- 1 The Mini Nutritional Assessment-Short Form and mortality in nursing home residents -
- 2 Results from the INCUR study

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- 13 Abstract
- 14 **OBJECTIVES:** To examine whether the Mini Nutritional Assessment-Short Form (MNA-
- 15 SF) score and its individual items are predictors of mortality in a nursing home population.
- 16 **DESIGN:** Prospective, secondary analysis from the Incidence of pNeumonia and related
- 17 ConseqUences in nursing home Residents (INCUR) study with 1-year follow-up.
- 18 **PARTICIPANTS:** A total of 773 older persons (women 74.4%) living in 13 French nursing
- 19 homes.

- 20 **MEASUREMENTS:** At baseline, nutritional status was assessed with the MNA-SF. Overall
- 21 mortality rate was measured over a 12-month follow-up period after the baseline assessment
- visit. Cox proportional hazard models were performed to test the predictive capacity of the
- 23 MNA-SF score and its single components for mortality.
- 24 **RESULTS:** Mean age of participants was 86.2 (standard deviation, SD 7.5) years. Mean
- MNA-SF score was 9.8 (SD 2.4). Among participants, 198 (25.6%) presented a normal
- nutritional status (12-14 points), 454 (58.7%) were at risk of malnutrition (8-11 points), and
- 27 121 (15.7%) were malnourished. After one year of follow-up, 135 (17.5%) participants had
- died. Age, female gender, baseline weight, BMI and MNA-SF were significant predictors of
- 29 mortality whereas no specific chronic disease was. The total MNA-SF score was a significant
- predictor of mortality (Hazard Ratio=0.81; 95% CI 0.74-0.90; p<0.001), even after adjustment
- 31 for potential confounders. Four individual items: weight loss, mobility, recent stress and BMI
- were independent predictors of mortality.
- 33 **CONCLUSIONS:** The MNA-SF appears to be an accurate predictor of one-year mortality in
- nursing home residents. Thus, this tool may be regarded not only as a nutritional screening
- tool, but also as an instrument for identifying the most-at-risk individuals in this population.

37 **Key words:** Older age; Mini Nutritional Assessment; Nursing Homes.

Introduction

Malnutrition is associated with adverse health outcomes in older subjects. It predicts hospitalization, infectious diseases (1) and death (2,3). Poor nutritional status is also related to increased health care expenditures (2). On the other hand, nutritional interventions have proven beneficial effects on weight gain and malnutrition-related outcomes such as morbidity and mortality (4). Therefore, there has been a growing interest in assessing the nutritional status of elders in order to facilitate the early detection of malnutrition and structure a proper management.

Although many instruments have been developed and validated for nutritional assessment (e.g. involuntary weight loss, Body Mass Index [BMI], albumin concentration, Mini-Nutritional Assessment [MNA] (5)), these tools have rarely been explored in nursing home (NH) residents (6). This population represents a highly vulnerable part of the heterogeneous geriatric patients, characterized by a high prevalence of chronic diseases, impaired cognitive and physical functions and limitations of activities of daily living (7). Many risk factors may also increase the risk for malnutrition in these subjects, such as polypharmacy (8) and multiple comorbidities (9). Unsurprisingly, the prevalence of malnutrition in NH population has shown to reach 30% (3).

The MNA test is a very commonly used assessment tool of nutritional status (5). It has shown great sensitivity, specificity and predictive positive value for malnutrition in elderly subjects (96%, 98% and 97% respectively), but needs 15 minutes to be completed. The MNA short form (MNA-SF) consists of 6 items and takes less than 5 minutes to complete. It was originally elaborated as a first step in the screening of malnutrition. A score of 11/14 or lower indicates a risk for malnutrition and triggers the administration of the full MNA questionnaire. Nevertheless, the MNA-SF has also been validated as an independent tool for nutritional screening in older adults (10). Interestingly, the items composing the MNA-SF are related to

functional or cognitive performance, and thus potentially provide information on multiple health domains over and above the mere nutritional status.

