# **SCIENTIFIC** REPORTS

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## OPEN Dose-response-relationship of stabilisation exercises in patients with chronic non-specific low back pain: a systematic review with meta-regression

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Stabilization exercise (SE) is evident for the management of chronic non-specific low back pain (LBP). The optimal dose-response-relationship for the utmost treatment success is, thus, still unknown. The purpose is to systematically review the dose-response-relationship of stabilisation exercises on pain and disability in patients with chronic non-specific LBP. A systematic review with meta-regression was conducted (Pubmed, Web of Knowledge, Cochrane). Eligibility criteria were RCTs on patients with chronic non-specific LBP, written in English/German and adopting a longitudinal core-specific/ stabilising/motor control exercise intervention with at least one outcome for pain intensity and/or disability. Meta-regressions (dependent variable = effect sizes (Cohens d) of the interventions (for pain and for disability), independent variable = training characteristics (duration, frequency, time per session)), and controlled for (low) study quality (PEDro) and (low) sample sizes (n) were conducted to reveal the optimal dose required for therapy success. From the 3,415 studies initially selected, 50 studies (n = 2,786 LBP patients) were included. N = 1,239 patients received SE. Training duration was 7.0 ± 3.3 weeks, training frequency was 3.1 ± 1.8 sessions per week with a mean training time of 44.6 ± 18.0 min per session. The meta-regressions' mean effect size was d = 1.80 (pain) and d = 1.70 (disability). Total R<sup>2</sup> was 0.445 and 0.17. Moderate quality evidence (R<sup>2</sup> = 0.231) revealed that a training duration of 20 to 30 min elicited the largest effect (both in pain and disability, logarithmic association). Low quality evidence ( $R^2 = 0.125$ ) revealed that training 3 to 5 times per week led to the largest effect of SE in patients with chronic non-specific LBP (inverted U-shaped association). In patients with nonspecific chronic LBP, stabilization exercise with a training frequency of 3 to 5 times per week (Grade C) and a training time of 20 to 30 min per session (Grade A) elicited the largest effect on pain and disability.

Exercise is evident for the management of chronic, non-specific low back pain in therapy and rehabilitation<sup>1-4</sup>. In general, strength/resistance and coordination/stabilisation exercise programmes appear to be superior to other interventions in the treatment of chronic low back pain<sup>5</sup>. Specifically, the effects of motor control exercise therapies on the reduction of pain and disability, as well as on improvements in functional performance, are highlighted in numerous meta-analyses on chronic, non-specific low back pain, as an acute, long term<sup>2</sup>, and sustainable treatment<sup>6</sup>. These types of sensorimotor/stabilisation training are the most established therapy forms in low back pain treatment which aim to improve neuromuscular deficits<sup>2,5</sup>. The use of the following interventions indicate the sensorimotor training principles in the context of chronic, low back pain treatment: motor control, sensorimotor, perturbation, neuromuscular, core stability, stabilisation, Pilates-based stabilisation and instability training. The superordinate principle, musculoskeletal control by afferent sensory/proprioceptive

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Criterion	Inclusion	Exclusion
Study design	Randomised controlled	Case studies, case-control, controlled, cohort studies, reviews (e.g. with meta-analysis), protocols, non-controlled intervention studies
	Adults	
Population	Low back pain patients	< 18 years of age
	Pain duration: Sub-acute, chronic, chronic-recurrent	
Intervention	Motor control exercise Core-specific sensorimotor/neuromuscular/sensorimotor/perturbation/core stability/stabiliz(s)ation/stabiliz(s)ation exercises/training	Static (non-dynamic) (motor control) exercises
	Duration of at least 2 weeks	
	Active (any type of exercise, stretching, general strengthening)	
Control/Comparator	Passive comparators (e.g. manual therapy)	
Control/Comparator	Advice to stay active, Usual care	
	Real control (inactive, waiting control)	
Outcome	At least one measure of pain (e.g. VAS, NRS, Korff) and/or disability (e.g. ODI, RMDQ, KORFF)	
Outcome	Outcome assessment at baseline and at least once at 2 week to 24 week post- intervention-initiation	
	Publication or e-pub before 30th March 2020	
Other	Language: German & English	
	Full-text availability	

Table 1. Inclusion and exclusion criteria for both the studies and participants.

input, central nervous system integration of the afferences and optimal stabilisation to ensure functional dynamic joint stability during perturbative situations, are key components of all the above mentioned training forms<sup>7</sup>. The meta-analyses on the effects of these training forms<sup>2–4,8,9</sup> have not pointed out training characteristics (period, duration, frequency, intensity, etc.) for the likely largest effect. The optimal dose for the maximal treatment success-response relationship is, thus, still unknown<sup>1,10</sup>.

It is evident that the success of exercise interventions in the therapy of musculoskeletal disease (including non-specific low back pain) is dependent on the high adherence of the patients to their therapy plan. Regarding the therapy of chronic, non-specific low back pain, the dose-response relationship between stabilisation exercise interventions and pain reduction is of great interest to policy makers, clinicians and individuals. van Tulder et al.<sup>4</sup> reported in their systematic review that a high training dosage ( $\geq 20$  h) is more effective in exercise interventions to improve pain and function in chronic, non-specific low back pain patients. More information on the period, duration, frequency and intensity were not presented. Saragiotto et al.<sup>2</sup> reported a wide range in the duration of the applied motor control intervention programmes in the studies included in their metaanalysis of 20 days to 12 weeks. The number of treatment sessions per week ranged from one to five sessions. Consequently, as a result of, inter alia, this variance in training scheduling, a large heterogenity was found in the meta-analyses highlighted above. Decreasing this heterogeneity would, on the one hand, increase the level of evidence of the stabilisation exercises' effects on low back pain patients. On the other hand, with a much higher impact on clinical and scientific practice, the determination of an optimal dose-response relationship with the thereof derived recommendations on how an intervention needs to be structured in terms of training type, duration, frequency and intensity, is of great relevance. As an impact of a high risk of bias<sup>11</sup> and a low sample size<sup>12</sup> of the studies included into meta-analyses is known, these potential confounders should be considered in dose-response-analyses, likewise.

The purpose of this systematic review with meta-regressions was to (1) delineate the dose-response-relationship of stabilisation exercises and (2) derive recommendations for the stabilisation exercises' training specifics that could maximise the reduction of pain and disability in chronic, non-specific low back pain patients.

#### Methods

The presented systematic review with meta-regression was conducted in accordance with the recommendations of the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA)<sup>13</sup>.

**Literature research.** The literature research was performed using the digital peer review-based databases PubMed (Medline), Web of Knowledge and the Cochrane Library. The following Boolean search syntax was applied (example for the PubMed-search): (stabili\* OR sensorimotor OR "motor control" OR neuromuscular OR perturbation) AND (exercise OR training OR therapy OR intervention OR treatment) AND ("low back pain" OR lumbalgia OR "lower back pain" OR dorsalgia OR backache OR lumbago OR LBP OR "back pain").

Two reviewers (JM & DN) independently conducted the literature research. Consequently, the identified studies were screened for eligibility, using firstly the titles and secondly the abstracts. Afterwards, the remaining full texts were assessed for eligibility by applying the inclusion and exclusion criteria (Table 1). A consensus was used to address any disparities; a third reviewer (N.N.) was planned to be asked, if necessary, to address any disparities. After study retrieval, additional studies were identified by manually searching through the reference

list (cross-referencing) of the selected articles. The search was limited to full-text availability, publication up to the 30th of March 2020 and in the languages of English or German (Table 1).<sup>6</sup>

**Inclusion and exclusion criteria.** The inclusion and exclusion criteria were defined with respect to population, intervention, control/comparator and outcome (PICO). The detailed criteria for both the participants and studies are displayed in Table 1.

**Data extraction.** The common effect estimators for pain intensity and disability were retrieved from each study. The intervention group baseline-to-post effects sizes (Cohens d) were calculated as the change in mean values from baseline to post intervention assessment divided by the baseline standard deviation values for the respective scale. All data of interest were retrieved from the individual study data; for this purpose, a data extraction form designed for this review was used. Data on training dose and frequency were retrieved according to the TIDieR checklist. One researcher recorded all the pertinent data from the included articles and the other author independently reviewed the extracted data for its relevance, accuracy and comprehensiveness. A consensus was used to address any disparities; a third reviewer (N.N.) was asked, if necessary, to address any disparities. Authors of those studies included in this review who had not reported sufficient details in the published manuscript, were personally addressed by e-mail requesting the provision of further data. The effect estimators for pain intensity and disability were calculated using either the visual analogue scale (VAS), the numeric rating scale (NRS) or the sum score, inherent of the scale/assessment tool (0-10, 0-24 or 0-100), as the calculation of the standard mean differences is scale independent. For such data, only the direction (lower values mean less pain, less disability) was normalised. For scale-dependent calculations (inverse weighting, calculated as sample size divided by the squared standard deviation of the baseline-to-post difference), z-transformed (0-10) variables were used. Missing standard deviations for the differences were imputed according to the procedure described by Follmann et al.<sup>14</sup>.

