The Baseline Nutritional Status Predicts Long-Term Mortality in Patients Undergoing Endovascular Therapy

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Received: 3 July 2019; Accepted: 26 July 2019; Published: 29 July 2019

Abstract: Introduction: Peripheral artery disease (PAD) occurs at an advanced stage of atherosclerosis and its comorbidities are associated with poor prognoses. Malnutrition is related to the severity of atherosclerosis in patients with cardiovascular disease and it predicts mortality. The Controlling Nutritional Status (CONUT) score is calculated from serum albumin concentration, peripheral lymphocyte count and total cholesterol concentration, and it robustly represents the nutritional status of hospitalized patients. This study aimed to determine the prognostic value of the CONUT score in patients with peripheral artery disease (PAD) who were undergoing endovascular therapy (EVT). METHODS and RESULTS: This study included 628 PAD patients who underwent EVT between 2013 and 2017 and were assigned to low (CONUT score 0: n = 81), mild (CONUT score 1–2: n = 250), moderate (CONUT score 3–4: n = 169), and high (CONUT score ≥ 5: n = 128) risk groups. The study’s primary endpoint was any death. Patients in the groups with higher CONUT scores were more likely to have chronic kidney disease (p < 0.001), impaired left ventricular ejection fractions (p < 0.001), and critical limb ischemia (p < 0.001) on admission. During follow-up, 95 patients (15%) died. Kaplan–Meier analyses revealed that the patients with higher CONUT scores had lower survival rates (p < 0.001; log-rank trend test). Multivariate Cox regression analyses showed that following adjustments for the confounding factors, a higher CONUT score was significantly associated with any death (hazard ratio, 1.15; 95% confidence interval, 1.03–1.30). CONCLUSION: The simple index CONUT score at the time of EVT may predict long-term mortality in PAD patients.

Keywords: peripheral artery disease; nutrition; endovascular therapy; all-cause mortality

1. Introduction

Peripheral artery disease (PAD) occurs at an advanced stage of atherosclerosis, and its risk factors are associated with poor prognoses. The presence of risk factors of atherosclerosis including diabetes, hypertension, dyslipidemia, and smoking habit is developing the status of PAD, coronary artery disease (CAD), and ischemic stroke [1–4]. However, over 50% of PAD patients are asymptomatic [4,5]; therefore, they do not receive preventive intervention. Successful endovascular therapy (EVT) can restore blood flow to the distal regions of the limbs, but it cannot overcome the high adverse event rate, and the prognostic indicators for patients with PAD after EVT remain unclear. Recently, malnutrition on admission has been shown to be an independent predictor of mortality, and it is related to the severity of the atherosclerosis in patients with coronary artery disease (CAD) and in those with heart failure [6–9]. The evaluation methods of nutritional status in patients suffering various diseases have been proposed as the Controlling Nutritional Status (CONUT) score and Prognostic Nutritional Index (PNI) [10–14]. The CONUT score is calculated from the serum albumin level, peripheral lymphocyte count, and the
total cholesterol (TC) concentration. This score can be calculated only by the blood test, and it robustly represents the nutritional status of patients with CAD [10–12,15]. Accordingly, the CONUT scoring system is superior in terms of its convenience and had comparable prognostic accuracy compared to the PNI scoring system [16]. Based on the background above, we hypothesized that malnutrition is associated with poor clinical prognoses in patients with PAD who progressively develop arteriosclerosis. The aim of this study was to determine the CONUT score’s prognostic impact on patients with PAD who were undergoing EVT.

2. Materials and Methods

2.1. Study Design and Study Population

This single-center cohort study included consecutive patients who underwent EVT between April 2013 and June 2017. Patients were excluded if the laboratory data required to calculate the CONUT score were missing. The CONUT score was calculated based on a patient’s serum albumin level, peripheral lymphocyte count, and TC level (Supplemental Table S1). On admission, the enrolled patients were assigned to 4 risk groups based on their CONUT scores, as follows: low-risk group (CONUT score = 0), mild-risk group (CONUT score = 1–2), moderate-risk group (CONUT score = 3–4), and high-risk group (CONUT score > 5) [8]. A baseline CONUT score of >2 is defined as malnutrition, and among hospitalized patients who had acute heart failure or CAD, those with CONUT scores of >3 were considered high risk [17,18]. This study was carried out in accordance with the principles of the Declaration of Helsinki. Our institution’s ethics committee approved the study’s protocol. Before study entry, the patients provided written informed consent that allowed the use of the data from their medical records only. The study was registered with the University Hospital Medical Information Network Clinical Trials Registry (UMIN000029848).

