Research paper

The SHARE Frailty Instrument for primary care predicts incident disability in a European population-based sample

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ABSTRACT

Background The adoption of a frailty paradigm in primary care would be helpful to identify adults who need priority access to specialised resources. The frailty phenotype by Fried *et al*¹ is a popular operationalisation of frailty, but it is not easily applicable in routine primary care practice. We recently created and validated a frailty instrument based on the *Survey of Health, Ageing and Retirement in Europe* (SHARE-FI),² in order to provide primary care practitioners with an easy, reliable and freely accessible tool for the assessment and monitoring of frailty in community dwelling adults over the age of 50 years (www.biomedcentral.com/1471–2318/10/57).

Aim To provide further prospective validation of SHARE-FI, with a focus on disability.

Methods *Design*: longitudinal study (wave 1: 2004–2006; mean follow-up: 2.4 years). *Setting*: European population-based survey (12 countries). *Subjects*: 17 567 community dwelling participants (mean age 63.3 years), of whom 13 378 (76.2%) were non-frail, 3438 (19.6%) pre-frail and 751 (4.3%) frail. *Main*

outcome measures: number of difficulties with basic (ADL) and instrumental (IADL) activities of daily living. *Statistical analyses*: repeated measures ANOVA with adjustment for baseline age.

Results By wave 2, 3.6% of the non-frail, 12.2% of the pre-frail and 30.4% of the frail had increased their number of ADL difficulties by at least one. Likewise, 6.6% of the non-frail, 20.4% of the pre-frail and 36.6% of the frail had, by wave 2, increased their number of IADL difficulties by at least one. Table 1 shows the repeated measures ANOVA suggested.

Conclusion SHARE-FI may contribute to quality in primary care by offering a quick and reliable way to assess and monitor frailty in community dwelling individuals over the age of 50 and prioritise their access to resources, and it serves as a novel tool for audit and research.

Keywords: diagnostic tests, frail elderly, persons with disabilities, primary health care, validation studies

How this fits in with quality in primary care

What do we know?

Frailty is a predictor of adverse outcomes, including premature death and disability. The phenotype by Fried *et al*¹ is a popular operationalisation of frailty, but in its original form it cannot be applied to routine primary care practice. The Frailty Instrument for primary care from the *Survey of Health, Ageing and Retirement in Europe* (SHARE-FI)² was created to facilitate the adoption of the frailty paradigm in primary care.

What does this paper add?

SHARE-FI is a significant predictor of incident disability. SHARE-FI is a novel tool for the assessment and monitoring of frailty, and for audit and research.

Introduction

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Frailty in older adults is a key clinical concept characterised by dysregulation of multiple biological systems, accumulation of deficits, vulnerability to stressors and increased risk of adverse outcomes.^{3,4} Even though there is still no international consensus on the definition of frailty,⁵ one of the most widely accepted operationalisations is that of Fried *et al*, who defined it as a clinical syndrome in which three or more of the following criteria are present: self-reported exhaustion, unintentional weight loss, weakness (by grip strength), slowness (by walking speed) and low physical activity.¹

Frailty, comorbidity (i.e. multiple chronic conditions) and disability (i.e. difficulties with activities of daily living) are commonly used interchangeably to identify vulnerable older adults, but there is a growing consensus that these are distinct clinical entities that are causally related.^{6,7} Consistently in longitudinal studies, the frailty phenotype by Fried *et al* has been found to be a significant predictor of new-onset disability in older adults.^{8–10} The frailty phenotype also predicts falls,^{8,11,12} hospitalisation,¹ institutionalisation¹³ and death.^{8,12,14,15}

Because of its predictive abilities, frailty is an emerging concept in primary care that may provide commissioners of health care with a clinical focus for targeting resources at an ageing population.¹⁶ However, operationalising the frailty phenotype on an individual patient requires complex calculations on a reference sample, which is not practical in the context of primary care. Indeed, family physicians and community practitioners are in need of easy instruments for determining frailty.¹⁷

In order to provide European community practitioners with an easy frailty metric, we recently developed a frailty instrument for primary care (SHARE-FI) based on the *Survey of Health, Ageing and Retirement in Europe.*² SHARE-FI is freely accessible on the website of *BMC Geriatrics* (www.biomedcentral.com/ <u>1471–2318/10/57</u>) and its use is intended (via downloadable calculators) for community dwelling adults over the age of 50 years. We previously found that SHARE-FI has sufficient concurrent validity and is a powerful predictor of mortality.² In this report we further validate SHARE-FI as a predictor of incident disability.