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However, there are still uncertainties regarding the ability of the MNA-SF to predict mortality in older adults. In a systematic review the MNA-SF (as well as the full MNA) was associated with higher mortality (Dent E, Visvanathan R, Piantadosi C, Chapman I Nutritional screening tools as predictors of mortality, functional decline, and move to higher level care in older people: a systematic review. J Nutr Gerontol Geriatr. 2012;31(2):97-145.) In a recent population-based study involving elders from Taiwan with a 4-year follow-up, the MNA-SF also appeared as an effective predictor of mortality. (Wang JY, Tsai AC. The short form Mini Nutritional Assessment is as effective as the full-Mini Nutritional Assessment in predicting follow-up 4-year mortality in elderly Taiwanese. J Nutr Health Aging 2013;17: 594–598.) On the other hand, another study found that the MNA-SF is not suitable to provide prognostic information in older adults with multiple comorbidities (Vischer UM, Frangos E, Graf C et al. The prognostic significance of malnutrition as assessed by the Mini Nutritional Assessment (MNA) in older hospitalized patients with a heavy disease burden. Clin Nutr 2012;31:113-117.) In the present study, we conducted longitudinal analyses aimed at examining the relationship between the MNA-SF and mortality in a sample of NH residents, over one year of follow-up. We also studied which items of the MNA-SF may independently explain this association.

Methods

Study design and participants

Data were from participants recruited as part of the Incidence of pNeumonia and related ConseqUences in nursing home Residents (INCUR) study, a prospective observational cohort study of 800 NH residents. The INCUR rationale, study design, and methodology have been previously described (11). The primary aim of INCUR was to estimate the incidence of pneumonia and the associated health-related expenditures in this population. The 6-month recruitment period started in February 2012. The INCUR project ended on June 2013 after all participants had been followed-up over 12 months.

Main eligibility criteria of INCUR included: age of 60 years and older; a functional status ranging from 2 to 5 at the Autonomie Gérontologie - Groupes Iso-Ressources (AGGIR) scale (i.e. the nationally recognized functional scale on which the allocation of social support is decided by public health authorities in France; a score between 2 and 5 excluded totally disabled patients as well as subjects with no impairment in basic activities of daily living) (12) residents living in the NH for more than 30 days. The design of the INCUR project was consistent with the Declaration of Helsinki and the study protocol was approved by the local Ethics Committee.

Two follow-up visits were scheduled after 6 and 12 months from the baseline visit. At these visits, besides of repeating the same multidimensional evaluation conducted at the baseline, the possible onset of major health-related events occurred during the past 6 months was ascertained. The present analyses were conducted in 773 subjects, after exclusion of 27 subjects with missing key data.

Variables of interest

At baseline, socio-demographic information, medical history, and comorbidities were recorded. Chronic diseases of interest were: atrial fibrillation, heart failure, coronary heart disease, respiratory conditions, history of stroke and stroke-related impairment, cancer, diabetes, Parkinson's disease and dementia. Weight and height were measured and BMI was calculated. Current smoking and oxygen therapy were also recorded. Cognitive function was assessed with the Abbreviated Mental Test scale (13). Depression was assessed with the 10-item Geriatric Depression Scale (14).

MNA-SF assessment

The MNA-SF consists of the first six items, also known as the "screening part" of the full MNA. Briefly these items are: A) food intake; B) involuntary weight loss; C) mobility; D) recent psychological stress or acute disease; E) neuropsychological problem (i.e. dementia or depression); and F) BMI. In case of missing value for this item (as frequently occurring in bed-ridden residents), the BMI item can be replaced by the calf circumference (measured with a tape). The MNA-SF score can range between 0 and 14 points with higher values indicating better nutritional status. The MNA-SF score is also usually categorized into three groups defining "normal" (12-14 points), "at risk" (8-11 points), and "malnutrition" (0-7 points) statuses.

Statistical analyses

Chi-squared tests and t-tests were used to describe the categorical and continuous characteristics of the study sample according to the outcome of interest, respectively. Cox proportional hazard models were used to evaluate the relationships of the MNA-SF score (as both continuous and categorical variable) and its composing items with mortality. Results are presented as hazard ratios (HR) and 95% confidence intervals (95% CI). Secondary analyses were also conducted using the single items composing the MNA-SF as independent variables of interest in the prediction of mortality. Although weight and height were significantly different between deceased subjects and survivors, these two variables were not included in the adjusted model because strongly correlated with the independent variables of interest. All statistical analyses were performed using SPSS statistical software version 18.0.0 (IBM Corp, New York). Statistical significance was defined as P<0.05. For all the single items significantly associated with mortality, sensitivity, specificity, positive and negative predictive values and positive and negative likelihood ratios were calculated.

