Review Article

The Effect of Kinesitherapy on Bone Mineral Density in Primary Osteoporosis: A Systematic Review and Meta-Analysis of Randomized Controlled Trial

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Objective. Osteoporosis (OP) is a well-established age-related disease, pathologically characterized by bone microarchitectural deterioration, increased fragility, and low BMD. Primary osteoporosis (POP) is the most common type of OP. Methods. Publications pertaining to the effectiveness of kinesitherapy on BMD in POP from PubMed, SCI, Cochrane Library, Embase, VIP, CNKI, and Wanfang Database were retrieved from their inception to October 2019. Results. A total of 21 studies with 1840 participants were included. The results of the meta-analysis revealed that kinesitherapy plus antiosteoporosis medications had a positive effect on lumbar spine BMD when the duration of intervention was 6 months (MD = 0.11 g/cm²; 95% CI: 0.06–0.15; \( P < 0.0001 \)) or >6 months (MD = 0.04 g/cm²; 95% CI: 0.02–0.06; \( P < 0.0001 \)) compared with antiosteoporosis medications alone. Additional kinesitherapy plus antiosteoporosis medications were associated with improved femoral neck BMD compared with antiosteoporosis medications alone (MD = 0.09 g/cm²; 95% CI: 0.03–0.16; \( P = 0.004 \)). Conclusions. Kinesitherapy plus antiosteoporosis medications significantly improved lumbar spine and femoral neck BMD in the current low-quality evidence. Additional high-quality evidence is required to confirm the effect of kinesitherapy on BMD in patients with POP.

1. Introduction

With age, bone mass declines, and an accelerated loss of bone mineral density (BMD) occurs by 50 years of age [1]. Osteoporosis (OP) is a well-established age-related disease pathologically characterized by bone microarchitectural deterioration, increased fragility, and low BMD [2]. The diagnosis of OP is based on measurements of BMD and is defined by the World Health Organization as BMD ≥ 2.5 standard deviations (SD) below the average value for young healthy individuals [3]. Moreover, OP is associated with high morbidity and mortality, as well as reduced quality of life, which in turn increases the rate of fractures, healthcare costs, and social economic stress [4–9]. In North America [10, 11], over 55 million people are at risk of developing OP or osteopenia. OP is typically classified into three main categories: (1) primary; (2) secondary; and (3) idiopathic. Primary osteoporosis (POP) refers to the natural aging process of human tissue and organ systems, and the symptoms are associated with degenerative changes in the skeletal system. Moreover, POP is the most common type of OP, accounting for 90% of OP cases, and includes women with postmenopausal osteoporosis (PMOP) and senile osteoporosis (SOP) [12, 13]. PMOP is primarily related to postmenopausal estrogen deficiency, whereas SOP is associated with increased age [14]. Thus, as the proportion of elderly populations increases throughout the world, the number of POP cases will also increase gradually. In the European Union, according to relevant statistics [15, 16], 22 million women and 5.5 million men over the age of 50 suffer from osteoporosis. And that number is expected to increase by 23 percent by 2025, according to a study. One study found that the overall prevalence rate of osteoporosis was 32.1% in at least one measurement site (28.5% in the lumbar and
14.5% in the femoral region), while 49.7% of elderly people suffer from decreased bone mass (osteopenia) in Amirkola, north of Iran [17]. According to the latest epidemiological results of osteoporosis in China, the prevalence of osteoporosis at the age of 40–49 is 3.2%, including 2.2% in males and 4.3% in females. The prevalence rate of osteoporosis over the age of 50 was 19.2%, including 6.0% in males and 32.1% in females. And the prevalence of osteoporosis over the age of 65 was 2.0%, with 0.7% of men and 51.6% of women [18].

