

The effect of local muscle vibration on clinical and biomechanical parameters in people with knee osteoarthritis: A systematic review

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Received: 19 Oct 2020

Published: 27 Sep 2021

Abstract

Background: To identify and synthesize available published studies on the effect of local muscle vibration (LMV) on pain, stiffness, and function in individuals with knee OA.

Methods: Five databases were searched to find relevant papers on April 29, 2020, including, PubMed, Scopus, EMBASE (Ovid), Science Citation Index, and COCHRANE Central Register for Controlled Trials (CENTRAL). Randomized controlled trials (RCTs) and nonrandomized-controlled-trials (non-RCTs), such as interrupted time series and prospective cohort studies were included. Two independent reviewers screened articles and assessed inclusion through predefined criteria. Participants' characteristics, study design, intervention characteristics, outcomes, and main results were collected independently by 2 reviewers. The risk of bias assessment of included studies was conducted using Cochrane risk of bias tools for RCTs and non-RCTs.

Results: Six studies were included: 3 RCTs and 3 non-RCTs. The risk of bias in included studies was generally moderate to high. Improvement of pain, stiffness, and function following the application of LMV were reported in all studies.

Conclusion: This review revealed the promising effect of LMV on pain, stiffness, function, and knee range of motion (ROM) improvements for individuals with knee Osteoarthritis (OA). However, further well-designed studies are required to have a convincing conclusion on the effect of LMV in individuals with knee OA.

Keywords: Knee Osteoarthritis, Local Muscle Vibration, Pain

Conflicts of Interest: None declared

Funding: This study was financially supported by Iran University of Medical Sciences (Grant No. 97-4-6-13280) and the Iranian National Science Foundation (Grant No. 96012163).

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Cite this article as: Barati K, Esfandiari E, Kamyab M, Ebrahimi Takamjani I, Atlasi R, Parnianpour M, Yazdi H, Shahali Sh, Bidari Sh. The effect of local muscle vibration on clinical and biomechanical parameters in people with knee osteoarthritis: A systematic review. *Med J Islam Repub Iran.* 2021 (27 Sep);35:124. <https://doi.org/10.47176/mjiri.35.124>

Introduction

Osteoarthritis (OA) is one of the most common knee joint degenerative diseases (1). The prevalence of knee

OA in the United States was estimated to be 27 million in 2005 and is expected to be 67 million in 2030 (2, 3). The

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↑What is “already known” in this topic:

Knee osteoarthritis is associated with pain, joint stiffness, functional disability, and loss of knee joint range of motion. The application of local muscle vibration is reported to be effective on biomechanical parameters in asymptomatic healthy people. However, there is contradictory evidence on the effect of local muscle vibration on clinical and biomechanical parameters in individuals with knee OA.

→What this article adds:

The results of this study showed that the application of local muscle vibration improves pain, stiffness, function, and knee ROM in people with knee osteoarthritis. However, further well-designed studies are required to have a convincing conclusion on the effect of local muscle vibration in people with knee osteoarthritis.

knee OA causes pain, stiffness, functional disability, loss of knee joint range of motion (ROM), and decreased quadriceps muscle activity (4, 5).

The conservative treatments for rehabilitation of individuals with knee OA consist of weight loss, patient education on coping strategies, exercises, orthotic treatments, and neuromuscular training (6). Neuromuscular training, such as whole-body vibration (WBV) and local muscle vibration (LMV), have been suggested over the past 2 decades to improve muscular activity and functional capacity in knee OA (7-10). The WBV is the most commonly used method and is reported to be effective in the management of complications of knee OA (7, 9, 11). However, a decrease in vibration energy before attaining the target muscle, the high cost of associated equipment, and the limited portability of the WBV platform, are reasons to limit its application in clinical settings (12, 13). The LMV is a method of neuromuscular training that could be directly applied to the targeted muscle belly or muscle-tendon via a lightweight handheld applicator and might be considered as a cost-effective alternative for WBV.

