Risk Factors of Frailty in Patients with Distal Radius Fractures

Geriatric Orthopaedic Surgery & Rehabilitation Volume 13: 1–8 © The Author(s) 2022 Article reuse guidelines: sagepub.com/journals-permissions DOI: 10.1177/21514593221094736 journals.sagepub.com/home/gos

Jeong-Hyun Kang, DDS, PhD¹ and Seok Woo Hong, MD, PhD²¹⁰

Abstract

Aim: The aim of this study was to determine risk factors for the incidence of frailty in patients with distal radius fractures (DRFs). **Methods:** In total, 116 patients (mean age, 66.3 ± 7.7 years) with DRFs were recruited. The participants were categorized into two groups, "frail" and "non-frail," according to the presence or absence of frailty, respectively. The areal bone mineral densities (aBMDs) of the total hip, femoral neck, and lumbar spine were measured using dual-energy x-ray absorptiometry. The participants' levels of resilience, depression, anxiety, nutritional intake, oral health-related quality of life, and social support were evaluated by self-reported questionnaires. The participants' grip strength, gait speed, number of teeth present in their oral cavities, circumference of their upper arms and calves, and serum levels of vitamin D were also assessed. **Results:** The participants in the "frail" group seemed to have lower aBMDs and muscle function and mass than those in the "non-frail" group. There were significant differences in grip strength, calf circumference, gait speed, and aBMD of the total hip, femoral neck, and lumbar spine between the groups. There were also significant differences in the levels of sarcopenia, malnutritional status, and aBMDs of the total hip and femoral neck had significant relationships with the development of frailty in patients with DRFs. **Conclusions:** An interdisciplinary approach involving the management of osteoporosis, sarcopenia, oral health, social relationships, and psychological support would be required for the proper management of DRF patients in preventing frailty.

Keywords

frailty, osteoporosis, sarcopenia, fracture, distal radius

Introduction

Frailty is commonly regarded as a type of geriatric syndrome featuring age-associated declines in physiological reserves and function across multi-organ systems, leading to increased vulnerabilities to adverse health outcomes.¹ In aged societies, the increasing numbers of frail elderly have become major issues owing to high social and medical costs. Therefore, identifying the risk factors for frailty and facilitating proper interventions to minimize frailty in elder populations are the main concerns of clinicians and policy makers of aged societies.

Fractures are considered to be important issues in geriatric medicine as they may lead to frailty, which, in turn, may increase the potential for falls and subsequent fractures.² Osteoporosis is one of the most common diseases in elderly populations and strongly correlates

¹Clinic of Oral Medicine and Orofacial Pain, Institute of Oral Health Science, Ajou University School of Medicine, Suwon, Korea ²Department of Orthopedic Surgery, Kangbuk Samsung Hospital, Sungkyunkwan University School of Medicine, Seoul, Korea

Submitted February 23, 2022. Accepted March 30, 2022

Corresponding Author:

Seok Woo Hong, MD, PhD, Division of Hand Surgery, Department of Orthopedic Surgery, Kangbuk Samsung Hospital, SungKyunKwan University School of Medicine, 29, Saemunan-ro, Jongno-gu, Seoul 03181, South Korea. Email: poisoxic@naver.com



Creative Commons Non Commercial CC BY-NC: This article is distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 License (https://creativecommons.org/licenses/by-nc/4.0/) which permits non-commercial use, reproduction and distribution of the work without further permission provided the original work is attributed as specified on the

SAGE and Open Access pages (https://us.sagepub.com/en-us/nam/open-access-at-sage).

with an increased incidence of fractures owing to low bone mass and the microarchitectural deterioration of bone tissues.³⁻⁵ Frailty, osteoporosis, and sarcopenia share various risk factors and pathophysiological pathways such as aging, low body mass, decreased physical activities, low muscle mass, inflammation, and vitamin D deficiencies.^{2,6-8} Many previous reports have focused on the increased probability of frailty in patients with osteoporotic fractures,^{9,10} and other studies have suggested that frailty itself could cause an increased prevalence of osteoporotic fractures.¹¹⁻¹⁶ Therefore, determining the risk factors of frailty in patients with osteoporotic fractures and the confounding factors correlating with the occurrence of osteoporotic fractures in the frail elderly is an important consideration of geriatric medicine in order to reduce the social and medical burdens of elderly populations.

