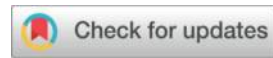




Summary of Best Evidence for Pelvic Floor Muscle Training During Pregnancy

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Abstract

Background: Pelvic floor muscle training during pregnancy is a first-line, non-invasive intervention for the prevention of pelvic floor dysfunction. Although its benefits have been widely recognized, existing evidence remains fragmented across different sources, and standardized clinical recommendations are lacking. A comprehensive synthesis of the best available evidence is urgently needed to guide clinical practice.

Objectives: To provide an evidence - based approach for the clinical practice of pelvic floor muscle training, thereby promoting pregnancy health management and long - term quality of life in women.

Methods: Guided by the 6S evidence pyramid model, a top-down search was conducted from database inception to December 30, 2025. Two reviewers independently screened, extracted, and synthesized evidence following the Joanna Briggs Institute methodology, and graded the evidence level and recommendation strength.

Results: A total of 33 publications were included (8 guidelines, 3 clinical decisions, 1 expert consensus, 4 evidence summaries, and 17 systematic reviews). Following a comprehensive analysis, 26 evidence-based recommendations were developed, which were categorised into 7 aspects: overall suggestion, cost-effectiveness, efficacy, safety, timing, protocol, and adherence.

Conclusions: This study summarised the best evidence of pelvic floor muscle training in pregnant women. By adhering to the best available evidence, healthcare providers and pregnant women can work cooperatively to promote delivery outcomes, enhance the pelvic floor muscle, and reduce the incidence of pelvic floor dysfunction.

Keywords: pelvic floor muscle training, best evidence, intervention, pregnancy

Brief Summary: This study systematically summarizes the best evidence for pelvic floor muscle training during pregnancy for clinical application.

Introduction

Over the past few decades, significant progress has been made globally in reducing maternal mortality rates. However, this achievement has been accompanied by a relative neglect of postpartum morbidity arising from pregnancy and childbirth (e.g., pelvic floor dysfunction and depression). These conditions can impose substantial social, economic, and health burdens.

As stated in the *Yellow Emperor's Inner Classic*, “The superior physician treats disease before it arises.” Pregnancy and childbirth are independent risk factors for pelvic floor dysfunction (PFD) [Error! Unknown switch argument.]. Pelvic floor muscle training (PFMT) is the first-line method for preventing PFD, which is effective at all stages of tertiary prevention. This indicates that it is extremely necessary to take proactive intervention measures during pregnancy [Error! Unknown switch argument.-Error! Unknown switch argument.].

Currently, there is no clear clinical guidance on how to perform PFMT during pregnancy, nor is there a standardised operating procedure. This directly results in significant heterogeneity in actual clinical practice and relevant research. Key challenges to PFMT include the following: (a) Unclear start timing; (b) Suboptimal adherence; (c) Inconsistent training efficacy; (d) Insufficient health education efforts by medical staff [Error! Unknown switch argument.,Error! Unknown switch argument.-Error! Unknown switch argument.]. This heterogeneity potentially increases women's susceptibility to additional health risks and associated complications. To address this critical evidence-practice gap, relevant domestic and international clinical decisions, guidelines, systematic reviews, and other studies were retrieved to form the best evidence, facilitating the translation of evidence into clinical practice for PFMT during pregnancy to promote women's health.

Method

Problem establishment

The PIPOST (P [population], pregnant women; I [intervention], PFMT; P [professional], medical staff; O [outcome], PFMT-relevant outcomes; S [setting], hospitals, communities, and families; and T [type of evidence], guideline, systematic review, expert consensus, evidence

summary, best practice, and clinical decision) method was applied to define the key research questions for this study [Error! Unknown switch argument.] .

Search strategy

Evidence retrieval is searched from the top-down according to the ‘6S’ evidence resource model [Error! Unknown switch argument.].The following databases were searched: UpToDate; BMJ Best Practice; National Institute of Health and Clinical Excellence (NICE); Scottish Intercollegiate Guidelines Network; New Zealand Guidelines Group; Guideline International Network; JBI Database; Cochrane Database; Society of Obstetricians and Gynaecologists of Canada (SOGC); Polish Society of Gynecologists and Obstetricians (PSGO); American Urological Association (AUA); European Association Of Urology (EAU); Canadian Urological Association (CUA); Department of Health and Aged Care (DHAC); International Consultation on Incontinence (ICI); International Continence Society (ICS); Registered Nurses’ Association of Ontario (RNAO); Chinese Nursing Association; Embase; PubMed; Web of Science; SinoMed; CINAHL; Yiigle; China National Knowledge Infrastructure (CNKI); Weipu; and Wanfang. The development of search queries involved a hybrid approach, utilising both MeSH terms and free words. Keywords were captured using terms like “pelvic floor muscle training / pelvic floor muscle exercise / conservation management / non-surgical treatment / physical therapy / Kegel exercise” and “pregnancy / gestation / gravidity / pregnant women / delivery / labour / antenatal / perinatal”. The search deadline is December 30,2025.

Literature inclusion and exclusion criteria

The selection criteria were as follows: (i) pregnant women; (ii) relevant to PFMT; (iii) guidelines, clinical decisions, evidence summaries, best practices, systematic reviews, or expert consensus; (iv) Published in Chinese or English. Grounds for exclusion were: (i) literature presented as guideline interpretations, republished, or updated versions and (ii) incomplete information, unavailable full text, or low quality.

Study selection and data extraction

The screening phase involved two independent reviewers assessing all retrieved records based on the inclusion and exclusion criteria. Any disagreements that arose were settled either by consensus following discussion or through decision by a third party. Following screening, the same reviewers proceeded to data extraction. This extraction was conducted independently using a

standardised template, with reviewers blinded to each other's work to minimise bias. Specific data collected encompassed study characteristics such as the first author, publication year, country, source, evidence type, and the title.