2.2. Treatment Protocols and Definitions

EVT was scheduled for patients who were diagnosed with PAD and who had a proven angiographic stenosis or an occlusion in a peripheral artery that was verified using imaging modalities, for example, enhanced computed tomography, magnetic resonance imaging or angiography. The EVT procedures were, in principle, performed using 6-Fr (guiding) sheaths for target lesions within the aortoiliac to popliteal arteries and 4.5-Fr guiding sheaths for isolated below-the-knee (BTK) lesions. Provisional stenting was performed on aortoiliac to popliteal lesions when flow limitations arose after balloon angioplasty that were caused by major arterial dissections, and balloon angioplasty alone was performed on BTK lesions in accordance with the healthcare reimbursement system in Japan. EVT that achieved at least “one straight line” to the vascular beds below the ankle was considered successful in the patients with critical limb ischemia (CLI). To prevent contrast-induced nephropathy, all patients with chronic kidney disease (CKD), except those on maintenance hemodialysis (HD), were given saline at a rate of 1.0 mL/kg/h for at least 12 h before the contrast was administered. EVT with carbon dioxide injections was considered on non-HD patients with end-stage renal dysfunction. All patients received aspirin (100 mg) and/or clopidogrel (75 mg) daily for at least 1 month post-EVT. In principle, patients with newly implanted stents received both aspirin and clopidogrel. The durations of the administration of the antiplatelet and other medical treatment regimens were at the discretion of the attending physicians. Dermatologists and plastic surgeons independently undertook wound care and made decisions about amputations during their regular clinic visits. CLI was defined as chronic ischemic rest pain, ulcers, or gangrene that was attributable to objectively proven arterial occlusive disease [9]. Diabetes was defined as a fasting plasma blood glucose level of ≥126 mg/dL on 2 separate occasions or ≥200 mg/dL on any single occasion.
2.3. Data Collection and Assessment

We evaluated the patients’ baseline characteristics, including their ages, sexes, body mass indexes (BMIs), complete blood counts, which comprised the total white blood cell, neutrophil, and lymphocyte counts, and the hemoglobin levels, blood chemistry test results, which included the albumin, blood urea nitrogen (BUN), creatinine, sodium, C-reactive protein (CRP), brain natriuretic peptide (BNP), lipids, and hemoglobin A1c levels, and the estimated glomerular filtration rates (eGFRs), and the oral medications administered at the time of EVT. Data describing the patients’ Rutherford classifications and histories of smoking, hypertension, diabetes, dyslipidemia, CKD, and regular HD were collected from their medical records, and the target EVT lesions were documented based on the angiography findings collected from the medical records. The participants’ left ventricular ejection fractions (LVEFs) were determined using contrast ventriculography or echocardiography. The follow-up data were obtained from inpatient and outpatient medical records.

2.4. Study Endpoint

We compared the groups’ clinical and angiographic profiles, medications at the time of EVT, and long-term prognoses. The study’s primary endpoint was any death. The patients who had and did not have CLI and those who did and did not undergo regular HD were compared in relation to the study’s primary endpoint.

2.5. Statistical Analyses

Data are expressed as the means and the standard deviations or as numbers and percentages. After testing the distribution of each parameters by the Shapiro–Wilk test, none of them were normally distributed. Therefore, the non-parametric equivalent Mann–Whitney-U test were used to compare the two groups, and the analysis of variance and Kruskal–Wallis test were used to compare among the ≥3 groups with respect to the continuous variables, and the Chi-squared test and Fischer’s exact test were used to compare the groups with respect to the categorical variables, as appropriate. Kaplan–Meier analyses were performed to evaluate the differences among the groups in relation to all-cause mortality. Univariate and multivariate Cox regression analyses were performed to evaluate the associations between the baseline parameters, including the CONUT score categories, and the prognosis. For the multivariate analyses, age, sex, the laboratory data, including hemoglobin, eGFR, BUN, CRP, and BNP, LVEF, and CLI were used to adjust the models. Further, the additive discriminative ability of the CONUT score was evaluated by comparing the areas under the receiver operating characteristic (AUROCs) curves using binary techniques that involved 2 models that predict the primary endpoint, based on the method described by DeLong et al. [19]. To compare the AUROCs between the CONUT score and each of its components, the Z-test was applied. Two-tailed p-values < 0.05 were considered statistically significant. Statistical analyses were performed using R software version 3.3.0 (R Foundation for Statistical Computing, Vienna, Austria).

