

Cognitive frailty in relation to adverse health outcomes independent of multimorbidity: results from the China health and retirement longitudinal study

Chen Chen^{1,2}, JuYoung Park³, Chenkai Wu^{4,5}, QianLi Xue⁶, George Agogo^{7,8}, Ling Han⁸, Emiel O. Hoogendijk⁹, Zuyun Liu^{8,10}, Zunyou Wu¹

¹National Center for AIDS/STD Control and Prevention, Chinese Center for Disease Control and Prevention, Beijing, China

²National Institute of Environmental and Health, Chinese Center for Disease Control and Prevention, Beijing, China

³Florida Atlantic University, Phyllis and Harvey Sandler School of Social Work, Boca Raton, FL 33431, USA

⁴Global Health Research Center, Duke Kunshan University, Kunshan, Jiangsu, China

⁵Duke Global Health Institute, Duke University, Durham, NC 27710, USA

⁶Division of Geriatric Medicine and Gerontology and Center on Aging and Health, Johns Hopkins Medical Institute, Baltimore, MD 21224, USA

⁷Centers for Disease Control and Prevention (CDC), Village Market, Nairobi, Kenya

⁸Department of Internal Medicine, Yale School of Medicine, New Haven, CT 06510, USA

⁹Department of Epidemiology and Biostatistics, Amsterdam Public Health Research Institute, Amsterdam UMC-location VU University Medical Center, Amsterdam, Netherlands

¹⁰Department of Big Data in Health Science, School of Public Health and the Second Affiliated Hospital, Zhejiang University School of Medicine, Hangzhou, China

Correspondence to: Zuyun Liu, Zunyou Wu; **email:** Zuyun.liu@outlook.com, <https://orcid.org/0000-0001-6120-5913>; wuzy@263.net, <https://orcid.org/0000-0002-0839-4548>

Keywords: older adults, cognitive frailty, disability, death, multimorbidity

Received: June 15, 2020

Accepted: September 5, 2020

Published: November 18, 2020

Copyright: © 2020 Chen et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/3.0/) (CC BY 3.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

ABSTRACT

Our objectives were to evaluate: 1) the associations of cognitive frailty with various health outcomes including disability, hospitalization, and death; 2) whether the associations differed by multimorbidity. We included data of 5113 Chinese older adults (aged 60+ years) who had baseline cognition and physical frailty assessments (2011 wave) and follow-up for 4 years. About 16.0% (n=820) had cognitive impairment; 6.7% (n=342) had physical frailty; and 1.6% (n=82) met criteria for cognitive frailty. Both cognitive impairment (odds ratios (ORs) range: 1.41 to 2.11) and physical frailty (ORs range: 1.51 to 2.43) were independently associated with basic activities of daily living (BADL), instrumental ADL (IADL), mobility disability, hospitalization, and death among participants without that corresponding outcome at baseline, even after accounting for covariates. Relative to participants who had normal cognition and were nonfrail, those with cognitive frailty had the highest risk for IADL disability (OR=3.40, 95% CI, 1.23–9.40) and death (OR=3.89, 95% CI, 2.25–6.47). We did not find significant interaction effects between cognitive frailty and multimorbidity ($P_{interactions}>0.05$). Overall, cognitive frailty was associated with disability and death, independent of multimorbidity. This highlights the importance of assessing cognitive frailty in the community to promote primary and secondary preventions for healthy aging.

INTRODUCTION

Both cognitive impairment and physical frailty are important indicators of the aging process, and are associated with adverse health outcomes including disability, hospitalization, and death [1, 2]. Due to the potential interaction between these two aging indicators, they have been incorporated together as a new concept, cognitive frailty, defined as the simultaneous presence of both cognitive impairment and physical frailty in non-demented older adults, proposed in 2013 by an (I.A.N.A./I.A.G.G.) international consensus group [3]. Cognitive frailty describes a preclinical cognitive status caused by physical frailty instead of neurodegenerative disorders [4, 5]. The fundamental hypothesis about cognitive frailty is the direction of causality between physical frailty and cognitive impairment, i.e. physical frailty precedes and causes cognitive impairment [4]. Cognitive frailty is different from the frailty index (FI), which is based on the degree of accumulation of health deficits and represents an alternative instrument of frailty that incorporates more health dimensions, including comorbidities, psychological factors, symptoms, and disabilities [6]. However, the FI does not underline the temporal relationship between physical frailty and cognitive impairment. So the main benefit of the construction of cognitive frailty is to explain heterogeneity in etiology of cognitive impairment so as to improve intervention target.

A few studies have reported that cognitive frailty is more strongly associated with adverse health outcomes, such as disability in basic activities of daily living (BADL) [7–9], low quality of life [9], and death [8–14], than cognitive impairment or physical frailty alone [7–10, 15]. Nearly all of these studies were conducted in developed countries, with one exception (in China) [14], which is limited in study setting (e.g., one province) and small in sample size [14] (For detailed information see Supplementary Table 1). To date, little is known about whether cognitive impairment and physical frailty jointly associate with adverse outcomes in nationwide samples of older adults from developing countries like China. The investigation of this topic in different countries is crucial due to differences in population characteristics (e.g., lifestyle factors, pattern of chronic diseases). Moreover, in the context of dramatic population aging in China, early screening and intervention of at-risk older adults [16] may result in substantial reductions in healthcare expenditures, one of the key challenges in the near future.

Early identification of high-risk older adults is challenging partly due to the complexity of health issues that one older adult has. One key facet of such complexity is multimorbidity, characterized as co-

existence of multiple chronic diseases/conditions within one individual [17]. It has been reported that 42.4% of older adults in China have multimorbidity [18], which is associated with worse functioning [19], poor quality of life [19, 20], and high mortality [21]. Cognitive frailty has become a novel target for achieving successful ageing [22]; therefore, confirming whether cognitive frailty is associated with adverse health outcomes independent of multimorbidity is of great importance to prioritize potential preventive interventions; however, it has not been investigated in previous studies in both developed countries [7–13, 15] and in China [14].

To address these questions, we used data from the China Health and Retirement Longitudinal Study (CHARLS), a large cohort study with a nationally representative sample of middle-aged and older adults in China. The objective of this study was two-fold: first, to evaluate the associations of cognitive frailty with an array of health outcomes including disability (BADL, instrumental ADL [IADL], and mobility), hospitalization, and death; and second, to determine whether the associations differed by multimorbidity.

RESULTS

Characteristics of study participants

Among the 5113 participants at baseline, mean (SD) age was 68.1 (6.5) years, and 49% (n=2479) were women (Table 1). The majority of participants at baseline had less than six years of education (including no formal education, did not finish primary school and elementary school), were currently married, lived in rural areas, and were more likely to be non-smokers and non-drinkers.

At baseline, about 16.0% (n=820) of all participants had cognitive impairment; 6.7% (n=342) had physical frailty; and 1.6% (n=82) had both components, i.e., met criteria for cognitive frailty. As expected, the oldest-old (aged 80+ years) were more likely to have normal cognition and physical frailty (Group 3) or cognitive frailty (Group 4). A higher proportion of women were found in the nonfrail group with cognitive impairment (Group 2) and in the group with cognitive frailty (Group 4), which mainly represent older adults with cognitive impairment. About 44% of all participants had two or more chronic diseases (i.e., multimorbidity). Similarly, about 44% of participants in Group 4 had multimorbidity. A little higher proportion of participants in Group 3 (59%) had multimorbidity. Overall, we did not find obvious differences in the proportion of multimorbidity between the four cognitive frailty groups.

Table 1. Baseline characteristics of the study participants, CHARLS 2011.

