on the quality of the input. To be used correctly, the definitions, time of data collection, rules for missing data, and so on must exactly match those applied when building the model. The reported reliability of the systems (intra- and inter-observer) must also be taken into account.

Third, there is an inherent bias in many of the derived equations used to predict mortality in that they are created from a limited population of patients from ICUs that are specifically interested in measuring (and improving) ICU performance.

Fourth, the outcome used in all these instruments is the vital status at hospital discharge; consequently, the use of other outcome measures (such as the vital status at
ICU discharge) will compromise the accuracy of the predictive equations. Nevertheless, some models have additional equations to assess use of resources, usually measured as risk-adjusted, weighted, ICU- or hospital days [9,13,18].

Fifth, the statistical methodology used to assess calibration of a predictive model, most commonly the Hosmer-Lemeshow statistic, may be influenced by various factors, including the number of covariates being assessed, the manner in which observations with equal probabilities of outcome are sorted, and the sample size (both small and large) [25]. Interpretation of the accuracy of predictive models should, therefore, include some knowledge of the statistical tests used. Different statistical techniques may be required for the larger models increasingly used to develop predictive models, such as the use of calibration graphs and, more recently, the Cox test of calibration and related statistics [26].

Sixth, despite the fact that predictive models have been developed in large populations, in almost all cases when they are applied to new populations calibration deteriorates, although discrimination hardly changes. Two recent examples of this effect were given in validation studies of SAPS 3 in Austria and in Italy [22,27].

Seventh, the use of automatic patient data management systems can, by changing the sampling rate for the physiological variables, change the accuracy of the model. Bosman and colleagues [28] reported that predicted mortality was greater with data management charting than with manual charting for APACHE II, SAPS II, and MPM II.

Organ dysfunction scores
Organ failure scores are primarily designed to describe the degree of organ dysfunction rather than to predict survival. The severity of organ dysfunction varies widely among individuals and within an individual over time and organ failure scores must be able to take both time and severity into account. Many organ dysfunction scores have been developed over the past few decades, but we will limit our discussion to three of the scores most commonly used in general ICU patients: the Logistic Organ Dysfunction System (LODS) [29], MODS [30], and SOFA [31] (Table 2).

Logistic Organ Dysfunction Score
The LODS was developed using a database of 13,152 admissions to 137 ICUs in 12 countries [29]. Using multiple logistic regression, 12 variables were selected to represent the function of six organ systems (neurologic, cardiovascular, renal, pulmonary, hematologic, hepatic). The worst value for each variable in the first 24 hours of admission is recorded, and for each system, a score of 0 (no dysfunction) to 5 (maximum dysfunction) is awarded.

LOGIC REGRESSION
Variables used to assess organ dysfunction:
- Neurologic
  - Glasgow Coma Scale
- Cardiovascular
  - Heart rate, systolic blood pressure
- Renal
  - Serum urea or urea nitrogen, creatinine, urine output
- Respiratory
  - PaO\textsubscript{2}/FiO\textsubscript{2} ratio, mechanical ventilation
- Hematologic
  - White blood cell count, platelet count
- Hepatic
  - Serum bilirubin, prothrombin time

Logistic Organ Dysfunction Score (LODS)

| Characteristics                          | LODS [29]                  | MODS [30]                  | SOFA [31]                  |
|------------------------------------------|----------------------------|----------------------------|----------------------------|
| Year of publication                      | 1996                       | 1995                       | 1996                       |
| Selection of variables and their weights | Multiple logistic regression| Literature review and logistic regression | Panel of experts |
| Variables used to assess organ dysfunction|                            |                            |                            |
| Neurologic                               | Glasgow Coma Scale         | Glasgow Coma Scale         | Glasgow Coma Scale         |
| Cardiovascular                           | Heart rate, systolic blood pressure | Pressure-adjusted heart rate | Mean arterial blood pressure, vasopressor use |
| Renal                                    | Serum urea or urea nitrogen, creatinine, urine output | Serum creatinine | Serum creatinine, urine output |
| Respiratory                              | PaO\textsubscript{2}/FiO\textsubscript{2} ratio, mechanical ventilation | PaO\textsubscript{2}/FiO\textsubscript{2} ratio, mechanical ventilation | PaO\textsubscript{2}/FiO\textsubscript{2} ratio, mechanical ventilation |
| Hematologic                              | White blood cell count, platelet count | Platelet count | Platelet count |
| Hepatic                                  | Serum bilirubin, prothrombin time | Serum bilirubin | Serum bilirubin |

LODS, Logistic Organ Dysfunction Score; MODS, Multiple Organ Dysfunction Score; SOFA, Sequential Organ Dysfunction Score.
LODS was accurate in characterizing the progression of organ dysfunction during the first week of ICU stay [33].