According to the International Classification of Functioning, Disability, and Health (ICF) (14), the main aim of rehabilitation of individuals with knee OA is to decrease the level of disability by improving 'body function' and 'activity and participation components'. Body function is defined as what an individual can do in a standard environment. Additionally, activity is the execution of a task by an individual, and participation is contribution and involvement in a life situation (15). To encompass all aspects of disability, it may be necessary to employ outcome measures in the 2 components. Pain is the most common consequence of knee OA (16) and is associated with increased joint stiffness, decreased knee ROM (17), and decreased muscle function (18). All of these complications are associated with a decrease in physical function (19-21). As a result, these outcomes should be assessed as the main outcomes following intervention in the rehabilitation of individuals with knee OA. In most studies, vibration has been applied in sitting positions (22-26). However, the application of LMV in standing and walking positions has recently been suggested (27). To decide on the application of the LMV in the standing and walking conditions for the rehabilitation of individuals with knee OA, it must be ensured that it does not adversely affect the postural control.

The effect of LMV on biomechanical parameters, such as peak torque, muscle function, and muscle strength, was reported in asymptomatic healthy individuals (13, 27-34). One systematic review synthesized the effect of LMV on muscle strength in asymptomatic healthy individuals (10) and showed that LMV could enhance muscle strength. However, there is contradictory evidence on the effect of LMV on clinical and biomechanical parameters in individuals with knee OA (22, 23, 25, 26). For example, Benedetti et al showed that the application of LMV for 2 weeks would increase the muscular activity in individuals with knee OA. However, Rice et al (26) reported no statistically significant change in the produced torque and quadriceps and hamstring muscle strength.

Therefore, we conducted a systematic review of the peer-reviewed publications to identify and synthesize the available peer-reviewed publications on the effect of LMV on knee joint pain, stiffness, physical function, postural balance, and muscular activity in individuals with knee OA. In addition, we aimed to identify the most important LMV characteristics, such as frequency, the amplitude of displacement, application location, and position of the application.

Methods

Protocol and Registration

The protocol of this systematic review has been formerly issued in PROSPERO with registration No. CRD42019116321. The PRISMA (Preferred Reporting Items for Systematic reviews and Meta-Analyses) guidelines were followed to perform the current review (35).

Eligibility Criteria

The published studies with the following criteria were included in our review: population: individuals with knee OA, grade II to III based on the Kellgren and Lawrence (K&L) grading system (36), aged between 35 to 75 years; intervention: LMV, which was the local application of vibration to the muscle belly or the tendon via an applicator; comparator: fake vibration, exercise, and no treatment; outcomes: pain, stiffness, and function, ROM, electromyography (EMG), and postural response; study designs: randomized controlled trials (RCTs), interrupted time series, and prospective cohort studies. Studies that evaluated the effect of WBV were excluded.

Information Sources

Five databases were searched to find relevant papers on April 29, 2020, including, PubMed, EMBASE (Ovid), Scopus, Science Citation Index, and COCHRANE Central Register for Controlled Trials (CENTRAL). We conducted a manual search of Google scholar with "Allintitle" to find more papers that are relevant. The registry of clinical trials (clinicaltrials.gov), and the World Health Organization (15) international clinical trials registry platform (<http://www.who.int/ictrp/en/>) were searched to ascertain the comprehensiveness of the search. First, the first author searched the PubMed database to develop a search template using key concepts, Medical Subject Headings (MeSH) terms, and keywords. Then, the second author checked the template for comprehensiveness. The search template was used to complete the searches in the remaining databases. We searched the reference list of included papers to find more papers that are relevant. The search was limited to the English language.

Search

The search strategy used for the PubMed search is presented in Table 1. This search strategy was used as a template to do the searches in the remaining databases.

Study Selection

Two independent reviewers (first and second authors) screened the titles and abstracts at screening level 1. The

