



Accuracy, feasibility and predictive ability of different frailty instruments in an acute geriatric setting

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Key Summary Points

Aim To investigate the feasibility and accuracy of four frailty instruments: FRAIL, Clinical Frailty Scale (CFS), hand grip strength (HGS) and the Spanish Frailty-VIG; and to evaluate their ability to predict adverse outcomes in an acute care setting (ACS).

Findings The four instruments had high feasibility but provided variable prevalence of frailty. FRAIL and CFS predicted well for three-month mortality, and FRAIL also for length of stay. However, none of the instruments predicted for the other outcomes.

Message The FRAIL and CFS may be of value in diagnostic and therapeutic decision-making in an acute geriatric setting, given their prognostic ability and feasibility. Further studies are needed to identify the best frailty instrument in an ACS.

Abstract

Purpose To analyze the feasibility, accuracy and the ability of different frailty instruments to predict adverse outcomes.

Methods A prospective cohort study was conducted in patients ≥ 70 years admitted to the acute care setting (ACS). Feasibility and prevalence of frailty were assessed by FRAIL, Clinical Frailty Scale (CFS), hand grip strength (HGS) and the Spanish Frailty-VIG. Receiver operator characteristic (ROC) curves and area under the curve (AUC) were performed to identify frailty according to each instrument, setting VIG as the reference. For each instrument, multiple logistic regressions were used to examine the effect of frailty on primary outcome (i.e., three-month mortality) and secondary outcomes (i.e., in-hospital mortality, length of stay, institutionalization, functional decline and 30-day readmission).

Results A total of 185 patients were included, with a median age of 89 years. The feasibility of the instruments was 100%, except for HGS (67%). The prevalence of frailty varied from 65.2% (FRAIL) to 86.7% (VIG). AUCs against VIG ranged from 0.69 (95% confidence interval [CI] 0.57–0.81; FRAIL) to 0.77 (95% CI 63.5–90.2; CFS). Frail patients defined by FRAIL were 2.7 times more likely to have a prolonged length of stay than non-frail patients (95% CI 1.385–5.416). Three-month mortality occurred more among frail patients, either defined by FRAIL (OR 2.5; 95% CI 1.072–5.881) or CFS (OR 3.7; 95% CI 1.255–10.812), than in non-frail patients.

Conclusion The four instruments had high feasibility providing variable prevalence of frailty. FRAIL and CFS predicted well for three-month mortality, and FRAIL also for length of stay. However, none of the instruments predicted for the other secondary outcomes of the study.

Keywords Frailty · Accuracy · Feasibility · Acute care setting

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Introduction

Frailty is a biological syndrome consisting of a decreased functional reserve caused by the decline of multiple physiological systems, leading to a loss of homeostatic capacity and making individuals more vulnerable to adverse events [1]. The frail older people living in the community have a higher risk of hospitalization, mortality, dependence, falls or institutionalization [2], whereas this condition leads the hospitalized older patients to worse outcomes in terms of in-hospital and long-term mortality, functional decline or institutionalization [3, 4]. There are two complementary views of this syndrome: (1) the phenotypic model of frailty, which is particularly useful for population screening to identify pre-disability states, and, (2) the deficit accumulation approach, that considers frailty as a quantifiable continuum of age-related health deficits [2]. In this regard, the Linda Fried Frailty Phenotype instrument is highly feasible to measure in nursing homes, primary care and outpatient settings frailty according to the phenotypic model, but its feasibility is significantly lower in those admitted to an AGS [5, 6]. The hand grip strength (HGS) as a measurement of muscle strength is a dimension of the Linda Fried Frailty Phenotype along with unintentional weight loss, self-reported exhaustion, sedentary behavior and slow gait speed, and HGS has shown predictive validity for decline in cognition, mobility, functional status and mortality in older people living in the community [7]. On the other hand, the Frail-VIG Index (VIG) is based on the Comprehensive Geriatric Assessment (CGA), measures 22 dichotomic variables to assess deficit accumulation in several domains, and has been recently validated in an AGS in Spain [8]. This and other instruments within deficit accumulation model [9, 10] are long, time-consuming and complex to apply, and therefore simpler and shorter screening instruments such as the Clinical Frailty Scale (CFS) [11] and the FRAIL questionnaire have been developed and are being increasingly used in AGS [12]. The CFS has strong correlation with the VIG and other instruments within the deficit accumulation model, and the FRAIL questionnaire is considered a mixed test as it is composed of four items of physical frailty and one of comorbidity [12, 13].

Frailty instruments are of outmost value in acute geriatric units, as they not only allow the clinician to establish a prognosis, but also to personalize the goals of care and tailor the diagnostic and therapeutic interventions [4]. However, there is no consensus on which frailty instrument is better to be applied at AGS [14]. Consequently, we sought to evaluate four existing instruments with different frailty approaches for the detection of this syndrome in AGS. This study aims to (1) identify the feasibility of these different frailty instruments (HGS, VIG, CFS, FRAIL); (2) compare their accuracy to identify frailty using the VIG as the reference and,

(3) evaluate their ability to predict adverse outcomes among hospitalized older adults admitted to an AGS.

