

**Table 2** Mean values (SD) of bone mineral density (g/cm<sup>2</sup>) and change rate (%) at lumbar spine L2–L4 and femoral neck over 3, 7, and 10 years, classified by age and gender

Birth cohort	Age-group (years)	L2–L4								Femoral neck						
		Baseline		2nd visit (3-year follow-up)		3rd visit (7-year follow-up)		4th visit (10-year follow-up)		Baseline	2nd visit		3rd visit		4th visit	
		n	BMD (g/cm <sup>2</sup> )	n	Change rate (%/3 years)	n	Change rate (%/7 years)	n	Change rate (%/10 years)	n	BMD (g/cm <sup>2</sup> )	Change rate (%/3 years)	Change rate (%/7 years)	Change rate (%/10 years)		
1943–1952	40–49	50	1.05 (0.15)	48	0.6 (3.8)	46	-0.6 (5.1)	43	-0.2 (5.8)	0.86 (0.09)	0.3 (4.6)	-1.8 (4.8)	-1.5 (10.9)			
1933–1942	50–59	50	0.98 (0.17)	47	1.0 (3.3)	46	-0.0 (6.3)	46	1.6 (8.0)	0.80 (0.13) <sup>a</sup>	-0.2 (4.9)	0.7 (10.0)	-3.0 (6.8)			
1923–1932	60–69	50	1.04 (0.21)	49	1.3 (3.6)	47	1.4 (7.1)	41	2.3 (9.4)	0.77 (0.11) <sup>a</sup>	1.0 (7.0)	-0.1 (9.3)	0.3 (12.5)			
1913–1922	70–79	50	0.97 (0.19)	37	0.1 (5.3)	31	-1.2 (7.9)	23	-1.5 (9.2)	0.71 (0.08) <sup>a,b,c</sup>	0.9 (6.3)	4.6 (10.2) <sup>a</sup>	6.6 (16.2) <sup>b</sup>			
1913–1952	40–79	200	1.01 (0.18)	181	0.8 (4.0)	170	0.0 (6.6)	153	0.8 (8.1)	0.79 (0.12)	0.5 (5.7)	0.5 (8.9)	-0.3 (11.7)			

SD standard deviation, BMD bone mineral density, n number of participants

<sup>a</sup> Significantly different ( $P < 0.05$ ) from values of subjects in their forties

<sup>b</sup> Significantly different ( $P < 0.05$ ) from values of subjects in their fifties

<sup>c</sup> Significantly different ( $P < 0.05$ ) from values of subjects in their sixties

BMD changes at the lumbar spine and femoral neck, beta values of FT for the rate of change for BMD at the lumbar spine at the first 3, 7, and 10 years were 0.08, 0.08, and 0.03, respectively, and those at the femoral neck were 0.21, 0.14, and 0.06, respectively. Mean FT levels were significantly related to the rate of change for BMD at the femoral neck during the first 3 years ( $R^2 = 0.05$ ,  $P < 0.01$ ), but could not predict bone change at any site at 7 or 10 years.

**Discussion**

The present study examined endogenous hormone levels among men in Japan, measuring changes in BMD over spans of 3, 7, and 10 years. The present study clarified the age distribution of endogenous sex steroids, and a significant trend was seen toward low FT levels with age. FT tended to be significantly lower in the sixties and older when compared with levels in the forties in the present study. Our results support the findings of other reports. Orwoll et al. [26] showed that testosterone levels, particularly FT levels, for 2,623 men 65 years or older were associated with increasing age. Similar findings have been described in other cross-sectional and longitudinal studies [27–29]. Based on these results, we concluded that older men tended to show lower testosterone levels than younger men, similar to the situation with E<sub>2</sub> in women. Some men might display testosterone insufficiency, as seen in women with E<sub>2</sub> insufficiency. However, we do not yet have enough evidence regarding normal ranges in young men and thresholds for testosterone insufficiency. In addition, levels of testosterone may vary among individuals and be influenced by body composition such as adipose tissue, muscle, and bone.

In contrast to testosterone, no significant age-related trend in E<sub>2</sub> was found in the present study. Little information is available regarding E<sub>2</sub> levels in older men. Orwoll et al. [26] reported that E<sub>2</sub> concentrations decreased as age increased, and similar findings have been described in various reports [30–33]. However, other studies have noted stable [34–36] or rising [37] E<sub>2</sub> levels with increasing age. Although the reasons for these discrepancies are unclear, E<sub>2</sub> levels may vary among individuals and may be influenced by body composition such as adipose tissue, muscle, and bone, as well as testosterone.

Regarding the ethnic variations in serum sex steroid levels, as most previous reports have been based on studies of Caucasian men, ethnic variations in FT levels among men remain unclear. To the best of our knowledge, the Osteoporotic Fractures in Men Study (MrOS) is the only study in which a sufficient number of Asian men have participated [26]. For reasons of differences in measurement methods, direct comparison of the present results and

those from the MrOS study is inappropriate, but FT levels among Japanese men tended to be lower than those in MrOS participants, although no significant difference in  $E_2$  levels was apparent. Orwoll et al. [26] analyzed ethnic differences in the MrOS study and stated that FT levels were lower in Asian men than in other races such as Caucasian, African-American, and Hispanic subjects, but no such differences were seen for  $E_2$ . The present results support these findings.

The present study found that serum levels of FT could offer a useful predictor of bone loss at the femoral neck within 3 years, but this effect was diluted with longer observation. Regarding the effects of testosterone on bone loss at the hip, Cauley et al. [38] reported, in an epidemiological study of 1,327 men  $\geq 65$  years old, that men in the lowest FT category experienced greater hip bone loss over 1.8 years. In addition, Ensrud et al. [39] reported that among men with weight loss, the rate of decline in total hip BMD showed a stepwise increase in magnitude with greater decreases in bioavailable testosterone from baseline. In the present study, the effect of FT levels on bone loss within the relatively short term up to 3 years was observed at the femoral neck, independent of age and BMI, supporting previous reports. Although reasons for site-specific differences in the predictive capacity of FT remain uncertain, we have already reported that bone loss rate differs depending on the site involved in another cohort study [40]. We have also reported that characteristics differ between fast bone losers at the lumbar spine and femoral neck [41]. One reason for site-specific differences might be because degenerative changes that increase BMD, such as osteophytosis or sclerotic change, are observed more frequently at the lumbar spine than at the femoral neck. These results suggest that the predictive capacity of FT might differ according to the sites involved.

A recent study showed that older men with total testosterone or  $E_2$  deficiency were more likely to be osteoporotic [19], but no report evaluated the capacity of serum sex steroids to predict occurrence of OP. Regarding the relationship between testosterone and fracture risk, Mellstrom et al. [42] reported that FT within the normal range was independently associated with the presence, but not occurrence, of osteoporotic fracture in elderly men. In contrast, an analysis from the Rotterdam Study failed to confirm any association between testosterone and fracture risk [43]. Data from the Framingham study indicated that men with low serum testosterone and  $E_2$  levels were at increased risk for incident hip fractures [44]. A recent report from the Dubbo osteoporosis epidemiology study revealed that in men older than 60 years, serum testosterone is independently associated with the risk of osteoporotic fracture [45]. We also tried to evaluate the predictive capacity of serum levels of sex steroids and occurrence of

OP based on WHO criteria [46] and osteoporotic fractures, but only identified 7 cases of OP and 10 cases of osteoporotic fractures including 1 vertebral fracture, 1 hip fracture, 2 wrist fractures, 3 costal fractures, 2 ankle fractures, and 1 finger fracture. After analysis using Cox proportional hazards models adjusted for age and BMI, serum levels of FT were significantly related to incidence of OP (hazard ratio, 0.42; 95% confidence interval, 0.19–0.90), but not to incidence of osteoporotic fractures. This analysis suggests the possibility of serum FT as a predictor for OP occurrence over 10 years. However, the number of occurrences of OP seems to be too small to reach any conclusion regarding the presence or absence of associations between sex steroids and OP or osteoporotic fractures.

There are several limitations in the present study. First, the small sample size seemed to be the most severe weakness. In fact, as already noted, only 7 cases of OP and 10 cases of osteoporotic fractures were accumulated during the 10 years of the study. Longer observation in the present cohort might be required to confirm the association between sex steroids and OP or osteoporotic fracture. Second, the dropout rate over 10 years for patients in their seventies was considerably high (54.0%). This high dropout rate might cause bias. In fact, the tendency toward an increase in BMD at the femoral neck for patients in their seventies was skewed by withdrawal bias. On the basis of this hypothesis, we reanalyzed the multivariate regression analysis to assess the change rate of BMD at the femoral neck and serum FT with exclusion of subjects in their seventies. However, the results were similar, with serum levels of FT predicting bone loss at the femoral neck within 3 years ( $\beta = 0.17$ ,  $P = 0.05$ ), but diluted effects with longer observation (7 years:  $\beta = 0.8$ ,  $P = 0.38$ ; 10 years:  $\beta = 0.03$ ,  $P = 0.77$ ). Third, all serum samples were extracted between 0900 and 1500, not at a fixed time in the morning, although samples for measurement of FT are recommended to be collected in the morning. Serum levels of testosterone tend to increase toward night, peaking in the early morning, then decreasing rapidly and reaching a nadir between 1300 and 2300. We collected samples when FT levels would probably have been decreasing toward the nadir. The present study might thus have underestimated FT values compared to collection at a fixed time in the morning.

Conversely, the study design shows several notable strengths. In this population-based cohort study, subjects were selected randomly from the resident registration list. BMD was carefully measured by a single observer (N.Y.), and measurements were repeated 3, 7, and 10 years later with high participation rate by the same device and same observer. Another strength was that the effect of serum levels of sex steroids on changes in BMD could be estimated directly.

In conclusion, we clarified that serum levels of FT could predict bone loss within 3 years, but not longer. Further observations are required to confirm the relationship between FT,  $E_2$ , and spinal OP and osteoporotic fractures. Other environmental and genetic factors should also be evaluated to develop strategies for the early prevention of OP.

