

The development and validation of the Satisfaction Questionnaire for Osteoporosis Prevention in Malaysia

Li Shean Toh¹
 Pauline Siew Mei Lai²
 David Bin-Chia Wu³
 Kok Thong Wong¹
 Bee Yean Low¹
 Alexander Tong Boon Tan⁴
 Claire Anderson⁵

¹School of Pharmacy, University of Nottingham, Semenyih, Malaysia; ²Department of Primary Care Medicine, University of Malaya Primary Care Research Group, University of Malaya, Kuala Lumpur, Malaysia; ³School of Pharmacy, Monash University Malaysia, Bandar Sunway, Malaysia; ⁴Department of Medicine, University of Malaya, Kuala Lumpur, Malaysia; ⁵Division of Social Research in Medicine and Health, School of Pharmacy, University of Nottingham, Nottingham, UK

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Correspondence: Pauline Siew Mei Lai
 Department of Primary Care Medicine,
 University of Malaya Primary Care
 Research Group (UMPCRG), Faculty of
 Medicine, University of Malaya, Lembah
 Pantai, Kuala Lumpur 59100, Malaysia
 Tel +60 3 794 9230
 Fax +60 3 7957 7941
 Email plai@ummc.edu.my

Purpose: To develop and validate the English version of the Satisfaction Questionnaire for Osteoporosis Prevention (SQOP) in Malaysia.

Methods: The SQOP was modified from the Osteoporosis Patient Satisfaction Questionnaire and developed based on literature review and patient interviews. Face and content validity were established via an expert panel. The SQOP consists of two sections: clinical services and types of counseling. There were 23 items in total, each with a five-point Likert-type response. Satisfaction score was calculated by converting the total score to a percentage. A higher score indicates higher satisfaction. English speaking, non-osteoporotic, postmenopausal women aged ≥ 50 years were included in the study. Participants were randomized to either the control or intervention group. Intervention participants were provided counseling, whereas control participants received none. Participants answered the SQOP at baseline and 2 weeks later.

Results: A total of 140 participants were recruited (control group: $n=70$; intervention group: $n=70$). No significant differences were found in any demographic aspects. Exploratory factor analysis extracted seven domains. Cronbach's α for the domains ranged from 0.531–0.812. All 23 items were highly correlated using Spearman's correlation coefficient 0.469–0.996 ($P<0.05$), with no significant change in the control group's overall test-retest scores, indicating that the SQOP achieved stable reliability. The intervention group had a higher score than the control group (87.91 ± 5.99 versus 61.87 ± 8.76 ; $P<0.05$), indicating that they were more satisfied than control participants. Flesch reading ease was 62.9.

Conclusion: The SQOP was found to be a valid and reliable instrument for assessing patients' satisfaction towards an osteoporosis screening and prevention service in Malaysia.

Keywords: patient satisfaction, randomized controlled trial, postmenopausal women, screening

Introduction

Patient satisfaction may potentially be used to evaluate current preventive efforts and to predict patients' adherence to preventive advice.¹ The shift from health care provider-centered care to a more patient-centered care emphasizes the need to evaluate humanistic outcomes such as patient satisfaction.² The rating of satisfaction by patients is a personal evaluation of health care services and providers.^{3,4} Patient satisfaction serves as an important determinant of the viability and sustainability of health care services.⁵

Due to the asymptomatic nature of osteoporosis, women who have osteoporosis are often not aware that they are at an increased risk of sustaining a fracture.⁶ Prevention measures and screening that aid in early detection are the most cost-effective ways to reduce the number of hospital admissions due to osteoporotic fractures. Evidence shows

that satisfied patients are more likely to continue using health care services, value and maintain relationships with health care providers, follow the advice of the health care professionals, adhere to treatment, and have better health outcomes, which in this case is a reduction in fracture rates.^{1,7}

Evaluating satisfaction can also assist health care professionals to provide health care services more effectively. Patient evaluations will help identify patients' needs, perceptions, concerns, and areas of service failure. This in turn may encourage health care providers to be accountable for the quality of service delivered which ensures continuous monitoring and improvement in health care delivery.⁸

A search of published literature found several randomized controlled trials for osteoporosis screening services. These randomized controlled trials were conducted by various health care professionals such as primary care physicians,^{9,10} orthopedic surgeons,^{10,11} pharmacists,^{12–14} and nurses.¹⁵ All interventions demonstrated a positive effect towards bone mineral density scanning and osteoporosis treatment post fracture.^{9–15} However, only one study assessed patient satisfaction, highlighting the fact that many studies did not assess patient satisfaction as an outcome.¹⁶

A validated tool is necessary as it ensures the validity and the reliability of the results. Validity refers to the questionnaire measuring the objective it was designed to measure. Reliability refers to the ability of the questionnaire to produce the same results under the same conditions. Three tools have been developed globally to assess satisfaction pertaining to osteoporosis and its treatment,^{17–19} two of which were developed and validated in the US^{17,18} and one was developed in Malaysia.¹⁹ However, these satisfaction tools were not suitable to assess the satisfaction of patients from an osteoporosis screening and prevention program, as their focus was on osteoporosis and its treatment.^{17–19} To date, no instrument has been developed and validated specifically to assess patients' satisfaction towards an osteoporosis screening and prevention program in Malaysia. Therefore, the aim of this study was to develop and validate the English version of the Satisfaction Questionnaire for Osteoporosis Prevention (SQOP) to evaluate patients' satisfaction towards an osteoporosis screening and prevention program in Malaysia.

Methods

Development of the SQOP

Despite Malay being the national language of Malaysia, postmenopausal women aged 50 years residing in Malaysia are more fluent in English, as schooling was only conducted in the English language back then. Hence, the SQOP was

developed in English based on modifications from the Osteoporosis Patient Satisfaction Questionnaire (OPSQ) and findings from a qualitative study that examined the barriers and needs towards an osteoporosis screening and prevention service in Malaysia.

Of the 16 items in the OPSQ, nine items were removed. Eight of these items were on satisfaction related to osteoporosis medication, whilst one item was on a follow-up visit. Four items from the OPSQ were rephrased. In item two, the word “session” was used instead of “appointment” as appointments were not made with the participants. The question in item seven – “How useful was the service provided by the pharmacist in this study?” – was considered leading and was rephrased to “How would you rate the advice given by the pharmacist?”. The word “overall” was added to item nine – “How would you rate the overall quality of service that was given by the pharmacist to you?” – to make the question more specific. The original question for item 18 – “How would you rate your understanding of osteoporosis since you participated in the study?” – was modified to “How would you rate your understanding of osteoporosis now?” as the questionnaire was going to be used in clinical practice after completion of the study. Three items were retained from the OPSQ without any modifications.

Results from the qualitative findings suggest that the following domains may influence patients' satisfaction when utilizing an osteoporosis screening and prevention service: outcomes/efficacy, accessibility/convenience, technical quality, interpersonal relationship, finance, and continuity. A literature search noted that these domains were similar to those recommended by Ware et al except that Ware et al had an additional domain (ie, physical condition)⁴. Hence, the domain of physical condition was also included in the SQOP. Therefore, 16 new items were added. The final SQOP consisted of 23 items, and was divided into seven domains. Each item had a five-point Likert-type response. Please refer to the Supplementary material for the full questionnaire.

Validation of the SQOP

The validation process was a prospective, randomized controlled trial conducted from September to December 2013 at a primary care clinic of a tertiary hospital in Malaysia. Institutional Medical Ethics approval was obtained prior to the study (reference number 920.27).