Methods

Population and study design

The study population consisted of older patients who were consecutively admitted to the Department of Geriatric Medicine at an academic tertiary care hospital, from 1 June 2019 to 31 December 2020 with a follow-up period of three months. This hospital covers a population of 312,000 inhabitants in the north of Madrid where around 500 older patients are attended per year in the acute geriatric wards.

Patients aged 70 and older were eligible for enrollment in the prospective cohort study if they were admitted to the Geriatric acute care wards and provided (themselves or a legal representative) a signed written informed consent document within 24 h of admission. Patients who suffered from COVID-19 within three months of discharge were excluded from the analysis of mortality and readmission rates.

Patients were assisted according to the principles of acute geriatric units, i.e. comprehensive geriatric assessment and care focused on the needs of the patients, interdisciplinary work carried out by a core team of professionals (geriatrician, nursing staff trained in geriatrics and social worker), and early discharge planning [15, 16]. In addition, a geriatrician, a specialist geriatric nurse and a geriatric student were responsible for the administration of the written informed consent document to participants. Data collection and assessment of frailty and comprehensive geriatric assessment were obtained within 24 h of admission. Our research interviewers contacted participants (or their legally acceptable representative) at three months after enrollment to determine their mortality and 30-day readmission.

Frailty assessment

In supplementary Fig. 1 and supplementary tables 1 and 2, we summarize the main features of the different instruments used to assess frailty in this study. The Rockwood Clinical Frailty Scale (CFS) considers the pre-existing level of function and mobility and classifies patients from (1) very fit to (9) terminally ill based on easy-to-understand pictograms and descriptors. When the score is ≥ 4 , the patient is considered frail [17].

The HGS in kilograms (kg) was measured in the self-reported stronger arm using a Jamar Hand Dynamometer, with participants seated in a chair and the higher value of two trials was used for data analysis [7]. We considered as a cut-off point for frailty in men a HGS < 23 kg and < 12 kg in women, according to the normal values from the Frailty

and Dependency Study Cohort (FRADEA) conducted in our country [18].

The FRAIL is a short interview-based tool (1–3 min) designed to assess fatigue, endurance, ambulation, weight loss and illness, with score range from 0 to 5. Values ≥ 3 identify the individual as being frail [13].

The Frail-VIG is a multidimensional index based on the accumulation of deficits extracted from the CGA, measuring 22 variables grouped in 8 domains: functional, nutritional, cognitive, emotional, social, geriatric syndromes prior to admission, symptoms with criteria of severity and the presence of chronic diseases. The Index range is from 0 to 1 point, coming from dividing the total sum of the points of the variables into 25, considering individuals with VIG scores ≥ 0.2 as being frail [19]. This instrument has been validated in the Spanish population in patients admitted to acute geriatric wards, contrary to other instruments that in addition are mostly applied in other settings. Therefore, VIG instrument was considered as the reference in this study [8, 19].

Feasibility was assessed on the percentage of patients with all composites of the VIG, CFS, and FRAIL scales completed; if any scale item was missing, it was considered incomplete. In the case of the HGS, feasibility was evaluated on the percentage of patients able to understand and coordinate the action of pressing the dynamometer.

Other variables of the study

In addition to the aforementioned variables, we also recorded age (years), sex (male or female), ability to perform activities of daily living before admission and at discharge according to the usual cut-off points of the Barthel Index [20], percentage of patients with dementia (considered when the diagnosis of dementia was previously made in an outpatient clinic) [21] and their stages according to the Global Deterioration Scale (i.e., mild-moderate cognitive impairment, equivalent to Global Deterioration Scale 4–5 or severe-very severe cognitive impairment, equivalent to Global Deterioration Scale 6–7) [22], place of residence before admission (i.e. nursing home or community-dwelling) and in-hospital diagnostics, grouped as cardiovascular, digestive, respiratory infections, neurological disorders, nephro-urinary diseases and others.

Adverse outcomes among hospitalized older adults

The primary outcome measure was three-month mortality. Secondary outcome measures included (1) in-hospital mortality, (2) prolonged length of stay (defined as higher than 6 days), (3) new institutionalization at discharge, (4) functional decline at discharge (defined as

a worsening of ≥ 5 points in Barthel Index at discharge compared with pre-morbid) [20, 23], (5) 30-day readmission and, (6) a composite adverse outcome that combines prolonged length of stay or functional decline or new institutionalization.

Statistical analyses

Baseline characteristics of the sample were presented as mean values \pm standard deviation (SD), or median value and interquartile range (IQR) for continuous variables according to parametric test results, and as absolute and relative frequencies for categorical variables.