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# Prevalence of Falls and the Association With Knee Osteoarthritis and Lumbar Spondylosis As Well As Knee and Lower Back Pain in Japanese Men and Women

SHIGEYUKI MURAKI,<sup>1</sup> TORU AKUNE,<sup>1</sup> HIROYUKI OKA,<sup>1</sup> YOSHIO EN-YO,<sup>2</sup> MUNEHITO YOSHIDA,<sup>2</sup> KOZO NAKAMURA,<sup>1</sup> HIROSHI KAWAGUCHI,<sup>1</sup> AND NORIKO YOSHIMURA<sup>1</sup>

**Objective.** There is little information on falls by sex and age strata in Japan, and few factors associated with falls have been established. However, the association between bone and joint diseases and falls remains unclear. We examined prevalence of falls by sex and age strata, determined its association with radiographic osteoarthritis (OA) of the knee and lumbar spine, and determined knee and lower back pain after single and multiple falls.

**Methods.** A questionnaire assessed the number of falls during 12 months preceding baseline. Knee and lumbar spine radiographs were read by Kellgren/Lawrence (K/L) grade; radiographic knee OA and lumbar spondylosis were defined as a K/L grade of 3 or 4. Knee and lower back pain were estimated by an interview.

**Results.** A total of 587 men and 1,088 women (mean  $\pm$  SD age 65.3  $\pm$  12.0 years) were analyzed. During 1 year, 79 (13.5%) men and 207 (19.0%) women reported at least 1 fall. With increasing age, the prevalence of multiple falls was higher in women, but lower in elderly men age >60 years. In men, few factors were significantly associated with falls. In women, radiographic knee OA and lumbar spondylosis, as well as knee and lower back pain, were significantly associated with multiple falls without adjustment. Lower back pain and knee pain were independently associated with multiple falls in women after adjustment.

**Conclusion.** Lower back pain and knee pain were significantly associated with multiple falls in women.

## INTRODUCTION

Falls are one of the main causes of injury, disability, and death among the elderly (1,2). In Japan, according to the

recent National Livelihood Survey of the Ministry of Health, Labour and Welfare, fall and fracture are ranked fifth among diseases that cause disabilities and subsequently require support with activities of daily living (3). However, there have been few population-based studies for prevalence of fall based on sex and age strata. Further, in terms of factors associated with falls, muscle strength, balance, vision, and functional capacities, there are traits that diminish with aging, and these factors have been suggested as predictive risk factors for falls and fractures (4). Cognitive impairment has also been established as a risk factor for falls (5), but the association of bone and joint diseases, especially osteoarthritis (OA), with falls remains unclear.

The representative sites of OA are the knee and lumbar spine. Knee OA and lumbar spondylosis (LS) are major public health issues since they cause chronic pain and disability (6–11). The prevalence of radiographic knee OA and LS is high in Japan (12,13), with 25,300,000 and 37,900,000 subjects ages  $\geq$ 40 years estimated to experience radiographic knee OA and LS, respectively (14). The National Livelihood Survey ranked OA fourth among diseases that cause disabilities and subsequently require sup-

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<sup>1</sup>Shigeyuki Muraki, MD, PhD, Toru Akune, MD, PhD, Hiroyuki Oka, MD, Kozo Nakamura, MD, PhD, Hiroshi Kawaguchi, MD, PhD, Noriko Yoshimura, MD, PhD: University of Tokyo, Tokyo, Japan; <sup>2</sup>Yoshio En-yo, MD, Munehito Yoshida, MD, PhD: Wakayama Medical University, Wakayama, Japan.

Address correspondence to Shigeyuki Muraki, MD, PhD, Department of Clinical Motor System Medicine, 22nd Century Medical and Research Center, Faculty of Medicine, University of Tokyo, Hongo 7-3-1, Bunkyo-ku, Tokyo 113-8655, Japan. E-mail: murakis-ort@h.u-tokyo.ac.jp

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### Significance & Innovations

- During 1 year, 13.5% of men and 19.0% of women reported at least 1 fall.
- With increasing age, prevalence of multiple falls was higher in women, but lower in elderly men age >60 years.
- Lower back pain and knee pain were independently associated with multiple falls in women.

port with activities of daily living (3), but there have been few studies of the association between falls and OA (15,16). In previous studies, knee OA was assessed only by interview and not by radiography. The principal clinical symptom of knee OA is pain (17), but its correlation with the radiographic severity of knee OA is not as strong as expected (12,18–20). In fact, in a study in Japan, ~20% of the subjects without knee OA had knee pain, and 30% of the subjects with severe knee OA had no knee pain (12). Therefore, knee OA diagnosed by interview could be limited by variable accuracy. In addition, men and women were not examined separately in these previous studies, although sex differences have been found in the prevalence of knee OA (12). Furthermore, knee OA is conventionally defined according to Kellgren/Lawrence (K/L) grade (21), and our previous study showed that the association of a K/L grade of 2 (knee OA with pain) was weak, but that a K/L grade of 3 or 4 (knee OA with pain) was strong (12); therefore, the association of knee OA with falls may be different between a K/L grade of 2 for knee OA and a K/L grade of 3 or 4 for knee OA. However, there are no population-based studies on the association of severity of knee OA with falls. With regard to LS, to the best of our knowledge, there have been no population-based studies regarding its association with falls.

Previous studies have shown that associations between individual risk factors and a single fall are few in number and weak compared to risk factors for multiple falls (16), indicating that single and multiple falls may have different backgrounds. Therefore, to determine factors associated with falls, single and multiple falls should be analyzed separately.

The objectives of this study were to clarify prevalence of single and multiple falls by sex and age strata in Japan using a large-scale, population-based cohort study known as Research on Osteoarthritis/osteoporosis Against Disability (ROAD). Further, we examined the associations of radiographic knee OA and LS, as well as knee and lower back pain, with single and multiple falls in Japanese men and women.

### PATIENTS AND METHODS

**Patients.** The ROAD study is a nationwide prospective study designed to establish epidemiologic indexes for evaluation of clinical evidence for the development of a disease-modifying treatment for bone and joint diseases

(OA and osteoporosis are the representative bone and joint diseases, respectively). It consists of population-based cohorts in 3 communities in Japan. A detailed profile of the ROAD study has been described elsewhere (12–14,22); a brief summary is provided here. To date, we have completed the creation of a baseline database that includes clinical and genetic information for 3,040 subjects (1,061 men and 1,979 women) with a mean age of 70.6 years (range 23–95 years), who were recruited from resident registration listings in 3 communities: an urban region in Itabashi, Tokyo; a mountainous region in Hidakagawa, Wakayama; and a coastal region in Taiji, Wakayama.

Residents of these regions were recruited from the resident registration list of the relevant region. The participants in the urban region were recruited from a randomly selected cohort from the Itabashi-ward residents' registration database (22). The participation rate was 75.6%. The participants in mountainous and coastal regions were also recruited from the resident-registration lists, and the participation rates in these 2 areas were 56.7% and 31.7%, respectively. The inclusion criteria, apart from residence in the communities mentioned above, were the ability to 1) walk to the survey site, 2) report data, and 3) understand and sign an informed consent form. The baseline survey of the ROAD study was completed in 2006. All participants provided their written informed consent, and the study was conducted with the approval of the ethics committees of the University of Tokyo and the Tokyo Metropolitan Institute of Gerontology.

**Falls assessment.** All subjects were interviewed with regard to falls and fractures by experienced interviewers and were asked the following questions: "Have you experienced falls during the 12 months preceding baseline, and if yes, how many falls did you experience?" and "Have you experienced any fractures when you fell?" According to a previous study on falls (23), a fall is defined as a sudden, unintentional change in position causing an individual to land at a lower level on an object, the floor, or the ground, other than as a consequence of a sudden onset of paralysis, epileptic seizure, or overwhelming external force.

**Pain assessment.** All subjects were also interviewed by experienced orthopedists (SM and HO) with regard to knee pain and lower back pain and were asked the following questions: "Have you experienced knee pain on most days in the past year, in addition to now?" and "Have you experienced lower back pain on most days in the past year, in addition to now?" Those who answered yes were defined as having pain.

**Radiographic assessment.** All participants underwent radiographic examination of both knees using anteroposterior and lateral views with weight-bearing and foot map positioning; radiographic examination of the anteroposterior and lateral views of the lumbar spine, including intervertebral levels L1/2 to L5/S, was also performed. Knee and lumbar spine radiographs were read without the knowledge of participant clinical status by a single, experienced orthopedist (SM) using the K/L radiographic atlas

(21) to determine the severity of K/L grading. Radiographs were scored as grade 0 through 4, with higher grades being associated with more severe OA. We defined knee OA and LS as a K/L grade of  $\geq 3$  in at least 1 knee and 1 intervertebral level, respectively. To evaluate the intraobserver variability of K/L grading, 100 randomly selected radiographs of the knee and the lumbar spine were scored by the same observer more than 1 month after the first reading. One hundred other radiographs were also scored by 2 experienced orthopedic surgeons (SM and HO) using the same atlas for interobserver variability. The intra- and interobserver variabilities evaluated were confirmed by kappa analysis to be sufficient for assessment (0.86 and 0.80 for knee OA, and 0.84 and 0.76 for LS, respectively).

**Covariates.** Anthropometric measurements included height, weight, and body mass index (BMI;  $\text{kg}/\text{m}^2$ ). Grip strength was measured on bilateral sides using a TOEI LIGHT handgrip dynamometer, and the best measurement was used to characterize maximum muscle strength. To measure physical performance, the time taken to walk 6 meters at normal walking speed in a hallway was recorded. Subjects were told to walk from a marked starting line to a 6-meter mark as if they were walking down their hallway at home. Time was measured in seconds with a stopwatch and rounded to the nearest hundredth of a second. The average of 2 trials was recorded. These gait-speed trial measurements are considered highly reliable in community-dwelling elderly subjects (24–27).

The time taken for 5 consecutive chair rises without the use of hands was also recorded. Hands were folded in front of the chest with feet flat on the floor, following the protocol described by Guralnik et al (28) and used by other researchers (25,29,30). Time was measured in seconds with a stopwatch and rounded to the nearest hundredth of a second. Timing began with the command “go” and ended when the buttocks contacted the chair on the fifth landing. The reliability of this protocol is adequate (25,28,29). Cognition was also evaluated for all subjects using a Mini-Mental State Examination, and a cutoff score of  $< 24$  was used to select participants with cognitive impairment (31).

