



### Efficacy of the Pilates method for pain and disability in patients with chronic nonspecific low back pain: a systematic review with meta-analysis

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ABSTRACT | **Objective:** To systematically review the available evidence on the efficacy of the Pilates method in patients with chronic nonspecific low back pain. **Method:** Searches were performed in MEDLINE, EMBASE, PEDro, SciELO, LILACS, CINAHL and CENTRAL in March 2013. Randomized controlled trials that tested the effectiveness of the Pilates method (against a nontreatment group, minimal intervention or other types of interventions) in adults with chronic low back pain were included regardless the language of publication. The outcome data were extracted from the eligible studies and were combined using a meta-analysis approach. **Results:** The searches identified a total of 1,545 articles. From these, eight trials were considered eligible, and seven trials were combined in the meta-analysis. The comparison groups were as follows: Pilates versus other types of exercises (n=2 trials), and Pilates versus no treatment group or minimal intervention (n=4 trials) for short term pain; Pilates versus minimal intervention for short-term disability (n=4). We determined that Pilates was not better than other types of exercises for reducing pain intensity. However, Pilates was better than a minimal intervention for reducing short-term pain and disability (pain: pooled mean difference=1.6 points; 95% CI 1.4 to 1.8; disability: pooled mean difference=5.2 points; 95% CI 4.3 to 6.1). **Conclusions:** Pilates was not better than other types of exercises with chronic low back pain. Pilates was not better than other types of exercises with chronic low back pain. Pilates was not better than other types of pain and disability in patients with chronic low back pain. Pilates was not better than other types of exercises for reducing low back pain. Pilates was better than a minimal intervention for reducing pain and disability in patients with chronic low back pain. Pilates was not better than other types of exercises for short-term pain and disability in patients with chronic low back pain. Pilates was better than a minimal intervention

Keywords: backache; exercise therapy; rehabilitation.

#### HOW TO CITE THIS ARTICLE

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#### Introduction

Low back pain is a major health and socioeconomic problem and one of the main causes of disability and absenteeism<sup>1.4</sup>. It is estimated that between 11 and 84% of adults will experience an episode of low back pain at least once in their lives<sup>5</sup>, and approximately 40% of these patients will develop chronic low back pain lasting more than three months<sup>4,6,7</sup>. A systematic review of the prognosis of patients with low back pain has shown that both patients with acute low back pain and those with persistent low back pain improved within the first six weeks. However, after this period, the improvement slowed down, and low to moderate levels of pain and disability were present at one year, especially in patients with persistent pain<sup>8</sup>. Estimates show that low back pain leads to an expenditure of nine billion Australian dollars in Australia and 12 billion British pounds in the United Kingdom when direct and indirect costs are taken into account<sup>9</sup>.

The effect of the treatments recommended by guidelines for chronic low back pain is moderate<sup>4,10</sup>. Exercise is the physical therapy treatment for chronic low back pain that shows the most lasting and positive effects<sup>10</sup>. Pilates method exercises involve contractions of the deep abdominal muscles. When used in physical therapy practice, the method requires modifications, such as adaptations and simplifications of the traditional method. Such exercises are thus described as Pilates method-based exercises<sup>11</sup>.

For some time, Pilates method-based exercises were considered to have unknown value in the

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treatment of low back pain<sup>10</sup>. However, randomized controlled studies on this method began to be published in 2006<sup>12-14</sup>. Two systematic reviews without meta-analysis have been published on the use of the Pilates method to treat patients with chronic low back pain<sup>3,15</sup>, and the results showed improvement in pain and disability. In 2011, two systematic reviews with meta-analyses were published regarding Pilates method-based exercises<sup>16,17</sup>. The results of the first review<sup>16</sup> showed that Pilates method-based exercises are superior to a minimal intervention for reducing chronic nonspecific low back pain after intervention. Conversely, the second<sup>17</sup> review showed no significant improvement in pain and disability when groups that participated in Pilates methodbased exercises were compared with a control group or group performing lumbar stabilization exercises.

Although four systematic reviews have been published recently, the authors included studies that had not yet been published as journal articles indexed in databases that go through rigorous peer review (especially master dissertations or doctoral theses)<sup>16,17</sup>. They also included studies of patients with discopathies<sup>15</sup>, which can directly interfere with the results of systematic reviews. The objective of this study was therefore to systematically review randomized controlled trials comparing the effectiveness of the Pilates method with a control group, other types of intervention or other types of exercises in terms of pain and disability in patients with chronic nonspecific low back pain.

#### Method

#### **Inclusion criteria**

The inclusion criteria for the analysis were studies that were randomized controlled trials of adults with chronic nonspecific low back pain that evaluated pain and/or disability; studies in which the primary treatment was based on Pilates method exercises compared with no treatment, minimal intervention, other types of intervention or other types of exercises; studies published in scientific journals between 1980 and 2013 and without restriction on language of publication.

#### Search strategy

Searches were conducted in the MEDLINE, EMBASE, PEDro, SciELO [Scientific Electronic Library Online], LILACS [Literatura Latino Americana em Ciências da Saúde (Latin American and Caribbean Health Sciences Literature)], CINAHL [Cumulative Index to Nursing and Allied Health Literature] and CENTRAL [Cochrane Central Register of Controlled Trials] databases. The terms used in the searches were based on the search strategies of the *Cochrane Back Review Group* (Appendix 1). The last day of the search for articles was March 10, 2013. The searches were adjusted for each database because of the differences between the search engines of each of those databases. These adjustments followed the recommendations of the Cochrane Collaboration<sup>18</sup>.

