Letter to the Editor

Open Access

Test-retest reliability of the Mini Nutritional Assessment– Short Form (MNA-SF) in older patients undergoing cardiac rehabilitation

Fritz Kather^{1,*}, Miralem Hadzic^{1,*}, Teresa Hehle¹, Sarah Eichler¹, Julia Klein¹, Heinz Völler^{1,2},

Annett Salzwedel^{1,#}

¹Department of Rehabilitation Medicine, University of Potsdam, Faculty of Health Sciences Brandenburg, Germany ²Klinik am See, Rehabilitation Center for Internal Medicine, Rüdersdorf, Germany

J Geriatr Cardiol 2020; 17: 574-579. doi:10.11909/j.issn.1671-5411.2020.09.007

Keywords: Cardiac rehabilitation; Malnutrition; Octogenarians; Test-retest reliability

Malnutrition is described as a state of insufficient intake of energy, protein and other nutrients leading to changes in body composition (weight loss, reduced fat-free mass) as well as adverse functional and clinical outcomes.^[1] Depending on the assessments and definition used, the prevalence in older patients ranges between 12% in communitydwelling adults to 60% of patients in geriatric care facilities.^[2–5] Older populations are at high risk of nutritional deficiencies because of risk factors such as multimorbidity, polypharmacy, cognitive and physical decline, poor appetite, depressive syndromes and socioeconomic changes.^[4,6]

In particular, malnutrition often affects patients with cardiovascular diseases, e.g., chronic artery disease, aortic stenosis and other valve diseases.^[7–9] A recent study reported that 11.3% of older patients (84 ± 1 years of age) with severe aortic stenosis are malnourished and 42.3% are at risk of malnutrition.^[9] In addition, it is reported that 44% to 47% of patients over 80 years are at risk of malnutrition or malnourished after transcatheter aortic valve implantation (TAVI), the treatment of choice in the case of aortic stenosis in older people.^[10,11] Furthermore, 21.2% to 28.3% of patients over 65 years and 31.7% of older patients (83 ± 6 years) with coronary artery disease undergoing percutaneous coronary intervention (PCI) are malnourished or at risk of malnutrition.^[12,13]

Due to reduced muscle protein synthesis and insufficient protein intake, malnourished older patients often suffer from reduced muscle mass and, therefore, have a higher risk of sarcopenia.^[14–16] Even if malnutrition can lead to sarcopenia, both syndromes can also co-occur, as they share a similar pathophysiology.^[16,17] As a result, these patients experience

extensive consequences, e.g., a higher rate of postoperative complications, impaired immune function and wound healing, decreased functional status and higher risk of mortality compared to patients with normal nutritive status.^[7,12,18] Accordingly, early and accurate identification of patients who are in need of nutritional care is important to provide prompt nutrition counseling.^[1,18]

In Germany, cardiac rehabilitation is highly recommended for patients after PCI or transcatheter heart valve interventions.^[19] Currently, no assessments are used in clinical routine to identify patients at risk of malnutrition, which means that malnutrition often remains undiagnosed.^[8]

Various assessments have been developed to evaluate patients' nutritional status.^[20,21] The Mini Nutritional Assessment (MNA) is one of the most widely used tools especially in older patients.^[22,23] Its short form (MNA-SF) with six items has been shown to have a comparable validity to the original version,^[24] and high sensitivity and specifity of over 90%.^[25] Due to its validity and recommendations from different guidelines,^[21,26] the MNA-SF has the potential to become a standard screening tool in cardiac rehabilitation. However, information is lacking on its test-retest reliability.^[25,27]

Hence, we aimed to investigate the between-day test-retest reliability of the MNA-SF in older patients in cardiac rehabilitation who underwent catheter-based interventions. A prospective monocentric reliability study was carried out in a rehabilitation center for internal medicine in Rüdersdorf (Germany) between October 2018 and July 2019. Patients in cardiac rehabilitation after transcatheter aortic valve implantation (TAVI), intervention at the mitral or tricuspid valve (AVI) or percutaneous coronary intervention (PCI) with a minimum age of 75 years were enrolled. Exclusion

Å

^{*}The two authors contributed equally to this work.

