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# Assessment of physical fitness during pregnancy: validity and reliability of fitness tests, and relationship with maternal and neonatal health – a systematic review

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## ABSTRACT

**Objectives** To systematically review studies evaluating one or more components of physical fitness (PF) in pregnant women, to answer two research questions: (1) What tests have been employed to assess PF in pregnant women? and (2) What is the validity and reliability of these tests and their relationship with maternal and neonatal health?

**Design** A systematic review.

**Data sources** PubMed and Web of Science.

**Eligibility criteria** Original English or Spanish full-text articles in a group of healthy pregnant women which at least one component of PF was assessed (field based or laboratory tests).

**Results** A total of 149 articles containing a sum of 191 fitness tests were included. Among the 191 fitness tests, 99 (ie, 52%) assessed cardiorespiratory fitness through 75 different protocols, 28 (15%) assessed muscular fitness through 16 different protocols, 14 (7%) assessed flexibility through 13 different protocols, 45 (24%) assessed balance through 40 different protocols, 2 assessed speed with the same protocol and 3 were multidimensional tests using one protocol. A total of 19 articles with 23 tests (13%) assessed either validity (n=4), reliability (n=6) or the relationship of PF with maternal and neonatal health (n=16).

**Conclusion** Physical fitness has been assessed through a wide variety of protocols, mostly lacking validity and reliability data, and no consensus exists on the most suitable fitness tests to be performed during pregnancy.

**PROSPERO registration number** CRD42018117554.

### WHAT IS ALREADY KNOWNWHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ The assessment of physical fitness during pregnancy requires special considerations to preserve fetal and maternal health.
⇒ Although physical fitness during pregnancy has been assessed inconsistently across studies, these tests have not been systematically compiled to date.
⇒ The validity and reliability of the variety of tests used to assess physical fitness during pregnancy has not been comprehensively reviewed.

### WHAT THIS STUDY ADDS

⇒ During pregnancy, physical fitness including cardiorespiratory fitness, muscular strength, flexibility and balance have been assessed inconsistently, using a wide variety of protocols.
⇒ Most of the tests used to assess physical fitness during pregnancy lack validity and reliability data.
⇒ Higher physical fitness might be associated with better maternal and neonatal health, although further research is needed.

### HOW THIS STUDY MAY AFFECT RESEARCH, PRACTICE AND POLICY

⇒ The extent to which the data derived from current physical fitness tests during pregnancy is valid and reliable is still unclear and, therefore, should be interpreted with caution.
⇒ Developing a battery of fitness tests to assess the different fitness components during pregnancy must be set as a priority for relevant institutions.
⇒ An expert consensus to develop a battery of physical fitness tests is recommended.

## BACKGROUND

Physical fitness (PF) has been defined as the ability to carry out daily tasks with vigour and alertness, without undue fatigue and with ample energy to enjoy leisure-time pursuits and meet unforeseen emergencies.[1 2] PF is considered a powerful marker of health that is associated with a lower risk of cardiovascular events, cancer and all-cause mortality in all ages.[3–7] In pregnant individuals, some studies have recently highlighted the potential impact of PF on maternal and fetal health.[8–15] Low PF levels are associated with low infant birth weight,[8] increased risk of gestational diabetes mellitus,[9 10] poor postpartum recovery[11] and

worse delivery outcomes.[12 13] Moreover, the anatomical, biomechanical, physiological and psychological changes during the pregnancy might compromise PF levels.[16–18] Consequently, it is of clinical and public health interest to assess PF during the pregnancy, and to understand which available tests are best to assess PF during this critical period of life.

Two categories of PF components have been defined as follows: (1) health-related components (cardiorespiratory fitness (CRF), muscular fitness, muscular endurance and flexibility) and (2) skill-related components (ability, coordination, balance, power, reaction time and speed).[1 2] These PF components can be assessed subjectively through questionnaires,[15] objectively and accurately through laboratory tests and efficiently, economically and easily through field-based tests. During the pregnancy, a wide variety of fitness tests have been used to assess PF, although a compilation of these tests has not been published to date. Compiling all fitness tests performed in pregnant women would help practitioners to select the most useful test according to their purpose. It is also important to note that, although laboratory tests are generally the gold standard for assessing PF, these tests are not accessible to everyone because they need sophisticated and expensive equipment, and it is not possible to evaluate a relatively large sample in a short period of time. As an alternative, a number of field tests exist that provide an opportunity to assess PF in a more accessible way.[2] However, there is no consensus on which fitness tests should be used to assess PF in pregnant individuals, and the validity and reliability of many of the tests used to assess PF during the pregnancy are unknown.[19]

Since the assessment of PF in pregnancy requires special consideration to preserve fetal and maternal health,[18 20 21] understanding which fitness tests are valid, reliable, and associated with maternal and neonatal health outcomes, would provide a framework for improving PF assessment during pregnancy and also for improving exercise prescription in this population.

The aims of this systematic review were to: (1) describe which fitness tests have been used to evaluate PF in pregnant individuals; and (2) to evaluate the validity and reliability of the fitness tests, and their relationship with maternal and neonatal health.

## METHODS

### Registration and review guidelines and checklist

This systematic review was prospectively registered at PROSPERO (CRD42018117554; available at http://www.t.ly/fS6a). In addition, the review followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines[22] and the PRISMA checklist[23] is included as online supplemental material 1, table S1. (1) .

### Search strategy

Articles were searched by two independent reviewers from two major databases, MEDLINE (PubMed) and the Web of Science (WOS) from inception to January 2021. For the search strategy undertaken in PubMed Medical Subject Heading, (MeSH) terms were used. All terms were combined using the connector OR for similar criteria the connector 'AND' was used to combine population group (ie, pregnant women), to delimit date of publication ('0001/01/01'(PDat): '2021/01/15'(PDat)), to include full text papers, and to include studies performed in humans.

A similar search strategy and term combination was undertaken in the WoS (online supplemental material 2, table S2), although MeSH terms and its appropriate terms connection were not used as they are exclusive for PubMed. The complete search strategy and further details are presented in online supplemental material 2, tables S1 and S2.

### Inclusion criteria

The inclusion criteria were as follows: (1) healthy pregnant individuals (no restriction regarding gestational week); (2) at least one component of PF assessed either through field based or laboratory tests; (3) access to full text; (4) only one original article from the same study/project using the same test were included and (5) text in English or Spanish.

### Quality assessment of the articles

To assess quality of the articles included in aim 2, three quality scores were applied. To assess validity and reliability, authors adapted two quality scores ad hoc previously used in two different systematic reviews following the same goal as the present review, however, undertaken in different populations.[24 25] To assess the association of PF with health-related outcomes the Effective Public Health Practice Project was used.[26] All procedures are comprehensively described in online supplemental material 3, tables S3–S5.

### Process and data extraction

After checking title and abstract, only the studies meeting all inclusion criteria were introduced in a reference manager software (Mendeley). In the event of disagreement between the two independent reviewers concerning the inclusion/exclusion of an article, a consensus was reached (there was no need of a third person). The snowball strategy was also used. Information including reference, age, sample size and fitness test description are summarised in online supplemental material 5, table S6.

## RESULTS

A comprehensive PRISMA flow diagram is presented in figure 1.

