Association between the modified Nutrition Risk in Critically Ill (mNUTRIC) score and clinical outcomes in the intensive care unit: A secondary analysis of a large prospective observational study

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Abstract

Background

Malnutrition in intensive care unit (ICU) patients is associated with adverse clinical outcomes.

The nutrition risk in the critically ill score (NUTRIC) was proposed as an appropriate nutritional assessment tool in critically ill patients. This score uses interleukin-6 (IL-6), a biomarker that
is not always available. This prospective observational study was conducted to identify the nutritional risk in ICU patients using the modified NUTRIC (mNUTRIC) score (which does not include IL-6) and to explore the relationship between 28-day mortality and high mNUTRIC scores.

**Methods**

The data were extracted from The Beijing Acute Kidney Injury Trial (BAKIT). This trial was a prospective, observational, multi-centre study conducted in 30 ICUs at 28 tertiary hospitals in Beijing, China, from March 1 to August 31, 2012. In total, 9049 patients were admitted consecutively, and 3107 patients with complete clinical data were included in this study. The predictive capacity of the mNUTRIC score was studied by receiver operating characteristic (ROC) curve analysis. The significance level was set at 5%.

**Results**

Among the 3107 patients, the 28-day mortality rate was 17.4% (540 patients died). High nutritional risk patients were older (P<0.001), with higher illness severity scores than low nutritional risk patients. Multivariate analysis revealed that the mNUTRIC score was an independent risk factor for 28-day mortality and mortality increased with increasing scores (p = 0.000). The calculated area under curve (AUC) for the mNUTRIC score was 0.763 (CI 0.740 - 0.786).

**Conclusions**

Nearly 28.2% of patients admitted to the ICU were at risk of malnutrition, and a high mNUTRIC score was associated with increased ICU length of stay and higher mortality.
**Trial Registration**

This study was registered at www.chictr.org.cn (registration number ChiCTR-ONC-11001875).

Registered on 14 December 2011.

**Key words**

The modified nutrition risk in critically ill score, Intensive care unit, Mortality

**Background**

Malnutrition is common in intensive care unit (ICU) patients, it is associated with a variety of adverse outcomes, including higher complication rates, prolonged mechanical ventilation, prolonged hospitalization, and higher mortality[1, 2]. For critically ill patients, we need to assess their nutritional status and provide adequate nutritional support[3], so effective tools are needed to determine which ICU patients need nutritional support and the benefits of nutritional support. However, traditional methods of nutrition assessment are limited in the hospital setting.

Recently, Heyland et al[4] published the first nutritional risk assessment tool specifically designed for critically ill patients: the NUTRIC score.

The NUTRIC score includes age, the Acute Physiology and Chronic Health Evaluation II (APACHE II) score[5], the Sequential Organ Failure Assessment (SOFA) score[6], comorbidities, days from hospitalization to ICU admission, and the interleukin-6 (IL-6) level, which was developed to link starvation, inflammation, and clinical outcomes[4]. Patients are scored from 0 to 10, a score of 6 or greater indicates a high nutritional risk.

The NUTRIC score can predict 28-day mortality in a medical-surgical ICU population, high-risk patients who stayed in the ICU for more than 3 days benefited more from nutritional support.
than low-risk patients[4]. But the use of original NUTRIC score is limited by the availability of IL-6, which is not readily available in many institutions. Another study evaluated a modified NUTRIC score excluding IL-6 and found that a score of 5 or higher still indicated a high risk of malnutrition[7]. Moreover, Heyland et al.[4] stated that IL-6 only increased the C-index by 0.007 (from 0.776 to 0.783), with no statistical difference. Therefore, they suggested that in settings in which IL-6 is not available, it could be omitted from the NUTRIC score. This adjusted score is called the modified NUTRIC score (mNUTRIC). Rahman et al[8] evaluated this modified NUTRIC score and found that mortality increased by 1.4 % (95% CI, 1.3-1.5) for every point increase in the mNUTRIC score.

Using appropriate nutrition screening and assessment tools will help identify effective strategies that reduce the negative impact of malnutrition. Our study was conducted to identify the prevalence of nutritional risk in general ICU patients based on mNUTRIC scores.

