

Original Article

Frailty Syndrome among oldest old Individuals, aged ≥ 80 years: Prevalence & Correlates

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Abstract

Objectives: Objectives were to study prevalence of frailty among Indian oldest old population, and to detect its correlates. **Methods:** A cross sectional community based study was done including 200 healthy participants aged ≥ 80 years, randomly sampled from Hyderabad city in India. They completed an administered questionnaire and physical function tests including SPPB, grip strength. Cognitive function was assessed using MMSE and depression using GDS. Blood pressure, haemoglobin, and fasting blood sugar were measured for all participants. Frailty was defined using Fried phenotype criteria. Logistic regression was done to identify independently associated correlates. **Results:** The prevalence of frailty syndrome was 83.4% in our study population. Frailty among men was 80.3% and among women was 84.7%, and it increased with increasing age. The independent correlates which increased the odds of frailty were poor physical performance (SPPB) (OR: 4.21; 95%CI: 1.12-15.83), depression (OR: 3.35; 95%CI: 1.29-8.73), chronic joint pains (OR: 4.90; 95%CI: 1.97-12.18) and COPD (OR: 3.01; 95%CI: 1.03-8.78), while hypertension showed inverse association (OR: 0.33; 95%CI: 0.11-0.94). **Conclusion:** The prevalence of frailty among the oldest old is very high. Geriatric medicine protocols must include routine screening for frailty, while also including early detection of poor physical performance, depression, COPD and osteoarthritis.

Keywords: COPD, Depression, Frailty, Hypertension, Oldest old

Introduction

The number of older people, aged ≥ 65 years of age is expected to rise up to 1.5 billion by the year 2050 from current 703 million in 2019 in the world. The largest increase is expected to occur in Eastern and South-Eastern Asia, from 261 million (2019) to 573 million by 2050¹. Global increase of oldest old persons, aged ≥ 80 years of age, will be three times from 137 million in 2017 to 425 million in 2050. India will be the second largest contributor of oldest old population to the world¹. Hence it is crucial to study health problems and correlated factors of older population of India.

Frailty syndrome, a clinical geriatric syndrome, is associated with high risk of adverse health outcomes in older age² like function disabilities, falls, mobility disabilities, frequent hospitalization, cardiovascular diseases, diabetes, and activities of daily living (ADL) as reported from developed countries³. Mortality is reportedly high in frail elders compared to non-frail elders⁴. Theoretically frailty

is defined as a clinically recognizable state of increased vulnerability, resulting from aging-associated decline in reserve and function across multiple physiologic systems such that the ability to cope with every day or acute stressors is compromised⁵. In 2001, Fried and colleagues have defined operational definition of frailty⁶. Age-related changes to various physiological systems, particularly the

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neuromuscular, neuroendocrine and immunological systems, are fundamental to the development of frailty⁷. Frailty has been considered synonymous with disability, comorbidity, and other characteristics, but it is recognized that it may have a biologic basis and be a distinct clinical syndrome⁸. There are no proven pharmacological interventions yet for treatment of frailty⁹. Only some evidence of improvements have been reported from physical activity intervention studies, but more research is required to establish preventive and therapeutic interventions. It is noteworthy that the lack of a universal definition of frailty, underscores the difficulty in synthesizing the intervention literature for frail adults.

Prevalence of frailty is reported as 7% to 12% in USA in ≥ 65 year old and 25% in ≥ 85 year old age¹; African American 13%⁶; Mexican American 7.8%⁴; wide variations have been observed in European countries from 5.8% Switzerland to 27% in Spain^{10,11} high in southern than northern European countries after adjusting to age and sex; from 5 large Latin American and Caribbean cities the prevalence of frailty was reported as 30% to 48%¹². All the studies reported prevalence was more in women than men. These wide variations in the geographic prevalence speculates that there may be difference in socio-cultural characteristics influencing frailty¹⁰.

The data from developing countries are sparse on frailty and its correlates. The dynamics of health changes more rapidly in older ages of life. To our knowledge no population based studies exploring frailty syndrome is available, focussed on oldest old population aged ≥ 80 years from India. Hence we explored prevalence of frailty syndrome and correlated factors in this analysis. Evaluating the causes and natural history is therefore critical for identifying newer avenues for frailty and treatment.