Distal radius fractures (DRFs) are the most common upper extremity fractures in women and represent 17.5% of all types of fractures.^{17,18} Patients with DRFs showed an increased risk of subsequent fractures including hip and spine fractures, which could lead to locomotive problems, functional declines, and frailty.^{19,20} There are a limited number of studies that have investigated the comprehensive and interdisciplinary relationship between DRFs and frailty. One study examined the incidence rate of frailty in DRF patients, but this reports included only a small number of samples, which inevitably compromised the validity of the results.²¹ Another study suggested the value of frailty evaluation criteria as predictive factors of preoperative complications and increased lengths of hospital stay for DRF patients who underwent surgical procedures.²² However, this study utilized retrospective data and simplified frailty evaluation criteria which could have undermined the study's value. Hence, the aim of the present study was to reveal the relationships between DRFs and frailty and determine the risk factors for the incidence of frailty in DRF patients.

Materials and Methods

Participants

In total, 116 patients (mean age, 66.3 ± 7.7 years; age range: 55–93) with DRFs were recruited from a tertiary care hospital from September, 2020 to December, 2021. A single orthopedic surgeon interviewed all the participants and gathered information regarding the participants' underlying diseases, number of teeth present in their oral cavity, and current and past medications. Participants with histories of taking bone active medications including bisphosphonates, denosumab, and hormone modulating agents, and participants who were uncommunicative were excluded. Body mass index (BMI), upper arm circumference, and calf circumference were measured by a trained nurse. The research protocol was approved by the Institutional Review Board of the University Hospital. Informed consents were obtained from all participants.

Assessment of Frailty

Frailty was assessed in accordance with the criteria suggested by Fried.²³ The Fried frailty index is comprised of five criteria: weight loss, exhaustion, low physical activity, decreased walking speed, and grip strength. Unintentional weight loss and exhaustion were assessed by self-administered questionnaires. Walking speed over 4 m was measured using a timer with acceleration and deceleration phases of 1.5 m. The mean values were selected from three independent measurements. The lowest 20% of gait speed adjusted by sex and height based on the Korean frailty and aging cohort study was used to determine cut-off values.² Handgrip strength was measured by a hand dynamometer (Jamar[®] 5030J1 hydraulic hand dynamometer, Sammons Preston Rolyan, Bolingbrook, IL, USA) on the uninjured contralateral hand during the patient's initial hospital visit. Handgrip strength was taken in a sitting position with a 90° elbow flexion and a neutral forearm position. The cut-off values of grip strength were determined from 20th percentile of grip strength stratified by sex and BMI quartiles based on the Korean frailty and aging cohort study.²⁴ Measurement of walking speed and grip strength were conducted by a trained orthopedic surgeon. Energy expenditure estimates (kcal/week) were calculated using the international physical activity questionnaire and metabolic equivalent scores were derived from vigorous, moderate, and mild activities from the self-administered questionnaires. A low physical activity level was defined as less than 495 kcal for men and 283.50 kcal for women.²⁵ If a participant showed three positive criteria on the Fried's index, this participant was classified as "frail," if a participant showed 1-2 positive criteria, the participant was classified as "pre-frail," and finally, if a participant showed no positive criteria, the participant was classified as "robust." We categorized participants into two groups, "frail" (N = 40, mean age = 71.5 \pm 8.5 years) and "non-frail" (N = 76, mean age = $63.5 \pm$ 5.5 years) including pre-frail and robust participants.

We also used the Kihon checklist to investigate the frailty status, which consist of seven domains including instrumental activities of daily living, physical function, nutritional status, oral function, homebound status, cognitive function, and mood.²⁶ The Kihon checklist is a self-administered questionnaire, consisting of 25 items concerning 7 domains.

Although not a disease, frailty is a multifactorial condition involving multi-organs and is related with to the general mental and physical health of the elderly.¹ Therefore, concerning diverse aspects of the elderly's lives such as psychological status, social relationships, resilience, nutritional status, oral health status, and physical function are essential to reveal the risk factors for frailty. We adopted the Brief Resilience Scale (BRS),²⁷ the short form Geriatric Depression Scale (GDS),²⁸ the Geriatric Anxiety Inventory (GAI),²⁹ the Mini-nutritional Assessment (MNA),³⁰ the Oral Health Impact Profile-14 (OHIP-14),^{31,32} the Enhancing Recovery in Coronary Heart Disease (ENRICHD),³³ and SARC-F³⁴ to assess levels of resilience, depression, anxiety, nutritional intake, oral health-related quality of life, and social support and sarcopenia screening, respectively.

Evaluation of Bone Mineral Density

The areal bone mineral densities (aBMDs, in grams per square centimeter) of the total hip, femoral neck, and lumbar spine (L1-L4) were measured using dual-energy x-ray absorptiometry (DEXA) with a Hologic device (Horizon-W; Hologic Inc., Bedford, MA, USA). T scores of the aBMDs of the total hip, femoral neck, and lumbar spine were evaluated based on the value of aBMDs.