### Overall results, quality assessment and gestational week

The search identified 2617 studies, of which 149 were included (figure 1). These articles contained 191 fitness tests, using 149 different protocols that were included for Aim 1. A summary of the number of articles that

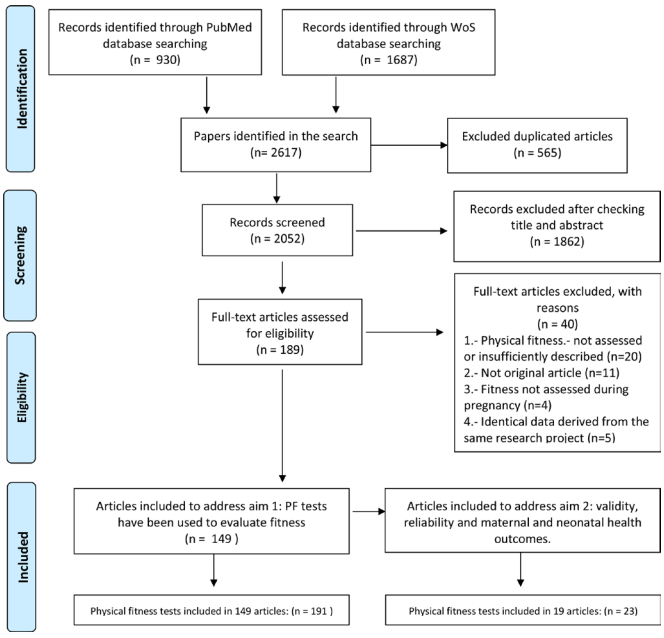


**Figure 1** Flow chart of the literature search and paper selection process.

assessed PF during the pregnancy and the protocols used for its assessment is presented in figure 2. This has been organised based on each of the different PF components assessed in those articles. Moreover, a comprehensive diagram of the fitness tests and the different protocols performed to date, organised by PF component, is presented in figure 3.

Regarding aim 1, 99 tests (including 75 different protocols) were used to assess CRF,[8 12 13 18 27–108] 28 (including 16 different protocols) to assess muscular fitness,[8 12 13 61 86 109–122] 14 (including 13 different protocols) to assess flexibility,[12 13 110 114 123–127] 45 tests (including 40 different protocols) to assess balance,[110 116 128–167] 2 tests using the same protocol to assess speed[168 169] and 3 tests using the same protocol were multidimensional.[168–170] No results were found for other PF components such as agility or coordination.

Regarding aim 2, a total of 19 articles (13% of the total number of articles included) assessed at least validity (n=3) and reliability (n=4) of fitness tests. These articles are summarised in table 1. Of the three articles[74 75 169] that assessed validity, two articles were classified as low quality[74 169] and one as high quality.[75] Of the four articles that assessed reliability criteria, three were considered high quality[74 117 168] and one low quality.[121] The relationship of PF with maternal and neonatal health outcomes (n=16 tests) are summarised in table 2. Of these 16 tests, 11 were classified as very low quality[13 57 68 95 108 111 126 157 158] and 5 were classified as low quality.[8 63 115 128 170]

The gestational week at PF assessment ranged from 8 to 41 across articles. Some articles assessed PF at different time points throughout pregnancy; therefore, we divided pregnancy into two stages. Early pregnancy (ie, from week 0 to week 20 of gestation) and late pregnancy (ie, from week 21 to week 40). Using this approach, 11 articles (7%) were performed in early pregnancy; 57 articles (38%) were performed in late pregnancy; 55 articles (37%) were performed several times (ie, range 2–5 times) throughout pregnancy; 7 (5%) articles specified a range of weeks that included early to late pregnancy; 14 articles (9%) reported only the trimester without specifying gestational week; 4 articles (3%) provided no information and 1 article (1%) assessed PF on the day of labour.

## Aim 1: fitness tests used to evaluate PF in pregnant women

### Cardiorespiratory fitness

We identified 99 tests assessing CRF, of which 61 (62%) were performed on a cycle ergometer, 25 (25%) on a treadmill, 10 (10%) on a track and 3 (3%) used step