POP treatment is a long-term process and may not 100% prevent the development or reverse the symptoms of the disease [19]. In addition, exercise is one of the key recommendations for the prevention and treatment of bone loss [20, 21]. Several studies have demonstrated that exercise can prevent bone loss [22] and improve calcium absorption, bone formation [23], and the secretion of sex hormones [24, 25], which then promotes BMD [26]. Kinesitherapy, as a part of physical therapy, is a comprehensive exercise that represents one of the most important aspects of medical rehabilitation. Kinesitherapy involves the movement of various parts of the body or the entire body using exercises to maintain, establish, develop, and change the function of the locomotor apparatus and organs in locomotion. Kinesitherapy for the treatment of POP primarily includes routine static training, walking training, grip training, outbreak and endurance exercise training, push-ups, stretching, or isometric exercise. In addition, multiple reviews have confirmed that exercise reduces bone loss and increases BMD in postmenopausal women or PMOP [27–29]. A meta-analysis also demonstrated that exercise can improve functional outcomes, including mobility, balance, and self-reported measures of functioning in persons with OP [30]. However, there was no systematic review to evaluate the effect of kinesitherapy on BMD in patients with POP. Therefore, the aim of this study was to conduct a systematic review and meta-analysis to assess the effect of kinesitherapy in persons with POP on lumbar spine and femoral neck BMD via conducting a maximal search of both Chinese and English databases.

2. Methods

2.1. Eligibility Criteria. Available human, clinical, or community studies with a randomized controlled trial published in English or Chinese were included in this review. The participants consisted of patients with POP who had no thoracolumbar vertebral fracture and other complications such as heart, vein, lung, liver, and kidney as well as metabolic diseases and were not taking drugs affecting bone metabolism. The age and gender of the subjects were not limited. The included studies focused on the effect of kinesitherapy plus antosteoporosis medications therapy as a kinesitherapy group compared with antosteoporosis medications therapy as a control group for the BMD of POP (SOP and SMOP). Those which compared kinesitherapy alone with another exercise or any other antosteoporosis intervention were excluded. The kinesitherapy should include weight-bearing, impact, resistance, endurance training, or a combination of these types of training, and only single-motion experiments will be ruled out. Health education can be added to all patients, and all inpatients can be given routine care. The outcomes included at least lumbar spine or femoral neck BMD.

2.2. Data Sources and Searches. The original research articles were obtained after a search of six electronic English and Chinese databases, which included PubMed, Science Citation Index (SCI), EMBASE, Chinese Scientific Journal Database (VIP), China National Knowledge Information (CNKI) database, and Wanfang from their inception to October 3, 2019. We used the following search strategy (kinesitherapy OR exercise) AND osteoporosis AND bone mineral density) in all the English and Chinese databases.

2.3. Study Identification and Quality Assessment. Two reviewers (WSX and LSZ) independently screened and reviewed the title and abstract of the searched studies using NoteExpress V3.2.0.6992 software. The full text of the studies that potentially met the eligibility criteria was obtained, and any potentially relevant references were retrieved according to predefined eligibility criteria. The data were extracted by one reviewer (WSX) using the prepared forms and checked for accuracy by the second reviewer (LSZ). The details extracted from the eligible literature included the language of publication, participant characteristics, sample size, methodological information, participant demographics, experimental and control interventions (category, intensity, frequency, duration, and details of antosteoporosis medication treatment), outcomes, and adverse effects [31]. Studies published in multiple reports were only included once to avoid duplication in this review. Disagreements were resolved through discussion, and the original author was contacted if an agreement could not be reached. The primary outcomes were lumbar spine BMD and femoral neck BMD, which were expressed as g/cm² assessed by dual X-ray absorptiometry or dual photo absorptiometry. The baseline and follow-up data pertaining to BMD were calculated. If follow-up data could not be obtained, the data at the end of the intervention were used instead.

The quality of the studies was independently evaluated by two reviewers (WSX and LSZ) using the Cochrane Collaboration’s tool for assessing the risk of bias [32]. The following recommended domains were considered: selection bias (random sequence generation and allocation concealment); performance bias (blinding of participants and personnel); detection bias (blinding of outcome assessment); attrition bias (incomplete outcome data); reporting bias (selective reporting); and other bias, each of which was rated according to the level of bias and categorized as either low, high, or unclear.

