INTRODUCTION

South Korea is one of the fastest aging societies worldwide; moreover, the estimated life expectancy of Korean people is the highest (in 2030) among people from developed countries. However, according to the 2018 Korean Geriatrics Fact Sheet, 89.5% of older Korean adults had one or more chronic diseases and the annual national medical expenses for older adults accounted for 39.9% of the total medical expenses.1 With these changes in population, there is an overarching need to identify vulnerable older adults at high risk for conditions such as multimorbidity, polypharmacy, low economic status, cognitive impairment, and functional decline; moreover, growing evidence suggests that frailty is a core geriatric syndrome that is highly correlated with human aging, multimorbidity, and disability.2-7 Frailty is defined as a state of decreased physiological reserve associated with human aging and is a strong predictor of adverse health outcomes such as falls, fractures, loss of independence, disability, institutionalization, and death.8,9 Therefore, screening for frailty in older populations is a crucial step in delivering tailored multifaceted care for highly heterogeneous populations.
Currently, the most commonly used frailty definitions in research are the frailty phenotype and frailty index. These tools are less feasible for use in mass populations as they are resource intensive, research oriented, and especially troublesome to adapt to policies of resource-limited community settings. As a screening measure, the Fatigue, Resistance, Ambulation, Illnesses, and Loss of weight (FRAIL) scale, a self-reported screening instrument for frailty developed by Morley et al., is a valid method for use in communities. This short questionnaire is easy to administer, requires minimal time and resources, and has comparable performance in screening frailty. It has been validated in various populations and is a useful screening tool for identifying frail older adults. The Korean version, the K-FRAIL scale, has been validated in the Korean population, especially in older adults in urban communities. The FRAIL scale has been shown to have predictive validity for mortality in various studies. However, other health outcomes such as disability or institutionalization require further investigation.

Thus, this study examined the association between geriatric conditions and health-related outcomes such as mortality, institutionalization, and disability using the five-item FRAIL scale in rural community-dwelling Korean older adults.

**MATERIALS AND METHODS**

**Study Population**

We analyzed the records of participants in the Aging Study of Pyeongchang Rural Area (ASPHA) cohort, a population-based, prospective cohort study originally established in 2014 that included community-dwelling adult residents of Pyeongchang-gun, Gangwon Province, South Korea who were aged 65 years and older. Detailed information on the original study design and measurements in the ASPHA has been published elsewhere. During annual assessments, the study population continuously increased with the expansion of the covered region of ASPHA among Pyeongchang-gun, with the inclusion and exclusion criteria remaining the same. Individuals were eligible for participation if they were (1) aged 65 years or older, (2) registered in the Community Healthcare Service, (3) ambulatory with or without an assistive device, (4) living at home, and (5) able to provide informed consent by themselves or their legal proxies. We excluded participants who were (1) living in nursing homes, (2) hospitalized, or (3) bedridden and receiving nursing home level care at home at the time of enrollment.

Between December 2014 and December 2017, 1,295 community-dwelling individuals were enrolled in the ASPHA cohort, which included over 90% of eligible people living in the study area. Among these older adults, we used records of 1,292 people who completed the K-FRAIL questionnaire. All participants underwent a comprehensive geriatric assessment every 12 months, while health-related outcomes, including death, institutionalization, and disability, were recorded every 3 months. The characteristics of the original ASPHA population were generally comparable to those of the general Korean rural population based on the Korea National Health and Nutrition Examination Survey. The study protocol of the ASPHA cohort was approved by the Institutional Review Board of the Asan Medical Center (No. 2015-0673). Written informed consent was obtained from all participants.

**Frailty Screening**

We used the K-FRAIL questionnaire as a screening measure for frailty status. The K-FRAIL is based on the original FRAIL scale, which has been validated in various populations and translated into Korean. This simple screening tool of five self-reported items can be assessed in 2–3 minutes with no requirement for special equipment or sophisticated training. The questionnaire comprises five items:

1. Fatigue (responses of “all of the time” or “most of the time” to the following question): “How much of the time during the past 4 weeks did you feel tired?”
2. Resistance (positive response to the question): “By yourself and not using aids, do you have any difficulty in walking up 10 steps without resting?”
3. Ambulation (positive response to the question): “By yourself and not using aids, do you have any difficulty in walking 300 m?”
4. Illnesses (five or more self-reported physician diagnoses of hypertension, diabetes, cancer excluding minor skin cancer, chronic lung disease, heart attack, congestive heart failure, angina, asthma, arthritis, stroke, or kidney disease), and
5. Loss of weight (> 5% in the past year).