Methods

The full methodology for the development of SHARE-FI is detailed in our main publication,² which departs from the premise that frailty is a complex, multidimensional concept that cannot be defined with a single measurement or variable. Frailty was therefore constructed as an unobserved (latent) variable that is indicated by five different (but related) observed variables; the latter were selected by Santos-Eggimann *et al*¹⁸ from the SHARE questionnaire as being closest to those identified by Fried *et al*:¹

- Exhaustion was identified as a positive response to the question: 'In the last month, have you had too little energy to do the things you wanted to do?'
- The weight loss criterion was fulfilled by reporting a 'Diminution in desire for food' in response to the question: 'What has your appetite been like?' or, in the case of a non-specific or uncodeable response to this question, by responding 'Less' to the question: 'So, have you been eating more or less than usual?'
- Weakness was assessed by handgrip strength (Kg) using a Smedley dynamometer (S Dynamometer, TTM, Tokyo, 100 Kg), according to the following measurement protocol: participants were instructed to stand (preferably) or sit, with the elbow at 90°, the wrist in neutral position, keeping the upper arm tight against the trunk, and the inner lever of the dynamometer adjusted to suit the hand; and they were then instructed to squeeze as hard as possible for a few seconds.¹⁹ Two consecutive measurements were taken from the left and right hands. The highest of the four was selected. This variable was kept continuous.
- Slowness was defined as a positive answer to either of the following two items: 'Because of a health problem, do you have difficulty (expected to last more than three months) walking 100 metres?' or '... climbing one flight of stairs without resting?'
- The low activity criterion was assessed by the question: 'How often do you engage in activities that require a low or moderate level of energy such as gardening, cleaning the car, or doing a walk?' This variable was kept ordinal: 1 = 'More than once a week'; 2 = 'Once a week'; 3 = 'One to three times a month' and 4 = 'Hardly ever or never'.

SHARE-FI was created via estimation of a discrete factor (DFactor) model based on the above five frailty variables, using the LatentGOLD® statistical package (version 4.5.0). A single DFactor with three ordered levels or latent classes (non-frail, pre-frail and frail) was modelled for each gender.² Based on these analyses, two web-based calculators (one for each gender) were created for easy retrieval of a subject's frailty class (relative to the SHARE sample) given any five indicator measurements. The SHARE-FI calculators use a linear combination of the standardised variables that have been entered, and incorporate the loadings of each of the indicator variables on the original DFactor.

The final formula for the predicted DFactor score (DFS) in females was:

DFS (females) = (2.077707 * Fatigue - 0.757295) * 0.4088+ (3.341539 * Loss of appetite - 0.332289) * 0.3325 +(0.132827 * Grip strength - 3.534515) * -0.4910 +(2.627085 * Functional difficulties - 0.461808) * 0.6012+ (0.918866 * Physical activity - 1.523633) * 0.4818

The predicted DFS formula for males was:

DFS (males) = (2.280336 * Fatigue – 0.592393) * 0.3762 + (4.058274 * Loss of appetite – 0.263501) * 0.3130 + (0.092326 * Grip strength – 3.986646) * –0.4653 + (3.098226 * Functional difficulties – 0.365971) * 0.6146 + (1.005942 * Physical activity – 1.571803) * 0.4680

The SHARE-FI calculators are freely available on <u>www.</u> <u>biomedcentral.com/1471–2318/10/57/additional/</u> and translated versions in various European languages can be found on <u>https://sites.google.com/a/tcd.ie/share-</u> frailty-instrument-calculators/home

As regards disability variables, SHARE participants were asked about the experience of difficulty in performing a series of tasks.^{20,21} The number of experienced difficulties with basic activities of daily living (ADL) was self-reported by the participants from the following list (minimum: 0; maximum: 6): 'Because of a health problem, do you have difficulty doing any of the following activities (exclude any difficulties you expect to last less than three months)?':

- dressing, including putting on shoes and socks
- walking across a room
- bathing or showering
- eating, such as cutting up the food
- getting in and out of bed
- using the toilet, including getting up or down.

The number of difficulties with instrumental activities of daily living (IADL) was self-reported by the participants from the following list (minimum: 0; maximum: 7): 'Because of a health problem, do you have difficulty doing any of the following activities (exclude any difficulties you expect to last less than three months)?':

- using a map to figure out how to get around in a strange place
- preparing a hot meal
- shopping for groceries
- making telephone calls
- taking medications
- doing work around the house or garden
- managing money, such as paying bills and keeping track of expenses.