Table 1. Search Strategies

Databases	Total Found Articles	Search
PubMed	100	("Vibration"[Mesh] OR Vibrat*[TIAB]) AND ("Osteoarthritis, Knee"[Mesh] OR ("Knee Joint"[Mesh] OR "Knee"[Mesh] OR Knee[TIAB] OR Knees[TIAB] OR femorotibial[TIAB]) AND ("Osteoarthritis"[Mesh] OR Osteoarth*[TIAB] OR arthros*[TIAB] OR "Arthritis"[Mesh] OR arthrit*[TIAB] OR osteo-arthritis[TIAB] OR osteo-arthritis[TIAB])) OR gonarthrosis[TIAB])
Scopus	157	TITLE-ABS-KEY((Vibrat*) AND (((Knee OR Knees OR femorotibial) AND (Osteoarth* OR arthros* OR arthrit* OR osteo-arthritis OR osteo-arthritis)) OR gonarthrosis))
Web of Science	176	TS=((Vibrat*) AND (((Knee OR Knees OR femorotibial) AND (Osteoarth* OR arthros* OR arthrit* OR osteo-arthritis OR osteo-arthritis)) OR gonarthrosis))
EMBASE	194	(Vibrat*:ab,ti OR 'vibration'/exp) AND ('knee osteoarthritis'/exp OR ('knee'/exp OR Knee:ab,ti OR Knees:ab,ti OR femorotibial:ab,ti) AND ('osteoarthritis'/exp OR 'arthrits'/exp OR Osteoarth*:ab,ti OR arthros*:ab,ti OR arthrit*:ab,ti OR osteo-arthritis:ab,ti OR osteo-arthritis:ab,ti)) OR gonarthrosis:ab,ti)

third reviewer adjudicated any unresolved discrepancies. Then, 2 reviewers screened the full-texts (level 2). The reason for the exclusion of the papers was recorded.

Data Collection Process

The data extraction was conducted by the first author and the second author reviewed the accuracy of the data. The missing records were requested from the corresponding author with email contact. If we received no response, the author was contacted twice, at 1-week intervals.

Data Extraction

The following information was extracted from each study. The first author, publication year, the country in which the study was conducted, study design, objectives, primary and secondary outcome measures, the main results, and conclusions, participants' characteristics (knee OA grade, number of samples, sex, and mean age), and vibration characteristics (frequency, amplitude of displacement, force, location of LMV application, and duration of LMV application).

Risk of Bias in Individual Studies

Two independent reviewers (first and second authors) performed the risk of bias assessment of the included studies. The disagreements among the reviewers were resolved by discussion with the third reviewer to reach a consensus.

The risk of bias in RCTs was assessed using the Cochrane Risk of Bias (RoB 2.0) tool (37). This tool consisted of 5 domains: randomization process, deviations from intended interventions, missing outcome data, measurement of outcomes, and selection of the reported result. Three judgments were possible for risk of bias: low, some concerns, and high. A summary of bias assessment of included randomized studies is provided in Table 2.

The nonrandomized studies were assessed using the Cochrane Risk Of Bias In Non-randomized Studies-of Interventions (ROBINS-I) tool (38). This tool consisted of 7 domains: confounding, selection of participants into the study, classification of the interventions, deviations from intended interventions, missing data, measurement of outcomes, and selection of the reported result. Five judgments were possible for risk of bias: no information, low, moderate, serious, or critical. A summary of bias assessment of included nonrandomized studies is provided in Table 3.

Synthesis of Results

We planned to conduct a meta-analysis if data were available. If pooling of data was not possible because of methodological heterogeneity (variability in trial design and quality) or clinical heterogeneity (variability in participants and intervention type) among studies (39), we planned to conduct a narrative synthesis of the findings.

Table 2. Risk of Bias Assessment for Randomized Controlled Studies Using the Cochrane Risk of Bias (RoB 2.0) Tool

Study (Year)	Randomization Process	Deviations From Intended Interventions	Missing Outcome Data	Measurement of Outcomes	Selection of the Reported Result	Overall
Benedetti et al	Low risk	Some concerns	Some concerns	Low risk	Some concerns	Some concerns
Rabini et al	Low risk	Low risk	Low risk	Low risk	Some concerns	Some concerns
Kitay et al	Low risk	High risk	Some concerns	Low risk	Some concerns	High risk

Table 3. Risk of Bias Assessment for Nonrandomized Studies Using the Cochrane Risk of Bias in Nonrandomized Studies of Interventions tool

Study (Year)	Confounding	Selection of Participants into the Study	Classification of the Interventions	Deviations From Intended Interventions	Missing Data	Measurement of Outcomes	Selection of the Reported Result	Overall
Rice et al	Serious	Low	Low	Low	Low	Moderate	Low	Serious
Shanahan et al	Low	Low	Low	Low	Low	Moderate	Low	Moderate
Lievens et al	Low	Low	Serious	Low	Low	Serious	Low	Serious

Results

Study Selection

A total of 676 studies were identified through 5 electronic databases (Fig. 1). After removing duplicates and review of titles and abstracts, 27 studies were considered eligible to be included in the full-text screening. Of these, 21 studies were excluded: 3 studies were not available in English, 7 used different interventions, 7 were conference abstracts, 2 used different populations, and 2 records were clinical trial registrations. No study was identified in a manual search of reference lists. Finally, 6 studies were included in the review.