**Study quality assessment.** The Physiotherapy Evidence Database (PEDro; 11 criteria) scale was used to assess the methodological quality of all trials included. The PEDro scale is a valid and reliable tool to rate the internal study validity and methodological quality of controlled studies<sup>15</sup>. If available, the validated rating scores of the articles were taken directly from the PEDro database (website; 35 out of 46 articles). If not, both authors evaluated the articles, each criterion was rated as 1 (definitely yes) or 0 (unclear or no); potential disagreements were discussed between the two authors and resolved. Overall, the scale ranges from 0 (high risk of bias) to 10 (low risk of bias) with a sum score of  $\geq 6$  representing a cut-off score for studies with a sufficient study quality. As study quality was considered as a potential explanator of the effect size homogeneity, all studies, irrespective of the quality, were analysed.

**Risk of bias within the studies.** The two review authors (JM and DN) independently rated the risk of bias of the outcomes pain and disability in the included studies by using the Cochrane Collaboration's tool Risk of Bias tool 2<sup>16,17</sup>. Studies' outcomes were graded for risk of bias in each of the following domains: sequence generation, allocation concealment, blinding (participants, personnel, and outcome assessment), incomplete outcome data, selective outcome reporting and other sources of bias. For the outcomes, each item was rated as "high risk", "low risk" or "unclear risk" of bias. Again, any disagreements were discussed between the raters. If a decision could not be reached after discussion, a third reviewer (N.N.), was included to resolve any conflicts. As the risk of bias was (indirectly, via the PEDro sum score) considered as a potential explanator of the effect size homogeneity, all studies, irrespective of the risk of bias, were analysed in the meta-regressions.

**Risk of bias across the studies.** The calculation of the risk of publication bias across all the studies was indicated by using funnel plots/graphs<sup>18</sup>. The Review Manager 5.3 (RevMan, Version 5.3, Copenhagen: The Nordic Cochrane Centre, The Cochrane Collaboration, 2014) was used for funnel plotting.

**Data processing and statistical analysis.** Data was initially plotted using scatterplot diagrams. The type of association between each independent and dependent variable was visually determined. In case of a linear association, data were processed as real values, thus, if a curve-linear association was determined, data were re-calculated using logarithmic transformations (log-association) and, respectively, Taylor-series (U-shaped-associations) to provide linearity for the regression calculation.

Sensitivity meta-regressions for dose-response analyses and the impact of study quality were conducted as described in Niederer & Mueller (2020)<sup>6</sup>. A syntax for SPSS (IBM SPSS 23; IBM, USA) was used (David B. Wilson; Meta-Analysis Modified Weighted Multiple Regression; MATRIX procedure Version 2005.05.23). Inverse variance weighted regression models with random intercepts (random effect model, fixed slopes model) with the dependent variables of pain intensity and disability effects (simple pre-post Cohen's ds) and the independent variables: intervention duration [weeks, U-shaped], intervention frequency [number of trainings/week, U-shaped], intervention duration [minutes, logarithmised], intervention total dose [minutes] were applied. The sample size (SE group) and the study quality PEDro sum score [points, linear] were considered as co-factors. Homogeneity analysis (Q- and p-values) and meta-regression partial coefficients B (95% confidence intervals and p-values) were calculated. All statistical analyses were tested against a 5% alpha-error probability level.

**Effect estimators' level of evidence.** The quality of the evidence revealed by the meta-analyses was graded using the tool established by the GRADE working group<sup>19</sup>. Quality evidence was categorised as "very



Figure 1. Research, selection and synthesis of included studies. n, number; Eng, English; Ger, German; WoK, web of knowledge.

low" (The estimate of effect is very uncertain), "low" (further research is likely to change the estimate), "moderate" (further research may change the estimate) or "high" (further research is very unlikely to change the estimate of effect) (plus interim values). The grading starts with the type of evidence (RCT = high, Observational = low, all other study types = very low) and is decreased or increased based on study limitations, inconsistencies, uncertainty about directness, imprecise data, reporting bias (decreasing items), or strong associations, dose-response findings, and confounder plausibility (increasing items)<sup>19</sup>.

Recommendations were derived using a clinical guideline developing tool<sup>20</sup>. Overall, four key factors were applied to determine the strength of the recommendations: Balance between desirable and undesirable effects (larger differences between desirable undesirable effects lead to stronger recommendations)—Quality of the available evidence—Values and preferences (higher variations lead to weaker recommendations)—costs (higher costs lead to weaker recommendations. Details that are more comprehensive can be found in<sup>21</sup>.

## Results

**Study selection.** The database search was completed in 03/2020. Figure 1 displays the research procedure and the flow of the study selection and inclusion.

**Study characteristics and individual studies' results.** Fifty (50) studies were included in the qualitative and in the quantitative analyses. Study characteristics and the main results are displayed in Table 2. For each of the studies included, methodological aspects, participants' characteristics and key results are presented. Overall, 2,786 participants, thereof n = 1,239 stabilisation exercise participants, were included in the analysis.

All included studies adopted a randomised controlled design (RCT). The main inclusion criterion was (chronic) non-specific low back pain  $\geq$  4 weeks<sup>22</sup>,  $\geq$  6 weeks<sup>23</sup>,  $\geq$  7 weeks<sup>24</sup>,  $\geq$  8 weeks<sup>25–27</sup>,  $\geq$  12 weeks<sup>28–55</sup>,  $\geq$  24 weeks<sup>56–58</sup> and  $\geq$  2 year history<sup>59</sup>, whilst in 11<sup>60–70</sup> studies this information was not presented. The baseline pain, effect sizes (Cohen's d, stabilisation exercise group only) for pain and disability are presented in Table 3.

**Study quality and risk of bias within studies.** Both the study quality and risk of bias ratings are presented in Table 2. The overall study quality ranged from 3/10 to 9/10 points, with a mean of  $5.7 \pm 1.4$  points on the Pedro scale.

**Individual studies' training characteristics.** Table 4 summarises the individual studies' training characteristics. All interventions and the comparators are described. The stabilisation exercises are called core stability exercise<sup>25,27,30-32,47,51,54,61-63,69</sup>, motor control exercise<sup>24,35,38,44,45,48,50</sup>, stabilisation<sup>23,26,28,34,41,52,55,60</sup>, lumbar stabilisation exercise<sup>39,46,56,64,67</sup>, spinal stabilisation<sup>33,37,68</sup>, sensorimotor training<sup>66,71</sup>, trunk stability exercise<sup>49,58</sup>, Swiss ball stabilisation<sup>43,65,70</sup>, perturbation training<sup>29</sup>, sling training<sup>53</sup>, McGill stabilisation exercise<sup>40,57</sup>, segmental stabilisation exercise<sup>36</sup>, neuromuscular exercise<sup>22</sup>, multifidus muscle retraining<sup>59</sup> and Pilates-based exercise<sup>42</sup>. The intervention period ranged between 2<sup>69</sup> and 24<sup>22</sup> weeks with a mean of 7.0 ± 3.3 weeks. Training frequency ranged from 1<sup>29</sup> to 12<sup>53</sup> times per week with a mean of  $3.1 \pm 1.8$  times; 3 studies<sup>24,55,63</sup> did not report on this information. Mean training time per session was  $44.6 \pm 18.0$  min with a range from  $15^{24}$  to 90 minutes<sup>29,33</sup> (9 studies<sup>35,47,49,54,62,63,65,67,68</sup> did not report on this aspect). The number of exercises practised per session varied between  $2^{35,47,49,54,62,63,65,67,68}$  to  $18^{29}$  exercises with a mean of  $7.2 \pm 3.9$  exercises; 13 studies<sup>30,32, 35,37,40,44,48,50,52,53,56,58</sup> did not report this information.

The qualitative analysis of the training volume revealed a range of  $1^{30,32,35,37,40,44,48,50,52,53,56,58,70}$  to  $10^{24,44,46,59,60}$  sets per exercise practiced with a mean of  $3.2 \pm 2.4$  sets, while  $28^{22,25,28,30-35,38-41,43,45,49-51,53,54,56,58,63,67-71}$  studies did not report any details on this aspect. In addition to this, only  $23^{22,25,28,30-35,38-41,43,45,49-51,53,54,56,58,63,67-71}$  studies reported on the number of repetitions per set per exercise, with a range of  $6^{23,24,26,27,36,38,42,44,46-48,50-52,55,57,59-61,64-66,69}$  to  $30^{66}$  repetitions (mean:  $13.6 \pm 5.6$  repetitions per set per exercise). In addition, only 12 studies<sup>29,30,42,46,48,57,59-62,64,65</sup> to  $300^{65}$  s (mean:  $106.3 \pm 86.5$  s).

**Meta-regression analysis.** The results of the meta-regressions are highlighted in Table 5. The total variance explanation was 44% for pain and 15% for disability. When all the other predictors were partialized, moderate quality evidence revealed that a training duration of 20 to 30 min elicits the largest impact on the effect sizes (both in pain and disability) of stabilisation exercise training in low back pain patients. Quality of evidence was downgraded due to risk of bias (-1), downgraded due to imprecise data (wide confidence intervals, -1), downgraded (-1) due to (some) uncertainty about directness, and upgraded due to dose-response-relationship (+1), upgraded due to: confounders were considered (+1).