Method

We enrolled 200 oldest old community-dwelling elders, 76 (38%) men and 124 (62%) women who completed 80 years of age or more, residing in south central part of India, Hyderabad city, Telangana state, in 2017. This cross-sectional research was designed to study frailty syndrome in oldest old urban population of south India. Ethical approval was obtained by institutional review board (IRB) of Medciti Institute of Medical Sciences, Hyderabad and Indian Council of Medical Research (ICMR), Delhi. The participants were enrolled from twelve residential townships (population ranging from 1000-6000) of Hyderabad city. Data was collected by trained investigator by home visits. Households having at least one age eligible person were enlisted and randomised with random number generator software. All age eligible persons in the selected households were contacted to participate in the study. Total of 218 individuals were contacted to reach required number of 200. The participation rate was 91.7%. Inclusion criteria were: Indian nationality, age >80 years, living in an urban area of Hyderabad city, possessing cognitive ability to understand the investigator's

instructions, and consenting to participate; persons with terminal illness or psychotic conditions were excluded from the study.

We collected data using structured questionnaire and forms prepared by adapting questions and measurement protocols from leading international studies: Mobility and Independent Living in Elders Study¹³ and the Lifestyle Interventions and Independence for Elders¹⁴ pilot study, WHO Study of Global AGEing and Adult Health¹⁵. Our questionnaires included information on age, sex, marital status, education, self-reported health status, medical history, physical functions, chronic diseases, medication inventory and functional disabilities. Other objective measurements included blood pressure, anthropometry (height, weight, waste and hip circumferences), grip strength, cognitive function, depression, haemoglobin and blood sugar.

Definitions and Measurements

Frailty Syndrome: Frailty syndrome was measured by Linda Fried's Frailty Criteria^{6,8} which includes examining 5 phenotypic criteria: low grip strength, low energy, slowed walking speed, low physical activity, and unintentional weight loss. Participants meeting 3 conditions of the 5 phenotypic criteria indicating compromised energetics were categorised as frail participant.

Cognitive function: Cognitive function was assessed by Mini Mental State Examination scale (MMSE)¹⁶. MMSE tests orientation, registration, attention and calculation, recall, language and praxis. A hindi (Indian language) version of this tool has been tested earlier in the Indian setting and validated. Single cut-off score <24 on MMSE was defined as cognitively impaired.

Activities of daily living (ADLs) constituted of 6 activities that included walking across the small room, bathing, grooming, dressing eating, getting out of bed and using toilet¹⁷. If any participant couldn't do even 1 activity out of above 6 then he was considered having poor activities of daily living.

Depression: The depression was assessed using 15 item Geriatric Depression Scale (GDS - 15)¹⁸. Of total score from 0 - 15, if participant gets score of >5 he was considered to be suffering from depression.

Blood pressure: Resting blood pressure was measured thrice with a 1 minute interval using an electronic sphygmomanometer (OMRON HEM 7120, OMRON HEALTHCARE Co., Ltd., Japan). Systolic and diastolic blood pressures were recorded on relaxed calm participants in the sitting position with their elbows raised at the level of their heart. They were instructed to abstain from eating, drinking alcohol or caffeinated drinks or exercise at least for 30 min before blood pressure measurement. The average of last two readings were used to define systolic and diastolic blood pressures¹⁹.

Grip strength: Hand grip strength was measured by JAMAR dynamometer using standard protocol. An

Characteristics	Women (N=124) (% & mean±sd)	Men (N=76) (% & mean±sd)	P value
Age (mean±sd)	83.21±3.77	83.69±4.01	0.40
Living single (%)	78.9	21.1	<0.001
No schooling (%)	69.4	30.6	0.03
Basal Metabolic Index (BMI) (mean)	23.25±4.76	22.31±4.31	0.17
Not working (%)	98.4	98.7	0.70
Depression (%)	68.9	38.1	0.02
Cognitive Impairment (%)	67.9	32.1	0.01

Table 1. Characteristics of the study population.

average of the last two readings out of three on the dominant hand was considered as the participant's grip strength²⁰. It was measured in kgs and expressed in mean and standard deviation.