Biochemical Evaluation

Peripheral venous blood samples from each participant were collected between 8:00 a.m. and 11:00 a.m. after overnight fasting, to minimize circadian rhythm variabilities. The concentrations of serum 25hydroxyvitamine D was assessed by high-performance liquid chromatography.

Statistical Analysis

Based on the Shapiro–Wilk normality test, data from the present study were normally distributed, and thus parametric tests were utilized. To compare the participants' demographic characteristics, including age, sex distribution, BMI, and history of underlying diseases with the presence of frailty, independent T-tests, and chi-square tests were applied for continuous and categorical variables, respectively. Differences in the parameters, including grip strength, circumferences of the upper arm and calf, total Kihon checklist score, walking speed, energy expenditure estimates, SARC-F, EN-RICHD, BRS, GDS, GAI, OHIP-14, number of teeth present, serum vitamin D levels, and aBMDs and T scores of the total hip, femur neck, and total lumbar were also compared between the two groups by independent T-tests and chi-square tests for continuous and categorical variables, respectively. A multivariate logistic regression analysis was used to analyze the potential risk factors of frailty in DRF patients adjusted for the potential confounders. Each variable with a significant outcome in the univariate analysis was integrated into the multivariate logistic regression analysis to identify interdependent contributions after adjusting for the presence of all variables to the dependent variable, the presence of frailty. Owing to the high collinearity among aBMDs of the total hip, femur neck, and lumbar spine, separate analyses of aBMDs of the total hip, femur neck, and lumbar spine were conducted. Gait speed and energy expenditure estimates were excluded from the logistic regression analysis because these two factors were already part of Fried's frailty index for determining frailty.

Results

The significant differences of age (P < .001), grip strength (P = .009), and calf circumference (P = .003) between non-frail group and frail group were detected. The background underlying diseases such as of hypertension (P < .001), diabetes mellitus (P < .001), and heart disease (P = .033) seemed to have significant associations with development of frailty in DRF patients. On the other hand, thyroid disorders (P = .506), kidney disorders (P = .466), rheumatoid arthritis (P =.301), liver disease (P = .685), and respiratory diseases (P = .089) might have little impact on occurrence of frailty on patients with DRF. The differences of upper arm circumference (P = .777) did not show statistical significance between non-frail group and frail group (Table 1).

The significant differences of total Kihon checklist score (P < .001), walking speed (P < .001), energy expenditure estimates (P = .018), SARC-F (P = .031), levels of social support (P = .003), resilience (P < .001), depression (P < .001), and aBMD and T scores of the total hip (P < .001), femur neck (P < .001), and lumbar spine between two groups were observed. The number of teeth present (P < .001) and nutritional uptake (P < .001) seemed to have significant influences on development of frailty, also. However, levels of anxiety (P = .528), serum vitamin D concentration (P = .926) did not show statistically significant differences between non-frail group and frail group (Table 2).

	Non-frail (N = 76)	Frail (N = 40)<	P Value
Age	63.5 ± 5.5	71.5 ± 8.5	<.001**
Sex (male/female) ^a	7/69	7/33	.193
BMI (kg/m ²)	24.6 ± 3.0	24.9 ± 3.2	.600
Comorbidities (yes/no) ^a			
Hypertension	I 6/60	24/16	<.001**
Diabetes mellitus	4/72	12/28	<.001**
Heart disease	4/72	7/33	.033*
Thyroid disorder	2/74	2/38	.506
Kidney disorder	1/75	0/40	.466
Rheumatoid arthritis	2/74	0/40	.301
Liver disease	3/73	1/39	.685
Respiratory disease	2/74	4/36	.089
Grip strength (kg)	22.8 ± 5.4	19.8 ± 6.1	.009*
Upper arm circumference (cm)	27.0 ± 3.3	26.8 ± 3.0	.777
Calf circumference (cm)	33.8 ± 3.3	32.0 ± 2.6	.003*

Table I. Demographic	Characteristic	of the	Participants.
----------------------	----------------	--------	---------------

BMI, body mass index

Data obtained from independent T-test and descriptive values are shown as mean ± SE.

^aData obtained from chi-square test.

*P < .05, **P < .001 by chi-square test or independent T-test.