**Methods**

**Study design and data collection**

This study used a database from a prospective, multi-centre, observational study that investigated the epidemiology of acute kidney injury (AKI) in critically ill patients in 30 ICUs at 28 tertiary hospitals in Beijing, China, from March 1 to August 31, 2012 (the Beijing Acute Kidney Injury Trial (BAKIT))[9]. (for a complete list of these hospitals and the persons responsible for the data acquisition, see Additional file 1). Study subjects included all adult patients (age≥18 years) admitted consecutively to the ICU. Only the initial ICU admission was considered in this study. The following patients were excluded: patients with preexisting end-stage chronic kidney disease, patients already receiving renal replacement therapy (RRT)
before admission to the ICU, and patients who had received kidney transplantation in the
previous 3 months[10]. Pre-existing comorbidities were diagnosed based on the International
Classification of Diseases (ICD-10) codes. Patients were followed up until death, until hospital
discharge, or for 28 days. Among the 9079 patients who were admitted consecutively, 3107
patients were included in our study (Figure 1).

Thorough follow-up of all patients included in the study was conducted in the first 10 days after
ICU admission. The collected data included demographics, anthropometrics, admission
diagnosis, comorbidities, daily vital signs and laboratory data, which were used to automatically
calculate the APACHE II score, the Simplified Acute Physiology Score II (SAPS II) score[11]
and the SOFA score, days from hospital to ICU admission, ICU length of stay (LOS), hospital
LOS, use of vasoactive drugs, and length of mechanical ventilation. RRT data were also
reported.

The patients were followed up until death, hospital discharge, or for 28 days.

**Nutritional support**

Nutritional support methods were based on the guidelines for enteral and parenteral nutrition
issued by the European and American Society of Enteroprotective Nutrition[12], combined
with our accumulated clinical experience, individualized nutritional support was given to all
patients. The patients began enteral nutrition (EN) 20-25 kcal/(kg.d) within 24-48 hours of
admission to the ICU (on average). If the patient was intolerant of EN or had contraindications
to EN, parenteral nutrition (PN) support was given within 24 - 48 hours. If EN could not fully
meet the nutritional needs of patients, appropriate intravenous supplementation with glucose,
amino acids, or fat emulsion was given, that is, the combination of EN and PN.
Definitions

We used the modified 9-point scale of the NUTRIC score, the mNUTRIC score. We defined the scores from 0 to 4 as “low scores”, which indicated a low level of risk of malnutrition, and the scores from 5 to 9 as “high scores”, which were associated with worse clinical outcomes.

Statistical analysis

Non-normally distributed continuous variables were expressed as the medians with interquartile ranges (IQRs) and were compared using the Mann-Whitney U test or Kruskal-Wallis analysis of variance with Bonferroni correction. Categorical variables were expressed as the number of cases and proportions and were compared using the Mantel-Haenszel Chi-square test.

A multivariate Cox regression analysis was performed using a backward stepwise selection method, with P value < 0.05 as the entry criterion, and P value ≥ 0.10 as the removal criterion. The assumption of proportional hazards was checked graphically using log (-log (survival probability)) plots and was found to be appropriate. Variables considered for multivariable analysis included age, sex, body mass index (BMI), illness severity scores, use of vasoactive drugs, mechanical ventilation and underlying diseases. We tested for collinearity among all variables using a Cox regression analysis to generate hazard ratios (HR) and 95% confidence intervals (CIs).

Receiver operator characteristic curve analysis was used to calculate the sensitivity and specificity for comparisons of outcomes and mNUTRIC scores. The 28-day survival stratified by low and high mNUTRIC scores was additionally evaluated graphically using the Kaplan-Meier product limit survival plot.
All statistical analyses were performed using SPSS software (IBM Corp., Statistics for
Windows, version 22.0, Armonk, NY, USA), with a two-sided P value < 0.05 considered
statistically significant.