We calculated the K-FRAIL score by combining the number of positive items and classified the participants as robust (0 points), prefail (1–2 points), or frail (3–5 points).

**Assessment of Geriatric Conditions**

Trained nurses performed comprehensive geriatric assessments (CGAs) to evaluate geriatric conditions. Multimorbidity was defined as having two or more of the 11 physician-diagnosed illnesses such as angina, arthritis, asthma, cancer, chronic lung disease, congestive heart failure, diabetes, hypertension, heart attack, kid-
ney disease, and stroke. Polypharmacy was defined as regularly taking five or more medications. Cognitive impairment was defined as a Korean version of the Mini-Mental State Examination score of < 24. Depressive mood was defined as a Korean version of the Center for Epidemiological Studies Depression scale score of > 20. Malnutrition risk was defined as a Mini-Nutritional Assessment-Short Form score of ≤ 11. To assess physical function, we measured handgrip strength (kg) using a dynamometer (T.K.K. 5401 Grip-D; Takei, Tokyo, Japan) and averaged two measurements of the dominant hand. We defined dysmobility as a slow gait (usual gait speed < 0.6 m/s from a timed 4-m walk).

We calculated the Short Physical Performance Battery (SPPB) score from three components: the ability to stand for up to 10 seconds with feet positioned in three ways (together side-by-side, semi-tandem, and tandem), the time to complete a 4-m walk, and the time to rise from a chair five times. The resultant SPPB scores ranged from 0 to 12, with higher scores indicating a higher level of function. Sarcopenia was defined according to the Asian Working Group for Sarcopenia as follows: decreased muscle mass (decreased handgrip strength of < 26 kg for men and < 18 kg for women) and decreased physical performance (slow gait of < 0.8 m/s). Disability was defined as requiring assistance in performing any of the seven activities of daily living (ADL; bathing, continence, dressing, eating, toileting, transferring, and washing face and hands) or 10 instrumental activities of daily living (IADL; food preparation, household chores, going out on a short distance, grooming, handling finances, laundry, managing own medications, shopping, transportation, and using a telephone).

Outcome Measurements
This study defined major adverse health outcomes as death, institutionalization, and incidence of disability during the 3-year follow-up period. We defined disability as the requirement for assistance from another person in performing any one of the ADL or IADL. Health-related quality of life was assessed during the follow-up period using the EuroQol-5D (EQ-5D). An EQ-SD score decline over 3 years of > 0.2 was considered clinically meaningful. To assess these outcome measures, all participants underwent structured interviews via phone calls every 3 months on institutionalization, death, and annualized follow-up assessments, including CGAs for ADL, IADL, and EQ-5D.

Statistical Analysis
To compare variables for baseline characteristics according to the frailty status derived from the K-FRAIL scale, we used Kruskal-Wallis and chi-square tests for continuous and discrete variables, respectively. To cross-sectionally assess the association between prevalent frailty and geriatric syndrome, we used logistic regression analysis with common geriatric syndromes as dependent variables and frailty status (robust, prefrail, and frail) as independent variables after adjusting for age and sex. For longitudinal outcome analysis, we used logistic regression analysis for health outcomes of mortality, long-term institutionalization, incidence of disability, and decline in EQ-5D score at 3 years from baseline, with frailty status (robust, prefrail, and frail) as dependent variables after adjusting age and sex. Mortality was further analyzed using Kaplan-Meier analysis, with log-rank tests and Cox proportional hazard regression analysis used to examine the statistical differences in survival according to the baseline frailty status. The restricted mean survival time was estimated as death at 40 months. Two-sided p-values < 0.05 indicated statistical significance in all analyses. Statistical analyses were performed using IBM SPSS Statistics for Windows, version 21.0 (IBM Corp., Armonk, NY, USA).