Literature quality evaluation criteria

We assessed the quality of the literature using a tiered strategy, where each document type was evaluated with its specific, corresponding critical appraisal instrument. (1) Guidelines were assessed with the AGREE II tool [Error! Unknown switch argument.]. (2) Systematic reviews were appraised with AMSTAR 2 [Error! Unknown switch argument.]. (3) The evidence summary and best practice were evaluated using the Critical Appraisal for Summaries of Evidence (CASE) tool [Error! Unknown switch argument.]. (4) For expert consensus, the standardised criteria from the JBI Centre were applied [Error! Unknown switch argument.]. (5) Evidence from authoritative databases (UpToDate and BMJ Best Practice) was regarded as high-quality evidence and adopted.

Literature quality evaluation process

To ensure methodological rigour, two independently working reviewers, both grounded in evidence-based nursing, performed the quality evaluations. Disagreements were adjudicated by a third party with expertise in evidence-based research. The guiding principle for inclusion was to resolve conflicts among the extracted evidence by prioritising those derived from explicit evidence-based processes, possessing higher quality assessments, and representing the most recent and authoritative sources.

Criteria for determining evidence and recommendation levels

The evaluation of the incorporated evidence was guided by the JBI grading system [Error! Unknown switch argument.]. This framework operates on a two-component principle: first, evidence is assigned to one of five levels (1-5) according to its research design; second, its recommendation strength is classified through the FAME structure (assessing Feasibility, Appropriateness, Meaningfulness, and Effectiveness) as either Grade A (strong) or Grade B (weak).

Ethics Approval

Not applicable. This review synthesizes published, publicly available literature; no participants or identifiable data were involved.

Results

General characteristics of the included literature

From an initial retrieval of 16,831 citations, a rigorous screening process involving deduplication and sequential review of titles, abstracts, and full texts yielded 33 articles for final inclusion. The composition of the evidence base included 8 guidelines [6,18-24], 3 clinical decisions [Error! Unknown switch argument.-Error! Unknown switch argument.], 1 expert consensus [Error! Unknown switch argument.], 2 best practices [Error! Unknown switch argument.,Error! Unknown switch argument.], 2 evidence summaries [29,30] and 17 systematic reviews [2,9,31-45]. The study selection flow is depicted in Figure 1, with the general information of the included studies provided in Table 1.

Quality evaluation results of the included literature

(1) Guidelines: The guideline appraisal encompassed eight documents [6,18-24]. Three received a Grade A recommendation for scoring $\geq 60\%$ on all six domains [18,19,22], while the rest were graded B [6,20,21,23,24]. The standardised domain scores and overall evaluations are available in Table 2. (2) Clinical decision: This study included three clinical decisions, all of which were from UpToDate, of high quality, and approved for inclusion [Error! Unknown switch argument.-Error! Unknown switch argument.]. (3) Expert Consensus: One expert consensus met all appraisal criteria ("yes" on all items) and was included [Error! Unknown switch argument.]. (4) Best Practices & Evidence Summaries: This paper included four best practice articles and evidence summaries, including 2 from JBI [7,8] and 2 from Yiigle [29,30]. The quality assessment results are shown in Table 3. (5) Systematic reviews: The final set contained 17 systematic reviews, among which 6 were from PubMed [2,9,31,40-42], 4 from Embase [39,43-45], 1 from Cochrane Library [Error! Unknown switch argument.], 2 from Yiigle [32,37], and 4 from CNKI [Error! Unknown switch argument.-Error! Unknown switch argument.]. Their quality evaluation results are shown in Table 4.

Summary and description of evidence

Data extraction was performed on the final included literature, and the body of evidence was subsequently assessed using the JBI evidence grading and recommendation system. Through a process of synthesis and integration, the evidence was organised into seven key categories, leading to the formulation of 26 best-practice recommendations (Table 5). The recommendations addressed 7 key categories: overall suggestion, cost-effectiveness, efficacy, safety, timing, protocol, and adherence.

Discussion

Overall suggestion

Preventing and managing PFD are essential for improving the quality of life among pregnant and elderly women while reducing the medical care burden on families and society. Numerous studies [6,18-24] indicated that pregnant women without contraindications should be encouraged to perform PFMT during pregnancy to prevent PFD and induce labour. Despite increasing attention to PFD and the rising popularity of PFMT during pregnancy, current management approaches remain unscientific and unstandardised [18,28,30]. It is recommended that a multidisciplinary team be formed to provide guidance on the assessment and treatment of PFD, including referral pathways to relevant specialists.

Cost-effectiveness

Globally, urinary incontinence (UI) is acknowledged as a widespread and costly condition affecting women. PFMT during pregnancy has been shown to be a cost-effective strategy for managing UI [18,31,45]. Brennen's team identified group-based PFMT as the most economically viable approach, with its value being sensitive to both attendance numbers and direct costs to participants [**Error! Unknown switch argument.**]. However, the current body of evidence supporting this has limitations, including relatively small studies and methodological constraints, resulting in evidence of mostly low-to-moderate quality. It is thus recommended that subsequent trials include comprehensive economic analyses or report sufficient intervention details to enable cost projections by other researchers and healthcare providers.

