Original Article

The Effect of *Curcumin* Ointment on the Quality of Life of Older Adults with Knee Osteoarthritis: A Randomized Placebo Clinical Trial

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Background: Knee osteoarthritis (KO) is one of the most common health problems affecting the quality of life (QOL) in older adults. Objective: Because improving the OOL is one of the important therapeutic goals, this study aimed to investigate the effect of curcumin ointment on the OOL in older adults with KO. Methods: This double-blind randomized clinical trial was performed from February 20, to June 22, 2019, on 72 older adults with KO referring to the office of a physician in Kashan, Iran. The individuals with eligibility criteria were recruited sequentially and were randomly assigned to two groups of 36 to apply either curcumin 5% ointment or Vaseline ointment twice daily for 6 weeks around the knee. The Osteoarthritis Knee and Hip Quality of Life questionnaire was used to measure the patients' QOL at the beginning of the study, and at the end of the 4th and 6^{th} weeks. The Fisher's exact test, independent samples t-test, Friedman test, and repeated-measures analysis were used for data analysis. Results: The mean baseline QOL scores in the intervention and the placebo groups were 55.98 ± 8.12 and 58.75 ± 7.86 (P = 0.160) which then changed to 59.21 ± 7.98 and 60.01 ± 8.23 at the end of the 4th week and to 65.83 ± 8.81 and 60.26 ± 7.25 at the end of the 6th week, respectively. Repeated-measures analysis showed that over time. the use of curcumin ointment could significantly increase the mean QOL score in the intervention group (P = 0.001). Conclusion: Topical administration of 5% curcumin ointment can significantly improve the OOL in older adults with KO. Therefore, this ointment might be considered a complementary therapy to improve the QOL of patients with KO.

KEYWORDS: Curcumin, Knee, Older adults, Osteoarthritis, Quality of life

Introduction

nee osteoarthritis (KO) is one of the most common health problems in older adults.^[1] Due to its progressive nature, KO causes joint stiffness, pain, muscle weakness, mobility impairment,^[2] disability, falls, depression, and impaired quality of life (QOL).^[3,4] The prevalence of osteoarthritis is 60%–90% in people 65 years and older.^[2] The prevalence of KO in Iranian older adults has also been reported as 63.1%.^[5] Iranian older adults are more prone to knee joints injuries due to habits and lifestyle factors such as sitting on the ground, the shape of toilets, inadequate exercise activities, and standing cooking and housekeeping practices.^[6] Joint



surgery, medications, and protective modalities are commonly used to treat KO.^[7] However, surgeries are expensive and medications have side effects – especially in older adults who suffer multiple organ dysfunctions.^[8] Besides, the effects of walking aids, physiotherapy, hydrotherapy, weight loss, and lifestyle

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modification, as well as complementary and alternative therapies, have been studied in the improvement of knee function, in older adults with KO.^[2,7,9]

Turmeric or *Curcumin longa* is one of the herbs in traditional Iranian medicine. It is a plant of the ginger family. Dried turmeric rhizomes are used for nutritional and medicinal purposes. Curcumin is the active ingredient of turmeric and chemically called diferuloylmethane.^[10,11]

Curcumin is powerful antioxidant and shows anti-cancer, anti-bacterial, antiviral, and anti-inflammatory effects. It also has shown beneficial effects in the liver, gastrointestinal, and cardiovascular disorders, and the improvement of Alzheimer's disease, rheumatoid arthritis, and diabetes mellitus. In traditional Iranian medicine, turmeric is commonly used as a topical poultice in the treatment of strain, sprain, and joint dislocations.[10]

Numerous studies are available on the beneficial effects of curcumin on diabetes, hyperlipidemia, [12] gastrointestinal disorders, [13] liver failure, premenstrual syndrome, [14] skin disorders [15] and episiotomy wound healing. [16] Some studies have also reported the effect of curcumin on the improvement of osteoarthritis in animal models. [17,18] Two studies also examined the effect of oral curcumin in human osteoarthritis. [19,20] A study also used a combination of herbs in older adults with joint pain. [21] However, none of the studies examined the impact of curcumin on the QOL of older adults with KO. In addition, citing the low bioavailability of oral forms of curcumin, a number of studies suggested the necessity of studying the effect of nonedible curcumin methods. [20,22,23]

Considering the lack of studies on its effect on the patients' QOL, this study aimed to investigate the effect of curcumin ointment on the QOL in older adults with KO