3. Results

3.1. Patients’ Baseline Characteristics

After excluding 77 patients whose laboratory data were incomplete, we evaluated data from 705 consecutive patients with PAD and for whom the CONUT scores could be calculated. Ultimately, 628 patients were enrolled to participate in this study (Figure 1). The patients were assigned to four risk groups based on their CONUT scores, as follows: low-risk group: n = 81 (12.9%); mild-risk group: n = 250 (39.8%); moderate-risk group: n = 169 (26.9%); and high-risk group: n = 128 (20.4%).
The patients’ mean CONUT score was 2.8 ± 2.2, and >67% of the patients had nutritional disturbances with CONUT scores of ≥2. Table 1 presents the patients’ baseline clinical characteristics. The patients’ mean age was 69 years. Male patients accounted for 69% and patients while CLI comprised 42% of the study population. Half of the study population underwent regular HD, and almost 70% of the patients had diabetes. In addition to differences with respect to the components of the CONUT score, significant differences were evident among the groups in relation to a range of parameters. The groups with higher CONUT scores contained more patients who were male, with lower BMIs, with CKD, who underwent HD, with impaired LVEFs, with histories of revascularization, and more patients with lower hemoglobin and low-density lipoprotein (LDL)-cholesterol levels and higher BUN, CRP, and BNP levels, and whose renal function was worse. The groups with higher CONUT scores had higher CLI rates. The administration of β-blockers and cilostazol at the time of EVT differed significantly among groups. Supplemental Table S2 exhibits baseline characteristics in patients with and without CLI. Patients with CLI also had poorer clinical backgrounds compared to those without CLI. In patients classified with CONUT score, different trends were observed in sex, dyslipidemia, prior revascularization, WBC, neutrophil numbers, and EVT in below-the-knee lesions. Supplemental Table S3 presents baseline clinical characteristics in patients with and without regular HD. Patients with HD generally had poorer clinical backgrounds compared to those without HD. When patients in each group were classified with CONUT score, different trends were observed in sex, BMI, prevalence of hypertension, impaired LVEF, neutrophil numbers, and prescription of β-blockers between patients with and without HD.
The Kaplan–Meier curves for all-cause mortality, with the curves representing the CONUT risk groups, that is, Low-Risk (CONUT score 0), Mild-Risk (CONUT score 1–2), Moderate-Risk (CONUT score 3–4), and High-Risk (CONUT score ≥ 5), are shown in Figure 2. The Kaplan–Meier survival analyses showed that in the presence or absence of CLI, the patients with higher CONUT scores had lower survival rates compared with those in the patients with lower CONUT scores ($p < 0.001$; log-rank trend test). The Kaplan–Meier survival analyses showed that in the presence or absence of CLI, the patients with higher CONUT scores had lower survival rates compared with those in the patients with lower CONUT scores ($p < 0.001$; log-rank trend test) (Figure 3). Compared with the patients in the low-, mild-, and moderate-risk groups who did not have CLI, the patients in the high-risk group who did not have CLI had lower survival rates compared with those in the patients with the low-risk group ($p = 0.001$; log-rank trend test).
not have CLI had a markedly lower survival rate. Supplemental Table S4 shows the causes of death. During the observation period, the death rate was more than two-fold higher for the patients with CLI (22.8%), compared with that for the patients who did not have CLI (9.6%). Systemic bacterial infection was the leading causes of death in both groups. Systemic bacterial infections, pneumonae and sudden death in patients with CLI were significantly higher than those in patients with non-CLI (Supplemental Table S4). Additionally, the CLI patients with higher CONUT score (CONUT >5 and 3–4) also had higher amputation rate than those with lower CONUT score (CONUT 0–2) (log-rank trend test: \( p < 0.001 \)) (Supplemental Figure S1).

Supplemental Figure S2 shows Kaplan–Meier curves for all-cause mortality in patients with or without HD. The highest CONUT group (CONUT ≥ 5) had a poor prognosis compared with lower CONUT groups (CONUT 0–4) in PAD patients without HD. However, the higher CONUT score PAD patients with HD had the poorer prognosis.
3.3. Contribution of Each Conut Score Component to the Clinical Outcomes

An evaluation of the contribution of each component of the CONUT score to the primary endpoint showed that patients with higher scores for each component had lower survival rates (all $p < 0.001$; log-rank test) (Supplemental Figure S3A–C). Additionally, comparisons of the AUROC curves showed that the predictive value of the CONUT score (0.67; $p < 0.001$) was superior to that of any of the individual components of the CONUT score, namely, the lymphocyte count (0.64; $p < 0.001$), albumin level (0.59; $p = 0.004$), and the TC level (0.60; $p < 0.001$) (Supplemental Figure S3D).

3.4. Predictors of All-Cause Mortality.

The univariate analysis determined values of $p < 0.10$ for age, sex, BMI, CLI, an LVEF $<40\%$, hemoglobin, BUN, BNP, and eGFR (Table 2). The multivariate Cox hazard analysis determined that the CONUT score remained an independent predictor of all-cause mortality even after adjusting the model for multiple covariates (hazard ratio, 1.14; 95% confidence interval, 1.02–1.30) (Table 2).