Face and content validity of the SQOP was established via consultation with an expert panel (a consultant endocrinologist and four pharmacists with many years of research and clinical experience). Comprehension of the questionnaire

was tested on ten postmenopausal women who understood English. This involved asking the participants for their opinions about the phrasing, format, and content of the instrument. This resulted in the change of item eight from “Has the advice given by the pharmacist affected your life in general?” to “How would you rate the advice given by the pharmacist?” as participants commented that it was difficult to gauge the effect of newly given advice.

Participants

To validate the final version of the SQOP, 140 postmenopausal women were recruited. Postmenopausal women aged 50 years and above who had not been diagnosed with osteoporosis/osteopenia were included. Women who did not speak English were excluded. Sample size was calculated based on a 5:1 subject-to-item ratio for factor analysis.²⁰ Allowing for a 20% loss at follow-up, the total number of participants required was 70 in each arm.

Intervention

Intervention participants received 30 minutes of verbal counseling and an osteoporosis booklet. Topics covered during the counseling session were the definition of osteoporosis, consequences of untreated osteoporosis, risk factors for osteoporosis, the role of the bone mineral density scan (its function, what the results mean, accessibility, and the frequency a patient has to go for a bone mineral density scan), other tests used in osteoporosis screening (quantitative ultrasound scanning, X-ray, blood test, and the Osteoporosis Screening Tool for Asians), lifestyle changes (calcium intake, vitamin D intake, weight bearing exercise, and fall prevention), and treatment of osteoporosis.

Control participants received standard care. Standard care involved a regular visit to the doctor. Any counseling on osteoporosis by the doctor was opportunistic and at the doctors' discretion.

Procedure

The researcher screened for potential participants by using a 1:2 systematic random sampling procedure (Figure 1). Participants were recruited in the clinic's waiting area while they were waiting to see their doctor. Randomization of participants to either the control or intervention group was performed by drawing pieces of paper stating control or intervention from a bag while participants were filling out the demographic form. Participants answered the questionnaire themselves. However, for those who experienced some difficulty in reading the questions themselves, the researcher

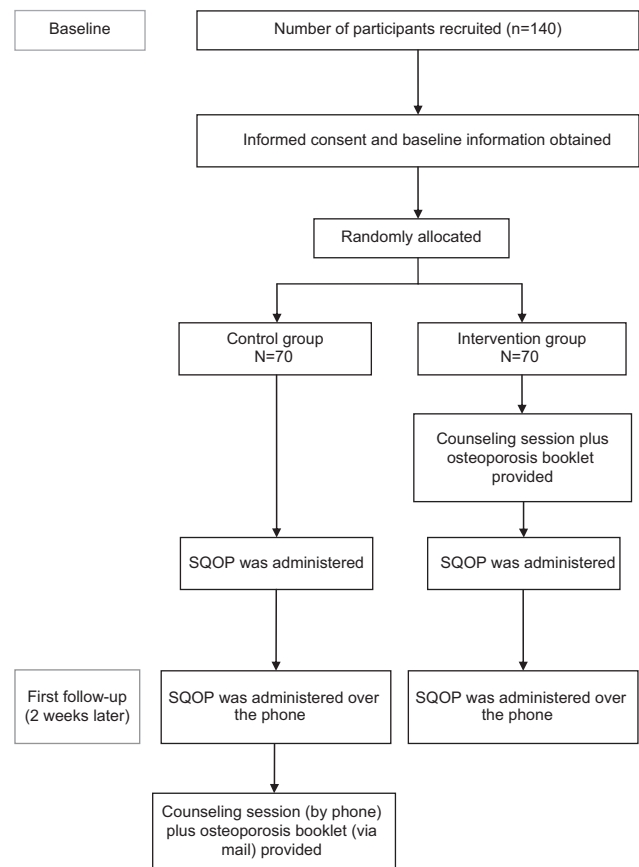


Figure 1 The validation procedure of the Satisfaction Questionnaire for Osteoporosis Prevention.

Abbreviation: SQOP, Satisfaction Questionnaire for Osteoporosis Prevention.

read the questions out for them and assisted them in filling in the questionnaire. The researcher ensured that all questions had been answered. Most participants required about 5 minutes to complete the questionnaire.

Data analysis

All data was entered into IBM® SPSS® version 20 (IBM Corporation, Armonk, NY, USA). Each item had a five-point Likert-type response. One indicates the lowest satisfaction for that item and five indicates the highest satisfaction. Scores ranged from 30–150, and were converted to a percentage, ranging from 0%–100%. Zero indicates the lowest level of satisfaction, whilst 100 indicates the highest. Baseline demographic data of the control and intervention group was compared using the chi-square test for categorical variables or the independent *t*-test for continuous variables. Nonparametric tests were used since the data obtained was not of normal distribution. A *P*-value <0.05 was considered statistically significant.

Face and content validity were used to ascertain analytical validity. Exploratory factor analysis (EFA) was used

Table 1 Baseline demographic characteristics of participants

Characteristics	Control (n=70)	Intervention (n=70)	χ^2/t -value ^{a,c}	P-value
Mean age \pm SD, years (range)/median	58.51 \pm 7.06 (50–77)/56.00	60.57 \pm 7.26 (50–77)/60.00	–1.700	0.091
Age range, n (%)				
<65 years	56 (80.0)	50 (71.4)	1.398	0.237
\geq 65 years	14 (20.0)	20 (28.6)		
Ethnicity, n (%)				
Malay	13 (18.6)	17 (24.3)		
Chinese	31 (44.3)	29 (41.4)	2.467	0.495
Indian	24 (34.3)	19 (27.1)		
Others ^b	2 (2.9)	5 (7.1)		
Mean BMI \pm SD/median	25.33 \pm 6.50/23.52	25.32 \pm 5.75/23.63	0.003	0.997
BMI, n (%)				
<18.5 (underweight)	5 (7.1)	5 (7.1)	0.833	0.841
18.5–24.9 (normal)	37 (52.9)	32 (45.7)		
25.0–29.9 (overweight)	17 (24.3)	19 (27.1)		
\geq 30.0 (obese)	11 (15.7)	14 (20.0)		
Level of education, n (%)				
Primary (6 years of education)	4 (5.7)	3 (4.3)		
Secondary (11–13 years of education)	33 (47.1)	28 (40.0)		
Diploma/technical (12–14 years of education)	15 (21.4)	14 (20.0)	1.727	0.631
Tertiary/postgraduate (15–21 years of education)	18 (25.7)	25 (35.7)		

Notes: ^aThe chi-square test was used for all categorical variables, whilst the independent t-test was used for all continuous variables. ^bIncludes four Eurasians, one Portuguese, and one Thai. ^cFisher's exact test was used as the number of cells with an expected count of less than five was more than 20% of the total number of cells.

Abbreviations: BMI, body mass index; SD, standard deviation.

to determine empirical validity. EFA was performed to provide information about the validity of the items within each domain and to explore the appropriateness of the factor structure of the current questionnaire. It is important to note that the control group was not required to answer items 19–23 as these items were specifically assessing the satisfaction of the intervention conducted. However, all items (ie, items 1–23) in the EFA were included for two reasons. Firstly, although the control group did not answer items 19–23, it still represented one of the seven domains. Secondly, if control participants were excluded from the EFA, then the minimum number of participants required would not have been satisfied (ie, 100 participants) as there

were more than six factors present.²¹ The extraction method used was maximum likelihood and the rotation method was promax. To determine how many factors were retained, a criterion of eigenvalue >1.0 was used. Corrected item-total correlations were used to identify items which did not measure the same main component as the other items. A value of <0.3 indicates that the item is measuring a different component from the scale as a whole.