**Statistical analyses.** The differences in age, anthropometric measurements, and physical performance measurements between men and women were examined by Student's unpaired *t*-test, and among groups of nonfallers, single fallers, and multiple fallers using one-way analysis of variance (ANOVA). The prevalence of cognitive impairment, radiographic knee OA and LS, and knee and lower back pain was compared between men and women, and among nonfallers, single fallers, and multiple fallers by using the chi-square test. The prevalence of single and multiple falls was also compared between men and women, among subjects with no knee OA (K/L grade 0 or 1), with K/L grade 2 for knee OA and K/L grade 3 or 4 for knee OA, and among subjects with no LS (K/L grade 0 or 1), with K/L grade 2 for LS, and K/L grade 3 or 4 for LS by using the chi-square test. The association of knee pain and lower back pain with physical performance was deter-

mined by logistic regression analysis. Multinomial logistic regression analysis was also used to determine the association of anthropometric measurements, physical performance, cognitive impairment, radiographic knee OA and LS defined as K/L grade 3 or 4, and knee and lower back pain, with single and multiple falls compared with nonfalls. Further, to determine the independent association of radiographic knee OA and LS, and knee and lower back pain with single and multiple falls compared with nonfalls, we first used multinomial logistic regression analysis with age, BMI, cognitive impairment, radiographic knee OA and LS, and knee and lower back pain as independent variables. In addition to the above independent variables, we additionally adjusted for grip strength, 6-meter walking time, and chair stand time. Data analyses were performed using SAS software, version 9.0.

## RESULTS

Of the 1,690 subjects in the mountainous and seaside cohorts at baseline, 15 subjects provided incomplete fall questionnaires, leaving a total of 1,675 subjects (587 men, 1,088 women). Table 1 shows the age, anthropometric measurements, and physical performance of the participants in the present study. Regarding physical performance, grip strength, 6-meter walking time, and chair stand time were significantly better in men than in women. The prevalence of cognitive impairment was not significantly different between men and women. The prevalence of radiographic knee OA and knee pain was significantly higher in women than in men, while that of LS and lower back pain was not different between men and women.

During the 12 months preceding the baseline examination, 79 men (13.5%, 95% confidence interval [95% CI] 10.9–16.5%) and 207 women (19.0%, 95% CI 16.8–21.5%) reported at least 1 fall, and 48 men (8.2%, 95% CI 6.2–10.7%) and 80 women (7.4%, 95% CI 5.9–9.1%) reported multiple falls. Chi-square test showed that the prevalence of single and multiple falls were significantly different between men and women ( $P < 0.0001$ ). Among 286 subjects with at least 1 fall, 6 subjects (2.1%) had a wrist fracture, 2 (0.7%) had a proximal humerus fracture, 1 (0.3%) had a vertebral fracture, and 12 (4.2%) had fractures at other sites. With increasing age, the prevalence of falls was lower in elderly men age  $> 60$  years; however, the prevalence of falls was higher in women with increasing age (Table 2). Moreover, with increasing age, the prevalence of multiple falls was also lower in elderly men age  $> 60$  years, but it was higher in women with increasing age (Table 2). The prevalence (95% CI) of a single fall (%) was similar among age strata in men and women (for men: 5.3% [1.8–14.4%], 6.8% [3.3–13.4%], 3.2% [1.4–7.3%], 5.5% [3.2–9.4%], and 7.4% [1.0–12.5%] in the age subgroups of  $< 50$  years, 50–59 years, 60–69 years, 70–79 years, and  $\geq 80$  years, respectively; for women: 11.9% [7.5–18.5%], 11.1% [7.5–16.1%], 12.0% [8.9–16.0%], 11.6% [8.6–15.6%], and 11.4% [6.7–18.9%] in the age subgroups of  $< 50$  years, 50–59 years, 60–69 years, 70–79 years, and  $\geq 80$  years, respectively).

Table 3 shows the age, anthropometric measurements,

	Overall	Men	Women
Subjects, no.	1,675	587	1,088
Age, years	65.3 ± 12.0	66.3 ± 11.7	64.7 ± 12.1†
Height, cm	155.1 ± 9.3	163.4 ± 7.2	150.6 ± 6.9†
Weight, kg	55.6 ± 10.8	62.3 ± 10.9	52.0 ± 8.9†
BMI, kg/m <sup>2</sup>	23.0 ± 3.4	23.3 ± 3.2	22.9 ± 3.5†
Grip strength, kg	27.4 ± 9.8	35.7 ± 9.3	22.9 ± 6.8†
6-meter walking time, seconds	5.5 ± 2.5	5.3 ± 2.2	5.6 ± 2.6†
Chair stand time, seconds	10.1 ± 4.4	9.7 ± 3.6	10.4 ± 4.8†
Cognitive impairment, %	4.5	5.2	4.2
Radiographic knee OA, %	20.3	15.0	23.0‡
Radiographic lumbar spondylosis, %	37.1	37.7	36.9
Knee pain, %	24.4	18.9	27.4‡
Lower back pain, %	20.1	21.7	21.2

\* Values are the mean ± SD unless indicated otherwise. BMI = body mass index; OA = osteoarthritis.  
† P < 0.05 vs. men by Student's unpaired t-test.  
‡ P < 0.05 vs. men by chi-square test.

physical performance, and prevalence of cognitive impairment among nonfallers, single fallers, and multiple fallers. One-way ANOVA showed that there were no significant associations of age, anthropometric measurements, physical performance, and prevalence of cognitive impairment with falls in men, while age and BMI were higher in multiple fallers than in nonfallers in women. With regard to physical performance, grip strength was lower and 6-meter walking time and chair stand time were longer in multiple fallers than in nonfallers and single fallers in women. Further, prevalence of cognitive impairment was also different among nonfallers, single fallers, and multiple fallers in women. Further, to determine the association of anthropometric measurements, physical performance, and cognitive impairment with single and multiple falls, we also used multinomial logistic regression analysis and found that age (odds ratio [OR] 1.04, 95% CI 1.02–1.06), BMI (OR 1.10, 95% CI 1.03–1.17), grip strength (OR 0.92, 95% CI 0.89–0.96), 6-meter walking time (OR 1.10, 95% CI 1.02–1.17), chair stand time (OR 1.06, 95% CI 1.02–1.10), and cognitive impairment (OR 3.86, 95% CI 1.67–3.83) were significantly associated with multiple falls in women.

To determine the association of the severity of knee OA with falls, we classified subjects as those with no knee OA (K/L grade 0 or 1), with K/L grade 2 for knee OA, and with K/L grade 3 or 4 for knee OA. The prevalence of falls in subjects with no knee OA, K/L grade 2 for knee OA, and

K/L grade 3 or 4 for knee OA was 11.8%, 17.1%, and 12.5%, and 17.7%, 17.6%, and 25.6% in men and women, respectively. There were no significant associations between falls and the severity of knee OA in men (chi-square test;  $P = 0.27$ ), while prevalence of falls was higher in women with K/L grade 3 or 4 for knee OA than those with no knee OA and K/L grade 2 for knee OA ( $P = 0.01$ ). Similar to knee OA, we classified subjects as those with no LS (K/L grade 0 or 1), those with K/L grade 2 for LS, and those with K/L grade 3 or 4 for LS. The prevalence of falls in subjects with no LS, K/L grade 2 for LS, and K/L grade 3 or 4 for LS was 16.3%, 11.3%, and 14.0%, and 17.0%, 20.5%, and 20.7% in men and women, respectively. There were no significant associations between falls and the severity of LS in men and women (chi-square test,  $P = 0.38$  and 0.32, respectively). We next used the chi-square test to determine the association of single and multiple falls with knee OA and LS defined as K/L grade 3 or 4 (Table 4). A chi-square test showed that no significant factors were associated with falls in men, but radiographic knee OA, knee pain, and lower back pain were significantly associated with falls in women.

Multinomial logistic regression analysis also showed that radiographic knee OA, LS, and knee and lower back pain were significantly associated with multiple falls in women (Table 5). Because knee pain and lower back pain were also significantly associated with grip strength, 6-meter walking time, and chair stand time in men and women

Age, years	Single fall		Multiple falls	
	Men	Women	Men	Women
<50	15.8 (8.5–27.4)	13.4 (8.7–20.2)	10.5 (4.9–21.1)	1.5 (0.4–5.3)
50–59	10.7 (6.1–18.1)	17.4 (12.8–23.1)	3.9 (1.5–9.6)	6.3 (3.7–10.4)
60–69	16.7 (11.6–23.3)	18.8 (14.9–23.4)	13.5 (9.0–19.7)	6.8 (4.5–10.1)
70–79	12.4 (8.7–17.5)	21.1 (16.9–25.9)	6.9 (4.2–11.1)	9.4 (6.7–13.1)
≥80	11.1 (5.2–22.2)	23.8 (16.7–32.8)	3.7 (1.0–12.5)	12.4 (7.4–20.0)

\* Values are the percentage (95% confidence interval).



**Table 3. Comparison of characteristics among nonfallers, single fallers, and multiple fallers in men and women\***

	Men				Women			
	Nonfallers	Single fallers	Multiple fallers	P	Nonfallers	Single fallers	Multiple fallers	P
Subjects, no.	508	31	48		881	127	80	
Age, years	66.4 ± 11.7	67.6 ± 11.9	64.6 ± 11.3	0.50	64.4 ± 12.1	64.3 ± 12.2	69.1 ± 10.4	0.004
Height, cm	163.5 ± 7.4	162.3 ± 6.3	162.9 ± 5.9	0.56	150.9 ± 6.8	150.7 ± 7.7	148.5 ± 7.0	0.01
Weight, kg	62.6 ± 11.1	60.7 ± 10.4	60.3 ± 9.0	0.27	51.8 ± 8.8	53.3 ± 9.2	52.8 ± 8.9	0.15
BMI, kg/m <sup>2</sup>	23.3 ± 3.2	23.0 ± 3.1	22.7 ± 2.8	0.27	22.7 ± 3.4	23.4 ± 3.6	23.9 ± 3.7	0.002
Grip strength, kg	35.8 ± 9.3	34.0 ± 9.6	35.5 ± 9.1	0.57	23.3 ± 6.8	22.6 ± 6.5	19.9 ± 5.3	< 0.001
6-meter walking time, seconds	5.2 ± 2.2	5.8 ± 2.5	5.6 ± 2.3	0.21	5.5 ± 2.6	5.7 ± 2.6	6.3 ± 2.7	0.03
Chair stand time, seconds	9.6 ± 3.6	10.3 ± 3.8	10.2 ± 3.3	0.30	10.2 ± 4.8	10.5 ± 4.6	11.9 ± 5.1	0.01
Cognitive impairment, %	4.6	6.5	10.6	0.26	3.3	5.6	11.7	0.008

\* Values are the mean ± SD unless indicated otherwise. One-way analysis of variance was used to determine the differences in age, height, weight, body mass index (BMI), grip strength, 6-meter walking time, normal step length, and chair stand time among nonfallers, single fallers, and multiple fallers. Chi-square test was used to determine the differences in prevalence of cognitive impairment among nonfallers, single fallers, and multiple fallers.