Efficacy

The integrated lifespan model proposed by De Lancey et al. provides a conceptual framework for understanding the impact of biological and lifestyle factors on pelvic floor integrity and performance [**Error! Unknown switch argument.**]. From a clinical perspective, the pelvic floor musculature possesses significant plasticity, allowing for targeted neuromuscular re-education. Such training can optimise its contribution to urethral closure mechanisms and substantially augment its functional reserve capacity [**Error! Unknown switch argument.**]. Handa et al. concluded that continent women who initiate PFMT during pregnancy are 30% less likely to develop UI up to 6 months following birth than women who do not engage in PFMT [**Error! Unknown switch argument.**]. There is insufficient high-quality evidence to show that PFMT before the birth reduces

the risk into the late postnatal period (i.e., > 6 months postpartum) [6,19,22,29]; it is reasonable to assume that to experience long-term benefits, any training must be continued long-term. Faecal incontinence (FI) is also painful and more common in parous women [Error! Unknown switch argument.]. Longitudinal research demonstrates a persistent pattern of FI following childbirth, affecting approximately 35% of women during pregnancy and 25% at one year postpartum, with a gradual decline to 12-14% over the subsequent 6-12 years. This long-term trajectory contrasts with findings from antenatal intervention [Error! Unknown switch argument.]. A systematic review by Woodley et al., which incorporated eight trials, did not identify a statistically significant reduction in FI prevalence during late pregnancy from antenatal PFMT (RR, 0.64; 95% CI, 0.36-1.14; moderate-quality evidence) [Error! Unknown switch argument.]. This conclusion aligns with the work of Zhang et al., who attribute the finding to the scarcity of FI-focused studies and a lack of large-sample data in the current evidence base [Error! Unknown switch argument.].

Pelvic organ prolapse (POP) is related to ageing, pregnancy, and childbirth. For elderly women, surgical treatment is usually required. Professors Hagen and Stark did a systematic review [Error! Unknown switch argument.], to compare how well conservative treatment works versus no treatment or other therapies for preventing and treating POP. Their findings showed that PFMT helps improve symptoms of POP. At this point, most guidelines [Error! Unknown switch argument.] suggest that pregnant women without exercise contraindications should do PFMT. Similarly, lumbopelvic pain [Error! Unknown switch argument.] is a frequently reported complaint during pregnancy, and for many women, this discomfort extends into the postpartum period. Vesentini's team conducted a meta-analysis of PFMT's effect on lumbopelvic pain [Error! Unknown switch argument.]. They pointed out that even though the results are statistically significant, the real efficacy is weak, and the trials exhibit high heterogeneity. But the impact of pain on women's daily lives and work is real [52,53], so its effectiveness still needs to be further explored.

In many countries, sexual dysfunction is wrapped in silence and stigma-due to cultural taboos, biological factors, and a history of sexual abuse. The topic has turned increasingly enigmatic. This societal silence has built an invisible wall, causing a lot of patients to struggle in the maze. Sexual dysfunction is relatively common among postpartum women. A prospective observational study noted that approximately 41%-83% of women experience sexual dysfunction within two or three months postpartum [Error! Unknown switch argument.], with around 64% still having it six

months later. Dr Jorge's meta-analysis of 4 studies noted that the PFMT group saw a 7.67-point increase in sexual function scores [Error! Unknown switch argument.], such as sexual arousal, orgasm, and satisfaction. As well as Dr Jorge's research, Zahra and other scholars conducted a systematic review to explore the impact of PFMT on sexual function and quality of life [Error! Unknown switch argument.]. The results indicated that PFMT can enhance sexual function in both primiparous and multiparous women. Beyond its benefits for sexual health, considerable research also points to a significant ancillary advantage: PFMT contributes to a reduction in the duration of the first and second stages of labour [8,9,32,33]. This finding suggests that continuous PFMT during pregnancy can help pregnant women improve the contraction and relaxation of pelvic floor muscle fibres. However, there is heterogeneity in the reduction of episiotomy and perineal tear rates. Accordingly, further high-quality research on episiotomy and perineal tears is needed in the future.

Safety

The meta-analysis by Woodley et al., which encompassed a substantial cohort of 10,832 women across 21 countries from 46 trials, affirmed the favourable safety of PFMT. Reports of adverse events were exceptionally rare, with only two participants withdrawing from a prevention study due to pelvic floor discomfort among 43 enrolled [Error! Unknown switch argument.]. No other trial documented any further adverse effects attributable to the intervention. The EAU guideline also pointed out that PFMT rarely causes adverse events, with no serious adverse events reported (1 level of evidence) [Error! Unknown switch argument.]. In addition, both studies indicate that pelvic floor training is unlikely to be harmful in most situations. Most adverse reactions are reversible and can be resolved by reducing the training dose [27,43]. Thus, medical staff should address concerns among pregnant women about PFMT during pregnancy in advance when providing health education.

Timing

It's generally accepted that pregnancy is a physical sensitivity phase, but there remains no definitive consensus on the best timing to start PFMT. NICE guidance emphasises that promoting pregnant women's awareness and training of PFMT during this period is crucial [Error! Unknown switch argument.]. Based on 3 considerations: the perinatal period serves as a unique intervention window, not only a critical time when potential symptoms first emerge and individual risk factors can be easily identified, but also the golden timing to initiate primary prevention strategies.

Although reported starting points in trials vary widely from the 11th to the 28th week of pregnancy [Error! Unknown switch argument.], Olander et al. reframe the entire pregnancy and postpartum journey as a series of discrete, opportune moments. Contacts such as the first prenatal visit and the postnatal check-up are thus seen as behavioural tipping points that can either facilitate or inhibit adherence to preventive training [Error! Unknown switch argument.]. Consequently, the optimal timing for pregnant women to begin training requires further high-quality randomised controlled trials to determine and further promote clinical practice.

Protocols

There is currently no standard PFMT protocol internationally, and there are many uncertainties regarding the specific implementation of PFMT in clinical practice, such as the intensity, frequency, duration, position, and whether auxiliary tools are needed. There is a lack of sufficient clinical data and expert consensus to support the efficacy of PFMT. Sun et al. emphasise that effective pelvic floor muscle training protocols should adhere to the FITT principle (Frequency, Intensity, Time, and Type) [Error! Unknown switch argument.]. Elaborating on this, Professor Kari Bø has outlined a minimal protocol for beginners: performing 2-3 sets daily (on at least 3 days per week) over 8-12 weeks, with each set comprising 8-10 near-maximal contractions held for 6-8 seconds [Error! Unknown switch argument.]. Practice guidelines further specify that a home-based regimen should incorporate basic voluntary contractions (1-2 second holds with equal rest) and sustained voluntary contractions (initially 6-10 seconds with equal rest). The duration of sustained contractions should be progressively increased up to 10 seconds once a baseline is established [Error! Unknown switch argument.].