^{*}Correspondence to: annett.salzwedel@fgw-brandenburg.de

http://www.jgc301.com; jgc@jgc301.com | Journal of Geriatric Cardiology

criteria were lack of comprehension (e.g., insufficient knowledge of the German language, dementia) or patients' refusal to participate in the study.

The MNA-SF was administered by trained raters within the first three days of cardiac rehabilitation followed by a second investigation one to two days afterwards. The questionnaire consists of six items (A: Decline of food intake over the past three months; B: Weight loss during the last three months; C: Mobility; D: Psychological stress or acute disease during the last three months; E: Neuropsychological problems; and F: Body Mass Index), which are answered by the patient or are evaluated using the data from the patients' records.^[24] Patients' neuropsychological decline was detected using the Mini-Mental State Examination with the cut-off \leq 24 points.^[28] Total scores of MNA-SF range from 0 to 14 points, with scores of 12 to 14 points indicating normal nutritional status, scores between 8 and 11 points identifying patients at risk of malnutrition, and scores < 8points diagnosing malnutrition.[29]

In addition, sociodemographic data (e.g., age, gender, body mass index) and clinical data (e.g., indications for rehabilitation, medication, comorbidities, left ventricular ejection fraction [LVEF], heart rhythm, New York Heart Association [NYHA] Functional Classification, laboratory data) were documented to describe the observed population. Laboratory data included hemoglobin, albumin and protein serum levels, as well as the estimated glomerular filtration rate (eGFR).

All patients gave their written informed consent to participate in the investigation, which was part of the FuNCaRe ("Functional and nutritional status of older patients in cardiac rehabilitation") study and had been approved by the ethics committee of the University of Potsdam (No.39/2018) and was registered in the German Register of Clinical Trials (DRKS00015176).

Continuous variables are displayed as mean \pm SD. Categorical variables are expressed as absolute values and percentages. The intra-class correlation coefficient (ICC) with 95% confidence interval (CI) was calculated in order to assess the test-retest reliability of the MNA-SF.^[30] The ICC presents values ranging from 0 to 1, with values < 0.5 indicating 'poor' reliability, 0.5 to 0.75 'moderate' reliability, 0.75 to 0.9 'good' reliability and values > 0.9 'excellent' reliability.^[31] The ICC is an indicator of relative reliability and is only related to group differences. Therefore, to give an indication of the accuracy of individual scores (absolute reliability), the minimal detectable change in percent (MDC%) was calculated based on the standard error of measurement (SEM).^[32,33] MDC% provides an estimate of the random

measurement error of the MNA-SF and, hence, determines whether a change in the score of individual patients is real at the 95% confidence level.^[34] A MDC% value under 30 is rated as acceptable and under 10 as excellent.^[35,36]

An analysis of variance (ANOVA) was used to compare patient characteristics, laboratory and clinical data and nutritional status between patient populations after TAVI, AVI and PCI. Statistical significance was set at P < 0.05(two-sided). Calculations were carried out using SPSS Version 25.0 (IBM) and Microsoft Excel 2013 (Microsoft Corporation).

A total sample of 122 patients (47.6% female, mean age 82.3 \pm 3.5 years) was included in the analysis (TAVI: n = 58; AVI: n = 21; PCI: n = 43). Patients' body mass index was 27.0 \pm 4.7 kg/m². In total, patients suffered from 5 \pm 2 comorbidities and took 9 \pm 3 pharmaceuticals, whereby patients after AVI had a significantly higher number of comorbidities compared to patients after TAVI and PCI (P < 0.001). In addition, patients after AVI showed significantly lower values in LVEF compared to the other groups (P < 0.001). Patient characteristics are presented in Table 1.

At baseline, patients reached a mean MNA-SF score of 11.7 \pm 2.3 points, whereby 75 patients (61.5%) showed normal nutritional status, 41 (33.6%) were at risk of malnutrition and 6 patients (4.9%) were malnourished. At the repeat measurement, 1.8 \pm 1.2 days after the baseline, the mean MNA-SF was 11.8 \pm 2.3 points, with an unchanged number of patients with normal nutritional status and (risk of) malnutrition (Table 2). Patients after AVI showed significantly different MNA-SF scores compared to patients after TAVI and PCI at both measurement points (P < 0.05, Table 2). An ICC value of 0.93 (95% CI: 0.90-0.95) and a MDC% of 15 was achieved for the MNA-SF score in the total sample (Table 2).