(logistic regression analysis;  $P < 0.05$ ); to examine the independent association between radiographic knee OA, knee pain, radiographic LS, and lower back pain in women, we first used multinomial logistic regression analysis with age, BMI, cognitive impairment, radiographic knee OA, knee pain, radiographic LS, and lower back pain as independent variables (Table 5). In this analysis, only lower back pain was independently associated with multiple falls in women. In addition to the above independent variables, we also adjusted for grip strength, 6-meter walking time, and chair stand time, and found that the significant association of lower back pain with multiple falls disappeared, while knee pain was independently associated with multiple falls in women (Table 5).

**DISCUSSION**

The present study is the first large-scale population-based cohort study of the prevalence of single and multiple falls and their association with radiographic knee OA and LS, as well as pain in Japanese men and women. We found

that lower back pain and knee pain were independently associated with multiple falls in women.

There were distinct associations between age strata and single and multiple falls. We found that several factors were associated with multiple falls in women, but no factors were associated with a single fall in women. Previous studies have shown that associations between individual risk factors and a single fall are few in number and weak compared with risk factors for multiple falls (16). A single fall in a year could be accidental and occur due to individual as well as environmental factors, which may partly explain why there were no factors significantly associated with a single fall in our study. In contrast, several factors were associated with multiple falls in the present study, indicating that multiple falls may occur primarily due to individual factors.

In women, the prevalence of multiple falls was higher with increasing age, but in men, the prevalence of multiple falls was lower in subjects ages >60 years, although this could be a random error because of small prevalence, particularly in men. This may be partly explained by the

**Table 4. Comparison of radiographic knee OA and LS, as well as knee and lower back pain, among nonfallers, single fallers, and multiple fallers in men and women\***

	Men				Women			
	Nonfallers	Single fallers	Multiple fallers	P	Nonfallers	Single fallers	Multiple fallers	P
Subjects, no.	508	31	48		881	127	80	
Radiographic knee OA†	77/507 (15.2)	4/31 (12.9)	7/47 (14.9)	0.9417	186/875 (21.3)	31/127 (24.4)	33/79 (41.8)	0.0002
Knee pain‡	97/508 (19.1)	3/31 (9.7)	11/48 (22.9)	0.3268	224/880 (25.5)	37/127 (29.1)	37/80 (46.3)	0.0003
Radiographic LS	190/508 (37.4)	12/31 (38.7)	19/48 (39.6)	0.9490	318/881 (36.1)	45/127 (35.4)	38/80 (47.5)	0.1210
Lower back pain§	99/508 (19.5)	10/31 (32.3)	9/48 (18.8)	0.2203	177/880 (20.1)	31/127 (24.4)	28/80 (35.0)	0.0062

\* Values are the number/total number (percentage) unless otherwise indicated. The chi-square test was used to determine the differences in radiographic findings and pain among nonfallers, single fallers, and multiple fallers. Radiographic knee OA and LS were defined as Kellgren/Lawrence grade 3 or 4. OA = osteoarthritis; LS = lumbar spondylosis.  
 † Nine subjects with total knee arthroplasty were excluded.  
 ‡ One subject with incomplete information regarding knee pain was excluded.  
 § One subject with incomplete information regarding lower back pain was excluded.

**Table 5. Association of radiographic knee OA and LS, as well as knee and lower back pain, with single and multiple falls in women\***

	Crude OR (95% CI)		Adjusted OR <sub>1</sub> (95% CI)†		Adjusted OR <sub>2</sub> (95% CI)‡	
	Single falls	Multiple falls	Single falls	Multiple falls	Single falls	Multiple falls
Radiographic knee OA	1.20 (0.76–1.83)	2.66 (1.64–4.26)	1.07 (0.63–1.82)	1.43 (0.78–2.61)	1.04 (0.60–1.77)	1.31 (0.70–2.43)
Knee pain	1.20 (0.79–1.81)	2.52 (1.58–4.02)	1.00 (0.62–1.61)	1.61 (0.92–2.79)	0.99 (0.60–1.61)	1.87 (1.06–3.28)
Radiographic LS	0.97 (0.65–1.43)	1.60 (1.01–2.54)	0.87 (0.57–1.32)	1.12 (0.68–1.85)	0.88 (0.57–1.33)	1.04 (0.61–1.74)
Lower back pain	1.28 (0.82–1.96)	2.14 (1.30–3.46)	1.34 (0.84–2.08)	1.72 (1.01–2.88)	1.33 (0.84–2.08)	1.58 (0.91–2.70)

\* Radiographic knee osteoarthritis (OA) and lumbar spondylosis (LS) were defined as Kellgren/Lawrence grade 3 or 4. Multinomial logistic regression analysis was used to calculate the odds ratio (OR) and 95% confidence interval (95% CI) compared with nonfallers. Eight subjects with total knee arthroplasty or incomplete information regarding pain were excluded.  
† Adjusted OR<sub>1</sub> was calculated using multinomial logistic regression analysis with age, body mass index, cognitive impairment, radiographic knee OA, knee pain, radiographic LS, and lower back pain as independent variables.  
‡ Adjusted OR<sub>2</sub> was calculated using multinomial logistic regression analysis with grip strength, 6-meter walking time, and chair stand time in addition to the above independent variables.

fact that elderly men generally retire from their occupations at approximately ages 60–70 years; therefore, their environment may change and men may become more sedentary as they age, leading to lower risks of falls. Women, however, must often continue to do household chores even after age 60 years, and their environment may therefore change to a smaller extent than that of men, but their health or muscle strength continues to decline (32), leading to the higher risk of falls.

Our study is the first population-based study to examine the association between knee OA and LS diagnosed by radiography and falls in Japanese men and women. Radiographic knee OA and LS were significantly associated with multiple falls in women, but not in men, although no significant association of radiographic knee OA or LS with falls may be due to the small number of falls in men. The sex differences identified in the association between radiographic knee OA and falls may be partly explained by the weaker quadriceps muscles and increased postural sway associated with knee OA (33,34), both of which are known to be independent risk factors for falls (16,35). In men, muscle strength was higher than that in women in all decades (32), which may obscure the association between radiographic knee OA and falls. LS was also significantly associated with falls in this study, but the OR was lower than that for knee OA. Therefore, falls may be more strongly associated with problems of the lower extremities rather than the trunk.

After adjustment for age, BMI, and cognitive impairment, lower back pain was independently associated with multiple falls, and after adjustment for age, BMI, grip strength, cognitive impairment, 6-meter walking time, and chair stand time, knee pain was independently associated with multiple falls. Given that the significant association of radiographic knee OA and LS with multiple falls disappeared after adjustment, multiple falls may occur due to symptoms such as pain caused by radiographic knee OA or LS rather than radiographic changes in the knee or lumbar spine itself. A previous study also suggested that subjects with knee pain had an increased risk of falls (15). In other words, falls may be preventable when pain is relieved by medical care, even if subjects have radiographic knee OA or LS.

The present study has several limitations. First, this is a

large-scale population-based study with a cross-sectional analysis of baseline data. Therefore, causal relationships could not be determined. The ROAD study is a longitudinal survey; therefore, further progress may help elucidate any causal relationships. Second, our subjects lived in the community, and therefore our findings may not apply to elderly persons residing in institutions. Third, we did not include other weight-bearing OA diseases, such as hip OA, in the analysis, although this disorder also affects falls (36). However, the prevalence of K/L grade 3 or 4 for hip OA is 1.4% and 3.5% in Japanese men and women (37), respectively, which is smaller than that of K/L grade 3 or 4 for knee OA in the present study. Therefore, it is possible that hip OA would not strongly affect the results of the present study. Fourth, the prevalence of fall was comparably small, particularly in men. Therefore, our results regarding the prevalence may include random error, but the present study is the first large-scale, population-based cohort study of the prevalence of falls in Japanese men and women.

In conclusion, the present cross-sectional analysis using a large-scale population from the ROAD study revealed the prevalence and factors associated with falls in men and women. In women, lower back pain and knee pain were significantly associated with multiple falls. Further studies, along with continued longitudinal surveys in the ROAD study, will help elucidate the background of knee OA and LS, and their relationship with falls.

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#### AUTHOR CONTRIBUTIONS

All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version to be submitted for publication. Dr. Muraki had full access to all of the data in the study and takes

responsibility for the integrity of the data and the accuracy of the data analysis.

**Study conception and design.** Muraki, Akune, Oka, En-yo, Yoshida, Nakamura, Kawaguchi, Yoshimura.

**Acquisition of data.** Muraki, Akune, Oka, En-yo, Yoshimura.

**Analysis and interpretation of data.** Muraki, Akune, Oka, Yoshimura.

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# Independent Association of Joint Space Narrowing and Osteophyte Formation at the Knee With Health-Related Quality of Life in Japan

## A Cross-Sectional Study

Shigeyuki Muraki,<sup>1</sup> Hiroyuki Oka,<sup>1</sup> Toru Akune,<sup>1</sup> Yoshio En-yo,<sup>2</sup> Munehito Yoshida,<sup>2</sup> Takao Suzuki,<sup>3</sup> Hideyo Yoshida,<sup>3</sup> Hideaki Ishibashi,<sup>3</sup> Fumiaki Tokimura,<sup>3</sup> Seizo Yamamoto,<sup>3</sup> Kozo Nakamura,<sup>1</sup> Hiroshi Kawaguchi,<sup>1</sup> and Noriko Yoshimura<sup>1</sup>

**Objective.** To clarify the individual associations of joint space narrowing (JSN) and osteophytosis at the knee with quality of life (QOL) in Japanese men and women using a large-scale population-based cohort from the Research on Osteoarthritis Against Disability (ROAD) study.