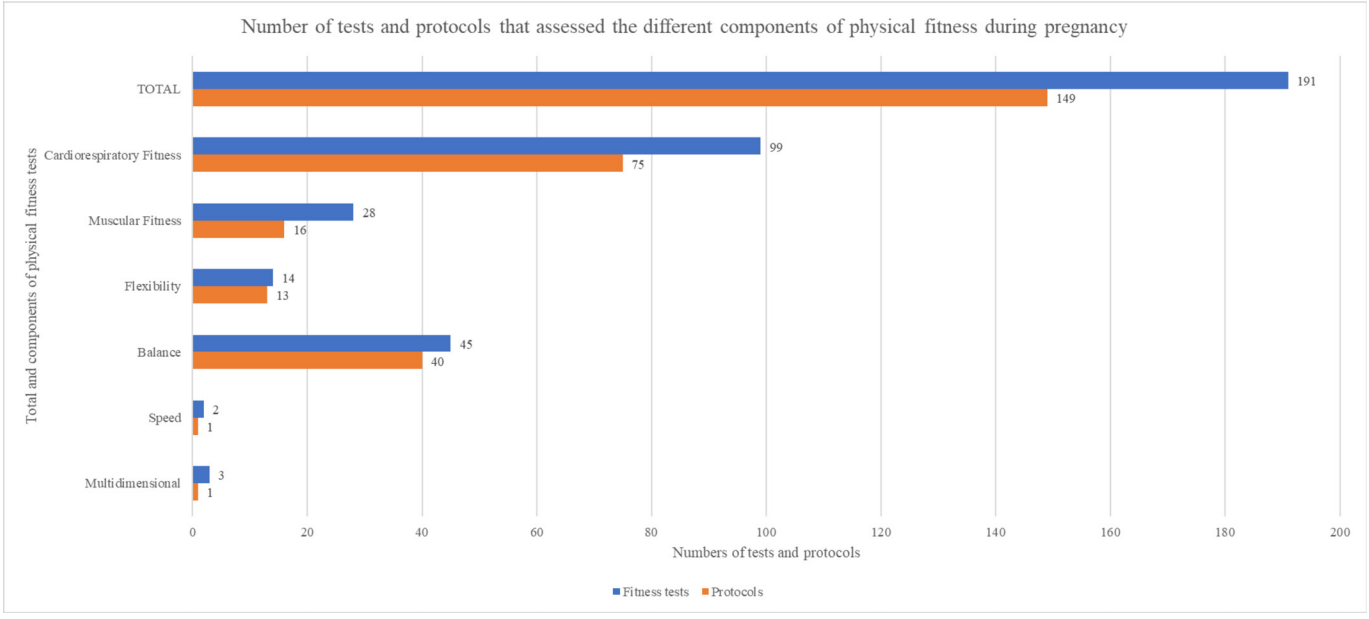


**Figure 2** Number of tests and protocols that assessed the different components of physical fitness during pregnancy.

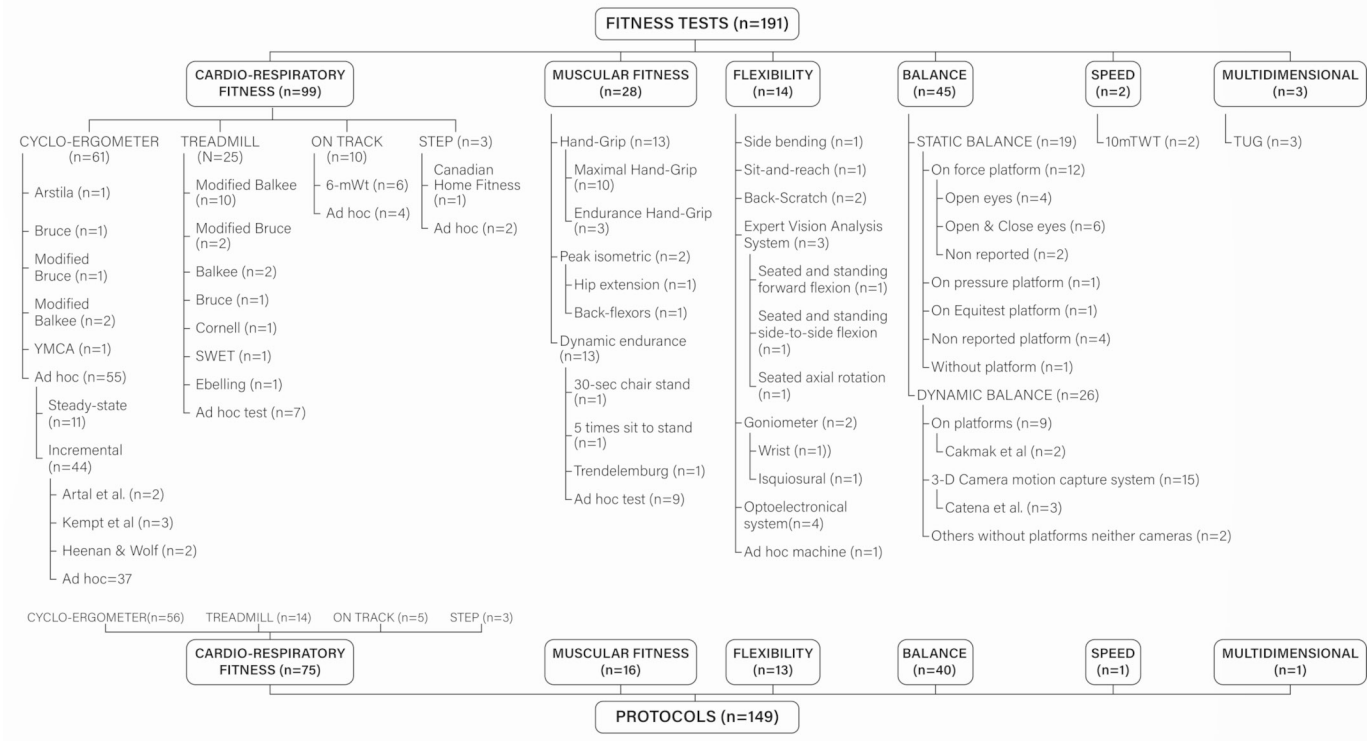


**Figure 3** Diagram of the fitness tests and the different protocols organised by PF component. PF, physical fitness.

protocols (figure 3). Of the 99 tests, a total of 75 corresponded to different protocols. For instance, there were 56 different protocols using a cycle ergometer, distributed as follows: only one article used the Arstila test[68]; one used the Bruce Protocol at 75% $HR_{max}$[27]; one applied the Modified Bruce ramp protocol at anaerobic threshold[104]; two employed the Modified Balke protocol at 70% $HR_{max}$[34 41]; one used a YMCA protocol;[106] The remaining of articles (n=55) used ad hoc tests (ie, specifically designed for the purpose of the investigation); 11 of which[32 37 38 41 45 57 64 79 107] used steady-state tests and 44[28–31 33 35 36 39–41 43 44 46–56 59–63 65–69 90 100–106 108 171–173] used incremental tests. When analysing the type of test based on intensity, we found that 13 tests were maximal tests,[31 43 44 47–49 59 60 67 103–105 171] 37 submaximal tests[29 30 35–40 42 45 46 50–52 54–57 62–66 68 69 79 90 100–102 106 108 172] and 3 used mixed tests[28 33 41] containing submaximal and maximal stages within the same protocol.

There were 25 treadmill tests that used 14 different protocols, distributed as follows: the Modified Balke protocol was used in 10 articles[8 31 71 73 75–78 82 96]; the Modified Bruce protocol in 2 articles[13 97] and the traditional Balke protocol used twice in the same article[70]; the traditional Bruce protocol,[98] the Cornell protocol,[74] the SWET protocol and the Ebbelling single-stage protocol[18] were each used in one article. There were seven ad hoc tests of which two were steady-state,[38 81] and five were incremental tests.[72 73 80 83 90] According to intensity, three were maximal tests[73 80 81] and four submaximal tests.[38 72 83 90]

Of the 10 tests performed on a track, 6 articles used the 6 min walk test protocol,[84 85 87–89 92] and 4 were ad hoc tests (ie, maximal and 4 were submaximal). With regard to the three step tests, one Canadian Home Fitness test[93] was used and two ad hoc incremental submaximal tests were used.[94 95]

## Muscular fitness

A total of 28 tests (ie, 14% of all included articles) that included 16 different protocols assessed muscular fitness, of which 10 performed maximal hand-grip strength tests,[8 12 13 86 109–115] performed endurance hand-grip tests, 2 for 3 min[118 120] and 1 for a 2 min period[119] (figure 3). In two of the articles conducting an endurance hand-grip test,[118 119] a hand-grip sphygmomanometer was used instead of dynamometry. On the other hand, one used a hand-held dynamometer fixed to a chair to assess quadriceps strength[116] and one used a toe-grip dynamometer.[116] Moreover, two ad hoc isometric tests were used to assess maximal voluntary hip extension and back flexors endurance in the same article.[174] Finally, 13 dynamic endurance tests were found, 9 were listed as ad hoc tests[12 112 122] and another 3 (30 s Chair Stand Test, 5 Times Sit to Stand test, Trendelenburg's test) were classified as 'other' dynamic tests.[13 112 117]

## Flexibility

Our search identified 14 (7%) tests that assessed flexibility using 13 different protocols, including the side bending test,[125] the sit-and-reach test,[12] the back-scratch test (twice),[13 110] the motion analysis (ie, including three different tests such as the seated and standing forward flexion, seated and standing side to side flexion and seated axial rotation)[123] and an optoelectrical system (ie, performing four different tests).[127] Goniometry