Table 2. Status of Frailty, Frai	ty Related Factors, and Bone	Mineral Density of the Participant
----------------------------------	------------------------------	------------------------------------

		Non-frail (N = 76)	Frail (N = 40)	P Value
Total Kihon checklist scor	·e	21.1 ± 2.4	18.9 ± 3.6	<.001**
Gait speed (m/sec)		1.16 ± .35	.78 ± .25	<.001**
Energy expenditure estimation	ates (kcal/week)	305.0 ± 669.2	48.9 ± 129.7	.018*
SARC-F		6.29 ± 2.18	7.15 ± 1.92	.031*
Enhancing Recovery in Co	oronary Heart Disease	24.8 ± 6.4	21.1 ± 5.8	.003*
Brief Resilience Scale		13.9 ± 4.2	17.7 ± 3.3	<.001**
Geriatric Depression Scale	e	2.38 ± 2.73	5.15 ± 2.79	<.001**
Geriatric Anxiety Inventor	ry	35.3 ± 6.0	34.5 ± 6.7	.528
Mini-nutritional Assessme	nt category ^a			<.001**
Normal/at risk/malnutri	tion	28/46/2	8/20/12	
Oral Health Impact Profile	e-14	4.63 ± 6.55	8.10 ± 6.40	.007*
Number of teeth ^a				<.001**
≥20/<20		68/8	24/16	
Serum vitamin D level (µg	I/L)	25.5 ± 14.4	25.7 ± 10.7	.926
Bone mineral density	Total hip aBMD (g/cm ²)	.81 ± .11	.71 ± .13	<.001**
	Total hip T score	65 ± .90	-1.41 ± 1.09	<.001**
	Femoral neck aBMD (g/cm ²)	.68 ± .09	.59 ± .11	<.001**
	Femur neck T score	−1.40 ± .74	-2.19 ± .71	<.001***
	LI-4 aBMD (g/cm ²)	.87 ± .16	.80 ± .13	.013*
	LI-4 T score	-1.45 ± 1.33	-1.97 ± 1.04	.037*

aBMD, areal bone mineral densities.

Data obtained from independent T-test and descriptive values are shown as mean \pm SE.

^aData obtained from chi-square test.

*P < .05, **P < .001 by chi-square test or independent T-test.

The multivariate logistic regression demonstrated that level of sarcopenia, malnutritional status, and aBMD of the total hip and femur neck had significant relationships with development of frailty in patients with DRF. On the other hand, the aBMD of the lumbar spine did not show significant relationship with presence of frailty in DRF patients (Table 3).

Total Hip			
	B (95% CI)	SE	P Value
Age	.139	.101	.167
Hypertension			
No		Reference	
Yes	1.241	1.175	.469
Diabetes mellitus			
No		Reference	
Yes	3.821	2.381	.108
Heart disease			
No		Reference	
Yes	-1.005	1.508	.505
Calf circumference	08I	2.207	.696
BRS	.254	.159	.111
SARC-F	1.109	.434	.011*
ENRICHD	—. 187	.114	.100
GDS	.199	.220	.366
OHIP-14	.122	.096	.203
Tooth no			
≥20		Reference	
<20	-1.300	1.434	.365
MNM category			
Normal		Reference	
At risk	-2.446	1.820	.122
malnutrition	-2.556	1.653	.047*
Total hip aBMD	-11. 480	6.366	.041*
Femoral neck			
	B (95% CI)	SE	P Value
Age	.109	.106	.301
Hypertension			
No		Reference	
Yes	1.209	1.751	.490
Diabetes mellitus			
No		Reference	
Yes	4.787	2.666	.073
Heart disease			
No		Reference	
Yes	2.142	1.839	.244
Calf circumference	—.I38	.202	.493
BRS	.179	.165	.278
SARC-F	1.369	.534	.010*
ENRICHD	—.160	.121	.187
GDS	.276	.248	.267
OHIP-14	.106	.097	.277
Tooth No			
≥20		Reference	
<20	-1.229	1.444	.395
MNM category			
Normal		Reference	
At risk	-2.45 I	1.1743	.160

Table 3. Adjusted Association Between Incidence of Frailty andRelated Risk Factors.

