Malnutrition as assessed by nutritional risk index is associated with worse outcome in patients admitted with acute decompensated heart failure: an ACAP-HF data analysis

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Abstract

Malnutrition is common at hospital admission and tends to worsen during hospitalization. This controlled population study aimed to determine if serum albumin or moderate and severe nutritional depletion by Nutritional Risk Index (NRI) at hospital admission are associated with increased length of hospital stay (LOS). Serum albumin levels and lymphocyte counts were retrospectively determined at hospital admission in 1740 consecutive patients admitted with primary and secondary diagnosis of ADHF. The Nutrition Risk Score (NRI) developed originally in AIDS and cancer populations was derived from the serum albumin concentration and the ratio of actual to usual weight, as follows: NRI = (1.519 × serum albumin, g/dL) + (41.7 × present weight (kg)/ideal body weight(kg)). Patients were classified into four groups as no, mild, moderate or severe risk by NRI. Multiple logistic regressions were used to determine the association between nutritional risk category and LOS.

Three hundred and eighty-one patients (34%) were at moderate or severe nutritional risk by NRI score. This cohort had lower BMI (24 ± 5.6 kg/m²), albumin (2.8±0.5 g/dL), mean NRI (73.5±9) and lower eGFR (50±33 mL/min per 1.73 m²). NRI for this cohort, adjusted for age, was associated with LOS of 10.1 days. Using the Multiple Logistic regression module, NRI was the strongest predictor for LOS (OR 1.7, 95% CI: 1.58-1.9; P=0.005), followed by TIMI Risk Score [TRS] (OR 1.33, 95% CI: 1.03-1.71; P=0.02) and the presence of coronary artery disease (OR 2.29, 95%CI: 1.03-5.1; P=0.04). Moderate and severe NRI score was associated with higher readmission and death rates as compared to the other two groups.

Nutritional depletion as assessed by Nutritional Risk Index is associated with worse outcome in patients admitted with ADHF. Therefore; we recommend adding NRI to further risk stratify these patients.

Introduction

Malnutrition is reported to be the leading cause of disease burden in developing countries with high morbidity and mortality rates. In 1974, Butterworth et al. documented malnutrition and its implications in hospitalized patients. Despite great advances in medical science and diagnostic techniques, little has changed during the last three decades. Even in North America, 30-50% of hospitalized patients are described as malnourished or at a risk of malnutrition, with higher rates of malnutrition reported specifically in elderly subjects. Associations have been reported between poor nutritional status and impaired wound healing, increased post-operative complications and mortality. Several studies have shown a relationship between the length of hospital stay and nutritional status, with malnutrition becoming more prevalent as the stay in hospital is prolonged. This led to further consideration of malnutrition given its contribution to the rising costs of our National Health Services and health care systems worldwide. Thus, there is a rationale to routinely screen patients for nutritional risk and risk for complications at hospital admission.

Numerous measures have been developed to assess the nutritional status of the hospitalized patients, among them, the Nutritional Risk Index (NRI) which was developed by the Veterans Affairs Total Parenteral Nutrition Cooperative Study Group. The NRI gained in popularity as it uses an objective (serum albumin and percent usual body weight) rather than subjective measurements to determine nutritional risk in hospitalized patient populations. It has been successfully modified from its original form to be used in various patient groups.

Among chronic diseases, chronic heart failure has already been shown to be associated with a poor prognosis with shortened survival, repeated hospitalizations and sizeable costs to society, estimated to be about $18.8 billion...
per year in the United States. Chronic heart failure (CHF) is increasingly recognized as a chronic multi-system disease with multiple comorbidities such as anemia, insulin resistance, autonomic dysfunction, or cardiac cachexia. Despite the fact that heart failure patients are more prone to experience malnutrition either due to symptomatic anorexia during an episode of ADHF, early satiety and ascites, pharmacological agents, psychological factors or catabolic/anabolic imbalance, the data regarding clinical significance of malnutrition in heart failure patients remains limited. Chronic heart failure patients routinely suffer from hypermetabolic states and pharmacologically induced diuretics can further compound the poor nutritional status of these patients.