**Table 1** Overview of studies that assessed the validity and/or reliability of fitness tests during pregnancy

| Reference (authors, year) | Validity | Reliability | Capacity evaluated, short test description and maternal and neonatal health outcomes or statistical results | Quality score |
|---|---|---|---|---|
| Cardio-respiratory fitness | | | | |
| Yeo *et al* (2005)[74] | Yes | Yes | Cornell Protocol on treadmill platform. Validity: Bland-Altman plots. The mean difference was 4.4±3.6 mL/kg/min. Data indicated that VO2000 overestimates $VO_2$ by an average of 4.4 mL/kg/min compared with CPX/D. Pearson correlation coefficient between the average and difference of paired measurements was close but not significant (r=0.48; p>0.01). Reliability: Paired t test (t (45)=3.9, p<0.001). Linear regression: y=0.96X-1.6; 95% CI for the slope: 0.94 to 1.1; $R^2$=0.91, p<0.001 | 4–8 |
| Mottola *et al* (2006)[75] | Yes | No | Modified Balke protocol on treadmill platform. Validity: Pearson Correlation: $R^2$=0.72, $R^2$ adjusted=0.71 and SEE=2.7 (The prediction equation was compared with cross validation (n=39; p=0.78). | 5 |
| Muscular fitness | | | | |
| Gutke *et al* (2008)[121] | No | Yes | Maximal voluntary isometric hip extension Reliability: Spearman's r and Intercorrelation coefficient (ICC). Right leg: r=0.82; ICC=0.87. Left leg: r=0.88; ICC=0.85 (both p value no reported). | 3 |
| Yenişehir *et al* (2020)[117] | No | Yes | Five Times Sit to Stand Test (5TSS) Reliability: Inter-rater reliability of 5TSS was excellent for subjects with and without pregnancy-related pelvic girdle pain (ICC ¼ 0.999, 95% CI ¼ 0.999 to 1.000; ICC ¼ 0.999, 95% CI ¼ 0.999 to 0.999, respectively). Test–retest reliability of 5TSS was also very high for subjects with and without PGP (ICC ¼ 0.986, 95% CI ¼ 0.959 to 0.995: ICC ¼ 0.828, 95% CI ¼ 0.632 to 0.920, respectively). | 5–7 |
| Flexibility | | | | |
| Lindgren and Kristiansson (2014)[126] | No | Yes | Ad hoc passive abduction of the left fourth finger. Reliability: Intraindividual coefficient of variance. (1) Between the first and second measurement=0.077; (2) Between the second and third=0.070 and between the third and fourth=0.071. | 2 |
| Speed | | | | |
| Evensen *et al* (2015)[168] | No | Yes | Ten metres Timed walk Test Reliability: ICC from a one-way random effects model and reporting the 95% CI. Coefficients for test–retest reliability for 10mTWT: (ICC=0.74; 95% CI=0.42 to 0.90; SEM=0.17 m/s; $MDC_{95}$=0.47 m/s) Coefficients for intertester reliability 10mTWT: (ICC=0.94; 95% CI=0.82 to 0.98; SEM=0.09 m/s; $MDC_{95}$=0.25 m/s). | 8 |
| Evensen *et al* (2016)[169] | Yes | No | Ten metres Timed walk Test Validity: Spearman correlation coefficient. Between the 10mTWT and ASLR (r=−0.65, p=0.003). Between the 10mTWT and PGQ (r=−0.25 to −0.56). | 3 |
| Multidimensional | | | | |
| Evensen *et al* (2015)[168] | No | Yes | Timed Up and Go Test (TUG) Reliability: ICC from a one-way random effects model and reporting the 95% CI. Coefficients for test–retest reliability TUG: (ICC=0.88; 95% CI 0.70 to 0.95; SEM=0.42 s; $MDC_{95}$=1.16 s.) Coefficients for intertester reliability TUG: (ICC=0.95; 95% CI 0.84 to 0.98; SEM=0.36 m/s; $MDC_{95}$=1.00 m/s). | 8 |
| Evensen *et al* (2016)[169] | Yes | No | TUG Validity: Spearman correlation coefficient. Between the TUG and ASLR ($r_s$=0.73, p=0.001). Between the TUG and ASLR ($r_s$=0.73, p=0.001). Between the TUG and PGQ ($r_s$=0.41 to 0.52). | 3 |

ASLR, Active Straight Leg Raised; MDC, minimal detectable change; 10mTWT, Ten-metre Timed Walk Test; PGP, Pelvic girdle pain; SEM, SE of measurement.

was used in two different articles to measure hamstring flexibility,[114] wrist flexion-extension and medial lateral deviation.[124] Only one article used an ad hoc machine to test passive abduction of the left fourth finger.[126]

## Balance

We identified 45 (24%) articles assessing balance of which 19 analysed static balance and 26 used dynamic balance with 40 different protocols. With regard to static balance, 18 were laboratory tests of which 12 assessed balance through stabilometry tests on a force platform,[129 131 132 138 149 158–160 162–164 167] one on a pressure platform[163] and another on an Equitest platform.[165] Four articles did not mention the type of platform used.[117 132 133 175] Regarding protocols, all articles conducted the tests with participants standing with bipedal support. However, standing position varied between articles. Ten articles maintained a standing posture with feet separated,[116 131 132 147 158 159 162 165–167]

**Table 2** Summary of studies assessing PF and its relationship with maternal and neonatal health outcomes

| Health-related outcome | Related to PF: Biblio no CRF | MF | Flexibility | Balance | Multidimensional | Assoc (-/+) | Statistics | Quality score (0–5) | Unrelated to PF: Biblio no | Quality score (0–5) |
|---|---|---|---|---|---|---|---|---|---|---|
| Maternal Health | | | | | | | | | | |
| Prepregnancy weight | 57 | | | | | – | r=–0.63, P=0.001 | 2 | | |
| Maternal HR at submaximal exercise | 108 | | | | | – | NR, P<0.05 | 1 | | |
| Duration of gestation | 63 | | | | | + | r=0.12, P=0.01 | 3 | | |
| | | | | | | | | | 108 | 1 |
| Physical activity practice | 95 | | | | | + | P=0.01 | 2 | | |
| Back pain | | | 126 | | | + | OR=1.09, 95% CI 1.01 to 1.17, P=0.022 | 2 | | |
| Anxiety | | | | 158 | | – | r=0.559, P=0.02 | 2 | | |
| Fall risk | | | | 128 | | – | P<0.0001 | 3 | | |
| | | | | 157 | | – | P<0.001 | 2 | | |
| Pelvic girdle pain | | | | | 170 | + | P<0.001 | 4 | | |
| Birth | | | | | | | | | | |
| Length of labour in nulliparas | 57 | | | | | – | r=–0.65, P=0.05 | 2 | | |
| Second stage of labour | 108 | | | | | – | NR | 1 | | |
| Caesarean | 13 | | | | | – | NR, P<0.001 | 2 | | |
| Pain during contractions | 111 | | | | | + | r=0.67, P<0.001 | 2 | | |
| Fetal and neonatal health | | | | | | | | | | |
| Fetal umbilical artery pH | 68 | | | | | + | NR, P<0.001 | 2 | | |
| | | | 13 | | | + | r=0.220, P<0.05 | 2 | | |
| Asphyxiated babies | 108 | | | | | – | NR, P<0.05 | 2 | | |
| Arterial umbilical cord PO2 | 13 | | | | | + | r=0.267, P<0.05 | 2 | | |
| | | | 13 | | | + | r=0.237, P<0.05 | 2 | | |
| Arterial umbilical cord PCO2 | | | 13 | | | – | R=0.331, P<0.01 | 2 | | |
| Neonatal birth weight | | 8 | | | | + | r=0.27, P=0.048 | 3 | | |
| | | 115 | | | | + | F (2182)=3.15, P=0.004 | 4 | | |
| | | 13 | | | | + | r=0.191, P<0.05 | 2 | | |
| | | | | | | | | | 93, 26 | 2,1 |
| New-born length | | | | | | | | | 93 | |
| New-born head circumference | | | | | | | | | 93 | |
| Apgar Score | | | | | | | | | 93, 26 | 2,1 |

Related to PF refers to those variables where authors found either a positive or negative association of the variable with PF levels. Unrelated to PF refers to those variables where authors could not find any association between the variable and PF.
+, direct association of the variable with PF; -, inverse association of the variable with PF; CRF, cardiorespiratory fitness; HR, heart rate; MF, muscular fitness; NR, not reported; PCO2, pressure of CO2; PF, physical fitness; PO2, pressure of O2.

one with feet together,[129] two used mixed protocols,[128 160] one with medial malleoli separated[130] and four did not mention the standing posture.[138 149 163 164] Moreover, three articles used protocols with eyes open[132 149 162] exclusively, eight articles used mixed protocols with eyes open and closed, one used visual target and visual tasks[164] and six did not specify whether participants kept their eyes closed or opened. Only one article used a field test, the one-legged standing protocol.[110] On the other hand, one test was a field-test without a platform.