Results

Descriptive characteristics of the study sample (n=773) according to the study outcome are presented in Table 1. One hundred and thirty five (17.4%) residents died during the 12 months of follow-up. Mean age of the study population was 86.1 (SD 7.5) years, with a higher prevalence (74.6%) of women. The mean MNA-SF score was 9.8 (SD 2.4). In the study sample, 198 persons (25.6%) had a normal nutritional status (MNA-SF 12-14 points), 454 (58.7%) were at risk of malnutrition (8-11 points), and 121 (15.7%) were malnourished.

Among the deceased residents, mean age was 88.5 (SD 6.9) vs. 85.7 (SD 7.5) in survivors (p<0.001). Women represented 76.7% of survivors vs. 63.1% of deceased (p=0.001). None of the chronic diseases was significantly associated with mortality. However, indicators of nutritional status were predictors of 1-year mortality: baseline weight and BMI were lower in NH residents who died (61.3 kg [SD 13.4] vs. 64.4 kg [SD 14.6], p=0.03; and 24.2 [SD 4.3] kg/m² vs. 25.4 [SD 5.3] kg/m², p=0.04 respectively) as well as the MNA-SF score (9.3 vs. 9.9, p=0.02; Table 1).

In Table 2, results from Cox-proportional hazard models examining the MNA-SF and one-year mortality were presented. The MNA-SF (continuous variable) was associated with a significantly lower risk of dying during the follow-up, even after adjustment for age and gender. When the MNA-SF score was categorized, malnourished subjects (0-7 points) showed a significantly higher risk of mortality (HR=4.64, 95%CI 1.79-12.0; p=0.002) compared to the reference group. A trend for association between being at risk of malnutrition and higher risk of mortality (HR=2.40; 95%CI 0.99-5.79; p=0.052) was also observed.

Similar results were found in secondary analyses exploring the individual components of the MNA-SF components and mortality (Table 3). Weight loss (p=0.02), BMI<21 kg/m² (or calf circumference<31 cm) (p=0.004), recent disease or psychological stress (p=0.01) and

lack of mobility (p=0.048) were all significant predictors of the studied outcome. Their sensitivity, specificity, positive and negative predictive values and likelihood ratios are respectively displayed in Table 4. When considered individually, the four latter items showed poor sensitivity for mortality. In contrast, the MNA-SF with a threshold of 12/14 had a correct sensitivity (88.5%) for mortality. Moreover, only a borderline significance was reported for the decrease in food intake item (p=0.053). The neuropsychological problem item was not associated with mortality (p=0.83).

Discussion

In the present prospective study, a low MNA-SF score was a strong predictor of death after one year of follow-up. A low BMI (or calf circumference) or recent weight loss were individual and significant predictors of mortality in our sample. Two other items: "functional impairment" and "recent acute stress", which are likely to reflect a more general status of frailty rather than malnutrition *sensu stricto*, were also significant predictors of mortality. On the other hand, education or clinical conditions, including depression and dementia were not.

Malnutrition dramatically affects the vulnerable older persons, in particular those living in institutions. Consistently to prior studies, only one quarter of our sample (25.6%) had a normal (i.e \geq 12) MNA-SF score whereas the other three quarters where either at risk of malnutrition (58.7%) or malnourished (15.7%) (15,16). A recent systematic review has examined the predictive validity of the available screening tools for malnutrition in NH populations (17). Authors concluded that none of them emerged as the gold-standard. Another study specifically assessed the usefulness of the MNA-SF for malnutrition screening in a NH population (18). This study considered a smaller sample (n=151) of institutionalized subjects compared to our work, and only 64.4% of undernourished patients were found to be correctly classified using this tool. Nevertheless, the MNA score demonstrated to be feasible and showed the best predictive capacity for survival (compared with Nutritional Risk Screening and the Malnutrition Universal Screening Tool) among well-nourished NH residents (15).

The use of the MNA-SF offers several advantages: this tool is standardized, reproducible, non-invasive, and takes only 5 minutes to be completed. Moreover, it is strongly correlated with the full MNA (19,20). Interestingly, the 6 items of the MNA-SF comprise three nutritional criteria (BMI, food intake and weight loss) as well as three criteria related to "geriatric conditions" (mobility, recent acute stress and neuropsychological disorder). Thus,

this tool may be specifically tailored for frail older persons and is likely to provide insights into the global health apart from the mere nutritional status.