#### **Study selection**

Two independent reviewers performed the first analysis based on the information provided by the title, abstract and keywords. In cases of disagreement between the reviewers, a third reviewer was consulted to achieve consensus.

## Evaluation of the methodological quality of the studies

The methodological quality and statistical description of the studies were measured using the PEDro quality scale<sup>19-21</sup> and these data were extracted from the PEDro database (www.pedro.org.au). The PEDro scale consists of the following items: 1) Were the eligibility criteria specified?, 2) Were the subjects randomly allocated to groups?, 3) Was the allocation of the subjects concealed?, 4) Were the groups similar at baseline with regard to the most important prognostic indicators?, 5) Was there blinding of all the subjects who participated in the study?, 6) Did all therapists who administered the therapy do so blindly?, 7) Was there blinding of all assessors who measured at least one key outcome?, 8) Were measurements of at least one key outcome obtained for more than 85% of subjects initially allocated to the groups?, 9) Did all subjects for whom outcomes were measured receive the treatment or control condition as allocated or, when this was not the case, was an "intention to treat" data analysis performed for at least one of the key outcomes?, 10) Were the results of between-group statistical comparisons reported for at least one key outcome?, and 11) Did the study have both point measures and measures of variability for at least one key outcome?<sup>22</sup>.

PEDro quality scale scoring is between 0 and 10 points; the first item, relating to external validity, is not used to calculate the PEDro score. The reliability of the items on this scale ranged from good to

excellent<sup>22,23</sup>. Studies that were not available on the PEDro database were evaluated by two independent reviewers. In cases of disagreement between the reviewers, a third reviewer was consulted to achieve consensus.

#### **Data extraction**

To analyze the studies, data were extracted regarding participants' characteristics (age and duration of symptoms), intervention, duration and frequency of treatment sessions based on the Pilates method, the control group (no treatment, minimal intervention, other types of intervention or other types of exercises) and pain and disability outcomes at all assessment time-points. Meta-analysis was not performed when there was only one study for comparison or when statistical heterogeneity was identified.

#### Statistical analysis of the meta-analysis

For the meta-analysis, data extraction was performed based on the data presented in the included articles, with data extracted from the text, tables or figures and expressed as mean and standard deviation (Corel Draw software Version 3 was used to extract data from figures). This procedure was performed for the outcomes short-term pain and disability; the Pilates method was compared with no treatment, minimal intervention, other types of intervention or other types of exercises. Sensitivity analyses were performed to identify studies with high statistical heterogeneity and to determine whether the methodological quality of the eligible articles and the number of sessions conducted influenced the size of the observed effects. The study was removed from the meta-analysis when the presence of statistical heterogeneity and/or an effect of methodological quality or number of sessions were detected. An analysis of publication bias was also performed via the visual inspection of funnel plots<sup>18</sup>. Studies with larger samples had a higher weight in the metaanalysis results<sup>24</sup>. Means and standard deviations were converted into scales from 0 to 10 points for the pain outcome and 0-100 points for disability, standardizing the scales that other studies used to perform the meta-analysis. I2 was calculated using Comprehensive Meta Analysis software Version 2 for a statistical analysis of heterogeneity, which describes the percentage of variation in the estimates of effects caused by heterogeneity rather than sampling error. A value above 50% can be considered substantial

heterogeneity. When the values were statistically homogeneous, the average effects (difference between the weighted averages) were calculated using a fixed-effect model (I<sup>2</sup><50%). When the values were statistically heterogeneous, the estimates of average effects (difference between the weighted averages) were obtained using a random-effects model (I<sup>2</sup>>50%). The standardized mean difference with a confidence interval of 95% was used. Forest plots, funnel plots and meta-analyses were also calculated using the Comprehensive Meta Analysis software.

#### Results

#### Study selection and characteristics

A total of 1,545 articles were found, of which eight studies<sup>13,14,25-30</sup> were considered eligible. Of these, seven<sup>13,14,26-30</sup> were included in the meta-analysis (Figure 1). Two studies were excluded from the meta-analysis: one study<sup>25</sup> evaluated three different regimens of the Pilates method and therefore had no comparison group for analysis, and another study<sup>13</sup> was excluded from the meta-analysis for the pain outcome based on the sensitivity analysis because its inclusion considerably increased the level of heterogeneity of the meta-analysis. In one study<sup>26</sup> comparing the vertical ground-reaction force between the control group (healthy subjects) and the group with low back pain (divided into two groups: Pilates group and no treatment group), only the groups of patients with low back pain were considered. Table 1 shows the characteristics of the studies in descriptive form, and Table 2 shows the details of the studies.

#### Methodological quality results

In evaluating the methodological quality using the PEDro scale, the scores for the seven articles were already available on the PEDro database<sup>13,14,25,26,28-30</sup>. One article<sup>27</sup> was evaluated by two independent reviewers because it was still not available. The scores ranged from 2 to 8 points on a scale of 0 to 10 points (Table 3). All of the studies lost points on items related to the blinding of the patient and therapist, and only four studies<sup>13,14,27,30</sup> blinded the assessor. A score of 2 points was observed in only one study<sup>25</sup>, and a score of 8 points, the maximum considered for this type of study as it is not possible to blind therapist and patients, was observed in two studies<sup>14,30</sup>.