2.4. Data Analysis. Review Manager 5.2 software from the Cochrane Collaboration was applied for the data analysis. Statistical heterogeneity among the studies was assessed using a chi-square test or by calculating Higgins $I^2$ values.
of the studies used a random number table, three of these studies described the generation of random sequences. Five of these studies used simple random methods, three studies used computer-generated random numbers, and the remaining one trial used the method of lottery. Three trials involve allocation concealment. However, the blind intervention associated with the intervention exercises cannot be implemented blindly. One study described that the data analysis was based on the author’s own statistics. One study described an exit from the case, but did not explain the reason. There were no dropouts indicated or explanations for withdrawal in the remaining studies. All of the included studies were considered to have a high risk of bias.

3. Results

3.1. Description of Studies. A detailed screening flowchart depicting the generation of eligible articles is presented in Figure 1. A total of 791 records were identified via database searches. After removing duplicates, 507 remained to be screened for eligibility. Consequently, 21 studies met the inclusion criteria and were included in the meta-analysis.

Table 1 presents the sample, intervention, and outcome characteristics. This review involved a total of 1840 POP patients (including SOP and PMOP). Part of the subjects were from outpatient clinics, inpatient settings, and community or physical examination centers, with the exception of eleven subjects for whom the sources were unknown. In the included studies, the subject type consisted of eleven PMOP patients, five POP patients, and five SOP patients. In one RCT, the diagnostic criteria for the study were in accordance with the WHO diagnostic criteria for osteoporosis. In the seven RCTs, the reference diagnostic criteria for osteoporosis were Chinese, with one diagnostic criterion for osteoporosis in Japan; the Chinese diagnostic criteria included the Chinese recommended diagnostic criteria for osteoporosis, the Chinese medical association osteoporosis diagnosis and treatment guidelines (a second draft) and a primary bone guide formulated by Chinese Medical Association Osteoporosis and Bone Mineral Disease Branch; one RCT used Western diagnostic criteria for disease; four RCTs only mentioned the diagnostic criteria in accordance with the diagnostic criteria of PMOP or POP; and five RCTs used the laboratory examination in which the T value of was less than or equal to −2.5 SD in at least one site; the diagnostic criteria in the remaining RCT did not elaborate. The whole RCTs compared kinesitherapy plus medication treatment with medication treatment alone. Kinesitherapy involves comprehensive exercises rather than individual exercises. Of those RCTs, only one RCT used traditional Chinese medicine (e.g., kidney method), and the remaining RCTs were treated with Western medicine. All of the RCTs included a measure of lumbar spine BMD, and five of the RCTs included lumbar spine and femoral neck BMD.

3.2. Methodological Quality. As shown in Figure 2, twelve studies described the generation of random sequences. Five of these studies used a random number table, three of these studies used simple random methods, three studies used computer-generated random numbers, and the remaining one trial used the method of lottery. Three trials involve allocation concealment. However, the blind intervention associated with the intervention exercises cannot be implemented blindly. One study described that the data analysis was based on the author’s own statistics. One study described an exit from the case, but did not explain the reason. There were no dropouts indicated or explanations for withdrawal in the remaining studies. All of the included studies were considered to have a high risk of bias.

3.3. Meta-Analysis of Lumbar Spine BMD. All the controls in the literatures were kinesitherapy plus antosteoporosis drugs versus antosteoporosis drugs. According to the duration of the intervention, the subgroups were divided into three groups based on an intervention duration of (1) less than 6 months; (2) 6 months; and (3) longer than 6 months.

3.3.1. Intervention Duration < 6 Months. Two trials [35, 53] compared the effect of kinesitherapy plus antosteoporosis medications with antosteoporosis medications alone on lumbar spine BMD when the duration of intervention was less than 6 months. The meta-analyses indicated that there was no significant difference between the two groups (MD = 0.02; 95% CI: −0.00–0.05; P = 0.10) (Figure 3).

3.3.2. Intervention Duration = 6 Months. Ten studies [37, 38, 41, 46, 51, 54] involving 699 participants reported that after 6 months, kinesitherapy had significantly increased lumbar BMD (MD = 0.11 g/cm²; 95% CI: 0.06–0.15; P < 0.0001). However, the heterogeneity among the studies was substantial with $I^2 = 83\%$, and no obvious changes were observed after the sensitivity analysis when any one or two of the studies were removed (Figure 4).