The mean score \pm standard deviation was calculated for each item. Cronbach's α and test–retest were used to assess the internal consistency and stable reliability of the questionnaire, respectively. The results for the intervention and control groups were combined to test for internal

Table 2 Eigenvalue of the Satisfaction Questionnaire for Osteoporosis Prevention and the proportion of variance explained using promax and maximum likelihood

Factor	Before rotation			After rotation		
	Eigenvalue	Percentage of variances	Cumulative percentage of variances	Eigenvalue	Percentage of variances	Cumulative percentage of variances
Factor 1	3.520	15.30	15.30	7.548	32.82	32.82
Factor 2	2.981	12.96	28.27	2.908	12.64	45.46
Factor 3	2.364	10.28	38.54	1.977	8.56	54.05
Factor 4	2.363	10.27	48.81	1.768	7.69	61.74
Factor 5	2.287	9.94	58.76	1.523	6.62	68.36
Factor 6	1.631	1.09	65.85	1.354	5.89	74.25
Factor 7	1.122	4.88	70.73	1.112	4.84	79.09

reliability (Cronbach's α). Cronbach's $\alpha \geq 0.70$ indicates good internal reliability.²² Microsoft Office® Word® 2007 (Microsoft Corporation, Redmond, WA, USA) was used to calculate the Flesch reading ease. Flesch reading ease was performed to assess the reading comprehension level necessary to understand the written document. An average document should have a score of 60–70.²³

External validity was determined using discriminant validity, which was performed on the control and intervention groups to assess if the SQOP was able to differentiate between the satisfaction levels of the two groups. Since, the control group was not required to answer items 19–23, the total score of both the control and intervention groups were changed into percentages. The Mann–Whitney *U*-test was then used to analyze the discriminant validity between the control and intervention groups using their percentages. To assess test–retest reliability, the Wilcoxon signed-rank test and Spearman's correlation were used.

Results

A total of 173 participants were approached: 33 declined and 140 participants (80.9%) were recruited (control group: $n=70$; intervention group: $n=70$). No significant differences were found between the control group and intervention group in all demographic aspects (Table 1).

Factor analysis and psychometric properties of the SQOP

EFA extracted seven domains with a criterion of eigenvalue >1.0 . This explains the 79.1% of the cumulative variances. The eigenvalue and the proportion of variance explained are shown in Table 2. The factor loading of the items to each factor are shown in Table 3 (0.118–0.977). All items had a factor loading of >0.3 except item 17, which had a factor loading of 0.118. However, this item was maintained due to its importance based on findings from the qualitative study. Correlations between the factors resulting from the rotation were similar and the residuals ranged from -0.184 – 0.167 (Table 4).

The items representing each domain are shown in Table 5. Cronbach's α of each domain are shown in Table 5 ranging from 0.531–0.812. All items had a corrected item-total correlation of >0.3 . Flesch reading ease was 62.9.

At retest, eight participants (5.7%) dropped out of the study: three (2.1%) were overseas and five (3.6%) could not be contacted, leaving 132 (94.3%) at retest. Results from the control group and intervention group were analyzed separately. All test–retest scores were significantly correlated for both the control ($P<0.05$) and intervention ($P<0.05$) group. No significant difference was found for all items in the control group except items five and six (Table 6). For the intervention group, no significant difference was found for all items except items

Table 3 Factor loading of items in the Satisfaction Questionnaire for Osteoporosis Prevention

Items	Factor 1	Factor 2	Factor 3	Factor 4	Factor 5	Factor 6	Factor 7
Q1	0.037	0.339	0.283	0.034	0.373	0.417	0.423
Q2	0.428	0.304	0.340	0.108	0.040	0.914	0.163
Q3	0.078	0.392	0.141	0.063	0.977	0.078	0.004
Q4	0.109	0.462	0.277	0.173	0.379	0.166	-0.120
Q5	0.122	0.046	-0.004	0.188	-0.058	0.082	0.451
Q6	-0.169	0.125	0.052	-0.035	0.826	-0.130	-0.001
Q7	0.502	0.427	0.298	0.387	-0.079	0.545	0.170
Q8	0.451	0.199	0.593	0.462	0.041	0.555	0.222
Q9	0.590	0.703	0.472	0.114	0.430	0.153	-0.035
Q10	0.761	0.232	0.403	0.049	0.066	0.258	0.008
Q11	0.788	0.225	0.603	0.433	0.036	0.472	0.384
Q12	0.519	0.250	0.383	0.367	-0.063	0.378	0.110
Q13	0.530	0.622	0.669	0.434	0.080	0.565	0.430
Q14	0.714	0.318	-0.035	0.338	-0.053	0.562	0.240
Q15	0.841	0.400	0.350	0.520	-0.019	0.365	0.210
Q16	0.432	0.315	0.563	0.910	0.164	0.094	0.010
Q17	-0.311	0.037	-0.633	-0.239	0.118	-0.112	0.106
Q18	0.363	0.675	0.338	0.164	0.227	0.404	0.243
Q19	0.331	0.701	0.159	0.178	0.257	-0.017	-0.286
Q20	-0.014	0.762	0.027	0.135	0.125	0.265	0.355
Q21	0.384	0.669	0.771	0.467	0.287	0.358	0.175
Q22	0.191	0.288	0.351	0.924	0.084	0.368	0.536
Q23	0.308	0.660	0.818	0.406	0.401	0.475	0.452

Table 4 Correlations between the factors resulting from the rotation using promax and maximum likelihood

