Poor Nutritional Status Is Associated with Arrhythmic Events on 24-Hour Holter Recording

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Highlights of the Study

- Malnutrition is associated with cardiovascular disease morbidity and mortality.
- It remains unclear whether the adverse outcomes occur because of arrhythmias in malnourished patients.
- Arrhythmias may be the cardiac consequence of malnutrition.

Keywords
Cardiology · Prognostic nutritional index · CONUT score · Arrhythmia

Abstract
Background: Malnutrition is associated with cardiovascular disease morbidity and mortality. Arrhythmias may be the cardiac consequences of malnutrition. Objectives: The objective of the study was to evaluate the association between prognostic nutritional index (PNI), Controlling Nutritional Status (CONUT) score, and arrhythmic events on 24-h electrocardiography (ECG) Holter recording in patients without manifested arrhythmia. Methods: In this retrospective analysis of 477 patients who underwent 24-h ECG Holter monitoring, PNI and CONUT score were calculated and patients were divided into tertiles according to PNI and into three groups according to CONUT score: 0: normal, 1–2: mild risk of malnutrition, ≥3: moderate-severe risk of malnutrition. Arrhythmic events were compared between PNI tertiles and CONUT score groups. Results: Total number of premature atrial contractions, premature ventricular contractions (PVCs), PVC burden, and incidence of paroxysmal atrial fibrillation (PAF) were significantly higher in patients within the lowest PNI tertile. Total number of PVCs, PVC burden, and incidence of PAF were significantly higher in patients with CONUT score ≥3. The cut-off value for PNI to predict the presence of PVC was defined as 39.41 using ROC curve analysis. The area under the curve was 0.650 (p < 0.001). Multivariate analysis showed that PNI was independent predictor of the presence of PVC and PAF. Also, CONUT score was independent predictor of the presence of PVC and PAF. Incidence of nonsustained ventricular tachycardia did not differ between PNI tertiles or CONUT score groups. Conclusion: Poor nutritional status, assessed by PNI and CONUT score, is associated with arrhythmic events on 24-h ECG Holter recording in patients without manifested arrhythmia.

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Introduction

Malnutrition is reported to be associated with a poor prognosis in patients with heart failure or atherosclerotic cardiovascular disease (CVD) [1, 2]. The prognostic nutritional index (PNI) and Controlling Nutritional Status (CONUT) score are easily calculated and established methods for evaluating the objective nutritional status. Numerous clinical studies have indicated that PNI and CONUT scores are associated with CVD morbidity and mortality in patients with heart failure and myocardial infarction or in patients undergoing percutaneous coronary intervention or coronary artery bypass grafting [3–6]. However, no previous study has exclusively investigated the predictive value of nutritional indices for arrhythmic events. Therefore, we examined the relationship between PNI and CONUT score and arrhythmias on 24-h electrocardiography (ECG) Holter recording in patients without a previous diagnosis of any arrhythmia.

Methods

Study Population

This study is a retrospective analysis derived from a single center. We analyzed the data of patients who had 24-h ECG Holter recording from consecutive outpatients who applied to our outpatient clinic with complaints of palpitation, irregular heartbeats, and arrhythmia between January 2019 and December 2020. Patients with missing serum albumin, total cholesterol, and total lymphocyte count were excluded from this study. Also excluded were patients with known malignancy, active infectious or inflammatory disease, severe renal or liver insufficiency, electrolyte abnormalities, hypothyroidism, left ventricular ejection fraction (LVEF) less than 50% (assessed by echocardiography), moderate or severe valvular heart disease, persistent or permanent atrial fibrillation on baseline ECG, previous diagnosis of arrhythmia, previous ablation for arrhythmia, receiving antiarrhythmic therapy, atrial fibrillation (PAF) and supraventricular tachycardia (SVT) or paroxysmal supraventricular tachycardia (SVT) or nonsustained ventricular tachycardia (NSVT) were obtained from the 24-h ECG Holter recording. Three or more consecutive PACs were considered as SVT. NSVT was defined as three or more consecutive regular ventricular beats at a rate of greater than 100 beats/minute with a duration of less than 30 s. The arrhythmic beats and events on Holter recording were verified by two experienced cardiologists.