The SHARE baseline sample (interviewed between 2004 and 2006) was composed of 28 361 community dwelling participants. After a mean follow-up period of 2.4 years, disability information was available for 17 567

respondents (i.e. 61.9% of the baseline sample, mean age 63.3 years), of whom 13 378 (76.2%) had been non-frail, 3438 (19.6%) pre-frail and 751 (4.3%) frail at baseline.

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Statistical analyses were conducted with SPSS 16.0. Descriptives were given as means and standard deviations (SD), or percentages (%), as appropriate. The Chi-square test was used to assess how participants with follow-up disability data differed from those without follow-up disability data, in terms of the baseline frailty status. The two-tailed Spearman's correlation coefficient (*rs*) was used to assess the correlation between baseline frailty status and the number of ADL and IADL difficulties, both at baseline and follow-up.

Analyses of frequencies were conducted to determine the proportions of non-frail, pre-frail and frail participants who had, by wave 2, increased at least one, two and three points in ADL and IADL disability. Separately for each frailty category, a repeated measures ANOVA was conducted to assess whether there were statistically significant differences in ADL or IADL scores between the two study waves. The analyses were repeated with the baseline age as covariate. In these analyses, ADL and IADL scores were used as continuous dependent variables.

The level of significance was established at P < 0.01 throughout.

Results

Both at baseline and at follow-up, baseline SHARE-FI had significant direct cross-sectional correlations with the number of ADL and IADL difficulties (all *r*s >0.3, P <0.001).

The analysis of frequencies revealed that, by wave 2, 3.6% of the non-frail, 12.2% of the pre-frail and 30.4% of the frail had increased the number of ADL difficulties by at least one.

By wave 2, 1.0% of the non-frail, 5.3% of the prefrail and 18.2% of the frail had increased the number of ADL difficulties by at least two.

By wave 2, 0.5% of the non-frail, 2.5% of the prefrail and 8.8% of the frail had increased the number of ADL difficulties by at least three.

Likewise, 6.6% of the non-frail, 20.4% of the prefrail and 36.6% of the frail had, by wave 2, increased the number of IADL difficulties by at least one.

By wave 2, 1.7% of the non-frail, 9.0% of the prefrail and 18.0% of the frail had increased the number of IADL difficulties by at least two.

By wave 2, 0.8% of the non-frail, 4.1% of the prefrail and 10.5% of the frail had increased the number of IADL difficulties by at least three. Appendix Table 1 shows the repeated measures ANOVA results, which suggest that the increment in disability at follow-up was directly correlated with the baseline frailty level. As regards ADL difficulties, the frail subgroup increased, on average, by 0.3 points, whilst the non-frail and pre-frail increased by 0.1 points. As regards IADL difficulties, the frail subgroup increased, on average, by 0.3 points, the pre-frail by 0.2 points, and the non-frail by 0.1 points. These differences were statistically significant, and most remained so after age adjustment (see Appendix Table 1).

Appendix Table 2 shows the frailty differences between participants with and without follow-up disability data. Participants with follow-up disability data were significantly less likely to have been non-frail at baseline, and more likely to have been pre-frail and frail (P < 0.001).

Discussion

The aim of this study was to provide further longitudinal validation (with a focus on disability) of a newly created Frailty Instrument intended for use in primary care, where reliable and easy-to-access frailty metrics are needed in order to enhance the process of identification of vulnerable adults who need priority access to resources.

Even after a relatively short follow-up (a mean of 2.4 years), the SHARE-FI classification was a significant predictor of incident disability, even after adjusting for age. This is consistent with the properties of the original Fried *et al*'s frailty phenotype^{8–10} and fulfils the validation aim of the present study.

From the point of view of primary care practice, SHARE-FI has the advantage of being a valid and simple tool for measuring the level of frailty in individuals aged \geq 50 years given five simple measurements, which leads to the identification of a high frailty group relative to a large population-based sample. The main potential use of SHARE-FI is the screening and monitoring of frailty in community dwelling adults to help prioritise secondary care referrals and/or early multidisciplinary case management. Other advantages of SHARE-FI are that it can be easily administered in the community by non-physicians (e.g. nurses, health visitors or other allied health professionals) and that it is a relatively brief instrument.