In relation to the 26 articles measuring dynamic balance, 9 assessed balance using platforms. Each of these articles used a different testing tool such as a balance master platform.[133] pressure platform,[163] force platform,[135] Equitest platform[134] and a movable platform, which was used in two articles.[136 137] Two of these articles were walking protocols,[135 163] one with translational perturbations,[157] one was standing with one knee flexed and arms across the chest.[136 137] Another 15 articles used three-dimensional (3-D) camera motion capture systems

using 13 different protocols. Twelve of the 15 articles were walking protocols[139 140 142–144 148 150 152–156 161] and 2 used a stand to sit motion protocol.[141 151] Moreover, one article used a triaxial accelerometer[146]; another article assessed balance through recording (without specification of camera type)[145] and another using instrumented insoles.[176] All three of these articles used walking protocols.

### Speed

The only protocol that was used to assess speed during pregnancy was the 10 m timed walk test (10mTWT). However, the same test was identified in two different articles.[168 169] In the 10mTWT, the participants commenced standing at a chair. When told to start, subjects walked as fast as possible along 14 m marked with white tape placed at 0 m, 2 m, 12 m and 14 m. The time (100th of a second) required to walk between the 2 m and 12 m markers was recorded and converted into speed in metres per second (m/sec).

### Agility and coordination

No articles of agility and coordination were identified.

### Multidimensional

Our search identified a walking multidimensional test that was used in three studies.[168–170] In the Timed Up and Go Test (TUG), the participant began seated in a chair with their arms on armrests and their toes against a start line. The purpose was to cross the front white line at 3 m away, turn around and walk back to the chair and sit down as fast as possible. The performance is measured in time (100th of a second).

## Aim 2: evaluation of the validity and reliability of the fitness tests, and their relationship with maternal and neonatal health

Articles assessing validity and reliability are summarised in table 1. Articles assessing PF and its relationship with maternal and neonatal health outcomes are presented in table 2 and follows a similar format as Sallis *et al*.[177]

### Cardiorespiratory fitness

We identified two articles examining validity.[74 75] Yeo *et al*[74] aimed to validate a portable metabolic testing system (VO2000) on healthy sedentary pregnant individuals. The VO2000 consistently overestimated $VO_2$ measurements, compared with the same manufacturer's reference system, by 4.4±3.6 SD mL/kg/min although the Pearson correlation was significant (r=0.48; p=0.01). When the VO2000 was used twice, the mean difference was statistically significant (1.0±1.8 mL/kg/min; t(45)=3.9, p<0.001). Mottola *et al*[75] provided a prediction equation for $VO_{2peak}$ in pregnant individuals between 16 and 22 weeks of gestation, using a modified Balke protocol. The results of this equation revealed an adjusted $R^2$ of 0.71 and differences between actual and predicted VO2 of 2.7 mL/kg/min. When the authors used this equation to predict $VO_{2peak}$ in a cross-validation group (n=39), they found a predicted value of 23.38±4.03 mL/kg/min, while the actual value was 23.54±5.9 mL/kg/min (p=0.78).

A total of six articles analysed the association of CRF with maternal and neonatal health outcomes. Pomerance *et al*[57] observed that $VO_{2max}$ was inversely associated with the length of labour in multiparas (r=−0.65; p=0.001) and prepregnancy weight (r=−0.63; p=0.001). However, $VO_{2max}$ was not correlated with newborn weight, length or head circumference, or with the 1 min Apgar scores (all p>0.05). In the same line, Wong and McKenzie[108] observed that fit mothers showed lower HR at submaximal exercise intensity (p<0.05) and the second stage of labour was shorter (no statistics reported) compared with unfit pregnant mothers. However, there was no difference between fit and unfit in the length of gestation or weight gained (no statistics reported). In the same article, the authors showed neither positive nor negative effects of maternal fitness on newborn weight or Apgar scores.

In addition, Erkkola and Rauramo[68] found that newborns from fit pregnant individuals had higher pH than fetuses of less physically fit women (p<0.01). In this article, participants with low physical performance were more likely to have asphyxiated neonates than neonates of physically fit women (p<0.05). In the same line, Baena-García *et al*[13] observed that maternal CRF at the 16th gestational week was related to higher arterial umbilical cord $PO_2$ (r=0.267, p<0.05), and those who had caesarean sections had significantly lower CRF compared with those who had vaginal births (p<0.001).

Moreover, Bisson *et al*[8] studied the association of CRF in early pregnancy with physical activity before and during early pregnancy. The authors found that a higher $VO_{2peak}$ in early pregnancy was positively associated with physical activity spent at sports and exercise before and during early pregnancy (p<0.001).

### Muscular fitness

Only two muscular fitness tests assessed reliability.[117 121] Yenişehir *et al*[117] analysed reliability and validity of Five Times Sit-to-Stand. Inter-rater reliability was excellent for subjects with and without pelvic girdle pain (PGP) (intraclass correlation coefficient, ICC=0.999, 95% CI 0.999 to 1.000: ICC=0.999, 95% CI 0.999 to 0.999, respectively). Test–retest reliability was also very high for subjects with and without PGP (ICC=0.986, 95% CI 0.959 to 0.995: ICC=0.828, 95% CI 0.632 to 0.920, respectively).

Gutke *et al*[121] analysed the reliability for an ad hoc test. This test consisted of a maximal voluntary isometric hip extension with a fixed sensor holding a sling around the thigh and pulling for 5 s during 3 reps with 5–10 s of rest (r=0.82 for the right leg and r=0.88 for the left leg; ICC=0.87 for the right leg and 0.85 for the left leg; with *p* value not reported).

Bisson *et al*[8] observed that hand-grip strength was positively associated with infant birth weight (r=0.34, p=0.0068) even after adjustment for confounders (r=0.27, p=0.0480). Żelaźniewicz and Pawłowski *et al*[115] observed that hand-grip strength was associated with offspring

birth weight when controlled for the newborn sex and gestational age at delivery (F(2.182)=3.15; p=0.04). Baena-García *et al*[13] found greater hand-grip strength weakly associated with greater neonatal birth weight (r=0.191, p<0.05). Wickboldt *et al*[111] found that hand-grip strength was moderately correlated with pain scores, where the mean hand-grip strength during contractions had the highest correlation coefficient (r=0.67; p<0.001) compared with peak hand-grip strength (r=0.56; p<0.001) and the area under the curve of hand-grip force (r=0.55; p<0.001).

### Flexibility

Lindgren and Kristiansson[126] designed an ad hoc machine to test passive abduction of the left fourth finger and its relationship with low-back pain during pregnancy and early postpartum. Abduction angle was measured at three different times throughout the pregnancy and once in the postnatal period. Reliability of the abduction angle was analysed by the intraindividual coefficient of variance. The coefficients of variance between the first and second measurement was 0.077, between the second and third 0.070 and between the third and fourth 0.071.

Only two flexibility tests evaluated associations with maternal and neonatal health outcomes. Lindgren and Kristiansson[126] found that women with greater passive abduction angle of the left fourth finger was associated with the highest back pain incidence (OR 1.09; 95% CI 1.01 to 1.17; p=0.022) and the highest number of previous pregnancies (OR 3.24; 95% CI 1.57 to 6.68; p=0.002). Baena-Garcia *et al*[13] found increased flexibility associated with a more alkaline arterial pH (r=0.220, p<0.05), higher arterial $PO_2$ (r=0.237, p<0.05) and lower arterial $PCO_2$ (r=−0.331, p<0.01) in the umbilical cord blood.