	Q1	Q2	Q3	Q4	Q5	Q6	Q7	Q8	Q9	Q10	Q11
Correlation											
Q1	10.000	0.320	0.351	0.336	-0.095	0.196	0.167	0.221	0.242	0.142	0.221
Q2	0.320	10.000	0.104	0.231	-0.116	-0.131	0.526	0.539	0.240	0.367	0.411
Q3	0.351	0.104	10.000	0.390	-0.055	0.808	-0.016	0.038	0.435	0.082	0.038
Q4	0.336	0.231	0.390	10.000	-0.110	0.170	0.193	0.154	0.448	-0.082	0.000
Q5	-0.095	-0.116	-0.055	-0.110	10.000	-0.136	0.198	0.137	0.021	-0.004	0.239
Q6	0.196	-0.131	0.808	0.170	-0.136	10.000	-0.155	-0.101	0.147	-0.090	-0.101
Q7	0.167	0.526	-0.016	0.193	0.198	-0.155	10.000	0.471	0.353	0.198	0.471
Q8	0.221	0.539	0.038	0.154	0.137	-0.101	0.471	10.000	0.243	0.438	0.576
Q9	0.242	0.240	0.435	0.448	0.021	0.147	0.353	0.243	10.000	0.521	0.398
Q10	0.142	0.367	0.082	-0.082	-0.004	-0.090	0.198	0.438	0.521	10.000	0.666
Q11	0.221	0.411	0.038	0.000	0.239	-0.101	0.471	0.576	0.398	0.666	10.000
Q12	0.221	0.382	-0.035	0.139	0.013	-0.122	0.462	0.440	0.341	0.531	0.440
Q13	0.238	0.470	0.064	0.127	0.228	-0.167	0.486	0.427	0.532	0.350	0.603
Q14	0.150	0.515	0.087	0.044	0.211	-0.161	0.436	0.283	0.288	0.511	0.524
Q15	0.019	0.317	0.051	0.202	0.226	-0.132	0.542	0.386	0.524	0.478	0.759
Q16	0.000	0.021	0.174	0.248	0.082	0.076	0.317	0.481	0.301	0.184	0.481
Q17	0.000	-0.207	0.190	-0.195	0.150	0.298	-0.113	-0.449	-0.218	-0.240	-0.449
Q18	0.360	0.368	0.247	0.131	0.081	-0.143	0.316	0.337	0.435	0.431	0.337
Q19	0.074	0.084	0.306	0.515	-0.115	0.084	0.352	0.051	0.574	0.204	0.051
Q20	0.216	0.109	0.152	0.140	0.051	0.031	0.237	-0.046	0.351	-0.064	-0.046
Q21	0.353	0.334	0.242	0.451	-0.027	0.155	0.348	0.420	0.557	0.375	0.420
Q22	0.188	0.078	0.109	0.109	0.317	0.047	0.324	0.429	0.031	-0.046	0.429
Q23	0.381	0.367	0.331	0.378	0.125	0.192	0.366	0.522	0.508	0.280	0.522
Reproduced correlation											
Q1	0.382	0.320	0.351	0.169	0.089	0.242	0.152	0.198	0.213	0.086	0.195
Q2	0.320	0.999	0.104	0.232	-0.115	-0.131	0.526	0.539	0.240	0.367	0.411
Q3	0.351	0.104	0.999	0.389	-0.055	0.808	-0.016	0.039	0.435	0.083	0.038
Q4	0.169	0.232	0.389	0.368	-0.150	0.238	0.179	0.159	0.365	0.072	0.027
Q5	0.089	-0.115	-0.055	-0.150	0.319	-0.046	0.061	0.036	-0.012	0.057	0.250
Q6	0.242	-0.131	0.808	0.238	-0.046	0.746	-0.209	-0.087	0.186	-0.095	-0.116
Q7	0.152	0.526	-0.016	0.179	0.061	-0.209	0.489	0.408	0.318	0.328	0.438
Q8	0.198	0.539	0.039	0.159	0.036	-0.087	0.408	0.594	0.228	0.347	0.579
Q9	0.213	0.240	0.435	0.365	-0.012	0.186	0.318	0.228	0.778	0.517	0.414
Q10	0.086	0.367	0.083	0.072	0.057	-0.095	0.328	0.347	0.517	0.679	0.641
Q11	0.195	0.411	0.038	0.027	0.250	-0.116	0.438	0.579	0.414	0.641	0.884
Q12	0.086	0.382	-0.035	0.111	0.064	-0.170	0.383	0.415	0.288	0.377	0.501
Q13	0.310	0.469	0.064	0.238	0.158	-0.124	0.505	0.522	0.504	0.422	0.627
Q14	0.135	0.515	0.087	0.040	0.213	-0.157	0.497	0.307	0.308	0.455	0.542
Q15	0.089	0.317	0.051	0.091	0.237	-0.160	0.493	0.431	0.489	0.584	0.738
Q16	0.008	0.021	0.174	0.258	0.086	0.076	0.318	0.480	0.302	0.182	0.480
Q17	0.037	-0.207	0.190	-0.071	0.099	0.186	-0.173	-0.434	-0.159	-0.310	-0.413
Q18	0.289	0.368	0.246	0.286	0.065	0.034	0.371	0.235	0.518	0.292	0.306
Q19	0.039	0.084	0.306	0.397	-0.123	0.094	0.258	0.025	0.597	0.203	0.031
Q20	0.263	0.109	0.153	0.249	0.130	0.004	0.255	-0.047	0.332	-0.076	-0.060
Q21	0.271	0.334	0.243	0.395	-0.002	0.069	0.398	0.481	0.565	0.309	0.455
Q22	0.187	0.078	0.109	0.108	0.317	0.047	0.324	0.429	0.031	-0.046	0.429
Q23	0.422	0.367	0.331	0.351	0.101	0.170	0.365	0.501	0.528	0.292	0.516

Q12	Q13	Q14	Q15	Q16	Q17	Q18	Q19	Q20	Q21	Q22	Q23
0.221	0.238	0.150	0.019	0.000	0.000	0.360	0.074	0.216	0.353	0.188	0.381
0.382	0.470	0.515	0.317	0.021	-0.207	0.368	0.084	0.109	0.334	0.078	0.367
-0.035	0.064	0.087	0.051	0.174	0.190	0.247	0.306	0.152	0.242	0.109	0.331
0.139	0.127	0.044	0.202	0.248	-0.195	0.131	0.515	0.140	0.451	0.109	0.378
0.013	0.228	0.211	0.226	0.082	0.150	0.081	-0.115	0.051	-0.027	0.317	0.125
-0.122	-0.167	-0.161	-0.132	0.076	0.298	-0.143	0.084	0.031	0.155	0.047	0.192
0.462	0.486	0.436	0.542	0.317	-0.113	0.316	0.352	0.237	0.348	0.324	0.366
0.440	0.427	0.283	0.386	0.481	-0.449	0.337	0.051	-0.046	0.420	0.429	0.522
0.341	0.532	0.288	0.524	0.301	-0.218	0.435	0.574	0.351	0.557	0.031	0.508
0.531	0.350	0.511	0.478	0.184	-0.240	0.431	0.204	-0.064	0.375	-0.046	0.280
0.440	0.603	0.524	0.759	0.481	-0.449	0.337	0.051	-0.046	0.420	0.429	0.522
10.000	0.411	0.452	0.411	0.374	-0.272	0.189	0.115	-0.017	0.509	0.292	0.316
0.411	10.000	0.370	0.640	0.455	-0.347	0.559	0.232	0.352	0.697	0.462	0.722
0.452	0.370	10.000	0.690	0.175	0.144	0.301	0.165	0.207	0.170	0.291	0.148
0.411	0.640	0.690	10.000	0.513	-0.250	0.285	0.301	0.110	0.397	0.433	0.382
0.374	0.455	0.175	0.513	10.000	-0.416	0.176	0.288	0.021	0.606	0.777	0.507
-0.272	-0.347	0.144	-0.250	-0.416	10.000	-0.026	-0.025	0.301	-0.289	-0.127	-0.319
0.189	0.559	0.301	0.285	0.176	-0.026	10.000	0.478	0.562	0.447	0.183	0.447
0.115	0.232	0.165	0.301	0.288	-0.025	0.478	10.000	0.472	0.395	0.018	0.292
-0.017	0.352	0.207	0.110	0.021	0.301	0.562	0.472	10.000	0.307	0.235	0.372
0.509	0.697	0.170	0.397	0.606	-0.289	0.447	0.395	0.307	10.000	0.432	0.804
0.292	0.462	0.291	0.433	0.777	-0.127	0.183	0.018	0.235	0.432	10.000	0.492
0.316	0.722	0.148	0.382	0.507	-0.319	0.447	0.292	0.372	0.804	0.492	1.000
0.086	0.310	0.135	0.089	0.008	0.037	0.289	0.039	0.263	0.271	0.187	0.422
0.382	0.469	0.515	0.317	0.021	-0.207	0.368	0.084	0.109	0.334	0.078	0.367
-0.035	0.064	0.087	0.051	0.174	0.190	0.246	0.306	0.153	0.243	0.109	0.331
0.111	0.238	0.040	0.091	0.258	-0.071	0.286	0.397	0.249	0.395	0.108	0.351
0.064	0.158	0.213	0.237	0.086	0.099	0.065	-0.123	0.130	-0.002	0.317	0.101
-0.170	-0.124	-0.157	-0.160	0.076	0.186	0.034	0.094	0.004	0.069	0.047	0.170
0.383	0.505	0.497	0.493	0.318	-0.173	0.371	0.258	0.255	0.398	0.324	0.365
0.415	0.522	0.307	0.431	0.480	-0.434	0.235	0.025	-0.047	0.481	0.429	0.501
0.288	0.504	0.308	0.489	0.302	-0.159	0.518	0.597	0.332	0.565	0.031	0.528
0.377	0.422	0.455	0.584	0.182	-0.310	0.292	0.203	-0.076	0.309	-0.046	0.292
0.501	0.627	0.542	0.738	0.480	-0.413	0.306	0.031	-0.060	0.455	0.429	0.516
0.370	0.442	0.379	0.480	0.377	-0.296	0.250	0.154	0.043	0.369	0.292	0.338
0.442	0.753	0.391	0.570	0.455	-0.321	0.515	0.291	0.368	0.672	0.462	0.716
0.379	0.391	0.836	0.673	0.174	0.076	0.344	0.188	0.205	0.131	0.291	0.134
0.480	0.570	0.673	0.806	0.515	-0.224	0.374	0.286	0.142	0.412	0.433	0.381
0.377	0.455	0.174	0.515	0.990	-0.418	0.174	0.288	0.020	0.603	0.777	0.507
-0.296	-0.321	0.076	-0.224	-0.418	0.612	-0.024	0.010	0.290	-0.397	-0.127	-0.335
0.250	0.515	0.344	0.374	0.174	-0.024	0.508	0.422	0.485	0.475	0.184	0.500
0.154	0.291	0.188	0.286	0.288	0.010	0.422	0.715	0.453	0.433	0.018	0.288
0.043	0.368	0.205	0.142	0.020	0.290	0.485	0.453	0.791	0.329	0.235	0.365
0.369	0.672	0.131	0.412	0.603	-0.397	0.475	0.433	0.329	0.784	0.432	0.770
0.292	0.462	0.291	0.433	0.777	-0.127	0.184	0.018	0.235	0.432	10.000	0.492
0.338	0.716	0.134	0.381	0.507	-0.335	0.500	0.288	0.365	0.770	0.492	0.865