Characteristic	Overall	Group 1 (Normal cognition & Nonfrail)	Group 2 (Cognitive impairment & Nonfrail)	Group 3 (Normal cognition & Frail)	Group 4 (Cognitive Frailty)	P-value
	N=5113	N=4033	N=738	N=260	N=82	
Age, years ^a	68.1 ± 6.5	67.9 ± 6.3	67.3 ± 6.0	73.0 ± 7.9	70.8 ± 7.4	<0.01
60-79 years old	4792 (93.7)	3804 (94.3)	714 (96.8)	205 (78.9)	69 (84.2)	<0.01
80+ years old	321 (6.3)	229 (5.7)	24 (3.3)	55 (21.2)	13 (15.9)	
Gender						<0.01
Women	2479 (48.5)	1758 (43.6)	541 (73.3)	123 (47.3)	57 (69.5)	
Men	2634 (51.5)	2275 (56.4)	197 (26.7)	137 (52.7)	25 (30.5)	
Education						<0.01
No formal education	1812 (35.4)	1077 (26.7)	550 (74.5)	117 (45.0)	68 (82.9)	
Did not finish primary school	1091 (21.3)	896 (22.2)	126 (17.1)	64 (24.6)	5 (6.1)	
Elementary school	1290 (25.2)	1179 (29.2)	51 (6.9)	52 (20.0)	8 (9.8)	
Middle school	614 (12.0)	578 (14.3)	9 (1.2)	26 (10.0)	1 (1.2)	
High school or above	306 (6.0)	303 (7.5)	2 (0.3)	1 (0.4)	0 (0)	
Marital status						<0.01
Currently married	4056 (79.3)	3274 (81.2)	559 (75.8)	175 (67.3)	48 (58.5)	
Others	1057 (20.7)	759 (18.8)	179 (24.3)	85 (32.7)	34 (41.5)	
Residence						<0.01
Rural	3233 (63.2)	2404 (59.6)	588 (79.7)	178 (68.5)	63 (76.8)	
Urban	1880 (36.8)	1629 (40.4)	150 (20.3)	82 (31.5)	19 (23.2)	
Smoking status						<0.01
Nonsmoker	2886 (56.5)	2152 (53.4)	545 (73.9)	137 (52.7)	52 (63.4)	
Ever smoker	623 (12.2)	549 (13.6)	34 (4.6)	35 (13.5)	5 (6.1)	
Smoker	1603 (31.4)	1331 (33.0)	159 (21.5)	88 (33.9)	25 (30.5)	
Alcohol consumption						<0.01
No	3019 (59.1)	2271 (56.4)	520 (70.6)	168 (64.6)	60 (73.2)	
Yes	2090 (40.9)	1759 (43.7)	217 (29.4)	92 (35.4)	22 (26.8)	
Body mass index ^a	22.9 ± 3.9	23.2 ± 3.9	22.4 ± 3.8	21.7 ± 4.7	21.4 ± 4.1	<0.01
Number of chronic diseases	1.5 ± 1.3	1.5 ± 1.3	1.4 ± 1.3	2.0 ± 1.4	1.5 ± 1.2	<0.01
Multimorbidity						<0.01
No	2856 (55.9)	2267 (56.2)	437 (59.2)	106 (40.8)	46 (56.1)	
Yes	2257 (44.1)	1766 (43.8)	301 (40.8)	154 (59.2)	36 (43.9)	

CHARLS, the China Health and Retirement Longitudinal Study.

Values are given as No. (percentages). ^a Values are given as Mean ± SD.

Percentages summed up by column, which may not sum to 100 because of rounding. There were missing data on smoking status (n=1), alcohol consumption (n=4), and body mass index (n=11).

Associations between cognitive frailty and adverse health outcomes

BADL, IADL, and mobility disability

Among the participants without BADL disability at baseline (n=3341), 33.6% (n=1121) reported BADL disability over 4-year follow-up. Similarly, the

percentages for incident IADL and mobility disability were 35.2% (n=1136) and 65.3% (n=1101), respectively.

Table 2 shows the individual and combined associations of cognitive impairment and physical frailty with adverse health outcomes, using the normal cognition

Table 2. Associations of cognitive impairment and physical frailty with disability (BADL, IADL, and mobility), hospitalization, and death in full sample, CHARLS 2011-2015.

		Model 1	Model 2
	No. of events/No. of participants	OR (95% CI)	OR (95% CI)
BADL disability			
Total	1121/3341		
Individual effect			
Cognitive impairment			
Normal cognition	924/2867	Ref.	Ref.
Cognitive impairment	197/474	1.41 (1.15–1.74)	1.38 (1.10–1.73)
Physical Frailty			
Nonfrail	1042/3207	Ref.	Ref.
Frail	79/134	2.38 (1.66–3.43)	2.16 (1.49–3.13)
Combined effect			
Group 1(Normal cognition & Nonfrail)	859/2758	Ref.	Ref.
Group 2(Cognitive impairment & Nonfrail)	183/449	1.43 (1.16–1.77)	1.40 (1.11–1.76)
Group 3(Normal cognition & Frail)	65/109	2.61 (1.74–3.90)	2.31 (1.53–3.48)
Group 4(Cognitive Frailty)	14/25	2.09 (0.93–4.71)	2.22 (0.97–5.08)
IADL disability			
Total	1136/3226		
Individual effect			
Cognitive impairment			
Normal cognition	924/2820	Ref.	Ref.
Cognitive impairment	212/406	2.11 (1.70–2.63)	1.79 (1.41–2.27)
Physical Frailty			
Nonfrail	1069/3114	Ref.	Ref.
Frail	67/112	2.43 (1.64–3.62)	2.13 (1.42–3.21)
Combined effect			
Group 1(Normal cognition & Nonfrail)	868/2725	Ref.	Ref.
Group 2(Cognitive impairment & Nonfrail)	201/389	2.13 (1.71–2.67)	1.81 (1.42–2.30)
Group 3(Normal cognition & Frail)	56/95	2.56 (1.66–3.94)	2.23 (1.43–3.47)
Group 4(Cognitive Frailty)	11/17	3.40 (1.23–9.40)	2.80 (1.00–7.87)
Mobility disability			
Total	1101/1685		
Individual effect			
Cognitive impairment			
Normal cognition	959/1499	Ref.	Ref.
Cognitive impairment	142/186	1.53 (1.06–2.21)	1.62 (1.09–2.40)
Physical Frailty			
Nonfrail	1075/1652	Ref.	Ref.
Frail	26/33	1.51 (0.64–3.59)	1.47 (0.61–3.53)
Combined effect			
Group 1(Normal cognition & Nonfrail)	937/1470	Ref.	Ref.
Group 2(Cognitive impairment & Nonfrail)	138/182	1.51 (1.04–2.18)	1.58 (1.06–2.36)
Group 3(Normal cognition & Frail)	22/29	1.38 (0.57–3.33)	1.29 (0.52–3.18)
Group 4(Cognitive Frailty)	4/4	1*	1*
Hospitalization			
Total	1075/3776		
Cognitive impairment			
Normal cognition	891/3164	Ref.	Ref.
Cognitive impairment	184/612	1.14 (0.94–1.39)	1.26 (1.02–1.56)
Physical Frailty			
Nonfrail	994/3561	Ref.	Ref.

Frail	81/215	1.37 (1.02–1.83)	1.34 (0.99–1.80)
Combined effect			
Group 1(Normal cognition & Nonfrail)	825/2997	Ref.	Ref.
Group 2(Cognitive impairment & Nonfrail)	169/564	1.17 (0.96–1.44)	1.29 (1.04–1.62)
Group 3(Normal cognition & Frail)	66/167	1.50 (1.08–2.07)	1.43 (1.03–2.00)
Group 4(Cognitive Frailty)	15/48	1.11 (0.60–2.08)	1.29 (0.68–2.44)
Death			
Total	434/5113		
Individual effect			
Cognitive impairment			
Normal cognition	353/4293	Ref.	Ref.
Cognitive impairment	81/820	1.58 (1.21–2.08)	1.49 (1.11–2.00)
Physical Frailty			
Nonfrail	363/4771	Ref.	Ref.
Frail	71/342	2.07 (1.53–2.81)	1.82 (1.33–2.48)
Combined effect			
Group 1(Normal cognition & Nonfrail)	303/4033	Ref.	Ref.
Group 2(Cognitive impairment & Nonfrail)	60/738	1.42 (1.05–1.92)	1.34 (0.97–1.85)
Group 3(Normal cognition & Frail)	50/260	1.82 (1.28–2.58)	1.59 (1.11–2.28)
Group 4(Cognitive Frailty)	21/82	3.89 (2.25–6.74)	3.49 (2.00–6.15)

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activity of daily living; IADL, instrumental activity of daily living; OR, odds ratio; CI, confidence interval.

As described in Methods, we ran a logistic regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (i.e., N=3341 participants without BADL disability at baseline). Odds ratios (ORs) and corresponding 95% confidence interval (CI) are presented. Model 1 adjusted for age and gender. Model 2 further adjusted for education, residence, marital status, smoking status, alcohol consumption, body mass index, and multimorbidity.

*Since all participants in this subgroup reported the health outcome over the follow-up period, we assigned 1 to them.

and nonfrail group (Group 1) as the reference group. After adjusting for age and gender, both cognitive impairment and physical frailty significantly increased the risk of developing BADL, IADL, and mobility disability, with ORs ranging from 1.41 (95% CI: 1.15–1.74) to 2.43 (95% CI: 1.64–3.62). In combined effect analyses, belonging to Group 4 (cognitive frailty) was strongly associated with an increased risk for BADL and IADL disability, with ORs of 2.09 and 3.40, respectively, although the OR for BADL was not statistically significant, due to the small sample size for Group 4. Only four participants were free of mobility disability at baseline, which all developed mobility disability during the follow-up period.