#### **Treatment effects**

The visual inspection of the funnel plot did not identify publication bias for any of the three metaanalyses. The sensitivity analyses considering the methodological quality and the number of sessions conducted did not significantly influence the results of the meta-analyses. Therefore, no study was removed from the meta-analyses. It was not possible to analyze the influence of the type of exercises on the metaanalyses because the descriptions of the exercises performed in eligible studies were very brief.

## *Comparison of the Pilates method with minimal intervention for pain outcome*

Four studies<sup>14,26,27,30</sup> evaluated pain before and after intervention (n=171). The study by Rydeard et al.<sup>14</sup> compared the Pilates method with primary care, such

as medical appointments when needed (n=39). Pilates method-based floor exercises were performed initially as static exercises; they progressed to exercises involving hip extension movements and then to exercises on the Reformer, with 12 one-hour sessions conducted in addition to a home-based program of 15 minutes, six days a week for four weeks. Da Fonseca et al.'s study<sup>26</sup> compared the Pilates method with no exercise (n=17). In that study, floor exercises were carried out based on the Pilates method. These exercises involved limb movement with a static back position progressing to a dynamic back position; onehour sessions were held twice a week for a total of 15 sessions. Quinn et al.'s study<sup>27</sup> compared the Pilates method with no treatment (n=29). In this study, Pilates exercises were performed based on O'Brien's

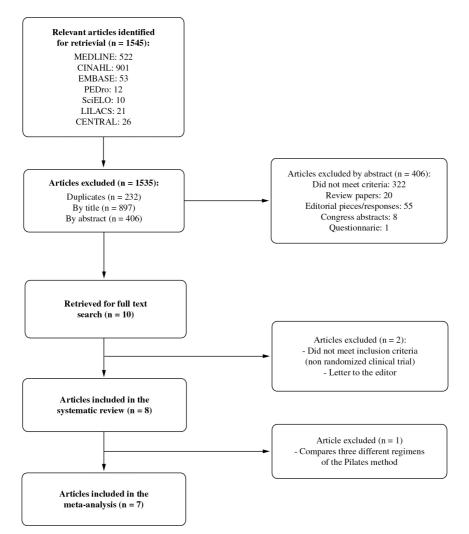


Figure 1. Selection process for studies included in the analysis.

Table 1.	Study	characteristics.

Authors	Participants	Intervention
Gladwell et al. <sup>13</sup> ***	Mean age: CG: 45.9 years; PG: 36.9 years n=34 Gender: CG: 4 male and 10 female; PG: 3 male and 17 female Duration of symptoms (mean±SD): CG: 139±148; PG: 115±101 months	s; CG: Normal activities and pain relief PG: Pilates method-based floor exercises with strengthening of abdominal, gluteus and back muscles and progression with movement of the limbs, 6 sessions with 1 hour of duration for 6 weeks
Rydeard et al. <sup>14*.***</sup>	Mean age: CG: 34 years; PG: 37 years; n=39 Gender: CG: 8 male and 13 female; PG: 6 male and 12 female Duration of symptoms (median (interquartile range)): CG: 108 (12-240) months; PG: 66 (6-324) months	CG: Normal care, such as medical appointments when necessary PG: Static Pilates method-based-exercises in floor with the following progression: addition of hip extension and exercises in the Reformer, 12 sessions with 1 hour of duration and a home- based program (duration of 15 minutes) for 4 weeks
Rajpal et al. <sup>28</sup> **	Mean age: CG: 21.6 years; PG: 22.1 years n=32 Gender: female Duration of symptoms: >3 months in both groups	CG: McKenzie for sitting and standing posture correction, 3 repetitions performed 15-20 times
Curnow et al. <sup>25</sup>	No specification of the participants characteristics; n=39	<ul> <li>18 sessions for 6 weeks</li> <li>The 3 groups performed 4 Pilates-based</li> <li>exercises on the floor; 2 exercises involved</li> <li>abdominal contraction, one involved lateral</li> <li>elevation of lower limbs in supine and one</li> <li>involved back muscles contraction, 40</li> <li>repetitions of each exercise.</li> <li>PG A: no additional exercises</li> <li>PG B: exercises and relaxation, during 3-5</li> <li>minutes</li> <li>PG C: exercises, relaxation (for 3-5 minutes)</li> <li>and posture training involving hip flexion and</li> <li>eccentric psoas contraction (20 repetitions)</li> </ul>
Da Fonseca et al. <sup>26</sup> *	Mean age: CG: 34.4 years; PG: 31.6 years n=17 Gender: 5 male and 12 female Duration of symptoms: >6 months in both groups	s; CG: No intervention PG: Pilates method-based floor exercises (with the activation of deep abdominal muscles), with the following progression: lower limb movements with static back position and dynamic back position, 15 sessions of 1 hour duration, 2 sessions per week
Quinn et al. <sup>27*,***</sup>	Mean age: CG: 44.1 years; PG: 41.8 years n=29 Gender: female Duration of symptoms (mean±SD): CG: 49.2 (49.2) months; PG: 57.6 (38.4) months	s; CG: No intervention PG: Pilates method-based exercises based on the Body Control Pilates program <sup>31</sup> , 6 to 8 1-hour sessions and 15 minutes at home, 5 times per week for 8 weeks

\*CG: control group; PG: Pilates group. \*Studies included in the comparison between Pilates and minimal intervention considering pain at short term. \*\*Studies included in the comparison between Pilates and other types of exercises considering pain at short term. \*\*\*Studies included in the comparison between Pilates and minimal intervention considering disability at short term.