Acknowledging the preference for easily remembered regimens, Woodley et al. advocate for a cognitively streamlined prescription. They propose an evidence-based minimum standard for pelvic floor muscle strengthening: a sequence of 8 contractions, each sustained for 8 seconds followed by an 8-second rest, performed 3 times daily on 3 days per week over a 3-month period [Error! Unknown switch argument.]. Another guideline from China recommends contracting the pelvic floor muscle for at least 3 s, relaxing for 2-6 s, and repeating this for 15-30 min, 3 times a day, or performing 150-200 contractions a day for ≥ 3 months [Error! Unknown switch argument.]. The use of different PFMT protocols can affect the efficacy of PFMT for pregnant women and subsequent treatment, so it is necessary to implement a standardised PFMT protocol.

Future research should strive to describe protocol details in a detailed and standardised manner to provide high-quality evidence for the collection of future pelvic floor rehabilitation big data.

Adherence

Viewed through the lens of behavioural science, PFMT adherence during pregnancy is a complex process that proves difficult to maintain long-term, even though it is essential for achieving therapeutic benefits [7,8,20,30]. Utilising the COM-B (Capability, Opportunity, Motivation, and Behaviour) framework, Olander et al. categorised the determinants of this behaviour [**Error! Unknown switch argument.**]. The model defines its initial component, ‘Capability’, as the individual’s psychological and physical capacity to execute the behaviour, which implies both adequate knowledge and practical skill. In a study of 633 pregnant women, Hill et al. reported that 41% believed UI during pregnancy is a normal phenomenon [**Error! Unknown switch argument.**]. A survey by Wang et al. of 1243 pregnant women showed that although 52% of women experienced UI symptoms during pregnancy, only 14.8% of pregnant women reported seeking help for UI [**Error! Unknown switch argument.**]. Therefore, the purpose of health education is not only to impart knowledge but more importantly to emphasise health awareness from the source of thought.

The ‘Opportunity’ dimension is constituted by a range of exogenous variables, encompassing both the social context and the physical environment that the woman encounters. The primary reason women who had previously engaged in PFMT ceased training during pregnancy was a concern that PFMT would increase the risk of miscarriage [**Error! Unknown switch argument.**]. Salmon et al. showed that pregnant women prefer to obtain pelvic floor rehabilitation information through consultations with physicians or nurses, but healthcare providers face highly competitive pressures and heavy workloads, which contribute to insufficient attention to this issue [**Error! Unknown switch argument.**]. Although Mason et al. introduced early evening classes to address the scheduling conflicts of working women, this accommodation did not translate into the anticipated participant turnout, with class attendance remaining lower than expected [**Error! Unknown switch argument.**]. Therefore, we can reasonably assume that pregnant women generally lack training time. If pregnant women can schedule classes on weekends, perhaps pregnant women will have more time to focus on PFMT.

The concept of ‘Motivation’ describes the degree of willingness an individual possesses to modify their actions. A study conducted in Nepal demonstrated that pregnant women who received

more support from their families were more willing to exercise and more likely to be compliant [Error! Unknown switch argument.]. Temtanakitpaisan et al. also found that the attitude of pregnant women toward PFMT is a key factor in the decision to engage in PFMT [Error! Unknown switch argument.].

In sum, synthesis of available data reveals that adherence to PFMT can be supported through digital tools, diary-keeping, communal exercise, health literacy programmes, and ongoing support [18,23,33,38,43]. Nevertheless, strategies for ensuring the long-term persistence of both clinical benefits and patient compliance constitute priority areas for further scientific inquiry.

Strength and limitations

This study conducted a comprehensive systematic search on PFMT during pregnancy, making the clinical practice recommendations developed in this study feature relatively high methodological quality and a solid evidence base, which can provide reliable support for the clinical promotion of PFMT during pregnancy. However, when using these recommendations, we need to consider their limitations. In existing trials, PFMT intervention protocols lack uniform standards for intensity, frequency, and duration, which leads to significant heterogeneity among studies and further affects the integration of results and the derivation of unified conclusions. While the literature included in this review covers different countries, the applicability of its findings in practice still requires further verification. At the same time, the impact of some supporting evidence is controversial, which also means that the interpretation of overall study results must be approached with caution.

Conclusions

As stated above, this paper has summarised the best evidence for PFMT during pregnancy from the following seven aspects: overall suggestion, cost-effectiveness, efficacy, safety, timing, protocol, and adherence. It therefore serves as a foundational reference for clinicians and community healthcare providers to guide clinical decision-making and deliver evidence-based care for pregnant women. Some of the evidence integrated in this review is derived from international studies. Therefore, when translating evidence into local clinical practice, it is advisable to make necessary adjustments to the relevant recommendations. Clinicians should take into account local healthcare resource availability, characteristics of the target population, and patient preferences to flexibly select and use the most relevant evidence. PFMT is a key measure to improve women's pelvic floor

health across their entire lifespan and has a clear role in preventing pregnancy-related UI, promoting birth outcomes, and decreasing the risk of PFD. Consequently, it is essential to promote in-depth collaboration between obstetrics and rehabilitation medicine, jointly establish evidence-based standardised PFMT implementation path, and build a sustainable long-term intervention mechanisms. Such measures will systematically enhance the quality of perinatal care and women's well-being.

Declarations

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Conflict of Interest: The authors declare that they have no Conflict of Interest.