A limitation of our study is that follow-up disability was available for only 61.9% of the baseline sample. In our previous study we demonstrated that missing wave 2 participants were more frail at baseline than non-missing participants² and this was also demonstrated here (Appendix Table 2). This is consistent with a known pattern in longitudinal studies by which frail people have higher dropout rates and are less likely to be recontactable.²² Therefore, and as we previously argued based on a sensitivity analysis,² results from this longitudinal sample are likely to represent an *underestimation* of the ability of SHARE-FI to predict incident disability in real life.

In terms of the clinical applicability of SHARE-FI, a potential limitation is that grip strength is not typically assessed in primary care, it can take some minutes to perform and how the test is administered (e.g. sitting versus standing, forearm position) may influence patient performance.²³ Because of this, and for purposes of consistency, we recommend that practitioners follow the grip strength measurement protocol adopted by SHARE.¹⁹

In order to assess the extent to which a reduced four-item version of SHARE-FI (i.e. without the inclusion of grip strength) would perform as a predictor of increased difficulties in ADL and IADL, we reclassified the longitudinal sample according to a new DFactor based on the four subjective frailty items only (i.e. exhaustion, weight loss, slowness and low physical activity). Appendix Table 3 shows these additional results, which suggest a significant loss of predictive validity of the instrument as compared to the results shown in Appendix Table 1. This is consistent with previous research highlighting grip strength as an important objective marker of frailty²⁴ and a determinant of disability in the older general population.²⁵ SHARE-FI calculators offer a meaningful framework where grip strength results can be integrated with other frailty markers without the need for additional norms or arbitrary cut-offs, which could otherwise limit its applicability in the primary care setting.

The development of SHARE-FI did not include any cognitive exclusion criteria, so in principle SHARE-FI can be applied to persons with cognitive impairment. However, because four out of the five SHARE-FI variables are based on self-report, in patients with severe dementia answers should be contrasted with those of a carer or informant.²⁶

In conclusion, SHARE-FI was found to be a significant predictor of incident disability in a European population-based sample. SHARE-FI may contribute to quality in primary care by offering a simple and reliable way to assess and monitor frailty in community dwelling adults over the age of 50 years, to prioritise their access to resources and serve as a tool for audit and research.

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REFERENCES

- Fried LP, Tangen CM, Walston J *et al.* Frailty in older adults: evidence for a phenotype. *Journals of Gerontology*. *Series A, biological sciences and medical sciences* 2001; 56:M146–56.
- 2 Romero-Ortuno R, Walsh CD, Lawlor BA and Kenny RA. A Frailty Instrument for primary care: findings from the Survey of Health, Ageing and Retirement in Europe (SHARE). *BMC Geriatrics* 2010;10:57.
- 3 Gobbens RJ, Luijkx KG, Wijnen-Sponselee MT and Schols JM. In search of an integral conceptual definition of frailty: opinions of experts. *Journal of the American Medical Directors Association* 2010;11:338–43.
- 4 Xue QL. The frailty syndrome: definition and natural history. *Clinics in Geriatric Medicine* 2011;27:1–15.
- 5 Conroy S. Defining frailty the Holy Grail of geriatric medicine. *Journal of Nutrition, Health and Aging* 2009; 13:389.
- 6 Polidoro A, Dornbusch T, Vestri A, Di Bona S and Alessandri C. Frailty and disability in the elderly: a diagnostic dilemma. *Archives of Gerontology and Geriatrics* 2010;52:e75–8.
- 7 Fried LP, Ferrucci L, Darer J, Williamson JD and Anderson G. Untangling the concepts of disability, frailty and comorbidity: implications for improved targeting and care. *Journals of Gerontology. Series A*, *biological sciences and medical sciences* 2004;59:255–63.
- 8 Ensrud KE, Ewing SK, Cawthon PM *et al.* A comparison of frailty indexes for the prediction of falls, disability, fractures, and mortality in older men. *Journal of the American Geriatrics Society* 2009;57:492–8.
- 9 Al Snih S, Graham JE, Ray LA, Samper-Ternent R, Markides KS and Ottenbacher KJ. Frailty and incidence of activities of daily living disability among older Mexican Americans. *Journal of Rehabilitation Medicine* 2009;41:892–7.
- 10 Boyd CM, Xue QL, Simpson CF, Guralnik JM and Fried LP. Frailty, hospitalization, and progression of disability in a cohort of disabled older women. *American Journal of Medicine* 2005;118:1225–31.
- 11 Nowak A and Hubbard RE. Falls and frailty: lessons from complex systems. *Journal of the Royal Society of Medicine* 2009;102:98–102.
- 12 Ensrud KE, Ewing SK, Taylor BC *et al.* Frailty and risk of falls, fracture, and mortality in older women: the study of osteoporotic fractures. *Journals of Gerontology. Series A, biological sciences and medical sciences* 2007;62:744–51.
- 13 Bandeen-Roche K, Xue QL, Ferrucci L *et al.* Phenotype of frailty: characterization in the women's health and aging studies. *Journals of Gerontology. Series A, biological sciences and medical sciences* 2006;61:262–6.
- 14 Graham JE, Snih SA, Berges IM, Ray LA, Markides KS and Ottenbacher KJ. Frailty and 10-year mortality in community-living Mexican American older adults. *Gerontology* 2009;55:644–51.