Results

Study population

Among the 9049 patients enrolled in the BAKIT study, 5942 were excluded for the reasons
shown in Figure 1, leaving 3107 patients for analysis. The characteristics of the entire cohort
are shown in Table 1. The median age was 64.0 (IQR: 51.0 - 77.0) years, and 61.5% were men.
The all-cause 28-day mortality rate was 17.4% and the median ICU LOS was 4.0 (IQR: 2.0 -
9.0) days. Among the included patients, the median BMI was 24.0 (IQR: 21.0 - 26.0) kg/m²,
the median APACHE II score was 14.0 (IQR: 10.0 - 20.0), the median SOFA score was 6.0
(IQR: 3.0 - 8.0), and the median number of comorbidities was 1 (IQR: 0 - 2). Mechanical
ventilation was used in 2021 (65.0%) patients, 1307 patients (42.1%) received vasopressors,
and 281 patients (9.0%) underwent RRT. A total of 876 patients (28.2%) had high mNUTRIC
scores.

Figure.1 Flowchart of validation cohort
Table 1 Patient characteristics by mNUTRIC score

| Characteristic               | All patients (n=3107) Median(IQR) Number (%) | Low nutrition risk(n=2231) Median(IQR) Number (%) | High nutrition risk(n=876) Median(IQR) Number (%) | P value |
|-----------------------------|---------------------------------------------|-------------------------------------------------|---------------------------------------------------|---------|
| Age(years)                  | 64(51-77)                                  | 60(47-72)                                      | 76(66-82)                                        | <0.001  |
| Male sex                    | 1912(61.5)                                 | 1378(61.8)                                    | 534(61.0)                                       | 0.919   |
| BMI                         | 24(21-26)                                  | 24(22-26)                                     | 23(21-26)                                       | 0.003   |
| Vasoactive therapy          | 1307(42.1)                                 | 954(42.8)                                     | 353(40.3)                                       | 0.457   |
| Mechanical ventilation      | 2021(65.0)                                 | 1354(60.7)                                   | 667(76.1)                                       | <0.001  |
| Sepsis                      | 896(28.8)                                  | 419(18.8)                                     | 477(54.5)                                       | <0.001  |
| **Severity of illness**     |                                             |                                                |                                                  |         |
| APACHEII                    | 14(10-20)                                  | 12(8-15)                                      | 23(19-28)                                       | <0.001  |
| SAPSII                      | 34(26-45)                                  | 30(23-38)                                     | 50(39-64)                                       | <0.001  |
| SOFA                        | 6(3-8)                                     | 4(3-7)                                        | 9(6-11)                                         | <0.001  |
| NUTRIC score                | 3(2-5)                                     | 3(2-3)                                        | 6(5-7)                                          | <0.001  |
| **Admission category**      |                                             |                                                |                                                  |         |
| medical                     | 1480(47.6)                                 | 878(39.4)                                     | 602(68.7)                                       | <0.001  |
| surgical                    | 1627(52.4)                                 | 1353(60.6)                                   | 274(31.3)                                       |         |
| **Comorbid diseases**       |                                             |                                                |                                                  |         |
| Cancer                      | 486(15.6)                                  | 297(13.3)                                     | 189(21.6)                                       |         |
| Hypertension                | 1222(39.3)                                 | 739(33.1)                                     | 483(55.1)                                       |         |
| Coronary disease            | 615(19.8)                                  | 293(13.1)                                     | 322(36.8)                                       |         |
| Chronic kidney disease      | 170(5.5)                                   | 63(2.8)                                       | 107(12.2)                                       |         |
| Diabetes                    | 532(17.1)                                  | 277(12.4)                                     | 255(29.1)                                       |         |
| COPD                        | 166(5.3)                                   | 89(4.0)                                       | 77(8.8)                                         |         |
| **Category of ICU admission** | **diagnosis**                             |                                                |                                                  |         |
| Cardiovascular              | 848(27.3)                                  | 681(30.5)                                     | 167(19.1)                                       |         |
| Respiratory                 | 548(17.6)                                  | 316(14.2)                                     | 232(26.5)                                       |         |
Neurologic  462(14.9)  321(14.4)  141(16.1)
Trauma  238(7.7)  191(8.6)  47(5.4)
Gastrointestinal  607(19.4)  413(18.5)  194(22.1)
Metabolic  77(2.5)  43(1.9)  34(3.9)