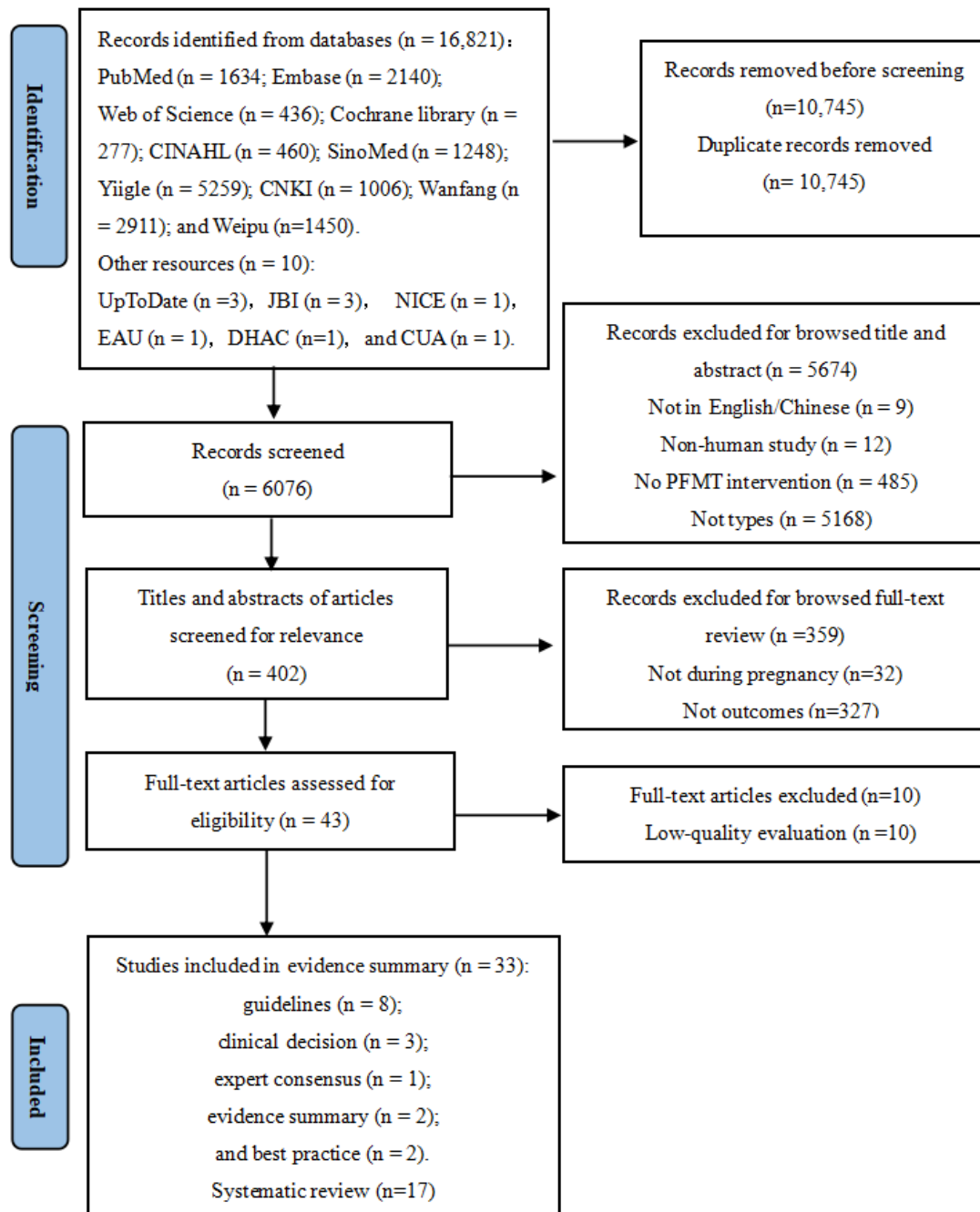


Figure 1. Flow diagram of literature search.

Section and Topic	Item #	Checklist item	Location where item is reported
TITLE			
Title	1	Identify the report as a systematic review.	NA
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	1
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	2
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	2
METHODS			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	2-3
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	2-3
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	2-3,15
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	3
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	3-4,15
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	4
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	16,17
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	3
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	4
Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #8)).	3
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	3
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	4
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	2, 15-18, 21-23
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	NA
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	4
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	NA
Certainty	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	3-4

Section and Topic	Item #	Checklist item	Location where item is reported
assessment			
RESULTS			
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	15
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	15
Study characteristics	17	Cite each included study and present its characteristics.	16-17
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	18-23
Results of individual studies	19	For all outcomes, present, for each study (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	16-17
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	16-23
	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision (e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	16-23
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	19-20
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	15-23
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	19-20
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	19-20
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	4-8
	23b	Discuss any limitations of the evidence included in the review.	8
	23c	Discuss any limitations of the review processes used.	8
	23d	Discuss implications of the results for practice, policy, and future research.	8-9
OTHER INFORMATION			
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	1
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	1
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	NA
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	9
Competing interests	26	Declare any competing interests of review authors.	9
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	NA

From: Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for

reporting systematic reviews. BMJ 2021;372:n71. doi: 10.1136/bmj.n71

Continued Table 5. Summary of best evidence for the current application of pelvic floor muscle training during pregnancy