METHODS

This study is a double-blind randomized, placebo trial performed from February 20 to June 22, 2019, on 72 older adults with KO aged 60 years and over. The study was conducted in 2019 on patients referring to the office of a physician in Kashan, Iran. Due to the lack of an earlier study on the effect of curcumin on the QOL of older adults, the sample size was estimated using the results of a pilot study in which we used the curcumin ointment on eight older adults with Knee OA. The mean baseline QOL was 58.48 ± 8.37 and changed to 64.52 ± 7.89 after six weeks of use. Therefore, considering the alpha of 0.05 and a power of 0.80, the

sample size was estimated to be 29 per group. However, considering a probable dropout of 20%, we recruited 36 participants in each group.

$$n = \frac{\left(z_{1-\frac{\alpha}{2}+z_{1-\beta}}\right)^{2} \left(\sigma_{1}^{2} + \sigma_{2}^{2}\right)}{\left(\mu_{1} - \mu_{2}\right)^{2}} = n = \frac{(1.96 + 0.84)^{2} \left(s_{3.37}^{2} + 7.89\frac{2}{2}\right)}{\left(58.48 - 64.52\right)^{2}} = 29$$

Inclusion criteria were age 60 years and over, lack of known allergy to the ginger family, lack of a known cognitive disorder, complete consciousness and ability to answer the questionnaire items, having a medical diagnosis of KO, and willingness to participate in the study. Exclusion criteria included any allergic reaction to curcumin ointment, discontinuation of the ointment prior to the end of the intervention, a decision to withdraw from the study, death, hospitalization, and being unavailable due to migration.

Totally, 85 patients were assessed for eligibility, among them 72 were recruited sequentially and were randomly allocated into the study groups [Figure 1]. Prior to sampling, a list of 72 individuals was prepared and then the study supervisor assigned them randomly into 18 quadruple blocks using an online block randomization software. Then, the number and the name of the treatments for each number (curcumin or placebo) were written on small cards and put in small opaque envelopes. The researcher who gathered the data and the physician were blind to the content of the sealed envelopes. Envelopes were arranged in order and provided to the physician. As each participant entered the study, the physician picked and unpacked an envelope and prescribed the predetermined treatment. Thus, the participants were assigned to one of the intervention or placebo groups.

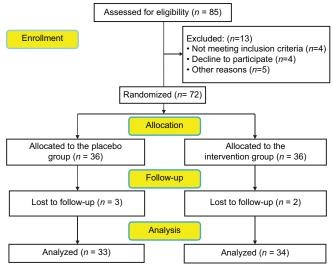


Figure 1: The study flow diagram

Data collection instruments

A two-part instrument was used for data collection. The first part included demographic characteristics such as age, sex, education level, duration of illness, weight (kg), height (cm), body mass index, regular physical activity (yes/no), and the use of prescribed and over-the-counter analgesics, sedatives, and remedies. In addition, there were two questions on perceived pain intensity and perceived knee stiffness that requested the patients to describe the severity of their knee pain and knee stiffness as mild, moderate, severe, or very severe.

The second part included the Osteoarthritis Knee and Hip Quality of Life questionnaire developed by Rat *et al.*^[24] This scale included 43 items in five domains namely physical activity (16 items), mental health (13 items), pain (4 items), social support (4 items), and social performance (3 items). Moreover, there were three independent items on personal relationships, sexual activity, and professional life. Each item is rated on a visual analog scale from 0 (worst condition) to 100 (best condition). To calculate the score for each domain as well as the total score, the sum of scores is divided by the number of questions. Higher scores indicate better OOL.

The Persian translation of this scale has been validated by Saffari *et al.* who examined the construct validity of the translated scale, through simultaneous administration of this scale and SF12 questionnaire on a sample of 434 patients and calculated the correlation coefficient of the subscales of the two instruments between 0.34 and 0.69. The Cronbach's alpha for its various domains has also been reported between 0.74 and 0.89 and its kappa agreement coefficient was 0.85.^[25] As some of the participants were illiterate or low-illiterate, the researcher read them the items and documented their answers into the scale.

Curcumin ointment preparation

For the preparation of curcumin ointment, 100 g of Vaseline and 5 g of the powdered extract of 90% curcumin (purchased from Barij Essence company, Kashan, Iran) were compounded by a pharmacist to make 5% curcumin ointment. Then, the ointment was prepared in 100 g cans similar – in shape and color – to the placebo (Vaseline).