### Balance

No validity or reliability assessments were performed regarding balance tests.

Three articles associated balance with neonatal and maternal health-related outcomes. Öztürk *et al*[128] observed that static balance decreased and fall risk increased in pregnant individuals with lower back pain (49.90±24.47 vs 28.47±19.60; p<0.0001). In relation to exercise, McCrory *et al*[157] showed that exercise may play a role in fall prevention in pregnancy (p=0.005) and they also found that dynamic balance is altered in pregnant individuals who have fallen compared with non-fallers and non-pregnant individuals (p<0.001). Nagai *et al*[158] studied the relationship between anxiety and balance. They concluded that when anxiety increases during pregnancy, the standing posture is destabilised (r=0.559, p=0.020), which may increase the chance of falling.

### Speed

Validity and reliability for 10mTWT was studied by Evensen *et al* in two different articles.[168 169] In 2015, Evensen *et al*[168] analysed the test–retest reliability of 10mTWT showing an ICC of (0.74). Intertester reliability was determined in the first 13 participants with strong correlation (ICC=0.94). In 2016,[169] the same authors analysed the convergent validity of 10mTWT by comparing performances with scores achieved on the Active Straight Leg Raise (ASLR) test and observed moderate positive correlations between 10mTWT and ASLR (r=0.65, p=0.003).

This systematic review did not find any articles that analysed the association of speed with maternal and neonatal health outcomes.

### Agility and coordination

No articles were identified.

### Multidimensional

Validity and reliability for TUG was analysed by Evensen *et al* in two different studies.[168 169] The TUG showed good test–retest reliability (ICC=0.88) and intertester reliability (ICC=0.95). Regarding reliability, strong correlations were found between the TUG and ASLR (r=0.73, p=0.001).

The time on TUG among pregnant individuals with PGP was significantly higher (mean (95% CI) 6.9 (6.5 to 7.3) seconds) than for asymptomatic pregnant (5.8 (5.5 to 6.0), p<0.001) and non-pregnant (5.5 (5.4 to 5.6), p<0.001) individuals.

## DISCUSSION

### Summary of the evidence

This systematic review revealed that PF has been assessed through a wide variety of tests during pregnancy. However, little is known on the validity and reliability of the tests performed, and the large variety of tests makes it challenging to compare results from different studies. Until a battery of specific fitness tests for pregnant women is developed and validated, the confidence of PF data obtained during pregnancy is limited and should be interpreted with caution. Consequently, the appropriateness of using this PF data to prescribe exercise during pregnancy could be questioned and is a matter that requires special attention. In this context, it is also difficult to evaluate the association of PF with maternal and neonatal health which, in fact, is of wide clinical and public health interest. However, some studies observed associations of PF with maternal and neonatal health outcomes, which needs to be replicated once a PF test battery is released. We strongly suggest that extensive research must be performed to validate such battery of PF tests.

### Cardiorespiratory fitness

This systematic review identified that a cycle ergometer has been the equipment most frequently used to assess CRF followed by treadmill and field tests, although step tests have also been conducted. There is a large disparity of protocols and wide variety of ad hoc tests used, which makes comparing results between studies difficult. However, the Modified Balke treadmill Protocol validated by Mottola *et al*[75] for pregnant women has been the most

frequently used test. There have been more incremental tests used for CRF tests during the pregnancy compared with steady-state tests and more submaximal compared with maximal tests. There is no consensus regarding test termination criteria for submaximal tests, which undoubtedly needs further research. Some articles used relative intensity using physiological variables such as %$HR_{max}$ or %$VO_{2max}$, and others used absolute intensity, such as specific HR (beats per minute). Among the studies that used %$HR_{max}$ as a test termination criterion, there was a variety of percentages such as 70%,[34 35 90 100] 75%[27 29 69 97] or 85%.[13 54 74] Among the studies that used %$VO_{2max}$, there were different percentages such as 40%,[38] 50%,[37 101] 60%[32 38] or 70%.[30] Among the studies that used absolute HR as a test termination criterion, the HR for finalising the tests were set either at 125,[61] 150,[36 45 62 108] 155,[94] 160[65] or 170[50 53 55 56] beats per minute. Some studies even used the rate of perceived exertion as complementary criteria[46 50 106] or peak aerobic power.[39] These complementary criteria have been recommended and studied in pregnant women by authors like Hesse *et al*[98] since the physical and emotional changes during pregnancy limit performance. It must be noted that the same equation was not used to estimate $HR_{max}$. Some articles used the traditional 220-age formula[29 35 54 69 97] while others used the Karvonen[74] or Tanaka[100] formulas. Some articles did not specify how $HR_{max}$ was estimated.[27 34 90] This heterogeneity could be due to the physiological complexity of pregnancy, in terms of cardiac changes and response to exercise and the lack of scientific information in this regard. Moreover, the gestational week could be a determinant for physiological responses since Bijl *et al*[100] observed a slower haemodynamic recovery and an increased ventilatory response to exercise in early pregnancy compared with non-pregnant women. With regard to maximal tests, different terms have been used for maximal criteria such as volitional fatigue,[30 33 43 44 47 48 98 103 105] exhaustion,[31] anaerobic threshold[73 80 104 171] and point of symptom limitation.[59 60 102]

This lack of consensus has many drawbacks that should be resolved in view of the need to accurately assess CRF during the pregnancy. We advocate for an expert consensus to be developed in the following years to achieve the goal of appropriate and effective CRF assessment during the pregnancy. In particular, it seems essential to develop a treadmill and a cycle ergometer submaximal test that reveals sufficient validity to confidently estimate $VO_{2max}$ throughout gestation.

### Muscular fitness

Muscular fitness tests included muscular strength, endurance and power.[2] The studies included in this systematic review show that muscular strength was the most frequently assessed component of muscular fitness, since only six studies[12 13 112 117 122 178 179] assessed endurance and none of them assessed power in pregnancy. In most studies, muscular strength was evaluated through hand-grip maximal strength using a dynamometer. However, two studies used a hand-grip sphygmomanometer test.[118 119] Some of the hand-grip tests were performed in a standing position,[8 109] while others used a sitting position[110] or supine position,[113] and others did not reveal the position used for the assessment.[86 112 114 115] Some tests were completed three times,[112] others twice[8 86 115] and others only once.[110 113 114] This clearly reveals a large methodological variability that might influence the results and make comparing results between studies difficult. Another limitation is the fact that the main strength outcome was hand-grip strength. While hand-grip strength is a good marker of health,[180] it is unclear whether hand-grip responds to changes following exercise interventions. Therefore, validating other muscular strength tests, including lower limb strength tests, is needed for researchers and practitioners to confidently assess muscular strength during the pregnancy.

There were no validity studies and the reliability was assessed only in one maximal isometric hip extension test.[121] This test has limitations since the pregnant abdomen must be on a bed and, as acknowledged by the authors, it cannot be performed during the third trimester. It must be noted that higher hand-grip strength was associated with higher birth weight.[8 115] Moreover, increased hand-grip strength was produced during uterine contraction.[111] The advantage of using hand-grip is that it represents an inexpensive, rapid and easy-to-use assessment with minimal training needed to appropriately administer. However, assessing the performance of pregnant athletes with this test seems clearly insufficient. More quality in tests employed is necessary since the association of muscular strength with maternal and neonatal health outcomes is of clinical importance. Moreover, other studies are needed to understand the extent to which preserving strength throughout pregnancy and post partum relates to clinical outcomes.