Table 2. Univariate and multivariate analyses of any death.

| Variables          | Univariate | Multivariate |
|--------------------|------------|--------------|
|                    | HR         | 95% CI       | $p$-Value | HR         | 95% CI       | $p$-Value |
| CONUT score        | 1.31       | 1.21–1.41    | $<0.001$ | 1.14       | 1.02–1.30    | 0.03      |
| Age                | 1.03       | 1.01–1.05    | 0.01     | 1.04       | 1.01–1.07    | 0.003     |
| Male               | 1.61       | 0.99–2.60    | 0.05     | 1.34       | 0.79–2.29    | 0.28      |
| BMI                | 0.93       | 0.88–0.99    | 0.02     | 0.99       | 0.93–1.06    | 0.93      |
| CLI                | 3.06       | 2.01–4.65    | $<0.001$ | 1.32       | 0.79–2.21    | 0.29      |
| Diabetes           | 1.08       | 0.70–1.69    | 0.72     |            |             |           |
| LVEF $<40\%$       | 3.23       | 2.09–5.17    | $<0.001$ | 1.98       | 1.20–3.29    | 0.008     |
| Hemoglobin         | 0.74       | 0.67–0.83    | $<0.001$ | 0.95       | 0.81–1.10    | 0.49      |
| eGFR               | 0.98       | 0.97–0.99    | $<0.001$ | 0.99       | 0.98–1.00    | 0.07      |
| CRP                | 1.01       | 0.99–1.03    | 0.24     |            |             |           |
| BNP log            | 4.51       | 3.09–6.38    | $<0.001$ | 2.78       | 1.72–4.44    | $<0.001$ |

BMI = body mass index; BNP = brain natriuretic peptide; BUN = blood urea nitrogen; CI = confidence interval; CLI = critical limb ischemia; CONUT = Controlling Nutritional Status; CRP = C-reactive protein; eGFR = estimated glomerular filtration rate; HR = hazard ratio; LVEF = left ventricular ejection fraction.

4. Discussion

The principal finding from this retrospective analysis of patients with PAD who were undergoing EVT was that patients with a worse nutritional status were at a higher risk of all-cause mortality. Malnutrition is related to the severity of the atherosclerosis in patients with CAD and it predicts their mortality. The differences among risk groups regarding death became apparent soon after EVT, and they continued to increase during the observation period. This finding was independent of the presence of CLI. The CONUT score robustly represents a patient’s nutritional status, and it is a prognostic predictor in hospitalized patients.

The CONUT score was also useful for predicting the long-term prognoses of patients who did and did not undergo regular HD. Evaluating the associations between nutritional status and serious clinical events in such a substantial number of high-risk patients, of whom 46% underwent regular HD and 71% had diabetes, may have helped to clarify the prognostic problems associated with EVT.

Prior studies investigated a variety of nutritional supplementation and indicators (BMI, Nutritional Risk Screening—NRS-2002, Malnutrition Universal Screening Tool—MUST, PNI, geriatric nutritional risk index—GNRI, CONUT score, among others) [20,21]. Guidelines recommend some of the above as standard nutritional indicators; however, these indices require many results from uncommon laboratory tests first, and little has been evaluated regarding their prognosis-predicting efficacy in atherosclerotic patients. Moreover, in terms of cost-effectiveness, the CONUT score only
needs parameters to be collected routinely, in spite of its comparable predicting efficacy for long-term prognosis to full nutritional assessment [10].

Since PAD is a polyvascular disease and at an advanced stage of atherosclerosis, the level of systemic inflammation on PAD is higher than that on CAD [22]. Additionally, developing PAD has multidisciplinary mechanisms involving inflammatory cytokines and oxidative stresses [23]. The presence of atherosclerosis causes malnutrition and the presence of malnutrition may be one of the risk factors developing atherosclerosis [20]. Therefore, the prognostic prediction of PAD patients should be assessed to evaluate both of nutritional status and the degree of inflammation. The CONUT score is not the nutritional guideline recommended tool [21]; however, this score might reflect both of the nutritional status and the degree of inflammation in atherosclerotic patients including PAD. As a result, this simple index successfully stratified short and long-term mortality risks of the advanced-staged atherosclerotic patients.

The AUROC curve analysis has shown that the predictive value of the CONUT score was better than that of its individual components. Lymphocyte risk score and TC risk score classification made differences in mortality rates among groups gradually, as time advanced after EVT. In contrast, the patients with high albumin risk scores showed marked declines in their mortality rates soon after EVT. Lymphocytes represent the systemic inflammatory status [24–26]. Many previous studies’ findings have shown that the lymphocyte count and the neutrophil/lymphocyte ratio can predict long-term mortality or amputations in CLI patients [26–31]. Our study’s findings concur with the findings from these studies, and they confirm the prognostic value of the lymphocyte count. The TC level, which was a complex of lipid profiles, might reflect the effect of not only chronic inflammation, but also a patient’s daily nutritional status, and it is related to digestive functions; therefore, it may have caused linear changes in the primary endpoint. Conversely, the serum albumin level may reflect multiple organ failure caused by excessive levels of free water and reductions in protein metabolism [32,33]. Therefore, patients in the high-risk albumin category may be at a higher risk of reduced survival rate during the early phase after EVT. A combination of parameters that predict short-term and long-term changes in the adverse event rate may be useful for risk stratifying PAD patients who have severe atherosclerotic profiles and whose prognoses are difficult to determine.