(Continued)

Table 4 (Continued)

	Q1	Q2	Q3	Q4	Q5	Q6	Q7	Q8	Q9	Q10	Q11
Residual											
Q1		0.000	0.000	0.167	-0.184	-0.047	0.015	0.023	0.029	0.056	0.026
Q2	0.000		0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Q3	0.000	0.000		0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Q4	0.167	0.000	0.000		0.040	-0.068	0.014	-0.005	0.083	-0.154	-0.027
Q5	-0.184	0.000	0.000	0.040		-0.090	0.137	0.101	0.033	-0.061	-0.011
Q6	-0.047	0.000	0.000	-0.068	-0.090		0.054	-0.013	-0.039	0.005	0.016
Q7	0.015	0.000	0.000	0.014	0.137	0.054		0.063	0.035	-0.130	0.033
Q8	0.023	0.000	0.000	-0.005	0.101	-0.013	0.063		0.016	0.091	-0.003
Q9	0.029	0.000	0.000	0.083	0.033	-0.039	0.035	0.016		0.004	-0.016
Q10	0.056	0.000	0.000	-0.154	-0.061	0.005	-0.130	0.091	0.004		0.024
Q11	0.026	0.000	0.000	-0.027	-0.011	0.016	0.033	-0.003	-0.016	0.024	
Q12	0.135	0.000	0.000	0.029	-0.051	0.049	0.079	0.025	0.054	0.153	-0.061
Q13	-0.072	0.000	0.000	-0.111	0.070	-0.042	-0.019	-0.095	0.028	-0.072	-0.024
Q14	0.016	0.000	0.000	0.003	-0.002	-0.005	-0.060	-0.023	-0.020	0.056	-0.018
Q15	-0.069	0.000	0.000	0.112	-0.011	0.028	0.049	-0.045	0.036	-0.106	0.021
Q16	-0.008	0.000	0.000	-0.009	-0.004	0.000	-0.001	0.001	-0.002	0.002	0.000
Q17	-0.037	0.000	0.000	-0.124	0.051	0.112	0.060	-0.014	-0.060	0.069	-0.036
Q18	0.071	0.000	0.001	-0.155	0.015	-0.178	-0.055	0.103	-0.082	0.139	0.031
Q19	0.035	0.000	0.000	0.117	0.008	-0.010	0.094	0.025	-0.023	0.001	0.020
Q20	-0.047	0.000	0.000	-0.109	-0.079	0.026	-0.018	0.001	0.019	0.012	0.014
Q21	0.082	0.000	0.000	0.056	-0.025	0.086	-0.050	-0.061	-0.008	0.066	-0.035
Q22	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Q23	-0.041	0.000	0.000	0.027	0.024	0.023	0.000	0.021	-0.020	-0.013	0.006

four, five, 13, 20, 21, and 23 (Table 6). The instrument in this study showed that the intervention group had a significantly higher patient satisfaction compared to the control group (88.10 ± 5.87 versus 61.87 ± 8.76 ; $P < 0.05$). Thus, the SQOP was able to discriminate between a group with higher satisfaction and lower satisfaction. This also demonstrated that the intervention had an effect on participants' satisfaction level.

Comparison of the SQOP with other validated instruments

The psychometric properties of the SQOP were similar to that of other validated instruments for measuring patient satisfaction (Table 7).

Discussion

The SQOP performed satisfactorily in both the EFA and psychometric properties. This indicates that the SQOP is suitable to assess patients' satisfaction towards an osteoporosis screening and prevention service in Malaysia.

EFA showed that there were seven domains of satisfaction being measured within the main component to assess patients' satisfaction towards an osteoporosis screening and prevention program in Malaysia. This was as expected as the researchers initially designed this tool to assess

seven domains, namely outcomes/efficacy, accessibility/convenience, technical quality, interpersonal relationship, finance, physical condition, and continuity. The seven domains were deemed to be significant to assess patients' satisfaction as they concurred with the themes from a previous qualitative study.⁴

The domains measured by SQOP were then compared to that of other satisfaction tools related to osteoporosis. For example, the OPSQ measured convenience, time, trust, and usefulness of the counseling session.¹⁹ The Osteoporosis Patient Treatment Satisfaction Questionnaire measured convenience, confidence with daily activities, side effects, and overall satisfaction.¹⁸ The Preference and Satisfaction Questionnaire on the other hand measured preference, bother, and satisfaction.¹⁷ It was difficult to compare the domains of the Osteoporosis Patient Treatment Satisfaction Questionnaire, Preference and Satisfaction Questionnaire, and OPSQ as these tools were targeted at osteoporosis treatment instead of prevention. However, there were similarities in terms of some of the domains measured such as convenience and usefulness.