Evidence items	Evidence content	Level of evidence	Recommended level
Protocol	Dose 1. A basic unsupervised PFMT protocol is 8-12 maximal voluntary contractions, three times a day, with adequate rest in between each contraction. Once a baseline sustained contraction is determined, the length of the sustained contraction should be increased gradually (to a maximum of 10 seconds) or do 150 to 200 repetitions daily for three months or longer, but no less than six weeks (ideally lifelong). ^[6,8,21,27,28,29,30,40] 2. There is no standard PFMT protocol, and there are variations in each protocol's training patterns. The protocol should include increasing pelvic floor muscle strength (type II muscle fibres), increasing endurance (type I muscle fibres), and coordinating muscle activity (knack) and respect the FITT principle, including exercise frequency (F) exercise intensity (I), exercise time (T), and exercise type (T). ^[2,6,7,8,9,28]	1a 5b	A A
	Position 1. Can be performed in any position (supine, sitting, standing), according to how you feel while practicing, and the supine position should not exceed 5 minutes after 13 weeks of pregnancy. ^[6,8,27,28,29]	5b	A
	Place 1. PFMT is not limited by time and place; once you have mastered it, incorporate it into your daily life. ^[27,28]	5b	A
Adherence	1. Use of mobile applications, training diaries, group training, health education, and follow-up can promote training continuity and adherence, but there is a lack of clinical data on adherence. ^[18,23,33,38,43] 2. Patients and their families participate together in discussions to develop a training protocol. ^[30] 3. The accuracy and adherence to training are important factors affecting training results. ^[7,8,20,30]	1a 5c 1a	A B A

Note: Absolute contraindications^a: premature rupture of membranes; threatened abortion; cervical insufficiency; persistent vaginal bleeding; placenta previa after 28 weeks of gestation; abnormal foetal movement; persistent pain or regular and painful contractions after exercise; uncontrolled type I diabetes mellitus, thyroid disease, blood and cerebrovascular diseases, respiratory diseases and other systemic diseases; arrhythmia, dizziness, or syncope; oedema or pain in the leg (excluding phlebitis); poor balance, difficulty in walking or difficulty maintaining balance. Relative contraindications^b: previous history of spontaneous abortion; premature birth; foetal growth restriction; after 28 weeks of gestation with twins; anaemia (haemoglobin < 100g/L); malnutrition or eating disorders; mild or moderate cardiovascular, cerebrovascular, and respiratory diseases; overheating during exercise. Stop signals^c: amniotic fluid leak or other loss of vaginal fluid including rupture of the membranes; pain or oedema of the calf (suspected deep vein thrombosis); chest pain; dizziness; syncope, or feeling of faintness that does not resolve at rest; headaches; muscle weakness affecting balance; regular painful uterine contractions; shortness of breath before exertion or persisting, excessive, and not resolving at rest; vaginal bleeding

Table 5. Summary of best evidence for the current application of pelvic floor muscle training during pregnancy

Evidence items	Evidence content	Level of evidence	Recommended level
Overall suggestion	1. It was particularly important to raise awareness in maternity services; this is when symptoms can first occur, when risk factors can be identified, and when prevention strategies can be started. ^[18,28,30] 2. Establish a specialized outpatient clinic for PFD care interventions and form a multidisciplinary team to jointly develop a PFMT intervention protocol. ^[18,29,30]	1a 5c	A A
	3. In the absence of contraindications, pregnant women are encouraged to engage in PFMT. ^[2,9,18,20,22,24,25,28,29,38,42]	1a	A
cost-effectiveness	1. PFMT during pregnancy for preventing postnatal urinary incontinence is clinically effective and cost-effective. ^[18,21] 2. Group-based interventions may offer a more cost-effective way to implement PFMT. ^[29,31,43]	1b 1b	A B
	1. There is no extra benefit of combining PFMT with biofeedback. ^[19,29] 2. Higher-intensity, supervised treatment regimens confer greater benefit in women receiving PFMT. ^[28,29,30] 3. PFMT in the antenatal period is associated with a reduced risk of urinary incontinence in the short term postnatally, but a long-term benefit has not been established (i.e. >6 months postpartum). ^[6,7,9,18,19,20,22,23,25,26,29,34,35,38,42,44]	1b 1a 1a	A A A
Efficacy	4. PFMT significantly shortened the first and second stage of labour in the primigravida, but it is not possible to recommend PFMT as an intervention to prevent perineal laceration. ^[8,9,32,33]	1a	A
	5. The efficacy of training interventions on natural childbirth, ranked from highest to lowest, is as follows: pelvic floor muscle training, exercise classes, gymnastics, aerobic exercise, birth balls, and yoga. ^[37]	1b	A
	6. PFMT is beneficial for improving female sexual function, thereby enhancing sexual health and quality of life. ^[34,43]	1b	A
	7. The evidence showed a benefit from pelvic floor muscle training for prolapse that does not extend more than 1 cm beyond the hymen, and the training period lasts for 16 weeks. ^[6,18,27]	1a	B
	8. Few data exist on faecal incontinence, it was uncertain whether PFMT reduced incontinence in the late postnatal period compared to usual care (very low-quality evidence). ^[18,27,35,38]	1b	B
	9. PFMT improves pain and function in pregnant and non-pregnant women with lumbopelvic pain, respectively (low and very low-quality evidence). ^[41]	1c	B
Safety	1. Pelvic floor muscle training exhibits very few adverse events and no serious adverse events. ^[19,27,42] 2. Absolute contraindications ^a ; Relative contraindications ^b . ^[20,24,28,30,37] 3. Stop signals ^c . ^[24,37]	1a 5b 5c	A A A
	1. The start time varies from 11 to 28 weeks. In China, the training usually starts from the 28th week of gestation. From week 20 of pregnancy, for pregnant women who have a first-degree relative with pelvic floor dysfunction. ^[9,18,28,32,33]	5b	A
	2. The start timing of training on natural childbirth, from highest to lowest, is as follows: >24 weeks, 13-24 weeks, ≤12 weeks. ^[37]	1b	B

meta-analysis was performed did the review authors use appropriate methods for statistical combination of results; ⑫ If meta-analysis was performed, did the review authors assess the potential impact of RoB in individual studies on the results of the meta-analysis or other evidence synthesis; ⑬ Did the review authors account for RoB in individual studies when interpreting/ discussing the results of the review; ⑭ Did the review authors provide a satisfactory explanation for, and discussion of, any heterogeneity observed in the results of the review; ⑮ If they performed quantitative synthesis did the review authors carry out an adequate investigation of publication bias (small study bias) and discuss its likely impact on the results of the review; ⑯ Did the review authors report any potential sources of conflict of interest, including any funding they received for conducting the review.