To assess the accuracy of frailty classification of each instrument, we derived receiver operator characteristic (ROC) curves for HGS, CFS, FRAIL, using VIG as the reference. A patient was considered to be frail when the VIG score was ≥ 0.2 . For each instrument, area under the curve (AUC) and its 95% confidence interval was calculated. An instrument was considered to acceptably diagnose frailty (according to VIG classification) when the AUC was greater than 0.7, and considered unacceptable when AUC was below 0.6 [24]. Additionally, ROC contrasts between each of the frailty instruments were performed to determine if there were statistical differences among AUC. In addition, the maximum likelihood cut-off point between sensitivity and specificity was calculated for each frailty instrument. To assess the effect of being frail, according to the usual definition for each instrument (i.e. CFS > 4 , VIG ≥ 0.2 , FRAIL ≥ 3 and HGS < 23 kg in men or < 12 kg in women) on the risk for each adverse outcome, multivariate logistic regressions were performed and the effect was adjusted for age, sex and principal diagnoses.

Finally, to discriminate the ability of each frailty instrument to predict adverse outcomes, we analyzed the ROC curves, calculating the AUC and its 95% confidence interval (CI). Frailty scales were considered as continuous variables and HGS was distributed by sex. For each ROC, curves a score greater than 0.7 was considered acceptable predictive ability and below 0.6 as unacceptable [24].

The existence of statistical significance was considered when the *p* value was less than 0.05. The analysis was performed with IBM SPSS 21.0

Ethical approval

The study complied with good clinical practice standards set forth in the Declaration of Helsinki of 1975 and was approved by the relevant institutional review board: Ethical and Research Committee of the Hospital.

Results

Baseline characteristics and feasibility of the scales

A total of 185 patients complied with the inclusion criteria of the study, and 171 could be analyzed for three-month mortality (Fig. 1). The median age of participants was 89 [85–93] years, a high percentage of them with dementia (63.8%), mostly in mild stage (43.8%), with a predominance of female participants (60.5%) and the majority was living in the community (60%). The most frequent principal diagnosis was respiratory infections (42.2%), followed by neurological disorders (17.8%) and nephro-urinary diseases (11.4%). The feasibility of the FRAIL, VIG and CFS scales was 100%, whereas in for the HGS it was 67% (Table 1).

Accuracy of frailty classification

The prevalence of frailty was 88.1%, 74.6%, 63.2% and 71.7%, when assessed using VIG, CFS, FRAIL and HGS, respectively (Table 1). AUCs for FRAIL, CFS and HGS against the reference VIG for diagnosis of frailty were 0.69 (95% confidence interval [CI] 0.57–0.81; $P=0.009$), 0.89 (95% CI 83.4–95.1; $P<0.001$) and 0.73 (95% CI (62.1–84); $P=0.001$), respectively. On ROC contrasts, the AUC was significantly different between CFS vs. FRAIL ($P=0.003$), whereas we did not find differences between the AUC of FRAIL vs. HGS ($P=0.517$), and the AUC of CFS vs. HGS ($P=0.054$) in the detection of frailty, as defined by VIG