Hospitalization

About 28.5% (n=1075) of the participants who did not report hospitalization at baseline reported a hospitalization during 4-year follow-up. In individual effect analyses, physical frailty (OR: 1.37, 95% CI: 1.02–1.83), but not cognitive impairment (OR: 1.14, 95% CI: 0.94–1.39), was independently associated with incidence of hospitalization. In combined effect analyses, compared with Group 1, the other three

groups were all associated with elevated risk for hospitalization, though only Group 3 (Normal cognition and frail; OR: 1.50, 95% CI, 1.08–2.07, Table 2) met statistical significance.

Death

Over 4-year follow-up, 434 of 5113 (8.5%) participants died. In individual effect analyses, both cognitive impairment and physical frailty were independently associated with an increased risk of death (OR=1.58 and 2.07, respectively). In combined effect analyses, all groups were associated with elevated risk of death (ORs range from 1.42 to 3.89). Compared with Group 1, Group 4 (cognitive frailty) had a nearly four-fold increased risk of death (OR=3.89, 95% CI, 2.25–6.74, Table 2),

For all (individual and combined) associations above, we did not observe substantial changes of the results when additionally adjusting for education, residence, marital status, smoking status, alcohol consumption, body mass index, and multimorbidity (Table 2, Model 2). Furthermore, additional analyses revealed that for the above individual effect analyses, mutually

controlled associations of cognitive impairment and physical frailty with all outcomes in logistic regression models did not affect the main model results (Supplementary Table 2).

Subgroup analyses by multimorbidity

We found that after adjusting for age and gender, multimorbidity significantly increased the risk of the five adverse health outcomes (Supplementary Table 3), which suggested that multimorbidity could play a role as moderator or confounder of the associations between cognitive frailty and adverse health outcomes.

Table 3 presents the associations of cognitive impairment and physical frailty with disability (ADL, IADL, and mobility), hospitalization, and death in older adults without and with multimorbidity. We did not find significant interaction effects between cognitive frailty and multimorbidity for these adverse outcomes (all interaction P values > 0.05). As shown in Table 3, we found that in participants without multimorbidity, the risk estimates were similar to those shown in the full sample, where cognitive frailty was a strong predictor for the health outcomes (ORs range from 3.59 to 4.47) except for hospitalization. In participants with multimorbidity, partly due to the small sample size, we did not observe statistically significant associations of cognitive frailty with the health outcomes, but the overall trend suggested that older adults with cognitive frailty were at high risk of IADL disability and death.

Sensitivity analyses

In sensitivity analyses, we found that: (1) the results were unchanged after removing 146 participants who reported having memory-related disease (dementia, Parkinson's disease, etc., Supplementary Table 4); (2) accounting for the competing risk of death did not substantially change the associations of cognitive frailty with disabilities and hospitalization appreciably (Supplementary Table 5); (3) the cross-sectional analyses obtained relatively higher risk estimates and further offered supportive evidence for the longitudinal relationships of cognitive impairment, physical frailty, and their combinations with adverse health outcomes (Supplementary Table 6).

DISCUSSION

To facilitate early identification of high-risk older adults in China, we evaluated the association of one important new construct, cognitive frailty, with a series of adverse outcomes in a large sample from CHARLS, a cohort study of middle-aged and older Chinese adults. We found that older adults with cognitive frailty have the

highest risks of BADL, IADL disability and death, relative to their counterparts who had normal cognition and were nonfrail, even after accounting for several covariates. Overall, these associations did not differ by multimorbidity. These findings suggest the possibility and importance of using cognitive frailty as a tool to help with risk stratification in the general Chinese older population.

In this sample of community-dwelling older adults, 1.6% met the criteria for cognitive frailty, in line with a recent literature review [5], which reported that the prevalence of cognitive frailty ranges from 1% to 5% in community-dwelling older adults worldwide. We acknowledged the influence of the exclusion of participants with missing data on cognition or physical frailty components, which is inevitable in large-scale epidemiological investigations. Nevertheless, this prevalence is remarkable when it comes to the very large population of older adults in China (i.e., 249 million aged 60+ years in 2018). That being said, approximately 3.9 million older adults may have cognitive frailty in China, and may require urgent and considerable attention.

Overall, our results of the combined associations of cognition and physical frailty with adverse outcomes were consistent with previous research (Supplementary Table 1), concerning death [7, 9, 10, 12–14, 23], BADL [7–9, 23], IADL [8, 9, 15, 23], mobility disability [8, 23], and hospitalization [8, 9, 14], although in the current study the estimated risk for cognitive frailty with regard to hospitalization was not statistically significant. These non-significant findings regarding hospitalization may reflect diverse pathways or risk factor profiles [23]. For example, hospitalization is often the result of acute illness or injury rather than the result of cognitive impairment or physical frailty itself.

To the best of our knowledge, this is the first study to explore the role of multimorbidity in the associations of cognitive frailty with various health outcomes in Chinese older adults. The comparable findings in subgroups with and without multimorbidity suggest that cognitive frailty and multimorbidity do not necessarily share the same underlying pathophysiologic pathways and mechanisms towards disability and death. Considering the complexity of aging, beyond the physical and cognitive dimensions (which cognitive frailty captures), multimorbidity per se could result from various factors such as nutrition [24] and mental health [25] (although these factors are also associated with cognitive frailty), and could lead to many adverse health outcomes. Finally, we acknowledge that the potential recall bias, mostly underreported, resulted from the self-report of multimorbidity, might introduce

Table 3. Associations of Cognitive Impairment and Physical Frailty with Disability, Hospitalization, and Death in persons without or with multimorbidity, CHARLS 2011-2015.

	BADL disability	IADL disability	Mobility disability	Hospitalization	Death
Without multimorbidity	N=2011	N=1913	N=1176	N=2212	N=2856
Group 1	Ref.	Ref.	Ref.	Ref.	Ref.
Group 2	1.44 (1.09–1.91)	2.60 (1.96–3.45)	1.52 (1.02–2.26)	1.22 (0.93–1.60)	1.86 (1.27–2.73)
Group 3	2.53 (1.42–4.52)	2.62 (1.37–5.00)	1.35 (0.41–4.46)	1.33 (0.80–2.22)	1.42 (0.80–2.52)
Group 4	3.68 (1.38–9.80)	4.47 (1.28–15.58)	1*	1.33 (0.59–2.98)	3.59 (1.67–7.80)
With multimorbidity	N=1330	N=1313	N=509	N=1564	N=2257
Group 1	Ref.	Ref.	Ref.	Ref.	Ref.
Group 2	1.71 (1.21–2.42)	1.86 (1.28–2.72)	2.34 (0.80–6.92)	1.21 (0.89–1.65)	0.98 (0.58–1.63)
Group 3	2.31 (1.31–4.09)	2.13 (1.19–3.81)	1.04 (0.27–3.94)	1.42 (0.93–2.18)	1.99 (1.26–3.12)
Group 4	0.79 (0.17–3.60)	2.58 (0.44–15.01)	–	1.91 (0.34–2.44)	4.34 (1.96–9.60)
Interaction P value:					
Cognitive frailty × multimorbidity	0.232	0.525	0.552	0.883	0.227

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activity of daily living; IADL, instrumental activity of daily living; Group 1, Normal cognition & Nonfrail; Group 2, Cognitive impairment & Nonfrail; Group 3, Normal cognition & Frail; Group 4, Cognitive Frailty.

As described in Methods, we ran a logistic regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (e.g., N=2011 participants without BADL disability at baseline and had no multimorbidity). Odds ratios (ORs) and corresponding 95% confidence interval (CI) are presented. All models were adjusted for age and gender.

*Since all participants in this subgroup reported the health outcome over the follow-up period, we assigned 1 to them.

exposure misclassification on multimorbidity and might obscure the true estimated associations, although the estimated prevalence of multimorbidity (43.8%) in this study was consistent with previous studies based on CHARLS (42.4%) [18] and other Chinese cohorts (44.4%) [26].