Intervention

Initially, each patient was examined by the physician for inclusion criteria. The patient was then introduced to the first researcher to complete the study instruments through a structured interview. Then, the patient was referred back to the doctor and he picked and unpacked an envelope, read the card inside it, and gave him/her

a can containing the predetermined treatment, and then referred them to the researcher (who was present in the office) to be trained on how to apply it. Patients in the intervention group received curcumin ointment and those in the placebo group received Vaseline ointment, which was prepared in 100-g containers of similar size and color (yellow) and identified by special codes. The use of both substances was similar in both groups. After curcumin ointment or Vaseline ointment was given to patients, the method of administration was individually taught to them. For the first time, the ointment was applied by the researcher on the patient's knee. First, the researcher washed the area with lukewarm water, dried it, and using a special cup took 1.5 mL of the ointment and applied it on and around the knee, and then, warp the area with a clean nonpenetrating cloth to prevent contamination of the patient's clothing. Then, the patient was asked to repeat the procedure on the other knee to ensuring both the researcher and the patient of the correct use. Patients were instructed to do so for 6 weeks, twice daily, once in the morning and once at night, before bed. A checklist was provided to each patient to mark it after each use.

All patients were instructed to report any allergic reaction, redness, or itching around the knee and to avoid any other topical remedies on their knees during the study. A telephone number was obtained from each patient, and the researcher followed the treatment process weekly by telephone. Then, at the end of the 4th and 6th weeks, when the patient was referred to the physician's office for follow-up, the QOL scale was again completed for them by a research assistant who was unaware of the type of medication they were receiving.

Ethical considerations

This study was reviewed and approved by the Research Ethics Committee of Kashan University of Medical Sciences (Code of Ethics: IR.KAUMS.NUHEPM. REC.1397.049, Project number 97157). Moreover, the study was registered in the Iranian Registry of Clinical Trials (code: IRCT20100403003618N6). All patients were informed that they would be contacted several times to evaluate the efficacy of the treatment. However, they did not know which group they were in. All patients were informed about voluntary participation, the right to withdraw at any time, confidential data management, and that the study would not cost them. All patients signed a written informed consent form and their rights were respected in accordance with the Helsinki declaration.

Data analysis

Data were entered into SPSS software version 16 (SPSS, Inc., Chicago, IL, USA), and descriptive methods such as frequency, percentage, mean and standard

deviation (SD) were used to describe the data. The normality of quantitative variables was assessed using the Kolmogorov–Smirnov test, and the normal distribution was confirmed. Chi-square and Fisher's exact tests were used to compare the two groups in terms of nominal and categorical variables, and independent samples *t*-test and analysis of variance with Tukey's *post hoc* test were used to compare the quantitative data. Friedman test was used to examine the changes in perceived knee pain and stiffness over time. Furthermore, repeated-measures analysis was used to compare changes in the QOL scores during the three measurement time points. The Greenhouse–Geisser estimation was also used to correct the degree of freedom. The level of significance was set at <0.05 in all testes.

RESULTS

Of the 72 patients who were recruited, two people of the intervention group and three ones from the placebo group withdrew after the 1st week. Of the total sample, 43 (64.2%) were women, 49 (73.1%) were illiterate or low-literate, and 18 (26.9%) were taking analgesics before starting the study. The mean \pm SD age of the patients was 68.21 ± 6.12 years in the intervention group and 67.02 ± 6.41 years in the placebo group (P = 0.439). The two groups were not significantly different in terms of their individual baseline characteristics, the perceived QOL, knee pain, and knee stiffness [Tables 1 and 2]. The Friedman test also showed that both perceived whee pain and perceived knee stiffness has improved overtime (P < 0.001 for both variables and the both group).

Repeated-measures analysis was performed to compare changes in the QOL scores over the three consecutive measurements. Mauchly's test showed that sphericity was not assumed ($\chi^2[2] = 65.75$, P = 0.001). Therefore, the degrees of freedom were corrected using the Greenhouse–Geisser test. The results showed that over time, the use of curcumin ointment could statistically significantly increase the mean QOL score in the intervention group (F = 50.20, df = 1.240, and P = 0.001).

Given the significant interaction between the measurement time and the type of intervention (group) (F = 38.859, df = 1.252, and P = 0.001), the *t*-test was used to conduct pair-wise comparisons between the two groups at the three measurement time points. The results illustrated that the mean QOL was significantly different between the two groups only at the third measurement (P = 0.006) [Table 3].

Figure 2 shows that the mean overall QOL scores in the intervention group increased over time but after the second measurement, it did not change significantly in the placebo group.