15 Cawthon PM, Marshall LM, Michael Y *et al.* Frailty in older men: prevalence, progression, and relationship with mortality. *Journal of the American Geriatrics Society* 2007;55:1216–23.

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- 16 De Lepeleire J, Iliffe S, Mann E and Degryse JM. Frailty: an emerging concept for general practice. *British Journal* of General Practice 2009;59:e177–82.
- 17 De Lepeleire J, Degryse J, Illiffe S, Mann E and Buntinx F. Family physicians need easy instruments for frailty. <u>Age</u> and Ageing 2008;37:484.
- 18 Santos-Eggimann B, Cuenoud P, Spagnoli J and Junod J. Prevalence of frailty in middle-aged and older communitydwelling Europeans living in ten countries. *Journals of Gerontology. Series A, biological sciences and medical sciences* 2009;64:675–81.
- 19 Mohd Hairi F, Mackenbach JP, Andersen-Ranberg K and Avendano M. Does socio-economic status predict grip strength in older Europeans? Results from the SHARE study in non-institutionalised men and women aged 50+. *Journal of Epidemiology and Community Health* 2010;64:829–37.
- 20 SHARE. Share 2004 Questionnaire Version 10 (manually edited April 2005). Available online: <u>www.share-project.</u> org/t3/share/new_sites/Fragebogen/ma-Gene.pdf
- 21 SHARE. *Release Guide 2.3.1 Waves 1 and 2*. Mannheim: Mannheim Research Institute for the Economics of Ageing, 2010.
- 22 Chatfield MD, Brayne CE and Matthews FE. A systematic literature review of attrition between waves in longitudinal studies in the elderly shows a consistent pattern of dropout between differing studies. *Journal of Clinical Epidemiology* 2005;58:13–19.
- 23 Roberts HC, Denison HJ, Martin HJ *et al.* A review of the measurement of grip strength in clinical and epidemiological studies: towards a standardised approach. *Age and Ageing* 2011;40:423–9.
- 24 Syddall H, Cooper C, Martin F, Briggs R and Aihie Sayer A. Is grip strength a useful single marker of frailty? <u>Age</u> and Ageing 2003;32:650–6.
- 25 den Ouden ME, Schuurmans MJ, Arts IE and van der Schouw YT. Physical performance characteristics related to disability in older persons: a systematic review. *Maturitas* 2011;69:208–19.
- 26 Farias ST, Mungas D and Jagust W. Degree of discrepancy between self and other-reported everyday functioning by cognitive status: dementia, mild cognitive impairment, and healthy elders. *International Journal of Geriatric Psychiatry* 2005;20:827–34.

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None.