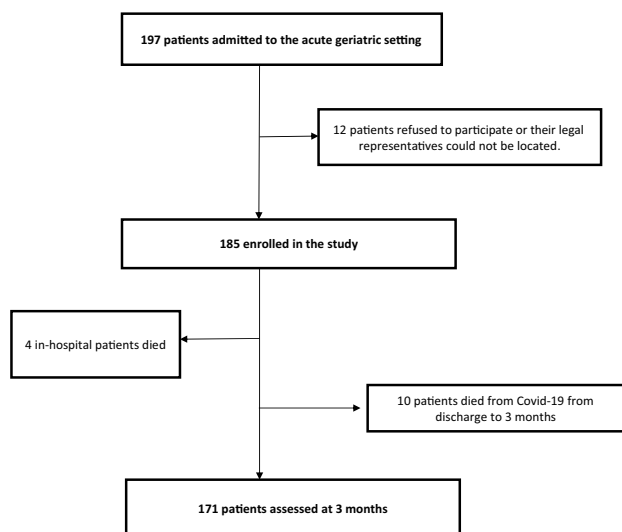


Fig. 1 Flow-chart of the study

Table 1 Baseline characteristics of study participants

| | <i>N</i> = 185 |
|--|------------------|
| Age, years | 89.0 (85.0–93.0) |
| Sex | |
| Females | 112 (60.5) |
| Males | 73 (39.5) |
| Prior Barthel index | 45.0 (16.3–73.8) |
| Barthel index at discharge | 40.0 (14.4–65.6) |
| Dementia | 118 (73.8) |
| Place of residence | |
| Nursing home | 74 (40.0) |
| Community-dwelling | 111 (60.0) |
| Frail VIG index | |
| Feasibility | 185 (100) |
| Median score | 0.4 (0.3–0.5) |
| < 0.2 (non-frail) | 22 (11.9) |
| ≥ 0.2 (frail) | 163 (88.1) |
| Clinical Frailty Scale | |
| Feasibility | 185 (100) |
| Median score | 6.0 (3.0–8.0) |
| < 4 (non-frail) | 47 (25.4) |
| ≥ 4 (frail) | 138 (74.6) |
| FRAIL scale | |
| Feasibility | 185 (100) |
| Median score | 3.0 (2.0–4.0) |
| < 3 (non-frail) | 68 (36.8) |
| ≥ 3 (frail) | 117 (63.2) |
| Hand grip strength | |
| Feasibility | 124 (67.0) |
| Median score in men | 17 (12–22) |
| Median score in women | 10 (6–14) |
| Frail according to hand grip strength ^a | 89 (71.7) |
| Principal diagnosis | |
| Cardiovascular | 18 (9.7) |
| Digestive | 11 (5.9) |
| Respiratory infections | 78 (42.2) |
| Nephro-urinary diseases | 21 (11.4) |
| Neurological disorders | 33 (17.8) |
| Others | 24 (13.0) |
| In-hospital mortality | 7 (3.8) |
| Length of stay | 6.0 (3.3–8.8) |
| Functional decline | 35 (19.7) |
| Instituzionalization at discharge | 9 (5.1) |
| Readmitted within 30 days | 61 (34.9) |
| 3-Month mortality | 47 (25.7) |

Results are expressed as *n* (%) or median (Q1–Q3)

^aMen with a hand grip strength < 23 kg and < 12 kg in women were considered frail

(Fig. 2). The CFS provided greater sensitivity (72.4%), whereas the CFS provided greater specificity (94.7%) in the diagnosis of frailty.

Predictive ability of the frailty instruments for adverse outcomes

Regarding negative outcomes, we found an in-hospital mortality of 3.8%, a median length of stay of 6 days (3.3–8.8), 19.7% developed functional decline, 9.1% need institutionalization at discharge, 34.9% were readmitted within 30 days, and 25.7% died within 3 months (Table 1).

Within the multivariate analysis adjusted for sex, age, and principal diagnosis, we found that frail patients (defined by FRAIL) were 2.7 times more likely to have a prolonged length stay than non-frail patients (58.1% vs. 39.7%; 95% CI of the OR: 1.385–5.416; $P=0.004$). Three-month mortality after discharged occurred more in frail patients than in non-frail patients, either defined by FRAIL or CFS (FRAIL: 31.9% vs. 14.9%, OR: 2.5; CI 95% 1.072–5.881; $P=0.034$; CFS: 25.7% vs. 11.1%, OR: 3.7; 95% CI 1.255–10.812; $P=0.018$). However, we did not find significant relation of the other negative outcomes, as shown in the Table 2.

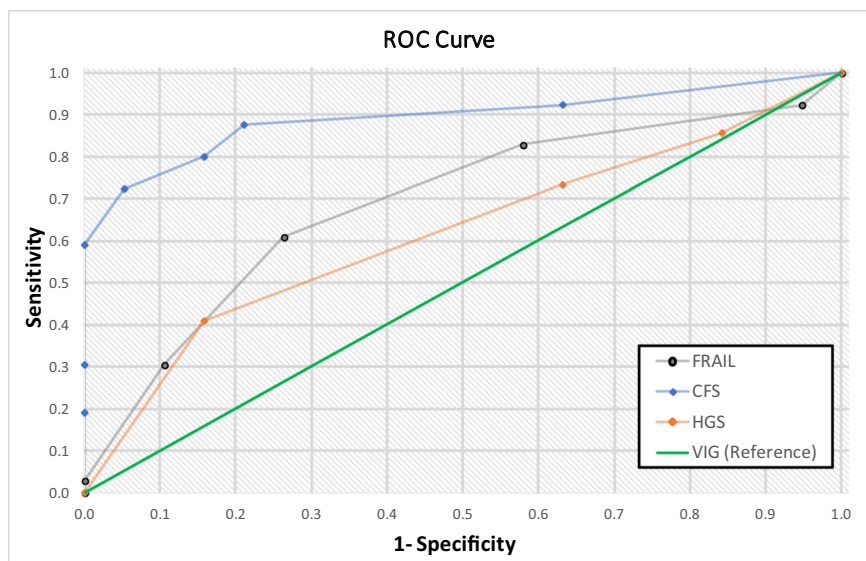
None of the frailty instruments resulted to be good predictors for any of the adverse outcomes (i.e. AUC-ROC above 0.7), as shown in Table 3 and Supplementary Fig. 2.

Discussion

We present a prospective cohort study of patients admitted to an AGS with a median age of almost 90 years aiming to evaluate the accuracy, feasibility and predictive ability of four different frailty instruments. Prevalence of frailty according to the different instruments varied from 62.2 to 86.7%. Therefore, FRAIL, HGS and CFS showed a variable prevalence and an acceptable ability to detect frailty when referenced with the VIG instrument. In addition, the feasibility of the instruments ranged from 67 to 100%. Regarding the predictive ability of the instruments, a frail patient by FRAIL and CSF was 2.5 and 3.7 times more likely to die at 3 months, respectively, than a non-frail. Moreover, patients classified as frail by FRAIL were more likely to stay in the hospital for more than 6 days. However, being classified as frail by any of the frailty instruments was not associated with in-hospital mortality, institutionalization or readmission.