DISCUSSION

At the beginning of the stay, no significant difference was found between the mean QOL scores of the two groups so that both groups possessed about half of the possible score. This finding can be attributed to the fact that most patients in both groups had moderate-to-severe knee pain and stiffness at the beginning of the study. Earlier studies have also shown that chronic pain from KO adversely affects on the older adults' QOL.[26] A study in Brazil also reported that about 65% of patients with KO experience degrees of dependence in life affairs and a decrease in QOL.[27] Another study has also shown that KO reduces the patients' QOL so that people with bilateral KO perceive a lower OOL than those with a unilateral disorder.^[28] It seems that knee pain resulted from KO can significantly decrease the older adults' QOL. This conclusion is consistent with previous studies that showed an inverse relationship between pain – especially chronic pain – and OOL.[29,30] Through appropriate patient care and education, nurses can play

Table 1: Between-group comparisons of the patients' individual and clinical characteristics

Variables	riables Group		
	Placebo	Intervention	
Age, mean \pm SD	67.02 ± 6.41	68.21 ± 6.12	0.439a
Weight (kg), mean \pm SD	75.01 ± 11.23	76.02 ± 12.62	0.647^{a}
Height (cm), mean \pm SD	165.24 ± 6.45	165.63 ± 7.04	0.833^{a}
BMI, mean \pm SD	27.48 ± 3.88	27.52 ± 3.40	0.964^{a}
Disease duration, year, mean \pm SD	6.12 ± 4.02	7.11 ± 4.33	0.336ª
Gender, n (%)			
Female	22 (66.7)	21 (61.8)	0.87^{b}
Male	11 (33.3)	13 (38.2)	
Education level, <i>n</i> (%)			
Illiterate or low-literate	23 (69.7)	26 (76.5)	0.72^{b}
Secondary school and higher	10 (30.3)	8 (23.5)	0.54 ^b
Regular physical activity,			
n (%)			
Yes	16 (48.5)	13 (38.2)	0.54^{b}
No	17 (51.5)	21 (61.8)	
Analgesic use (before the study), n (%)			
Yes	8 (24.2)	10 (29.4)	0.84^{b}
No	25 (75.8)	24 (70.6)	
Use of hypnotics, n (%)			
Yes	13 (39.4)	6 (17.6)	0.08^{b}
No	20 (60.6)	28 (82.4)	

^aT-test, ^bChi-square test. BMI: Body mass index, SD: Standard deviation

Table 2: Comparison of the perceived knee pain and stiffness between the intervention and placebo groups at the three measurements

	Baseline		P 4 th weel	week P	6 th week		P		
	Placebo	Intervention		Placebo	Intervention		Placebo	Intervention	
Perceived knee pain, n (%)									
Mild	2 (6.0)	3 (8.8)	0.790^{a}	1 (3.0)	4 (11.8)	0.120^{b}	3 (9.1)	14 (41.2)	0.009^{b}
Moderate	16 (48.5)	15 (44.1)		18 (54.6)	24 (70.6)		20 (60.6)	15 (44.1)	
Severe	9 (27.3)	12 (35.3)		12 (36.4)	5 (14.7)		8 (24.3)	5 (14.7)	
Very severe	6 (18.2)	4 (11.8)		2 (6.0)	1 (2.9)		2 (6.0)	0	
Perceived knee stiffness, n (%)									
Mild	1 (3.0)	1 (2.9)	0.116^{b}	1 (3.0)	4 (11.8)	0.188^{b}	9 (27.3)	19 (55.9)	0.023^{b}
Moderate	22 (66.7)	15 (44.2)		24 (72.7)	18 (52.9)		22 (66.7)	15 (44.1)	
Severe	8 (24.3)	17 (50)		8 (24.3)	12 (35.3)		2 (6.0)	0	
Very severe	2 (6.0)	1 (2.9)		0	0		0	0	

^aChi-square test, ^bExact χ²

Table 3: Comparisons of the mean quality of life scores at three measurement time points							
Time Me		± SD	P a	P, (RM ANOVA)			
	Intervention	Placebo		The effect of group	The interaction between time and group		
Start of the study	55.98 ± 8.12	58.75 ± 7.86	0.160	0.001	0.001		
After 4 weeks	59.21 ± 7.98	60.01 ± 8.23	0.687				
After 6 weeks	65.83 ± 8.81	60.26 ± 7.25	0.006				
P^{b}	0.0001	0.701					

 ^{a}T -test. $^{b}ANOVA$ (in *post hoc* analysis, in the intervention group showed that the scores at 1st and 2nd measurement did not significantly differ (P<0.102). However, the scores at 1st and 3rd measurements (P<0.001) and also at 2nd and 3rd measurements significantly (P<0.002) differed from each other). SD: Standard deviation