More detailed information on the meta-regressions are depicted in Fig. 2. The training period showed no systematic impact on the effect size for pain intensity (Fig. 2A). Training frequency showed an inverted U-shaped association with the effect size (13% variance explanation) (Fig. 2B), training duration showed a logarithmic association with the pain effect size (23% variance explanation; Fig. 2C). Low quality evidence suggested that training 3 to 5 times per week leads to the largest effect of stabilisation exercise in chronic, non-specific low back pain patients. Quality of evidence was downgraded due to risk of bias (-1), downgraded due to imprecise data (wide confidence intervals, -1), downgraded (-1) due to (some) uncertainty about directness, and upgraded due to dose-response-relationship (+1).

**Risk of bias across studies.** The risk of bias across studies (publication bias) is, by means of a funnel plot, highlighted in Fig. 3. It reveals an unclear, but rather low, risk of publication bias.

#### Discussion

This systematic review with meta-regression examined the dose-response-relationship of stabilisation exercise interventions in chronic, non-specific low back pain patients and, thus, derived recommendations for the stabilisation exercises' training characteristics in this special cohort.

**Summary of main results.** The main findings of the presented meta-regression are that: (1) moderate quality evidence indicates that a training duration of 20 to 30 min elicits the largest impact on the effect sizes on both pain and disability of core-specific stabilisation interventions in non-specific chronic low back pain patients, (2) low quality evidence advocates that training 3 to 5 times per week leads to the largest effect of core-specific stabilisation exercise in chronic, non-specific low back pain patients with an inverted U-shaped association with the effect size and (3) no systematic impact of the training period (duration of intervention in weeks) on the effect size for pain intensity was found.

**Comparison with other evidence.** Saragiotto et al.<sup>2</sup> reported a wide range of 20 days to 12 weeks in the period of the applied motor control intervention programmes in their meta-analysis. The number of treatment

	Peo	lro											Risk of bias	assessment					
Item/Study	1	2	3	4	5	6	7	8	9	10	11	Sum PEDro	Random sequence generation	Allocation concealment	Blinding of participants and personnel*	Blinding of outcome assessment*	Incomplete outcome data	Selective reporting	Other bias
Alp, 2014	1	1	0	1	0	0	1	1	0	1	1	6	Low	High	High	Low	Unknown	Low	Low
Alrwaily, 2019	1	1	1	1	0	0	0	1	0	1	1	6	Low	Low	High	High	Unknown	Low	Unknown
Andru- saitis, 2011	0	1	1	1	0	0	1	1	0	1	1	7	Low	Low	High	Low	Unknown	Low	High
Arampatzis, 2017	1	1	0	1	0	0	0	1	0	1	1	5	Low	High	High	High	Unknown	Low	Low
Areeudom- wong, 2019	1	1	1	1	0	0	1	1	1	1	1	8	Low	Low	High	Low	Low	Low	Unknown
Bae, 2018	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Low	Low
Bauer, 2019	1	1	1	1	0	0	1	0	1	1	1	7	Low	Low	High	Low	High	Low	Low
Brooks 2012	0	1	0	1	1	0	1	1	1	1	1	8	Unknown	High	High	Low	Low	Unknown	Unknown
Chung, 2018	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Unknown	Unknown
Critchley, 2007	1	1	1	1	0	0	1	0	1	1	1	7	Low	Low	High	Low	High	Low	Unknown
Da Luz, 2019	1	1	1	1	0	0	1	1	1	1	1	8	Low	Low	High	Low	Low	Low	Low
Demirel, 2019	0	1	1	1	0	0	0	1	0	1	1	6	Low	Low	High	High	Unknown	Low	Unknown
Ferreira 2007	1	1	1	1	0	0	1	1	1	1	1	8	Low	Low	High	Low	Low	Low	Low
Franca 2012	1	1	1	1	0	0	1	1	1	1	1	8	Unknown	Low	High	Low	Low	Low	Low
Ghorban- pour, 2018	0	1	0	1	0	0	0	1	0	1	1	5	Low	High	High	High	Unknown	Low	Low
Hosseinifar, 2013	1	1	0	1	0	0	1	0	0	1	1	5	Low	High	High	Low	High	Low	Unknown
Hwang, 2013	1	1	0	1	0	0	0	0	0	1	1	4	Unknown	High	High	High	High	Low	High
Ibrahim, 2018	1	1	1	1	0	0	1	1	1	1	1	8	Low	Low	High	Low	Low	Low	Low
Inani, 2013	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Unknown	High
Khodadad, 2019	1	1	1	1	0	0	1	1	1	1	1	8	Low	Low	High	Low	Low	Low	Unknown
Kim, 2018	1	1	0	1	0	0	0	0	1	1	1	5	Low	High	High	High	High	High	Unknown
Kim, 2019	1	1	0	1	0	0	0	0	1	1	1	5	Low	High	High	High	High	Low	Unknown
Ko, 2018	0	1	0	1	0	0	0	0	0	1	1	4	Unknown	High	High	High	High	Low	High
Kofotolis, 2016	1	1	1	1	0	0	0	0	0	1	1	5	Unknown	Low	High	High	High	Low	Low
Lee, 2014	1	1	0	1	0	0	0	1	0	1	1	5	Low	High	High	High	Unknown	Unknown	High
Lee, 2011	0	1	0	1	0	0	0	0	0	1	1	4	Unknown	High	High	High	High	Unknown	High
Letafatkar, 2017	0	1	0	1	0	0	0	0	0	1	1	4	Unknown	High	High	High	High	Low	Unknown
Liu, 2019	1	1	0	1	0	0	0	1	0	1	1	5	Low	High	High	High	Unknown	Low	Unknown
Lomond, 2015	1	1	0	1	0	0	1	1	0	1	0	5	Unknown	High	High	Low	Unknown	Low	Low
Macedo, 2012	1	1	1	1	0	0	1	1	1	1	1	8	Low	Low	High	Low	Low	Low	Low
Marshall, 2013	1	1	1	1	1	0	1	1	1	1	1	9	Unknown	Low	High	Low	Low	Low	Unknown
Miller, 2013	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Low	Unknown
Moon, 2013	1	1	0	1	0	0	1	1	0	1	1	6	Unknown	High	High	Low	Unknown	Low	Low
Noormo- hammad- pour, 2018	1	1	1	1	0	0	1	0	1	1	1	7	Unknown	Low	High	Low	High	Low	Low
Rabin, 2014	1	1	1	1	0	0	0	0	1	1	1	6	Unknown	Low	High	High	High	Low	Unknown
Rasmussen- Barr, 2003	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Low	Unknown
Rasmussen- Barr, 2009	1	1	1	1	0	0	0	1	1	1	1	7	Low	Low	High	High	Low	Low	Low
Rhee, 2012	1	1	1	1	0	0	0	0	0	1	1	5	Low	Low	High	High	High	Low	Unknown
Continued																			

	Pedro					Risk of bias	assessment												
Item/Study	1	2	3	4	5	6	7	8	9	10	11	Sum PEDro	Random sequence generation	Allocation concealment	Blinding of participants and personnel*	Blinding of outcome assessment*	Incomplete outcome data	Selective reporting	Other bias
Salamat, 2017	1	1	0	1	0	0	0	0	0	1	1	4	Unknown	High	High	High	High	Low	Unknown
Seo, 2019	0	1	0	1	0	0	0	0	0	1	1	4	Low	High	High	High	High	Low	Unknown
Shamsi 2017	0	1	0	1	0	0	0	1	0	1	1	5	Low	High	High	High	Unknown	Low	High
Shaugh- nessy, 2004	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Low	Unknown
Soundara- rajan, 2016	0	1	0	1	0	0	0	0	0	1	1	4	Unknown	High	High	High	High	Unknown	Low
Sung, 2013	1	1	1	1	0	0	1	0	0	1	1	6	Unknown	Low	High	Low	High	Low	Unknown
Ulger, 2017	1	1	0	1	0	0	1	0	0	1	1	5	Unknown	High	High	Low	High	Low	Unknown
Unsgaard- Tondel, 2010	1	1	1	1	0	0	0	1	1	1	1	7	Low	Low	High	High	Low	Low	Low
Vikranth, 2015	1	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	Unknown	Unknown
Waseem, 2018	1	1	1	0	0	0	0	1	0	1	0	4	Unknown	Low	High	High	Unknown	High	Unknown
Woo, 2016	0	1	0	1	0	0	0	1	0	1	1	5	Unknown	High	High	High	Unknown	High	High
Young, 2015	1	1	0	1	0	0	0	0	0	0	1	3	Unknown	High	High	High	High	High	High

Table 2. Study quality (Pedro scale) and risk of bias assessment.

sessions per week varied from 1 to 5. This partly covers the results of our presented meta-regressions. Nevertheless, a detailed analysis on the effect of training characteristics on pain reduction is missing in their systematic review<sup>2</sup>. The current evidence only proves the use of general and stabilisation exercise (covering sensorimotor, stabilisation and/or core stability) in the therapy of chronic non-specific low back pain<sup>2</sup>. Regarding the training period/duration (weeks of intervention), our results showed that the duration of intervention (in weeks) presented no systematic impact on the effect size for pain intensity. Taking the current knowledge on the effects and adaptation of sensorimotor training into account, a duration of about six weeks seems to be both feasible and effective. This is in accordance with our quantitative results (mean duration of  $7.0 \pm 3.3$  weeks). However, future research is required to define evidence-based recommendations of this aspect.