Frailty instruments must be selected according to the characteristics of the setting where they are subministered, the ability to complete the test and the time required to perform it [25]. In this sense, we apply three brief instruments i.e. CFS, FRAIL and HGS (90, 24 and 90–120 s, respectively) and one longer i.e. VIG (10 min) in our study to measure frailty [6, 26]. Therefore, the first finding worth mentioning is the high feasibility of the self-reported instruments chosen, i.e., FRAIL, CFS or VIG in our AGS, in contrast to those containing objective measures such as the HGS. This different feasibility found between

Fig. 2 The AUCs for FRAIL, CFS, and HGS against the reference VIG in diagnosis of frailty. ROC receiver operator characteristic, CFS Clinical Frailty Scale, HGS hand grip strength, AUC area under the curve



| Frailty instrument | AUC (95% CI) | P-value | Cut point, (Sensitivity, Specificity) |
|--------------------|--------------------|---------|---------------------------------------|
| FRAIL | 69.0 (56.8 - 81.2) | 0.009 | 2.5, (61-73.7%) |
| CFS | 89.2 (83.4 - 95.1) | < 0.001 | 4.5, (72.4-94.7%) |
| HGS | 73.1 (62.1 - 84.0) | 0.001 | 14.5, (69.5-73.7%) |

Table 2 Predictive ability of the frailty instruments for adverse outcomes: multivariate analysis adjusted for sex, age, and principal diagnosis

| | OR (CI 95%) associated with IF-VIG scale ≥ 0.2 (frail) | OR (CI 95%) associated with FRAIL scale ≥ 3 (frail) | OR (CI 95%) associated with Clinical Frailty Scale ≥ 4 (frail) | OR (CI 95%) associated with low ^b hand grip strength (frail) |
|--------------------------------|---|--|---|---|
| In-hospital mortality | 1.142 (0.115–11.359) | 1.620 (0.265–9.896) | 3.029 (0.292–31.468) | 2.028 (0.095–43.324) |
| Length of stay ≥ 6 | 1.518 (0.588–3.920) | 2.739 (1.385–5.416)* | 1.558 (0.748–3.246) | 1.122 (0.468–2.687) |
| Functional decline | 1.753 (0.398–7.731) | 0.891 (0.347–2.289) | 1.444 (0.484–4.307) | 2.039 (0.380–10.950) |
| 30-Days readmission | 1.608 (0.562–4.601) | 1.297 (0.641–2.626) | 0.932 (0.431–2.017) | 0.852 (0.345–2.100) |
| 3-Month mortality | 1.111 (0.362–3.408) | 2.511 (1.072–5.881)* | 3.684 (1.255–10.812)* | 3.780 (0.973–14.692) |
| New institutionalization | 0.366 (0.036–3.761) | 0.619 (0.124–3.097) | 0.217 (0.038–1.245) | 3.737 (0.196–71.245) |
| Composite outcome ^a | 1.645 (0.624–4.338) | 1.590 (0.868–2.915) | 1.626 (0.765–3.458) | 1.391 (0.586–3.304) |

P* value < .05^aComposite outcome: length of stay ≥ 6 or functional decline or new institutionalization^bMen with a hand grip strength < 23 kg and < 12 kg in women were considered frailTable 3** The area under the receiver operating characteristic curves for VIG, FRAIL, clinical frailty scale and hand grip strength in predicting adverse outcomes

| | IF-VIG scale AUC (95% CI) | FRAIL scale AUC (95% CI) | Clinical Frailty Scale AUC (95% CI) | Hand grip strength ^b AUC (95% CI) |
|--------------------------------|------------------------------|-----------------------------|--|---|
| In-hospital mortality | 0.659 (0.440–0.877) | 0.619 (0.428–0.811) | 0.568 (0.357–0.778) | 0.643 (0.350–0.937) |
| Length of stay ≥ 6 | 0.553 (0.470–0.635) | 0.591 (0.509–0.673) | 0.562 (0.479–0.645) | 0.520 (0.417–0.623) |
| Functional decline | 0.548 (0.444–0.651) | 0.550 (0.440–0.660) | 0.617 (0.524–0.709) | 0.570 (0.446–0.694) |
| 30-Day readmission | 0.508 (0.419–0.596) | 0.507 (0.418–0.596) | 0.508 (0.420–0.597) | 0.517 (0.406–0.627) |
| 3-Month mortality | 0.627 (0.534–0.719) | 0.644 (0.554–0.734) | 0.666 (0.576–0.757) | 0.563 (0.445–0.681) |
| New institutionalization | 0.603 (0.421–0.784) | 0.622 (0.471–0.773) | 0.623 (0.410–0.836) | 0.646 (0.437–0.856) |
| Composite outcome ^a | 0.529 (0.445–0.614) | 0.554 (0.469–0.638) | 0.534 (0.450–0.618) | 0.523 (0.420–0.625) |

^aComposite outcome: length of stay ≥ 6 or functional decline or new institutionalization^bHand grip strength was distributed by sex

these two types of instruments is similar to the one found in the study by Oviedo-Briones et al. in patients admitted to geriatric wards [6].