The clinical utility and relevance of our findings should be placed in the context of the large older population in China and relatively limited healthcare resources. As mentioned above, about 3.9 million older adults may suffer from cognitive frailty. We have demonstrated that cognitive frailty, relative to cognitive impairment and physical frailty individually, could be a better measure of vulnerability because it is more sensitive and thus captures more susceptible older adults who were currently non-disabled but at high risk of developing disability (such as, BADL) or other subsequent adverse consequences. Early identification of older adults with cognitive frailty has public health implications, in both clinical practice and large epidemiological screening settings, considering the fact that there are still no effective pharmacological treatments to improve physical frailty status [27] and cognitive impairment [28]. Targeting this specific population would be a priority, especially when considering that interventions may be more effective if applied at an early stage. Nowadays, many comprehensive geriatric assessments are being done in China [29, 30], which include detailed

physical examinations and offer an opportunity for the application of cognitive frailty. Since the assessment of cognitive frailty is not simple currently, appropriate implementation while conducting comprehensive geriatric assessment should be explored in the near future.

In the current study, data from a nationwide prospective cohort study in China and the availability of data on multiple health outcomes in a relatively large cohort provided a unique opportunity to evaluate the associations between cognition, physical frailty, and multiple adverse outcomes, of which some (e.g., IADL, mobility disability, and hospitalization) have not been investigated previously. Second, the high quality of the data, low rate of loss-to-follow-up, and comparable study design of the CHARLS (compared to other international Health and Retirement Study sister studies [31]) strengthen our findings. Future research could attempt to explore the associations of cognitive frailty and health outcomes in more countries and cohorts (e.g., UK BIOBANK).

The current study nevertheless has several limitations. First, we used a modified definition to construct the physical frailty phenotype compared with the one proposed by Fried (e.g., the use of a self-reported measure of activity intensity-whether walk 10 or more minutes continuously during a usual week-rather than

the original kilocalories expended index), which would influence the proportion of physical frailty to some extent. However, the utility and validity of this definition for physical frailty in identifying frail Chinese adults have been previously demonstrated [32]. Second, the timing of the occurrence of health outcomes was not available in this study. Third, the exclusion of participants with missing data on physical frailty or cognition and/or the restriction to participants without outcome at baseline (in Table 2) would have induced selection bias and resulted in limited sample size for some outcomes (e.g., mobility disability). Yet, the effects of this bias are probably relatively small since we observed similar results across these health outcomes while the missingness was randomized to some extent. Fourth, although the overall sample size is relatively large, some group (such as Group 4, cognitive frailty) are still not large enough in subgroup analysis, which may lead to false positive results. At the same time, we observed a consistent trend, that is, from Group 1 to Group 4, the risks of outcomes are gradually increasing, which has little to do with the power itself. Finally, the study attrition (e.g., death, loss-to-follow-up) could be problematic in the longitudinal cohort. However, when running competing risk Cox regression models to account for the influence of death, we did not find substantial changes to the results (Supplementary Table 5).

CONCLUSIONS

In a large sample of Chinese community-dwelling older adults, we demonstrated that overall, cognitive frailty was associated with disability and death, independent of multimorbidity. In the context of rapid population aging, these findings highlight the importance of assessing cognitive frailty in community to promote primary and secondary preventions for healthy aging.

MATERIALS AND METHODS

Study population

The CHARLS targeted Chinese community-dwelling adults aged 45 years and older and their spouses. The CHARLS used a multistage sampling strategy covering 28 provinces, 150 counties/districts, and 450 villages/urban communities across the country. Participants were first recruited in 2011/2012, and completed two follow-up visits biennially up to 2015/2016. Details of the CHARLS survey have been described elsewhere [31]. The CHARLS study offers a wide range of information on socioeconomic status and health, including demographic characteristics, family structure, health status and functioning, biomarkers,

health care and insurance, work, retirement and pension, income and consumption, and assets.

Out of 17,708 participants aged 45 years and older enrolled in the baseline survey (2011/2012), we excluded those with missing data on age (n=26), aged <60 years (n=10229), with missing data on cognition (n=623) and four or more physical frailty components (n=1717), leaving 5113 participants available (Figure 1). Finally, for different adverse health outcomes, we assembled various sets of analytical samples, as shown below:

Sample 1: for BADL disability: restricted to participants without BADL disability at baseline (n=3341);

Sample 2: for IADL disability: restricted to participants without IADL disability at baseline (n=3226);

Sample 3: for mobility disability: restricted to participants without mobility disability at baseline (n=1685);

Sample 4: for hospitalization: restricted to participants without hospitalization occurred at baseline (n=3776);

Sample 5: for death: all participants (n=5113).

Measures

Cognitive impairment

Following prior CHARLS publications [33, 34], we used three assessments of cognition: the Telephone Interview of Cognitive Status (TICS-10), word recall, and figure drawing. The overall cognition score is the sum score of the TICS-10 (orientation and attention, 0-10), word recall (episodic memory, 0-10), and figure drawing (visual spatial abilities, 0-1). The TICS is a well-established measure for assessing mental status [35]. In CHARLS, ten items from TICS were included: date (month, day, and year), day of the week, season of the year, and serial subtraction of 7 from 100 (up to five times). Summing the number of correct answers to these questions resulted in a TICS-10 score, ranging from 0 to 10. TICS-10 primarily assesses the executive functions of orientation to time and attention [36]. For word recall, participants were asked to memorize and immediately recall as many words as they could in any order immediately after interviewers read a list of 10 Chinese nouns (i.e., immediate recall). Four to 10 minutes later, participants were asked to recall as many of the original words as possible (i.e., delayed recall) [34, 35]. The average number of immediate and delayed word recalls ranged from 0 to 10, and is regarded as assessing episodic memory [37]. For figure drawing, participants were shown a picture of two overlapped pentagons and asked to draw a similar figure. Participants who successfully completed the task received a score of 1, and those who failed received a score of 0. This task assessed visuospatial abilities [38]. A summary score of the three

parts range from 0 to 21, with higher scores indicating better performance [33, 34]. According to the literature, participants were classified as having cognitive impairment if their summary score fell more than 1 SD below age-appropriate norms; otherwise, they were defined as normal cognition [39].

Physical frailty

Frailty was measured by an adapted version of the Fried physical frailty phenotype approach [40], and has been previously developed and validated in the CHARLS [32]. Five items/criteria are included: shrinking, weakness, exhaustion, slowness, and inactivity. Shrinking was established as self-reported loss of 5 or more kg in the previous year or having a body mass index (BMI) of 18.5 kg/m² or less. Weakness was defined as the lowest quintile on maximum handgrip strength (either hand; two trials for each; measured in a standing position with arm bended at 90 degrees) among the population, adjusting for gender and BMI. Exhaustion was determined according to two questions from the Center for Epidemiological Studies-Depression scale: “I could not get going” and “I felt everything I did was an effort”. Slowness was defined as the lowest quintile on

the average of two-timed walk tests over a 2.5-meter course, at usual pace, among the population, adjusting for gender and standing height. Inactivity was determined if participants self-reported that they did not walk 10 or more minutes continuously during a usual week. We modified the original physical frailty phenotype proposed by Fried et al. [40], due to limited availability of data in CHARLS. Participants who met three or more criteria were defined as having physical frailty (frail); otherwise, they were considered as having no physical frailty (nonfrail).

Cognitive frailty

In line with the definition by an (I.A.N.A./I.A.G.G.) international consensus group [3], cognitive frailty was defined as the simultaneous presence of both cognitive impairment and physical frailty [3], and has been previously validated [41–43]. Based on the two components—cognitive impairment and physical frailty, we defined four combined groups:

- Group 1: normal cognition and nonfrail
- Group 2: cognitive impairment and nonfrail
- Group 3: normal cognition and frail
- Group 4: cognitive impairment and frail (cognitive frailty)