Low quality evidence supports an inverted U-shaped association of the training frequency (sessions per week) with the effect size on improvement of pain and disability in chronic, non-specific low back pain patients. The overall relationship between (the amount of) physical activity and low back pain is considered to be U-shaped. This means that both the absence of exercise and extremely high levels of physical activity (elite sports) may lead to an increase in the risk of developing (low) back pain. In contrast, a "normal" (medium) level of physical activity shows the lowest risk and, therefore, appears to be protective<sup>2-4,8,9</sup>. In this context, our findings of adopting a dose of 3 to 5 sessions per week covers this. In addition, moderate quality evidence indicates that a training duration of 20 to 30 min elicits the largest impact on the effect sizes on pain and disability; this may correspond to the patients' essential need of achieving pain reduction with the minimum effort (time). Nevertheless, this is partly in contrast to van Tulder's result<sup>4</sup>. They concluded that exercise interventions with a high dosage (>20 h) have the highest effect. Van Tulder et al.<sup>4</sup> fail to point out how this dosage should be applied (duration, frequency). Supported by our findings, it may be more effective to reach this dosage with a high frequency, short bout type of intervention. One of the main reasons of failed treatment success in exercise therapy is the low adherence rate of the patients to their scheduled therapy<sup>4</sup>. Lack of time and long journey times to the therapy centre are commonly cited barriers to regularly participating in therapy sessions<sup>72</sup>. Therefore, patients and physiotherapists are constantly searching for the effective dose-response-relationship that could be reduced to the minimum required. Based on our results, we can recommend exercising for more than 2 sessions per week with a minimum of 20 to 30 min per session. Nevertheless, there is still a need for future research on the minimal dosage in the context of stabilisation exercise interventions for chronic, non-specific low back pain patients.

**Practical relevance and recommendations.** The training-dose and effect-response relationship between core-specific stabilisation exercise interventions and pain reduction or disability improvement in chronic, non-specific low back pain patients is of great interest to policy makers, health insurers and clinicians, as well as the persons affected. This review proved the (low to moderate) evidence, that a core-specific stabilisation intervention of 3 to 5 times per week, 20 to 30 min per session, has a positive effect on pain reduction and improvement of disability in low back pain patients. Conclusively, we suggest the following graded recommendations:

Grade A recommendation: At the group level, stabilisation exercise is likely to be most effective to treat non-specific low back pain when it is scheduled with a time per session of 20–30 min.

Study informat	ion		Population					Assessments	Outcomes	
First author, year	Citation number	Study design, no of study arms	Main inclusion criterion LBP (time, other)	N (Total, per grop) (SE, C, C2))	Age Mean±SD (years)	Sex (f/m)	Baseline- pain (Scale, mean, SD if not stated otherwise)	Measurement time points total (N: weeks (if not, stated otherwise) after Baseline)	Primary outcome pain, scale, Co-hens d, (M0-M1)	Primary outcome disability name, Cohens d,, (M0-M1)
Alp, 2014	56	RCT, 2 SE Ctrl	CLBP≥24 weeks	48, 24, 24	25–64, 48, 51	48/0	VAS (0–10), 6, range 4–9 6, range 1–10	2: 0; 12	VAS (0-10) SE: 0.8 Imputed from Saragiotto et al. <sup>2</sup>	RMDQ SE: 0.59 Imputed from Saragiotto et al. <sup>2</sup>
Alrwaily, 2019	28	RCT, 2 SE Ctrl	$CLBP \ge 12$ weeks, NPRS $\ge 3$ MODQ score $\ge 20\%$	30 15, 15	38.3±11.3, 33.4±9.0	19/11 11/4 8/7	$4.4 \pm 1.8$ $4.2 \pm 1.9$	2: 0; 6	NPRS (0–10) SE: 1.29	MODQ SE: 1.76
Andrusaitis, 2011	60	RCT, 2 SE Ctrl	nonspecific, CLBP	10, 5, 5	Range: 30–55	10/0 5/0 5/0	VAS (0–10), 4.83, range 4.3–5.5, 5.08, range 0.5–7.7	2: 0; 7	VAS (0–10) SE: 1.60	ODI SE: 1.68
Arampatzis, 2017	29	RCT, 2 SE Ctrl	LBP≥12 weeks	40, 20, 20	$31.9 \pm 6.0, \\ 31.4 \pm 5.5$	N.A	VAS (0-10), 3.96±1.41, 4.22±1.66	2: 0; 13	VAS (0–10) SE: 0.60	N.A
Areeudom- wong, 2019	30	RCT, 3 SE Ctrl 1 Ctrl 2	CLBP≥12 weeks	45 15 15 15	$\begin{array}{c} 24.08 \pm 1.00 \\ 24.00 \pm 8.47 \\ 24.36 \pm 9.97 \end{array}$	34/11 11/4 12/3 11/4	$\begin{array}{c} 4.40 \pm 1.40 \\ 4.13 \pm 0.92 \\ 4.07 \pm 1.28 \end{array}$	3: 0; 4; 12	NRS (0–10) SE: 2.61	Functional disability SE: 1.44
Bae, 2018	31	RCT, 2 SE Ctrl	LBP≥12 weeks	36, 18, 18	$32.7 \pm 6.1,$ $32.4 \pm 11.0$	18/20	VAS (0-10), 2.9±0.8, 3.0±1.3	4: 0; 4; 8; 16	VAS (0–10) SE: 1.0	ODI SE: 0.19
Bauer, 2019	22	RCT, 2 SE Ctrl	LBP≥4 weeks NRS≥3	83 42 41	$45.7 \pm 7.8$ $46.7 \pm 7.7$	83/0	$34.0 \pm 21.0$ $28.0 \pm 21.1$	3: 0; 24; 48	VAS (0–100) SE: 0.42	N.A
Brooks, 2012	32	RCT, 2 SE Ctrl	LBP≥12 weeks	64, 32, 32	$36.2 \pm 8.2,$ $36.3 \pm 6.3$	40/24	VAS (0-10) 3.6±2.1, 4.5±2.5	2: 0; 8	VAS (0–10) SE: 0.58	ODI SE: 1.08
Chung, 2018	61	RCT, 2 SE I SE II	CLBP	27, 14, 13	$32.47 \pm 7.89,$ $34.18 \pm 6.59$	17/10	VAS (0-10) 6.63±1.21, 6.55±1.09	2: 0; 6	VAS (0–10) SE I: 4.35 SE II: 2.95	Korean Ver- sion of ODI SE I: 3.22 SE II: 1.95
Critchley, 2007	33	RCT, 3 SE Ctrl 1 Ctrl 2	CLBP≥12 weeks	212 72 71 69	$44 \pm 13$ $45 \pm 12$ $44 \pm 12$	133/89	NRS (0–100), mean, 95%CI 67, 61–73 60, 54–66 59, 52–65	4: 0; 24; 48; 72	NRS (0–100) SE: 1.08	RMDQ SE: 0.23
da Luz, 2019	62	RCT, 3 SE Ctrl 1 Ctrl 2	CLBP VAS≥4	30 10 10 10	$26.40 \pm 3.41 \\ 25.50 \pm 5.28 \\ 27.10 \pm 4.95$	30/0	$6.4 \pm 0.8$ $6.6 \pm 1.1$ $6.8 \pm 0.4$	3: 0; 4; 24	VAS (0–10) SE: 5.12	ODI SE: 2.02
Demirel, 2019	34	RCT, 2 SE Ctrl	CLBP≥12 weeks	77 37 40	$45.59 \pm 12.32$ $44.25 \pm 8.71$	62/15 29/8 33/7	$2.62 \pm 2.23$ $2.92 \pm 2.65$	2: 0; 6	VAS (0–10) SE: 0.39	ODI SE: 0.75
Ferreira, 2007	35	RCT, 3 SE Ctrl. 1 Ctrl. 2	LBP≥12 weeks	240, 80, 80, 80	$51.9 \pm 15.3,$ $54.8 \pm 15.3,$ $54.0 \pm 14.4$	165/75	VAS (0-10), 6.3±2.0, 6.5±2.1, 6.2±2.0	4: 0; 8, 24; 48	VAS (0–10) 0.92	RMDQ SE: 1.15
Franca, 2012	36	RCT, 2 SE Ctrl	LBP≥12 weeks	30, 15, 15	$42.1 \pm 8.2, \\ 41.5 \pm 4.4$	N.A	VAS (0-10), 5.94±1.56, 6.35±1.51	2: 0; 6	VAS (0–10) SE: 3.77	ODI SE: 3.83
Ghorbanpour, 2018	57	RCT, 2 SE Ctrl	LBP≥24 weeks	30, 15, 15	$23.8 \pm 3.5,$ $20.9 \pm 1.2$	16/14	VAS (0-10), 29.5±4.8, 28.3±6.5	2: 0; 6	VAS (0–100) SE: 0.94	Persian version of the Quebec Low Back Pain Disability Scale Questionnaire SE: 0.33
Hosseinifar, 2013	37	RCT, 2 SE Ctrl	LBP≥12 weeks	30, 15, 15	40.1±10.8, 36.6±8.2	N.A	VAS (0–100), 4.33±1.58, 4.40±1.95	2: 0; 6	VAS (0–100) d=1.77	FRI question- naire d=1.45
Hwang, 2013	71	RCT, 3 SE Ctrl. 1 Ctrl. 2	LBP≥12 weeks	21, 7, 7, 7	45.7±8.5, 44.8±7.9, 45.8±9.2,	10/11	VAS (0-10), N.A., 5.83±0.38, 5.71±0.61	2: 0; 4	VAS (0–10) SE: 3.32	ODI SE: 1.18
Ibrahim, 2018	38	RCT, 3 SE Ctrl 1 Ctrl 2	LBP≥12 weeks	30 10 10 10	$\begin{array}{c} 48.5 \pm 14.9 \\ 50.3 \pm 9.09 \\ 49.9 \pm 8.82 \end{array}$	6/25 3/7 1/9 2/8	$\begin{array}{c} 6.00 \pm 1.41 \\ 6.00 \pm 1.41 \\ 6.80 \pm 1.31 \end{array}$	2: 0; 6	NPRS (0–10) SE: 2.13	ODI SE: 0.97
Continued										