3.3.3. Intervention Duration > 6 Months. Eleven studies [34, 36, 39, 40, 42–44, 48–50, 52] involving 1019 participants revealed that when the duration of treatment was longer than 6 months, the lumbar spine BMD in the kinesitherapy group significantly increased compared with the control group (MD = 0.04 g/cm²; 95% CI: 0.02–0.06; P < 0.0001) with high heterogeneity ($I^2 = 73\%$) (Figure 5). The heterogeneity was $I^2 = 23\%$ after the sensitivity analysis when any one study [40] was removed.

3.4. Meta-Analysis of Femoral Neck BMD. Five trials [43–45, 47, 53] involving 439 participants compared effect of kinesitherapy plus antosteoporosis medications with antosteoporosis medications alone on lumbar spine BMD. The meta-analysis revealed a significant antosteoporosis effect on lumbar spine BMD (MD = 0.09 g/cm²; 95% CI: 0.03–0.16; P = 0.004) but with high heterogeneity ($I^2 = 90\%$) (Figure 6). It showed low heterogeneity ($I^2 = 0\%$) after the sensitivity analysis when two studies were removed [45, 47].
284 records were excluded which were duplication (according to title and author names) or conference papers

469 articles were excluded by reading their abstract: unacceptable participants, \( n = 171 \); irrelevant intervention, \( n = 40 \); unqualified control type, \( n = 46 \); irrelevant outcome assessment, \( n = 67 \); narrative reviews, \( n = 109 \); zoopery, \( n = 19 \); and protocol, \( n = 17 \).

1 article identified from scanning references of relevant narrative review; 5 articles did not contain full text.

13 articles were excluded: data cannot be extracted, \( n = 6 \); incomplete data (undefined intervention duration), \( n = 5 \); and substandard intervention methods, \( n = 2 \).

Studies included in quantitative synthesis (meta-analysis), \( n = 21 \).