Evaluation of Nutritional Indices

The baseline PNI was calculated as 10 × serum albumin (g/dL) + 0.005 × total lymphocyte count (per mm³) [7]. Baseline CONUT score was calculated from serum albumin levels, total cholesterol levels, and total lymphocyte counts as previously reported [8]. CONUT scores range from 0 to 12. A person with normal nutritional status would have a CONUT score of 0, and the higher the score, the worse the nutritional status. Nutritional indices were categorized based on previous studies and defined classifications. Patients were divided into tertiles according to the PNI. Also, patients were classified according to CONUT score as follows: CONUT score of 0 – normal, 1–2 – mild risk of malnutrition, ≥ 3 – moderate-severe risk of malnutrition, as described [9].

Twenty-Four-Hour ECG Holter Recording

The number of premature atrial contractions (PACs) and premature ventricular contractions (PVCs), PVC burden (%) defined as total number of PVC/total beats, and presence of paroxysmal atrial fibrillation (PAF) and supraventricular tachycardia (SVT) or nonsustained ventricular tachycardia (NSVT) were obtained from the 24-h ECG Holter recording. Three or more consecutive PACs were considered as SVT. NSVT was defined as three or more consecutive regular ventricular beats at a rate of greater than 100 beats/minute with a duration of less than 30 s. The arrhythmic beats and events on Holter recording were verified by two experienced cardiologists.

Statistical Analysis

Continuous variables were expressed as mean ± standard deviation (SD) or median (interquartile range); categorical variables were defined as percentages. The distribution of continuous variables was considered as normal or not based on the Kolmogorov-Smirnov test. One-way ANOVA or Kruskal-Wallis tests were used to compare continuous variables. Conover-Inman or least significant difference tests were performed for the binary comparisons among the groups. Differences in the distribution of categorical variables were assessed by χ² test. Multiple logistic regression analysis with backward elimination was used to identify independent predictors of PVCs and PAF and was performed separately for PNI and CONUT scores. The selection of variables for the multivariate analysis was based on previous reports and clinical importance. Age, gender, LVEF, BMI, diabetes mellitus, hypertension, CAD, creatinine, the use of β-blockers, and the use of nondihydropyridine calcium channel blockers were included in the analysis as confounding variables. The area under the receiver operating characteristic (ROC) curve was used to indicate the predictive value of PNI for PVC. Pearson’s or Spearman’s correlation analysis was used to examine correlation between continuous variables. A p value of <0.05 was considered statistically significant. Data analyses were performed by using SPSS for Windows, version 22.0 (SPSS Inc., Chicago, IL, USA).
Results