Our study aimed to evaluate the role of nutritional evaluation of heart failure patients by using a modified NRI tool, as well as to study the impact of malnutrition severity on hospital length of stay (LOS), readmission rates and to assess existence between malnutrition and mortality in these patients.

Materials and Methods

Subjects

We identified 1,740 consecutive patients from the Advanced Cardiac Admission Program data (ACAP-HF) who were admitted with acute decompensated heart failure (ADHF). Patients with acute myocardial infarction (<3 days duration), hemodynamically significant valvular abnormalities, hemodynamic instability, patients taking any laxatives or weight loss medication or pregnant, were excluded from the study. Serum albumin levels and lymphocyte counts were determined at hospital admission. Questionnaires providing details of the past medical history and demographics were completed by the physicians involved in management during hospitalization of these patients on admission who were kept blinded about the study. Complete records were available for 1110 patients and/or death certificate. The Internal Review Board (IRB) of our Institute approved this study.

Measurements

Anthropometric measures

All measurements were taken at the hospital admission. Body weight was measured to the nearest 0.1 kg (Weight Tronix, New York, NY, USA) and height to the nearest 0.5 cm using a stadiometer (Holtain; Crosswell, Wales). Body Weight (BW) change was calculated as: (current BW in kilograms-ideal BW in kilograms)/ideal BW x 100. Ideal BW was calculated according to the Lorentz formula that takes into account patient’s height and sex as follows: IBW (kg) = height (cm) – 100 – (height (cm)-150)/4. The Body Mass Index (BMI: kilograms/meters²) was calculated as weight in kilograms divided by height (meters²).

Laboratory measures

Blood samples for serum albumin, serum brain natriuretic peptide (BNP) and serum creatinine levels were drawn on admission and before initiation of intravenous diuretics or intravenous fluids if needed. Albumin was measured by immunonephelometry (normal range 3.5-5.5 g/dL). An albumin level of less than 3.5 g/dL was set as the lower limit of normal in our study. Blood was collected in tubes containing potassium EDTA (1 mg/ml blood), for the Triage BNP test. (Biosite Diagnostics Inc., San Diego, California, USA).

Nutritional risk index

The NRI²⁸,²⁹ was originally derived from the serum albumin concentration and the ratio of present to usual weight. Faced with the difficulty in identifying the usual body weight of heart failure patients, we alternatively used ideal body weight instead of usual body weight in the NRI formula³⁰ as follows: NRI = (1.519 × serum albumin, g/dL) + [(41.7 × present weight (kg)/ideal body weight(kg)]. From these NRI values, we defined four grades of nutrition-related risk: i) major risk (NRI<83.5); ii) moderate risk (NRI 83.5-97.5); iii) mild risk (NRI 97.5-100); iv) No risk (NRI >100). The NRI cut-off values were determined according to weight losses of 5%, 10% or 20%. The weight loss norms of 5% and 10% have already been validated by the European Society of Parenteral and Enteral Nutrition (ESPEN) Guidelines for Nutritional Screening.³²

Length of stay

Hospital LOS was the actual number of days the patients remained in the hospital, obtained from the hospital chart after patients were discharged.

Follow up

Serial prospective follow up (mean 2.7±1.0 years) was obtained in all patients by means of a physician-directed telephone interview using a standardized questionnaire. Length of stay was calculated for the index admission, and all subsequent ADHF related readmissions were also documented. If the patient died during follow up, the closest surviving relative or the patient’s physician was interviewed to determine the cause of death. Cardiac death was confirmed by review of the hospital medical records and/or death certificate.

Results

The basic characteristics of our study participants (n=1110) are shown in Table 1. The patients were divided into four groups according to NRI scores identified as No Risk group (n=666) with mean age 68±14 years; 51% males, Mild Risk group (n=63) with mean age 72±14 years; 51% males, Moderate Risk group (n=213) with mean age 72±14 years; 59% males and Severe Risk group (n=168) with mean age 68±15 years; 56% males. As expected, there was a high prevalence of hypertension in the entire cohort (84%) followed by coronary heart disease (69%), diabetes mellitus (45%), hyperlipidemia (45%) and smoking (16%). In addition the BMI values were higher in the No Risk group (31±7 kg/m²) as compared to Mild Risk (25±4 kg/m²), Moderate Risk (23±5 kg/m²) or Severe Risk groups (24±7 kg/m²) with P=0.001.