Outcome data

|                      | ICU LOS(days) | Hospital LOS(days) | 28-day mortality | In-hospital mortality | AKI | RRT |
|----------------------|---------------|--------------------|------------------|----------------------|-----|-----|
| 4(2-9)               | 19(12-29)     | 540(17.4)          | 521(16.8)        | 1334(42.9)           | 281(9.0) | 108(4.8) |
| 4(2-7)               | 19(12-28)     | 208(9.3)           | 173(7.8)         | 752(33.7)            | 173(19.7) |
| 6(3-13)              | 21(11-34)     | 332(37.9)          | 348(39.7)        | 582(66.4)            |      |

Hospitalization expense ( thousand yuan )

|                      |               |                   |                   |                     |
|----------------------|---------------|------------------|-------------------|---------------------|
| 40(19-96)            | 34(17-87)     | 55(27-113)       | <0.001            |

Data are expressed as the median (interquartile range), and number (percentage). BMI, body mass index; SAPS II, Simplified Acute Physiology Score II; SOFA, Sequential Organ Failure Assessment; APACHE II, Acute Physiology and Chronic Health Evaluation II; NUTRIC score, the nutrition risk in the critically ill score; COPD, chronic obstructive pulmonary disease; LOS, length of stay; AKI, acute kidney injury; RRT, renal replacement therapy.

Characteristics of high nutritional risk patients

From Table 1, we can see high nutritional risk patients were older (P < 0.001), with higher illness severity scores than low nutritional risk patients. High nutritional risk patients were more likely to present with sepsis on ICU admission and had longer durations of ICU and hospital stays when compared to the low nutritional risk group. Furthermore, mechanical ventilation was more commonly used in high nutritional risk patients (76.1% vs 60.7%; P < 0.001). The 28-day mortality and in-hospital mortality rates were higher among high nutritional risk patients than low nutritional risk patients (P < 0.001).

28-Day mortality according to score
Our analysis showed that the 28-day mortality increased with higher mNUTRIC scores (Figure 2), and the 28-day mortality for the maximum mNUTRIC score was 67.4%.

Figure 2. The 28-day mortality according to modified NUTRIC score.

High mNUTRIC score and the 28-day mortality

In multivariate Cox regression analysis (Table 2), after adjusting for age, sex, BMI, illness severity scores, use of vasoactive drugs, mechanical ventilation and underlying diseases, the mNUTRIC score, sepsis and AKI were independent predictors of 28-day mortality. The presence of high mNUTRIC scores was associated with a higher risk of mortality (Figure 3).

Table 2  Multivariate Cox regression analysis of 28-day mortality in all patients

| Characteristic | Hazard ratio | 95%CI        | P    |
|----------------|--------------|--------------|------|
| mNUTRIC score  | 1.430        | 1.351–1.514  | 0.000|
| Sepsis         | 2.832        | 2.272–3.529  | 0.000|
| AKI            | 2.171        | 1.732–2.720  | <0.001|
mNUTRIC score, the modified nutrition risk in the critically ill score; AKI, acute kidney injury; CI, confidence interval

Figure 3  Survival curve of 28-day mortality stratified by mNUTRIC scores.

**Area under the curve of scores for predicting 28-day mortality**

We can see that in this cohort and each subgroup, the areas under the curve (AUCs) of the mNUTRIC score for predicting 28-day mortality indicated good predictive performance of the score (Figure 4). In the ROC curve for the mNUTRIC score, the best cut-off value was at 4 (sensitivity 61.48% and specificity 78.81%) in this cohort, and the Youden index was 0.4029.
Figure 4 Performance of mNUTRIC scores in predicting 28-day mortality. a. All patients (n=3107); b. All mechanical ventilation patients (n=2021); c. Medical mechanical ventilation patients (n=751); d. Sepsis patients (n=896); e. AKI patients (n=1334); f. CRRT patients (n=281)