**Methods.** The associations of minimum joint space width (JSW) and osteophyte area in the medial compartment of the knee with QOL parameters, such as the Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC), were examined. Minimum

JSW and osteophyte area in the medial compartment of the knee were measured using a computer-aided system for the diagnosis of knee osteoarthritis.

**Results.** Of the 3,040 participants in the ROAD study, the present study included 2,039 participants age 40 years or older who completed the questionnaires (741 men and 1,298 women with a mean  $\pm$  SD age of 68.6  $\pm$  10.9 years). Multiple regression analysis after adjustment for age and body mass index showed that minimum JSW was significantly associated with scores on the pain domains of the WOMAC in men and women, while osteophyte area was significantly associated with scores on the physical function domains of the WOMAC in men and women.

**Conclusion.** The findings of this cross-sectional study using a large-scale population from the ROAD study indicate that JSN and osteophytosis are independently associated with QOL.

Knee osteoarthritis (OA) is a major public health issue that causes chronic pain and disability (1–3). The prevalence of radiographic knee OA is high in Japan (4), with 25,300,000 persons age 40 years and older estimated to have radiographic knee OA (5). According to the recent National Livelihood Survey of the Ministry of Health, Labor, and Welfare of Japan, OA is ranked fourth among diseases that cause disabilities that subsequently require support with activities of daily living (6).

Knee OA is characterized by the pathologic features of joint space narrowing (JSN) and osteophytosis, but there is some controversy regarding whether osteophytosis affects knee symptoms or quality of life

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<sup>1</sup>Shigeyuki Muraki, MD, PhD, Hiroyuki Oka, MD, Toru Akune, MD, PhD, Kozo Nakamura, MD, PhD, Hiroshi Kawaguchi, MD, PhD, Noriko Yoshimura, MD, PhD: University of Tokyo, Tokyo, Japan; <sup>2</sup>Yoshio En-yo, MD, Munehito Yoshida, MD, PhD: Wakayama Medical University, Wakayama, Japan; <sup>3</sup>Takao Suzuki, MD, PhD, Hideyo Yoshida, MD, PhD, Hideaki Ishibashi, MD, PhD, Fumiaki Tokimura, MD, PhD, Seizo Yamamoto, MD, PhD: Tokyo Metropolitan Institute of Gerontology, Tokyo, Japan.

Patents for the Knee Osteoarthritis Computer-Aided Diagnosis (KOACAD) system are held by the University of Tokyo.

Address correspondence to Shigeyuki Muraki, MD, PhD, Department of Clinical Motor System Medicine, 22nd Century Medical and Research Center, Faculty of Medicine, University of Tokyo, Hongo 7-3-1, Bunkyo-ku, Tokyo 113-8655, Japan. E-mail: murakis-ort@h.u-tokyo.ac.jp.

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(QOL). Nevertheless, researchers examining the hand and hip have argued that the separate radiographic features should be recorded and may be more meaningful than overall composite scores such as the Kellgren/Lawrence (K/L) scale (7). Furthermore, a previous study showed that osteophytes were well correlated with knee symptoms and performed better as a primary diagnostic feature than JSN in cross-sectional epidemiologic studies of knee OA (8). However, most conventional systems for grading radiographic severity have consisted of categorical grades, such as the K/L scale (9), which is unable to individually assess JSN and osteophytosis. Several studies have shown that knee OA had a strong effect on QOL (10–13), but in those studies, knee OA was defined by categorical grades such as the K/L grade or the American College of Rheumatology grade (14), total knee arthroplasty, and self-questionnaire.

A radiographic atlas of individual features published by the OA Research Society International in 1995 (15) and revised in 2007 (16) allows JSN and osteophyte formation to be evaluated separately. However, the grading is still limited in reproducibility and sensitivity due to the subjective judgment of individual observers and the categorical classification into 4 grades (0–3). To overcome this problem, joint space width (JSW) and osteophyte area should be evaluated using a fully automatic system. To the best of our knowledge, no population-based studies have been conducted to separately measure JSW or osteophyte area in order to clarify the associations of JSN with QOL and of osteophytosis with QOL, despite the fact that the associations between these major features of knee OA and QOL are likely to be different.

Differences between the sexes have also been observed in knee OA. The prevalence of knee OA is higher in women than in men (4), and the association of knee pain with knee OA also differs by sex (4). Thus, the impact of JSN on QOL and of osteophytosis on QOL may also differ between the sexes. However, to the best of our knowledge, no population-based studies have been conducted to assess the associations of JSN and osteophytosis with QOL in men and women separately.

The objective of this study was therefore to separately clarify the association between JSN and QOL and the association between osteophytosis and QOL in Japanese men and women in a large-scale, population-based cohort from the Research on Osteoarthritis Against Disability (ROAD) study. A fully automatic system was used to measure JSW and osteophyte area. QOL was measured using disease-specific scales for

knee OA, such as the Western Ontario and McMaster Universities OA Index (WOMAC).

## SUBJECTS AND METHODS

**Participants.** The ROAD study is a nationwide prospective study designed to establish epidemiologic indexes for the evaluation of clinical evidence for the development of a disease-modifying treatment for bone and joint diseases (with OA and osteoporosis as the representative bone and joint diseases). It consists of population-based cohorts in several communities in Japan. The ROAD study has been described in detail previously (4,5,17). To date, we have completed the creation of a baseline database including clinical and genetic information for 3,040 participants (1,061 men and 1,979 women) ranging in age from 23 to 95 years (mean 70.6 years), who were recruited from resident registration listings in 3 communities: an urban region in Itabashi, Tokyo; a mountainous region in Hidakagawa, Wakayama; and a seacoast region in Taiji, Wakayama. All participants provided written informed consent, and the study was conducted with the approval of the ethics committees of the University of Tokyo and the Tokyo Metropolitan Institute of Gerontology. Height, weight, and body mass index (BMI) (weight [kg]/height [m<sup>2</sup>]) were measured. Among the 2,995 participants in the ROAD study who were age 40 years or older, 2,222 (74.2%) completed the WOMAC. The 2,222 participants who completed the WOMAC were younger than those who did not (mean age 68.9 years for those who completed the WOMAC versus 75.9 years for those who did not;  $P < 0.0001$ ). These 2,222 participants were also less likely to be women (63.8% of those who completed the WOMAC versus 68.3% of those who did not;  $P < 0.05$ ), and were less likely to have knee OA than the subjects who did not complete the WOMAC (54.1% versus 60.4%;  $P < 0.01$ ). Of the 2,222 subjects, 183 subjects with lateral knee OA or total knee arthroplasty were excluded. Therefore, a total of 2,039 participants (741 men and 1,298 women) age 40 years or older (mean  $\pm$  SD 68.6  $\pm$  10.9 years) who had completed the WOMAC were included in the present study.

**Radiographic assessment.** Radiographic examinations of both knees of all participants, using an anteroposterior view with weight-bearing and foot map positioning, were performed by experienced radiologic technicians. The beam was positioned parallel to the floor with no angle and aimed at the joint space. To visualize the joint space properly and to center the patella over the lower end of the femur, we used fluoroscopic guidance with an anteroposterior x-ray beam, and the images were downloaded into Digital Imaging and Communication in Medicine (DICOM) format files. Knee radiographs were read by a single experienced orthopedist (SM), who was blinded with regard to participant clinical status, using the K/L radiographic atlas for overall knee radiographic grades (9), and knee OA was defined as a K/L grade of 2 or severe. Minimum JSW in the medial compartment and osteophyte area at the medial tibia were measured by the Knee Osteoarthritis Computer-Aided Diagnosis (KOACAD) system, and for each subject the knee with the lower minimum JSW was defined as the designated knee. The KOACAD system is a fully automatic system that can quantify the major features of knee OA

on standard radiographs and allows for objective, accurate, and simple assessment of the structural severity of knee OA in general clinical practice. This system was programmed to measure minimum JSW in the medial and lateral compartments and osteophyte area at the medial tibia using digitized knee radiographs. The KOACAD system has been described in detail previously (18). The KOACAD system was applied to the DICOM data by the experienced orthopedist who developed this system (HO), and the reliability of measurement is good (18). Lateral knee OA was defined as a K/L grade of  $\geq 2$  with lower lateral minimum JSW than medial minimum JSW.

**QOL instrument.** To carry out the QOL assessment, we used the WOMAC. The WOMAC, a 24-item OA-specific index, consists of 3 domains: pain, stiffness, and physical function. Each of these 24 items is graded on either a 5-point Likert scale (scores of 0–4) or a 100-mm visual analog scale (19,20). In the present study, we used the Likert scale (version LK 3.0). The domain score ranges from 0 to 20 for pain, 0 to 8 for stiffness, and 0 to 68 for physical function. Japanese versions of the WOMAC have been validated (21).

**Statistical analysis.** Differences in age, height, weight, BMI, minimum JSW, osteophyte area, and QOL measurements between men and women were examined using Student's unpaired *t*-test. Associations of minimum JSW and osteophyte area with scores on the pain and physical function domains of the WOMAC were determined using multiple regression analysis without adjustment. To assess independent associations of minimum JSW and osteophyte area with QOL, multiple regression analysis was used with age, BMI, minimum JSW, and osteophyte area as independent variables. Data analysis was performed using SAS, version 9.0.

## RESULTS

The characteristics of the 2,039 participants in the present study are shown in Table 1. The minimum JSW was significantly lower and osteophyte area was significantly higher in women than in men. Scores on all domains of the WOMAC were significantly lower (indicating better status) in men than in women. Osteophyte

area was only moderately associated with minimum JSW on linear regression analysis ( $R^2 = 0.173$ ,  $P < 0.05$ ).