On the other hand, the prevalence of frailty reported in our study is higher than the 20–50% reported in most of the studies [27–29], but it is nevertheless in line with that demonstrated by Chong et al. with a sample of patients very similar to ours, i.e. very old patients with comorbidity burden, low baseline functional status, high prevalence of cognitive impairment and a considerable proportion of patients coming from nursing homes [3, 30]. Regarding the ability of frailty instruments to predict mortality [14, 31, 32], we should emphasize that in patients admitted to AGS, decisions about diagnosis, treatment, and intervention are often made without a strong evidence base. Therefore, some patients may be subjected to overtreatment or adverse effects of interventions that cause distress at the end of their lives. In contrast, other patients who would potentially benefit from the intervention may not undergo

these interventions only because of their advanced age. Hence, the ability to predict three-month mortality through FRAIL or CFS makes them useful tools for making tailored decisions in this setting. Moreover, the association between frailty according to the FRAIL and prolonged length of stay, in line with recent studies may also indicate that this scale adequately reflects the complexity of these patients in our sample [33]. With respect to HGS, it has been associated with functional status, mobility or mortality. However, our results do not support this correlation [7, 34]. A possible explanation for this lack of relation, could be that the population included in previous studies was younger and with better functional status than ours. On the other hand, the low feasibility of the test meant that we lost statistical power and excluded patients with a higher risk of mortality or functional impairment (i.e., those with more severe disease and worse functional status). With regards the capacity of detect frailty of the three instruments when referenced with the Spanish validated

cumulative deficit approach instrument, we found that all of them were almost acceptable, but variable. One likely explanation is that this variability reflects that different scales are measuring different components of the construct of frailty.

Regarding the limitations of the study, we should highlight that we selected the VIG instrument as a reference, because it was validated to predict survival at 1 and 2 years in a sample similar to ours. However, as we have demonstrated in this study, the instrument is not adequate for predicting adverse outcomes among hospitalized older adults or three-month mortality [8, 19]. Another limitation of the present study is the small sample size, which means that our results must be interpreted with caution and need to be reproduced in other populations. In addition, the scales applied assess previous frailty and not that acquired during acute admission (except for the assessment of HGS on admission). This assessment at a different time and in different circumstances limits the comparison of their discriminative ability. Finally, we should highlight as a limitation of the study that we canceled the follow-up, due to the mortality impact that COVID-19 could cause in a sample in which 40% of the patients came from nursing homes. In addition, the low in-hospital mortality we found probably prevented us from finding a valuable instrument to predict it.

Regarding the strengths of the study, first we emphasize that it was conducted in an AGS through validated instruments, given that most of the studies which focused on frailty are conducted in non-geriatric disciplines and two thirds of them identify participants as frail even without measuring frailty [14]. Second, we have evaluated and compared within the same population three important characteristics of a frailty instrument should present i.e. accuracy, feasibility and predictive ability to be applied.

Therefore and according our results we can recommend to administer FRAIL and CFS in geriatric wards due to their high feasibility, accuracy to detect frail population and their ability predict short adverse outcomes as 3-month mortality. In contrast and according to our outcomes VIG and HGS are not highly recommendable in this setting. However, more researches are needed to be conducted in AGS to find better instruments to predict in-hospital mortality, institutionalization or readmission. In this sense finding frailty biomarkers as reference to measure frailty in the future, may allow to find an accurate clinical instrument of measurement [35].

Conclusion

The prevalence of frailty in a sample of older patients admitted to an AGS according to different validated instruments is very high and varies from 63.2% to 74.6% when assessed

using FRAIL and CFS, respectively. The CFS predicted mortality at three months and the FRAIL scale as well as three-month mortality was associated with prolonged length of stay (≥ 6 days). The full feasibility of both scales together with their ability to predict short-term adverse outcomes make them recommendable in this setting. VIG and HGS were not associated with any outcome. Finally, none of the instruments were related to in-hospital mortality, 30-day readmission and functional decline, warranting further studies with any other validated instrument in this setting to predict these outcomes.

Supplementary Information The online version contains supplementary material available at <https://doi.org/10.1007/s41999-022-00645-1>.

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Author contributions Study concept and design: RB and FZ. Acquisition of data: CB, AL and RB, NA, CS and MN. Analysis and interpretation of data: IJT, CA-V. Drafting of the manuscript: EA, AL, PPCO. Critical revision of the manuscript for important intellectual content: RB, FZ. All authors saw and approved of the final version of the manuscript.

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Declarations

Conflict of interest The authors declare that they have no conflicts of interest.

Ethical approval All experimental procedures were performed in accordance with the principles of the Declaration of Helsinki. This study was approved by the institutional review board of the hospital.

Informed consent Written informed consent was given by each patient. All participants were provided the opportunity to have their data excluded from the study analysis.