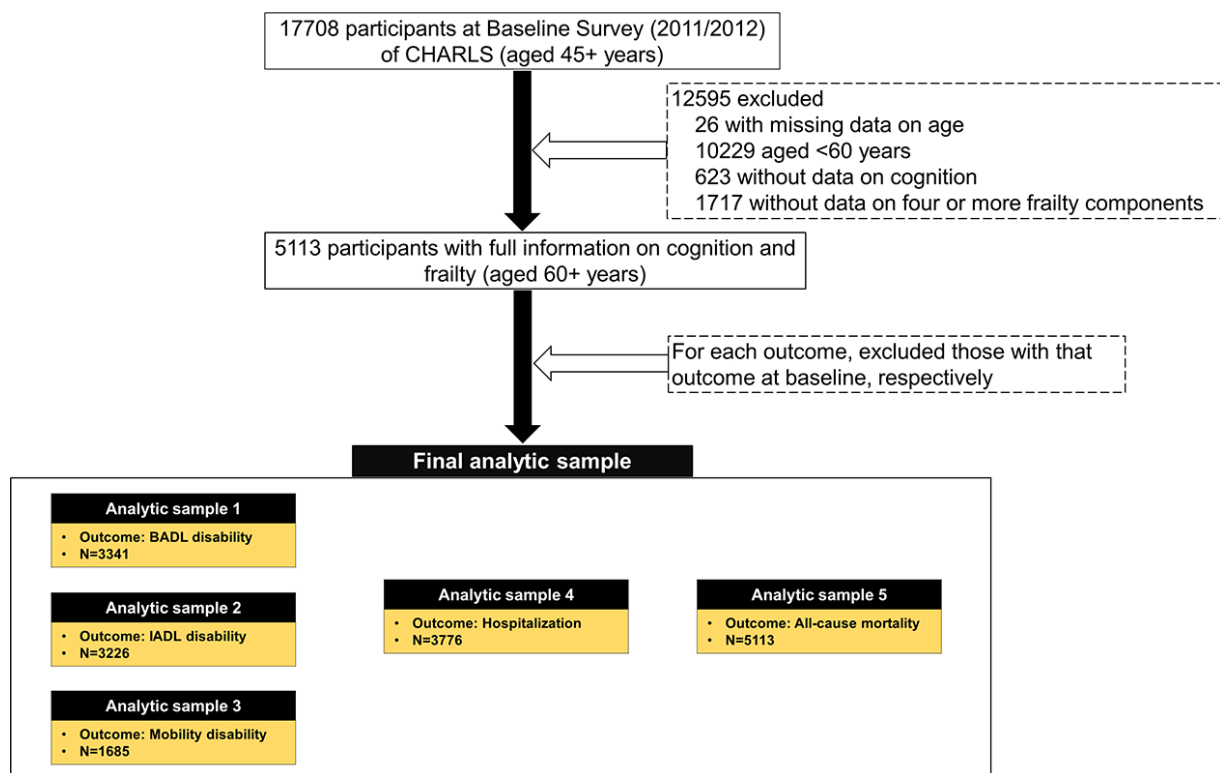


Figure 1. Flow chart of analytic sample. CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activity of daily living; IADL, instrumental activity of daily living.

Disability and hospitalization

We considered important patient-centered outcomes: disability and hospitalization, as we did before [23]. Three domains (BADL, IADL, and mobility) of disability were operationalized, including 5 BADL (dressing, bathing, eating, getting in/out of bed, and using the toilet), 5 IADL (managing money, taking medications, shopping for groceries, meal preparation, and cleaning house), and 7-item mobility activities (100m, climbing several flights of stairs, getting up from a chair, stooping or kneeling or crouching, extending arms up, lifting 11 lb, and picking up a small coin). For each task in the three domains, participants were asked, “Do you have difficulty in performing the task?” Those participants who need personal assistance in performing one or more of the corresponding activities in each domain were defined as disability. Hospitalization was derived from the participants’ self-report of whether they received any inpatient care in the past year. Since the timing of developing disability during the follow-up period was not available [44], we defined a binary outcome to denote the occurrence of disability over the 4-year follow-up. For hospitalization, we did a similar definition to denote whether the participant had hospitalization over the 4-year follow-up since baseline in this study.

Death

The death information in CHARLS was collected from the exit interview in 2013 and 2015 waves. Generally, a binary variable representing occurrence of death and a variable representing date of death were provided. However, date of death was only available in the 2013 wave according to the current dataset in the CHARLS website. Therefore, we constructed a binary variable to denote occurrence of death within the 4-year follow-up since baseline in this study.

Covariates

Demographic characteristics including age, gender, residence (urban vs rural), education, and marital status were collected at baseline. Five categories were considered for education: no formal education/illiterate, can read but did not finish elementary school, elementary school/traditional Chinese school, middle school, and high school or above. Marital status was defined as currently married (including partnered), and others (e.g., separated, divorced, widowed). Health behaviors included smoking status (never, ever, and currently smoking), alcohol consumption (yes/no), and body mass index (BMI, kg/m²).

Self-reported comorbidity included ten self-reported chronic diseases by asking “Have you been diagnosed

with the following conditions by a doctor”: hypertension; diabetes or high blood sugar; cancer or malignant tumor; chronic lung disease; heart problems; stroke; kidney disease; stomach or other digestive disease; arthritis or rheumatism; and asthma. The total number of chronic diseases was calculated. Multimorbidity was defined as co-existence of two or more chronic diseases within one older adult [18].

Statistical analyses

Characteristics of the study population were presented as means (\pm standard deviation [SD]) or frequencies (percentage) in the overall sample, and four combined groups (Group 1–4).

Logistic regression models were used to estimate the individual (i.e., cognitive impairment and physical frailty as independent categorical variable respectively) and combined associations (i.e., the I.A.N.A./I.A.G.G. criteria defined four cognitive frailty groups as independent categorical variable, with Group 1 as the reference group) of cognitive impairment and physical frailty with adverse health outcomes. Note that we ran a logistic regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (e.g., n=3341 participants without BADL disability at baseline). We adjusted for age and gender in model 1 and presented the odds ratios (ORs) and corresponding 95% confidence interval (CI). We further adjusted for other covariates including residence, education, marital status, smoking status, alcohol consumption, BMI, and multimorbidity in model 2.

To evaluate whether the associations of cognitive frailty with health outcomes differed by multimorbidity, we added the interaction terms: the four cognitive frailty groups \times multimorbidity in the model 1. We then repeated the above analyses in two subgroups: those with and without multimorbidity and presented the results separately.

Sensitivity analyses were performed to test the robustness of the results. First, as the information on dementia was not available in CHARLS, we used one item—self-reported memory related problem (dementia, Parkinson's disease, etc.) as a proxy for dementia. We excluded participants with a response of “Yes” to this question and rerun the model 1 above to testify whether the individual and combined associations of cognitive impairment and physical frailty with adverse health outcomes changed. Second, to address the concern about the attrition of study participants, particularly due to death, we used a competing risk Cox regression model [45] to re-examine the above individual and

combined associations of cognitive impairment and physical frailty with disability and hospitalization. To do so, we assumed that the date of incurring disability, hospitalization, or death was the time of the survey if the participant reported having that disability, hospitalization, or death (exit interview). The follow-up time was calculated from the baseline to the date of incurring disability (hospitalization) or death, whichever came first. Finally, we ran an additional analysis by examining the cross-sectional association of cognitive impairment, physical frailty, and their combinations with the health outcomes, in order to confirm the findings from the majority of the literature on a cross-section study design.

All statistical analyses were performed using SAS version 9.4 (SAS Institute, Cary, NC). $P < 0.05$ (two-tailed) was considered as statistically significant.

Abbreviations

CHARLS: China Health and Retirement Longitudinal Study; BADL: Basic activities of daily living; IADL: Instrumental activities of daily living; TICS: Telephone Interview of Cognitive Status.

AUTHOR CONTRIBUTIONS

ZY.W. and ZY.L. conceived and designed this research, ZY.L. and C.C. collected and managed the data, C.C. performed statistical analyses, ZY.L. supervised and validated the data analyses. C.C. wrote the initial draft of the paper, ZY.W., ZY.L., QL.X. and EO.H. provided constructive comments for the revision of the paper, JY.P., CK.W., G.A. and L.H. reviewed and edited the manuscript. All authors approved the final manuscript.

ACKNOWLEDGMENTS

We thank all participants in the China Health and Retirement Longitudinal Study. We thank the CHARLS research team for providing the data. We thank Jennifer M. McGoogan for comments and editing assistance.

CONFLICTS OF INTEREST

The authors declare that they have no conflicts of interest.