Lowering of LDL-cholesterol levels appears to be associated with more favorable clinical prognoses for patients with atherosclerosis, including those with PAD and CLI [34–36]. However, findings from recent studies have shown worse outcomes in patients with malnutrition whose lipid profiles were controlled, which included low TC and triglyceride levels [37–39]. Since the triglyceride levels might reflect dietary habits, patients with advanced atherosclerosis and malnutrition, and especially those who undergo HD, should be treated carefully with lipid-lowering drugs.

Like previously reported findings [40,41], the leading cause of death in our study was systemic bacterial infection, the rate of which reached 30%, and it occurred at 30% in the patients with CLI who died and 25% in those without CLI. Bacterial infection in patients with severe atherosclerosis might be associated with any death.

These results indicate that the impaired immune defenses caused by malnutrition may determine the long-term prognosis, irrespective of the severity of the atherosclerosis. If the nutritional status of the patients with PAD had been well controlled before the de novo development of CLI, their prognoses might have been more favorable. Therefore, the nutritional status of patients with PAD should be evaluated during the early stages of atherosclerosis, and risk stratifications for critical events should be performed properly.

The comorbidity associated with regular HD is one of the most prominent risk factors related to adverse clinical events [42,43]. Indeed, the large proportion of the patients (64.7%) who died in this study had undergone HD. HD patients are malnourished [44], which augments severe limb infections and leads to death or amputations. Predicting the prognoses of patients who undergo regular HD is difficult, because they have high rates of a diverse range of comorbidities. However, the CONUT score successfully stratified the risk of death in this study. In this very high-risk population, the prognostic
value of the CONUT score was evident, even when these patients were compared with the patients who did not undergo HD. To date, an effective intervention has not been described that improves the clinical prognoses of patients with high CONUT scores. Hence, after determining the risks of adverse clinical events, nutritional supplementation and close observation should be considered for high-risk patients. To confirm this study’s findings, further interventional studies are warranted that will lead to improved outcomes after EVT.

The study population consisted of large numbers of patients who received regular HD, mainly because our institute has a huge dialysis center. The higher CONUT score group had higher prevalence of HD patients, and HD patients had higher mortality in this study. Given that the current study included the population at an extremely higher risk compared to that of prior trials and observations, it included a possible patient-selection bias, and it is not possible to generalize the findings as they are. However, among HD patients, the CONUT score at the time of EVT could stratify the risk of death during the long-term observation period, and even in the non-HD patients, it successfully extracted the high-risk population.

Collectively, the risk stratification of PAD patients, which was the population at an advanced stage of atherosclerosis, is meaningful for a better long-term prognosis, and the CONUT scoring system may be a promising option, independent of clinical presentation or HD, even it does not perfectly represent the ‘nutritional’ status in PAD patients.

5. Study Limitations

This study was a single-center cohort with a relatively small sample size and a limited observation period as well as with a potential patient selection bias, and it included a limited population who had received EVT. The CONUT score was evaluated only at the time of EVT, and we did not assess the effect of the change of score during the observation period on the prognosis. Furthermore, it is unclear, based on this study’s results, whether the stratified risk classes provide an intervention.

6. Conclusions

Determining the nutritional status of patients by simply calculating the lymphocyte counts and the albumin and TC levels in patients undergoing EVT was associated with adverse clinical events, including all-cause mortality.

Supplementary Materials: The following are available online at http://www.mdpi.com/2072-6643/11/8/1745/s1, Table S1: CONUT score, Table S2: Baseline characteristics: CLI vs. Non-CLI, Table S3: Baseline characteristics: HD vs. Non-HD, Table S4: Cause of death. Figure S1: Major amputation after Endovascular Therapy in Patients with Critical Limb Ischemia, Figure S2: All-cause Mortality After Endovascular Therapy in Patients Who Did and Did Not Undergo Hemodialysis, Figure S3: Combined Outcome of Death from Any Cause and Major Amputation After Endovascular Therapy in Patients in Relation to Each CONUT Score Component.

Author Contributions: K.M., acquisition of data, drafting of the manuscript; K.J., conception and design of the research, critical revision of the manuscript for important intellectual content; Y.M., analysis and interpretation of the data; I.I.; M.N., acquisition of data; N.H., supervision

Acknowledgments: We would like to thank Editage (www.editage.jp) for English language editing.

Conflicts of Interest: This study was not financially supported by any company, grant, or fund. The authors declare that they have no conflicts of interest and no relationships with industry. All other authors have reported that they have no relationships relevant to the contents of this paper to disclose.