This study demonstrated 'excellent' test-retest reliability of the MNA-SF in patients after catheter-based interventions undergoing cardiac rehabilitation. According to MNA-SF, more than one-third of the patients investigated were malnourished or at risk of malnutrition, with small variations between the populations after TAVI, AVI or PCI.

In studies with patients after TAVI and of comparable age, similar mean total scores on the MNA-SF but a slightly higher percentage of patients with a risk of malnutrition were identified.^[10,11,37] These differences could be due to the heterogeneity of older populations regarding comorbidities, number of pharmaceuticals taken and functional status, which are all factors influencing the nutritional status.^[18,38] In addition, a recent study showed that one-fifth of patients

http://www.jgc301.com; jgc@jgc301.com | Journal of Geriatric Cardiology

Table 1. Patient characteristics.

	Total sample ($n = 122$)	TAVI (<i>n</i> = 58)	AVI $(n = 21)$	PCI $(n = 43)$	<i>P</i> -value
Age, yrs	82.3 ± 3.5	82.8 ± 3.3	82.8 ± 3.9	81.4 ± 3.6	0.145
Sex, female	58 (47.5%)	28 (48.3%)	10 (47.6%)	20 (45.5%)	0.985
BMI, kg/m ²	27.0 ± 4.7	27.1 ± 4.9	26.6 ± 4.4	27.1 ± 4.5	0.894
Comorbidities, no.	4.7 ± 2.0	4.7 ± 1.9	6.4 ± 2.2	3.7 ± 1.4	< 0.001
Arterial hypertension	100 (82.0%)	50 (86.2%)	16 (76.2%)	34 (79.1%)	0.491
Hyperlipidemia	49 (40.2%)	20 (34.5%)	8 (38.1%)	21 (48.8%)	0.339
Kidney disease	42 (34.4%)	17 (29.3%)	10 (47.6%)	15 (34.9%)	0.317
Diabetes mellitus	38 (31.1%)	17 (29.3%)	7 (33.3%)	14 (32.6%)	0.915
Orthopedic diseases	25 (20.5%)	8 (13.8%)	4 (19.0%)	13 (30.2%)	NA
Infections	15 (12.3%)	4 (6.9%)	7 (33.3%)	4 (9.3%)	NA
Cerebrovascular diseases	11 (9.0%)	6 (10.3%)	2 (9.5%)	3 (7.0%)	NA
Cancer	11 (9.0%)	4 (6.9%)	2 (9.5%)	5 (11.6%)	NA
Gastrointestinal diseases	12 (9.8%)	4 (6.9%)	2 (9.5%)	6 (14.0%)	NA
Pharmaceuticals, no.	8.9 ± 2.7	8.7 ± 2.6	9.1 ± 2.7	9.2 ± 2.9	0.734
Beta blockers	95 (77.9%)	42 (72.4%)	19 (90.5%)	34 (79.1%)	0.226
Diuretics	71 (58.2%)	34 (58.6%)	18 (85.7%)	19 (44.2%)	0.007
ACE inhibitors	56 (45.9%)	25 (43.1%)	8 (38.1%)	23 (53.5%)	0.428
Angiotensin II receptor antagonists	48 (39.3%)	25 (43.1%)	7 (33.3%)	16 (37.2%)	0.690
Opioid analgesics	15 (12.3%)	7 (12.19	3 (14.3%)	5 (11.6%)	NA
Anticonvulsants	7 (5.7%)	4 (6.9%)	1 (4.8%)	2 (4.7%)	NA
Antidepressants	7 (5.7%)	4 (6.9%)	2 (9.5%)	1 (2.3%)	NA
MMSE	25.9 ± 2.9	25.6 ± 3.2	26.7 ± 2.1	25.9 ± 2.7	0.298
NYHA					
Class I	59 (48.4%)	33 (56.9%)	8 (38.1%)	18 (41.9%)	0.074
Class II + III	63 (51.6%)	25 (43.1%)	13 (61.9%)	25 (58.1%)	
Echocardiography					
Sinus rhythm	81 (66.4%)	38 (65.5%)	5 (23.8%)	38 (88.4%)	< 0.001
Pacemaker/artrial fibrillation	41 (33.6%)	20 (34.5%)	16 (76.2%)	5 (11.6%)	
LVEF, %	53.7 ± 8.4	56.5 ± 6.2	49.3 ± 10.4	52.2 ± 8.7	< 0.001
Laboratory data					
Hemoglobin, mmol/L	7.5 ± 0.8	7.2 ± 0.7	7.4 ± 0.7	7.9 ± 0.9	< 0.001
Protein, g/dL	6.7 ± 0.4	6.7 ± 0.5	6.6 ± 0.6	6.5 ± 0.3	0.053
Albumin, g/dL	4.1 ± 0.2	4.1 ± 0.2	4.0 ± 0.3	4.1 ± 0.2	0.156
eGFR, mL/min per 1.73 m ²	65.9 ± 17.0	64.3 ± 14.7	67.4 ± 25.8	67.3 ± 14.5	0.612