Table 3. (continued)

Total Hip			
	B (95% CI)	SE	P Value
Malnutrition	-2.326	1.781	.043*
Femoral neck aBMD	- 16.198	7.755	.037*
Lumbar spine	B (95% CI)	SE	P Value
Age	.189	.098	.053
Hypertension			
No		Reference	
Yes	.318	1.377	.818
Diabetes mellitus			
No		Reference	
Yes	3.032	1.731	.080
Heart disease			
No		Reference	
Yes	—.536	1.455	.713
Calf circumference	—.27I	.181	.133
BRS	.267	.150	.075
SARC-F	.902	.324	.005*
ENRICHD	—. 176	.103	.088
GDS	.190	.208	.361
OHIP-14	.102	.088	.245
Tooth no			
≥20		Reference	
<20	1.025	1.262	.417
MNM category			
Normal		Reference	
At risk	-2.113	1.419	.137
Malnutrition	-2.309	1.668	.035*
Lumbar spine aBMD	.775	4.244	.855

B, unstandardized regression coefficient; SE, standard error; aBMD, areal bone mineral density; BRS, Brief Resilience Scale; ENRICHD, Enhancing Recovery in Coronary Heart Disease; GDS, Geriatric Depression Scale; MNA, Mini-nutritional Assessment; OHIP-14, Oral Health Impact Profile-14.

Data obtained from the multivariate logistic linear regression.

*P < .05, **P < .001 by the multivariate logistic linear regression.

Discussion

The relationship between osteoporotic fractures and frailty has been discussed in previous literature. Several studies revealed the increased probability of frailty in patients with osteoporotic fractures^{9,10} and other reports have proposed that frailty itself has impacts on increased occurrence of osteoporotic fractures also.¹¹⁻¹⁶ However, majority of these studies included patients with hip and spine fractures, which might directly correlate with locomotive problems and functional declines. Even though, DRF is not directly associated with locomotive problems and functional disabilities, it has a significant relationship with decreased bone microstructures and bone density, the main indicators of osteoporosis, and DRF patients showed increased risks

(continued)

for subsequent fractures including fractures of the hip and spine.^{19,20} To the best of our knowledge, there have been spares studies that have presented thorough and multidimensional insights into the relationship between DRF and frailty. Therefore, the aim of the present study was to reveal the relationships between DRF and frailty and determine the risk factors for the incidence of frailty in DRF patients.

In general, the elderly's medical conditions may not be the result of a single disorder but instead may be the results of chronic conditions involving multiple organs.¹ Frailty is multidimensional, involving not only physical factors but social and psychological factors, as well. In the present study, we included factors related with BMD, and muscle mass as well as nutritional uptake, oral health, underlying diseases, anxiety, depression, resilience, and social supports to reveal the risk factors for frailty. Aforementioned results showed that decreased BMD and muscle function showed significant relationships with the development of frailty as previous reports have suggested.35,36 Interestingly, resilience, social support, and depression also showed significant associations with frailty in DRF patients. Despite previous reports revealing the importance of management of oral health^{37,38} and psychological problems^{39,40} in the prevention of frailty in communitydwelling elderly, the significance of these factors' contribution to frailty in osteoporotic fracture patients has not yet been determined. Social isolation, depression, and poor oral health may lead to physical inactivity and malnutrition, which then can result in decreased bone mass and muscle atrophy⁴¹⁻⁴⁴ and subsequently DRF. Hence, for the prevention of frailty in patients with osteoporotic fractures and, DRFs, monitoring bone density and muscle mass as well as the encouragement of social relationships, proper oral healthcare, and psychological support should be recommended.

Adequate nutritional intake is important for improving frailty, osteoporosis, and sarcopenia.45-47 The results from present study demonstrated the potential contribution of proper nutritional uptake in DRF patients for preventing frailty. Nutrition plays a critical role in the pathophysiology of frailty and the maintenance of bone⁴⁷ and muscle health.^{45,48} One study suggested that malnutrition could play a role as a mediator of osteosarcopenia and sarcopenia in the development of frailty.⁴⁵ Furthermore, sufficient protein and vitamin D uptake in the elderly has been recommended for the prevention of muscle and bone loss as well as the maintenance of sufficient leg and grip strength.⁴⁹ Associations between nutritional uptake, especially protein and the occurrence of post-operative surgical complications in patients with hip fractures^{50,51} and the importance of nutritional status for the improvement of activities of daily living and the incidence of subsequent fracture in DRF patients⁵² have been proposed. However, the potential role of active nutritional intervention in terms of preventing frailty in DRF patients has not been suggested. Even though serum vitamin D levels did not show significant associations with frailty in aforementioned results, the role of proper nutritional uptake cannot be discounted in the management of DRF patients.

To the best of our knowledge, the present study is the first trial to reveal the risk factors of frailty in DRF patients with consideration of frailty's multidimensional nature. However, there are still several limitations of this study. First, owing to the cross-sectional study design, causal relationships could not be derived. Second, due to the relatively, small sample size, the significance of the results could be inevitably compromised. Finally, because of the predominance of females with osteoporotic fractures, this study could not provide sufficient information about the risk factors for frailty in male DRF patients. Future prospective study with large sample numbers including adequate numbers of both male and female participants are warranted.