Short Physical Performance Battery (SPPB): The Short physical performance battery (SPPB) consisting of a 4 m walk, repeated chair stands, and three hierarchical standing balance tests, was performed. Each of the three performance measures was assigned a categorical score ranging from 0 to 4, with 4 indicating the highest level of performance and 0 the inability to complete the test. A summary score ranging from 0 (worst performers) to 12 (best performers) was calculated by adding Gait speed, chair stands, and balance scores²¹.

Co-morbidities: Asthma, chronic obstructive pulmonary disease, cardiovascular disease, renal impairment, hypertension, diabetes, osteoarthritis, functional disabilities, vision and hearing were self-reported in the interview. We also measured biomarkers: haemoglobin and blood sugar.

Statistical analysis

Data was analyzed using SPSS, version 21 software (SPSS Inc., Chicago, IL, USA). Comparison of characteristics and correlates among frail individuals and non-frail individuals were done by using chi-square test for categorical variables and t-test for continuous variables. The variables those reported statistical significance (P value <0.05) in the univariate analysis were selected for multivariable model. We selected best fit model and checked for collinearity among the variables before running them in the logistic regression model. Specifically we examined Poor vision, osteoarthritis, hypertension, COPD, Body aches and pains, chronic joint pains, pain on standing/sitting down, depression, poor physical performances (SPPB), cognitive and impairment in the model, adjusting for age and sex. We excluded low physical activity, asthma and grip strength variables from the model due to high collinearity. We performed backward elimination logistic regression testing the deletion of each variable using a

chosen model comparison criterion, deleting the variable that improves the model the most and repeating this process until no further improvement was possible, to get final set of the independent predictor variables. Variables with $P < 0.05$ were retained in the final model. Results of logistic regression were expressed in odds ratios (OR) and 95% confidence intervals (95% CI). For continuous variables in logistic regression we expressed the OR per 1 standard deviation (SD) increase.

Results

Characteristics of Study population

In our study population 38.2% were men and 61.8% were women. The characteristics of the study population are described in Table 1. The mean ages and BMI of women and men were similar. A significantly higher proportion of women were living single, had no schooling, suffered with depression and impaired cognition, compared with men ($p < 0.05$).

Prevalence of Frailty syndrome

The prevalence of frailty syndrome in our study population was 83.4%, 80.3% of men and 84.7% of the women were frail ($p = 0.26$). The prevalence of frailty increased with increasing age and 100% were frail above the age of 95 years.

Correlates of frailty

The frail individuals were older, shorter, lighter, having a lower BMI than their non-frail counterparts, but the differences were not significant (Table 2). A significantly greater proportion of those having frailty had poor vision, osteoarthritis, hypertension, COPD, body pains, chronic joint pains, asthma, low physical activity compared with non-frail ($p < 0.05$). The frail group had more depression, low grip strength, poor SPPB, impaired cognitive function compared with the no frailty group ($p < 0.01$) (Table 2).

Upon logistic regression, the independently associated correlates of frailty were poor SPPB (OR: 4.21; 95%CI: 1.12-