**Table 1:** The characteristics of all the trails.

| Author, year | Participants (type, source, age, sample) | Duration (months) | Kinesitherapy group | Control group | Outcomes |
|--------------|------------------------------------------|-------------------|---------------------|---------------|----------|
| Iwamoto et al., 2001 [34] | PMOP, unspecified, 53–77 years, 28 (KT: 8, CON: 20) | 24 | Brisk walking (1000 steps in the first 7 days, increase the step count by 30%/week) + gymnastic training (no details provided) + CON | Calcium lactate (2.0 g, Qd) and 1α-hydroxyvitamin D3 (1 µg, Qd) | Lumbar spine BMD |
| Shen, 2003 [35] | POP, outpatient and inpatient, 45–80 years, 60 (KT: 30, CON: 30) | 3 | Aerobics, tai chi, dance, yangko, jogging, walking etc. (30–60 min/time, 5–7 times/week) + CON | Tonifying kidney granules (3 times/day, 1 dose/time) | Lumbar spine BMD |
| Zhu, 2007 [36] | SOP, outpatient, 60–72 years, 96 (KT: 48, CON: 48) | 12 | Walking, jogging, tai chi (30–60 min/time, 3–5 times/week) + CON | Calcium (600 mg/d) | Lumbar spine BMD |
| Liu et al., 2007 [37] | PMOP, outpatient, 48–61 years, 68 (KT: 36, CON: 32) | 6 | Draft movement, abdominal isometric exercises, flexion, and extension of the upper limbs (20 minutes each time, once every 3 days) + CON | Caltrate D (600 mg, Qd) | Lumbar spine BMD |
| Li et al., 2008 [38] | PMOP, unspecified, 48–61 years, 70 (KT: 38, CON: 32) | 6 | Draft movement, abdominal isometric exercises, flexion, and extension of the upper limbs (20 min/time, once every four days) + CON | Ossotide injection (50 mg, Qd, 20 days in total) | Lumbar spine BMD |
| Liu et al., 2009 [39] | SOP, outpatient service, 60–94 years, 60 (KT: 30, CON: 30) | 12 | Walking, jogging, tai chi (60 min/time, 1 time/day) + CON | Fosamax (10 mg, once a day) and calcium (600 mg/d) | Lumbar spine BMD |
| Author, year | Participants (type, source, age, sample) | Duration (months) | Intervention | Control group | Outcomes |
|--------------|-----------------------------------------|------------------|-------------|---------------|----------|
| Liu and Wang, 2012 [40] | SOP, unspecified, 60–81 years, 320 (KT: 162, CON: 158) | 12 | Tai chi and jogging (no less than 30 min/time, no less than 4 times/week) + CON | Calcium carbonate D3 (600 mg, Qd) | Lumbar spine BMD |
| Li et al., 2013 [41] | SOP, hospital, 67 ± 4 years, 86 (KT:43, CON: 43) | 6 | Progressive lumbar dorsal muscle function exercise includes sitting training, swallowing training and five-point support training (1-2 times/day) + CON | Lorelli calcium capsule (1 capsules, 1 time/d for 2 consecutive months) | Lumbar spine BMD |
| Ming, 2013 [42] | SOP, hospital, 60–78 years, 96 (KT: 48, CON: 48) | 12 | Walking, aerobics, running, and tai chi (5 to 7 times a week for 45 to 60 minutes) + CON | Calcium gluconate (20 ml/time, 3 times/day) and vitamin D (400 units, 2 times/day) | Lumbar spine BMD |
| Chen, 2015 [43] | PMOP, clinic, 53–70 years, 57 (KT: 27, CON: 30) | 12 | Brisk walking (15–30 minutes), resistance strength exercises (15–20 minutes), and balance and flexibility exercises (simplify tai chi and five birds, 15–20 min) + CON | Adequate calcium and vitamin D supplementation and bisphosphonate therapy | Lumbar spine and femoral neck BMD |
| Dischereit et al., 2016 [44] | PMOP, unspecified, 68 years, 42(KT: 25, CON: 17) | 24 | Endurance and strength training program (3 sessions once weekly, 65 min) + CON | | |
| Li et al., 2016 [45] | PMOP, unspecified, 52–76 years, 188 (KT: 94, CON: 94) | 6 | Mainly includes walking, aerobics, running, and tai chi (30–60 min/time, more than 3 times/week) + CON | Caltrate D (1000 mg, Qd), derivatives, vitamin D, and raloxifene (1 pill, Qd) | Lumbar spine and femoral neck BMD |
| Chen, 2016 [46] | PMOP, unspecified, 53–77 years, 65(KT: 33, CON:32) | 6 | Brisk walking and tai chi (30–50 min/time, 2-3 times/week) + CON | Alendronate (70 mg, Qd), caltrate D (600 mg, Qd), and alfacalcidol soft capsules (0.25 μg, Qd) | Lumbar spine BMD |
| Xu, 2017 [47] | PMOP, unspecified, 51–67 years, 100 (KT:50, CON:50) | 6 | Aerobics, tai chi, and jogging (more than 30 min, more than 3 times/week) + CON | Calcine D (2 times/day, 2pills/time) + estrogen (1 time/day, 60 mg/time) | Lumbar spine and femoral neck BMD |
| Chang, 2017 [48] | POP, unspecified, 60–79 years, 84 (KT: 42, CON: 42) | 12 | Aerobics, walking, swimming, jogging, and cycling (3–4 times, not less than 2 times, each exercise 30–60 minutes) + CON | Calcine D 600 (1 tablet once, 2 times a day) and health education | Lumbar spine BMD |
| Qi, 2017 [49] | PMOP, community healthcare center, 45–65 years, 56 (KT: 28, CON:28) | 12 | Aerobic exercise resistance group, load bearing, and stretching (30 min/time, 3–5 times/week, more than 1 h) + CON | Conventional treatment | Lumbar spine BMD |
| Wen, 2017 [50] | POP, unspecified, 60–78 years, 96 (KT: 48, CON: 48) | 12 | Jogging, tai chi, etc. (daily) + CON | Routine prevention and taking medicine | Lumbar spine BMD |
| Liu and Yang, 2018 [51] | POP, unspecified, 36–79 years, 80 (KT: 42, CON: 40) | 6 | Walking, fitness running, ballroom dancing, and swimming (at least 12 times a month, each time ≥30 min) + CON | Calcium and vitamin D3 | Lumbar spine BMD |
| Zheng et al., 2019 [52] | POP, unspecified, 53–77 years, 84 (KT: 42, CON: 42) | 12 | Walking, jogging, alternate running, cycling, etc. (3 to 4 times per week, the minimum 2 times, 30–60 min) + CON | Calcium (300 mg/tablet, 2 times/d, 1 tablet/time) | Lumbar spine BMD |
| Li et al., 2019 [53] | PMOP, unspecified, 50–65 years, 52 (KT: 26, CON: 26) | 3 | Brisk walking (30 min, once a day, 4d/week) and resistance training (week 1, 2 weekly complete 1 set (15 times/set), and then add 1 set every 2 weeks) + CON | Calcium carbonate D3 (600 mg, 1 time/d), alfacalcidol soft capsule (0.5 g, 1time/d), and sodium alendronate (70 mg, 1 time/d/weeks) for 3 months | Lumbar spine and femoral neck BMD |
| Yan et al., 2019 [54] | POP, unspecified, 53–77 years, 52 (KT: 26, CON: 26) | 6 | Flexible resistance exercise therapy to exercise the lumbar and dorsal muscles (5 times/week) + CON | Calcium carbonate D3 (600 mg), vitamin D3 (0.25 UG), health education, and routine nursing | Lumbar spine BMD |