Corrected item-total correlations showed that all items measured the same main component – satisfaction. SQOP was designed as a multidimensional tool, rendering an overall

Q12	Q13	Q14	Q15	Q16	Q17	Q18	Q19	Q20	Q21	Q22	Q23
0.135	-0.072	0.016	-0.069	-0.008	-0.037	0.071	0.035	-0.047	0.082	0.000	-0.041
0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
0.000	0.000	0.000	0.000	0.000	0.000	0.001	0.000	0.000	0.000	0.000	0.000
0.029	-0.111	0.003	0.112	-0.009	-0.124	-0.155	0.117	-0.109	0.056	0.000	0.027
-0.051	0.070	-0.002	-0.011	-0.004	0.051	0.015	0.008	-0.079	-0.025	0.000	0.024
0.049	-0.042	-0.005	0.028	0.000	0.112	-0.178	-0.010	0.026	0.086	0.000	0.023
0.079	-0.019	-0.060	0.049	-0.001	0.060	-0.055	0.094	-0.018	-0.050	0.000	0.000
0.025	-0.095	-0.023	-0.045	0.001	-0.014	0.103	0.025	0.001	-0.061	0.000	0.021
0.054	0.028	-0.020	0.036	-0.002	-0.060	-0.082	-0.023	0.019	-0.008	0.000	-0.020
0.153	-0.072	0.056	-0.106	0.002	0.069	0.139	0.001	0.012	0.066	0.000	-0.013
-0.061	-0.024	-0.018	0.021	0.000	-0.036	0.031	0.020	0.014	-0.035	0.000	0.006
	-0.031	0.074	-0.069	-0.003	0.024	-0.061	-0.038	-0.060	0.140	0.000	-0.022
-0.031		-0.021	0.070	0.001	-0.026	0.045	-0.059	-0.015	0.025	0.000	0.005
0.074	-0.021		0.018	0.001	0.068	-0.042	-0.022	0.002	0.039	0.000	0.014
-0.069	0.070	0.018		-0.002	-0.026	-0.089	0.015	-0.032	-0.015	0.000	0.001
-0.003	0.001	0.001	-0.002		0.001	0.003	0.000	0.001	0.003	0.000	0.000
0.024	-0.026	0.068	-0.026	0.001		-0.001	-0.035	0.011	0.108	0.000	0.016
-0.061	0.045	-0.042	-0.089	0.003	-0.001		0.056	0.076	-0.028	0.000	-0.052
-0.038	-0.059	-0.022	0.015	0.000	-0.035	0.056		0.020	-0.038	0.000	0.004
-0.060	-0.015	0.002	-0.032	0.001	0.011	0.076	0.020		-0.023	0.000	0.007
0.140	0.025	0.039	-0.015	0.003	0.108	-0.028	-0.038	-0.023		0.000	0.035
0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000		0.000
-0.022	0.005	0.014	0.001	0.000	0.016	-0.052	0.004	0.007	0.035	0.000	

Cronbach's α unsuitable. Hence, Cronbach's α was conducted for each domain. The physical and technical quality domains had a low Cronbach's α of 0.531 and 0.535, respectively, due to the small number of items in each domain.²⁴ Nonetheless, the other two domains had an acceptable and good Cronbach's α of 0.661 (accessibility/convenience) and 0.812 (outcomes/efficacy).²⁴ However, Cronbach's α was not computed for the interpersonal relationship, continuity, and finance domains as these domains only had one item. These domains comprise a total of four items out of the 23 items. Flesch reading ease was satisfactory at 62.9 as this represents the standard reading level of documents for adults.

The SQOP performed satisfactorily at test-retest, indicating that the SQOP achieved stable reliability. Only item five and item six were significantly different at test-retest for control participants. This may be because participants may have been more "satisfied with the service" provided by the pharmacist.

As for the intervention group, there was no significant difference in item scores between test and retest except for items four, five, 13, 20, 21, and 23. Item four was regarding the comfort of the location. This difference may be due to participants having more time to reflect on the program and realizing that they were "more satisfied", leading them

to view the overall comfort more positively. Item five was regarding whether participants would ask the pharmacist if they had any questions about osteoporosis. Several of the participants changed to a more positive answer during the retest and this may be because of their intervention experience with the researcher who is a pharmacist. Items 13, 20, 21, and 23 were regarding the amount of information provided, explanation of consequences of untreated osteoporosis, explanation on how osteoporosis can be prevented via lifestyle changes, and osteoporosis booklet provided, respectively. A possible explanation for this positive change could be because participants may have had more time to consider the information provided, found it useful, and hence were more satisfied. This in turn improved the overall score for the intervention participants significantly.

The SQOP was able to differentiate between patients who were expected to be more satisfied as they received an additional service (ie, the intervention) compared to those who did not receive the intervention, indicating that the items in SQOP were specific in addressing the different attributes of patients' satisfaction towards an osteoporosis screening and prevention service in Malaysia.

Currently, there are only a limited number of studies that have validated instruments to measure patient satisfaction.^{17-19,25-28}

Table 5 Psychometric properties of the Satisfaction Questionnaire for Osteoporosis Prevention

Items	Questions	Corrected item-total correlation	Cronbach's α if item deleted	Cronbach's α	Domains
1	The service was conducted at a time that _____ (fill in the blank) for you	0.508	0.527	0.661	Accessibility/ convenience
2	During the session, what did you think about the time given to discuss your problems with the pharmacist?	0.486	0.549		
3	How would you rate the location of this service?	0.432	0.618		
4	How would you rate the comfort of the location?	0.363	–	0.531	Physical
12	If you have questions about osteoporosis, would you ask the pharmacist?	0.363	–		
5	Was the pharmacist easy to talk to?	0.378	–	0.535	Technical quality
9	How would you rate the service provided by the pharmacist?	0.378	–		
6*	How would you rate the advice given by the pharmacist?				Interpersonal relationship
7	How would you rate the overall quality of service that was given by the pharmacist to you?	0.439	0.801	0.812	Outcomes/ efficacy
8	This pharmacist service should _____ (fill in the blank)	0.426	0.803		
10	What do you think about having the same pharmacist to see you for subsequent osteoporosis care?	0.263	0.813		
13	Pharmacist in other hospitals should _____ (fill in the blank) this service	0.574	0.788		
14	How would you rate the amount of information provided to prevent falls?	0.500	0.797		
15	How would you rate the amount of information provided to change your diet to prevent bone loss?	0.480	0.799		
18	How would you rate the amount of information provided on the exercises to help strengthen bones?	0.551	0.791		
19	Would you pay for a pharmacist counseling service?	0.421	0.802		
20	If yes, how much are you willing to pay for each visit to the pharmacist? If you are not willing to pay anything for the service, please proceed to question 18	0.509	0.794		
21	How would you rate your understanding of osteoporosis now?	0.440	0.801		
22	Explanation of osteoporosis	0.446	0.800		
23	Explanation of consequences of untreated osteoporosis	0.543	0.793		
11*	Explanation on how osteoporosis can be prevented via lifestyle change(s)				Continuity
16*	Explanation on the available methods to screen for osteoporosis				Finance
17#	Osteoporosis booklet provided				

Notes: *There was only one item in these domains, hence Cronbach's α could not be conducted. #This was an optional question, hence it was excluded from the calculation of Cronbach's α .

The psychometric properties of the SQOP were similar to previous satisfaction tools.

A limitation of this study was that SPSS Amos (IBM) was not used to conduct confirmatory factor analysis as five of the seven domains were developed with two items

or less (finance, physical condition, interpersonal relationship, continuity, and technical quality). This was because the questionnaire was designed so that participants would only take approximately 5 minutes to complete, to ensure the practicality of its usage in daily practice. The number

Table 6 Test and retest reliability of the Satisfaction Questionnaire for Osteoporosis Prevention