In our study, the MNA-SF appeared as a predictor of mortality. Not only the total score, but also the score categories (i.e. people at risk of malnutrition and malnourished subjects) as well as most of the subitems when individually considered. Our results are supported by the study of Tangvik and colleagues who recently investigated the association between nutritional status and clinical outcomes (21). Authors have found that the combination of four criteria from the ESPEN guidelines for nutrition screening 2002 (22) (BMI <20.5 kg/m2 / Weight loss within the last weeks / Reduced dietary intake during the last weeks / Severe illness) was accurate to predict mortality, morbidity and hospitalizations in Norway hospital in-patients. Interestingly, such criteria are very similar to 4 MNA-SF items we found in our analyses. Thus, we may draw two main conclusions. First, our results are consistent with the established relationship between nutritional status and survival in institutionalized elderly (23,24). Second, the MNA-SF, may be regarded as a multidimensional instrument for identifying the most vulnerable individuals of an elderly population.

On the other hand, the absence of chronic conditions related with enhanced mortality might be surprising. This result may be explained by the high prevalence of coexisting chronic diseases in our population. Further, we did not take into consideration the severity of the diseases. Obviously, severe heart failure or dementia are a heavier burden than mild stages of these conditions and increase the risk of poor outcomes. Yet, the simple MNA-SF showed an additional value to identify NH residents at higher risk of death whereas specific pathological conditions did not. Two NH residents with the same multiple (but often stable) clinical conditions may be at different risk of dying given their nutritional status assessed with the MNA-SF. But the death event can be the consequence of an impaired

response to an acute stressor (e.g. infection) in polypathological individuals (25). Thus, the MNA-SF may be considered as multidimensional assessment tool, resembling the frailty ones (26,27), thus overcoming the single nosological entities commonly used in the clinical setting. Consistently, the "mobility" and the "acute stress" items of the MNA-SF both reflect functional performances and were predictors of mortality in our study. As such, our results highlight the relationship between risk of death and functional status rather than comorbidities.

The main strengths of our study were the large sample size and the prospective design. The representativeness of our sample was good, with few missing data (23 subjects i.e. less than four percent) despite one year of follow-up. On the other hand, some limitations have to be acknowledged. This study did not analyze biological markers of protein malnutrition (e.g. albumin concentration) that are independent risk factor for mortality in NH residents. We did not consider either the causes of death or some other potential confounding factors to explain the death. Yet, comorbidities, depression and dementia were not significantly associated with death in our analyses. Moreover, the individual items of the MNA-SF have been dichotomized instead of examining each category for each question of the form. Nevertheless, we aimed at preserving the clinical meaningfulness when combining different categories of a single item.

Conclusion

The MNA-SF and most of its subitems, but not clinical conditions, were significant predictors of overall mortality in NH residents, independently of potential confounders. Our findings support the use of this simple test in this population, not only for malnutrition screening but also for obtaining an overview of the general risk profile of these complex older adults. Therefore, the MNA-SF may pave the way not only for nutritional assessment but also

for comprehensive geriatric assessment and management of these vulnerable elders.

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 Table 1. Baseline characteristics of our population

Variable, M±SD	Death			
variable, MESD	No (n=638)	Yes (<i>n</i> =135)	P	
Age (years)	85.7±7.5	88.5±6.9	< 0.001	
Gender (women)	76.7	63.1	0.001	
Current smoking	2.5	3.1	0.66	
Education (years)	8.5 ± 3.3	8.1 ± 3.1	0.28	
Height (cm)	159.2±8.6	159.1±7.9	0.89	
Weight (kg)	64.4±14.6	61.3±13.4	0.03	
Body mass index (kg/m ²)	25.4 ± 5.3	24.2 ± 4.3	0.04	
Clinical conditions				
Atrial fibrillation	12.3	17.0	0.25	
Heart failure	26.9	33.3	0.26	
Coronary heart disease	5.4	8.2	0.34	
Respiratory disease	9.6	14.2	0.11	
Stroke	7.7	11.9	0.38	
Cancer	12.6	9.0	0.13	
Diabetes	15.0	14.0	0.79	
Parkinson's disease	5.9	5.9	0.98	
Dementia	35.3	33.3	0.32	
O ₂ therapy	1.8	2.2	0.70	
Abbreviated Mental Test score (/10)	5.7±3.6	5.0±3.4	0.06	
10-item Geriatric Depression Scale	2.9 ± 2.4	2.9 ± 2.5	0.84	
MNA-SF score (/14)	9.9 ± 2.4	9.3 ± 2.4	0.02	

Results are presented as means \pm SDs, or percentages

Table 2. Relationships of the Mini Nutritional Assessment-Short Form (MNA-SF) score with mortality over one year of follow-up in nursing home residents.