FUNDING

The data used in this study are from China Health and Retirement Longitudinal Study (CHARLS). We thank the National Institute on Aging (NIA) in the United States (1-R21-AG031372-01, 1-R21-AG033675-01-A1, 1-R01-AG037031-01, and 1-R01-AG037031-03S1), the

National Natural Science Foundation of China (70773002, 70910107022, and 71130002), and the World Bank (7159234) for the support for the CHARLS. The current study was conducted partially at School of Public Health and the Second Affiliated Hospital, Zhejiang University School of Medicine (Hundred Talents Program). This study was supported by the Fundamental Research Funds for the Central Universities (China); and an NWO/ZonMw Veni fellowship (91618067). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

REFERENCES

1. Clegg A, Young J, Iliffe S, Rikkert MO, Rockwood K. Frailty in elderly people. *Lancet*. 2013; 381:752–62. [https://doi.org/10.1016/S0140-6736\(12\)62167-9](https://doi.org/10.1016/S0140-6736(12)62167-9) PMID:[23395245](https://pubmed.ncbi.nlm.nih.gov/23395245/)
2. Panza F, Lozupone M, Solfrizzi V, Sardone R, Dibello V, Di Lena L, D'Urso F, Stallone R, Petrucci M, Giannelli G, Quaranta N, Bellomo A, Greco A, et al. Different cognitive frailty models and health- and cognitive-related outcomes in older age: from epidemiology to prevention. *J Alzheimers Dis*. 2018; 62:993–1012. <https://doi.org/10.3233/JAD-170963> PMID:[29562543](https://pubmed.ncbi.nlm.nih.gov/29562543/)
3. Kelaiditi E, Cesari M, Canevelli M, van Kan GA, Ousset PJ, Gillette-Guyonnet S, Ritz P, Duveau F, Soto ME, Provencher V, Nourhashemi F, Salvà A, Robert P, et al, and IANA/IAGG. Cognitive frailty: rational and definition from an (I.A.N.A./I.A.G.G.) international consensus group. *J Nutr Health Aging*. 2013; 17:726–34. <https://doi.org/10.1007/s12603-013-0367-2> PMID:[24154642](https://pubmed.ncbi.nlm.nih.gov/24154642/)
4. Xue QL, Buta B, Ma L, Ge M, Carlson M. Integrating frailty and cognitive phenotypes: why, how, now what? *Curr Geriatr Rep*. 2019; 8:97–106. PMID:[31815092](https://pubmed.ncbi.nlm.nih.gov/31815092/)
5. Arai H, Satake S, Kozaki K. Cognitive frailty in geriatrics. *Clin Geriatr Med*. 2018; 34:667–75. <https://doi.org/10.1016/j.cger.2018.06.011> PMID:[30336994](https://pubmed.ncbi.nlm.nih.gov/30336994/)
6. Mitnitski AB, Mogilner AJ, Rockwood K. Accumulation of deficits as a proxy measure of aging. *ScientificWorldJournal*. 2001; 1:323–36. <https://doi.org/10.1100/tsw.2001.58> PMID:[12806071](https://pubmed.ncbi.nlm.nih.gov/12806071/)
7. Aliberti MJ, Cenzer IS, Smith AK, Lee SJ, Yaffe K, Covinsky KE. Assessing risk for adverse outcomes in older adults: the need to include both physical frailty and cognition. *J Am Geriatr Soc*. 2019; 67:477–83. <https://doi.org/10.1111/jgs.15683> PMID:[30468258](https://pubmed.ncbi.nlm.nih.gov/30468258/)

8. Avila-Funes JA, Amieva H, Barberger-Gateau P, Le Goff M, Raoux N, Ritchie K, Carrière I, Tavernier B, Tzourio C, Gutiérrez-Robledo LM, Dartigues JF. Cognitive impairment improves the predictive validity of the phenotype of frailty for adverse health outcomes: the three-city study. *J Am Geriatr Soc.* 2009; 57:453–61. <https://doi.org/10.1111/j.1532-5415.2008.02136.x> PMID:[19245415](https://pubmed.ncbi.nlm.nih.gov/19245415/)
9. Feng L, Zin Nyunt MS, Gao Q, Feng L, Yap KB, Ng TP. Cognitive frailty and adverse health outcomes: findings from the Singapore longitudinal ageing studies (SLAS). *J Am Med Dir Assoc.* 2017; 18:252–58. <https://doi.org/10.1016/j.jamda.2016.09.015> PMID:[27838339](https://pubmed.ncbi.nlm.nih.gov/27838339/)
10. St John PD, Tyas SL, Griffith LE, Menec V. The cumulative effect of frailty and cognition on mortality - results of a prospective cohort study. *Int Psychogeriatr.* 2017; 29:535–43. <https://doi.org/10.1017/S1041610216002088> PMID:[27903307](https://pubmed.ncbi.nlm.nih.gov/27903307/)
11. Cano C, Samper-Ternent R, Al Snih S, Markides K, Ottenbacher KJ. Frailty and cognitive impairment as predictors of mortality in older Mexican Americans. *J Nutr Health Aging.* 2012; 16:142–47. <https://doi.org/10.1007/s12603-011-0104-7> PMID:[22323349](https://pubmed.ncbi.nlm.nih.gov/22323349/)
12. Solfrizzi V, Scafato E, Lozupone M, Seripa D, Giannini M, Sardone R, Bonfiglio C, Abbrescia DI, Galluzzo L, Gandin C, Baldereschi M, Di Carlo A, Inzitari D, et al, and Italian Longitudinal Study on Aging Working Group. Additive role of a potentially reversible cognitive frailty model and inflammatory state on the risk of disability: the italian longitudinal study on aging. *Am J Geriatr Psychiatry.* 2017; 25:1236–48. <https://doi.org/10.1016/j.jagp.2017.05.018> PMID:[28689645](https://pubmed.ncbi.nlm.nih.gov/28689645/)
13. Solfrizzi V, Scafato E, Seripa D, Lozupone M, Imbimbo BP, D’Amato A, Tortelli R, Schilardi A, Galluzzo L, Gandin C, Baldereschi M, Di Carlo A, Inzitari D, et al, and Italian Longitudinal Study on Aging Working Group. Reversible cognitive frailty, dementia, and all-cause mortality. The italian longitudinal study on aging. *J Am Med Dir Assoc.* 2017; 18:89.e1–8. <https://doi.org/10.1016/j.jamda.2016.10.012> PMID:[28012505](https://pubmed.ncbi.nlm.nih.gov/28012505/)
14. Yu R, Morley JE, Kwok T, Leung J, Cheung O, Woo J. The effects of combinations of cognitive impairment and pre-frailty on adverse outcomes from a prospective community-based cohort study of older Chinese people. *Front Med (Lausanne).* 2018; 5:50. <https://doi.org/10.3389/fmed.2018.00050> PMID:[29600247](https://pubmed.ncbi.nlm.nih.gov/29600247/)
15. Shimada H, Makizako H, Lee S, Doi T, Lee S, Tsutsumimoto K, Harada K, Hotta R, Bae S, Nakakubo S, Harada K, Suzuki T. Impact of cognitive frailty on daily activities in older persons. *J Nutr Health Aging.* 2016; 20:729–35. <https://doi.org/10.1007/s12603-016-0685-2> PMID:[27499306](https://pubmed.ncbi.nlm.nih.gov/27499306/)
16. Wu Y, Dang J. China report of the development on aging cause. Social Sciences Academic Press: Beijing, China, 2013.
17. Organization WH, World report on ageing and health. 2015: World Health Organization.
18. Yao SS, Cao GY, Han L, Chen ZS, Huang ZT, Gong P, Hu Y, Xu B. Prevalence and patterns of multimorbidity in a nationally representative sample of older Chinese: results from CHARLS. *J Gerontol A Biol Sci Med Sci.* 2019. [Epub ahead of print]. <https://doi.org/10.1093/gerona/glz185> PMID:[31406983](https://pubmed.ncbi.nlm.nih.gov/31406983/)
19. Pati S, Swain S, Hussain MA, van den Akker M, Metsemakers J, Knottnerus JA, Salisbury C. Prevalence and outcomes of multimorbidity in South Asia: a systematic review. *BMJ Open.* 2015; 5:e007235. <https://doi.org/10.1136/bmjopen-2014-007235> PMID:[26446164](https://pubmed.ncbi.nlm.nih.gov/26446164/)
20. Gu J, Chao J, Chen W, Xu H, Zhang R, He T, Deng L. Multimorbidity and health-related quality of life among the community-dwelling elderly: a longitudinal study. *Arch Gerontol Geriatr.* 2018; 74:133–40. <https://doi.org/10.1016/j.archger.2017.10.019> PMID:[29096228](https://pubmed.ncbi.nlm.nih.gov/29096228/)
21. Prince MJ, Wu F, Guo Y, Gutierrez Robledo LM, O’Donnell M, Sullivan R, Yusuf S. The burden of disease in older people and implications for health policy and practice. *Lancet.* 2015; 385:549–62. [https://doi.org/10.1016/S0140-6736\(14\)61347-7](https://doi.org/10.1016/S0140-6736(14)61347-7) PMID:[25468153](https://pubmed.ncbi.nlm.nih.gov/25468153/)
22. Ruan Q, Yu Z, Chen M, Bao Z, Li J, He W. Cognitive frailty, a novel target for the prevention of elderly dependency. *Ageing Res Rev.* 2015; 20:1–10. <https://doi.org/10.1016/j.arr.2014.12.004> PMID:[25555677](https://pubmed.ncbi.nlm.nih.gov/25555677/)
23. Liu Z, Han L, Gahbauer EA, Allore HG, Gill TM. Joint trajectories of cognition and frailty and associated burden of patient-reported outcomes. *J Am Med Dir Assoc.* 2018; 19:304–09.e2. <https://doi.org/10.1016/j.jamda.2017.10.010> PMID:[29146224](https://pubmed.ncbi.nlm.nih.gov/29146224/)
24. Moraes MB, Araujo CF, Avgerinou C, Vidal EI. Nutritional interventions for the treatment of frailty in older adults: a systematic review protocol. *Medicine (Baltimore).* 2018; 97:e13773.