Linear regression analysis without adjustment showed that minimum JSW and osteophyte area were significantly associated with scores on the pain and physical function domains of the WOMAC in the overall population as well as in men and women analyzed separately (Table 2). To determine the independent associations of minimum JSW and osteophyte area with scores on the pain and physical function domains of the WOMAC, we used multiple regression analysis with age, sex, BMI, minimum JSW, and osteophyte area as independent variables in the overall population (Table 2). Minimum JSW and osteophyte area were independently associated with scores on the pain and physical function domains of the WOMAC ( $\beta$  coefficients  $-0.16$  and  $0.11$  for the association of pain domain score with minimum JSW and osteophyte area, respectively, and  $\beta$  coefficients  $-0.13$  and  $0.16$  for the association of physical function domain score with minimum JSW and osteophyte area, respectively).

When men and women were analyzed separately (Table 2), in men, minimum JSW was independently associated with the pain domain scores ( $\beta$  coefficient  $-0.13$ ), but not with the physical function domain scores ( $\beta$  coefficient  $0.07$ ) of the WOMAC, while osteophyte area was independently associated with the physical function domain scores ( $\beta$  coefficient  $0.14$ ), but not with the pain domain scores ( $\beta$  coefficient  $-0.07$ ) of the WOMAC. In women, both minimum JSW and osteophyte area were independently associated with scores on the pain and physical function domains of the WOMAC, and the absolute values of the beta values for minimum JSW for scores on the pain domains of the WOMAC

Table 1. Characteristics of the subjects\*

	Overall population (n = 2,039)	Men (n = 741)	Women (n = 1,298)
Age, years	68.6 $\pm$ 10.9	69.7 $\pm$ 10.5	67.9 $\pm$ 11.2†
Height, cm	154.7 $\pm$ 8.9	162.8 $\pm$ 6.5	150.1 $\pm$ 6.5†
Weight, kg	55.1 $\pm$ 10.4	61.4 $\pm$ 10.2	51.5 $\pm$ 8.6†
BMI, kg/m <sup>2</sup>	22.9 $\pm$ 3.3	23.1 $\pm$ 3.1	22.8 $\pm$ 3.4†
Minimum JSW, mm	2.61 $\pm$ 0.98	2.97 $\pm$ 0.92	2.40 $\pm$ 0.96†
Osteophyte area, mm <sup>2</sup>	2.99 $\pm$ 8.68	1.28 $\pm$ 4.46	3.98 $\pm$ 10.25†
Radiographic knee OA, %	50.2	39.0	56.8
WOMAC			
Pain	1.35 $\pm$ 2.42	1.10 $\pm$ 2.12	1.50 $\pm$ 2.57†
Stiffness	0.72 $\pm$ 1.25	0.63 $\pm$ 1.10	0.77 $\pm$ 1.33†
Function	3.99 $\pm$ 7.84	3.24 $\pm$ 6.69	4.42 $\pm$ 8.41†

\* Except where indicated otherwise, values are the mean  $\pm$  SD. BMI = body mass index; JSW = joint space width; OA = osteoarthritis; WOMAC = Western Ontario and McMaster Universities OA Index. †  $P < 0.05$  versus men, by Student's unpaired *t*-test.

**Table 2.** Associations of minimum JSW and osteophyte area with WOMAC domain scores\*

	Pain				Physical function			
	Crude regression coefficient (95% CI)	P	Adjusted regression coefficient (95% CI)†	P	Crude regression coefficient (95% CI)	P	Adjusted regression coefficient (95% CI)†	P
Overall population								
Minimum JSW	-0.71 (-0.81, -0.60)	<0.0001	-0.37 (-0.48, -0.25)	<0.0001	-2.33 (-2.66, -1.99)	<0.0001	-0.97 (-1.34, -0.59)	<0.0001
Osteophyte area	0.07 (0.05, 0.08)	<0.0001	0.03 (0.02, 0.04)	<0.0001	0.25 (0.21, 0.29)	<0.0001	0.14 (0.10, 0.18)	<0.0001
Men								
Minimum JSW	-0.47 (-0.64, -0.31)	<0.0001	-0.29 (-0.47, -0.11)	0.002	-1.34 (-1.86, -0.82)	<0.0001	-0.48 (-1.04, 0.08)	0.10
Osteophyte area	0.07 (0.04, 0.11)	<0.0001	0.03 (-0.005, 0.07)	0.09	0.30 (0.19, 0.41)	<0.0001	0.20 (0.09, 0.32)	0.0005
Women								
Minimum JSW	-0.83 (-0.97, -0.69)	<0.0001	-0.41 (-0.57, -0.25)	<0.0001	-2.89 (-3.35, -2.43)	<0.0001	-1.22 (-1.72, -0.72)	<0.0001
Osteophyte area	0.06 (0.05, 0.08)	<0.0001	0.03 (0.01, 0.04)	0.0001	0.24 (0.20, 0.29)	<0.0001	0.12 (0.08, 0.17)	<0.0001

\* WOMAC = Western Ontario and McMaster Universities Osteoarthritis Index; 95% CI = 95% confidence interval.

† Calculated by multiple regression analysis with age, sex, body mass index, minimum joint space width (JSW), and osteophyte area as the independent variables in the overall population and with age, body mass index, minimum JSW, and osteophyte area as the independent variables in the groups of men and women only.

were larger than those for osteophyte area (-0.15 and 0.11, respectively), while the absolute values of the beta values for minimum JSW for scores on the physical

function domains of the WOMAC were smaller than those for osteophyte area (-0.14 and 0.15, respectively). When the analysis was restricted to the partici-

**Table 3.** Associations of minimum JSW and osteophyte area with WOMAC domain scores in the subjects with knee OA\*

	Pain				Physical function			
	Crude regression coefficient (95% CI)	P	Adjusted regression coefficient (95% CI)†	P	Crude regression coefficient (95% CI)	P	Adjusted regression coefficient (95% CI)†	P
Overall population								
Minimum JSW	-0.81 (-0.97, -0.65)	<0.0001	-0.51 (-0.69, -0.33)	<0.0001	-2.77 (-3.32, -2.22)	<0.0001	-1.46 (-2.05, -0.87)	<0.0001
Osteophyte area	0.06 (0.04, 0.07)	<0.0001	0.03 (0.01, 0.04)	0.0007	0.22 (0.18, 0.27)	<0.0001	0.12 (0.07, 0.17)	<0.0001
Men								
Minimum JSW	-0.59 (-0.86, -0.31)	<0.0001	-0.42 (-0.72, -0.11)	0.009	-1.95 (-2.81, -1.08)	<0.0001	-0.97 (-1.97, -0.01)	0.05
Osteophyte area	0.07 (0.02, 0.11)	0.003	0.02 (-0.02, 0.07)	0.40	0.34 (0.21, 0.48)	<0.0001	0.24 (0.10, 0.39)	0.001
Women								
Minimum JSW	-0.89 (-1.09, -0.68)	<0.0001	-0.56 (-0.78, -0.34)	<0.0001	-3.00 (-3.71, -2.29)	<0.0001	-1.61 (-2.35, -0.88)	<0.0001
Osteophyte area	0.05 (0.04, 0.07)	<0.0001	0.03 (0.01, 0.04)	0.002	0.20 (0.15, 0.26)	<0.0001	0.11 (0.05, 0.16)	0.0001

\* Knee osteoarthritis (OA) was defined as a Kellgren/Lawrence grade of  $\geq 2$ . WOMAC = Western Ontario and McMaster Universities OA Index; 95% CI = 95% confidence interval.

† Calculated by multiple regression analysis with age, sex, body mass index, minimum joint space width (JSW), and osteophyte area as the independent variables in the overall population and with age, body mass index, minimum JSW, and osteophyte area as the independent variables in the groups of men and women only.

pants with knee OA, the results were almost the same (Table 3). In men with knee OA, minimum JSW was independently associated with pain domain scores ( $\beta$  coefficient  $-0.17$ ), but not with physical function domain scores ( $\beta$  coefficient  $0.05$ ). In women with knee OA, both minimum JSW and osteophyte area were independently associated with physical function domain scores, but the beta value for minimum JSW for physical function domain scores was smaller than that for osteophyte area ( $-0.12$  and  $0.20$ , respectively).

## DISCUSSION

This is the first study to separately examine the associations of JSN and osteophytosis with QOL, measured by a disease-specific scale such as WOMAC, using a large-scale population-based Japanese cohort. In addition, JSN and osteophytosis were estimated not by categorical grade but by continuous values such as minimum JSW and osteophyte area at the knee. In the present study, JSN as well as osteophytosis was independently associated with QOL.

The present study showed that both JSN and osteophytosis reduce QOL. Osteophytosis appears to begin with the activation of periosteal layers, with initial generation of chondrocytes and subsequent calcification to real osteophytes. The process is probably an adaptive reaction of the joint in order to cope with joint instability, and thus osteophyte area may indicate the severity of joint instability (22), which might lead to loss of QOL. When men and women were analyzed separately, minimum JSW was significantly associated with scores on the WOMAC pain domain but not the WOMAC physical function domain in men, while osteophyte area was associated with scores on the physical function domain but not the pain domain. According to the methodology of the WOMAC, pain domains estimate the severity of pain, indicating that JSN may be strongly associated with pain. In contrast, physical function domains assess difficulties in activities of daily living, indicating that osteophytosis may be mainly associated with activities of daily living, particularly in men.

Our findings also indicated differences between the sexes in the associations of JSN and osteophytosis with QOL. Minimum JSW was significantly associated with scores on the physical function domains of the WOMAC in women, but not in men. Similarly, osteophyte area was associated with scores on the pain domains of the WOMAC in women, but not in men. These differences may indicate that JSN and osteophytosis were more strongly associated with loss of QOL

in women than in men. Our previous study also showed that the odds ratio of knee pain for K/L grade 3 or 4 knee OA was approximately twice as high in women as in men (4). This may be partly explained by the lower muscle mass in women than in men. Previous reports have shown that muscle mass is also associated with QOL (23,24). In men, muscular strength may obscure the associations of JSN and osteophytosis with QOL loss; thus, these were not associated with some QOL parameters in men.