Study informat	ion		Population					Assessments	Outcomes	
First author, year	Citation number	Study design, no of study arms	Main inclusion criterion LBP (time, other)	N (Total, per grop) (SE, C, C2))	Age Mean±SD (years)	Sex (f/m)	Baseline- pain (Scale, mean, SD if not stated otherwise)	Measurement time points total (N: weeks (if not, stated otherwise) after Baseline)	Primary outcome pain, scale, Co-hens d, (M0-M1)	Primary outcome disability name, Cohens d., (M0-M1)
Inani, 2013	63	RCT, 2 SE Ctrl	diagnosed with non-specific LBP	30, 15, 15	$27.8 \pm 7.3,$ $32.9 \pm 64$	10/20	VAS (0–10), 6.3±1.8, 7.0±1.6	2: 0; 12	VAS (0–10) SE: 2.72	Modified ODI SE: 2.28
Khodadad, 2019	39	RCT, 3 SE Ctrl 1 Ctrl 2	LBP≥12 weeks	52 17 17 18	$\begin{array}{c} 42.2 \pm 3.78 \\ 44.3 \pm 1.43 \\ 44.4 \pm 2.17 \end{array}$		$\begin{array}{c} 6.2 \pm 1.48 \\ 5.5 \pm 1.03 \\ 5.6 \pm 1.45 \end{array}$	2: 0; 8	NRS (0–10) SE: 1.89	N.A
Kim, 2018	40	RCT, 2 SE I SE II	LBP > 12 weeks	30 15 15	N.A 22.31±1.6 22.92±1.55	30/0 15/0 15/0	N.A	2: 8	N.A	ODI SE I: 1.47 SE II: 1.64
Kim, 2019	41	RCT, 2 Ctrl SE	LBP≥12 weeks	48 24 24	N.A 26.0±3.82 28.79±9.05	7/15 15/11	NRS (0-10) 4.70±1.04 4.73±0.82	4: 4, 8, 24	NRS (0–10) SE: 3.22	ODI SE: 0.32
Ko, 2018	64	RCT, 3 SE Ctrl. 1 Ctrl. 2	CLBP	29, 10, 10, 9	$\begin{array}{c} 43.1 \pm 3.7, \\ 43.6 \pm 4.5, \\ 41.3 \pm 3.8 \end{array}$	N.A	NRS (0-10), 5.5±1.3, 5.3±1.3, 5.2±2.1	2: 0; 12	NRS (0–10) SE: 1.15	N.A
Kofotolis, 2016	42	RCT, 3 SE Ctrl. 1 Ctrl. 2	CLBP≥12 weeks	101, 28, 37, 36	$42.71 \pm 6.1,$ $41.22 \pm 8.49,$ $39.11 \pm 8.68$	101/0 37 36 28	SF-36 (bodily pain), 36.93±15.52, 38.51±12.62, 39.42±14.49	5: 0; 4; 8; 12; 20	SF-36 pain (0–100) SE: 1.9	RMDQ SE: 0.75
Lee, 2014	65	RCT, 2 SE Ctrl	CLBP	40, 20, 20	$34.20 \pm 0.69, \\ 34.75 \pm 0.85$	N.A	VAS (0-10), 7.85±1.00, 7.95±1.00	3: 0; 2, 4, 6	VAS (0–10) SE: 5.75	N.A
Lee, 2011	25	RCT, 2 SE Ctrl	LBP≥8 weeks	32, 13, 19	26-63, 50.4±9.1, 46.6±9.1	15/17	N.A	2: 0; 4	Million pain interfer- ence visual analogue scale MVAS (0–100 mm; 15 items) SE: 0.78	N.A
Letafatkar, 2017	66	RCT, 2 SE Ctrl	chronic non- specific LBP; scores > 4 in RMDQ	53, 27, 26	N.A., 36.86±7.16, 38.25±6.19	N.A	VAS (0–10), 6.90±1.87, 5.91±1.31	2: 0; 5	VAS (0–10) SE: 2.9 Imputed from graph	RMDQ: SE: 2.3 Imputed from graph
Liu, 2019	43	RCT, 3 Ctrl SE Ctrl	LBP > 12 weeks	43 15 15 13	N.A 58.13±5.38 58.4±5.08 60.67±2.58	35/8 12/3 12/3 11/2	VAS (0-10) 5.67±0.81 5.67±0.72 5.85±0.89	2,:12	VAS (0–19) SE: 1.92	N.A
Lomond, 2015	58	RCT, 2 SE Ctrl	LBP>24 weeks; ODI≥19%	33, 12, 21	$\begin{array}{c} 43.1 \pm 11.9, \\ 41.6 \pm 10.9 \end{array}$	15%male 6%male	NRS (0-10), 2.8±1.6, 3.6±1.6	2: 0; 7	NRS 0–100 SE: 1.1	ODI SE: 0.9
Macedo, 2012	44	RCT, 2 SE Ctrl	CLBP≥12 weeks	158, 76, 82	48.7±13.7, 49.6±16.3	57/19 45/37	NRS (0-10), 6.1±2.1, 6.1±1.9	4: 0; 8, 24; 48	NRS (0–10) SE: 1.05	RMDQ: SE: 0.81
Marshall, 2013	45	RCT,2 SE Ctrl	Ongoing recurrent LBP≥12 weeks	64, 32, 32	$18-50, \\ 36.2 \pm 8.2, \\ 36.2 \pm 6.2$	40/24	VAS (0-10), 3.6±2.1, 4.5±2.5	3: 0; 8; 24	VAS 0–10, SE: 0.9	ODI: SE: 0.93
Miller, 2013	24	RCT, 2 SE Ctrl	LBP≥7 weeks	29, 15, 14	19-87, 54±15, 44±16	14/15	VAS (0-10), 4.1±2.0, 3.0±2.0	2: 0; 6	VAS (0–10) SE: 0.5	N.A
Moon, 2013	46	RCT, 2 SE I SE II	LBP≥12 weeks	21, 11, 10	28.6±4.9, 28.4±5	7/14	VAS (0-100), 34.2±17.1, 33.5±18.4	2: 0; 8	VAS (0–100), SE: 0.78, SE II: 0.93	ODQ, SE: 0.84 SE II: 2.1
Noormoham- madpour, 2018	47	RCT, 2 SE Ctrl	CLBP≥12 weeks	20, 10, 10	$18-55, \\ 43.3 \pm 7.5, \\ 41.0 \pm 6.4$	20/0	VAS (0-100), 38.4±21.7, 36.2±27.2	N.A	VAS (0–100), SE: 1.6	RMDQ, SE: 2.0
Rabin, 2014	67	RCT, 2 SE Ctrl	CLBP	105, 48, 57	Range: 18–60	25/23, 31/26	NRS (0-10), 4,9±1.7, 5.3±1.7	2: 0; 8	NRS (0–10) SE: 1.5	MODI (0–100) SE: 2.0
Rasmussen- Barr, 2003	23	RCT, 2 SE Ctrl	LBP≥6 weeks	42, 22, 20	39±12, 37±10	17/7 18/5	VAS (0-100), 33, 32	4: 0; 6; 12; 24	VAS (0–100) SE: 0.95 Imputed from Saragiotto et al. <sup>2</sup>	ODI SE: 1.18 Imputed from Saragiotto et al. <sup>2</sup>