NUTRITIONAL RISK INDICES

The mean NRI score of the No Risk group was 119±29, Mild Risk group was 92±6.7, Moderate Risk group was 91±4 and Severe risk group was 61±16 (Table 1). Out of 1,110 total patients, 381 patients (34%) were at moderate or severe nutritional risk according to the NRI score. This cohort had a mean NRI score of 73±±9 (P<0.001), BMI value (24 ± 5.6 kg/m²; P<0.001), albumin level (2.8±0.5 g/dL; P<0.001, Lower eGFR (50 ± 33 mL/min per 1.73
m²; P<0.001) and median BNP level (1090 pg/mL; P<0.001) on admission.

Effects of NRI scores on length of hospital stay

Lower NRI scores were found to be strong predictors of extended LOS with HR 1.7 (95%CI: 1.58-1.9); P=0.005 (Table 2 and Figure 1). Patients with lower NRI scores allocating them in either the Moderate or the Severe Risk group were more likely to stay hospitalized for longer (Figure 2). This inverse correlation was highly significant (χ² = 36; P <0.001). The median LOS was found to be 5.8 days in the No Risk group, 7.5 days in the Mild Risk group, 10.1 days in the Moderate Risk group and 10.9 days in the Severe Risk group.

Effect of NRI scores on heart failure readmission rate and all cause mortality

Our study showed that ADHF readmission rates were significantly higher in the patients with lower NRI scores, as seen in Figure 3. The readmission rates were 35% in the No Risk group and 37% in the Mild Risk group as compared to 52% and 68% in the Moderate Risk and the Severe Risk groups, respectively (P<0.001). Similarly, all cause mortality was higher among patients with lower NRI scores as shown in Figure 4. The all cause mortality was found to be 6% in the No Risk group and 9% in the Mild Risk groups as compared to 15% and 19% in the Moderate and Severe Risk groups, respectively (P<0.001).

Kaplan-Meier survival analysis showed that patients with lower NRI scores and therefore in Moderate or Severe Risk groups were prone to have 6-fold higher event rate comprising either frequent readmissions or all cause mortality (HR=0.42; 95% CI 0.34-0.50 and P<0.0001) as compared to those with higher NRI scores and classified as in No or Mild Risk groups (Figure 5). Individually, Kaplan-Meier survival analysis to study free event survival as a function of NRI score showed that patients with Moderate or Severe Risk were prone to have three-fold higher event rate comprising either frequent readmissions or all cause mortality OR=1 (95% CI: 0.98-1.02; P = 0.69).

Association between NRI scores and BNP levels in ADHF patient

Another important finding of our study was that higher BNP levels among patients at moderate or severe nutritional risk were seen as compared to patients with no or mild nutritional risk on admission (Table 1). The median BNP levels were 698 in No Risk group, 902 in the Mild Risk group, 1096 in the Moderate Risk group and 1040 in the Severe Risk Group (P=0.001).

Table 1. Basic demographics and clinical parameters of ADHF patients.