PMOP, postmenopausal osteoporosis; POP, primary osteoporosis; SOP, senile osteoporosis; KT, kinesitherapy group; CON, control group; BMD, bone mineral density.
Study or subgroup | Kinesitherapy | Control | Weight (% | Mean difference | Mean difference |
|------------------|--------------|---------|-----------|----------------|----------------|
| Li R, et al. 2019 | 0.746        | 0.062   | 26        | 0.709          | 0.065          |
| Shen CF 2003     | 0.739        | 0.096   | 30        | 0.738          | 0.073          |
| Li CX, et al. 2013| 0.02         | 0.1     | 40        | 0.09           | 0.09           |
| Xu ML 2017       | 1.04         | 0.18    | 50        | 0.06           | 0.15           |
| Yan Y J, et al. 2019 | 0.841     | 0.075   | 21        | 0.771          | 0.069          |
| Total (95% CI)   |              |         | 56        | 100.0          | 0.02 [-0.00, 0.05] |

Heterogeneity: chi^2 = 1.63, df = 1 (P = 0.20); I^2 = 39%
Test for overall effect: Z = 1.67 (P = 0.10)

Figure 2: Risk of bias summary for each included study.

Study or subgroup | Kinesitherapy | Control | Weight (%) | Mean difference | Mean difference |
|------------------|--------------|---------|------------|----------------|----------------|
| Chen C, 2016     | 0.802        | 0.081   | 33         | 0.766          | 0.088          |
| Li CX, et al. 2013| 1.012        | 0.114   | 43         | 0.855          | 0.103          |
| Li HB, et al. 2008| 0.53         | 0.19    | 38         | 0.48           | 0.22           |
| Li XQ, et al. 2016| 0.02         | 0.1     | 94         | 0.84           | 0.16           |
| Liu HQ, et al. 2007 | 0.52      | 0.17    | 36         | 0.45           | 0.22           |
| Liu M, et al. 2018 | 0.83        | 0.11    | 40         | 0.75           | 0.09           |
| Xu ML, 2017      | 1.04         | 0.18    | 50         | 0.08           | 0.15           |
| Yan YJ, et al. 2019 | 0.841     | 0.075   | 21         | 0.771          | 0.069          |
| Total (95% CI)   |              |         | 355        | 344            | 0.11 [0.06, 0.15] |

Heterogeneity: tau^2 = 0.00; chi^2 = 41.00, df = 7 (P < 0.00001); I^2 = 83%
Test for overall effect: Z = 4.64 (P < 0.00001)

Figure 3: Kinesitherapy plus antiosteoporosis medications versus antiosteoporosis medications on lumbar spine BMD (intervention duration < 6 months).