The present study has several limitations. First, this is a large-scale, population-based study, with a cross-sectional study of baseline data. Thus, causal relationships could not be determined. The ROAD study is a longitudinal survey, so further progress may help elucidate any causal relationships. Second, we did not include other weight-bearing forms of OA, such as hip OA, in the analysis, although this disorder may also affect QOL. However, the prevalence of K/L grade 3 or 4 hip OA is 1.4% and 3.5% in Japanese men and women (25), respectively, which is lower than the prevalence of K/L grade 3 or 4 knee OA (13.5% and 24.6% in Japanese men and women, respectively) (4). Thus, it is possible that including hip OA would not strongly affect the results of the present study. Third, the QOL questionnaire was completed by 74.2% of all participants age 40 years or older in the ROAD study. Participants who completed the questionnaire were younger and more likely to have knee OA than the participants who did not complete the questionnaire, and thus the participants included in this study may have had better QOL than those who did not complete the questionnaire, and our results may have overestimated QOL. Fourth, although osteophytes may be even more pronounced in the contralateral tibiofemoral compartment (26), at present the KOACAD system can only measure medial osteophytes at the tibia. We are now developing the KOACAD system to measure osteophytes at other sites; thus, we may be able to clarify the association between osteophytes at other sites and QOL in the near future.

In conclusion, the present cross-sectional study using a large-scale population from the ROAD study revealed that JSN and osteophytosis are independently associated with QOL. Further studies, along with continued longitudinal surveys in the ROAD study, will help clarify the mechanisms of JSN and osteophytosis at the knee, and their relationship with QOL.

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#### AUTHOR CONTRIBUTIONS

All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version to be published. Dr. Muraki had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

**Study conception and design.** Muraki, Oka, Akune, En-yo, M. Yoshida, Suzuki, H. Yoshida, Ishibashi, Tokimura, Yamamoto, Nakamura, Kawaguchi, Yoshimura.

**Acquisition of data.** Muraki, Oka, Akune, En-yo, Yoshimura.

**Analysis and interpretation of data.** Muraki, Oka, Akune, Yoshimura.

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## Association of vitamin D status with knee pain and radiographic knee osteoarthritis

S. Muraki†\*, E. Dennison‡, K. Jameson‡, B.J. Boucher§, T. Akune†, N. Yoshimura||, A. Judge‡¶, N.K. Arden‡¶, K. Javaid‡¶, C. Cooper‡¶

†Department of Clinical Motor System Medicine, 22nd Century Medical & Research Center, Faculty of Medicine, University of Tokyo, Tokyo, Japan

‡MRC Lifecourse Epidemiology Unit, University of Southampton, Southampton General Hospital, UK

§Centre for Diabetes, Bart's & The London School of Medicine and Dentistry, Queen Mary University of London, UK

||Department of Joint Disease Research, 22nd Century Medical & Research Center, Faculty of Medicine, University of Tokyo, Tokyo, Japan

¶NIHR Oxford Biomedical Research Unit, University of Oxford, Nuffield Orthopaedic Centre, Oxford, UK

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### SUMMARY

**Objective:** The objective of the present study was to explore the association of serum vitamin D concentration and polymorphism in the vitamin D receptor (VDR), with knee pain and radiographic knee osteoarthritis (OA) among men and women in a large population-based UK cohort study.

**Methods:** Seven hundred and eighty-seven participants in the Hertfordshire Cohort Study (399 men, 388 women; mean age  $65.6 \pm 2.7$  years) underwent a questionnaire on knee pain and radiographic knee examination. This study examined the association of Fok1, Cdx2 and Apa1 polymorphism in the gene for the VDR and serum 25(OH)D concentration with knee pain and radiographic knee OA by a generalized estimating equations population averaged logistic regression analysis in the Hertfordshire Cohort Study. **Results:** There were no associations of Fok1, Cdx2 and Apa1 polymorphisms of the VDR with knee OA except for Aa for Apa1 compared with AA [Odds ratio (OR) 0.59, 95% confidence interval (CI) 0.36–0.95,  $P = 0.031$ ]. While, ff for Fok1 (OR 1.60, 95% CI 1.07–2.39,  $P = 0.022$ ) and AA for Cdx2 polymorphism (OR 2.21, 95% CI 1.07–4.56,  $P = 0.032$ ) was significantly associated with higher prevalence of knee pain compared with FF for Fok1 and GG for Cdx2, respectively. None of these are statistically significant after adjusting for the three polymorphisms tested. 25(OH)D level was not significantly associated with radiographic knee OA, while, low tertile of 25(OH)D level tended to be associated with knee pain compared with high tertile of 25(OH)D level.

**Conclusion:** The present cross-sectional study using a large-scale population from the Hertfordshire Cohort study indicated that vitamin D may be associated with pain rather than radiographic change, but the evidence for an association between vitamin D genetic variation and pain in knee OA is very weak in the present study. Further replication of our results will be required to elucidate the association of vitamin D and knee OA.

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### Introduction

Knee osteoarthritis (OA) is a major public health issue that causes chronic pain and disability<sup>1–3</sup>, although at present the pathogenesis of this condition remains largely unknown. Several environmental factors have been associated with OA, including obesity<sup>4–6</sup>, previous injury<sup>7</sup>, knee-bending occupations<sup>8,9</sup>, and

other metabolic factors<sup>10,11</sup>. A previous population-based UK study of twins has also demonstrated a clear genetic influence on radiologic knee OA in women, with up to 65% of the variance being explained by genetic factors<sup>12</sup>.

Vitamin D has been shown to stimulate synthesis of proteoglycan by mature articular cartilage *in vitro*<sup>13</sup>, and this suggests that vitamin D may directly affect articular cartilage metabolism. Vitamin D receptor (VDR) is found in many types of tissues, including chondrocytes<sup>14,15</sup>. A previous study showed that VDR gene polymorphism was associated with bone<sup>16</sup>, although it is still controversial<sup>17</sup>. The relationship between osteoporosis and OA suggests that VDR gene polymorphisms may be associated with both diseases<sup>18</sup>. However, the association of VDR gene polymorphisms with knee OA is

\* Address correspondence and reprint requests to: S. Muraki, Department of Clinical Motor System Medicine, 22nd Century Medical & Research Center, Faculty of Medicine, University of Tokyo, Hongo 7-3-1, Bunkyo-ku, Tokyo 113-8655, Japan. Tel: 81-3-5800-9178; Fax: 81-3-5800-9179.

E-mail address: murakis-ort@h.u-tokyo.ac.jp (S. Muraki).

controversial<sup>19–23</sup>. This may be partly due to different races, differences in environmental factors related to vitamin D metabolism, or the presence of other genetic factors that influence VDR function.

The association of vitamin D level with knee OA is also controversial<sup>24–28</sup>. In previous studies, McAlindon suggested that low serum levels of vitamin D were associated with progression of knee OA<sup>24</sup>. A recent study has also shown that serum 25(OH)D levels were associated with decreased knee cartilage loss<sup>28</sup>, but Hunter *et al.* found that there was no significant association between vitamin D levels and knee osteophytes after adjusting for age, body mass index (BMI) and relatedness<sup>25</sup>. The Framingham study also found no association of vitamin D levels with knee OA worsening<sup>26</sup>. This may be partly explained by VDR gene polymorphism, because vitamin D exerts its endocrine and autocrine/paracrine local effects upon binding to and activating its intracellular receptor VDR. In other words, the association of vitamin D level with knee OA may be different by VDR gene polymorphisms, but, to the best of our knowledge, there were no studies investigating the association of vitamin D level with knee OA by VDR gene polymorphisms.

The principal clinical symptom of knee OA is pain<sup>29</sup>, but the correlation between pain and radiographic severity is inconsistent<sup>4,30–32</sup>. Fewer studies have addressed factors which might influence knee pain<sup>32–35</sup>; among these, older age, female gender, and physically demanding work, have all been proposed<sup>30–33</sup>. Previous studies, however, have not addressed the role of vitamin D status or fixed genetic variation in the VDR.

The objective of this study was to clarify the association of VDR gene polymorphism with knee pain and radiographic knee OA among men and women in the general population, as well as to examine the association between circulating vitamin D concentration and these indices of OA.

## Subjects and methods

### Subjects

The Hertfordshire Cohort Study is a population-based cohort study in the UK. Details of the study design have been published previously<sup>36</sup>, thus, a brief summary is provided here. The selection procedure was as follows: using the National Health Service Central Registry at Southport, and Hertfordshire Family Health Service Association, we traced men and women who were born during 1931–1939 in Hertfordshire, and still lived there during the period 1998–2003. After obtaining written permission from each subject's general practitioner (GP), we approached each person by letter, asking him or her if they would be willing to be contacted by one of our research nurses. If subjects agreed, a research nurse performed a home visit and administered a structured questionnaire. This included information on socioeconomic status, medical history, drug history, cigarette smoking, alcohol consumption, and reproductive variables in women.

At a subsequent clinic, height was measured to the nearest 0.1 cm using a Harpenden pocket stadiometer (Chasmors Ltd, London, UK) and weight to the nearest 0.1 kg on a SECA floor scale (Chasmors Ltd). Fasting venous whole-blood samples were taken at this clinic visit. Eligible subjects were then invited to book a return visit for knee radiography. Weightbearing anteroposterior and lateral semiflexed radiographs of both knees were taken at the same hospital using the same radiographic equipment; a standard tube to film distance of 100 cm was used. Radiographs were performed at a median duration of 6 months [interquartile range (IQR) 4.8–7.2] after the clinic visit. Radiographs were graded at the tibiofemoral joints using the Kellgren Lawrence (KL) grade<sup>37</sup>. One trained reader graded the radiographs; KL grade  $\geq 2$  was the threshold for a definition of knee OA. Subjects were also asked

“Have you had any pain in or around your right knee on most days in the last month?” and “Have you had any pain in or around your left knee on most days in the last month?” Knee pain reported in this way was defined as having knee pain. A total of 498 men and 468 women completed a home questionnaire, attended clinic, and underwent knee radiography.

A fasting morning blood sample was obtained from all subjects at the first clinic visit, and the serum separated and stored at  $-70^{\circ}\text{C}$ . 25(OH)vitamin D was assayed using a DiaSorin Liason automated chemiluminescent assay with equal specificity for both D2 and D3 (coefficient of variation for vitamin D across the assays was 10–12% for within batch and 10–15% between batch).