RESULTS

Baseline Characteristics according to Frailty Status based on the FRAIL Scale
As shown in Table 1, 471 (36.5%) participants were classified as robust, 524 (40.5%) as prefrail, and 297 (23%) as frail according to the K-FRAIL scale. The frail group was older, more likely to be female, more likely to be living alone, and had lower income than the other two groups (p < 0.001). The frail group also had a higher burden of multimorbidity and polypharmacy and was more likely to have depression and cognitive impairment than the other two groups (p < 0.001). Physical performance was worse and the prevalence of disability was higher in the frail group than in the other two groups (p < 0.001).

Associations between Frailty Status and Geriatric Conditions
In Table 2, frailty according to the FRAIL scale was associated with geriatric conditions after adjusting for age and sex. The frail group showed a significant association with multimorbidity (odds ratio [OR] = 2.24; 95% confidence interval [CI], 1.57–3.21), dysmobility (OR = 5.35; 95% CI, 3.58–7.97), malnutrition (OR = 4.09; 95% CI, 2.83–5.90), cognitive impairment (OR = 3.52; 95% CI, 2.36–5.25), depression (OR = 27.50; 95% CI, 11.09–68.19), and disability (OR = 3.12; 95% CI, 2.18–4.46).

Associations between FRAIL Scale and Health Outcomes
Longitudinal analysis showed that baseline frailty status significantly increased the risks of mortality (OR = 3.57; 95% CI, 1.69–7.54), institutionalization (OR = 2.42; 95% CI, 1.30–4.51), and disability at 3 years (OR = 3.04; 95% CI, 1.91–4.85) but not the
Table 1. Baseline characteristics according to frailty status in the Korean version of the FRAIL scale

|                      | Robust (n=471) | Prefrail (n=524) | Frail (n=297) | p-value$^a$ |
|----------------------|---------------|------------------|--------------|-------------|
| Age (y)              | 72.2 ± 5.2    | 74.6 ± 6.1       | 78.3 ± 6.3   | <0.001      |
| Sex, female          | 188 (40.0)    | 312 (59.5)       | 217 (73.1)   | <0.001      |
| Living alone         | 91 (19.3)     | 149 (28.4)       | 101 (34.0)   | <0.001      |
| Low income           | 21 (4.5)      | 37 (7.1)         | 30 (10.1)    | 0.010       |
| BMI (kg/m$^2$)       | 24.6 ± 3.3    | 24.5 ± 3.7       | 24.3 ± 3.8   | 0.534       |
| Multimorbidity       | 139 (29.5)    | 210 (40.1)       | 168 (56.6)   | <0.001      |
| Polypharmacy         | 73 (15.5)     | 118 (22.5)       | 107 (36.0)   | <0.001      |
| Cognitive dysfunction| 62 (13.2)     | 164 (31.3)       | 157 (52.9)   | <0.001      |
| Depressive mood      | 6 (1.3)       | 49 (9.4)         | 87 (29.3)    | <0.001      |
| At risk of malnutrition| 1 (0.0)     | 20 (3.8)         | 43 (14.5)    | <0.001      |
| SPPB score           | 9.7 ± 2.3     | 8.0 ± 2.8        | 5.0 ± 3.1    | <0.001      |
| Dysmobility          | 61 (13.0)     | 133 (25.4)       | 169 (56.9)   | <0.001      |
| Sarcopenia           | 78 (16.6)     | 185 (35.3)       | 145 (48.8)   | <0.001      |
| ADL disability       | 17 (3.6)      | 76 (14.5)        | 86 (29.0)    | <0.001      |
| IADL disability      | 111 (23.6)    | 149 (28.4)       | 176 (59.3)   | <0.001      |

Values are presented as mean±standard deviation or number (%).
FRAIL scale, Fatigue, Resistance, Ambulation, Illnesses, & Loss of Weight scale; BMI, body mass index; SPPB, Short Physical Performance Battery; ADL, activity of daily living; IADL, instrumental activity of daily living.