Study Characteristics

The studies were conducted Australia (24), Belgium (40), Italy (22, 23), New Zealand (26), and United States (25). The sample size ranged from 8 to 71, with a mean age of 58.6 to 74.4 years (Table 4). Table 4 details the characteristics of participants.

Four studies included individuals with knee OA (22, 23, 25, 40), and 2 included 2 groups of healthy asymptomatic

individuals and individuals with knee OA (24, 26). Also, 4 studies included established knee OA (≥ 2 based on the K&L grading system) (23-26), and 2 studies did not report knee OA grade (22, 40).

The characteristics of LMV (frequency, displacement amplitude, force, duration, and location of application) used in the studies are summarized in Table 4. The setups of LMV were different; the amplitude and frequency of the LMV were 0.1 (25) to 1.5 mm (26) and 10 (25) to 150 Hz (22), respectively. The LMV application time was 15 seconds (24) to 20 minutes (22, 25, 26, 40). The application of LMV was on the muscles' belly (22, 24), on the infra-patellar tendon (23, 26), and the knee joint (25, 40). The force of LMV was reported to be 20 (22) to 30 Newton (26). The force was not reported in 4 studies (23-25, 40).

Risk of Bias in Individual Studies

Based on the RoB 2 Tool, one of the RCTs was at high risk of bias. Based on the Robins-I tool, 2 non-randomized studies were at serious risk of bias.