Baseline Characteristics

The mean age of study population was 60.0 ± 17.8 years, and 42.2% of the 477 patients were males. Baseline characteristics are presented in Table 1. Mean LVEF was 61.4 ± 6.1%, mean PNI was 38.7 ± 4.9, and mean CONUT score was 1.36 ± 1.2. The baseline clinical and laboratory characteristics of the study patients according to the PNI tertiles are presented in Table 1. Patients in the lowest PNI tertile were significantly older and had a higher prevalence of hypertension, diabetes mellitus, and history of CAD as well as lower LVEF, hemoglobin, and higher CONUT score. Patients in the lowest PNI tertile more commonly received β-blockers. Total number of PACs, PVCs, and PVC burden were higher in the patients within the lowest PNI tertile. PAF and SVT were more commonly detected on 24-h ECG Holter recording in patients within lowest PNI tertile. The incidence of NSVT did not differ among the groups. The baseline clinical and laboratory characteristics of the study patients according to the CONUT score are presented in Table 2. Patients with higher CONUT scores were significantly older and had higher prevalence of hypertension and history of CAD as well as lower LVEF, PNI, triglyceride,
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Hemoglobin, and higher creatinine. The use of β-blockers and nondihydropyridine calcium channel blockers did not differ among the three groups. Patients with higher CONUT scores more commonly received statins. Total number of PACs was lowest in the patients with CONUT score 0. Total number of PVCs and PVC burden were highest in patients with CONUT score ≥3. PAF was more commonly detected on 24-h ECG Holter recording in patients with higher CONUT scores. The incidence of SVT and NSVT did not differ among the groups. The use of statin affects serum total cholesterol, and total cholesterol is a component of CONUT score. So, we excluded the patients receiving statins (18.2%) and reanalyzed the data of patients not receiving statins. Patients with higher CONUT score more commonly had hypertension, PAF, and SVT (p = 0.012, p = 0.010, and p = 0.035, respectively). Total number of PVCs and PVC burden were highest in the patients with highest CONUT scores (p < 0.001, p = 0.005, and p < 0.001, respectively). Total number of PACs was lowest in the patients with CONUT score 0. The incidence of NSVT did not differ among the groups (p = 0.171).

Correlations between PNI, CONUT Score, and Arrhythmic Events

PNI was strongly and negatively correlated with CO-NUT score (r = −0.767, p < 0.001). The number of PVCs positively correlated with number of PACs (r = 0.260, p < 0.001). Correlations between PNI and number of PACs (r = −0.244, p < 0.001), number of PVCs (r = −0.335, p < 0.001), and PVC burden (r = −0.342, p < 0.001) were sig-

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Table 2. Clinical and laboratory characteristics of patients stratified by CONUT score

| Variable                        | CONUT score = 0 (N = 204) | CONUT score = 1–2 (N = 175) | CONUT score ≥3 (N = 98) | p value |
|---------------------------------|---------------------------|-----------------------------|-------------------------|---------|
| Age                             | 58.0±16.7                 | 58.5±18.8                   | 67.6±16.5               | <0.001<sup>b, c</sup> |
| Male, %                         | 35.3                      | 49.1                        | 46.7                    | 0.117   |
| Hypertension, %                 | 47.3                      | 52.9                        | 74.2                    | <0.001  |
| Diabetes mellitus, %            | 19.4                      | 23.3                        | 28.1                    | 0.252   |
| CAD, %                          | 14.4                      | 29.1                        | 38.2                    | <0.001  |
| Smoking, %                      | 20.9                      | 24.4                        | 11.2                    | 0.042   |
| BMI, kg/m²                      | 24.6±3.3                  | 24.6±2.3                    | 23.4±1.3                | 0.030<sup>b, c</sup> |
| LVEF, %                         | 62.4±5.3                  | 61.4±5.9                    | 59.1±7.7                | 0.002<sup>b, c</sup> |
| Beta blocker, %                 | 31.3                      | 38.4                        | 40.4                    | 0.215   |
| Nondihydropyridine calcium channel blockers, % | 2.5                      | 4.1                          | 5.6                     | 0.402   |
| Statin, %                       | 11.9                      | 22.1                        | 28.1                    | <0.001  |
| PNI                             | 41.0±3.2                  | 39.3±4.1                    | 32.5±4.4                | <0.001<sup>a, b, c</sup> |
| Serum albumin, g/dL             | 4.03±0.42                 | 3.91±0.42                   | 3.1±0.53                | <0.001<sup>a, b, c</sup> |
| Total cholesterol, mg/dL        | 225.2±34.7                | 176.6±41.9                  | 167.5±44.0              | <0.001<sup>a, b</sup> |
| LDL cholesterol, mg/dL          | 146.3±32.2                | 106.7±35.2                  | 103.9±32.2              | <0.001<sup>a, b</sup> |
| HDL cholesterol, mg/dL          | 51.9±15.8                 | 49.2±14.2                   | 45.7±13.9               | 0.006<sup>b</sup> |
| Triglyceride, mg/dL             | 160.9±30.3                | 126.6±66.6                  | 123.5±50.3              | <0.001<sup>a, b</sup> |
| Creatinine, mg/dL               | 0.92±0.6                  | 1.04±0.74                   | 1.29±0.9                | <0.001<sup>a, b</sup> |
| White blood cell count, ×10⁹/L  | 7.45±2.5                  | 7.76±2.3                    | 8.19±4.2                | 0.685   |
| Hemoglobin, g/dL                | 14.0±1.5                  | 14.4±6.5                    | 12.7±3.1                | <0.001<sup>a, b, c</sup> |
| Lymphocyte count                | 2.42±0.7                  | 2.03±0.7                    | 1.52±0.7                | <0.001<sup>a, b, c</sup> |
| Heart rate (mean±SD), bpm       | 76.99±7.11                | 76.45±7.42                  | 77.84±6.35              | 0.379   |
| PACs, mean ± SD, n              | 638.2±2,322.6             | 1,075.2±2,944.8             | 1,562.6±4,080.7         | <0.001<sup>a, b</sup> |
| Median (interquartile range)    | 17 (127)                  | 37 (377)                    | 108 (1,012)             | <0.001<sup>a, b</sup> |
| PVCs, mean ± SD, n              | 1,422.2±5,055.8           | 1,826.0±8,414.9             | 1,802.5±3,813.9         | <0.001<sup>a, b</sup> |
| Median (interquartile range)    | 1 (276)                   | 11 (728)                    | 179.5 (1,963)           | <0.001<sup>a, b, c</sup> |
| PVC burden, mean ± SD, %        | 1.43±4.8                  | 1.84±4.6                    | 1.97±4.2                | <0.001<sup>a, b</sup> |
| Median (interquartile range)    | 0.00 (0.48)               | 0.01 (1)                    | 1 (2.1)                 | <0.001  |
| NSVT, %                         | 3.5                       | 8.4                         | 20.7                    | <0.001  |
| PAF, %                          | 8.4                       | 6.6                         | 20.7                    | <0.001  |
| SVT, %                          | 21.8                      | 23.4                        | 34.6                    | 0.070   |