$^a$Kruskal–Wallis test.

Table 2. Associations between frailty status and geriatric conditions

| Outcome                      | Level       | Cases | OR$^b$ | 95% CI       | p-value |
|------------------------------|-------------|-------|--------|--------------|---------|
| Living alone (n = 341)       | Robust (Ref)| 91    | -      | -            | -       |
|                             | Prefrail    | 149   | 1.13   | 0.82–1.56    | 0.462   |
|                             | Frail       | 101   | 1.08   | 0.73–1.61    | 0.702   |
| Low income (n = 88)          | Robust (Ref)| 21    | -      | -            | -       |
|                             | Prefrail    | 37    | 1.43   | 0.81–2.52    | 0.220   |
|                             | Frail       | 30    | 1.61   | 0.83–3.13    | 0.158   |
| Multimorbidity (n = 517)     | Robust (Ref)| 139   | -      | -            | -       |
|                             | Prefrail    | 210   | 1.33   | 1.01–1.75    | 0.047   |
|                             | Frail       | 168   | 2.24   | 1.57–3.21    | <0.001  |
| Dysmobility (n = 363)        | Robust (Ref)| 61    | -      | -            | -       |
|                             | Prefrail    | 133   | 1.53   | 1.07–2.19    | 0.02    |
|                             | Frail       | 169   | 5.35   | 3.58–7.97    | <0.001  |
| Malnutrition (n = 64)        | Robust (Ref)| 1     | -      | -            | -       |
|                             | Prefrail    | 20    | 2.29   | 1.72–3.06    | <0.001  |
|                             | Frail       | 43    | 4.09   | 2.83–5.90    | <0.001  |
| Impaired cognition (n = 383)  | Robust (Ref)| 62    | -      | -            | -       |
|                             | Prefrail    | 164   | 1.96   | 1.38–2.79    | <0.001  |
|                             | Frail       | 157   | 3.52   | 2.36–5.25    | <0.001  |
| Depressive mood (n = 142)    | Robust (Ref)| 6     | -      | -            | -       |
|                             | Prefrail    | 49    | 6.24   | 2.62–14.9    | <0.001  |
|                             | Frail       | 87    | 27.50  | 11.09–68.19  | <0.001  |
| Disability (n = 615)         | Robust (Ref)| 128   | -      | -            | -       |
|                             | Prefrail    | 225   | 1.12   | 0.84–1.50    | 0.448   |
|                             | Frail       | 262   | 3.12   | 2.18–4.46    | <0.001  |

OR, odds ratio; CI, confidence interval.

$^b$According to age, sex-adjusted logistic regression analysis.
deterioration of the quality of life (Table 3). The differences in 3-year mortality according to the FRAIL scale are shown in Fig. 1. The frail group had significantly lower estimated median survival (203 months) than the prefrail (410 months) and robust (357 months) groups at 30 months follow-up (all p < 0.05).

DISCUSSION

The results of this prospective study showed that frailty according to the K-FRAIL scale was associated with accompanying geriatric conditions and with 3-year health-related outcomes such as mortality, institutionalization, and disability in community-dwelling Korean older adults.

The K-FRAIL scale showed high sensitivity and negative predictive value in differentiating vulnerability from robustness; however, the quantification of frailty status with respect to various health-related outcomes has been understudied. The clinical implications of our findings are that frail individuals identified using the K-FRAIL scale have an elevated disability, institutionalization, and mortality profile. Therefore, these frail individuals might benefit from further comprehensive geriatric assessments and multidomain interventions. The K-FRAIL scale is correlated with the previously validated frailty index and other frailty measures.\(^\text{(16)}\) Our results provide evidence for the adequacy of the FRAIL scale, including its associations with health-related outcomes, as well as identifying individuals with a high burden of geriatric conditions in rural communities.