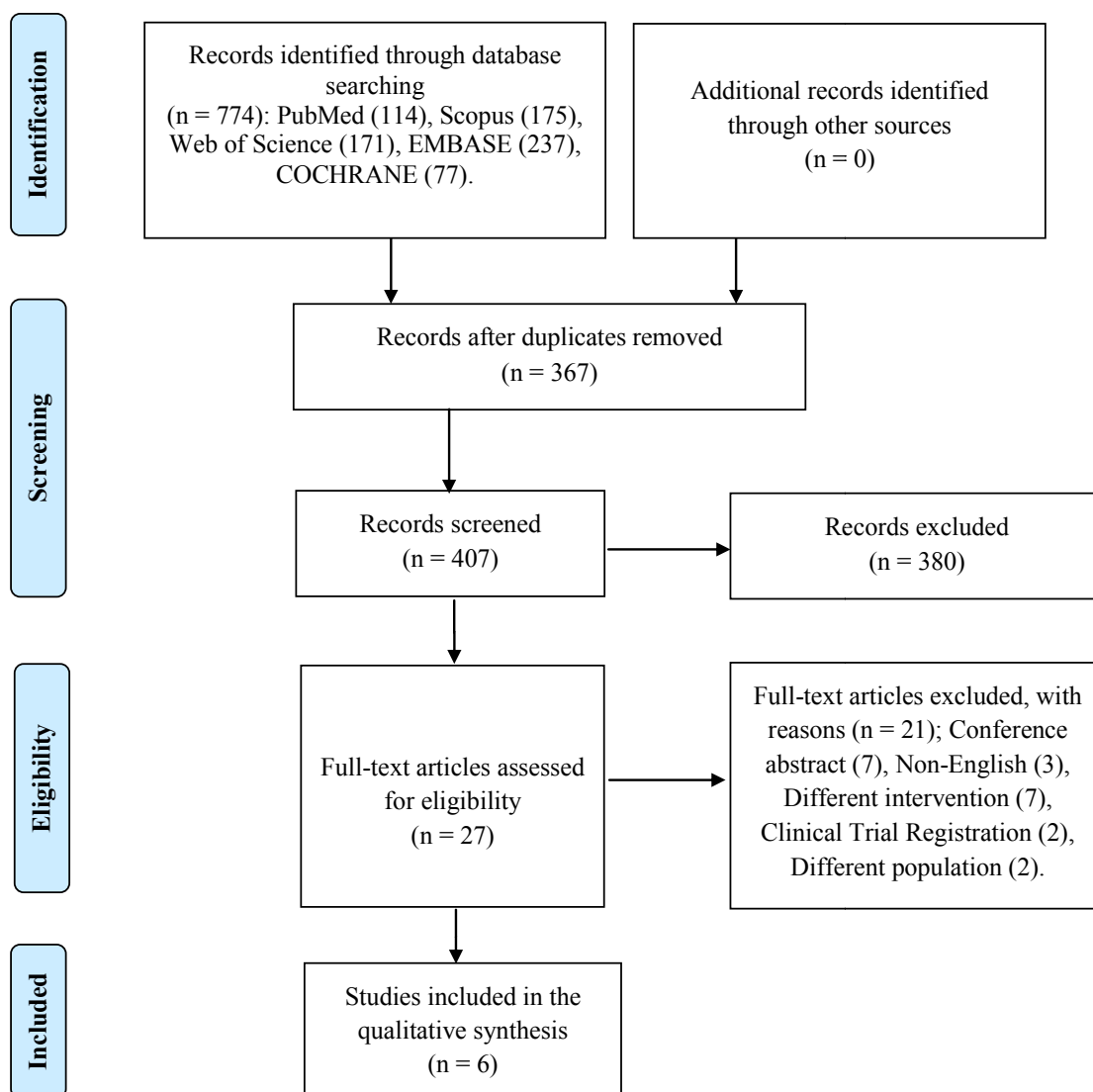


Fig. 1. The PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow diagram)

Table 4. Overview of the Studies Concerning the Effect of Local Muscle Vibration in Patients With Knee Osteoarthritis. Description of Interventions in Included Studies

Author/ Year/ Country	Mean age (years)	Participants	Control Intervention	Vibration Protocol		
				Vibration Parameters	Location of Vibration Application	Duration of Vibration Application
Benedetti et al. 2017, Italy	58.6	- Knee OA participants in the intervention group (n=15) - Knee OA participants in the control group (n=15)	NMES application	F= 150 Hz A= N-m Force= 20-25 N	- Vastus lateralis muscle belly - Vastus medialis muscle belly - Rectus femoris muscle belly	2 weeks, 5 days per week, 1 time each day, 20 min each time.
Rabini et al. 2015, Italy	74.4	- Knee OA participants in the intervention group (n=25) - Knee OA participants in the control group (n=25)	Sham treatment	F= 100 Hz A= 0.2 to 0.5 mm Force= N-m	Close to infrapatellar tendon	3 consecutive days, 3 times each day, 10 min each time, 1 min interruption.
Shanahan et al. 2014, Australia	66	- Knee OA participants treated with LMV (n=30) - Health group treated with LMV (n=30)	Pre/ post vibration application	F= 60 Hz A= 0.5 mm Force= N-m	- Quadriceps muscle belly. - Triceps surae tendon. - Tibialis anterior muscle belly.	1 day, 2 times each day, 15 s each time.
Rice et al. 2011, New Zealand	62.7	- Knee OA participants treated with LMV (n=15) - Health group treated with LMV (n=15)	Pre/ post vibration application	F= 50 Hz A= 1.5 mm Force= 25-30 N	Infrapatellar tendon.	1 day, 1 time each day, 20 min each time.
Kitay et al. 2009, United States	61.4	- Knee OA participants in the intervention group (n= 34) - Knee OA participants in the control group (n=37)	Sham treatment	F= 10Hz, 27 Hz, 42 Hz A= 0.1 mm Force=N-m	- Just above the knee joint. - Just below the knee joint.	4 weeks, 7 days each week, 2 times each day, 20 min each time.
Lievens et al. 1984, Belgium	N-m	- Knee OA participants in the intervention group (n= 8) - Knee OA participants in the control group (n=8)	No-treatment	F= N-m A= .5-1 mm Force= N-m	Just above the knee.	10 days, 1 time each day, 20 min each time.

LMV, local muscle vibration; NMES, neuromuscular electrical stimulation; F, frequency; A, amplitude; Hz, Hertz; min, minute; N, newton; s, second; mm; millimeter; N-m, not mentioned.