**Multiple Organ Dysfunction Score**

The development of the MODS was based on a literature review of 30 publications that had characterized organ dysfunction [30,34]. Seven organ systems were then selected for further consideration (respiratory, cardiovascular, renal, hepatic, hematological, central nervous system, gastrointestinal), and variables for each organ system were chosen according to a set of ‘ideal descriptor’ criteria (Table 3). No accurate descriptor of gastrointestinal function could be identified, so this system was not included in the final model. For the cardiovascular system, Marshall and colleagues [30] created a composite variable, the pressure-adjusted heart rate (heart rate × central venous pressure/mean arterial pressure); in patients without a central line, this variable is assumed to be normal. For each of the six organs, the first parameters of the day are used to calculate the score and a score of 0 (normal) to 4 (most dysfunction) is awarded, giving a total maximum score of 24. The score was developed in 336 patients admitted to one surgical ICU and validated in 356 patients admitted to the same ICU [30]. Although not designed to predict ICU mortality, increasing MODS values do correlate with ICU outcome [30]. ICU mortality also increases with increasing numbers of failing organ systems [30,35]. The delta MODS, defined as the difference between the MODS at admission and the maximum score, may be more predictive of outcome than individual scores [30].

**Sequential Organ Failure Assessment**

The SOFA was developed in 1994 during a consensus conference [31]. Six organ systems (respiratory, cardiovascular, renal, hepatic, central nervous, coagulation) were selected based on a review of the literature, and the function of each is scored from 0 (normal function) to 4 (most abnormal), giving a possible score of 0 to 24. Unlike the MODS score in which the first value of each day is used, for the SOFA score, the worst value on each day is recorded. Another key difference is in the cardiovascular component; instead of the composite variable, the SOFA score uses a treatment-related variable (dose of vasoressor agents). This is not ideal, as treatment protocols vary among institutions, among patients and over time, but it is difficult to avoid, especially for the cardiovascular system.

The SOFA was initially validated in a mixed, medical-surgical ICU population [31,36] and has since been validated and applied in various patient groups [37-39]. In a prospective analysis of 1,449 patients, a maximum total SOFA score greater than 15 correlated with a mortality rate of 90% [40]. Changes in SOFA score over time are also useful in predicting outcome. In a prospective study of 352 ICU patients, an increase in SOFA score during the first 48 hours in the ICU, independent of the initial score, predicted a mortality rate of at least 50%, while a decrease was associated with an ICU mortality rate of just 27% [41]. In a prospective observational study of 1,340 patients with multiple organ dysfunction syndrome, Cabrè and colleagues [42] reported 100% mortality for patients with age over 60 years, a total maximum SOFA greater than 13 on any of the first 5 days of ICU admission, minimum SOFA greater than 10 at all times, and a positive or unchanged SOFA trend over the first 5 days of ICU admission.

**Discussion**

Several studies have directly compared the various organ dysfunction scoring systems. Pettilä and colleagues [43] reported comparable discriminative power of APACHE III, LODS, SOFA, and MODS to predict hospital mortality in a single center study. Peres Bota and colleagues [44] reported no significant differences between MODS and SOFA for mortality prediction in 949 general ICU patients. However, when using the cardiovascular component, outcome prediction was better for the SOFA score at all time intervals compared to the MODS, a finding confirmed by other studies [45]. In a multicenter study, Timsit and colleagues [33] reported good accuracy and internal consistency for both the SOFA and LODS. However, in a Canadian study of 1,436 ICU patients [45], SOFA and MODS had only a modest ability to discriminate between survivors and non-survivors. More recently, SOFA was reported to have superior discriminative ability for