Table 4. Methodologic quality evaluation results of systematic reviews

systematic review	Evaluation entry																Overall quality
	①	②	③	④	⑤	⑥	⑦	⑧	⑨	⑩	⑪	⑫	⑬	⑭	⑮	⑯	
Zhang et al. ^[9]	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	Partial Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	High
Brennen et al. ^[31]	Yes	Yes	Yes	Partial Yes	No	No	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Moderate
Yang et al. ^[32]	Yes	Yes	Yes	Partial Yes	No	Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	Moderate
Wang et al. ^[33]	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	Partial Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	No	Moderate
Wang et al. ^[34]	Yes	Partial Yes	Yes	Partial Yes	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	High
Zhang et al. ^[35]	Yes	Partial Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	High
Al et al. ^[36]	Yes	Partial Yes	Yes	Partial Yes	Yes	Yes	Yes	Partial Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	No	Moderate
Ji et al. ^[37]	Yes	Partial Yes	Yes	Partial Yes	No	Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	High
Woodley et al. ^[38]	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	High
Jorge et al. ^[39]	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	High
Lopes et al. ^[40]	Yes	Partial Yes	Yes	Partial Yes	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	High
Vesentini et al. ^[41]	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	High
Davenport et al. ^[2]	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	High
Lu et al. ^[42]	No	Partial Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	High
Zahra et al. ^[43]	Yes	Partial Yes	Yes	Partial Yes	Yes	Yes	Yes	Partial Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	High
Santos et al. ^[44]	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Moderate
Yang et al. ^[45]	Yes	Yes	Yes	Partial Yes	Yes	Yes	Yes	Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	High

Note: ①Did the research questions and inclusion criteria for the review include the components of PICO; ②Did the report of the review contain an explicit statement that the review methods were established prior to the conduct of the review and did the report justify any significant deviations from the protocol; ③Did the review authors explain their selection of the study designs for inclusion in the review; ④Did the review authors use a comprehensive literature search strategy; ⑤Did the review authors perform study selection in duplicate; ⑥Did the review authors perform data extraction in duplicate; ⑦Did the review authors provide a list of excluded studies and justify the exclusions; ⑧Did the review authors describe the included studies in adequate detail; ⑨Did the review authors use a satisfactory technique for assessing the risk of bias (RoB) in individual studies that were included in the review; ⑩Did the review authors report on the sources of funding for the studies included in the review; ⑪If

Table 3. Methodologic quality evaluation of evidence summaries and best practices

Evidence summary	Evaluation entry										Overall quality
	①	②	③	④	⑤	⑥	⑦	⑧	⑨	⑩	
Zhang et al. ^[29]	Yes	Yes	Yes	Not completely	Yes	Yes	Yes	Yes	Yes	Yes	Medium
Yang et al. ^[30]	Yes	Yes	Yes	Not completely	Yes	Yes	Yes	Yes	Yes	Yes	Medium
Viñaspre et al. ^[7]	Yes	Yes	Yes	Not completely	No	Not completely	Yes	Yes	Yes	Yes	Medium
Xing et al. ^[8]	Yes	Yes	Yes	Not completely	Yes	Not completely	Yes	Yes	Yes	Yes	Medium

Note: ①Is the summary specific in scope and application? ②Is the authorship of the summary transparent? ③Are the reviewer(s)/editor (s) of the summary transparent? ④Are the search methods transparent and comprehensive? ⑤Is the evidence graded and is the system transparent and translatable? ⑥Are the recommendations clear? ⑦Are the recommendations appropriately cited? ⑧Are the recommendations current? ⑨Is the summary free of possible bias? ⑩Can this summary be applied to your patient(s).

Table 2. Methodologic quality evaluation results of the guidelines

Included literature	Percentage of field standardization %						≥60% field number (n)	≥30% field number (n)	Level
	Scopes and object	Participant	Rigour of the guidelines	Clarity of guidelines	Application of guidelines	Independence of the guide			
NICE ^[18]	97-22%	72-22%	89-58%	80-55%	83-33%	79-16%	6	0	A
Okeahialam et al. ^[6]	88-88%	66-66%	29-16%	66-66%	62-50%	16-66%	4	0	B
Harding et al. ^[19]	94-44%	83-33%	95-83%	100%	60-41%	100%	6	0	A
Mottola et al. ^[20]	100%	100%	83-33%	94-44%	54-16%	50-00%	4	0	B
Brown et al. ^[22]	100%	66-66%	87-50%	77-77%	89-58%	100%	6	0	A
Carlson et al. ^[23]	100%	52-77%	79-16%	88-88%	20-83%	16-66%	3	2	B
Boisseau et al. ^[24]	100%	66-66%	50%	72-22%	20-83%	50%	3	1	B
Urogynocology Subgroup et al. ^[21]	63-88%	61-11%	10-41%	66-66%	58-33%	16-66%	3	2	B

Note: Standardization percentage of each field=(obtained score-minimum possible score)/(maximum possible score-minimum possible score) × 100%; Recommendation level: if the standardized percentage of 6 fields is > 60%, it is highly recommended (level A); if > 3 areas have a standardized percentage > 30% and < 60% are recommended (level B); if there are ≥ 3 areas with a standardized percentage < 30%, it is not recommended (level C).