References
1. Gerhard-Herman, M.D.; Gornik, H.L.; Barrett, C.; Barshes, N.R.; Corriere, M.A.; Drachman, D.E.; Fleisher, L.A.; Fowkes, F.G.R.; Hamburg, N.M.; Kinlay, S.; et al. 2016 AHA/ACC Guideline on the management of patients with lower extremity peripheral artery disease: A report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. Circulation 2017, 135, e726–e779. [PubMed]
2. Golomb, B.A.; Dang, T.T.; Criqui, M.H. Peripheral arterial disease: Morbidity and mortality implications. Circulation 2006, 114, 688–699. [CrossRef] [PubMed]
3. Hankey, G.J.; Norman, P.E.; Eikelboom, J.W. Medical treatment of peripheral arterial disease. *JAMA* 2006, 295, 547–553. [CrossRef] [PubMed]

4. Shu, J.; Santulli, G. Update on peripheral artery disease: Epidemiology and evidence-based facts. *Atherosclerosis* 2018, 275, 379–381. [CrossRef] [PubMed]

5. Hoyer, C.; Sandermann, J.; Petersen, L.J. The toe-brachial index in the diagnosis of peripheral arterial disease. *J. Vasc. Surg.* 2013, 58, 231–238. [CrossRef]

6. Kunimura, A.; Ishii, H.; Uetani, T.; Aoki, T.; Harada, K.; Hirayama, K.; Negishi, Y.; Shibata, Y.; Sumi, T.; Kawashima, K.; et al. Impact of nutritional assessment and body mass index on cardiovascular outcomes in patients with stable coronary artery disease. *Int. J. Cardiol.* 2017, 230, 653–658. [CrossRef] [PubMed]

7. Nochioka, K.; Sakata, Y.; Takahashi, J.; Miyata, S.; Miura, M.; Takada, T.; Fukumoto, Y.; Shiba, N.; Shimokawa, H.; CHART-2 Investigators. Prognostic impact of nutritional status in asymptomatic patients with cardiac diseases: A report from the CHART-2 Study. *Circ. J.* 2013, 77, 2318–2326. [CrossRef]

8. Lin, H.; Zhang, H.; Lin, Z.; Li, X.; Kong, X.; Sun, G. Review of nutritional screening and assessment tools and clinical outcomes in heart failure. *Heart Fail. Rev.* 2016, 21, 549–565. [CrossRef]

9. Sargento, L.; Longo, S.; Lousada, N.; dos Reis, R.P. The importance of assessing nutritional status in elderly patients with heart failure. *Curr. Heart Fail. Rep.* 2014, 11, 220–226. [CrossRef]

10. Igncio de Ulibarri, J.; González-Madroño, A.; de Villar, N.G.; González, B.; Mancha, A.; Rodriguez, F.; Fernández, G. CONUT: A tool for controlling nutritional status. First validation in a hospital population. *Nutr. Hosp.* 2005, 20, 38–45.

11. Sun, X.; Luo, L.; Zhao, X.; Ye, P. Controlling Nutritional Status (CONUT) score as a predictor of all-cause mortality in elderly hypertensive patients: A prospective follow-up study. *BMJ Open* 2017, 7, e015649. [CrossRef] [PubMed]

12. Nishi, I.; Seo, Y.; Hamada-Harimura, Y.; Sato, K.; Sai, S.; Yamamoto, M.; Ishizu, T.; Sugano, A.; Obara, K.; Wu, L.; et al. Nutritional screening based on the controlling nutritional status (CONUT) score at the time of admission is useful for long-term prognostic prediction in patients with heart failure requiring hospitalization. *Heart Vessels* 2017, 32, 1337–1349. [CrossRef] [PubMed]

13. Ahn, S.S.; Yoo, J.; Jung, S.M.; Song, J.J.; Park, Y.B.; Lee, S.W. Comparison of the Clinical Implications among five different nutritional indices in patients with lupus nephritis. *Nutrients* 2019, 11, 1456. [CrossRef] [PubMed]

14. Shirakabe, A.; Hata, N.; Kobayashi, N.; Okazaki, H.; Matsushita, M.; Shibata, Y.; Nishigori, S.; Uchiyama, S.; Asai, K.; Shimizu, W. The prognostic impact of malnutrition in patients with severely decompensated acute heart failure, as assessed using the Prognostic Nutritional Index (PNI) and Controlling Nutritional Status (CONUT) score. *Heart Vessels* 2018, 33, 134–144. [CrossRef] [PubMed]

15. Wada, H.; Dohi, T.; Miyauichi, K.; Doi, S.; Konishi, H.; Naito, R.; Tsuboi, S.; Ogita, M.; Kasai, T.; Okazaki, S.; et al. Prognostic impact of nutritional status assessed by the Controlling Nutritional Status score in patients with stable coronary artery disease undergoing percutaneous coronary intervention. *Clin. Res. Cardiol.* 2017, 106, 875–883. [CrossRef] [PubMed]

16. Basta, G.; Chatzianagnostou, K.; Paradossi, U.; Botto, N.; Del Turco, S.; Taddei, A.; Berti, S.; Mazzzone, A. The prognostic impact of objective nutritional indices in elderly patients with ST-elevation myocardial infarction undergoing primary coronary intervention. *Int. J. Cardiol.* 2016, 221, 987–992. [CrossRef] [PubMed]