Study informat	ion		Population					Assessments	Outcomes	
First author, year	Citation number	Study design, no of study arms	Main inclusion criterion LBP (time, other)	N (Total, per grop) (SE, C, C2))	Age Mean±SD (years)	Sex (f/m)	Baseline- pain (Scale, mean, SD if not stated otherwise)	Measurement time points total (N: weeks (if not, stated otherwise) after Baseline)	Primary outcome pain, scale, Co-hens d, (M0-M1)	Primary outcome disability name, Cohens d,, (M0-M1)
Rasmussen- Barr, 2009	26	RCT, 2 SE Ctrl	LBP≥8 weeks	71, 36, 35	36±10, 40±12	18/18, 18/17	VAS (0–100), 32, range 18–59, 38, range 10–47	5: 0; 8; 12; 24; 144	VAS (0-100) SE: 0.99 Imputed from Saragiotto et al. <sup>2</sup>	Oswestry Low Back Pain Questionnaire (OSD), n SE: 1.11 Imputed from Saragiotto et al. <sup>2</sup>
Rhee, 2012	68	RCT, 2 SE Ctrl	LBP	42, 21, 21	53.09±9.04, 50.90±5.24	11/10, 10/11	Million Visual VAS (0–100), 42.7±13.8 32.8±10.9	2: 0; 4	MVAS (0–100) SE: 0.66	ODI SE: 1.14
Salamat, 2017	48	RCT, 2 SE I SE II	extension related non-specific CLBP≥12 weeks	24, 12, 12	$35.83 \pm 9.31,$ $36.09 \pm 9.6$	N.A	VAS (0–10), 5.16±1.74, 5.9±1.9	2: 0; 4	NRS (0–10) SE I: 1.3 SE II; 1,8	ODI SE I: 0.66 SE II: 0.76
Seo, 2019	49	RCT, 2 Ctrl SE	LBP≥12 weeks	26 13 13	$\begin{array}{c} 22.62 \pm 1.58 \\ 22.31 \pm 1.60 \\ 22.92 \pm 1.55 \end{array}$	15/11 7/6 8/5	N.A:	2:4	N.A	ODI SE: 0.86
Shamsi, 2017	50	RCT, 2 SE I SE II	LBP≥12 weeks, VAS 3–6	51, 27, 24	38.9±12.2, 47.0±9.9	33/18,	VAS (0-100), 52.4±9.2, 53.0±9.2	2: 0; 6	VAS (0–100), SE I: 4.0 SE II: 3.1	ODI SE I: 1.3 SE II: 1.1
Shaughnessy, 2004	51	RCT, 2 SE Ctrl	LBP≥12 weeks	41, 20, 21	43±9, 46±11	27/14, 14/6, 13/8	Sf-36 (bodily pain), 31±12, 32±13	2: 0; 10	Sf-36 (bodily pain), SE: 0.9	ODI SE: 0.85
Soundarara- jan, 2016	59	RCT, 2 SE Ctrl	2-year history Of CLBP	30, 15, 15	26.87±2.17, 27.1±2.09	12/18, 6/9, 6/9	VAS (0-10), 6.27±0.70, 6.6±0.74	2: 0; 6	VAS (0–10) SE: 5.06	MODQ SE: 3.3
Sung, 2013	27	RCT, 2 SE Ctrl	Recurrent LBP≥8 weeks	50, 25, 25	Range 27–63, 47.7±8.9, 53.1±9.1	20/30 10/15, 10/15	N.A	2: 0; 4	N.A	ODI SE: 0.26
Ulger, 2017	52	RCT, 2 SE Ctrl	LBP≥12 weeks	113, 57, 56	Range 20–73, 41.6±12.9, 43.1±14.3	67/46, 35/22, 32/24	VAS (0-10), 6.69±1.6, 3.0±2.43	2: 0; 6	VAS (0–10) SE: 2.3	ODI SE: 1.2
Unsgaard- Tondel, 2010	53	RCT, 3 SE Ctrl 1 Ctrl 2	CLPB≥12 weeks	109, 36, 36, 37	Range 19–60, 40.9±11.5, 43.4±10.2, 36.0±10.3	76/33 29/7 23/13 24/13	NRS (0-10), 3.31±1.42, 3.61±1.75, 3.30±1.74	3: 0; 8; 48	NRS (0–10) SE: 0.37	ODI SE: 0.28
Vikranth, 2015	69	RCT, 2 SE Ctrl	mechanical low back pain VAS<5	30, 15, 15	Range 30–45, 37.0±2.76, 37.1±3.51	11/19 5/10, 6/9	VAS (0-10), 3.8±0.83, 3.73±1.06	2: 0; 2	VAS (0–10) SE: 0.5	ODI SE: 0.9
Waseem, 2018	54	RCT, 2 SE Ctrl	LBP≥12 weeks	108, 53, 55	Range 20–60, 46.39±7.43, 45.5±6.61	37/71, 18/35, 19/36	N.A	4: 0; 2; 4; 6	N.A	ODI SE: 1.8
Woo, 2016	55	RCT, 2 SE I SE II	$LBP \ge 12$ weeks	30, 15, 15	N.A., 39.8, 40.1	N.A	N.A	2: 0; 4	N.A	ODI, SE I: 1.85 SE II: 2.37
Young, 2015	70	RCT, 2 SE Ctrl	СВР	48, 24, 24	N.A	N.A	VAS (0-10), 4.3±1.26, 4.0±1.38	2: 0; 6	VAS (0–10) SE: 0.43	N.A

**Table 3.** Study characteristics (left columns) and the individual studies' results (right columns). For each of the studies included, the methodological aspects, participants' characteristics and key results are displayed. Legend: RCT, randomized controlled trial; T, total, E, exercise, SE, stabilisation exercise, Ctrl, control or comparison group; CLBP, chronic low back pain; N, number; f, female; m, male; SD, standard deviation; Mx, measurement visit number, VAS, visual analogue scale; NRS, numeric rating scale; NPRS, numeric pain rating scale; ODI, owestry disability index, RMDQ, Roland Morris disability questionnaire

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Grade C recommendation: At the group level, stabilisation exercise to treat non-specific low back pain is potentially most helpful when it is scheduled three to five times a week.

**Future study.** Nevertheless, the evidence of more detailed training specifica (training intensity: number of exercises per session, repetitions per exercise, sets per exercise, rest after exercise, etc.) remains unclear. Furthermore, the minimal clinically relevant dosage of core-specific stabilisation interventions in chronic, non-specific low back pain patients remains unclear; this may define a future area of low back pain research as there exists a societal pressure of consistently high low back pain prevalence across all lifespans.

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Alp, 2014	56	Lumbar core stabilization exercise (SE)	N.A	Conventional home-based exercise (HE)	6	SE: 3	45–60 (30 MCE)	SE: N.A HE: 1	SE: N.A HE: 20	N.A., N.A
Alrwaily, 2019	28	Stabilization exercise	5; Abdominal bracing (supine), Abdominal bracing (supine) with heel slide, Abdominal bracing (supine) with leg lifts, Abdominal brac- ing (supine) with bridging, Bracing with single leg bridging	Stability exercise combined with neuromuscu- lar electrical stimulation	6	2	20	N.A	N.A	N.A.; N.A
Andrusaitis, 2011	60	Stabilization	2: dorsal decubitus, ventral decubitus	Strengthening	7	3	40	1	6—10	N.A.; 30 sec
Arampatzis, 2017	29	Perturbation- based core training	15–18: 3 different perturbation exer- cises in half-seated position, classical core stability exer- cises on unstable surfaces	No specific training, nor- mal routine	13	2	90	3	60 sec	180—300; 120
Areeudom- wong, 2019	30	Core stabilisa- tion exercise	N.A.: Practiced recruitment of deep trunk muscles, particu- larly transversus abdominis (TrA) and lumbar multifidus (LM) muscles, together with the dia- phragm and pelvic floor muscles, reducing superficial trunk muscle activity in order to improve function of deep trunk muscles and control inter- segmental lumbar spine movement during activities Exercise difficulty was increased by integrating deep muscle cocontrac- tion with control- ling movement of extremities and heavier loading positions, such as bridging, bird-dog position and single knee to chest	Proprioceptive Neuromus- cular Facilitation Training Inactive con- trol group	4	3	30	N.A	N.A	N.A.; 60
Bae, 2018	31	Core stability exercises	6: Abdominal drawing-in in 4-point kneeling and supine posi- tion, Opposite upper and lower extremity lift in quadruped posi- tion, Straight leg raise exercise in prone position, Supine lower extremity extender in supine position, Straight leg raise exercise in supine position, Horizon- tal side-support exercise in side lying position	Assisted sit-up exercise (SUE)	4	3	30	N.A	N.A	N.A.; N.A

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Bauer, 2019	22	Neuromuscu- lar exercise	9; Modified curl up, Bird dog, Side bridge/Mermaid, Single leg stretch, Shoulder bridge, Weight transfer side lunge and one leg stand, "Tai chi warrior", Lifting up an imaginary Ball, To achieve normal range of motion in thoracic region, and hip and ankle joints,	Inactive con- trol group	24	2	60	N.A	N.A	N.A.; N.A
Brooks, 2012	32	Specific trunk exercise group (SEG)	N.A: Included skilled cognitive activation of the trunk muscles in addition to a num- ber of other best practice exercises: Skilled abdominal contractions and postural training, Side lying trunk exercises (mat- based), Prone lying trunk exercises (mat-based; Hip- specific exercises, Upper and lower limb-focused exercises, Full- body exercises (reformer-based)	Seated cycling	8	3	50-60	N.A	N.A	N.A.; N.A
Chung*, 2018	61	Core stability exercises with flexi bar	4: Abdominal drawing-in maneu- ver in standing, hook-lying, quadruped, and prone positions by maintaining each motion for 10 s. It was used both hands holding the FB	No further, both groups SE	6	3	30	3	10	180; N.A:
		Core stability exercises	4: Abdominal drawing-in maneu- ver in standing, hook-lying, quadruped, and prone positions by maintaining each motion for 10 s	No further, both groups SE	6	3	30	3	10	180; N.A:
Critchley, 2007	33	Spinal stabili- zation (SS)	5: individual trans- versus abdominis and lumbar multifidus muscle training followed by group exercises that challenged spinal stability	Physio Pain Management	8	8	90	Individual	Individual	individual
Da Luz, 2019	62	Core stability exercise	4; prone bridge, supine bridge, side bridge, bird dog with lower limb elevation As the partici- pants progressed throughout the program, the degree of difficulty of the exercises increased	Core stabil- ity exercise combined with neuromuscu- lar electrical stimulation; neuromuscu- lar electrical stimulation only	4	3	N.A	10	N.A	N.A., 60