Frailty is a chronic and multidimensional condition in elderly and a comprehensive understanding of its features could be important for managing frail elderly population to develop strategies for preventing frailty. An interdisciplinary approach including psychology, dentistry, orthopedics, and endocrinology is essential for the proper management of DRF patients in order to prevent frailty.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This study is supported by National Research Foundation of Korea (2020R111A1A01071537).

ORCID iD

Seok Woo Hong D https://orcid.org/0000-0003-4059-1557

References

- Chen X, Mao G, Leng SX. Frailty syndrome: An overview. *Clin Interv Aging*. 2014;9:433-441. doi:10.2147/CIA.S45300.
- Rolland Y, Abellan van Kan G, Benetos A, et al. Frailty, osteoporosis and hip fracture: Causes, consequences and therapeutic perspectives. *J Nutr Health Aging*. 2008;12(5): 335-346. doi:10.1007/BF02982665.
- Yeung SSY, Reijnierse EM, Pham VK, et al. Sarcopenia and its association with falls and fractures in older adults: A systematic review and meta-analysis. J Cachexia Sarcopenia Muscle. 2019;10(3):485-500. doi:10.1002/jcsm.12411.

- Gauthier A, Kanis JA, Jiang Y, et al. Epidemiological burden of postmenopausal osteoporosis in the UK from 2010 to 2021: Estimations from a disease model. *Arch Osteoporos*. 2011;6:179-188. doi:10.1007/s11657-011-0063-y.
- NIH consensus development anel on osteoporosis prevention, diagnosis, and therapy, March 7-29, 2000: Highlights of the Conference. *South Med J.* 2001;94(6):569-573.
- Clegg A, Young J, Iliffe S, Rikkert MO, Rockwood K. Frailty in elderly people. *Lancet*. 2013;381(9868):752-762. doi:10.1016/S0140-6736(12)62167-9.
- Sternberg SA, Levin R, Dkaidek S, Edelman S, Resnick T, Menczel J. Frailty and osteoporosis in older women–a prospective study. *Osteoporos Int*. 2014;25(2):763-768. doi: 10.1007/s00198-013-2471-x.
- Walston J, McBurnie MA, Newman A, et al. Frailty and activation of the inflammation and coagulation systems with and without clinical comorbidities: Results from the cardiovascular health study. *Arch Intern Med.* 2002;11162(20): 2333-2341. doi:10.1001/archinte.162.20.2333.
- Gajic-Veljanoski O, Papaioannou A, Kennedy C, et al. Osteoporotic fractures and obesity affect frailty progression: a longitudinal analysis of the Canadian multicentre osteoporosis study. *BMC Geriatr.* 2018;18(1):4. doi:10.1186/ s12877-017-0692-0.
- Li G, Papaioannou A, Thabane L, Cheng J, Adachi JD. Frailty change and major osteoporotic fracture in the elderly: Data from the global longitudinal study of osteoporosis in women 3-year hamilton cohort. *J Bone Miner Res.* 2016; 31(4):718-724. doi:10.1002/jbmr.2739.
- Bartosch P, Malmgren L, Kristensson J, McGuigan FE, Akesson KE. In community-dwelling women frailty is associated with imminent risk of osteoporotic fractures. *Osteoporos Int.* 2021;32(9):1735-1744. doi:10.1007/ s00198-021-05886-7.
- Kojima G. Frailty as a predictor of fractures among community-dwelling older people: A systematic review and meta-analysis. *Bone*. 2016;90:116-2122. doi:10.1016/j. bone.2016.06.009.
- Li G, Compston JE, Leslie WD, et al. Relationship between obesity and risk of major osteoporotic fracture in postmenopausal women: Taking frailty into consideration. J Bone Miner Res. 2020;35(12):2355-2362. doi:10.1002/ jbmr.4139.
- Li G, Ioannidis G, Pickard L, et al. Frailty index of deficit accumulation and falls: Data from the global longitudinal study of osteoporosis in women (GLOW) hamilton cohort. *BMC Musculoskelet Disord*. 2014;15:185. doi:10.1186/ 1471-2474-15-185.
- Li G, Thabane L, Papaioannou A, Adachi JD. Comparison between frailty index of deficit accumulation and fracture risk assessment tool (FRAX) in prediction of risk of fractures. *Bone*. 2015;77:107-114. doi:10.1016/j.bone.2015.04. 028.