17. Shimokawa, H.; Miura, M.; Nochioka, K.; Sakata, Y. Heart failure as a general pandemic in Asia. *Eur. J. Heart Fail.* 2015, 17, 884–892. [CrossRef]

18. Nakagomi, A.; Kohashi, K.; Morisawa, T.; Kosugi, M.; Endoh, I.; Kusama, Y.; Aitarashi, H.; Shimizu, W. Nutritional status is associated with inflammation and predicts a poor outcome in patients with chronic heart failure. *J. Atheroscler. Thromb.* 2016, 23, 713–727. [CrossRef]

19. DeLong, E.R.; DeLong, D.M.; Clarke-Pearson, D.L. Comparing the areas under two or more correlated receiver operating characteristic curves: A nonparametric approach. *Biometrics* 1988, 44, 837–845. [CrossRef]

20. Diana, P.B.; Alan, T.H.; Tadie, C.C.; Mindy, S.K. The role of nutrition and body composition in peripheral arterial disease. *Nutr. Res.* 2012, 32, 634–643.

21. Volkert, D.; Beck, A.M.; Cederholm, T.; Cruz-Jentoft, A.; Goisser, S.; Hooper, L.; Kiesswetter, E.; Maggio, M.; Raynaud-Simon, A.; Sieber, C.C. ESPEN guideline on clinical nutrition and hydration in geriatrics. *Clin. Nutr.* 2019, 38, 10–47. [CrossRef] [PubMed]
22. Reis, P.; Saely, C.H.; Silbernagel, G.; von Bank, A.; Mathies, R.; Drexel, H.; Baumgartner, I. Systemic inflammation is higher in peripheral artery disease than in stable coronary artery disease. *Atherosclerosis* 2015, 239, 299–303. [CrossRef] [PubMed]

23. Ross, R. Atherosclerosis: An inflammatory disease. *N. Engl. J. Med.* 1999, 340, 115–126. [CrossRef] [PubMed]

24. Horne, B.D.; Anderson, J.L.; John, J.M.; Weaver, A.; Bair, T.L.; Jensen, K.R.; Renlund, D.G.; Muhlstein, J.B. Which white blood cell subtypes predict increased cardiovascular risk? *J. Am. Coll. Cardiol.* 2005, 45, 1638–1643. [CrossRef] [PubMed]

25. Stabile, E.; Kinnaird, T.; la Sala, A.; Hanson, S.K.; Watkins, C.; Campia, U.; Shou, M.; Zbinden, S.; Fuchs, S.; Kornfeld, H.; et al. CD8+ T lymphocytes regulate the arteriogenic response to ischemia by infiltrating the site of collateral vessel development and recruiting CD4+ mononuclear cells through the expression of interleukin-16. *Circulation* 2006, 113, 118–124. [CrossRef] [PubMed]

26. Gary, T.; Pichler, M.; Belaj, K.; Hafer, F.; Gerger, A.; Froehlich, H.; Eller, P.; Pilger, E.; Brodmann, M. Neutrophil-to-lymphocyte ratio and its association with critical limb ischemia in PAOD patients. *PLoS ONE* 2013, 8, e56745. [CrossRef] [PubMed]

27. Chan, C.; Puckridge, P.; Ullah, S.; Delaney, C.; Spark, J.I. Neutrophil-lymphocyte ratio as a prognostic marker of outcome in infrapopliteal percutaneous interventions for critical limb ischemia. *J. Vasc. Surg.* 2014, 60, 661–666. [CrossRef] [PubMed]

28. Teperman, J.; Carruthers, D.; Guo, Y.; Barnett, M.P.; Harris, A.A.; Sedlis, S.P.; Pilling, M.; Babaev, A.; Staniloae, C.; Attubato, M.; et al. Relationship between neutrophil-lymphocyte ratio and severity of lower extremity peripheral artery disease. *Int. J. Cardiol.* 2017, 228, 201–204. [CrossRef]

29. González-Fajardo, J.A.; Brizuela-Sanz, J.A.; Aguirre-Gervás, B.; Merino-Díaz, B.; Del Río-Solá, L.; Martín-Pedrosa, M.; Vaquero-Puerta, C. Prognostic significance of an elevated neutrophil-lymphocyte ratio in the amputation-free survival of patients with chronic limb ischemia. *Ann. Vasc. Surg.* 2014, 28, 999–1004. [CrossRef]

30. Erturk, M.; Cakmak, H.A.; Surgit, O.; Celik, O.; Aksu, H.U.; Akgül, O.; Gurdogan, M.; Bulut, U.; Ozalp, B.; Akbay, E.; et al. The predictive value of elevated neutrophil to lymphocyte ratio for long-term cardiovascular mortality in peripheral arterial occlusive disease. *J. Cardiol.* 2014, 64, 371–376. [CrossRef]