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**Table 3. ‘Ideal’ descriptors of organ dysfunction in ICU patients**

| Descriptor                                                                 | Characteristics                                                                 |
|----------------------------------------------------------------------------|---------------------------------------------------------------------------------|
| Simple and inexpensive                                                    | Routinely available in all ICUs                                                 |
| Reliable (intra and inter-observer)                                        | Objective (that is, observer independent)                                       |
| Reflect acute dysfunction of the organ in question                        | Reproducible in large, heterogeneous groups of ICU patients                   |
| Not affected by transient, reversible abnormalities associated with       | Reproducible in several types of ICUs from different regions of the globe     |
| therapy or practical interventions                                        | Abnormal in one direction only                                                  |
| Reflect acute dysfunction of the organ in question but not chronic         | Using continuous rather than dichotomous variables                             |
| dysfunction                                                                |                                                                                 |

Modified from [34].
hospital mortality and unfavorable neurologic outcome compared to MODS in patients with brain injury [46].

Severity assessment based on nursing workload use
The Therapeutic Intervention Scoring System (TISS)

TISS was originally developed in 1974 to assess severity of illness and compare patient care based on the measurement of nursing workload [47]. The original score included 57 therapeutic activities with points assigned for each activity conducted during a 24-hour period; higher values were given for more specialized or time-consuming activities. In 1983, the score was updated and expanded to include 76 items [48]. However, TISS-76 was criticized for being too time-consuming and cumbersome, and in 1996, a simplified version was devised using advanced statistical analysis [49]. TISS-28 includes just 28 items, divided into 7 groups: basic activities, ventilatory support, cardiovascular support, renal support, neurological support, metabolic support, and specific interventions. The scoring is weighted to give a total score of 78. TISS-28 was validated in 22 Dutch ICUs [49] and in 19 ICUs in Portugal [50]. According to this system, each nurse can provide care for 46.35 TISS-28 points per shift, with each TISS-28 point requiring 10.6 minutes of each nurse’s shift. This information can be useful for planning the allocation of nursing manpower, to evaluate the efficacy in the use of nursing workload use and to objectively classify ICUs based on the amount (and not the complexity) of care provided [51].

Nine Equivalents of Nursing Manpower Use Score

NEMS was derived from the TISS-28 with the aim of creating a simpler system that would be more widely used [52]. Nursing activities are separated into nine categories: basic monitoring, intravenous medication, mechanical ventilatory support, supplementary ventilatory care, single vasoactive medication, multiple vasoactive medication, dialysis techniques, specific interventions in the ICU, specific interventions outside the ICU. Each of these is awarded weighted points, giving a maximum score of 56. NEMS has been validated in large cohorts of ICU patients and is easy to use with almost no interrater variability [53]. Again, this system can be used to evaluate the efficacy of nursing workload use at the ICU level so as to objectively classify ICUs based on the amount (and not only on the complexity) of care provided [51].

Nursing Activities Score

Based on the TISS-28, the Nursing Activities Score (NAS) includes several additional nursing activities not necessarily related to the severity of illness of the patients [54]. The list of items was developed by consensus. The average time consumption of the activities was determined from a 1-week observational cross-sectional study and the results compared with those of the TISS-28 items in a cohort of 99 ICUs in 15 countries. At the end of this process, a total of five new items and 14 sub-items describing nursing activities in the ICU (for example, monitoring, care of relatives, administrative tasks) were added to the TISS-28 list. The new activities accounted for 60% of the average nursing time; and in the development study, NAS activities accounted for 81% of the nursing time (versus 43% in TISS-28) [54].

Discussion

These scores have been used mainly to assess nurse staffing in the ICU, although higher scores are associated with worse outcomes [55,56]. All the scores are limited by the items included, and can be prone to subjective interpretation and influenced by patient case-mix, local admission and discharge policies, and local management protocols. Use of these scores to compare units may, therefore, be difficult; however, within a unit they can provide a valuable indication of changing workload needs. These scores may also be used to estimate overall costs for groups of ICU patients, although they are less reliable on an individual patient basis [57]. Instruments, such as the Work Utilization Ratio, which evaluates the total number of points actually scored divided by the total possible points, have been proposed to evaluate the effectiveness of the use of nursing workload resources [51]. A recent position statement by the European Federation of Critical Care Nursing Associations recommends that all units use such a system on a regular basis to monitor the efficiency of the use of nursing manpower [58].