#### Table 1. Continued ...

Authors	Participants	Intervention
Wajswelner et al. <sup>29</sup> **	Mean age: CG: 48.9 years; PG: 49.3 years n=87 Gender: CG: 20 male and 23 female; PG: 19 male and 25 female Duration of symptoms (mean±SD): CG:14.2±12.7; GP: 13.6±14.2 years	s; 12 sessions, 1-hour duration, for 6 weeks, with exercises prescribed at home CG: General exercises used in the treatment (e.g., stationary bike, stretching, resistance training) and 4 exercises performed at home PG: 6 to 12 Pilates method-based exercises in the Reformer and Cadillac and 1 to 4 Pilates method-based exercises performed at home on the floor or with support of a chair or wall, involving back exercises in all planes of movement
Miyamoto et al. <sup>30*,***</sup>	Mean age: CG: 38,3 years; PG: 40,7 years n=86 Gender: CG: 9 male and 34 female; PG: 7 male and 36 female Duration of symptoms (mean±SD): CG: 56.7±53.5; PG: 73.3±79.6 months	s; 12 sessions for 6 weeks, 2 times per week GC: Educational booklet and 12 telephone calls during a 6-week period GP: Pilates method-based floor exercises involving deep abdominal muscle contraction with stretching or strengthening of the back and lower limb muscles; 12 sessions, 1-hour duration, for 6 weeks

\*CG: control group; PG: Pilates group. \*Studies included in the comparison between Pilates and minimal intervention considering pain at short term. \*\*Studies included in the comparison between Pilates and other types of exercises considering pain at short term. \*\*\*Studies included in the comparison between Pilates and other types of exercises considering pain at short term. \*\*\*Studies included in the comparison between Pilates and other types of exercises considering pain at short term. \*\*\*Studies included in the comparison between Pilates and other types of exercises considering pain at short term. \*\*\*Studies included in the comparison between Pilates and other types of exercises considering pain at short term. \*\*\*Studies included in the comparison between Pilates and minimal intervention considering disability at short term.

Body Control Pilates program<sup>31</sup>, which consists of six to eight sessions lasting one hour and 15 minutes at home, five times a week at home for eight weeks. The study by Miyamoto et al.<sup>30</sup> compared Pilates method treatment with giving patients an educational booklet (n=86). The Pilates method-based floor exercises involved deep abdominal muscle activation exercises and exercises to strengthen and stretch the spinal and lower limb muscles, with 12 one-hour sessions over six weeks.

For the meta-analysis of this comparison, a fixed-effect model was used (I<sup>2</sup>=0%,  $\chi^2$ =1.16, df=3, P=0.76) in which a statistically significant difference was found between the groups when the Pilates method was compared with no treatment or minimal intervention for pain in the short term (difference between means=1.6 points, 95% CI 1.4 to 1.8; Figure 2A).

## Comparison of Pilates method with other types of exercises for the pain outcome

For this analysis, two studies<sup>28,29</sup> were included in the meta-analysis (n = 119). Rajpal et al.'s study<sup>28</sup> compared the Pilates method with the McKenzie method (n=32). In this study, Pilates method-based floor exercises were used and were associated with breathing exercises with deep abdominal muscle contraction in the supine, quadruped and sitting positions, with 10 repetitions of 10 seconds performed in each position. These were compared with McKenzie exercises, which involve sitting and standing posture correction, with three repetitions performed 15 to 20 times per day. Wajswelner et al.'s study<sup>29</sup> compared the Pilates method with general exercises (stationary bike, stretching and resistance training; n=87), in which six to 12 supervised Pilates method-based exercises were performed on the Reformer and Cadillac, and one to four floor exercises were performed at home over 12 one-hour sessions for six weeks.

For the meta-analysis of this comparison, a fixed-effect model was used (I<sup>2</sup>=0%,  $\chi^2$ =0.69, df=1, P=0.41). At short term, no statistically significant difference was found between exercises for pain (difference between means=0.1 points, 95% CI -0.3 to 0.6; Figure 2B).

# *Comparison of the Pilates method with minimal intervention for the disability outcome*

For disability, four studies that compared the Pilates method with minimal intervention<sup>13,14,27,30</sup> using pre- and postintervention evaluations were included in the meta-analysis (n=188). The study by