The overall risk of bias assessment showed that the risk of bias was high (or serious) in 3 studies (25, 26, 40); of the other 3 studies, 2 (22, 23) had a 'some concerns' risk of bias, and 1 (24) had a moderate risk of bias. In RCTs, the 'randomization process' and 'measurement of outcomes' domains were dominant with a low risk of bias (Table 2). In nonrandomized papers, the 'selection of participants into the study,' 'deviations from intended interventions,' and 'missing data' domains were dominant with a low risk of bias (Table 3). The most common shortcomings in RCTs and nonrandomized studies were the measurement of outcomes and selection of the reported result (Table 2 and Table 3).

Results of Studies

We reviewed the results of studies with respect to therapeutic and biomechanical outcomes: pain, stiffness, function, EMG, ROM, and postural response. A detailed summary of the study results is provided in Table 4 and Table 5.

Pain

Pain was reported in 3 studies (22, 23, 25) using the Western Ontario and McMaster universities arthritis index (WOMAC). Two 'some concern quality' studies found that 3 consecutive days to the 2-week application of LMV improved pain in individuals with knee OA in comparison with individuals with knee OA who received sham treat-

ment (23) or neuromuscular electrical stimulation (NMES) application (22). One 'high-risk quality' study also showed that a 4-week application of LMV improved pain in individuals with knee OA compared to individuals with knee OA who received sham treatment (25). The knee OA grade of participants was ≥ 2 based on the K&L grading system.

Stiffness

The effect of LMV on stiffness was assessed in 3 studies using the WOMAC index (22, 23, 25). One 'some concern quality' study found that 3-day application of LMV improved stiffness in individuals with knee OA compared to individuals high-risk OA who received sham treatment (23). A similar finding was reported in a 'high risk quality' study which assessed the 4-week effect of LMV in individuals with knee OA (25). The stiffness score was not reported separately in 1 'some concern quality' study (22). However, the improvement of the WOMAC total score following a 2-week application of LMV in individuals with knee OA compared to individuals with knee OA who received NMES was reported. The knee OA grade of participants was ≥ 2 based on the K&L grading system (Table 4) (22, 23, 25).

Physical Function

Change in physical function following application of LMV was investigated in 3 studies using the WOMAC

Table 5. Overview of the Results and Conclusion of the Studies Concerning the Effect of Local Muscle Vibration in Patients With Knee Osteoarthritis

Outcome Measure	Studies	Mean (SD)	Mean Difference	P Value	CI	Conclusions
Pain	Kitay et al.	10.5 (3.3)	-	0.001	-	After 4 weeks of application of LMV, pain decreased.
	Benedetti et al.	4 (1.8)	-1.4 (1.6)	0.02	-2.63	After 2 weeks of application of LMV, pain decreased.
	Rabini et al.	2.6 (1.5)	-	-	-	After 3 days of application of LMV, pain decreased.
		57.68 (18.99)	-18.48	0.001	28.85	
Stiffness	Kitay et al.	39.20 (17.45)	-	<	-	After 4 weeks of application of LMV, stiffness decreased.
	Benedetti et al.	4.6 (1.4)	-8.7 (6.9)	0.001	-	After 2 weeks of application of LMV, stiffness decreased.
	Rabini et al.	57.68 (18.99)	-18.48	0.001	21.13	After 3 days of application of LMV, stiffness decreased.
		39.20 (17.45)	-	-	28.85	
Physical function	Kitay et al.	29.1 (1.8)	-	<	-	After 4 weeks of application of LMV, function increased.
	Benedetti et al.	53.1 (17.5)	-8.7 (6.9)	0.001	-	After 2 weeks of application of LMV, function increased.
	Rabini et al.	44.4 (15.7)	-18.48	0.001	21.13	After 3 days of application of LMV, function increased.
		57.68 (18.99)	-	-	28.85	
EMG	Rice et al.	0.13 (0.06)	1.4 %	0.94	-0.04	After 20 min application of LMV, EMG amplitude and isometric torque of quadriceps and hamstring muscles were unchanged.
	Benedetti et al.	0.13 (0.06)	0.2 (0.3)	0.40	-0.27	After 2 weeks of application of LMV, the involvement of type II muscle fiber increased.
	Lievens et al.	3.0 (0.4)	6.2	<0.05	-	After 10 days of application of LMV, active and passive RoM of knee increased.
		3.2 (0.8)	-	-	-	
Knee RoM	Kitay et al.	149.5 (not-reported)	-	0.02	-	After 4 weeks of application of LMV, RoM increased.
	Benedetti et al.	155.7 (not-reported)	4.3	0.001	-1.55	After 2 weeks of application of LMV, RoM increased.
	Shanahan et al.	1.6 (24)	0.2	0.76	-	- CoP displacement increased after application of LMV to triceps surae muscle.
		129.7 (8.5)	-	-	-	- CoP displacement was unchanged after LMV application to quadriceps and tibialis anterior muscles
Postural response		134.0 (7.1)	-	-	-	
		1.9 (not-reported)	-	-	-	
		2.1 (not-reported)	-	-	-	