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Demirel, 2019	34	Stabilization exercise	4–5; The TA and multifidus muscles were contracted together with diaphragm respira- tion appropriately in basic positions (supine, prone, standing, sitting and crawling posi- tions) Progress over the six weeks included different positions, use of resistance bands	Yoga exercises	6	3	60	N.A	N.A	N.A.; N.A
Ferreira, 2007	35	Motor control exercise	N.A.: Improving function of specific trunk muscles thought to control inter-segmental movement of the spine, includ- ing transversus abdominis, mul- tifidus, the dia- phragm and pelvic floor muscles	General exercise Spinal manipulation therapy	8	12	N.A	N.A	N.A	N.A.; N.A
Franca, 2012	36	Segmental stabilization exercises (SSEs)	4: exercises for the TrA in 4 point kneeling, exercises for the TrA in dor- sal decubitus with flexed knees, exer- cises for the LM in ventral decubitus, Cocontraction of the TrA and LM in the upright position	Stretching (ST)—focused on stretching the erector spinae, ham- strings, and triceps surae	6	2	30	3	15	N.A.; N.A
Ghorbanpour, 2018	57	McGill stabilization exercises group	3: Curl up, Side Bridge, Bird Dog with one hand or one foot and one hand and the opposite leg	Conven- tional physio (strengthening, stretching, flexibility)	6	3	30	3	10	N.A.; 120
Hosseinifar, 2013	37	Spinal stabali- zation seercise	N.A	McKenzie Method	6	3	60	10	N.A	N.A.; N.A
Hwang, 2013	71	Sensorimotor training	6: Hollowing exercise, Single leg raising in the quadruped posi- tion, contralateral arm and leg raising in the quadru- ped position, abdominal bracing Holding a bridging position, single leg raising in the bridging position	2 Group: 1 healthy controls ©, 1 lbp physical therapy (C LBP)	4	5	40	N.A	N.A	N.A.; N.A
Ibrahim, 2018	38	Motor control exercise	4–12; Abdominal drawing in in supine, in quadru- ped, in sitting, in standing, in supine with heel slide, in supine with leg lift (each leg), in supine with bridg- ing, in supine with single-leg bridge, with curl-up, hori- zontal side support with knees bent, in quadruped with leg raise, etc	Motor control exercise plus patient educa- tion; Patient Education only	6	2	30	N.A	10	N.A.; N.A

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Inani, 2013	63	Core stability exercies	4: Slow curl ups, sit ups, oblique plank/side bridge, bird dog	Conventional Exercise	12	N.A	N.A	N.A	N.A	N.A.; N.A
Khodadad, 2019	39	Lumbar Stabi- lization	5; Elbow-Toe, Back Bridge, Hand- Knee, Side Bridge, Curl up	Cognitive functional treatment; Inactive con- trol group	8	3	60	N.A	N.A	N.A.; N.A
Kim 2018*	40	McGill's exercise; Sahrmann 0–5 level Exercise	N.A.; curl up, side bridge, and bird dog	No further, both groups SE	8	3	30	N.A	N.A	N.A.; N.A
Kiiii, 2018		Stabilization exercise	N.A.; Pro bal- ance trainer and dynamic air cush- ion training	No further, both groups SE	8	3	30	N.A	N.A	N.A.; N.A
Kim, 2019	41	Stabilization exercise	4: supine pelvic lift, supine and prone bridging exercise, and side-lying hip abduction	Simulated horseback riding	8	2	30	N.A	N.A	N.A.; N.A
Ko, 2018	64	Lumbar stabi- lization (LS)	8: sit up, superman, quadruped arm & leg raise, squat, lower body fixation plank, upper body fixation plank, side plank, hip bridge	2 Groups: Sling, Control	12	3	60 min (40 min MCE)	3	10	60; 60
Kofotolis, 2016	42	Pilates	16: Roll down, mermaid, spine stretching, pelvic curl, criss-cross, double leg stretch, hundreds, double knee folds, table top, swimming, swan, catstretch, child's pose, hips stretch	General strengtheing/ stabilisation exercise, control	8	3	60	Progressive: 2 (until week 4), then 3	Progressive: 15 (week 1–2), 20 (w 3–4), 15 (5–6), 20 (7–8)	120; 30
Lee, 2014	65	Ball exercise group	10: exercises on swiss ball from sit- ting to bridging	PNF pattern group	6	4	N.A	2	20	15; N.A
Lee, 2011	25	Core stability exercises	5: upper body extension in prone position, alternate arm and leg lift in quadruped position, alternate arm and leg lift in prone position, diagonal curl-up and straight curl- up in supine position, quad- ruped exercises, performed from an all-fours position with the arms and legs extending	Control	4	4	20	N.A	N.A	N.A.; N.A
Letafatkar, 2017	66	SMT- Perturbation with HUBER machine	10: upright stance, push and pull with oscillatory perturbative move- ments of variable amplitude and speed	Control	5	2	30-45	2-4	2-6	N.A.; 300
Continued										

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Liu, 2019	43	Core Stabiliza- tion Exercise on Swiss ball	6: Glute Bridge Pose, Single Leg Bridge, Bridge and Double Knee Flex, Single Leg Bridge and Double Knee Flex, Reverse Bridge, Reverse Bridge and Hip and Knee Flex	Chen-Style Tai Chi	12	3	60	N.A	N.A	N.A.; N.A
Lomond, 2015	58	Trunk stabili- zation	N.A.: 3 compo- nents of spinal stability	Movement System Impair- ment (MSI)	6	1	45-60	N.A	N.A	N.A.; N.A
Macedo, 2012	44	MCE	N.A	Graded activity	8	2 (4 weeks), 1 (4 weeks)	60	1	10	N.A.; N.A
Marshall, 2013	45	MCE, Pilates	8: Whole body stretching, Skilled abdominal con- tractions and pos- tural training, side lying trunk, prone lying trunk, hip specific exercises, upper and lower limb, full body exercises, whole body stretching	Stretching and cycling	8	3	55	N.A	N.A	N.A.; N.A
Miller, 2013	24	Stabilzing MCE	10: Phase one: Prone, Supine, Quadruped; Phase two: Supine leg machine, Quad- ruped - Alternate arm lifts, Alternate leg lifts, standing; Phase three: Quadruped-Alter- nate arm and leg lifts, Standing with rotation, Bridging	McKenzie	6	N.A	10-15	1	10-50	N.A.; N.A
Moon*, 2013	46	Lumbar stabilization exercises,	16: aimed to strengthen the deep lumbar stabilizing muscles: the transversus abdominis, lumbar multifidi, and internal obliques	No further, both groups SE	8	2	60 (35 min LSE)	1	10	N.A.; 60
		Lumbar dynamic strengthening exercises	14: activated the extensor (erector spinae) and flexor (rectus abdominis) muscle groups	No further, both groups SE	8	2	60 (35 min LDSE)	1	10	N.A.; 60
Noormoham- madpour, 2018	47	Multi-step core stability exercise	4: 2 on floor; 2 on swiss ball	Waiting list	8	3		3	10	N.A.; N.A
Rabin, 2014	67	Lumbar stabilization exercise	4: Quadruped, sidelying, supine, and standing posi- tions	Manual therapy	8	Supervised: 2×first 4 weeks; 1×week 5–8;	N.A	N.A	N.A	N.A.; N.A
Rasmussen- Barr, 2003	23	Stabilizing training	6-8: motor control, supine crooked- lying, four-point kneeling, prone, sitting and stand- ing	Manual therapy	6	1 supervised, 1 homebased	45 supervised, 10–15 unsu- pervised	3	15	N.A.; N.A
Rasmussen- Barr, 2009	26	Graded stabi- lizing exercise	7: supine crooked- lying, four-point kneeling, prone, sitting, standing	30-min walk every day	8	1 supervised, 1 homebased	45 supervised, 10–15 unsu- pervised	3	15	N.A.; N.A
Continued										