Discussion

This study was a secondary analysis of a prospective observational study in surgical-medical ICUs. We used a validated nutrition assessment tool in an attempt to demonstrate an association between malnutrition and 28-day mortality. We found a high incidence of malnutrition in ICU patients, and malnutrition was associated with a poor prognosis.
In the present study, 28.2% of the critically ill patients admitted to the ICU were at high nutritional risk and had mNUTRIC scores $\geq 5$. These findings were similar to the results of a study conducted in Turkey[13], in which 22.4% patients were evaluated as having high scores (between 5 and 9). Lew et al[14] also demonstrated that the prevalence of malnutrition in the ICU was 28% using the 7-point Subjective Global Assessment (7-point SGA) to determine patients’ nutritional status. Recently, a study[15] reported that 45% of mechanically ventilated patients admitted to the ICU were at high nutritional risk. Similarly, Kalaiselvan et al.[16] reported that 42.5% of mechanically ventilated patients had NUTRIC scores $\geq 5$. Our study is more generalizable because of the inclusion of both medical and surgical patients. The aforementioned studies included only patients on mechanical ventilation, and patients on mechanical ventilation were more seriously ill than those not on mechanical ventilation. The differences among studies are mainly the result of different populations and nutrition screening tools.

In our study, the 28-day mortality associated with the maximum mNUTRIC score was 67.7%, which is similar to the finding in the study by Jeong[17], in which this rate was 62.5%. Compared with patients with a low NUTRIC score, patients with high NUTRIC score had a higher mortality rate and longer ICU LOS. Similar results were reported by Mendes et al[1], in which the NUTRIC score was used in an ICU population, and the findings are consistent with those in the study by Heyland[4].

The mortality rate in our study was 17.4%, which was lower than the rate reported in the second validation study of the NUTRIC score (29%) by Rahman et al[8]. This difference may be
because our study included many postoperative care patients. In this study, we found that the mNUTRIC score was a good prognostic predictor in critically ill patients and that high mNUTRIC scores were associated with an elevated risk of death at 28 days (HR=1.430, 95% CI=1.351 to 1.514, P=0.000). This finding is consistent with those of prior studies [1, 18, 19]. Several studies have shown that the beneficial effects of adequate nutritional support are more evident in high-risk patients than in low-risk patients [20, 21]. The mNUTRIC score may be helpful in guiding clinicians in providing adequate nutritional support to ICU patients.

The mNUTRIC score was found to have a fair predictive performance for 28-day mortality in this cohort (AUC 0.763; 95% CI 0.740 - 0.786) and each subgroup. These results are in line with those of the initial validation study by Heyland et al. (AUC: 0.783)[4] and a recently published validation study of the mNUTRIC score by Mukhopadhyay et al. (AUC 0.71)[22]. Recently, a study[17] showed that the AUC of the NUTRIC score for the prediction of 28-day mortality was 0.762 (95% CI: 0.718-0.806), while that of the mNUTRIC score was 0.757 (95% CI: 0.713-0.801). There was no significant difference between the two scores (p = 0.45). The mNUTRIC score is a good nutritional risk assessment tool for critically ill patients.

We found that the best cut-off value for the mNUTRIC score was $> 4$ (sensitivity 61.48% and specificity 78.81%) in this cohort, and the Youden index was 0.4029, which is consistent with previous work by de Vries et al[23]. However, in another study, the best cut-off value was at 6 (sensitivity 75% and specificity 65%), and the Youden index was 0.401[17]. Jung et al reported that patients were considered to be at high risk of malnutrition and to benefit from aggressive nutritional support when their mNUTRIC score was $\geq 5[24]$. Our study included patients with...
various diseases, while Jung’s study population was limited to patients with sepsis. Further investigation is needed to find the best cut-off value of the mNUTRIC score to define the high-risk group.

The limitations of our study stem mainly from the fact that it is a secondary analysis of an original database that lacked data on inflammation indicators such as IL-6. Therefore, we could not calculate the NUTRIC score to verify the differences between the two scores. Second, nutrition history and feeding parameters were not available in our cohort, so the associations among nutritional adequacy, mNUTRIC score and mortality could not be confirmed by our results.