- <https://doi.org/10.1097/MD.00000000000013773>
PMID:30593155
25. Brown PJ, Roose SP, Fieo R, Liu X, Rantanen T, Sneed JR, Rutherford BR, Devanand DP, Avlund K. Frailty and depression in older adults: a high-risk clinical population. *Am J Geriatr Psychiatry*. 2014; 22:1083–95. <https://doi.org/10.1016/j.jagp.2013.04.010>
PMID:23973252
26. Zhou B, Chen K, Wang J, Wang H, Zhang S, Zheng W. Quality of life and related factors in the older rural and urban Chinese populations in Zhejiang province. *Journal of Applied Gerontology*. 2011; 30:199–225. <https://doi.org/10.1177/0733464810361346>
27. Cesari M, Vellas B, Hsu FC, Newman AB, Doss H, King AC, Manini TM, Church T, Gill TM, Miller ME, Pahor M, and LIFE Study Group. A physical activity intervention to treat the frailty syndrome in older persons—results from the LIFE-P study. *J Gerontol A Biol Sci Med Sci*. 2015; 70:216–22. <https://doi.org/10.1093/gerona/glu099>
PMID:25387728
28. Klimova B, Valis M, Kuca K. Cognitive decline in normal aging and its prevention: a review on non-pharmacological lifestyle strategies. *Clin Interv Aging*. 2017; 12:903–10. <https://doi.org/10.2147/CIA.S132963> PMID:28579767
29. Ma L, Li Z, Tang Z, Sun F, Diao L, Li J, He Y, Dong B, Li Y. Prevalence and socio-demographic characteristics of disability in older adults in China: findings from China comprehensive geriatric assessment study. *Arch Gerontol Geriatr*. 2017; 73:199–203. <https://doi.org/10.1016/j.archger.2017.07.029>
PMID:28822922
30. Cui Y, Gong D, Yang B, Chen H, Tu MH, Zhang C, Li H, Liang N, Jiang L, Chang P. Making the CARE comprehensive geriatric assessment as the core of a total mobile long term care support system in China. *Stud Health Technol Inform*. 2018; 247:770–74. PMID:29678065
31. Zhao Y, Hu Y, Smith JP, Strauss J, Yang G. Cohort profile: the China health and retirement longitudinal study (CHARLS). *Int J Epidemiol*. 2014; 43:61–68. <https://doi.org/10.1093/ije/dys203> PMID:23243115
32. Wu C, Smit E, Xue QL, Odden MC. Prevalence and correlates of frailty among community-dwelling Chinese older adults: the China health and retirement longitudinal study. *J Gerontol A Biol Sci Med Sci*. 2017; 73:102–08. <https://doi.org/10.1093/gerona/glx098>
PMID:28525586
33. Li J, Cacchione PZ, Hodgson N, Riegel B, Keenan BT, Scharf MT, Richards KC, Gooneratne NS. Afternoon napping and cognition in Chinese older adults: findings from the China health and retirement longitudinal study baseline assessment. *J Am Geriatr Soc*. 2017; 65:373–80. <https://doi.org/10.1111/jgs.14368> PMID:27995615
34. Huang W, Zhou Y. Effects of education on cognition at older ages: evidence from China’s great famine. *Soc Sci Med*. 2013; 98:54–62. <https://doi.org/10.1016/j.socscimed.2013.08.021>
PMID:24331882
35. Lei X, Smith JP, Sun X, Zhao Y. Gender differences in cognition in China and reasons for change over time: evidence from CHARLS. *J Econ Ageing*. 2014; 4:46–55. <https://doi.org/10.1016/j.jeoa.2013.11.001>
PMID:25530942
36. Pangman VC, Sloan J, Guse L. An examination of psychometric properties of the mini-mental state examination and the standardized mini-mental state examination: implications for clinical practice. *Appl Nurs Res*. 2000; 13:209–13. <https://doi.org/10.1053/apnr.2000.9231>
PMID:11078787
37. McArdle JJ, Fisher GG, Kadlec KM. Latent variable analyses of age trends of cognition in the health and retirement study, 1992–2004. *Psychol Aging*. 2007; 22:525–45. <https://doi.org/10.1037/0882-7974.22.3.525>
PMID:17874952
38. Mathuranath PS, Nestor PJ, Berrios GE, Rakowicz W, Hodges JR. A brief cognitive test battery to differentiate Alzheimer’s disease and frontotemporal dementia. *Neurology*. 2000; 55:1613–20. <https://doi.org/10.1212/01.wnl.0000434309.85312.19>
PMID:11113213
39. Jak AJ, Bondi MW, Delano-Wood L, Wierenga C, Corey-Bloom J, Salmon DP, Delis DC. Quantification of five neuropsychological approaches to defining mild cognitive impairment. *Am J Geriatr Psychiatry*. 2009; 17:368–75. <https://doi.org/10.1097/JGP.0b013e31819431d5>
PMID:19390294
40. Fried LP, Tangen CM, Walston J, Newman AB, Hirsch C, Gottdiener J, Seeman T, Tracy R, Kop WJ, Burke G, McBurnie MA, and Cardiovascular Health Study Collaborative Research Group. Frailty in older adults: evidence for a phenotype. *J Gerontol A Biol Sci Med Sci*. 2001; 56:M146–56. <https://doi.org/10.1093/gerona/56.3.m146>
PMID:11253156
41. Chu NM, Bandeen-Roche K, Tian J, Kasper JD, Gross AL, Carlson MC, Xue QL. Hierarchical Development of Frailty and Cognitive Impairment: Clues Into Etiological

- Pathways. *J Gerontol A Biol Sci Med Sci*. 2019; 74:1761–1770.
<https://doi.org/10.1093/gerona/glz134>
PMID:[31120105](https://pubmed.ncbi.nlm.nih.gov/31120105/)
42. Shimada H, Doi T, Lee S, Makizako H, Chen LK, Arai H. Cognitive Frailty Predicts Incident Dementia among Community-Dwelling Older People. *J Clin Med*. 2018; 7:250.
<https://doi.org/10.3390/jcm7090250> PMID:[30200236](https://pubmed.ncbi.nlm.nih.gov/30200236/)
43. Sargent L, Brown R. Assessing the current state of cognitive frailty: measurement properties. *J Nutr Health Aging*. 2017; 21:152–60.
<https://doi.org/10.1007/s12603-016-0735-9>
PMID:[28112769](https://pubmed.ncbi.nlm.nih.gov/28112769/)
44. Hardy SE, Dubin JA, Holford TR, Gill TM. Transitions between states of disability and independence among older persons. *Am J Epidemiol*. 2005; 161:575–84.
<https://doi.org/10.1093/aje/kwi083> PMID:[15746474](https://pubmed.ncbi.nlm.nih.gov/15746474/)
45. Fine JP, Gray RJ. A Proportional Hazards Model for the Subdistribution of a Competing Risk. *Journal of the American Statistical Association*. 1999; 94:496–509.
<https://doi.org/10.2307/2670170>

SUPPLEMENTARY MATERIALS

Supplementary Tables

Supplementary Table 1. Summary of the compatible studies mentioned in our study.

First author	Year	Country	N	Age	study types	Frailty criteria	Cognition	BADL	IADL	Mobility disability	Hospitalization	Mortality
Aliberti MJ[7]	2019	USA	7338	≥65	6 years longitudinal	Fried PF	HRS tests	↑	/	/	/	↑
Avila-Funes JA[8]	2009	French	6030	65-95	4 years longitudinal	Fried PF	MMSE	↑	↑	↑	↑	→
Feng L[9]	2017	Singapore	2375	≥55	6 years longitudinal	Fried PF	MMSE	↑	↑	/	→	↑
John PDS[10]	2017	Canada	1751	≥65	5 years longitudinal	FI	MMSE	/	/	/	/	↑
Cano C[11]	2012	USA	1815	≥67	10 years longitudinal	Fried PF	MMSE	/	/	/	/	→
Solfrizzi V[12]	2017	Italian	2373	65-84	4 years longitudinal	Fried PF	MMSE	/	/	/	/	↑
Solfrizzi V[13]	2017	Italian	2150	65-84	7 years longitudinal	Fried PF	MMSE	/	/	/	/	↑
Yu R[14]	2018	China	3491	≥65	12 years longitudinal	Fried PF	MMSE	/	/	↑	↑	↑
Shimada H[15]	2016	Japan	8864	≥65	cross-sectional	Fried PF	NCGG-FAT	/	↑	/	/	/
Z Liu[23]	2018	USA	754	≥70	11 years longitudinal	Fried PF	MMSE	↑	↑	↑	→	
Roppolo M[S1]	2016	Italian	594	≥65	cross-sectional	Fried PF	MMSE	/	/	↑	/	/

BADL, basic activity of daily living; IADL, instrumental activity of daily living; Fried PF, Fried physical frailty; FI, frailty index; MMSE, Mini-mental State Examination; NCGG-FAT, National Center for Geriatrics and Gerontology-Functional Assessment Tool; MCA, Montreal Cognitive Assessment.