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Appendix

Table 1 Rep	eated measur	es ANOV	A results (orig	inal SHAI	RE-FI)					
Variable	Category	N ₁ (total)	Value _{1 (total)}	N_2	Value ₂	N (long- itudinal)	Value ₁ (long- itudinal)	Value ₂ (long- itudinal)	Unadjusted <i>P</i> (longitudinal sample)	Age-adjusted P (long- itudinal sample)
ADL difficulties	Non-frail Pre-frail Frail	20 937 5896 1528	$\begin{array}{c} 0.0 \ (0.2) \\ 0.3 \ (0.7) \\ 1.1 \ (1.5) \end{array}$	13 379 3438 751	$\begin{array}{c} 0.1 \ (0.4) \\ 0.4 \ (1.0) \\ 1.3 \ (1.7) \end{array}$	13 378 3438 751	$\begin{array}{c} 0.0 & (0.3) \\ 0.3 & (0.7) \\ 1.0 & (1.4) \end{array}$	$\begin{array}{c} 0.1 \ (0.4) \\ 0.4 \ (1.0) \\ 1.3 \ (1.7) \end{array}$	<0.001 [*] <0.001 [*] <0.001 [*]	<0.001 [*] 0.282 [*] <0.001 [*]
IADL difficulties	Non-frail Pre-frail Frail	20 937 5896 1528	$\begin{array}{c} 0.1 \ (0.3) \\ 0.5 \ (1.0) \\ 1.8 \ (1.8) \end{array}$	13 379 3438 751	$\begin{array}{c} 0.1 \ (0.5) \\ 0.6 \ (1.3) \\ 2.0 \ (2.1) \end{array}$	13 378 3438 751	$\begin{array}{c} 0.1 \ (0.3) \\ 0.4 \ (0.9) \\ 1.7 \ (1.8) \end{array}$	$\begin{array}{c} 0.1 \ (0.5) \\ 0.6 \ (1.3) \\ 2.0 \ (2.1) \end{array}$	<0.001 [#] <0.001 [#] <0.001 [#]	$< 0.001^{*}$ $< 0.001^{*}$ 0.001^{*}
# Repeated meas ADL: basic activi N: number of pa.	ures ANOVA – test d ties of daily living; L rticipants	of within-subj ADL: instrum	jects contrasts (wave nental activities of da	e * frailty cate ily living	egory)					

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	Persons with follow-up data	Persons without follow- up data	Significance of the difference (<i>P</i>)
Non-frail (%)	13 378 (76.2)	7559 (70.0)	<0.001*
Pre-frail (%)	3438 (19.6)	2458 (22.8)	<0.001*
Frail (%)	751 (4.3)	777 (7.2)	<0.001*

Table 2 Comparisons between participants with and without follow-up disability data, in terms of their frailty status

Table 3 Rep	eated measure	s ANOVA	A results (reduc	ed four-	item version o	of SHARE-FI wi	thout inclusion	of grip streng	jth)	
Variable	Category	N ₁ (total)	Value ₁ (total)	N_2	Value ₂	N (long- itudinal)	Value ₁ (long- itudinal)	Value ₂ (long- itudinal)	Unadjusted <i>P</i> (longitudinal sample)	Age-adjusted <i>P</i> (longitudinal sample)
ADL difficulties	Non-frail Pre-frail Frail	17 819 8567 1975	$\begin{array}{c} 0.1 & (0.5) \\ 0.2 & (0.7) \\ 0.2 & (0.8) \end{array}$	$11 \ 429 \\ 5069 \\ 1070$	$\begin{array}{c} 0.1 \ (0.6) \\ 0.2 \ (0.8) \\ 0.3 \ (1.0) \end{array}$	11 428 5069 1070	$\begin{array}{c} 0.1 \ (0.4) \\ 0.2 \ (0.7) \\ 0.2 \ (0.7) \end{array}$	$\begin{array}{c} 0.1 & (0.6) \\ 0.2 & (0.8) \\ 0.3 & (1.0) \end{array}$	$\begin{array}{c} 0.019^{*} \\ 0.334^{*} \\ 0.004^{*} \end{array}$	0.390* 0.846* 0.038*
LADL difficulties	Non-frail Pre-frail Frail	17 819 8567 1975	$\begin{array}{c} 0.2 \ (0.7) \\ 0.4 \ (1.0) \\ 0.4 \ (1.0) \end{array}$	11 429 5069 1070	$\begin{array}{c} 0.2 \ (0.8) \\ 0.4 \ (1.1) \\ 0.5 \ (1.3) \end{array}$	11 428 5069 1070	$\begin{array}{c} 0.2 \ (0.6) \\ 0.3 \ (0.9) \\ 0.4 \ (0.9) \end{array}$	$\begin{array}{c} 0.2 \ (0.8) \\ 0.4 \ (1.1) \\ 0.5 \ (1.3) \end{array}$	<0.001# 0.002 [#] 0.002 [#]	$\begin{array}{c} 0.010^{*} \\ 0.104^{*} \\ 0.041^{*} \end{array}$
# Repeated measu ADL: basic activit N: number of part	res ANOVA – test of ies of daily living; IAI icipants	within-subje JL: instrume	ects contrasts (wave * ental activities of dail	frailty cate y living	gory)					

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