Conclusion

Patients were considered to be at high risk of malnutrition when their mNUTRIC score was > 4. The mNUTRIC score is a practical, easy-to-use tool based on variables that are easy to obtain in the critical care setting.

Abbreviations

ICU: intensive care unit; NUTRIC: the nutrition risk in the critically ill score; IL-6: interleukin-6; mNUTRIC: the modified nutrition risk in the critically ill score; ROC: receiver operating characteristic; AUC: area under curve; AKI: acute kidney injury; BAKIT: the Beijing Acute Kidney Injury Trial; BMI: body mass index; RRT: renal replacement therapy; APACHE II: acute physiology and chronic health evaluation II; SAPS II: the simplified acute physiology score II; SOFA: sequential organ failure assessment; LOS: length of stay; EN: enteral nutrition; PN:
parenteral nutrition; IQR: interquartile range; HR: hazard ratio; CI: confidence interval; COPD: chronic obstructive pulmonary disease

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Authors’ contributions

NW and MPW designed and carried out the study, NW performed the statistical analysis, and drafted the manuscript. LJ and BD were involved in design and in acquisition of data and helped to revise the manuscript critically for important content. BZ was involved in the design and the statistical analysis. The Beijing Acute Kidney Injury Trial (BAKIT) Workgroup participated in acquisition and interpretation of data. XX conceived of the study, participated in its design, and helped to revise manuscript. All authors read and approved the final manuscript.

Conflict of interest

The authors declare that they have no conflict of interest.

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Availability of data and materials

The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.
Ethics approval and consent to participate

This study was approved by the Institutional Review Boards of the Ethics Committees of the lead study centre (Fu Xing Hospital, Capital Medical University, China) and all other participating hospitals (Additional file 2). We confirm that all methods were carried out in accordance with relevant guidelines and regulations. Being an observational study, written informed consent from participants to partake into the study was not necessary. Hence, we obtained an informed consent waiver from the above ethical committees.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

Reference

1. Correia MI, Waitzberg DL. The impact of malnutrition on morbidity, mortality, length of hospital stay and costs evaluated through a multivariate model analysis. Clin Nutr 2003;22(3):235–239.

2. Heyland DK, Cahill N, Day AG. Optimal amount of calories for critically ill patients: depends on how you slice the cake! Crit Care Med 2011;39(12):2619-26.

3. Al-Dorzi HM, Al barrak A, F erwana M, et al. Lower versus higher dose of enteral caloric intake in adult critically ill patients: a systematic review and meta-analysis. Crit Care 2016;20(1):358.
4. Heyland DK, Dhaliwal R, Jiang X, et al. Identifying critically ill patients who benefit the most from nutrition therapy: the development and initial validation of a novel risk assessment tool. Crit Care 2011;15(6):R268.

5. Knaus WA, Draper EA, Wagner DP, et al. APACHE II: a severity of disease classification system. Crit Care Med 1985;13(10):818-29.

6. Vincent JL, de Mendonca A, Cantraine F, et al. Use of the SOFA score to assess the incidence of organ dysfunction/failure in intensive care units: Results of a multicenter, prospective study. Critical Care Medicine 1998;26(11):1793-1800.

7. Heyland DK, Dhaliwal R, Wang M, et al. The prevalence of iatrogenic underfeeding in the nutritionally 'at-risk' critically ill patient: Results of an international, multicenter, prospective study. Clin Nutr 2015;34(4):659-66.

8. Rahman A, Hasan RM, Agarwala R, et al. Identifying critically ill patients who will benefit most from nutritional therapy: Further validation of the "modified NUTRIC" nutritional risk assessment tool. Clin Nutr 2016;35(1):158-62.

9. Luo X, Jiang L, Du B, Wen Y, et al. A comparison of different diagnostic criteria of acute kidney injury in critically ill patients. Crit Care 2014; 18(4):R144.