Characteristics & Correlates	Non – Frail individuals (N=34) (% & mean±sd)	Frail individuals (N=166) (% & mean±sd)	Total (N=200) (% & mean±sd)	P value
Demographic				
Age (years) (mean)*	82.69±3.46	83.46±4.01	83.34±3.93	0.31
Height (cm) (mean)*	156.72±11.19	153.96±9.56	154.35±9.82	0.71
Weight (kg) (mean)*	60.07±4.77	56.63±35.76	57.12±33.59	0.62
BMI (kg/m ²) (mean)*	24.19±3.99	22.67±4.67	22.88±4.60	0.11
Marital status (single) (%)	54.5	63.9	62.3	0.20
No education (%)	39.4	52.1	50.0	0.12
Medical history & Chronic diseases				
Poor self-reported health	29.4	23.5	24.5	0.29
Smoking (%)	34.4	35.5	35.0	0.50
Alcohol (%)	29.4	30.7	30.5	0.54
Poor Vision (%)	47.1	67.5	64.0	0.02
Osteoarthritis (%)	26.5	46.4	43.0	0.02
Stroke (%)	11.8	3.6	5.0	0.06
Hypertension (%)	79.4	61.4	64.5	0.03
Cardiovascular disease (%)	85.3	90.4	89.5	0.27
Chronic Obstructive Pulmonary Diseases (COPD) (%)	17.6	36.7	33.5	0.02
Diabetes (%)	20.6	21.5	21.3	0.55
Body aches and Pains (muscular) (%)	60.6	80.7	77.4	0.03
Low physical activity (no exercise/no sports) (%)	70.6	93.4	89.5	0.001
Asthma (%)	11.8	26.5	24.0	0.04
Falls (%)	26.5	38.8	36.7	0.12
Chronic Joint pains (%)	47.1	81.3	75.5	<0.001
Pain on Long standing (%)	73.5	84.3	82.5	0.10
Pain on standing or sitting down (%)	58.8	77.7	74.5	0.02
Objective measurements				
Depression: >5 (GDS-15 point scale) (%)	20.6	58.4	52.0	<0.001
Grip strength - kg (mean)*	7.56±10.33	4.54±6.45	5.06±7.31	0.02
Low Grip strength – kg (%)	70.6	98.8	94.0	<0.001
Poor Short Physical Performance Battery (score ≤ 9) (%)	79.4	97.0	94.0	0.001
Impaired Cognitive function (MMSE <23) (%)	47.1	69.9	66.0	0.01
Systolic Blood pressure (mmHg)(mean)*	145.68±24.07	142.52±25.05	143.05±24.85	0.52
Diastolic blood pressure (mmHg)(mean)*	84.41±14.76	82.21±12.21	82.15±12.61	0.30

Table 2. Characteristics and Correlates of Frail and Non – Frail individuals.

15.83), depression (OR: 3.35; 95%CI: 1.29-8.73), chronic joint pains (OR: 4.90; 95%CI: 1.97-12.18) and COPD (OR: 3.01; 95%CI: 1.03- 8.78). Hypertension was negatively associated (OR: 0.33; 95%CI: 0.11-0.94). (Table. 3)

Discussion

We found a prevalence of 83.4% frailty syndrome in our oldest old population; 80% among men and 85% among women. The global prevalence of frailty among

Correlates	Odds Ratio (OR)	95% Confidence Interval (95% CI)	
		Lower	Upper
Short Physical Performance Battery (SPPB)***	4.21	1.12	15.83
Depression (%)***	3.35	1.29	8.73
Chronic Joint pains (%)***	4.90	1.97	12.18
Hypertension (%)**	0.33	0.11	0.94
Chronic Obstructive Pulmonary Disease (COPD)(%)**	3.01	1.03	8.78

*Backward stepwise logistic regression: **p<0.05; ***p<0.01.*

Table 3. Logistic regression predicting odds of having frailty by risk factors.

octogenarians is reported much lower at 23%²². Overall, the few independent studies on community dwelling oldest old reported lower prevalence of frailty than ours; 20.5% in Spain²³, 28% in California (among ages 90+ years)²⁴, 19.5% in Jerusalem²⁵, 21% and 34% (among 90-94 years and 95 years +) in Ireland²⁶, 34.9% in Japan²⁷, and 39.8% in Brazil²⁸. Most of these studies used the frailty phenotype criteria. Most studies on frailty on community dwelling elderly population aged 55 years and older, using varied frailty measurement criteria, reported prevalence rates ranging from 9.9% in developed countries²⁹ to 5.4-44% in developing countries³⁰. A study from Brazil reported higher frailty, 65.2% among individuals aged 80 years and older³¹.