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Rhee, 2012	68	Specific local- ized exercises aimed at restoring the stabilizing protective function of the spinal muscles around the spinal joint	5: Upper-body extension, alternate arm and leg lift, alternate arm and leg extension on all fours, diagonal curl-up, curl-up	Advice regard- ing bed rest, absence from work, prescrip- tion medica- tions, and resuming normal activity as tolerated	4	5	N.A	N.A	N.A	N.A.; N.A
Salamat*, 2017	48	Movement control	N.A.: The aim of the intervention was to normal- ize the abnormal movement patterns and postures and to relax trunk muscles. Exercises involved train- ing to modify pain provoca- tive postures and movement patterns in order to decrease pain while performing the task	No further, both groups SE	4	2	45	3	15–30	60 – 120; 300
		Stabilization exercise	N.A.: Exercises involved coor- dinated training and independent activity of deep trunk muscles including trans- versus abdominis and multifidus in pain-free positions and movements	No further, both groups SE	4	2	45	3	15-30	60 - 120; 300
Seo, 2019	49	Trunk stability exercise	16: nine move- ments of mat- based trunk stabil- ity exercises and seven movements of Swiss ball trunk stability exercises	Gyrotonic exercise	4	3	N.A:	N.A	N.A	N.A.; N.A
		MCE	N.A.: Progressive classic stabilization	No further, both groups SE	6	3	20	N.A	10	N.A.; N.A
Shamsi*, 2017	50	Core	N.A.: Exercises were performed in a lying position starting with simple movements and advancing to more difficult exercises (e.g. on a Swiss ball)	No further, both groups SE	6	3	20	N.A	10	N.A.; N.A
Shaughnessy, 2004	51	Core	3: Prone lying, kneeling, supine	Standard physiotherapy	10	2 (week 1–2), 1 (week 3–10)	60 week 1, else 30	N.A	Max. 10	N.A.; N.A
Soundarara- jan, 2016	59	Multifidus muscle retraining	8: Bridging, lying prone, quadruple, prone lying, leg extension, sitting, standing, shoulder flexion	Traditional back exercises (strength and stretching)	6	3	20	1	20	N.A.; 120–240
Sung, 2013	27	Core	5: Knee to chest for each leg in supine position, double leg knee to chest in supine position, prayer stretch on all fours, leaning forward position while sitting, lat- eral side stretch in standing position	Flexibility	4	1 supervised, 6 homebased	20	2	15	N.A.; N.A

First author, year	Citation number	Type intervention (MCE, Core)	Exercises (No; Name): (Description/ Name of exercises)	Type comparator(s)	Training period (weeks)	Training frequency (sessions per week) scheduled, real	Training duration (minutes per session)	Sets (number per exercise)	Repetitions (per set per exercise)	Rest (between sets per exercise; between exercises in seconds)
Ulger, 2017	52	Stabilization	N.A: Increasing intensity and changing exercises once/week	Manipulation	6	3	60	3	10	N.A.; N.A
Unsgaard- Tondel, 2010	53	Sling	N.A: Sling	Low-load MCE (feedback) and General exercise	8	1	40	N.A	N.A	N.A.; N.A
Vikranth, 2015	69	Core stabiliza- tion	8: Week 1: Trans- versus abdominus activation, trans- versus abdominus marching, pelvic tilt, segmental bridge; Week 2: Fall out, modified crunch, cat stretch, back extension	MCE (passive)	2	5	35		Week 1: 8; week 2: 15	120; N.A
Waseem, 2018	54	Core stabiliza- tion	7: Pressure feed- back core exercise, multifidus exercise, frontal and side plank exercise, pel- vic floor exercises, diaphragmatic strengthening, single leg standing on foam, tandem standing with perturbation	Routine exercise	6	1 supervised, 2 homebased	N.A	N.A	N.A	N.A.; N.A
Woo*, 2016	55	Lumbar stabilization exercise	6: Lower extremity lifting in a bridge posture, lower extremity lift in a prone position on a ball, upper extrem- ity lift in a prone position on a ball, moving the body forward grasping a sling in a kneeling position, lifting the buttocks with the lower extrem- ity hooked on a sling in a supine position;	No further, both groups SE	4	N.A	40 (30 min MCE)	Group A: 4 Group B: 2	10-12	N.A.; N.A
		Lumbar stabilization exercise with thoracic exten- sion exercise	10: Lower extrem- ity lifting in a bridge posture, lower extremity lift in a prone position on a ball, upper extremity lift in a prone position on a ball, moving the body forward grasping a sling in a kneeling position, lifting the buttocks with the lower extremity hooked on a sling in a supine posi- tion; plus thoracic extension exercise	No further, both groups SE	4	N.A	40 (30 min MCE)	Group A: 4 Group B: 2	10-12	N.A.; N.A
Young, 2015	70	Swiss ball stabilization	N.A	PNF	6	3	50	N.A	N.A	N.A.; N.A

**Table 4.** Individual studies' training specifications. All interventions and the respective comparators are described. exercises, stabilisation exercise; N.A., not applicable. \*Both groups were included into quantitative analysis (meta-regression).

	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	в	95% CI. LL, UL	p-value
A Pain		1	I	1		I	
Intervention: duration [weeks]					009	1, .08	.8
Intervention: frequency [N <sub>Trainings</sub> / week] data transformed from U-shaped association	1				.164	239, .567	.4
Intervention: Time per session [minutes] Data transfomed from negative log associa- tion	.445	1.8	40	31	- 1.75	- 2.61,879	.0001
PEDro sum score [points]					17	36, .016	.07
Sample size (MCE)					.005	016, .026	.6
	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	в	95% CI. LL, UL	p-value
B Disability	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	В	95% CI. LL, UL	p-value
B Disability Intervention: duration [weeks]	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	<b>B</b>	<b>95% CI. LL, UL</b> - 0.3, 0.95	p-value
B Disability Intervention: duration [weeks] Intervention: frequency [N <sub>Trainings</sub> / week] data transformed from U-shaped association	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	в .1 .26	<b>95% CI. LL, UL</b> - 0.3, 0.95 61, 1.1	<b>p-value</b> .3 .6
B Disability         Intervention: duration         [weeks]         Intervention:         frequency [N <sub>Trainings</sub> /         week]         data transformed from         U-shaped association         Intervention: time per         session [minutes]         Data transformed from         negative log associa-         tion	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	B .1 .26 - 1.0	<b>95% CI. LL, UL</b> - 0.3, 0.9561, 1.1 - 3.1, .95	<b>p-value</b> .3 .6 .3
B Disability         Intervention: duration         [weeks]         Intervention:         frequency [N <sub>Trainings</sub> /         week]         data transformed from         U-shaped association         Intervention: time per         sesion [minutes]         Data transformed from         negative log associa-         tion         PEDro sum score         [points]	Model R <sup>2</sup>	Mean effect size	N effect sizes included	Homogeneity Q	B .1 .26 - 1.0 04	95% CI. LL, UL - 0.3, 0.95 61, 1.1 - 3.1, .95 37, .30	p-value           .3           .6           .3           .6           .3           .8

**Table 5.** Outcomes of the sensitivity meta-regressions. For each single analysis, effect sizes, number of included effect sizes, homogeneity, the regression coefficient B, its confidence interval (CI) and the corresponding p-value are displayed. Legend: LL, lower level, UL, upper level.

#### Limitations

**Limitations at the study and outcome levels.** A common limitation in exercise trials is the limited possibility to blind the participants. This limitation is increased by the self-reported assessment of pain and pain-related function.

**Limitations at the review level.** We only screened the databases PubMed (Medline), Web of Knowledge and the Cochrane Library. Considering the topic of our review, almost all manuscripts of interest should be found therein<sup>73–75</sup>. However, expanding the search to even more databases, like EMBASE, PEDro, CINAHL; AMED, and CENTRAL may would have led to slightly more hits.

The advantage of meta-regressions are, inter alia, that the interventional effect sizes are compared to each other to find a dose-response-relationship, the effect sizes are thus relativized to each other. The estimates found are valid for the isolated intervention group effects comparisons, given by the meta-regression. The mean effects are, given by the nature of the meta-regression, absolute and not in comparison to a control/comparator. The mean effect sizes (refer to the study description and meta-regressions) are thus not directly comparable to those found in meta-analyses where the effects are calculated in comparison to a control/comparator group.

The funnel plot analysis revealed an unclear, but rather low, risk of publication bias within our review. The findings of our (retrospective) meta-regression should be confirmed prospectively, at best adopting a prospective meta-analysis.

Sensitivity of the interventions' name. The interventions of the studies included into our meta-analysis are defined as stabilization exercise. Motor control exercises are classically defined as core-specific dynamic stabilization exercises with an a priori education on deep trunk muscles activation and/or the control of deep muscles activation during exercising. We only included studies with dynamic/exercise parts. When solely stabilisation exercises without pre-conditioning are performed, they are often called "coordination", "stabilisation"<sup>5</sup>, "senso-rimotor"<sup>76</sup> or even as well "motor control"<sup>2</sup> exercise. As described above, the term "motor control exercise" may be slightly too sensitive for the interventions included into our review. In contrary, the terms "sensorimotor", "coordination" and "stabilisation" training/exercise may be too general. Consequently, we name the intervention "stabilisation exercise" to highlight that the stabilisation/active/dynamic parts of the originally described



**Figure 2.** Meta-regression bubble plots for the dependent variable Cohens d (pain), independent variable training period (weeks, A), training frequency (times/week, B) and training duration (minutes, C). The weighting is illustrated by the size of the bubbles.





as "motor control exercise"-theorem are adopted. Nevertheless, the intervention could also be called "motor control stabilization exercise" or "sensorimotor exercise".

#### Conclusions

A training frequency of 3 to 5 times per week (low quality evidence) with a training duration of 20 to 30 min (moderate quality evidence) per session causes the largest impact on the effect sizes (both in pain and disability) of stabilisation exercise in low back pain patients. However, the training period showed no systematic impact on the effect size for pain intensity. Future work is required to enhance the quality of the evidence of our findings, possibly focussing on the definition of a minimum dosage.

Received: 29 May 2020; Accepted: 23 September 2020 Published online: 09 October 2020

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## Author contributions

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## Funding

Open Access funding enabled and organized by Projekt DEAL. This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

## **Competing interests**

The authors declare no competing interests.

## Additional information

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