### Flexibility

Although there were seven studies assessing flexibility, none of them used the same protocol. Once again, this reflects a lack of agreement when assessing the same component of PF. Moreover, Lindgren and Kristiansson[126] found that higher flexibility showed higher low back pain. Despite the limitation of a finger laxity test, we considered these findings an interesting association that warrants further investigation since passive stretching is one of the most common practical prescriptions for exercise professionals instead of mobility and breathing exercises. On the other hand, the results of Baena-García *et al*[13] are very relevant to fetal health since flexibility was associated with a better pH, $PO_2$ and $PCO_2$ in umbilical cord blood. Hence, more research about flexibility tests, their outcomes and their prescription are needed.

### Balance

We identified that balance was the second PF component most frequently evaluated during pregnancy, following CRF. This makes sense since the centre of gravity changes during pregnancy as a result of expansion of the

uterus and the risk of falls increases. However, there is high heterogeneity between the protocols employed in different studies. For static balance, the protocol most frequently used was stabilometry on a force platform with bipedal support and eyes open and eyes closed within the same test. For dynamic balance, there was a greater heterogeneity across protocols both in the platform used and in the movements over the platforms. Regarding the assessment tool, the 3-D camera was the device most frequently used.[139–142 144 165] Likewise, we observed differences between the number of platform pieces, trials and Hz used. Some protocols were performed on two piece platforms,[130 131 149] others on one piece platforms[129 132 138 158 160 166 167] and others did not specify the type of platform.[163–165] Although the number of trials and the frequency of recording (ie, Hz) are important protocol parameters that should be carefully documented, only 5 (out of 13) articles described the number of trials[131 138 166 167 175] and 1 described frequency of recording.[149] The usefulness of these tests is restricted to the research area and all of them use expensive technological tools; therefore, it is difficult to extrapolate these tests to fitness centres or clinical settings. Falls during pregnancy could be prevented if balance was easily assessed. For this reason, it is necessary to develop an inexpensive and easy-to-use balance field test.

### Validity and reliability of PF tests, and association with maternal and neonatal health

Unfortunately, studies that examine validity and reliability of PF tests are scarce. The PF component most frequently studied was CRF. However, we only found two studies that analysed the validity of the CRF tests, and no studies examined the reliability of these tests. On a treadmill platform, Mottola *et al*,[75] validated a special equation for modified Balke protocol that has been used by numerous other authors. In contrast, Yeo *et al*[74] aimed to validate a portable metabolic testing system (mod. VO2000) but it overestimated $VO_2$ measurements for pregnant individuals compared with non-pregnant females and males.

Regarding muscular fitness, the hand-grip test was most commonly used; this test was used as the gold standard for muscular fitness during pregnancy. Only Gutke *et al*[121] studied the reliability of a test for hip extension. However, the *p* value was not reported, and the position adopted in the test could be uncomfortable for pregnant participants. Finally, the studies evaluating validity and reliability of speed and multidimensional tests of PF have been researched by Evensen *et al*.[168 169] They demonstrated that TUG and 10mTWT are reliable and valid tests for use during the pregnancy.

The validity and reliability of balance (without tests), agility and/or coordination tests has not been investigated to date.

We suggest that specific tests to be performed in pregnancy are needed and their validity and reliability must be assessed to understand the extent to which one might rely on such measures when prescribing exercise, or making clinical recommendations.

Regarding the association of PF with maternal and neonatal health outcomes, we conclude that more research is also necessary. Nevertheless, from this review we can highlight some interesting associations with different fitness components. A better CRF was associated with a shorter labour[57 108] and a lower risk of caesarean section.[13] However, no association was found regarding other fetal outcomes such as Apgar scores or the newborn anthropometrics.[57 108] By contrast, muscular strength was associated with optimum infant birth weight.[8 13 115] Other neonatal outcomes like fetal umbilical cord pH were positively associated with maternal CRF.[68] On the other hand, better balance scores were associated with lower risk of falls,[128 158 181] which is of particular interest for exercise professionals, who might include balance as a component of exercise programs for pregnant women. Finally, Evensen *et al*[169] found that PGP could be a limiting factor to assess PF in pregnant individuals since the time of TUG was significantly higher in those with pain than in asymptomatic pregnant and non-pregnant individuals.

None of the studies reviewed in this article have described adverse events during PF assessment. Moreover, official bodies such as the American College of Obstetricians and Gynecologists, the Canadian Society of Exercise Physiology and the Society of Obstetricians and Gynaecologists of Canada have highlighted the benefits of an adequate PF assessment, and assert the need of consensus in PF assessment during the pregnancy.[182] Consequently, the findings from this study have important research and clinical implications.

### Limitations and strengths

A limitation of this article is that, although PubMed and WOS are among the most relevant databases in the medical literature, the possibility that a small number of studies have been overlooked cannot be discarded. Nevertheless, these two databases are the biggest databases in sports medicine and sports sciences and, therefore, include the vast majority of studies.

A strength of this systematic review is the fact that, to the best of our knowledge, this is the first article to comprehensively analyse PF assessments, the validity and reliability of fitness tests, and their relationship with maternal and neonatal health outcomes during the pregnancy. The results from this systematic review provide an overall picture of how PF is being assessed in this population, what type of tests are being performed, their specific characteristics, whether these tests have been tested for validity and/or reliability; and whether PF is associated with maternal and neonatal health outcomes. All this information is of wide and undoubted clinical interest.

## CONCLUSIONS

The main finding of this systematic review is that PF has been assessed through a wide variety of protocols, mostly lacking validity and reliability data, and that no

consensus exists on the most suitable fitness tests to be performed during pregnancy. In addition, the available evidence regarding the association of PF with maternal and neonatal health outcomes is scarce and is a matter of further investigation. Provided the need to assess PF during the pregnancy and the importance not only to understand the physical state of the pregnant individual but also to precisely prescribe exercise in this population, extensive research is needed to design and validate a battery of fitness tests to be used for the safe and effective assessment of PF during pregnancy. We advocate for an expert consensus panel to develop a battery of PF tests to assess the different PF components during pregnancy.

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*Assessing physical fitness during pregnancy: validity and reliability of fitness tests, and relationship with maternal and neonatal health– a systematic review*

**Online Supplemental Material 1, table S1.** PRISMA Checklist 2020

| Section and Topic | Item # | Checklist item | Location where item is reported |
|---|---|---|---|
| **TITLE** | | | |
| Title | 1 | Identify the report as a systematic review. | 1 |
| **ABSTRACT** | | | |
| Abstract | 2 | See the PRISMA 2020 for Abstracts checklist. | 2 |
| **INTRODUCTION** | | | |
| Rationale | 3 | Describe the rationale for the review in the context of existing knowledge. | 3 |
| Objectives | 4 | Provide an explicit statement of the objective(s) or question(s) the review addresses. | 3-4 |
| **METHODS** | | | |
| Eligibility criteria | 5 | Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses. | 4 |
| 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. | 4 |
| Search strategy | 7 | Present the full search strategies for all databases, registers and websites, including any filters and limits used. | 4 |
| 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. | 4 |
| 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. | 5 |
| 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. | 5 |
| | 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. | 5 |
| 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. | 5 |
| 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. | - |
| 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 #5)). | - |