We attribute our high frailty prevalence to higher levels of low education, low physical activity and lower working status of our population in addition to poor nutrition. A recent study from Thanjavur district of south India also showed that lower education, lower physical activity, and lower working status, higher age increases the odds of frailty in older individuals⁷¹. Although their study population was younger than ours i.e. ≥ 60 years of age, but they also reported higher prevalence of frailty compared to age matched population from other parts of the world. Other studies also reported lower levels of physical activity and working status in our older population^{16,21}. Our high prevalence of frailty may also be explained by varying lifestyles, health statuses, demographic and socioeconomic characteristics across countries as well as lower life expectancy in India.

The prevalence of frailty among women and men was alike in our study. Some others have mirrored this finding showing no difference in frailty prevalence by gender^{23,32}. Global literature, however abounds with findings of women being more commonly frail than men^{33,34}. Alvarado³³ and colleagues used a life course design to explain that women had more likelihood of frailty than men due to lack of schooling, a manual occupation, being a housewife, and perceived economic hardship later in life. They further discussed that social and chronic conditions did not totally explain the higher odds of frailty in women, thereby

suggesting the presence of unmeasured underlying factors such as differences in biological factors, muscle strength, and other life-course exposures including low food intake and poor nutrition because of lack of social support and networks, low rates of physical activity related to lack of physical activity at younger ages, poorer perception of health, lack of self-sufficiency, and greater exposure to unsafe neighborhoods. Mitnitski et al.³⁴ used a novel theory of accumulated deficits to explain the increasing risk of frailty in community dwelling elders wherein a frailty index was constructed as a proportion of all potential deficits (symptoms, signs, laboratory abnormalities, disabilities) expressed in a given individual. Their model also showed that at given age, women were frailer. Since women live longer and generally have a larger number of comorbidities³⁵ a higher prevalence of frailty is expected among them.

Among functional parameters assessed, SPPB was independently associated with frailty in our population, posing a four folds increase in risk. Frail individuals have lower physical and muscular performances. Previous studies have shown associations between frailty and lower levels of lower limb muscle strength³⁶. Moreover, SPPB is commonly used as an indicator of frailty³⁷; whereby the high prevalence of frailty in our population detected using frailty phenotype criteria explained and justified a strong association with poor SPPB. We could not find any studies that showed this association specifically among the oldest old. Fried et al. theorized the loss of the muscle mass playing an important aetiologic role in the frailty process of elderly subjects⁹. Sarcopenia is frequently associated with poor endurance, physical inactivity and slow gait speed^{38,39}. It is plausible that the oldest old, as a consequence of ageing, have greater sarcopenia compared with the younger elders, thereby having poorer SPPB scores and higher frailty.

Independently associated clinical parameters included depression, joint pains and COPD. Depression was associated with a 3 folds risk for frailty. Previous literature has also consistently reported this relationship between frailty and depression^{28,40}. It has been earlier suggested that this association might be linked to characteristics present in

both conditions, such as inactivity, weight loss, low level of physical activity and exhaustion⁴⁰. Others have suggested a complex and two-way causal nature between frailty and depression, that needs further exploration. While on one hand the presence of depressive symptoms may constitute a risk factor for this syndrome with changes in behavior, activity level or social commitment contributing to the decline of one's functional state and frailty, often referred to as 'psychosocial frailty'⁴¹. Depressive symptoms on the other hand, may represent the early manifestation of frailty, worsening one's mood and depression due to the syndrome⁴². Researchers have also suggested a considerable overlap between these two conditions^{28,43}. A recent systematic review on co-occurrence of depressive symptoms and frailty among elderly aged 55 years and older, indicated a prospective relationship between depressive symptomatology and increased risk of incident frailty⁴⁴.