Results are presented as mean \pm SD or *n* (%). ACE: angiotensin-converting enzyme; AVI: atrioventricular valve intervention; eGFR: estimated glomerular filtration rate; LVEF: left ventricular ejection fraction; MMSE: Mini-Mental State Examination; NA: not applicable; NYHA: New York Heart Association Functional Classification; PCI: percutaneous coronary intervention; TAVI: transcatheter aortic valve implantation.

after TAVI are referred to geriatric rehabilitation, which is administered for multimorbidity patients, and these patients have a lower mobility and nutritional status compared to patients referred to cardiac rehabilitation.^[39] Hence, mainly healthier patients were examined in this study, indicating a positive selection bias. Regarding patients after PCI, our results are in accordance with a recent investigation by Calvo *et al.* with older patients undergoing primary PCI.^[13] Remarkably, patients after AVI reached a statistically significant lower score on the MNA-SF and showed a significantly higher number of comorbidities as well as lower LVEF values compared to patients after TAVI or PCI. If patients suffer from a high number of comorbidities, it is more likely that they will have had an acute illness in the last three months, which is related to one item of the MNA-SF.^[24] Indeed, the number of comorbidities was negatively correlated to the MNA-SF score in our dataset. Furthermore, Fukui, *et al.*^[9] demonstrated that lower LVEF as a measure of heart function is related to malnutrition. However, our data did not confirm this association.

Journal of Geriatric Cardiology | jgc@jgc301.com; http://www.jgc301.com

MNA-SF results	Total sample ($n = 122$)	TAVI $(n = 58)$	AVI (<i>n</i> = 21)	PCI $(n = 43)$	<i>P</i> -value
First measurement					
MNA-SF score	11.7 ± 2.3	12.0 ± 1.8	10.3 ± 3.0	12.1 ± 2.2	0.006
Normal nutritional status	75 (61.5%)	36 (62.1%)	9 (42.9%)	30 (69.8%)	0.115
Risk of malnutrition	47 (38.5%)	22 (37.9%)	12 (47.1%)	13 (30.2%)	
Second measurement					
MNA-SF score	11.8 ± 2.3	12.0 ± 1.8	10.4 ± 3.1	12.0 ± 2.2	0.014
Normal nutritional status	75 (61.5%)	34 (58.6%)	11 (52.4%)	30 (69.8%)	0.336
Risk of malnutrition	47 (38.5%)	24 (41.4%)	10 (47.6%)	13 (30.2%)	
Test-retest reliability					
ICC (95% CI)	0.93 (0.90-0.95)	0.89 (0.83-0.94)	0.95 (0.89-0.98)	0.92 (0.86-0.96)	
MDC	15%	13%	15%	15%	

 Table 2.
 Nutritional status and test-retest reliability.

Results are presented as mean \pm SD or *n* (%). CI: confidence interval; ICC: intraclass correlation coefficient with < 0.5 "poor", 0.5–0.75 "moderate", 0.75–0.9 "good", > 0.9 "excellent"; MDC: minimal detectable change; MNA-SF: Mini Nutritional Assessment-Short Form with < 12 points "(risk of) malnutrition".