Abbreviations as in Table 1. Significant differences were found between <sup>a</sup> CONUT score = 0 versus CONUT score = 1–2, <sup>b</sup> CONUT score = 0 versus CONUT score ≥3, and <sup>c</sup> CONUT score = 1–2 versus CONUT score ≥3.
Correlations between CONUT score and number of PACs ($r = 0.215$, $p < 0.001$), number of PVCs ($r = -0.206$, $p < 0.001$), and PVC burden ($r = -0.220$, $p < 0.001$) were significant.

**Multivariate and ROC Analyses**

Multivariate analysis revealed PNI as an independent predictor of the presence of PVC (OR = 0.936, 95% CI: 0.891–0.984, $p = 0.009$) and PAF (OR = 0.911, 95% CI: 0.853–0.973, $p = 0.005$) (Table 3). Also, multivariate analysis revealed CONUT score as an independent predictor of the presence of PVC (OR = 1.177, 95% CI: 1.010–1.372, $p = 0.037$) and PAF (OR = 1.210, 95% CI: 1.016–1.440, $p = 0.032$) (Table 4). The ROC curve of the PNI as a marker to predict the presence of PVC is illustrated in Figure 1. The cut-off value was 39.41 as identified by ROC. The area under the curve of the PNI for predicting the presence of PVC was 0.650 (95% CI: 0.599–0.701, $p < 0.001$). A PNI value of 39.41 had a sensitivity of 62% and specificity of 66.5%.