ROM, range of motion; CoP, center of pressure; LMV, local muscle vibration; EMG, electromyography; SD, standard deviation.

index (22, 23, 25). One 'some concern quality' study on 3-consecutive days of LMV application showed the improvement of physical function in individuals with knee OA compared to individuals high-risk OA who received sham treatment (23). Similar results were reported in a 'high risk quality' study with 4-weeks of LMV application (25). The physical function score was not reported separately in 1 'some concern quality' study (22). However, they reported improvement in the WOMAC total score following a 2-week application of LMV in individuals with knee OA compared to individuals with knee OA who received NMES. The knee OA grade of participants was ≥ 2 based on the K&L grading system.

Electromyography

One 'some-concern quality' study (22) (based on RoB 2.0 tool) and 1 'serious-risk quality' study (26) (based on ROBINS-I tool) evaluated the muscular activity of quadriceps and hamstrings muscles following application of LMV using EMG. Rice et al (26) reported that 1-day application of LMV (20 minutes) to infer-patellar tendon has no significant effect on the EMG amplitude of quadriceps muscle in individuals with knee OA compared to asymptomatic healthy individuals who received LMV. Benedetti et al (22) reported that 2-week application of LMV to vastus lateralis could lead to a decrease in mean frequency of the power spectrum and muscle fiber conduction veloc-

ity of individuals with knee OA compared to individuals with knee OA who received NMES.

Range of Motion

One 'some-concern quality' (22), 1 'high-risk quality' study (25) (based on RoB 2.0 tool), and 1 'serious risk quality' study (40) (based on ROBINS-I tool) evaluated the effect of LMV on knee ROM in individuals with knee OA. Consistently across 3 studies, knee ROM was increased following the application of LMV. Application of LMV to quadriceps muscle belly induced an increase in knee ROM in individuals with knee OA compared to individuals with knee OA who received NMES (22). Application of LMV to surfaces above and below of the knee joint induced an increase in knee ROM in individuals with knee OA compared to individuals with knee OA who received sham treatment (25). Application of LMV to only 1 surface above the knee joint induced an increase in knee ROM in individuals with knee OA compared to individuals with knee OA who received no treatment (40). The duration of the intervention varied from 3 consecutive days (3 times each day) (22) to 10 days (1 time each day) (40), and 4 weeks (14 times each week) (25).

Postural Response

Only one 'moderate-risk quality' study evaluated the effect of LMV on postural control in individuals with knee

OA using the displacement of the center of pressure (CoP) (24). The results suggested that anterior-posterior displacement of CoP had no change following 2-time (15 seconds each time) application of LMV to quadriceps and tibialis anterior muscles in individuals with knee OA compared to asymptomatic healthy individuals who received LMV. However, the application of LMV on triceps surae would result in more CoP displacement in individuals with knee OA compared to asymptomatic healthy individuals.

Discussion

Summary of Evidence

This systematic review synthesized the evidence on the clinical and biomechanical effect of LMV in individuals with knee OA. This review highlighted that the number of published studies and the quality of studies assessed the effect of LMV on pain, stiffness, function, EMG, and knee ROM in individuals with knee OA are limited and generally in weak quality. As a result, we cannot draw any definite conclusion. More studies with a stronger level of evidence are required to evaluate the effects of LMV on people with knee OA.