Laboratory parameters such as albumin, hemoglobin or protein are commonly used as markers to assess patients' nutritional status. These parameters are influenced by various factors, such as inflammatory processes, hydration status or renal impairment and therefore their eligibility in identifying malnutrition is controversial.^[11,40–42] More than one-third of the patients in this study had a kidney disease and suffered on average from five comorbidities, which might have had an impact on these parameters. Furthermore, these parameters can only detect patients who are already malnourished, which could be too late for optimal treatment strategies as malnutrition is reversible if detected early.^[38]

In contrast, the MNA-SF has the advantage of identifying patients at risk of malnutrition and is a reliable assessment according to ICC and MDC%.

These results are comparable to previous research by Bleda *et al.* and Lin *et al.*, who examined the test-retest reliability of the original version (MNA) in institutionalized elderly people and patients with stroke, respectively.^[32,43] The high reliability of both assessment tools can be explained by the way they are carried out. Firstly, the questionnaires are interviewer-administered, so questions can be clarified immediately in the case of misunderstanding by the older patients, which is an advantage over other methods (e.g., self-reporting).^[32,44] On the other hand, the data do not only rely on the patient's statement, but can partly be obtained from the patient's record, resulting in a high degree of objectivity.

Our study has some limitations. The targeted interval of one to two days between the first and second measurements of the nutritional status with the MNA-SF could not always be achieved because of patients' individual therapeutic plans during the rehabilitation process. Due to the short interval between the measurements of 1.8 days, it is possible that patients remembered their answers from the baseline investigation. Even if no interval between measurements is recommended, in comparable studies seven or more days are common.^[32,43] However, similar results were found between the cited studies and our results. Furthermore, differences between the defined patient groups should be interpreted with caution due to the small sample size.

In conclusion, the MNA-SF established good to excellent test-retest reliability for assessing nutritional status in older patients after catheter-based interventions. A high number of patients after AVI were affected by probable malnutrition, highlighting the importance of the assessment of malnutrition in older populations for timely implementation of preventive treatment. The MNA-SF can be highly recommended as a reliable screening tool for identifying patients at risk of malnutrition in cardiac rehabilitation.

Acknowledgements

We would like to thank all patients who participated in this study for their time and patience, as well as all physicians and study nurses in the cardiac rehabilitation center. All authors (FK, MH, TH, SE, JK, HV, AS) have no conflict of interest to declare. The authors declare that the study procedures comply with current ethical standards for research involving human participants in Germany. The study had been approved by the ethics committee of the University of Potsdam (No.39/2018). Written informed consent was obtained from all participants.

References

 Cederholm T, Jensen GL, Correia MITD, *et al.* GLIM criteria for the diagnosis of malnutrition - A consensus report from the global clinical nutrition community. *Clin Nutr* 2019; 38: 1–9.