### Table 3. Multivariate analysis including PNI

| Predictors of PVCs | Odds ratio | 95% confidence interval | $p$ value |
|--------------------|------------|-------------------------|-----------|
| PNI                | 0.936      | 0.891–0.984              | 0.009     |
| Age                | 1.033      | 1.019–1.047              | <0.001    |
| LVEF               | 0.873      | 0.825–0.924              | <0.001    |
| Predictors of PAF  |            |                         |           |
| PNI                | 0.911      | 0.853–0.973              | 0.005     |
| LVEF               | 0.904      | 0.858–0.954              | <0.001    |
| CAD                | 1.267      | 1.106–1.671              | 0.005     |

Abbreviations as in Table 1.

### Table 4. Multivariate analysis including CONUT score

| Predictors of PVCs | Odds ratio | 95% confidence interval | $p$ value |
|--------------------|------------|-------------------------|-----------|
| CONUT score        | 1.177      | 1.010–1.372              | 0.037     |
| Age                | 1.037      | 1.023–1.051              | <0.001    |
| LVEF               | 0.881      | 0.833–0.932              | <0.001    |
| Predictors of PAF  |            |                         |           |
| CONUT score        | 1.210      | 1.016–1.440              | 0.032     |
| Age                | 1.041      | 1.016–1.066              | <0.001    |
| LVEF               | 0.908      | 0.861–0.957              | <0.001    |
| CAD                | 1.279      | 1.109–1.714              | 0.008     |

Abbreviations as in Table 1.

The present study showed that low PNI and high CONUT score were significantly associated with frequent PACs, frequent PVCs, and PAF detected on 24-h ECG Holter recording in patients without manifested arrhythmia. To the best of our knowledge, this is the first study to evaluate relationship between nutritional status, estimated by PNI and CONUT score, and arrhythmic events on 24-h ECG Holter recording. The findings of this study suggest that evaluation of nutritional risk may be important for predicting arrhythmic events in patients without manifested arrhythmia. The PNI and CONUT score were demonstrated to be objective and simple tools for assessment of nutritional status [7, 8]. Subsequently, a number of clinical studies were conducted to evaluate the association of the PNI and CONUT score with CVD morbidity and mortality in many patient cohorts. The PNI and CONUT score have been shown to predict adverse outcomes in patients with acute and chronic heart failure [1, 3, 10]. Also, PNI and CONUT score were found to be independent predictors of adverse outcomes in patients with CAD [4, 9, 11, 12]. Whether the adverse outcomes in these study cohorts are related to arrhythmias is still unclear.
Malnutrition is commonly prevalent in patients with several CVDs [1, 13, 14]. So far, however, not many studies have evaluated the relationship between nutritional indices, malnutrition, and arrhythmia. More data have recently been provided on the relationship between atrial fibrillation and nutritional indices. Zhu et al. [15] reported that poor nutritional status assessed by CONUT score is associated with recurrence of AF following ablation. In older patients with nonvalvular AF, moderate to severe malnutrition assessed by CONUT score and PNI was found to be significantly associated with adverse outcomes including thromboembolic events and death [14]. In previous studies using BMI as a tool to estimate nutritional status, being underweight was reported to be a risk factor for developing AF [16, 17]. In our study, incidence of PAF detected on 24-h ECG Holter recording was higher in those with lower PNI and higher CONUT score compared with patients with higher PNI and lower CONUT score. The main differences in our study are using simple, validated tools for estimating nutritional status, and study population consists of younger, ambulatory patients without manifest ed arrhythmia. In a meta-analysis including 198,000 patients without history of AF, PACs on 24–48-h ECG Holter recording were significantly associated with AF, first stroke, and mortality [18]. In our study, number of PACs and incidence of PAF were higher in patients with poor nutritional status.