Items	Control group							Spearman's correlation test*
	Test (n=70)		Retest (n=70)		Wilcoxon–signed rank test			
	Mean ± SD	Median	Mean ± SD	Median	Mean/rank	z score	P-value	
1	3.17±0.72	3.00	3.13±0.78	3.00	4.00/4.00	−1.134	0.257	0.938
2	2.90±0.98	3.00	2.91±1.03	3.00	1.00/2.00	−0.447	0.655	0.978
3	3.73±1.23	4.00	3.74±1.21	5.00	4.00/4.00	−0.378	0.705	0.949
4	3.59±1.17	4.00	3.63±1.17	5.00	9.33/8.00	−0.688	0.491	0.877
5	3.26±1.37	3.00	3.31±1.37	4.25	0.00/2.50	−2.000	0.046*	0.983
6	3.49±1.13	4.00	3.23±1.22	4.00	8.32/4.50	−2.508	0.012*	0.745
7	3.59±0.81	4.00	3.63±0.85	4.00	3.00/3.00	−1.342	0.180	0.958
8	3.29±0.95	3.00	3.13±1.06	4.00	8.61/5.50	−1.639	0.101	0.726
9	3.14±0.97	3.00	3.11±0.93	4.00	3.50/3.50	−0.816	0.414	0.955
10	3.36±0.84	3.00	3.41±0.83	4.00	3.50/5.10	−1.100	0.271	0.908
11	3.91±1.03	4.00	3.86±0.92	5.00	5.25/4.50	1.155	0.248	0.925
12	3.33±0.78	3.00	3.33±0.78	3.25	1.50/1.50	0.000	1.000	0.996
13	2.41±1.10	3.00	2.31±0.96	3.00	11.09/8.50	−1.170	0.242	0.802
14	2.20±0.97	2.00	2.19±1.01	3.00	9.00/8.00	−0.229	0.819	0.862
15	2.20±0.97	2.50	2.13±0.95	3.00	11.82/12.28	−0.923	0.356	0.796
16	2.01±1.35	1.00	1.99±1.29	3.00	3.33/2.50	−0.707	0.480	0.947
17	3.25±1.71	4.00	3.73±1.61	5.00	0.00/1.00	−1.000	0.317	0.726
18	3.01±0.12	3.00	3.00±0.00	3.00	1.00/0.00	−1.000	0.317	0.981
19#								
20#								
21#								
22#								
23#								
Total	61.87±8.76	61.18	61.23±8.96	61.77	29.79/24.41	−1.580	0.114	0.941

Items	Intervention group							Spearman's correlation test
	Test (n=70)		Retest (n=62)		Wilcoxon–signed rank test			
	Mean ± SD	Median	Mean ± SD	Median	Mean/rank	t-value	P-value	
1	4.49±0.79	5.00	4.55±0.92	5.00	7.50/5.00	−1.387	0.166	0.818
2	4.47±0.85	5.00	4.50±0.99	5.00	4.50/4.50	−1.414	0.157	0.849
3	4.67±0.76	5.00	4.69±0.78	5.00	3.50/3.50	−0.816	0.414	0.868
4	4.49±0.68	5.00	4.63±0.73	5.00	5.50/5.50	−2.530	0.011*	0.794
5	4.27±1.01	5.00	4.35±0.98	5.00	0.00/3.50	−2.333	0.020*	0.900
6	4.97±0.17	5.00	4.97±0.18	5.00	1.50/1.50	0.000	1.000	0.483
7	4.60±0.49	5.00	4.68±0.47	5.00	4.00/4.00	−1.890	0.059	0.769
8	4.83±0.38	5.00	4.84±0.37	5.00	2.50/2.50	0.000	1.000	0.762
9	4.31±0.60	4.00	4.39±0.58	4.00	3.00/3.60	−1.667	0.096	0.815
10	4.81±0.46	5.00	4.79±0.48	5.00	2.50/2.50	0.000	1.000	0.794
11	4.51±0.78	5.00	4.42±0.90	5.00	2.50/1.00	−1.069	0.285	0.910
12	4.66±0.66	5.00	4.69±0.64	5.00	3.00/3.00	−1.342	0.180	0.822
13	4.43±0.75	5.00	4.50±0.74	5.00	0.00/3.50	−2.449	0.014*	0.869
14	4.40±0.75	5.00	4.45±0.74	5.00	5.50/5.50	−1.265	0.206	0.751
15	4.46±0.76	5.00	4.47±0.76	5.00	2.50/2.50	−1.000	0.317	0.915
16	2.76±1.47	3.00	2.28±1.48	3.00	0.00/1.00	−1.000	0.317	0.967
17	3.25±1.48	4.00	3.23±1.42	3.50	1.00/0.00	−1.000	0.317	0.987
18	4.64±0.54	5.00	4.69±0.50	5.00	0.00/2.00	−1.732	0.083	0.929
19	4.20±4.67	4.00	4.29±0.49	4.00	3.50/3.50	−1.633	0.102	0.785
20	4.06±0.59	4.00	4.19±0.62	4.00	6.00/6.00	−2.111	0.035*	0.798
21	4.26±0.50	4.00	4.37±0.52	4.00	4.50/5.06	−2.309	0.021*	0.699
22	4.19±0.55	4.00	4.23±0.64	4.00	3.50/3.50	−1.633	0.102	0.849
23	4.44±0.50	4.00	4.77±0.42	5.00	0.00/11.50	−4.690	0.000*	0.459
Total	88.10±5.87	88.18	89.36±5.98	89.09	13.79/25.79	−4.995	0.000*	0.948

Notes: *Statistically significant at $P < 0.05$. **Item 17 was excluded from the total score calculation as it was an optional question. #The control group was not required to answer items 19–23 as these items were specifically assessing the satisfaction of the intervention conducted.

Abbreviation: SD, standard deviation.

Table 7 Comparison of psychometric properties of the Satisfaction Questionnaire for Osteoporosis Prevention to other validated patient satisfaction instruments

	SQOP	OPSQ ¹⁹	OPSAT-Q ¹⁸	PSQ ¹⁷	PSQ-An ²⁸	DMET ²⁶	DDSM-Q ²⁵	PEQD ²⁷
Number of subjects	140	180	104	1,583	312	202	114	1,472
Age,* years	59.5±7.2	65.6±9.3	65.1±10.3	65.4	60.2±11.8	55.7	61.9±9.4	51.0±16.0
Type of study	RCT	RCT	Observational	RCT	Observational	Observational	Observational	RCT
Number of items	23	16	16	20	11	73	31	12
Number of domains	7	2	4	6	2	2	3	4
Cronbach's α	0.81	0.86	0.72–0.89	0.61–0.93	0.83	0.79–0.95	–	0.73–0.84
Factor analysis: number of components	7	1	–	5	–	13	3	–
Number of times administered	Twice	Once	Twice	Once	Twice	Once	Once	Twice (n=202)
Interval between administration	2 weeks	NA	2 weeks	NA	4 weeks	NA	NA	66±14 days*
Test–retest reliability: intraclass correlation	0.46–0.98	NA	0.62–0.81	NA	0.45–0.67	NA	NA	0.66–0.83

Note: *Data are presented as mean \pm standard deviation.

Abbreviations: DDSM-Q, Diabetes Disease State Management Questionnaire; DMET, Diabetes Management Evaluation tool; NA, not applicable; OPSAT-Q, Osteoporosis Patient Treatment Satisfaction Questionnaire; OPSQ, Osteoporosis Patient Satisfaction Questionnaire; PEQD, Patients' Evaluation of the Quality of Diabetes Care; PSQ, Preference and Satisfaction Questionnaire; PSQ-An, Patient Satisfaction Questionnaire for Anemia Treatment; RCT, randomized controlled trial; SQOP, Satisfaction Questionnaire for Osteoporosis Prevention.

of items per domain was determined based on the patients' emphasis towards the domain during the in-depth interviews. Nonetheless, EFA showed that there were seven components measuring the patients' satisfaction.

Another limitation of this study was that SQOP was designed to measure the satisfaction of patients towards a pharmacist-conducted osteoporosis screening and prevention service. Hence, wordings such as “Was the pharmacist easy to talk to?” was used. Hence, the implementation of this questionnaire is limited to services provided by a pharmacist. Minor modifications of the SQOP will be required if used to assess satisfaction provided by other health care professionals.