31. Spark, J.I.; Sarveswaran, J.; Blest, N.; Charalabidis, P.; Asthana, S. An elevated neutrophil-lymphocyte ratio independently predicts mortality in chronic limb ischemia. *J. Vasc. Surg.* 2010, 52, 632–636. [CrossRef] [PubMed]

32. Van Hemelrijck, M.; Harari, D.; Garmo, H.; Hammar, N.; Walldius, G.; Lambe, M.; Jungner, I.; Holmberg, L. Biomarker-based score to predict mortality in persons aged 50 years and older: A new approach in the Swedish AMORIS study. *Int. J. Mol. Epidemiol. Genet.* 2012, 3, 66–76. [PubMed]

33. Wang, B.; Chen, G.; Cao, Y.; Xue, J.; Li, J.; Wu, Y. Correlation of lactate/albumin ratio level to organ failure and mortality in severe sepsis and septic shock. *J. Crit. Care* 2015, 30, 271–275. [CrossRef] [PubMed]

34. González, L.; Helkin, A.; Gahtan, V. Dyslipidemia part 2: Review of dyslipidemia treatment in patients with noncoronary vascular disease. *Vasc. Endovasc. Surg.* 2016, 50, 119–135. [CrossRef] [PubMed]

35. Drozda, J.P.; Jr; Ferguson, T.B., Jr; Jneid, H.; Krumholz, H.M.; Nallamothu, B.K.; Olin, J.W.; Ting, H.H. 2015 ACC/AHA focused update of secondary prevention lipid performance measures: A report of the American College of Cardiology/American Heart Association Task Force on Performance Measures. *J. Am. Coll. Cardiol.* 2016, 67, 558–587. [CrossRef] [PubMed]

36. O'Donnell, T.F.X.; Deery, S.E.; Darling, J.D.; Shean, K.E.; Mittleman, M.A.; Yee, G.N.; Dernbach, M.R.; Schermerhorn, M.L. Adherence to lipid management guidelines is associated with lower mortality and major adverse limb events in patients undergoing revascularization for chronic limb-threatening ischemia. *J. Vasc. Surg.* 2017, 66, 572–578. [CrossRef] [PubMed]

37. Dias, C.B.; Moughan, P.J.; Wood, L.G.; Singh, H.; Garg, M.L. Postprandial lipemia: Factoring in lipemic response for ranking foods for their healthiness. *Lipids Health Dis.* 2017, 16, 178. [CrossRef] [PubMed]

38. Mente, A.; Dehghan, M.; Rangarajan, S.; McQueen, M.; Dagenais, G.; Wielgosz, A.; Lear, S.; Li, W.; Chen, H.; Yi, S.; et al. Association of dietary nutrients with blood lipids and blood pressure in 18 countries: A cross-sectional analysis from the PURE study. *Lancet Diabetes Endocrinol.* 2017, 5, 774–787. [CrossRef]

39. Julius, U. Influence of plasma free fatty acids on lipoprotein synthesis and diabetic dyslipidemia. *Exp. Clin. Endocrinol. Diabetes* 2003, 111, 246–250. [CrossRef] [PubMed]
40. Frykberg, R.G. An evidence-based approach to diabetic foot infections. Am. J. Surg. 2003, 186, 44S–54S. [CrossRef]

41. Jaccard, Y.; Waltner, S.; Anderson, S.; Tauber, M.; Kummer, O.; Baumgartner, R.; Diehm, N.; Dörrfler-Melly, J.; Baumgartner, I. Influence of secondary infection on amputation in chronic critical limb ischemia. Eur. J. Vasc. Endovasc. Surg. 2007, 33, 605–609. [CrossRef] [PubMed]

42. Buronova, N.; Kamishima, K.; Yamaguchi, J.; Jujo, K.; Watanabe, E.; Inagaki, Y.; Kishi, S.; Wada, A.; Hatakeyama, S.; Watanabe, M.; et al. Effect of hemodialysis on 7-year clinical outcomes after sirolimus-eluting stent implantation. Circ. J. 2015, 79, 2169–2176. [CrossRef] [PubMed]

43. Okai, I.; Dohi, T.; Okazaki, S.; Jujo, K.; Nakashima, M.; Otsuki, H.; Tanaka, K.; Arashi, H.; Okabe, R.; Nagura, F.; et al. Clinical characteristics and long-term outcomes of rotational atherectomy—J2T multicenter registry. Circ. J. 2018, 82, 369–375. [CrossRef] [PubMed]

44. Sueta, D.; Hokimoto, S.; Sakamoto, K.; Akasaka, T.; Tabata, N.; Kaikita, K.; Honda, O.; Naruse, M.; Ogawa, H. Validation of the high mortality rate of Malnutrition-Inflammation-Atherosclerosis syndrome: Community-based observational study. Int. J. Cardiol. 2017, 230, 97–102. [CrossRef] [PubMed]

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