- Middleton R, Poveda JL, Orfila Pernas F, et al. Mortality, falls and fracture risk are positively associated with frailty: A SIDIAP cohort study of 890,000 patients. *J Gerontol A Biol Sci Med Sci.* 2022;77:148-154. doi:10.1093/gerona/glab102.
- Court-Brown CM, Caesar B. Epidemiology of adult fractures: A review. *Injury*. 2006;37(8):691-697. doi:10.1016/j. injury.2006.04.130.
- Nellans KW, Kowalski E, Chung KC. The epidemiology of distal radius fractures. *Hand Clin*. 2012;28(2):113-125. doi: 10.1016/j.hcl.2012.02.001.
- Shin YH, Hong WK, Kim J, Gong HS. Osteoporosis care after distal radius fracture reduces subsequent hip or spine fractures: A 4-year longitudinal study. *Osteoporos Int*. 2020; 31(8):1471-1476. doi:10.1007/s00198-020-05410-3.
- Klotzbuecher CM, Ross PD, Landsman PB, Abbott TA III, Berger M. Patients with prior fractures have an increased risk of future fractures: A summary of the literature and statistical synthesis. *J Bone Miner Res.* 2000;15(4):721-739. doi:10.1359/jbmr.2000.15.4.721.
- Caliskan H, Igdir V, Ozsurekci C, Caliskan E, Halil M. Frailty and sarcopenia in patients with distal radius fracture: A geriatric perspective. *Geriatr Orthop Surg Rehabil*. 2020; 11:2151459320906361. doi:10.1177/2151459320906361.
- Wilson JM, Holzgrefe RE, Staley CA, Schenker ML, Meals CG. Use of a 5-item modified frailty index for risk stratification in patients undergoing surgical management of distal radius fractures. *J Hand Surg Am.* 2018;43(8): 701-709. doi:10.1016/j.jhsa.2018.05.029.
- Fried LP, Tangen CM, Walston J, et al. Frailty in older adults: Evidence for a phenotype. *J Gerontol A Biol Sci Med Sci*. 2001;56(3):M146-M157. doi:10.1093/gerona/56.3.m146.
- Won CW, Lee S, Kim J, et al. Korean frailty and aging cohort study (KFACS): Cohort profile. *BMJ Open*. 2020;10(4): e035573. doi:10.1136/bmjopen-2019-035573.
- 25. Welfare MoHa. *Living Profiles of Older People Survey in Korea*. Sejong. Korea: Korean Institute for Health and Social Affairs, Ministry of Health and Welfare; 2008.
- Sewo Sampaio PY, Sampaio RA, Yamada M, Arai H. Systematic review of the kihon checklist: Is it a reliable assessment of frailty? *Geriatr Gerontol Int.* 2016;16(8): 893-902. doi:10.1111/ggi.12833.
- Smith BW, Dalen J, Wiggins K, Tooley E, Christopher P, Bernard J. The brief resilience scale: Assessing the ability to bounce back. *Int J Behav Med.* 2008;15(3):194-200. doi:10. 1080/10705500802222972.
- Yesavage JA, Brink TL, Rose TL, et al. Development and validation of a geriatric depression screening scale: A preliminary report. *J Psychiatr Res.* 1982;17(1):37-49. doi: 10.1016/0022-3956(82)90033-4.
- Pachana NA, Byrne GJ, Siddle H, Koloski N, Harley E, Arnold E. Development and validation of the geriatric anxiety inventory. *Int Psychogeriatr.* 2007;19(1):103-114. doi:10.1017/S1041610206003504.