Participants in this study were mainly Chinese (42.9%) and Indians (30.7%). This does not represent the ethnic distribution of Malaysia. It only represents the ethnicity of patients who sought treatment at the study site, meaning that the results cannot be considered population based. Future validation studies of the tool to Malay and Mandarin, and enrollment of participants from multiple sites would be more representative of the Malaysian population.

Conclusion

The English version of the SQOP was found to be a reliable and valid instrument for assessing patients' satisfaction towards an osteoporosis screening and prevention program in Malaysia. Future studies should include the translation of the SQOP into Malay and Mandarin to assess patient satisfaction for Malaysians that are not fluent in English.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary material

The full questionnaire of the English version of the Satisfaction Questionnaire for Osteoporosis Prevention.

Satisfaction Questionnaire for Osteoporosis Prevention (SQOP) self/assisted

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This questionnaire is to assess your satisfaction about a pharmacist-conducted osteoporosis prevention program. Filling out this questionnaire will provide information for us to further improve our services.

Please tick the answer that best suits your opinion.

A. Clinical services

1. The service was conducted at a time that _____
(fill in the blank) for you
 - ☐ ₅ Was definitely suitable
 - ☐ ₄ Was probably suitable
 - ☐ ₃ Made no difference
 - ☐ ₂ Was probably unsuitable
 - ☐ ₁ Was definitely unsuitable
2. During the session, what did you think about the time given to discuss your problems with the pharmacist?
 - ☐ ₅ Definitely enough time
 - ☐ ₄ Probably enough time
 - ☐ ₃ No difference
 - ☐ ₂ Probably insufficient time
 - ☐ ₁ Definitely insufficient time
3. How would you rate the location of this service?
 - ☐ ₅ Very convenient
 - ☐ ₄ Somewhat convenient
 - ☐ ₃ No difference
 - ☐ ₂ Somewhat inconvenient
 - ☐ ₁ Very inconvenient
4. How would you rate the comfort of the location?
 - ☐ ₅ Very comfortable
 - ☐ ₄ Somewhat comfortable
 - ☐ ₃ No difference
 - ☐ ₂ Somewhat uncomfortable
 - ☐ ₁ Very uncomfortable
5. If you have questions about osteoporosis, would you ask the pharmacist?
 - ☐ ₅ Yes, I would definitely trust the answer given by the pharmacist
 - ☐ ₄ Yes, I would probably trust the answer given by the pharmacist
 - ☐ ₃ No comment
 - ☐ ₂ No, I probably would not trust the answer given by the pharmacist
 - ☐ ₁ No, I definitely would not trust the answer given by the pharmacist

6. Was the pharmacist easy to talk to?
- ☐ ₅ Definitely approachable
 - ☐ ₄ Probably approachable
 - ☐ ₃ No difference
 - ☐ ₂ Probably unapproachable
 - ☐ ₁ Definitely unapproachable
7. How would you rate the service provided by the pharmacist?
- ☐ ₅ Definitely useful
 - ☐ ₄ Useful
 - ☐ ₃ No difference
 - ☐ ₂ Not useful
 - ☐ ₁ Definitely not useful
8. How would you rate the advice given by the pharmacist?
- ☐ ₅ Definitely helpful
 - ☐ ₄ Probably helpful
 - ☐ ₃ No difference
 - ☐ ₂ Probably not helpful
 - ☐ ₁ Definitely not helpful at all
9. How would you rate the overall quality of service that was given by the pharmacist to you?
- ☐ ₅ Excellent
 - ☐ ₄ Good
 - ☐ ₃ Satisfactory
 - ☐ ₂ Fair
 - ☐ ₁ Poor
10. This pharmacist service should _____ (fill in the blank)
- ☐ ₅ Definitely be continued
 - ☐ ₄ Probably be continued
 - ☐ ₃ No comment
 - ☐ ₂ Probably be discontinued
 - ☐ ₁ Definitely be discontinued
11. What do you think about having the same pharmacist to see you for subsequent osteoporosis care?
- ☐ ₅ Yes, I would definitely like that
 - ☐ ₄ Yes, I would probably like that
 - ☐ ₃ No difference
 - ☐ ₂ No, I probably would not like that
 - ☐ ₁ No, I definitely would not like that
12. Pharmacist in other hospitals should _____ (fill in the blank) this service
- ☐ ₅ Definitely provide
 - ☐ ₄ Probably provide
 - ☐ ₃ No comment
 - ☐ ₂ Probably not provide
 - ☐ ₁ Definitely not provide
13. How would you rate the amount of information provided to prevent falls?
- ☐ ₅ Definitely enough
 - ☐ ₄ Probably enough
 - ☐ ₃ No difference
 - ☐ ₂ Probably not enough
 - ☐ ₁ Definitely not enough

14. How would you rate the amount of information provided to change your diet to prevent bone loss?
15. How would you rate the amount of information provided on the exercises to help strengthen bones?
16. Would you pay for a pharmacist counseling service?
17. If yes, how much are you willing to pay for each visit to the pharmacist?
If you are not willing to pay anything for the service, please proceed to question 18
18. How would you rate your understanding of osteoporosis now?
- ☐ ₅ Definitely enough
- ☐ ₄ Probably enough
- ☐ ₃ No difference
- ☐ ₂ Probably not enough
- ☐ ₁ Definitely not enough
- ☐ ₅ Definitely enough
- ☐ ₄ Probably enough
- ☐ ₃ No difference
- ☐ ₂ Probably not enough
- ☐ ₁ Definitely not enough
- ☐ ₅ Yes, definitely
- ☐ ₄ Yes, probably
- ☐ ₃ No difference
- ☐ ₂ No, probably not
- ☐ ₁ No, definitely not
- ☐ ₅ RM1–5
- ☐ ₄ RM6–10
- ☐ ₃ RM11–15
- ☐ ₂ RM16–20
- ☐ ₁ More than RM20
- ☐ ₅ Much better than before
- ☐ ₄ Slightly better than before
- ☐ ₃ Same as before
- ☐ ₂ Slightly worse than before
- ☐ ₁ Much worse than before

B. Types of counseling

Please indicate how you found the following information which the pharmacist may have provided. If you were not provided with any counseling, please omit this section.

19. Explanation of osteoporosis
20. Explanation of consequences of untreated osteoporosis
- ☐ ₀ Not applicable
- ☐ ₁ Definitely not useful
- ☐ ₂ Not useful
- ☐ ₃ No difference
- ☐ ₄ Useful
- ☐ ₅ Extremely useful
- ☐ ₀ Not applicable
- ☐ ₁ Definitely not useful
- ☐ ₂ Not useful
- ☐ ₃ No difference
- ☐ ₄ Useful
- ☐ ₅ Extremely useful

21. Explanation on how osteoporosis can be prevented via life-style change(s)
22. Explanation on the available methods to screen for osteoporosis
23. Osteoporosis booklet provided
- ☐ ₀ Not applicable
- ☐ ₁ Definitely not useful
- ☐ ₂ Not useful
- ☐ ₃ No difference
- ☐ ₄ Useful
- ☐ ₅ Extremely useful
- ☐ ₀ Not applicable
- ☐ ₁ Definitely not useful
- ☐ ₂ Not useful
- ☐ ₃ No difference
- ☐ ₄ Useful
- ☐ ₅ Extremely useful
- ☐ ₀ Not applicable
- ☐ ₁ Definitely not useful
- ☐ ₂ Not useful
- ☐ ₃ No difference
- ☐ ₄ Useful
- ☐ ₅ Extremely useful

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