- 2 Sánchez-Rodríguez D, Marco E, Ronquillo-Moreno N, et al. Prevalence of malnutrition and sarcopenia in a post-acute care geriatric unit: Applying the new ESPEN definition and EWGSOP criteria. *Clin Nutr* 2017; 36: 1339–1344.
- 3 Kaiser MJ, Bauer JM, Rämsch C, *et al.* Frequency of malnutrition in older adults: a multinational perspective using the mini nutritional assessment. *J Am Geriatr Soc* 2010; 58: 1734–1738.
- 4 Fávaro-Moreira NC, Krausch-Hofmann S, Matthys C, *et al.* Risk factors for malnutrition in older adults: a systematic review of the literature based on longitudinal data. *Adv Nutr* 2016; 7: 507–522.
- 5 Hegendörfer E, VanAcker V, Vaes B, Degryse JM. Malnutrition risk and its association with adverse outcomes in a Belgian cohort of community-dwelling adults aged 80 years and over. *Acta Clin Belg* 2020: 1–8.
- 6 Morais C de, Oliveira B, Afonso C, et al. Nutritional risk of European elderly. Eur J Clin Nutr 2013; 67: 1215–1219.
- 7 Emami S, Rudasill S, Bellamkonda N, *et al.* Impact of Malnutrition on Outcomes Following Transcatheter Aortic Valve Implantation (from a National Cohort). *Am J Cardiol* 2020; 125: 1096–1101.
- 8 Müller-Werdan U, Norman K. Aktuelle Aspekte kardiologischer Therapien bei älteren Menschen. Dtsch Med Wochenschr 2018; 143: 460–464.
- 9 Fukui S, Kawakami M, Otaka Y, *et al.* Malnutrition among elderly patients with severe aortic stenosis. *Aging Clin Exp Res* 2019; 32: 373–379.
- 10 Stortecky S, Schoenenberger AW, Moser A, et al. Evaluation of multidimensional geriatric assessment as a predictor of mortality and cardiovascular events after transcatheter aortic valve implantation. JACC Cardiovasc Interv 2012; 5: 489–496.
- 11 Goldfarb M, Lauck S, Webb JG, *et al.* Malnutrition and mortality in frail and non-frail older adults undergoing aortic valve replacement. *Circulation* 2018; 138: 2202–2211.
- 12 Wada H, Dohi T, Miyauchi K, *et al.* Combined effect of nutritional status on long-term outcomes in patients with coronary artery disease undergoing percutaneous coronary intervention. *Heart vessels* 2018; 33: 1445–1452.
- 13 Calvo E, Teruel L, Rosenfeld L, *et al.* Frailty in elderly patients undergoing primary percutaneous coronary intervention. *Eur J Cardiovasc Nurs* 2019; 18: 132–139.
- 14 Cruz-Jentoft AJ, Sayer AA. Sarcopenia. *Lancet* 2019; 393: 2636–2646.
- 15 Pacifico J, Geerlings MAJ, Reijnierse EM, et al. Prevalence of sarcopenia as a comorbid disease: A systematic review and meta-analysis. *Exp Gerontol* 2020; 131: 110801.
- 16 Reijnierse EM, Trappenburg MC, Leter MJ, *et al.* The association between parameters of malnutrition and diagnostic measures of sarcopenia in geriatric outpatients. *PloS One* 2015; 10: e0135933.
- 17 Vandewoude MFJ, Alish CJ, Sauer AC, Hegazi RA. Malnutrition-sarcopenia syndrome: is this the future of nutrition

screening and assessment for older adults? *J Aging Res* 2012; 2012: 651570.

- 18 Norman K, Pichard C, Lochs H, Pirlich M. Prognostic impact of disease-related malnutrition. *Clin Nutr* 2008; 27: 5–15.
- 19 S3-Leitlinie zur kardiologischen Rehabilitation (LL-KardReha) im deutschsprachigen Raum Europas Deutschland, Österreich, Schweiz (D-A-CH) (Langversion–Teil 1), 2019. www.awmf. org (accessed Aug 26, 2020).
- 20 Slee A, Birch D, Stokoe D. A comparison of the malnutrition screening tools, MUST, MNA and bioelectrical impedance assessment in frail older hospital patients. *Clin Nutr* 2015; 34: 296–301.
- 21 Schütz T, Valentini L, Plauth M. Nutritional screening according to the ESPEN Guidelines 2002. Akt Ernaehr Med 2005; 30: 99–103.
- 22 Guigoz Y, Vellas B, Garry PJ. Assessing the nutritional status of the elderly: The Mini Nutritional Assessment as part of the geriatric evaluation. *Nutr Rev* 1996; 54: S59–S65.
- 23 Phillips MB, Foley AL, Barnard R, *et al.* Nutritional screening in community-dwelling older adults: A systematic literature review. *Asia Pac J Clin Nutr* 2010; 19: 440–449.
- 24 Rubenstein LZ, Harker JO, Salvà A, et al. Screening for undernutrition in geriatric practice: Developing the short-form mini-nutritional assessment (MNA-SF). J Gerontol A Biol Sci Med Sci 2001; 56: M366–M372.
- 25 Skipper A, Ferguson M, Thompson K, *et al.* Nutrition screening tools: an analysis of the evidence. *JPEN J Parenter Enteral Nutr* 2012; 36: 292–298.
- 26 Volkert D. Ernährungszustand, Energie- und Substratstoffwechsel im Alter: Leitlinie Enterale Ernährung der DGEM und DGG. Ernährungs-Umschau 2004; 51(10).
- 27 Skipper A, Coltman A, Tomesko J, *et al.* Adult Malnutrition (Undernutrition) Screening: An Evidence Analysis Center Systematic Review. *J Acad Nutr Diet* 2019; 120: 669–708.
- 28 Folstein MF, Folstein SE, McHugh PR. "Mini-mental state": A practical method for grading the cognitive state of patients for the clinician. *J Psychiatr Res* 1975; 12: 189–198.
- 29 Kaiser MJ, Bauer JM, Ramsch C, *et al.* Validation of the Mini Nutritional Assessment short-form (MNA-SF): a practical tool for identification of nutritional status. *J Nutr Health Aging* 2009; 13: 782–788.
- 30 Koo TK, Li MY. A Guideline of Selecting and Reporting Intraclass Correlation Coefficients for Reliability Research. J Chiropr Med 2016; 15: 155–163.
- 31 Lahey MA, Downey RG, Saal FE. Intraclass correlations: There's more there than meets the eye. *Psychological Bulletin* 1983; 93: 586–595.
- 32 Lin SC, Lin KH, Lee YC, *et al.* Test-retest reliability of the Mini Nutritional Assessment and its relationship with quality of life in patients with stroke. *PloS One* 2019; 14: e0218749.
- 33 Weir JP. Quantifying test-retest reliability using the intraclass correlation coefficient and the SEM. J Strength Cond Res 2005; 19: 231–240.
- 34 Chen HM, Hsieh CL, Lo Sing K, et al. The test-retest reliabil-