| Parameter | No Risk (n=63) | Mild Risk (n=213) | Moderate Risk (n=168) | Severe Risk (n=168) | P |
|-----------|----------------|------------------|-----------------------|---------------------|---|
| Demographics | | | | | |
| Age | 68±14 | 72±14 | 72±14 | 68±15 | 0.006 |
| Sex, % men | 318 (53) | 205 (66) | 205 (66) | 205 (66) | 0.0004 |
| Hypertension (%) | 562 (84) | 61 (96) | 172 (80) | 134 (79) | 0.08 |
| Diabetes mellitus (%) | 342 (48) | 26 (41) | 81 (38) | 50 (30) | 0.001 |
| Smoking (%) | 96 (14) | 8 (12) | 42 (19) | 24 (13) | 0.08 |
| BMI, kg/m² | 31±7 | 25±4 | 23±5 | 24±7 | 0.001 |
| Hyperlipidemia (%) | 278 (42) | 22 (35) | 70 (33) | 61 (36) | 0.07 |
| History of heart failure | 464 (69) | 42 (66) | 149 (70) | 112 (66) | 0.55 |
| History of CAD | 195 (29) | 16 (24) | 62 (29) | 45 (27) | 0.63 |
| Albumin | 3.9±0.8 | 3.4±0.4 | 3.2±0.4 | 2.9±0.4 | 0.001 |
| Nutrition risk score (NRI) | 0.99±0.7 | 0.97±0.7 | 0.95±0.7 | 0.96±0.7 | 0.001 |
| Heart failure indices | | | | | |
| NYHA Class | 2.5±0.9 | 2.6±0.7 | 2.6±0.7 | 2.8±0.7 | 0.07 |
| Median BNP, pg/mL | 698 | 902 | 1096 | 1040 | 0.001 |
| eGFR, mL/min per 1.73 m² | 68±45 | 53±27 | 44±29 | 58±36 | 0.001 |
| Kidney function | | | | | |
| Creatinine, mg/dL | 2.0±1.2 | 1.6±1.3 | 2.1±1.9 | 1.9±1.9 | 0.54 |
| eGFR, mL/min per 1.73 m² | 68±45 | 53±27 | 44±29 | 58±36 | 0.001 |
| Table 2. Cox proportional-hazards regression predictors for the composite endpoint of all-cause mortality and heart failure readmission. |

| Covariate | Exp (b) | Univariate 95% CI | P | Multivariate 95% CI | P |
|------------|---------|-------------------|---|---------------------|---|
| Age | 1.01 | 1.00-1.02 | <0.0001 | 1.01 | 1.01-1.02 | 0.02 |
| BMI | 0.96 | 0.94-0.98 | <0.0001 | 1.01 | 0.98-1.02 | 0.69 |
| eGFR | 0.99 | 0.98-0.99 | <0.0001 | 1.00 | 0.99-1.00 | 0.29 |
| Known CAD | 1.61 | 1.24-2.10 | 0.0003 | 1.48 | 1.11-1.98 | 0.006 |
| Known HF | 1.25 | 0.97-1.61 | 0.08 | 1.1 | 0.83-1.43 | 0.47 |
| NRI score | 3.03 | 2.33-3.94 | <0.0001 | 3.1 | 2.34-4.22 | <0.0001 |
| > 3 Risk factors | 1.41 | 1.11-1.81 | 0.004 | 1.37 | 1.05-1.79 | 0.017 |

Figure 1. Effect of nri on length of hospital stay.

Figure 2. Effect of nri on heart failure readmission.
Discussion

This is the first study to describe a prognostic nutritional risk index (NRI), which enables quantitative determination of the risk of malnutrition related morbidity and mortality in cardiac failure patients. Our study demonstrates that nutritional depletion as assessed by NRI is associated with worse outcome in patients admitted with ADHF (Figures 4 and 5). NRI can be used as a valid expedient of nutritional status of the hospitalized chronic heart failure patients like other patient populations.

Heart failure patients with lower NRI scores have a greater chance of prolonged hospitalization (Figure 2), higher readmission rates with episodes of ADHF (Figure 3) and higher mortality in subsequent years (Figure 4). The results of our study display the high prevalence of malnutrition in chronic heart failure patients presenting with episodes of ADHF in the United States.

The results of our study conform with the studies in the past displaying higher risk of various complications in malnourished patients as compared to non-malnourished patients. With the recent emerging evidence of the prevalence of cardiac cachexia in heart failure patients and the associated proposed neuro-endocrine and metabolic components involved, assessment of nutritional status in heart failure patients now seems more important than ever. There appears to be an intricate interrelationship between nutritional risk, severity of illness and clinical outcomes. The higher median values of BNP levels in moderate and severe risk malnourished patients in our study certainly point towards a more comprehensive measure, which can be conducted by tools like NRI at hospital admission can certainly help in prompt identification of the additional danger at which these patients are at risk.