Genomic DNA was extracted from whole-blood samples according to standard procedures. VDR genotype was determined by the polymerase chain reaction (PCR) and restriction fragment length polymorphism (RFLP) analysis, and three VDR polymorphic sites (Fok1, Cdx2 and Apa1) were analyzed.

Ethical permission for the study was granted by the East and North Hertfordshire Ethical Committees. All participants gave written informed consent.

### Statistical analysis

To assess gene polymorphism effects on radiographic knee OA and knee pain, indicator variables were created for Fok1 (FF and ff), Cdx2 (GA and AA) and Apa1 (AA and aa) polymorphism. As both knees have a pain score and a radiographic grade, a generalized estimating equations (GEE) population averaged logistic regression model was used to adjust for clustering of knees within patients. To examine 25(OH)D levels and their association with knee OA and knee pain, we classified subjects into three categories; high tertile ( $>51.5$  nmol/l), middle tertile (35.5–51.5 nmol/l) and low tertile ( $<35.5$  nmol/l). A GEE population averaged logistic regression analysis was used to determine the association of vitamin D level with knee OA and knee pain with and without adjustment for age, gender, BMI, season of the clinic visit and KL grade. To decide whether statically significant associations between VDR polymorphisms and knee outcomes are noteworthy, we used Wacholder's method to calculate the False Positive Report Probability (FPRP) [Wacholder in JNCI 2004]. Data analyses were performed using SAS version 9.0 (SAS Institute, Cary, NC, USA) and Stata version 11.2 (Stata, College Station, TX, USA).

## Results

Of 984 subjects, 170 (17.3%) provided incomplete pain questionnaires. A further 19 (1.9%) lacked genotypic information. We also excluded eight subjects with total knee arthroplasty, leaving 787 (399 males and 388 females) participants in this analysis. Comparison between the 787 subjects with complete information and those without complete information revealed no statistically significant differences in mean age (responders 65.6 years, nonresponders 65.7 years;  $P=0.66$ ), sex (responders 80.3% women, nonresponders 79.8% women;  $P=0.87$ ), BMI (responders 27.0, nonresponders 27.4;  $P=0.19$ ) or prevalence of knee OA (responders 15.2% women, nonresponders 16.3% women;  $P=0.74$ ). The characteristics of these participants are shown in Table I. The men were slightly younger than the women, and they had a lower mean BMI; serum vitamin D concentration was also significantly higher among men than women. There were no significant differences in mean values [IQR] of 25(OH)D concentration (nmol/l) among VDR gene polymorphisms of Fok1 [FF 45.5 (30.0–56.0), Ff 47.1 (31.5–56.8), ff 51.3 (31.0–69.0)], Cdx2 [GG 45.9 (30.6–56.0), AG 47.5 (31.2–60.2), AA 54.1 (36.7–68.0)] and Apa1 [AA 48.4 (32.9–61.3), Aa 47.5 (30.0–59.1), aa 42.7 (31.0–50.9)]. There were no significant

**Table I**  
Characteristics of participants

	Overall	Men	Women	P-value
Number of subjects	787	399	388	
Age, years	65.6 (2.7)	64.8 (2.6)	66.4 (2.6)	<0.001
BMI, kg/m <sup>2</sup>	27.0 (4.3)	26.8 (3.6)	27.2 (4.9)	0.22
25(OH)D level, nmol/l* means, (IQR)	42.5 (30.8, 57.3)	44.4 (34.7, 64.2)	41.0 (28.3, 54.1)	<0.001
Radiographic knee OA, n, (%)	120 (15.3)	70 (17.5)	50 (12.9)	0.069
Knee pain, n, (%)	309 (39.3)	147 (36.8)	162 (41.8)	0.16

Except where indicated otherwise, values represent means (standard deviation). The differences in age, BMI and 25(OH)D level between men and women were examined by the non-paired Student's *t*-test. The differences in prevalence of radiographic knee OA and knee pain between men and women were examined by chi square test.

\* Of 787 subjects, 25(OH)D was measured in 683 subjects.

differences in the prevalence of radiographic knee OA and knee pain between genders. Of 120 subjects with radiographic knee OA, 79 (65.8%) had knee pain, while, of 667 subjects without radiographic knee OA, 230 (34.5%) had knee pain. Knee pain was significantly associated with radiographic knee OA after adjustment for age, gender and BMI [Odds ratio (OR); 3.03, 95% confidence interval (CI); 1.98–4.68].

We examined the association of VDR gene polymorphisms and radiographic knee OA (Table II). There were no associations of Fok1, Cdx2 and Apa1 polymorphisms of the VDR with knee OA except for Aa for Apa1 compared with AA after adjustment for age, gender and BMI, and FPRP values were low for association of Apa1 (Aa) on radiographic knee OA suggesting this association may be noteworthy. We also examined the associations of the alleles with knee OA. *f* for Fok1 tended to associate with higher prevalence of knee OA than *F* ( $P=0.06$ ). The alleles for Cdx2 and Apa1 were not significantly associated with knee OA ( $P=0.94$  and  $0.64$ ).

We also examined the association of VDR gene polymorphisms and knee pain (Table III). Unlike radiographic knee OA, Fok1 and Cdx2 polymorphism was significantly associated with prevalence of knee pain after adjustment for age, gender BMI and KL grade, and FPRP values were low for association of Fok1 (*ff*) for knee pain, suggesting this association may be noteworthy. There were no associations of Apa1 polymorphisms with knee pain. When analyzed in men and women separately, Fok1 polymorphism was significantly associated with knee pain after adjustment for age, BMI and KL grade in women (Ff: OR; 1.17, 95% CI; 0.75–1.81,

$P=0.486$ , ff: OR; 2.46, 95% CI; 1.38–4.39,  $P=0.002$ , compared with FF), while, not in men (Ff: OR; 1.10, 95% CI; 0.71–1.73,  $P=0.649$ , ff: OR; 1.01, 95% CI; 0.58–1.76,  $P=0.98$ , compared with FF). We also examined the associations of the alleles with knee pain. *f* for Fok1 had significantly associated with higher prevalence than *F* ( $P=0.01$ ). The alleles for Cdx2 and Apa1 were not significantly associated with knee pain ( $P=0.49$  and  $0.64$ , respectively).

We next examined the association of 25(OH)D level and knee OA (Table IV). GEE logistic regression analysis showed that 25(OH)D level was not significantly associated with radiographic knee OA. For knee pain effect of vitamin D level was non-linear, so we classified subjects into three groups; high tertile (>51.5 nmol/l,  $n=225$ ), middle tertile (35.5–51.5 nmol/l,  $n=229$ ) and low tertile (<35.5 nmol/l,  $n=229$ ); low tertile of 25(OH)D level tended to be associated with knee pain compared with high tertile of 25(OH)D level after adjustment for age, gender, BMI, season of the clinic visit and KL grade (Table IV).

## Discussion

This is the first study to examine the association of radiographic knee OA and knee pain with vitamin D level and VDR gene polymorphism at the same time. A Fok1 polymorphism of the VDR was significantly associated with radiographic knee OA and knee pain. There were no associations between radiographic knee OA and 25(OH)D level, while 25(OH)D level tended to be associated with knee pain.

The association of VDR gene polymorphism with OA is controversial<sup>19–23</sup>. In previous studies, a nested case-control study in Britain showed the 'T' allele was associated with knee OA in women<sup>19</sup>. The Rotterdam Study showed that the 'bAT' haplotype was associated with reduced prevalence of OA<sup>20</sup>. While, The Framingham study found no evidence for an association of the VDR gene with knee OA<sup>23</sup>. In a case-control study in Japan, there was also no significant association between VDR gene polymorphism and knee OA<sup>21</sup>, although cases were sampled from hospital attenders in the study and controls did not undergo X-rays, causing the inevitable selection bias to occur. This inconsistency may also be due to differences in the relative importance of this gene in different races, differences in environmental factors related to vitamin D metabolism, or the presence of other genetic factors that influence VDR function. Further, the association of genetic factors with knee OA is diminishing later in life due to the effects of lifestyle factors, thus it may be difficult to find out their association in the elderly. In the present study, a Fok1 polymorphism of the VDR was significantly associated with radiographic knee OA. Vitamin D has been shown to stimulate synthesis of proteoglycan by mature

**Table II**  
Association of VDR gene polymorphisms and radiographic knee OA

	Total	Number (%) with knee OA	Crude OR (95% CI)	P-value	Adjusted OR (95% CI)*	P-value	Power†	FPRP prior probability	
								0.1	0.01
Fok1									
FF	328	39 (11.9)	1.00		1.00				
Ff	333	60 (18.0)	1.50 (0.96, 2.34)	0.072	1.54 (0.96, 2.46)	0.071	0.47	0.58	0.94
ff	108	18 (16.7)	1.38 (0.75, 2.57)	0.301	1.56 (0.82, 2.94)	0.173	0.27	0.85	0.98
Cdx2									
GG	491	75 (15.3)	1.00		1.00				
AG	248	37 (14.9)	1.03 (0.66, 1.59)	0.903	0.94 (0.60, 1.48)	0.781	0.49	0.93	0.99
AA	29	5 (17.2)	1.30 (0.48, 3.57)	0.605	1.09 (0.43, 2.74)	0.858	0.11	0.99	1.00
Apa1									
AA	213	36 (16.9)	1.00		1.00				
Aa	388	51 (13.1)	0.64 (0.40, 1.03)	0.068	0.59 (0.36, 0.95)	0.031	0.39	0.42	0.89
aa	166	31 (18.7)	1.12 (0.65, 1.91)	0.687	1.04 (0.60, 1.81)	0.884	0.28	0.97	1.00

Of 787 subjects, genotyping was completed for 769, 768 and 767 with Fok1, Csk2 and Apa1 polymorphism of the VDR, respectively.

\* As both knees have a radiographic grade, GEE population averaged logistic regression analysis after adjustment for age, gender and BMI was used to calculate adjusted OR.

† To detect an OR of 1.5, we are looking for a difference in proportions of 15.3% vs 21.3% for radiographic knee OA.