Chronic joint pains were independently associated 4 folds with frailty in our study. Earlier studies on elderly corroborated with this finding of ours. The EPOSA study from six European countries on 2,455 individuals aged 65-85 years showed odds of frailty 2.96 (95 % CI:2.11-4.16) as high among OA individuals than those without. They further found the association to strengthen with increasing number of joints involved than with individual joints⁴⁵. In another independent study with more than 52% participants aged 80 years and above, Coelho et al. showed that pain predicted physical frailty; through mechanisms that were linked to mobility limitations, fatigue, disturbed sleep and decreased nutritional intake⁴⁶. Some hypothesized that frailty is associated with physiological changes (in brain, immune system, endocrine system, skeletal system) which could alter pain perception or exacerbate pain in an older person⁴⁷. Data on chronic pain and frailty among community dwelling elderly are very limited^{48,49} and missing for 80 years and older population. These limited studies showed older people reporting pain to be statistically more likely to be frail. The direction of the relationship however, couldn't be affirmed due to the lack of prospective data. Additionally, these conditions may overlap considerably, indicating that somatic complaints are symptoms of diseases associated with the frailty syndrome⁵⁰.

Frailty in our population was also independently associated 3 folds with COPD. Of the very few studies to elaborate on similar association, the Rotterdam study found elderly participants (having mean age 74 years) with COPD showed a more than twofold increased prevalence of frailty (odds ratio 2.2, 95% CI: 1.34-3.54). A recently published meta-analysis on relationship between COPD and frailty among older adults including 27 studies found that patients with COPD had a two-fold increased odds of frailty (pooled OR: 1.97; 95% CI: 1.53-2.53). Three longitudinal studies included in the review suggested a bidirectional association between COPD and frailty⁵¹. This association is biologically plausible, since complex age related changes occur in the

ageing lung characterized by a decline in FEV1, a reduction in muscular strength and an increase in inflammatory cells in bronchial tissue; these changes are aggravated in frail individuals. Furthermore, the predilection for managing COPDs with steroids predisposes ageing individuals to steroid related multi morbidities and resultant frailty⁵².

Hypertension emerged as negatively associated with frailty upon logistic regression. The protective effects of hypertension on cardiovascular and all cause mortality has been shown earlier among the oldest old populations^{53,54}. Wong et al. also similarly noted that cross-sectional analysis could not fully exclude the possibility of reverse causation for hypertension and frailty⁵⁵, with many observational studies indicating weaker associations of blood pressure with CVD in older individuals compared with the middle ages. Our population was however, much older compared with these studies. These earlier findings and the HYVET study recommendations later⁵⁶ questioned the effect of antihypertensive treatment and the ideal blood pressure goal of treatment in the frail elderly⁵⁷. De Sousa et al., in their frailty prediction model for Brazilian oldest old, failed to find an association with cardiovascular diseases, inspite of CVD being the most commonly reported disease and antihypertensive being the predominant medications⁵⁸. The mechanisms for negative association of hypertension with frailty, or conversely, positive association of low blood pressure with frailty among the oldest old were well debated⁵⁹. Few studies proposed that older or frail elders were more likely to have chronic health conditions resulting in low blood pressure⁶⁰. Others explained this association with increased vascular stiffness resulting from atherosclerosis, arterial calcification, endothelial dysfunction, and smooth muscle cell fibrosis leading to isolated systolic hypertension, more pronounced on the frail oldest old⁶¹.

We showed that frailty was significantly associated with cognitive impairment upon univariate analysis. Studies on ageing and frailty have reported this association very often^{62,63}. In a comparative study, Marcucci et al. showed a 5 fold odds (OR=5.15, 95% CI=2.58-10.26) of mortality among the frail oldest old women with cognitive decline compared with the younger elderly (relatively healthy) without cognitive impairment⁶⁴. The characteristic structural and physiological changes in the brain associated with ageing, which place disproportionate stress on neurons with high metabolic demands, such as the hippocampal pyramidal neurons that are important mediators in pathophysiology of cognitive decline^{65,66}, explain this association. Additionally, several factors that are related to physical frailty also are related to cognitive impairment, including inflammatory markers, diabetes, congestive heart failure, and stroke. In our population, however, most of these factors did not show any association, barring pain on movement which underlines inflammation. Interestingly, stroke showed an inverse relationship with frailty that was nearly significant in this population. These inverse associations with cardiovascular

parameters (hypertension and stroke) lead us to hypothesize that cardiovascular factors may protect the oldest old Indians against frailty. Moreover, it may be assumed that frail elderly who have survived the age of 80 years may have several other compensatory mechanisms in place that mask the risk posed due to adverse cardiovascular events. This remains amenable to testing using more objective criteria and possibly biomarkers⁶⁷, since the phenotypic criteria (based on measures of mobility and motor performance) used in our study were not derived from stroke patients (who were originally excluded from the CHS).