- Barone L, Milosavljevic M, Gazibarich B. Assessing the older person: Is the MNA a more appropriate nutritional assessment tool than the SGA? *J Nutr Health Aging*. 2003; 7(1):13-17.
- Allen F, Locker D. A modified short version of the oral health impact profile for assessing health-related quality of life in edentulous adults. *Int J Prosthodont*. 2002;15(5):446-450.
- Slade GD, Spencer AJ. Development and evaluation of the oral health impact profile. *Community Dent Health*. 1994; 11(1):3-11.
- Mitchell PH, Powell L, Blumenthal J, et al. A short social support measure for patients recovering from myocardial infarction: the ENRICHD social support inventory. *J Cardiopulm Rehabil*. 2003;23(6):398-403. doi:10.1097/ 00008483-200311000-00001.
- Chen LK, Woo J, Assantachai P, et al. Asian working group for sarcopenia: 2019 consensus update on sarcopenia diagnosis and treatment. *J Am Med Dir Assoc.* 2020;21(3): 300-307. doi:10.1016/j.jamda.2019.12.012.
- Wilson D, Jackson T, Sapey E, Lord JM. Frailty and sarcopenia: The potential role of an aged immune system. *Ageing Res Rev.* 2017;36:1-10. doi:10.1016/j.arr.2017.01.006.
- Greco EA, Pietschmann P, Migliaccio S. Osteoporosis and sarcopenia increase frailty syndrome in the elderly. *Front Endocrinol (Lausanne)*. 2019;10:255. doi:10.3389/fendo. 2019.00255.
- Castrejon-Perez RC, Jimenez-Corona A, Bernabe E, et al. Oral disease and 3-year incidence of frailty in Mexican older adults. *J Gerontol A Biol Sci Med Sci.* 2017;72(7):951-957. doi:10.1093/gerona/glw201.
- Iwasaki M, Yoshihara A, Sato N, et al. A 5-year longitudinal study of association of maximum bite force with development of frailty in community-dwelling older adults. *J Oral Rehabil.* 2018;45(1):17-24. doi:10.1111/joor.12578.
- Hoogendijk EO, van Hout HP, Heymans MW, et al. Explaining the association between educational level and frailty in older adults: Results from a 13-year longitudinal study in the Netherlands. *Ann Epidemiol.* 2014;24(7): 538-544. doi:10.1016/j.annepidem.2014.05.002.
- Woods NF, LaCroix AZ, Gray SL, et al. Frailty: Emergence and consequences in women aged 65 and older in the women's health initiative observational study. *J Am Geriatr Soc.* 2005; 53(8):1321-1330. doi:10.1111/j.1532-5415.2005.53405.x.
- Kobayashi LC, Steptoe A. Social isolation, loneliness, and health behaviors at older ages: Longitudinal cohort study. *Ann Behav Med.* 2018;52(7):582-593. doi:10.1093/abm/kax033.

- Hawkley LC, Thisted RA, Cacioppo JT. Loneliness predicts reduced physical activity: Cross-sectional & longitudinal analyses. *Health Psychol.* 2009;28(3):354-363. doi:10. 1037/a0014400.
- Yoshida M, Hiraoka A, Takeda C, et al. Oral hypofunction and its relation to frailty and sarcopenia in communitydwelling older people. *Gerodontology*. 2021;39:26-32. doi:10.1111/ger.12603.
- Ohta M, Imamura Y, Chebib N, et al. Oral function and nutritional status in non-acute hospitalised elders. *Gerodontology*. 2021;39:74-82. doi:10.1111/ger.12612.
- Chew J, Yeo A, Yew S, et al. Nutrition mediates the relationship between osteosarcopenia and frailty: A pathway analysis. *Nutrients*. 2020;12(10):2957. doi:10.3390/ nu12102957.
- 46. Kositsawat J, Duque G, Kirk B. Nutrients with anabolic/ anticatabolic, antioxidant, and anti-inflammatory properties: Targeting the biological mechanisms of aging to support musculoskeletal health. *Exp Gerontol.* 2021;154:111521. doi:10.1016/j.exger.2021.111521.
- Siddique N, O'Donoghue M, Casey MC, Walsh JB. Malnutrition in the elderly and its effects on bone health - A review. *Clin Nutr ESPEN*. 2017;21:31-39. doi:10.1016/j. clnesp.2017.06.001.
- De Rui M, Inelmen EM, Pigozzo S, Trevisan C, Manzato E, Sergi G. Dietary strategies for mitigating osteosarcopenia in older adults: A narrative review. *Aging Clin Exp Res.* 2019; 31(7):897-903. doi:10.1007/s40520-019-01130-9.
- Cramer JT, Cruz-Jentoft AJ, Landi F, et al. Impacts of high-protein oral nutritional supplements among malnourished men and women with sarcopenia: A multicenter, randomized, double-blinded, controlled trial. *J Am Med Dir Assoc.* 2016;17(11):1044-1055. doi:10.1016/j.jamda.2016. 08.009.
- Bonjour JP, Schurch MA, Rizzoli R. Nutritional aspects of hip fractures. *Bone*. 1996;18(3 suppl):139S-144S. doi:10. 1016/8756-3282(95)00494-7.
- Myint MW, Wu J, Wong E, et al. Clinical benefits of oral nutritional supplementation for elderly hip fracture patients: A single blind randomised controlled trial. *Age Ageing*. 2013;42(1):39-45. doi:10.1093/ageing/afs078.
- 52. Nagai T, Tanimoto K, Tomizuka Y, Uei H, Nagaoka M. Nutrition status and functional prognosis among elderly patients with distal radius fracture: A retrospective cohort study. *J Orthop Surg Res.* 2020;15(1):133. doi:10.1186/ s13018-020-01657-y.