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References

- Campbell AJ, Buchner DM (1997) Unstable disability and the fluctuations of frailty. *Age Ageing* 26:315–318
- Vermeiren S, Vella-Azzopardi R, Beckwée D, Habbig A-K, Scafoglieri A, Jansen B et al (2016) Frailty and the prediction of negative health outcomes: a meta-analysis. *J Am Med Dir Assoc* 17:1163.e1–1163.e17. <https://doi.org/10.1016/j.jamda.2016.09.010>
- Chong E, Ho E, Baldevarona-Llego J, Chan M, Wu L, Tay L (2017) Frailty and risk of adverse outcomes in hospitalized older adults: a comparison of different frailty measures. *J Am Med Dir Assoc* 18:638.e7–638.e11. <https://doi.org/10.1016/j.jamda.2017.04.011>
- Hubbard RE, Peel NM, Samanta M, Gray LC, Mitnitski A, Rockwood K (2017) Frailty status at admission to hospital predicts multiple adverse outcomes. *Age Ageing* 46:801–806. <https://doi.org/10.1093/ageing/afx081>
- Fried LP, Tangen CM, Walston J, Newman AB, Hirsch C, Gottdiener J et al (2001) Frailty in older adults: evidence for a phenotype. *J Gerontol A Biol Sci Med Sci* 56:M146–M156
- Oviedo-Briones M, Laso AR, Carnicero JA, Cesari M, Grodzicki T, Gryglewska B et al (2020) A Comparison of frailty assessment instruments in different clinical and social care settings: the Frailtools Project. *J Am Med Dir Assoc*. <https://doi.org/10.1016/j.jamda.2020.09.024>
- Rijk JM, Roos PRKM, Deckx L, Van den Akker M, Buntinx F (2016) Prognostic value of handgrip strength in people aged 60 years and older: a systematic review and meta-analysis. *Geriatr Gerontol Int* 16:5–20. <https://doi.org/10.1111/ggi.12508>
- Amblàs-Novellas J, Martori JC, Molist Brunet N, Oller R, Gómez-Batiste X, Espauella PJ (2017) Frail-VIG index: design and evaluation of a new frailty index based on the comprehensive geriatric assessment. *Rev Esp Geriatr Gerontol* 52:119–127. <https://doi.org/10.1016/j.regg.2016.09.003>
- Jones DM, Song X, Rockwood K (2004) Operationalizing a frailty index from a standardized comprehensive geriatric assessment. *J Am Geriatr Soc* 52:1929–1933. <https://doi.org/10.1111/j.1532-5415.2004.52521.x>
- Soong J, Poots AJ, Scott S, Donald K, Bell D (2015) Developing and validating a risk prediction model for acute care based on frailty syndromes. *BMJ Open* 5:1–12. <https://doi.org/10.1136/bmjopen-2015-008457>
- Rockwood K, Song X, MacKnight C, Bergman H, Hogan DB, McDowell I et al (2005) A global clinical measure of fitness and frailty in elderly people. *CMAJ* 173:489–495. <https://doi.org/10.1503/cmaj.050051>
- Moreno-Ariño M, Torrente Jiménez I, Cartanya Gutiérrez A, Oliva Morera JC, Comet R (2020) Assessing the strengths and weaknesses of the Clinical Frailty Scale through correlation with a frailty index. *Aging Clin Exp Res* 32:2225–2232. <https://doi.org/10.1007/s40520-019-01450-w>
- Kojima G (2018) Frailty defined by FRAIL scale as a predictor of mortality: a systematic review and meta-analysis. *J Am Med Dir Assoc* 19:480–483. <https://doi.org/10.1016/j.jamda.2018.04.006>
- Theou O, Squires E, Mallery K, Lee JS, Fay S, Goldstein J et al (2018) What do we know about frailty in the acute care setting? A scoping review. *BMC Geriatr* 18:1–20. <https://doi.org/10.1186/s12877-018-0823-2>
- Baztan JJ, Suarez-Garcia FM, Lopez-Arrieta J, Rodriguez-Manas L, Rodriguez-Artalejo F (2009) Effectiveness of acute geriatric units on functional decline, living at home, and case fatality among older patients admitted to hospital for acute medical disorders: meta-analysis. *BMJ* 338:b50–b50. <https://doi.org/10.1136/bmj.b50>
- Ellis G, Whitehead MA, Robinson D, O'Neill D, Langhorne P (2011) Comprehensive geriatric assessment for older adults admitted to hospital: meta-analysis of randomised controlled trials. *BMJ* 343:d6553
- Ellis HL, Wan B, Yeung M, Rather A, Mannan I, Bond C et al (2020) Complementing chronic frailty assessment at hospital admission with an electronic frailty index (FI-Laboratory) comprising routine blood test results. *CMAJ* 192:E3–8. <https://doi.org/10.1503/cmaj.190952>
- Abizanda Soler P, López-Torres Hidalgo J, Romero Rizo L, Sánchez Jurado PM, García Noguera I, Esquinas Requena JL (2012) Normal data of functional assessment tools of the elderly in Spain: the FRADEA Study. *Aten Primaria* 44:162–171. <https://doi.org/10.1016/j.aprim.2011.02.007>
- Amblàs-Novellas J, Martori JC, Espauella J, Oller R, Molist-Brunet N, Inzitari M et al (2018) Frail-VIG index: a concise frailty evaluation tool for rapid geriatric assessment. *BMC Geriatr* 18:1–12. <https://doi.org/10.1186/s12877-018-0718-2>
- Shah S, Vanclay F, Cooper B (1989) Improving the sensitivity of the Barthel Index for stroke rehabilitation. *J Clin Epidemiol* 42:703–709
- Seitz DP, Gill SS, Gruneir A, Austin PC, Anderson GM, Bell CM et al (2014) Effects of dementia on postoperative outcomes of older adults with hip fractures: a population-based study. *J Am Med Dir Assoc* 15:334–341. <https://doi.org/10.1016/j.jamda.2013.12.011>
- Reisberg B, Ferris SH, De Leon MJ, Crook T (1982) The global deterioration scale for assessment of primary degenerative dementia. *Am J Psychiatry* 139:1136–1139. <https://doi.org/10.1176/ajp.139.9.1136>
- Baztan JJ, Suárez-García FM, López-Arrieta J, Rodríguez-Mañas L, Rodríguez-Artalejo F (2009) Effectiveness of acute geriatric units on functional decline, living at home, and case fatality among older patients admitted to hospital for acute medical disorders: meta-analysis. *BMJ* 338:334–336. <https://doi.org/10.1136/bmj.b50>
- Wians FH (2009) Clinical laboratory tests: Which, why, and what do the results mean? *Lab Med* 40:105–113. <https://doi.org/10.1309/LM4O4L0HHUTWWUDD>
- Walston J, Buta B, Xue QL (2018) Frailty screening and interventions: considerations for clinical practice. *Clin Geriatr Med* 34:25–38. <https://doi.org/10.1016/j.cger.2017.09.004>
- Roberts HC, Denison HJ, Martin HJ, Patel HP, Syddall H, Cooper C et al (2011) A review of the measurement of grip strength in clinical and epidemiological studies: towards a standardised approach. *Age Ageing*. <https://doi.org/10.1093/ageing/afr051>
- Pilotto A, Rengo F, Marchionni N, Sancarlo D, Fontana A, Panza F et al (2012) Comparing the prognostic accuracy for all-cause mortality of frailty instruments: a multicentre 1-year follow-up in hospitalized older patients. *PLoS ONE*. <https://doi.org/10.1371/journal.pone.0029090>
- Joosten E, Demuyneck M, Detroyer E, Milisen K (2014) Prevalence of frailty and its ability to predict in hospital delirium, falls, and 6-month mortality in hospitalized older patients. *BMC Geriatr*. <https://doi.org/10.1186/1471-2318-14-1>
- Hogan DB, Maxwell CJ, Afilalo J, Arora RC, Bagshaw SM, Basran J et al (2017) A scoping review of frailty and acute care in middle-aged and older individuals with recommendations for future research. *Can Geriatr J* 20:22–37. <https://doi.org/10.5770/cgj.20.240>
- Chong E, Ho E, Baldevarona-Llego J, Chan M, Wu L, Tay L et al (2018) Frailty in hospitalized older adults: comparing different frailty measures in predicting short- and long-term patient