4. Discussion

4.1. Summary. POP is a worldwide health problem that primarily impacts postmenopausal women and senile individuals. Moreover, POP is often related to physical frailty, an increased risk of falls, substantial morbidity, mortality, and impairment in quality of life [55]. The aim of OP treatment is to improve BMD and prevent fractures. Nonpharmacological treatment includes a healthy diet, prevention of falls, and physical exercise programs. Pharmacological treatment involves calcium, vitamin D, and medications for activating bone tissue (e.g., antiresorptives, bone formers, and mixed agents) [56]. In addition, exercise is considered important for maintaining bone health. Individuals with OP are strongly recommended to regularly engage in multicomponent exercise programs [57]. Moreover, several studies have confirmed that exercise can increase BMD at the femoral neck and the lumbar spine in elderly women with osteoporosis [58, 59]. This review is the first systematic review and meta-analysis to evaluate the effect of kinesitherapy on BMD on the lumbar spine and femoral neck in persons with POP from RCTs. This study
involved 21 RCTs that included a total of 1840 subjects with POP (including SOP and PMOP). The duration of treatment varied from 3 to 24 months. The outcome measures primarily consisted of lumbar spine and femoral neck BMD. The results of the meta-analysis showed that there were no statistically significant differences between kinesitherapy plus antosteoporosis medications versus antiosteoporosis medications alone on lumbar spine BMD when the duration of intervention was less than 6 months (MD = 0.02; 95% CI: −0.00–0.05; \( P = 0.10 \)). However, kinesitherapy had a positive effect on lumbar spine BMD when the duration of intervention was 6 months (MD = 0.11 g/cm²; 95% CI: 0.06–0.16; \( P < 0.0001 \)) or more than 6 months (MD = 0.04 g/cm²; 95% CI: 0.02–0.06; \( P < 0.0001 \)) compared with antosteoporosis medications alone. Furthermore, kinesitherapy had a remarkable effect on femoral neck BMD (MD = 0.09 g/cm²; 95% CI: 0.03–0.16; \( P < 0.004 \)) when compared with antosteoporosis medications alone.

4.2. Limitations and Suggestions for Future Research. A total of 21 RCTs were included in this review, which showed that kinesitherapy had a favourable effect on lumbar spine and femoral neck BMD in patients with POP. Nevertheless, the interpretation and generalization of this systematic review and meta-analysis are subject to some limitations. According to the Cochrane Collaboration’s tool, low-quality evidence, which included studies with a high risk of bias, resulted in a high heterogeneity of the meta-analysis results and favoured the positive effect of kinesitherapy on BMD in patients with POP. There were eleven RCTs that did not report the random sequence generation, and the remaining RCTs were lacking detailed descriptions of randomization, which could result in selection bias. The performance bias was high since the positive effect of kinesitherapy on BMD in patients with POP was observed in eleven RCTs that did not report the random sequence generation, and the remaining RCTs were lacking detailed descriptions of randomization, which could result in selection bias. The performance bias was high since the positive effect of kinesitherapy on BMD in patients with POP was observed in eleven RCTs that did not report the random sequence generation, and the remaining RCTs were lacking detailed descriptions of randomization, which could result in selection bias.

**Figure 5:** Kinesitherapy plus antosteoporosis medications versus antosteoporosis medications on lumbar spine BMD (intervention duration > 6 months).

**Figure 6:** Meta-analysis of femoral neck BMD.
studies; thus, the reliability of the treatment effects of kinesitherapy on femoral neck BMD is reduced. Therefore, more multicenter, larger sample, long-term, single-blind RCTs are required to assess the effect of kinesitherapy on BMD in patients with POP.

5. Conclusion

The meta-analyses in this study suggest that kinesitherapy plus antiosteoporosis medications can significantly improve lumbar spine BMD when the duration of intervention is longer than 6 months compared with antiosteoporosis medications alone in the current low-quality evidence. More high-quality evidence in the form of multicenter, larger sample, long-term, single-blind, randomized controlled trials is required to confirm the effect of kinesitherapy on BMD in patients with POP.

Conflicts of Interest

The authors have no conflicts of interest to declare.

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