Table 2. Details of the e	extraction of data	Table 2. Details of the extraction of data from studies, in mean and standard deviation.	tion.		
Authors	Participants	Before treatment	After treatment	6 months after randomization	Results
Gladwell et al. <sup>13</sup>	CG: 14 PG: 20	CG: Pain RMVAS: 2.4 (0.9); Disability OSWDQ: 24.1 (13.4) PG: Pain RMVAS: 2.7 (0.9); Disability OSWDQ: 19.7 (9.8)	CG: Pain RMVAS: 2.4 (0.8); Disability OSWDQ: 18.1 (13.0) CG: Pain RMVAS: 2.2 (0.9); Disability OSWDQ: 18.1 (11.2)	ł	Statistically significant improvement in pain (p <0.05) but not disability
Rydeard et al. <sup>14</sup>	CG: 18 PG: 21	CG: Pain: NRS-101: 30.4 (1.8); Disability RM: 4.2 (3.4) PG: Pain NRS-101: 23.0 (1.8); Disability RM: 3,1 (2.8)	CG: Pain NRS-101: 33.9 (1.5); Disability RM: 3.2 (0.4) PG: Pain NRS-101: 18.3 (1.5); Disability RM: 2.0 (0.3)	ł	Statistically significant improvement in pain (p=0.002) and disability (p=0.023)
Rajpal et al. <sup>28</sup>	CG: 15 PG: 17	CG: Pain VAS: 4.4 (1.5) PG: Pain VAS: 5.6 (0.7)	CG: Pain VAS: 2.1 (0.8) PG: Pain VAS: 2.1 (0.7)		Statistically significant improvement in pain (p<0.05)
Curnow et al. <sup>25</sup>	PG A: 13 PG B: 14 PG C: 12	Without values	Without values	-	Only statistically significant difference between Group A and B, with greater improvement in Group B (p=0.02)
Da Fonseca et al. <sup>26</sup>	CG: 9 PG: 8	CG: Pain VAS: 6.1 (1.8) PG: Pain VAS: 5.9 (2.0)	CG: Pain VAS: 4.9 (2.5) PG: Pain VAS: 3.0 (3.4)	1	Statistically significant improvement in pain (p<0.05)
Quinn et al. <sup>27</sup>	CG: 14 PG: 15	CG: Pain VAS -101: 39.9 (19.9); Disability RM: 7.7 (5.0) PG: Pain VAS -101: 40.4 (14.6); Disability RM: 6.9 (4.6)	CG: Pain VAS -101: 44.6 (14.7); Disability RM: 7.5 (1.7) PG: Pain VAS -101: 30.9. (15.2); Disability RM: 5.4 (1.0)	-	Statistically significant improvement in pain (p<0.05) but not disability
Wajswelner et al. <sup>29</sup>	CG: 43 PG: 44	CG: Pain NRS: 4.6 (1.8); Disability Quebec: 23.9 (14.0) PG: Pain NRS: 4.9 (1.6); Disability Quebec: 28.1 (11.4)		CG: Pain NRS: 3.2 (2.1); Disability CG: Pain NRS: 2.2 (1.7); Disability No statistically significant difference Quebec: 17.1 (13.4) Quebec: 13.0 (11.4) for any comparison at short and PG: Pain NRS: 2.8 (1.6); Disability PG: Pain NRS: 2.5 (1.8); Disability Quebec: 15.3 (9.1) Quebec: 14.1 (10.4)	No statistically significant difference for any comparison at short and medium term
Miyamoto et al. <sup>30</sup>	CG: 43 PG: 43	GC: Pain NRS: 6.5 (1.7); Disability RM: 10.5 (5.4) GP: Pain NRS: 6.6 (1.5); Disability RM: 9.7 (4.5)	<ul> <li>GC: Pain NRS: 6.5 (1.7); Disability GC: Pain NRS: 5.2 (2.3); Disability GC: Pain NRS: 5.3 (2.3); Disability RM: 10.5 (5.4)</li> <li>RM: 10.5 (5.4)</li> <li>RM: 7.1 (5.7)</li> <li>RM: 6.7 (5.6)</li> <li>GP: Pain NRS: 6.6 (1.5); Disability GP: Pain NRS: 3.1 (2.3); Disability GP: Pain NRS: 4.5 (2.2); Disability RM: 9.7 (4.5)</li> </ul>	GC: Pain NRS: 5.3 (2.3); Disability Statistically significant improvement RM: 6.7 (5.6) in pain and disability (p<0.05) at GP: Pain NRS: 4.5 (2.2); Disability short term but not at medium term RM: 4.5 (4.5)	Statistically significant improvement in pain and disability (p<0.05) at short term but not at medium term
*CG: control group; PG: pain rating visual analogi visual analogue scale (0-	Pilates group; V A ue scale (0-10) for 100) for pain evalı	*CG: control group; PG: Pilates group; VAS: visual analogue scale (0-10) for pain evaluation; OSWDQ: Oswestry Low Back Pain Disability Questionnaire (0-100) for disability evaluation; RMVAS: Roland Morris pain rating visual analogue scale (0-10) for pain evaluation; NRS: 101-point numerical rating scale (0-100) for pain evaluation; RM: Roland Morris Disability Questionnaire (0-24) for disability evaluation; VAS-101: visual analogue scale (0-100) for pain evaluation; RM: Roland Morris Disability Questionnaire (0-24) for disability evaluation; VAS-101: visual analogue scale (0-100) for pain evaluation; NRS: numerical rating scale (0-10) for pain evaluation; RM: Roland Morris Disability Questionnaire (0-24) for disability evaluation; VAS-101: visual analogue scale (0-100) for pain evaluation; NRS: numerical rating scale (0-10) for pain evaluation.	luation; OSWDQ: Oswestry Low Back Pair rating scale (0-100) for pain evaluation; RN or pain evaluation.	n Disability Questionnaire (0-100) for disa 4: Roland Morris Disability Questionnaire	(0-24) for disability evaluation; RMV AS: Roland Morris (0-24) for disability evaluation; VAS-101:

logical quality.			86		Da			05
Criteria	Gladwell et al.	Gladwell et al. <sup>13</sup> Rydeard et al. <sup>14</sup> Rajpal et al. <sup>28</sup>	<sup>+</sup> Rajpal et al. <sup>28</sup>	Curnow et al. <sup>25</sup>	Fonseca et al. <sup>26</sup>		Quinn et al. <sup>27</sup> Wajswelner et al. <sup>29</sup> Miyamoto et al. $^{30}$	amoto et al. <sup>3(</sup>
1. Eligibility criteria were specified	yes	yes	yes	ou	yes	yes	yes	yes
2. Subjects were randomly allocated to groups (in a crossover study, subjects were randomly allocated an order in which treatments were received)	yes	yes	yes	yes	yes	yes	yes	yes
3. Allocation was concealed	ou	yes	no	0U	00	yes	yes	yes
4. The groups were similar at baseline regarding the most important prognostic indicators	yes	yes	Ю	Ю	yes	yes	yes	yes
5. There was blinding of all subjects	ou	0U	no	0U	ou	ou	no	ou
6. There was blinding of all therapists who administered the therapy	no	ои	no	ou	ои	NO	Ou	no
7. There was blinding of all assessors who measured at least one key outcome	yes	yes	Ш	ou	ou	yes	ОП	yes
8. Measures of at least one key outcome were obtained from more than $85\%$ of the subjects initially allocated to groups	оп	yes	yes	оп	yes	ОП	yes	yes
9. All subjects for whom outcome measures were available received the treatment or control condition as allocated or, when this was not the case, data for at least one key outcome was analysed by "intention to treat"	ои	yes	оп	оц	оц	yes	yes	yes
10. The results of between-group statistical comparisons are reported for at least one key outcome	yes	yes	yes	yes	по	yes	yes	yes
11. The study provides both point measures and measures of variability for at least one key outcome	yes	yes	yes	оп	yes	yes	yes	yes
Score 0/10	ŝ	8	4	7	4	7	7	8
*yes=1 point; no=0; Criterion 1 was not used to calculate the PEDro score.	lculate the PED	ro score.						

Gladwell et al.<sup>13</sup> compared the Pilates method with normal activities (n=34). The exercises were basic Pilates method-based floor exercises that involved strengthening the abdominal muscles, gluteus muscles and spinal erectors and progressed to limb exercises, with a total of six sessions lasting one hour over six weeks. The other studies<sup>14,27,30</sup> have been described above.

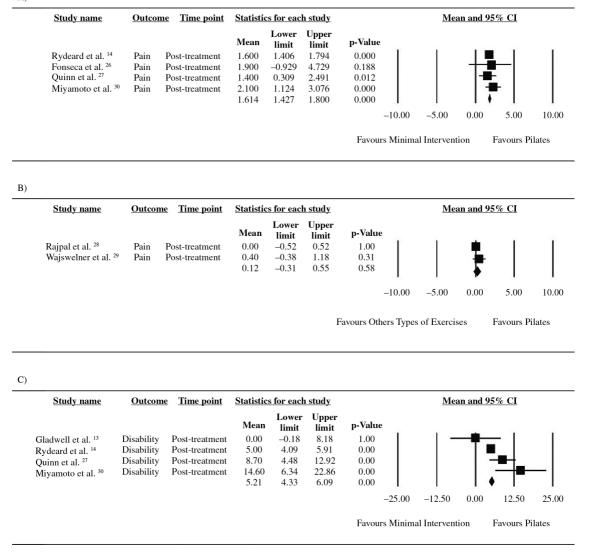
Figure 2C shows the results of this comparison for the disability outcome, using a random-effects model. A statistically significant improvement in

A)

disability favoring the Pilates method was observed at short term (difference between means=5.2 points, 95% CI 4.3 to 6.1), with high level of heterogeneity ( $1^2$ =68%,  $\chi^2$ =9.35, df=3, P=0.02).

#### Discussion

This systematic review shows evidence that Pilates method-based exercises are more effective than no treatment or minimal intervention in the treatment of chronic nonspecific low back pain for the pain



**Figure 2.** Forest plot of the results of the meta-analysis. A) Comparison of the Pilates method with minimal intervention for pain. The values shown are the average effects (difference between means) and the 95% confidence interval. The average effect was calculated using a fixed-effects model for short-term pain relief. B) Comparison of the Pilates method with other types of exercises for pain. The values shown are the average effects (difference between means) and the 95% confidence interval. The average effect was calculated using a fixed-effects model for short-term pain relief. C) Comparison of the Pilates method with minimal intervention for disability. The values shown are the average effects (difference between means) and the 95% confidence interval. The average effect was calculated using a fixed-effects model for short-term pain relief. C) Comparison of the Pilates method with minimal intervention for disability. The values shown are the average effects (difference between means) and the 95% confidence interval. The average effect was calculated using a fixed-effects model for short-term pain relief. C) Comparison of the Pilates method with minimal intervention for disability. The values shown are the average effects (difference between means) and the 95% confidence interval. The average effect was calculated using a random-effects model for short-term changes in disability.

outcome. However, the difference between the means of this comparison was 1.6 points (on a scale of 0 to 10 points). This value does not represent a clinically significant improvement, as a minimum difference of 2 points is required<sup>32</sup>. The results show that Pilates method-based exercises and other types of exercises are equally effective for pain relief. These results are similar to other reviews of specific exercises<sup>10,33</sup> that compare them with other types of exercises or other conservative interventions. For the disability outcome, greater improvement was observed for the Pilates method-based exercises compared with minimal intervention at short term. However, the difference between the means in this comparison was approximately 5 points (on a scale of 0 to 100 points), which does not represent a clinically significant improvement. A minimum difference of 10 points is required for there to be a clinical improvement<sup>32</sup>. There are few studies<sup>14,30</sup> of high methodological quality with representative samples regarding the effectiveness of the Pilates method to treat chronic nonspecific low back pain; for this reason, the results should be interpreted with caution.