We also found an association between frailty and low vision, which has been reported earlier by very few studies. One Chinese study reported low vision to be an independent predictor of frailty among elderly aged 60 years and older. They attributed the association to the increased risk of falls and fractures leading to frailty, consequent to visual impairment⁶⁸. Lang et al. described sensory impairment in relation to and as a manifestation of frailty earlier⁶⁹. This relationship may be independent due to ageing impacting both physical frailty and visual impairment consequent to degenerative conditions including cataract formation.

All the positive associations of risk factors with frailty in our study point to the chronic nature of these diseases which call for early detection and control. Additionally, it is worthwhile to note that most of these conditions are not treated appropriately, and elderly continue to survive in the absence of medical care. The draft National Program for the Health Care of the Elderly (NPHCE)⁷⁰ has attempted to identify some of these problems at grass root level but focused interventions on frailty seem missing. Developed countries, on the contrary have already identified frailty as a major problem, with guidelines for preventive interventions instituted. For instance, the revised contract for General Practitioners in the UK for 2017–18 includes advice for frailty screening of older people using electronic frailty questionnaire and advocates strategies for annual review of medication for frail older people⁵⁵. Such screening may include opportunities to review lifestyle advice and medication to optimise control of risk factors, CVD risk factors in particular, for prevention of disability and death in frail older people. Similarly, Arantes P et al.⁷² in their systematic review have compared and verified the effect of seven different types of interventions namely, 1) muscle strengthening; 2) exercises for muscle strengthening, balance, coordination, flexibility, reaction time and aerobic training; 3) functional training; 4) physical therapy; 5) at-home physical therapy; 6) environment adaptation and prescription of assistive device; 7) water exercise. de Labra et al.⁷³ also reported the benefits of exercise interventions in their systematic review that included studies on multi-component exercise interventions (aerobic and resistance training not coexisting in the intervention), physical comprehensive training, and exercises based on strength training, compared with a control group

receiving no treatment, maintaining their habitual lifestyle or using a home-based low level exercise program. The benefits reported in the review ranged from positive impact of exercise interventions on falls, enhancements in several measurements of mobility, enhanced balance, and functional ability, increases in muscle strength, improvements in body composition, and improvement in frailty. They however, concluded that though exercise interventions had demonstrated improvement in different outcome measurements in frail older adults, there were large differences between studies with regard to effect sizes, thereby inconclusive regarding an optimal program.

We recommend further studies of similar kind from different regions that can reflect the actual burden and the presumed effect of interventions for frailty prevention. We further recommend that the current geriatric medicine protocols include multidisciplinary- group-based rehabilitation interventions in order to reduce the progression and the impact of frailty in this population. The current geriatric medicine protocols of the country should be revisited to address these problems using active ageing and rehabilitation medicine for elderly⁷⁴⁻⁷⁶.

Strengths and limitations

The major strength of this study is the large number of community dwelling participants, aged 80 years and older used to estimate the prevalence of frailty, which provides a stable estimation of prevalence. This is by far, the largest study in India, and in South East Asia, to use the phenotypic definition of frailty to analyze prevalence in this age group, allowing further analysis according to age and sex specific groups. The sample of 200 participants from South India however, may not sufficiently reflect the burden in rural parts of the country, and surrounding states and countries with varying population characteristics and health services. We did not calculate required sample size to state the risk factors due to absence of comparative literature. A post hoc power analysis showed that the included number could detect the prevalence and correlates with sufficient power of more than 90%.

Conclusion

Prevalence of frailty was very high in our oldest old population. Frailty was similar in women and men in our population. The independent positively associated correlates of frailty-poor physical performance, depression, COPD and chronic joint pains were chronic health problems of the oldest old, which suggest the need for early detection and prevention measures in order to control frailty. The current geriatric medicine protocols of the country should be revisited to address these problems using a holistic approach.

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