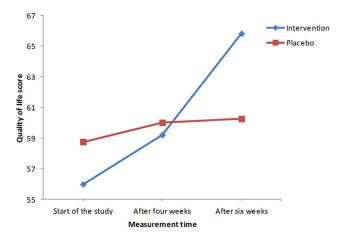


Figure 2: Variations of the overall quality of life at three measurements in the two groups

an important role in the improvement of knee pain in older adults with KO, which consequently improves their QOL.^[31]

The Australian Department of Health emphasized the role of nurses in enhancing the QOL of patients with osteoarthritis, including the role of nurses in helping patients to set realistic short-term and long-term goals; educating patients and the community about osteoarthritis and methods of prevention and treatment; prevention and reduction of adverse drug reactions; follow-up of

the treatment and care recommendations; introduction and recommendation of complementary pain relief therapies appropriate to the patient's condition such as heat therapy, local remedies, sports and appropriate exercises, mind and body therapies, diet and weight loss; use of auxiliary equipment and protective footwear; and introduction of patients to arthritis support groups.^[32]

The present study showed that the QOL scores of both groups increased slightly in the 4th week, although the increase in the intervention group was more than the placebo group. However, after the 4th week, the mean QOL in the placebo group did not change significantly, whereas in the intervention group, it continued to increase by more than six points. Thus, while the QOL score in the placebo group increased only 1.51 from the beginning to the end of the study, the amount of increase was 9.85 in the intervention group. It seems that the topical application of curcumin ointment can gradually increase the OOL in older adults with KO. This effect might be attributable to the gradual effect of curcumin ointment in decreasing knee inflammation and pain. Curcumin is an antioxidant, anti-inflammatory, and anti-swelling agent.[11] Its short-term anti-inflammatory effect is reported to be equal to hydrocortisone and phenylbutazone. [33] It seems that the gradual reduction of inflammation, pain, and swelling of the knee, reduced

its stiffness and improved patients' mobility which consequently improved the QOL of the intervention group.

In the present study, a noticeable change in the QOL of the intervention group began after the 4th week. This can be attributed to the fact that QOL is a mental perception that develops gradually over time and it takes several weeks to observe tangible changes in it. For this reason, QOL studies often question one's QOL over the past few weeks. On the other hand, the delayed changes in the QOL can be attributed to a gradual decrease in patients' perceived knee pain and stiffness. A study of 55 older adults showed that >96% and 64% of them had hip and KO, respectively, and possessed only 55% of the QOL score.^[34]

Pain and decreased QOL compel patients with osteoarthritis to try different methods along with medical treatments to alleviate these problems. A review of 29 studies has shown that these patients use a variety of methods such as acupuncture, tai chi, yoga, and massage therapy.^[35] Some studies have also reported the effects of turmeric^[23] or its extract on pain and other symptoms of arthritis[36] however; they did not focus on the QOL of these patients. A study also examined the effects of a combination of glucosamine sulfate, methyl sulfonyl methane, willow bark extract, ginger, boswellic acid, turmeric root extract, hyaluronic acid and cayenne on pain, joint stiffness, QOL, and physical functioning in middle-aged and older adults with osteoarthritis. This compound was effective in reducing the pain severity and improving the physical functioning but had no significant effect on the patients' QOL.[21] However, the present study showed that topical use of curcumin ointment for 6 weeks could significantly increase the QOL of the intervention group.

This is one of the few studies on the effect of curcumin on knee pain in patients with KO. However, we only studied the patients who referred to a physician's office. Therefore, further multicenter studies with larger samples are recommended. We only followed the patients for 6 weeks, therefore, longer follow-ups are suggested to assess the duration of the effects of curcumin. Although the researcher emphasized to the patients not to disclose their treatment to others, this issue cannot be fully guaranteed.

CONCLUSION

The present study showed that topical administration of 5% curcumin ointment can significantly improves the QOL in older adults with KO. Therefore, the use of this ointment in the treatment of older adults with KO is suggested. Given the side effects of the synthetic drugs used in these patients, curcumin ointment may possibly be used as a supplement to conventional medications to

help reduce the dose of painkillers and anti-inflammatory drugs and improve their pain and QOL.

This study was performed on a small sample of older adults. Given the scarcity of studies on the effect of topical administration of curcumin on QOL, it is recommended to repeat the same studies on larger samples with longer follow-up.

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Conflicts of interest

There are no conflicts of interest.

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