A previous systematic review and meta-analysis of 3–5 studies demonstrated high ADL (pooled OR = 9.82, compared with robustness) and IADL (pooled OR = 2.50, compared with robustness) disability risks for frailty according to the FRAIL scale.\(^\text{(20)}\)

Table 3. Association between FRAIL scale and health outcomes

| Outcome                  | Level       | OR\(^a\) | 95% CI   | p-value |
|--------------------------|-------------|----------|----------|---------|
| Mortality                | Robust (Ref)| -        | -        |         |
|                          | Prefrail    | 1.53     | 0.78–2.98| 0.214   |
|                          | Frail       | 3.57     | 1.69–7.54| <0.001  |
| Institution              | Robust (Ref)| -        | -        |         |
|                          | Prefrail    | 1.69     | 0.96–2.95| 0.068   |
|                          | Frail       | 2.42     | 1.30–4.51| 0.005   |
| Incidence of disability  | Robust (Ref)| -        | -        |         |
|                          | Prefrail    | 1.58     | 1.10–2.27| 0.013   |
|                          | Frail       | 3.04     | 1.91–4.85| <0.001  |
| Mortality or institution | Robust (Ref)| -        | -        |         |
|                          | Prefrail    | 1.70     | 1.09–2.67| 0.02    |
|                          | Frail       | 3.14     | 1.90–5.18| <0.001  |
| Mortality, institution, or incident disability | Robust (Ref)| -        | -        |         |
|                          | Prefrail    | 1.67     | 1.22–2.29| 0.001   |
|                          | Frail       | 3.78     | 2.52–5.66| <0.001  |
| EQ-SD decrease of ≥ 0.2  | Robust (Ref)| -        | -        |         |
|                          | Prefrail    | 1.44     | 0.67–3.09| 0.350   |
|                          | Frail       | 0.57     | 0.19–1.69| 0.315   |

FRAIL scale, Fatigue, Resistance, Ambulation, Illnesses, & Loss of Weight scale; OR, odds ratio; CI, confidence interval; EQ-SD, EuroQol-SD.

\(^a\)For 3-year outcomes according to age, gender-adjusted logistic regression analysis.
Our results using the K-FRAIL scale were consistent with previous results. Similar to our results, another meta-analysis of three studies showed that both frailty (pooled OR = 3.53, compared with robustness) and prefrailty (pooled OR = 1.75 compared with robustness) were significantly associated with higher mortality risk. Although previous studies less often focused on institutionalization as an outcome of the FRAIL scale, we found that frailty according to the K-FRAIL scale was associated with an increased risk of long-term institutionalization. The results of our cross-sectional analysis showed that frailty was highly associated with depressed mood, dysmobility, risk of malnutrition, and multimorbidity. The fact that the FRAIL scale encompasses comorbidities should be considered when interpreting multimorbidity rates.

However, the longitudinal analysis showed that baseline frailty status was not associated with future QOL. Although the mechanism is not clear in the current study, the lack of association between the FRAIL scale and future QOL decline might be due to the characteristics of the FRAIL scale rather than the clinical construct of frailty itself, with a previous study being able to produce meaningful changes in frailty status from the same study population using EQ-5D as an anchor.

Our study has several strengths and limitations. The limitation was that the results were obtained from underserved older populations in rural areas, which restricted their generalizability to the general population of Korea. However, this cohort study included more than 90% of eligible older adults, with high completion rates of geriatric parameters in the study population. The outcomes of functional changes were available in our study population because the study population remained relatively stable, with < 5% of older adults immigrating or emigrating from the area except for morbidity or mortality, with the study area rather distant from major cities in Korea. Residents received medical and preventive care almost exclusively from regional community health posts operated by the National Health Service. Therefore, we could examine long-term clinical outcomes such as mortality and institutionalization.

In conclusion, frailty according to the five-item FRAIL scale was associated with geriatric syndromes and 3-year health-related outcomes in rural community-dwelling Korean older adults. In the context of public health, this simple screening tool for frailty assessment might be applicable to older populations in Korea.

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CONFLICT OF INTEREST

The researchers claim no conflicts of interest.

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AUTHOR CONTRIBUTION

Conceptualization, HL, JYC, HWJ, IYJ; data acquisition, IYJ; data analysis, HL, JYC, IYJ; writing, original draft, HL, JYC, HWJ, IYJ; writing review and editing, HL, JYC, HWJ, JYB, EL, IYJ.

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