- outcomes. *J Am Med Dir Assoc* 19:450–457.e3. <https://doi.org/10.1016/j.jamda.2017.10.006>
31. Buta BJ, Walston JD, Godino JG, Park M, Kalyani RR, Xue Q-L et al (2016) Frailty assessment instruments: systematic characterization of the uses and contexts of highly-cited instruments. *Ageing Res Rev* 26:53–61. <https://doi.org/10.1016/j.arr.2015.12.003>
 32. Church S, Rogers E, Rockwood K, Theou O (2020) A scoping review of the Clinical Frailty Scale. *BMC Geriatr*. <https://doi.org/10.1186/s12877-020-01801-7>
 33. Gleason LJ, Benton EA, Alvarez-Nebreda ML, Weaver MJ, Harris MB, Javedan H (2017) FRAIL questionnaire screening tool and short-term outcomes in geriatric fracture patients. *J Am Med Dir Assoc* 18:1082–1086. <https://doi.org/10.1016/j.jamda.2017.07.005>
 34. Dodds RM, Syddall HE, Cooper R, Kuh D, Cooper C, Sayer AA (2016) Global variation in grip strength: a systematic review and meta-analysis of normative data. *Age Ageing* 45:209–216. <https://doi.org/10.1093/ageing/afv192>
 35. Picca A, Coelho-Junior HJ, Calvani R, Marzetti E, Vetrano DL (2022) Biomarkers shared by frailty and sarcopenia in older adults: a systematic review and meta-analysis. *Ageing Res Rev*. <https://doi.org/10.1016/J.ARR.2021.101530>

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