Journal of Geriatric Cardiology | jgc@jgc301.com; http://www.jgc301.com

ity of 2 mobility performance tests in patients with chronic stroke. *Neurorehabil Neural Repair* 2007; 21: 347–352.

- 35 Huang SL, Hsieh CL, Wu RM, *et al.* Minimal detectable change of the timed "up & go" test and the dynamic gait index in people with Parkinson disease. *Phys Ther* 2011; 91: 114–121.
- 36 Smidt N, van der Windt DA, Assendelft WJ, et al. Interobserver reproducibility of the assessment of severity of complaints, grip strength, and pressure pain threshold in patients with lateral epicondylitis. Arch Phys Med Rehabil 2002; 83(8): 1145–1150.
- 37 Eichler S, Salzwedel A, Harnath A, *et al.* Nutrition and mobility predict all-cause mortality in patients 12 months after transcatheter aortic valve implantation. *Clin Res Cardiol* 2017; 107: 304–311.
- 38 Bauer JM, Wirth R, Volkert D, *et al.* Malnutrition, Sarkopenie und Kachexie im Alter--Von der Pathophysiologie zur Therapie. Ergebnisse eines internationalen Expertenmeetings der BANSS-Stiftung. *Dtsch Med Wochenschr* 2008; 133: 305–310.
- 39 Eichler S, Völler H, Reibis R, et al. Geriatric or cardiac reha-

bilitation? Predictors of treatment pathways in advanced age patients after transcatheter aortic valve implantation. *BMC Cardiovasc Disord* 2020; 20: 158.

- 40 Durán Alert P, Milà Villarroel R, Formiga F, et al. Assessing risk screening methods of malnutrition in geriatric patients: Mini Nutritional Assessment (MNA) versus Geriatric Nutritional Risk Index (GNRI). Nutr Hosp 2012; 27: 590–598.
- 41 Pirlich M, Schwenk A, Müller MJ. DGEM-Leitlinie Enterale Ernährung: Ernährungsstatus. Akt Ernaehr Med 2003; 28: 10–25.
- 42 Valentini L, Volkert D, Schütz T, *et al.* Leitlinie der Deutschen Gesellschaft für Ernährungsmedizin (DGEM). Aktuel Ernahrungsmed 2013; 38: 97–111.
- 43 Bleda MJ, Bolibar I, Parés R, Salvà A. Reliability of the mini nutritional assessment (MNA) in institutionalized elderly people. *J Nutr Health Aging* 2002; 6: 134–137.
- 44 Donini LM, Marrocco W, Marocco C, Lenzi A. Validity of the Self- Mini Nutritional Assessment (Self- MNA) for the Evaluation of Nutritional Risk. A Cross-Sectional Study Conducted in General Practice. *J Nutr Health Aging* 2018; 22: 44–52.

http://www.jgc301.com; jgc@jgc301.com | Journal of Geriatric Cardiology