This systematic review only included studies published in full article form in journals indexed in databases that undergo rigorous peer review, which is usually performed by experts in the field. One study was excluded from the meta-analysis because it compared three different regimens of the Pilates method<sup>25</sup> and had no comparison group for analysis. Other systematic reviews with meta-analysis included three<sup>17</sup> and four<sup>16</sup> master dissertations and/or doctoral theses, studies that are difficult for the public to access and can present results without clinical relevance. Studies that have not been subjected to peer review may have a higher risk of bias or negative results and are not recommended for inclusion in systematic reviews<sup>34</sup>.

The present study's results are similar to those of another systematic review with meta-analysis<sup>16</sup> on the same topic because the results show a significant difference in pain when the Pilates method is compared with minimal intervention and no statistically significant difference between groups for the same outcome when the Pilates method is compared with other types of exercises. However, the results do not corroborate the disability outcome, for which the present study shows a statistically significant difference in the short term. The difference between the results can be explained by the fact that the abovementioned systematic review<sup>16</sup> included four studies that were not published in journals indexed in databases<sup>31,35-37</sup>. In addition, three studies that were included<sup>12,36,37</sup> had low methodological quality (three to four points on a nine-point scale, the Delphi List), and the effect of the methodological quality of those studies on the meta-analyses was not assessed.

The results of the present study are in opposition to those obtained in another systematic review with meta-analysis<sup>17</sup>, as our results showed a significant difference for pain and disability when the Pilates method and minimal intervention were compared. However, the results corroborate the comparison between the Pilates method and other types of exercises. In the other systematic review<sup>17</sup>, the authors included three studies that had not been published as articles in journals indexed in databases<sup>31,35,36</sup>. In their systematic review<sup>17</sup>, the authors evaluated the risk of bias according to the scale proposed by the Cochrane Back Review Group; however, they did not describe having performed a sensitivity analysis that considered the methodological quality of studies, although they did assess the risk of bias.

The results obtained in this review are related to effects that were analyzed after intervention because there are insufficient studies to conduct a meta-analysis comparing the effects of the Pilates method over time. Only one study30 has evaluated the Pilates method by comparing it with minimal intervention in the medium term, and it did not find a significant difference between the groups. One study<sup>29</sup> evaluated the Pilates method by comparing it to other types of exercises in the medium term, but it also failed to find a difference between the groups. The Pilates method has been used in rehabilitation programs, primarily to treat patients with chronic nonspecific low back pain. However, the effects of the Pilates method are only proven for patients with chronic nonspecific low back pain in the short term (immediately after intervention), and it is still not possible to make inferences regarding the effects of treatment over time. For this reason, more studies evaluating the effectiveness of this method for this population should be conducted. Furthermore, there are no studies that establish the standardized treatment duration, weekly frequency, intensity and type of exercises appropriate for patients with chronic nonspecific low back pain. The analyzed studies show some differences in terms of the choice of exercises because some used Pilates method-based floor exercises<sup>13,25-28,30</sup> and others also used exercise equipment<sup>14,29</sup>. All of the studies opted to begin sessions with basic exercises, but the duration and frequency of sessions differed among the studies.

Studies of the Pilates method should better describe the goals of the exercises used in treatment. Few studies have explored the effect of springs and the variations in Pilates exercises that are methodologically different<sup>34,38</sup>. Perhaps for this reason, it is difficult to standardize an effective exercise program for rehabilitating specific groups of patients. Because there are few studies with a low risk of bias published on this topic, it is suggested that more studies on the effects of chronic nonspecific low back pain and that follow methodological quality criteria be undertaken and published.

Although it does not represent a direct limitation of this review, only a small number of studies were included because there are few indexed articles on the topic, and some studies were excluded because they offered comparisons that differed from those analyzed in the study. This small number of published articles reflected directly on the meta-analysis, as the analyses were performed using only a few articles. However, many studies are currently being conducted in different countries<sup>39,40</sup>, so this limitation should be overcome within a few years.

#### Conclusions

The results of this systematic review suggest that Pilates method exercises are more effective than minimal intervention in improving pain and disability in the short term. However, they are not more effective than other types of exercise in reducing pain. Pilates method exercises can therefore be recommended for the improvement of pain and disability, but no definitive conclusion can be made regarding the analyzed outcomes (pain and disability) in the medium term. Further studies with a low risk of bias and larger samples should be published to obtain greater statistical power in the analyses.