10. Piccinni P, Cruz DN, Gramaticopol S, et al. Prospective multicenter study on epidemiology of acute kidney injury in the ICU: a critical care nephrology Italian collaborative effort (NEFROINT). Minerva Anestesiol 2011;77(11):1072-83.
11. Le Gall JR, Lemeshow S, Saulnier F. A new Simplified Acute Physiology Score (SAPS II) based on a European/North American multicenter study. JAMA 1993;270(24):2957-63.

12. McClave SA, Taylor BE, Martindale RG, et al. Guidelines for the Provision and Assessment of Nutrition Support Therapy in the Adult Critically Ill Patient: Society of Critical Care Medicine (SCCM) and American Society for Parenteral and Enteral Nutrition (A.S.P.E.N.). JPEN J Parenter Enteral Nutr 2016; 40(2):159-211.

13. Ozbilgin S, Hanci V, Omur D, et al. Morbidity and mortality predictivity of nutritional assessment tools in the postoperative care unit. Medicine (Baltimore) 2016;95(40):e5038.

14. Lew CCH, Wong GJY, Cheung KP, et al. Association between Malnutrition and 28-Day Mortality and Intensive Care Length-of-Stay in the Critically ill: A Prospective Cohort Study. Nutrients 2017;10(1).

15. Ata Ur-Rehman HM, Ishtiaq W, Yousaf M, et al. Modified Nutrition Risk in Critically Ill (mNUTRIC) Score to Assess Nutritional Risk in Mechanically Ventilated Patients: A Prospective Observational Study from the Pakistani Population. Cureus 2018;10(12):e3786.

16. Kalaiselvan M, Renuka M, Arunkumar A. Use of nutrition risk in critically ill (nutric) score to assess nutritional risk in mechanically ventilated patients: A prospective observational study. Indian J Crit Care Med 2017;21(5):253-56.
17. Jeong DH, Hong SB, Lim CM, et al. Comparison of Accuracy of NUTRIC and Modified NUTRIC Scores in Predicting 28-Day Mortality in Patients with Sepsis: A Single Center Retrospective Study. Nutrients 2018;10(7).

18. Mendes R, Policarpo S, Fortuna P, et al. Portuguese NUTRIC Study Group. Nutritional risk assessment and cultural validation of the modified NUTRIC score in critically ill patients – A multicenter prospective cohort study. J Crit Care 2017;37:249.

19. Oliveira ML, Heyland DK, Silva FM, et al. Complementarity of modified NUTRIC score with or without C-reactive protein and subjective global assessment in predicting mortality in critically ill patients. Rev Bras Ter Intensiva 2019;31(4):490-496.

20. Alberda C, Gramlich L, Jones N et al. The relationship between nutritional intake and clinical outcomes in critically ill patients: results of an international multicenter observational study. Intensive Care Med 2009;35:1728-1737.

21. Artinian V, Krayem H, DiGiovine B. Effects of early enteral feeding on the outcome of critically ill mechanically ventilated medical patients. Chest 2006;129:960-967.

22. Mukhopadhyay A, Henry J, Ong V et al. Association of modified NUTRIC score with 28-day mortality in critically ill patients. Clin Nutr 2017;36(4):1143-1148.

23. de Vries MC, Koekkoek WK, Opdam MH et al. Nutritional assessment of critically ill patients: validation of the modified NUTRIC score. Eur J Clin Nutr 2018;72:428-435.
24. Jung YT, Park JY, Jeon J et al. Association of Inadequate Caloric Supplementation with 30-Day Mortality in Critically Ill Postoperative Patients with High Modified NUTRIC Score. Nutrients 2018;10:1589.

Table 1 Patient characteristics by mNUTRIC score

Table 2 Multivariate Cox regression analysis of 28-day mortality in all patients

Figure 1 Flowchart of validation cohort

Figure 2. The 28-day mortality according to modified NUTRIC score.

Figure 3 Performance of mNUTRIC scores in predicting 28-day mortality. a. All patients (n=3107); b. All mechanical ventilation patients (n=2021); c. Medical mechanical ventilation patients (n=751); d. Sepsis patients (n=896); e. AKI patients (n=1334); f. CRRT patients (n=281)

Figure 4 Survival curve of 28-day mortality stratified by mNUTRIC scores.