↑, refers to cognitive frailty was associated with increased risk of adverse health outcomes; →, refers to no statistically significant association; /, refers to do not mention.

[7–15, 23] keep the same citation with Manuscript Text.

Supplementary Table 2. Mutually controlled associations of cognitive impairment and physical frailty with disability (BADL, IADL, and mobility), hospitalization, and death in full sample, CHARLS 2011-2015.

	BADL disability	IADL disability	Mobility disability	Hospitalization	Death
Individual effect/ No.	3341	3226	1685	3776	5113
Cognition					
Normal cognition	Ref.	Ref.	Ref.	Ref.	Ref.
Cognitive impairment	1.39 (1.13–1.71)	2.10 (1.69–2.61)	1.53 (1.06–2.21)	1.13 (0.93–1.38)	1.53 (1.16–2.01)
Physical frailty					
Nonfrail	Ref.	Ref.	Ref.	Ref.	Ref.
Frail	2.34 (1.63–3.37)	2.39 (1.60–3.57)	1.52 (0.64–3.60)	1.36 (1.01–1.82)	2.01 (1.48–2.72)

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activities of daily living; IADL, instrumental activities of daily living.

As described in Methods, we ran a logistic regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (i.e., N=3341 participants without BADL disability at baseline). Odds ratios (ORs) and corresponding 95% confidence interval (CI) are presented. The model adjusted for age, gender, cognition, and physical frailty.

Supplementary Table 3. Associations of multimorbidity with disability (BADL, IADL, and mobility), hospitalization, and death in full sample, CHARLS 2011-2015.

	BADL disability	IADL disability	Mobility disability	Hospitalization	Death
Multimorbidity /No.	3341	3226	1685	3776	5113
No	Ref.	Ref.	Ref.	Ref.	Ref.
Yes	2.10 (1.81–2.44)	1.92 (1.65–2.23)	1.95 (1.54–2.47)	1.81 (1.57–2.09)	1.26 (1.03–1.54)

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activities of daily living; IADL, instrumental activities of daily living.

As described in Methods, we ran a logistic regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (i.e., N=3341 participants without BADL disability at baseline). Odds ratios (ORs) and corresponding 95% confidence interval (CI) are presented. The model adjusted for age and gender.

Supplementary Table 4. Associations of cognitive impairment and frailty with disability, hospitalization, and death in sample after removing those with memory problem, CHARLS 2011-2015.

	BADL disability	IADL disability	Mobility disability	Hospitalization	Death
Individual effect					
Cognition					
Normal cognition	Ref.	Ref.	Ref.	Ref.	Ref.
Cognitive impairment	1.39 (1.13–1.71)	2.11 (1.69–2.63)	1.54 (1.07–2.23)	1.11 (0.91–1.36)	1.57 (1.19–2.07)
Physical frailty					
Nonfrail	Ref.	Ref.	Ref.	Ref.	Ref.
Frail	2.34 (1.61–3.38)	2.48 (1.66–3.72)	1.46 (0.61–3.49)	1.38 (1.03–1.87)	2.05 (1.50–2.80)
Combined effect					
Normal cognition & Nonfrail	Ref.	Ref.	Ref.	Ref.	Ref.
Cognitive impairment & Nonfrail	1.40 (1.13–1.73)	2.12 (1.69–2.65)	1.52 (1.05–2.20)	1.15 (0.93–1.41)	1.41 (1.04–1.92)
Normal cognition & Frail	2.48 (1.64–3.73)	2.54 (1.65–3.92)	1.33 (0.54–3.22)	1.53 (1.09–2.15)	1.80 (1.25–2.60)
Cognitive impairment & Frail	2.32 (1.01–5.32)	4.13 (1.40–12.15)	1*	1.06 (0.56–2.02)	3.76 (2.14–6.61)

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activities of daily living; IADL, instrumental activities of daily living.

As described in Methods, we ran a logistic regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (i.e., N=3341 participants without BADL disability at baseline). Odds ratios (ORs) and corresponding 95% confidence interval (CI) are presented. The model adjusted for age and gender.

*Since all participants in this subgroup reported the health outcome over the follow-up period, we assigned 1 to them.

Supplementary Table 5. Associations of cognitive impairment and frailty with disability and hospitalization accounting for the competing risk of death, CHARLS 2011-2015

	BADL disability	IADL disability	Mobility disability	Hospitalization
Individual effect				
Cognition				
Normal cognition	Ref.	Ref.	Ref.	Ref.
Cognitive impairment	1.27 (1.10–1.47)	1.63 (1.42–1.86)	1.17 (1.02–1.34)	1.06 (0.86–1.32)
Physical frailty				
Nonfrail	Ref.	Ref.	Ref.	Ref.
Frail	1.57 (1.28–1.93)	1.82 (1.48–2.24)	1.17 (0.89–1.55)	1.40 (1.06–1.87)
Combined effect				
Normal cognition & Nonfrail	Ref.	Ref.	Ref.	Ref.
Cognitive impairment & Nonfrail	1.31 (1.13–1.52)	1.65 (1.44–1.90)	1.17 (1.02–1.34)	1.09 (0.87–1.36)

Normal cognition & Frail	1.75 (1.40–2.19)	1.92 (1.53–2.40)	1.16 (0.84–1.60)	1.49 (1.09–2.04)
Cognitive impairment & Frail	1.24 (0.77–2.01)	2.16 (1.32–3.52)	1.46 (1.10–1.94)	1.18 (0.62–2.24)

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activities of daily living; IADL, instrumental activities of daily living.

As described in Methods, we ran a competing risk Cox regression model for each health outcome (e.g., BADL disability) in participants who did not have exposure for that outcome at baseline (i.e., N=3341 participants without BADL disability at baseline). Hazard ratios (HRs) and corresponding 95% confidence interval (CI) are presented. The model adjusted for age and gender.

*Since all participants in this subgroup reported the health outcome over the follow-up period, we assigned 1 to them.

Supplementary Table 6. Cross-sectional associations of cognitive impairment and frailty with disability and hospitalization at baseline, CHARLS 2011.

	BADL disability	IADL disability	Mobility disability	Hospitalization
No.	1130	1289	3199	602
Individual effect				
Cognition				
Normal cognition	Ref.	Ref.	Ref.	Ref.
Cognitive impairment	1.88 (1.58–2.22)	2.51 (2.14–2.96)	1.67 (1.41–1.99)	0.96 (0.75–1.22)
Physical frailty				
Nonfrail	Ref.	Ref.	Ref.	Ref.
Frail	3.83 (3.04–4.82)	4.58 (3.62–5.79)	5.03 (3.51–7.20)	1.61 (1.20–2.17)
Combined effect				
Normal cognition & Nonfrail	Ref.	Ref.	Ref.	Ref.
Cognitive impairment & Nonfrail	1.84 (1.53–2.21)	2.46 (2.07–2.92)	1.59 (1.33–1.90)	0.93 (0.72–1.21)
Normal cognition & Frail	3.90 (3.00–5.07)	4.53 (3.47–5.91)	4.57 (3.11–6.72)	1.60 (1.14–2.25)
Cognitive impairment & Frail	5.57 (3.55–8.74)	9.34 (5.66–15.42)	11.14 (4.06–30.57)	1.57 (0.87–2.82)

CHARLS, the China Health and Retirement Longitudinal Study; BADL, basic activities of daily living; IADL, instrumental activities of daily living.

As described in Methods, we ran a logistic regression model for each health outcome (e.g., BADL disability) at baseline. Odds ratios (ORs) and corresponding 95% confidence interval (CI) are presented. All models were adjusted for age and gender.

Supplementary References

1. Roppolo M, Mulasso A, Rabaglietti E. Cognitive Frailty in Italian Community-Dwelling Older Adults: Prevalence Rate and Its Association with Disability. *J Nutr Health Aging*. 2017; 21:631–636. <https://doi.org/10.1007/s12603-016-0828-5> PMID:28537326