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Search Strategy for EMBASE (OVID) Search Strategy for MEDLINE (OVID) 1. clinical article/ 1. randomized controlled trial, pt. 2. exp clinical study/ 2. controlled clinical trial.pt. 3. randomized.ab. 3. clinical trial/ 4. placebo.ab,ti. 4. controlled study/ 5. drug therapy.fs. 5. randomized controlled trial/ 6. randomly.ab,ti. 6. major clinical study/ 7. trial.ab.ti. 7. double blind procedure/ 8. groups.ab,ti. 8. multicenter study/ 9. or/1-8 9. single blind procedure/ 10. (animals not (humans and animals)).sh. 10. phase 3 clinical trial/ 11.9 not 10 11. phase 4 clinical trial/ 12. dorsalgia.ti,ab. 12. crossover procedure/ 13. exp Back Pain/ 13. placebo/ 14. or/1-13 14. backache.ti.ab. 15. exp Low Back Pain/ 15. allocate\$.mp. 16. assign\$.mp. 16. (lumbar adj pain).ti,ab. 17. blind\$.mp. 17. coccyx.ti.ab. 18. (clinic\$ adj25 (study or trial)).mp. 18. coccydynia.ti,ab. 19. sciatica.ti.ab. 19. compar\$.mp. 20. sciatica/ 20. control\$.mp. 21. cross?over.mp. 21. spondylosis.ti,ab. 22. lumbago.ti,ab. 22. factorial\$.mp. 23. or/12-22 23. follow?up.mp. 24. placebo\$.mp. 24. pilates.mp 25. prospectiv\$.mp. 25. exercise therapy.mp 26. random\$.mp. 26. motor control exercise.mp 27. ((singl\$ or doubl\$ ou trebl\$ or tripl\$) adj25 27. pilates based exercises.mp (blind\$ or mask\$)).mp. 28. exercise movement technics.mp 28. trial.mp. 29. exercises movement techniques.mp 29. (versus or vs).mp. 30. or/24-29 30. or/15-29

31. 11 and 23 and 30

31. 14 and 30	S5. "triple-blind"
32. human/	S6. (MH "Placebo Effect")
33. nonhuman/	S7. (MH "Placebos")
34. exp ANIMAL/	S8. "placebo*"
35. animal experiment/	S9. "random*"
36. 33 or 34 or 35	S10. (MH "Random Sample+")
37. 32 not 36	S11. (MH "Study Design+")
38. 31 not 36	S12. "latin square"
39. 37 and 38	S13. (MH "Comparative Studies")
40. 38 or 39	S14. (MH "Evaluation Research+")
41. dorsalgia.mp.	S15. (MH "Prospective Studies+")
42. back pain.mp.	S16. "follow-up stud*"
43. exp LOW BACK PAIN/	S17. "control*"
44. exp BACKACHE/	S18. prospectiv*
45. (lumbar adj pain).mp.	S19. "volunteer*"
46. coccyx.mp.	S20. S19 or S18 or S17 or S16 or S15 or S14 or S13
47. coccydynia.mp.	or S12 or S11 or S10 or S9 or S8 or S7 or S6 or S5 or S4 or S3 or S2 or S1
48. sciatica.mp.	S21. (MH "Animals+")
49. exp ISCHIALGIA/	S22. S20 not S21
50. spondylosis.mp.	S23. Dorsalgia
51. lumbago.mp.	S24. (MH "Back Pain+")
52. or/41-50	S25. (MH "Low Back Pain")
53. pilates	S26. "backache"
54. exercise therapy	S27. "lumbar pain"
55. motor control exercises	S28. (MH "Coccyx")
56. pilates based exercises	S29. (MH "Sciatica")
57. exercise movement technics	S30. "coccyx"
58. exercise movement techniques	S31. "sciatica"
59. or/53-58	S32. Coccydynia
60. 40 and 52 and 59	S33. (MH "Lumbar Vertebrae")
Search Strategy for CINAHL (EBSCO) S1. (MH " Clinical Trials+")	S34. (MH "Thoracic Vertebrae")
	S35. (MH "Spondylolisthesis")
32 "randomi?ed controlled trial*"	555. (Will Spondytonsulesis )
52. "randomi?ed controlled trial*"	S36. (MH "Spondylolysis")

- S38. S37 or S36 or S35 or S34 or S33 or S32 or S31 or S30 or S29 or S28 or S27 or S26 or S25 or S24 or S23
- S39. pilates
- S40. exercise therapy
- S41. motor control exercise
- S42. Pilates based exercises
- S43. (MH "Therapeutic Exercise+")
- S44. S43 or S42 or S41 or S40 or S39
- S45. S22 and S38 and S44

Search Strategy for CENTRAL – ONLINE (Cochrane Library)

- 1. MeSH descriptor Back explode all trees in MeSH products
- 2. MeSH descriptor Buttocks, this term only in MeSH products
- 3. MeSH descriptor Leg, this term only in MeSH products
- 4. MeSH descriptor Back Pain explode tree 1 in MeSH products

- 5. MeSH descriptor Back Injuries explode tree 1 in MeSH products
- 6. MeSH descriptor Low Back Pain, this tterm only in MeSH products
- 7. (low next back next pain) in All Fields, from 1800 to 2011 in all products
- 8. Ibp in All Fields, form 1800 to 2011 in all products
- 9. MeSH descriptor Sciatica, this term only in MeSH products
- 10. (#1 or #2 or #3 or #4 or #5 or #6 or #7 or #8 or #9)
- Search Strategy for PEDro
- 1. Low back pain
- 2. Pilates
- 3. Clinical trial
- 4.1 and 2 and 3
- Search Strategy for SciELO and LILACS
  - 1. Pilates
  - 2. Low back pain
  - 3.1 and 2