	Unadjusted HR (95% CI)	P	Adjusted* HR (95% CI)	P
MNA-SF score (continuous), n/N=52/773	0.83 (0.75, 0.91)	< 0.001	0.81 (0.74, 0.90)	< 0.001
MNA-SF score categories				
Normal nutritional status (12-14 points), n/N=6/198	1 (Reference group)		1 (Reference group)	
At risk of malnutrition (8-11 points), n/N=31/454	2.30 (0.96, 5.51)	0.06	2.40 (0.99, 5.79)	0.052
Malnourished (0-7points), n/N=15/121	4.31 (1.67, 11.10)	0.003	4.64 (1.79, 12.00)	0.002

CI: confidence interval; HR: Hazard Ratio; MNA-SF: Mini Nutritional Assessment-Short Form; n: number of deceased subjects/N: total number of subjects; *Adjusted for age and gender

	Unadjusted HR for mortality (95% CI)	P	Adjusted* HR for mortality (95% CI)	P
Decrease in food intake over the past 3 months				
No decrease in food intake, n/N=36/627	1 (Reference group)		1 (Reference group)	
Moderate and severe decrease in food intake, n/N=16/146	1.97 (1.09, 3.55)	0.02	1.82 (0.99, 3.34)	0.053
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Weight loss over the past 3 months				
No weight loss, n/N=26/513	1 (Reference group)		1 (Reference group)	
Weight loss between 1 and 3 kg, and greater than 3 kg n/N=25/248	2.04 (1.18, 3.54)	0.01	1.93 (1.10, 3.39)	0.02
Mobility				
Goes out, n/N=25/468	1 (Reference group)		1 (Reference group)	
Able to get out of bed/chair but does not go out, and bed or chair bound,	r (reconstruct growp)		r (restored group)	
n/N=27/305	1.68 (0.97, 2.89)	0.06	1.75 (1.00, 3.06)	0.048
Acute disease or psychological stress over the past 3 months				
No, $n/N=31/593$	1 (Reference group)		1 (Reference group)	
Yes, $n/N=21/180$	2.30 (1.32, 4.00)	0.003	2.12 (1.20, 3.74)	0.01
Neuropsychological problems				
No psychological problems, n/N=17/226	1 (Reference group)		1 (Reference group)	
Mild and severe dementia or depression, n/N=35/547	0.85 (0.47, 1.51)	0.58	0.94 (0.52, 1.70)	0.83
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Body mass index (BMI, kg/m2) or calf circumference (CC, cm)**				
BMI \geq 21 or CC \geq 31, n/N=33/603	1 (Reference group)		1 (Reference group)	
BMI $<$ 21 or CC $<$ 31, n/N=19/170	2.08 (1.18, 3.66)	0.01	2.34 (1.31, 4.17)	0.004

BMI: Body Mass Index; CC: Calf circumference; HR: Hazard Ratio; CI: confidence interval n: number of deceased subjects/N: total number of subjects; *Adjusted for age and gender; ** if BMI was not available, the CC was used at its place to define the item

Table 4. Sensitivity, Specificity, Predictive values and Likelihood ratios for mortality of the MNA-SF and its significant items

	Sensitivity (%)	Specificity (%)	Positive Predictive Value (%)	Negative Predictive Value (%)	Positive Likelihood Ratio	Negative Likelihood Ratio
MNA-SF items						
• BMI < 21 or CC <31	36.5	79.1	11.2	94.5	1.74	0.80
• Acute disease / stress	40.4	78.0	11.7	94.8	1.83	0.76
• Weight loss ≥1 kg	49.0	68.6	10.1	94.9	1.56	0.74
• Impaired mobility	51.9	61.4	8.9	94.7	1.35	0.78
MNA-SF <12	88.5	26.6	8.0	97.0	1.21	0.43
MNA-SF <8	28.9	85.2	12.3	94.3	1.95	0.84

³⁷³ BMI: Body Mass Index; CC: Calf circumference; MNA-SF: Mini Nutritional Assessment-Short Form

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