The WOMAC questionnaire could be used to evaluate pain, stiffness, and physical function in individuals with knee OA (41). All 3 studies on the WOMAC score consistently reported an improvement in pain, stiffness, and physical function following the application of LMV (22, 23, 25). The previous investigation suggested that gains in clinical and biomechanical parameters following application of the LMV are results of enhanced spinal-mediated reflex activity (42) and cortical excitability (27). The results of our study demonstrate that application of LMV showed the promising effect on clinical outcomes for individuals with knee OA, despite the difference in vibration frequency, duration, and displacement amplitude. The results suggest that the difference in intervention characteristics may have a minimum impact on LMV efficacy on pain, stiffness, and function as long as the vibration frequency is in the range of 10 Hz to 150 Hz, the displacement amplitude in the range of 0.2 to 1.5 mm, and the duration of application is in range of 1 to 20 minutes. While the risk of bias was generally moderate to high, the results should be interpreted with caution.

The quadriceps weakness is reported in people with knee OA (43, 44). Physiotherapy and land-based exercise have been reported to improve muscle strength in people with knee OA, but the presence of pain limits active exercise (45, 46). Physical modalities, such as LMV, have been suggested to facilitate physiotherapy interventions to improve muscle activation and strength. The previous study reported that the quadriceps muscle activation level remains elevated for 5 minutes after application of the LMV (13). The controversies in the effect of LMV on muscle strength could be related to the sample of the study and the duration of intervention (22, 26). Given the equal vibration duration (20 minutes) (22, 26), different follow-up duration, vibration frequency, vibration force, vibration displacement amplitude, and location of application could

explain the controversy among the results. However, it can be suggested that improving muscle performance in individuals with knee OA needs prolonged application of LMV to gain a positive result. In summary, given the limited number of studies that assessed the effect of LMV on muscle strength, it is unclear which parameters in the LMV application are more decisive to improve muscle function in individuals with knee OA, and further well-designed studies are required to have a definitive conclusion.

The promising effect of LMV on the improvement of knee ROM was shown for individuals with knee OA (22, 25, 40). While the limitation of ROM is associated with an increasing level of disability in individuals with knee OA (47), it could be suggested that the application of LMV can lead to a reduced level of disability in individuals with knee OA (22, 25, 40). However, 3 studies with moderate to high risk of bias are not sufficient to make a conclusive decision regarding the effect of LMV application on knee ROM in individuals with knee OA. These studies provide preliminary evidence to pursue further research.

Application of vibration to tendon or muscle belly may cause the illusion of movement (48, 49), or falling when applied to leg muscles in standing position or during walking (50, 51). Therefore, the application of LMV may be associated with compensatory strategies for compromised postural control for individuals with knee OA (51). On the other hand, the application of LMV to knee muscles in standing and walking conditions is suggested for the rehabilitation of people with knee OA (27). As a result, evaluating the effect of LMV on postural control has a key role in deciding whether to use LMV in walking/standing conditions for the rehabilitation of people with knee OA.

The available evidence reported no significant change in anterior-posterior displacement of CoP following application of LMV to quadriceps and tibialis anterior muscles. However, increased CoP perturbation was detected following the application of LMV to the triceps surae muscle. These results suggest that the application of LMV to the quadriceps muscle, as the most predictive muscle in the progression of knee OA (52), in standing and walking conditions could be implemented for the rehabilitation of people with knee OA. However, there is only 1 study with a moderate risk of bias available. Thus, caution is needed when interpreting these results.

Limitations

There are some limitations in this systematic review: First, despite the inclusion of 5 databases to identify the relevant evidence, we may have missed some studies; second, because of the limited number of studies and heterogeneity of study designs, knee OA grades, and the outcomes, we were not able to provide a quantitative synthesis of the evidence; third, limiting this systematic review to studies in the English language excluded evidence in other languages.

Conclusion

This review revealed the promising effect of LMV on

pain, stiffness, physical function, and knee ROM improvements for individuals with knee OA. However, this review highlighted a knowledge gap for high-quality evidence on the effect of LMV on clinical and biomechanical outcomes in individuals with knee OA.

Funding

This study was financially supported by Iran University of Medical Sciences (Grant No. 97-4-15-14030) and the Iranian National Science Foundation (Grant No. 96012164).

Conflict of Interests

The authors declare that they have no competing interests.

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