There is no universally accepted gold standard for defining malnutrition. Many physicians consider measurement of serum proteins such as albumin to be an adequate assessment of nutritional status. However, in hospitalized patients, albumin is influenced by multiple factors as a sufficient measure of nutritional status. Reduction of albumin in ADHF patients may reflect impaired metabolic conditions including hepatic dysfunction, dehydration, protein loss, renal dysfunction, infection and inflammation. Utility of measuring either serum albumin or ideal body weight alone in assessment of nutritional status in different patient populations has been questioned. Kyle et al. reported that in hospitalized patient albumin understimates the prevalence of malnutrition if taken alone as an indicator of malnutrition. It is important to note that the development of the cachectic state in CHF is a process that can only be proven by a documented weight loss measured in a non-edematous state. This weight loss should usually be observed over a period of more than six months. The use of ideal body weight in NRI formula rather than usual body weight certainly solves this problem and gives a more comprehensive measure, which can be easily calculated in ADHF patients. Serum albumin is negatively correlated with increased extracellular fluid volume. Weight is also affected by hydration status, but variations in hydration status contrast strongly with variations in albumin concentrations. The use of both indicators in the NRI minimizes confounding variables such as hydration status, an important fluctuating variable observed in heart failure patients. The Nutritional Risk Index appears to capture both nutritional risk severity in ADHF patients. The Nutritional Risk Index seems to capture both nutritional risk and poor clinical outcome in hospitalized patients independent of disease severity. Significant correlation was observed between severity score and anthropometric measurements. This observation conflicts with the results of Bouillane et al. but agrees with Buzby et al. The patients at moderate or severe nutritional risk by the NRI scoring system in our study were found to have lower BMI values (24±6 kg/m²) as compared to no or mild risk groups as has been shown in other studies. A noteworthy finding in our study was lower mean BMI values (23±5) in the Moderate Risk group as compared to mean BMI values (24±7) in the Severe Risk group (P=0.001).

Our study indicates that lower NRI scores on admission can provide considerable information about risks of extended hospitalization in ADHF patients. The present study agrees with previously reported results, that moderate or severe nutritional depletion by the NRI was sig-
nificantly associated with increased LOS. Our study results also indicate that in ADHF patients the LOS increased significantly with decreasing NRI scores (Figure 2). In the past, studies have shown that serum albumin at hospital admission correlated significantly with ICU days and LOS but reductions in serum albumin may reflect metabolic or inflammatory conditions rather than nutritional status. There were no significant associations between low BMI and low albumin levels with increased LOS as was previously observed by Engelman et al. Our study demonstrates that NRI discriminates nutritional risk far better than weight loss, albumin, or BMI alone or combined in ADHF patients. Various studies have also shown that LOS was associated with NRI regardless of underlying disease or even treatment. Associations between moderate and severe NRI scores and LOS were significant with and without adjustment for age. The left ventricular ejection fraction (LVEF %) or NYHA classes among groups were not significantly different so could not be considered confounders to this association (Table 1).

Clinical implications of nutritional risk and hospital cost

Our simulated hospital costs increased significantly with increased nutritional risk. We found hospital costs (missing data) were 5 times higher in patients with severe nutritional risk compared with patients with no risk. Actual hospital costs would most likely be greater than the estimated cost in severely malnourished patients because we did not account for costs of complications. Thus, nutritional risk contributed to increased hospital cost.

Conclusion

Nutritional depletion as assessed by Nutritional Risk Index is associated with worse outcome in patients admitted with ADHF. The NRI is a more reliable prognostic indicator of morbidity and mortality in hospitalized ADHF patients than are indices that use albumin or BMI alone. Therefore, we recommend the use of NRI to further risk stratify these patients for nutritional depletion assessment. In our study, 34% of the hospitalized ADHF patients had moderate or severe nutrition-related risk and would be suitable for nutritional supplementation. Whether this would have an effect on outcomes awaits further study.

Future direction

Proper assessments of nutritional status of heart failure patients on hospital admission can help to minimize if not prevent complications, readmissions and mortality rates. Demonstrating that treating malnutrition in chronic heart failure patients improves outcomes would be the most convincing evidence. However, evidence that aggressive treatment of malnutrition improves outcomes is surprisingly scarce and requires further exploration and research.

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