Other uses of scoring systems

In addition to their use in outcome prediction, organ function assessment, and nursing workload evaluation discussed above, scoring systems have several other potential uses, including use in clinical trials for case-mix comparisons and use in the assessment and comparison of ICU quality and performance.

Clinical trials

Scoring systems are increasingly being incorporated into clinical trial design. Outcome prediction scores, such as APACHE and SAPS, have been used for some time to compare patient populations in clinical trials and even for the identification of eligible patients for inclusion. The analysis of results from one recent randomized controlled study [59], which showed improved outcomes in patients with higher APACHE II scores, led to the drug under investigation, drotrecogin alfa (activated), being licensed in the United States for use only in patients with severe sepsis who are at a high risk of death, that is, those with
an APACHE II score above 25. However, this is a controversial approach and these scores were not designed for this purpose [60].

The realization that mortality alone is inadequate as an outcome measure for interventional studies in ICU patients has led many trials, especially in sepsis, to include an organ dysfunction score as part of ongoing patient assessment so that effects on morbidity can also be evaluated. Increased economic pressure has also led to greater concerns about cost-effectiveness of new and established interventions and nursing workload scores are also being incorporated into clinical trial design, particularly for interventions likely to impact on nursing workload.

Assessment of ICU performance
Costs of care for an ICU patient have been estimated as being three times the costs of care for a general ward patient [61]. Monitoring ICU performance is, therefore, increasingly important in the fight to control hospital expenses. While crude mortality data may offer some global guidance as to ICU performance, adjusting mortality rates according to disease severity, by using outcome prediction models such as the APACHE score to calculate the standardized mortality ratio, can help improve quality assessment. Such severity-adjusted indicators can be used to assess performance of a single ICU over time or to compare several or more units. However, this approach has several limitations, including potential effects of pre-ICU admission factors, implications of different ICU discharge policies [62], and effects of different patient case-mix and hence disease severity between units or in the same unit at different times [63]. Nevertheless, there are large variations in risk-adjusted mortality rates among hospitals [20] and repeated quality assessment may help determine the reasons underlying these differences and enable programs to be developed to improve performance.

Conclusions
General illness severity scores are widely used in the ICU to assess resource use, predict outcome, and characterize disease severity and degree of organ dysfunction. All the scores were developed to be used in mixed groups of ICU patients and their accuracy in subgroups of patients can be questioned; disease-specific scoring systems are increasingly being developed. As ICU populations change and new diagnostic, therapeutic and prognostic techniques become available, all the scoring systems will need to be updated. Importantly, the different scoring systems have different purposes and measure different parameters; we believe they should be seen as complementing each other, rather than competing with one another. For example, outcome prediction models cannot be used to assess the severity of individual organ dysfunctions or to monitor patient progress over time. Although organ dysfunction scores correlate with outcomes, this is not what they were developed for and outcome prediction should be left to scores such as the APACHE and SAPS systems. The workload scores complete the picture by offering information on how the patient’s disease will impact on staffing requirement and resource use. We envisage that, increasingly, all patients will be initially evaluated using a general outcome prediction model computed on admission or within the first 24 hours, then by repeated organ failure (for example, SOFA) and nursing workload (for example, TISS–28) scores during their ICU stay. When used together, these three approaches could provide a more accurate indication of disease severity and prognosis, which could be of help both to the clinician in charge of the patient and to the manager involved in resource allocation and performance assessment.

Abbreviations
APACHE = Acute Physiology and Chronic Health Evaluation; LODS = Logistic Organ Dysfunction Score; MOIDS = Multiple Organ Dysfunction Score; MPM = Mortality Probability Model; NAS = Nursing Activities Score; NEMS = Nine Equivalents of Nursing Manpower Use Score; SAPS = Simplified Acute Physiology Score; SOFA = Sequential Organ Failure Assessment; TISS = Therapeutic Intervention Scoring System; WHD-94 = Weighted Hospital Days scale.

Author details
1 Department of Intensive Care, Erasme University Hospital, Route de Lennik 808, 1070 Brussels, Belgium. 2 Department of Intensive Care, Hospital de St Antonio dos Capuchos, Centro Hospitalar de Lisboa Central, EPE, 1169-050 Lisbon, Portugal.

Competing interests
The authors declare that they have no competing interests.

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