Continued Table 1. General information of the included literature

Included literature	Year	Country	Literature sources	Type of evidence	Topic of the literature
Yang et al. ^[32]	2017	China	Yiigle	Systematic review	Effect of continuous pelvic floor muscle training during pregnancy on delivery outcome: a Meta-analysis
Wang et al. ^[33]	2023	China	CNKI	Systematic review	Intervention time of pelvic floor muscle training for prevention or treatment of postpartum urinary incontinence: a Meta-analysis
Wang et al. ^[34]	2019	China	CNKI	Systematic review	Meta-analysis of effect of pelvic floor muscle training during pregnancy to prevent or treat urinary incontinence in primipara
Zhang et al. ^[35]	2024	China	CNKI	Systematic review	Meta-analysis of the effectiveness of pelvic floor muscle training during pregnancy in preventing or treating urinary and faecal incontinence
Al et al. ^[36]	2023	China	CNKI	Systematic review	Effect of pelvic floor muscle training on female sexual function: a systematic review and Meta-analysis
Ji et al. ^[37]	2022	China	Yiigle	Systematic review	Influence of Physical Exercise Interventions during Pregnancy on Natural Childbirth: a Meta-analysis
Woodley et al. ^[38]	2020	New Zealand	Cochrane Library	Systematic review	Pelvic floor muscle training for preventing and treating urinary and faecal incontinence in antenatal and postnatal women
Jorge et al. ^[39]	2024	Brazil	Embase	Systematic review	Pelvic floor muscle training as treatment for female sexual dysfunction: a systematic review and meta-analysis
Lopes et al. ^[40]	2022	Brazil	PubMed	Systematic review	Can pelvic floor muscle training prevent perineal laceration? A systematic review and meta-analysis
Vesentini et al. ^[41]	2020	Brazil	PubMed	Systematic review	Pelvic floor muscle training for women with lumbopelvic pain: A systematic review and meta-analysis
Davenport et al. ^[2]	2018	Canada	PubMed	Systematic review	Prenatal exercise (including but not limited to pelvic floor muscle training) and urinary incontinence during and following pregnancy: a systematic review and meta-analysis
Lu et al. ^[42]	2021	China	PubMed	Systematic review	Meta-analysis of Perinatal Pelvic Floor Muscle Training on Urinary Incontinence
Zahra et al. ^[43]	2019	Iran	Embase	Systematic review	Effect of pelvic floor muscle training on postpartum sexual function and quality of life: A systematic review and meta-analysis of clinical trials
Santos et al. ^[44]	2023	Brazil	Embase	Systematic review	Effectiveness of group aerobic and/or resistance exercise programs associated with pelvic floor muscle training during prenatal care for the prevention and treatment of urinary incontinence: A systematic review
Yang et al. ^[45]	2022	China	Embase	Systematic review	The effectiveness of group-based pelvic floor muscle training in preventing and treating urinary incontinence for antenatal and postnatal women: a systematic review

Note: NICE: National Institute for Health and Clinical Excellence; EAU: European Association of Urology; DHAC: Department of Health and Aged Care; CUA: Canadian Urological Association; JBI: Joanna Briggs Institute; CNKI: China National Knowledge Infrastructure

Table 1. General information of the included literature

Included literature	Year	Country	Literature sources	Type of evidence	Topic of the literature
NICE ^[18]	2021	Britain	NICE	Guideline	Pelvic floor dysfunction: prevention and non-surgical management
Okeahialam et al. ^[6]	2022	Britain	Embase	Guideline	Pelvic floor muscle training: A practical guide
Harding et al. ^[19]	2025	European	EAU	Guideline	EAU Guidelines on management of Non-Neurogenic Female Lower Urinary Tract Symptoms
Mottola et al. ^[20]	2018	Canada	PubMed	Guideline	2019 Canadian guideline for physical activity throughout pregnancy
Brown et al. ^[22]	2020	Australia	DHAC	Guideline	Evidence-based physical activity guidelines for pregnant women
Carlson et al. ^[23]	2024	Canada	CUA	Guideline	2024 Canadian Urological Association guideline: Female stress urinary incontinence
Boisseau et al. ^[24]	2022	France	PubMed	Guideline	Physical Activity During the Perinatal Period: Guidelines for Interventions During the Perinatal Period from the French National College of Midwives
Urogynecology Subgroup ^[21]	2017	China	Yiigle	Guideline	Update of guideline on the diagnosis and treatment of female stress urinary incontinence (2017)
Artal et al. ^[25]	2025	America	Up to Date	Clinical decision	Exercise during pregnancy and the postpartum period
Handa et al. ^[26]	2025	America	Up to Date	Clinical decision	Effect of pregnancy and childbirth on urinary incontinence and pelvic organ prolapse
Brubaker et al. ^[27]	2025	America	Up to Date	Clinical decision	Patient education: Pelvic floor muscle exercises (Beyond the basics)
Sun et al. ^[28]	2024	China	CNKI	Expert consensus	Expert consensus on primary prevention strategy of pelvic floor dysfunction based on pregnancy (2024 edition)
Zhang et al. ^[29]	2024	China	Yiigle	Evidence summary	Summary of the best evidence for pelvic floor muscle training in the prevention and treatment of postpartum urinary incontinence
Yang et al. ^[30]	2023	China	Yiigle	Evidence summary	Summary of the best evidence for prevention and management of stress urinary incontinence in pregnant and postpartum patients
Viñaspre et al. ^[7]	2021	Spain	JBI	Best practice	Training women's pelvic floor muscles during pregnancy and postpartum at primary health centres: a best practice implementation project
Xing et al. ^[8]	2017	China	JBI	Best practice	Pelvic floor muscle training for the prevention of urinary incontinence in antenatal and postnatal women: a best practice implementation project
Zhang et al. ^[9]	2024	Spain	PubMed	Systematic review	Influence of pelvic floor muscle training alone or as part of a general physical activity program during pregnancy on urinary incontinence, episiotomy and third- or fourth-degree perineal tear: Systematic review and meta-analysis of randomized clinical trials
Brennen et al. ^[31]	2021	Australia	PubMed	Systematic review	Group-based pelvic floor muscle training for all women during pregnancy is more cost-effective than postnatal training for women with urinary incontinence: cost-effectiveness analysis of a systematic review

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