Studies evaluating the relationship between malnutrition and arrhythmias are scarce. Even fewer studies have focused on the impact of malnutrition on arrhythmias other than AF. Malnutrition represented as low geriatric nutritional risk score was significantly associated with mortality in patients who underwent pacemaker implantation for bradycardia [19]. Recently, the relationship between BMI and ventricular arrhythmias has been studied. In a study including patients with implantable cardioverter-defibrillator and ischemic cardiomyopathy, patients with normal BMI compared to overweight and obese patients were found to have more recurrent ventricular arrhythmias and higher mortality [20]. But underweight patients were not included in this study. In a small study including hospitalized patients, malnourished patients (BMI <18.5) had longer QTc interval on ECG than nonmalnourished patients [21]. In patients with cardioverter-defibrillator, the frequency of arrhythmic events was significantly increased and the incidence of ventricular tachycardia/ventricular fibrillation was markedly higher in patients with stage IV cancer than in patients with earlier stages [22]. Cancer is a well-known risk factor for malnutrition, and there may be a link between malnutrition and ventricular arrhythmias in cancer patients, but we can only speculate about this link based on this study because of the lack of data on the nutritional status of participants. In our study, although the number of PVCs was higher in patients with poor nutritional status than in patients with better nutritional status, incidence of NSVT did not differ between groups.

Although there is increasing evidence on the relationship between malnutrition and arrhythmias, there is no clear explanation for the underlying pathophysiology. One possible mechanism for this relationship is inflammation, as malnutrition is associated with chronic inflammation and studies have suggested that there is a relationship between inflammation and arrhythmias [23–25]. Other possible mechanisms that may explain the increased risk of having arrhythmic event in the presence of malnutrition are electrolyte imbalance and deficiency of trace elements and vitamins [26]. Endothelial dysfunction is present in patients with arrhythmias and is associated with adverse outcomes, and malnutrition has been shown to be associated with endothelial dysfunction [27, 28]. It has been reported that malnutrition affects cardiac autonomic system in children and some studies in animals have suggested that protein malnutrition causes changes in renin-angiotensin system and sympathetic activity [29–31]. These changes may cause formation of an arrhythmia substrate or predisposition to arrhythmias.

This study has several limitations. One important limitation is that approximately 9 months of the study included the period of the COVID-19 pandemic. During this period, the increased stress and anxiety levels of the patients and changing lifestyles may have triggered changes in their nutritional status and development of possible arrhythmias. Effects of all potential confounding factors on arrhythmia cannot be controlled because of the retrospective design of the study. Also, there is the possibility of bias for nutritional indices from unmeasured confounders such as dietary habits or undiagnosed systemic illness. The observational nature of this study did not allow us to make a cause-and-effect explanation for the nutritional status that was associated with arrhythmic events. There is known daily variability in PAC and PVC frequency, and this might have affected the findings of our study. Further studies are needed to validate the clinical application of nutritional indices in the diagnosis and management of arrhythmias.
Conclusions

In our study, poor nutritional status, represented by low PNI and high CONUT score, was found to be associated with PACs, PVCs, and PAF on 24-h ECG Holter recording in patients without manifested arrhythmia. No single nutritional index to date has been integrated into routine cardiology practice, and findings of our study suggest that nutritional indices might be useful for predicting arrhythmic events.

Statement of Ethics

The research protocol was approved by the Mersin University Clinical Research Ethics Committee: Reference number: 76/2021.

Conflict of Interest Statement

The authors have no conflicts of interest to declare.

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Data Availability Statement

Derived data supporting the findings of this study are available from the corresponding author Ozge Kurmus Ferik on request.

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Author Contributions

Concept – Ozge Kurmus Ferik; design – Ozge Kurmus Ferik, Kursat Akbuga, Hatice Tolunay, and Turgay Aslan; supervision – Murat Eren, Aycan Fahrî Erkan, and Berkay Ekici; materials – Ozge Kurmus Ferik, Aycan Fahrî Erkan, Ebru Akgül Erkan, and Celal Kervancioglu; data collection and processing – Ozge Kurmus Ferik, Kursat Akbuga, Turgay Aslan, and Murat Eren; analysis and interpretation – Ozge Kurmus Ferik, Berkay Ekici, Murat Eren, and Hatice Tolunay; literature search – Ozge Kurmus Ferik; writing – Ozge Kurmus Ferik; critical review – Ebru Akgül Erkan, Celal Kervancioglu, and Hatice Tolunay.
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