*Assessing physical fitness during pregnancy: validity and reliability of fitness tests, and relationship with maternal and neonatal health– a systematic review*

| Section and Topic | Item # | Checklist item | Location where item is reported |
|---|---|---|---|
| | 13b | Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions. | - |
| | 13c | Describe any methods used to tabulate or visually display results of individual studies and syntheses. | - |
| | 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. | 4 |
| | 13e | Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression). | - |
| | 13f | Describe any sensitivity analyses conducted to assess robustness of the synthesized results. | - |
| Reporting bias assessment | 14 | Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases). | ESM2-3 |
| Certainty assessment | 15 | Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome. | ESM2-3 |
| **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. | 5 |
| | 16b | Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded. | 5 |
| Study characteristics | 17 | Cite each included study and present its characteristics. | 5-12 |
| Risk of bias in studies | 18 | Present assessments of risk of bias for each included study. | 5-12 |
| 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. | 5-12 |
| Results of syntheses | 20a | For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies. | 5-12 |
| | 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. | 5-12 |
| | 20c | Present results of all investigations of possible causes of heterogeneity among study results. | 5-12 |
| | 20d | Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results. | 5-12 |
| Reporting biases | 21 | Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed. | - |
| Certainty of evidence | 22 | Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed. | - |

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| Section and Topic | Item # | Checklist item | Location where item is reported |
|---|---|---|---|
| **DISCUSSION** | | | |
| Discussion | 23a | Provide a general interpretation of the results in the context of other evidence. | 12-16 |
| | 23b | Discuss any limitations of the evidence included in the review. | 17 |
| | 23c | Discuss any limitations of the review processes used. | 17 |
| | 23d | Discuss implications of the results for practice, policy, and future research. | 17 |
| **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. | 2 |
| | 24b | Indicate where the review protocol can be accessed, or state that a protocol was not prepared. | 4 |
| | 24c | Describe and explain any amendments to information provided at registration or in the protocol. | 4 |
| Support | 25 | Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review. | 17 |
| Competing interests | 26 | Declare any competing interests of review authors. | 36 |
| 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. | ESM5 |

*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
ESM=Electronic Material Suplement

For more information, visit: http://www.prisma-statement.org/

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**Online Supplemental Material 2.** Search strategy in **PubMed** and **Web of Science.**

For PubMed, we used Medical Subject Heading (MeSH) terms. This is a powerful method to enhance the quality of the search. In addition, all MeSH terms were included without the command MeSH attached, to consolidate our results and avoid losing those papers not included in MeSH database. This is because some MeSH terms were introduced in a specific date (e.g., 'physical fitness' was included in 1996). Hence papers published in a previous date would be lost. The same process was developed with terms not available in the MeSH database such as agility, aerobic capacity, etc. (see ESM 2-Table-S1) for search criteria and related terms.

All terms were combined using the connector OR for similar criteria. The connector 'AND' was used to combine population group (i.e., pregnant women), to delimit date of publication ("0001/01/01"[PDat]: "2021/01/15"[PDat]), to include full text papers, and to include studies performed in humans. A similar search strategy and terms combination was undertaken in WoS (ESM 2-Table-S2), although MeSH terms and its appropriate terms connection were not used as they are exclusive for PubMed.

The first step of the search was to look for systematic reviews and meta-analysis within the field of this systematic review. Since there was no such article published regarding our topic, the research team agreed on starting the search with no limit on the publication date. Then, an initial search was undertaken in both databases following the strategy explained in ESM 2-Table-S1 and ESM 2-Table-S2 for PubMed and WoS database respectively. The results from both, were merged.



Romero-Gallardo L, *et al. BMJ Open Sp Ex Med* 2022; 8:e001318. doi: 10.1136/bmjsem-2022-001318

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**Online Supplemental Material 2, Table S2.** Search strategy used and number of articles found in **Pubmed**.

| ***Search Strategy*** | | | |
|---|---|---|---|
| ("Pregnant Women"[Mesh] OR "Pregnant Women" OR "Pregnancy"[Mesh] OR "Pregnancy") AND ("Physical Fitness"[Mesh] OR "Physical Fitness" OR "Physical Conditioning" OR "Exercise Test"[Mesh] OR "Exercise Test" OR "Fitness Trackers"[Mesh] OR "Fitness Trackers" OR "Muscle Strength"[MeSH] or "Muscle Strength" OR "Muscular fitness" OR "Range of motion, articular"[Mesh] OR "Range of motion, articular" OR "Postural Balance"[MeSH] OR "Postural Balance" OR "Walk Test"[Mesh] OR "Walk Test" OR "Cardiorespiratory Fitness" [Mesh] OR "Cardiorespiratory Fitness" OR "Agility" OR "running speed" OR "aerobic fitness" OR "aerobic capacity" OR "maximal oxygen consumption" OR "V02max" OR "Physical function") AND full text[sb] AND ( "0001/01/01"[PDat] : "2021/01/15"[PDat] ) AND Humans[Mesh] | | | |
| **Search criteria 1** | **MeSH Entry Terms for Criteria 1** | **Search criteria 2** | **MeSH Entry Terms for Criteria 2** |
| Pregnant Women (MeSH) | Women, Pregnant<br>Pregnant Woman<br>Woman, Pregnant | Physical fitness (MeSH) | Fitness, Physical |
| Pregnancy (MeSH) | | Exercise Test (MeSH) | Exercise Tests<br>Test, Exercise<br>Tests, Exercise<br>Arm Ergometry Test<br>Arm Ergometry Tests<br>Ergometry Test, Arm<br>Ergometry Tests, Arm<br>Test, Arm Ergometry<br>Tests, Arm Ergometry<br>Bicycle Ergometry Test<br>Bicycle Ergometry Tests |

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| | | |
|---|---|---|
| | | Ergometry Test, Bicycle<br>Ergometry Tests, Bicycle<br>Test, Bicycle Ergometry<br>Tests, Bicycle Ergometry<br>Fitness Testing<br>Fitness Testings<br>Testing, Fitness<br>Testings, Fitness<br>Step Test<br>Step Tests<br>Test, Step<br>Tests, Step<br>Stress Test<br>Stress Tests<br>Test, Stress<br>Tests, Stress<br>Treadmill Test<br>Test, Treadmill<br>Tests, Treadmill<br>Treadmill Tests<br>Physical Fitness Testing<br>Fitness Testing, Physical<br>Fitness Testings, Physical<br>Physical Fitness Testings<br>Testing, Physical Fitness<br>Testings, Physical Fitness<br>Cardiopulmonary Exercise Test<br>Cardiopulmonary Exercise Tests<br>Exercise Test, Cardiopulmonary<br>Exercise Tests, Cardiopulmonary |

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| | |
|---|---|
| | Test, Cardiopulmonary Exercise |
| Fitness Trackers (MeSH) | Fitness Tracker<br>Tracker, Fitness<br>Trackers, Fitness<br>Physical Fitness Trackers<br>Fitness Tracker, Physical<br>Fitness Trackers, Physical<br>Physical Fitness Tracker<br>Tracker, Physical Fitness<br>Trackers, Physical Fitness<br>Activity Trackers<br>Activity Tracker<br>Tracker, Activity<br>Trackers, Activity<br>Personal Fitness Trackers<br>Fitness Tracker, Personal<br>Fitness Trackers, Personal<br>Personal Fitness Tracker<br>Tracker, Personal Fitness<br>Trackers, Personal Fitness |
| Muscle Strength (MeSH) | Strength, Muscle |
| Muscle strength dynamometer (MeSH) | Dynamometer, Muscle Strength<br>Dynamometers, Muscle Strength<br>Muscle Strength Dynamometers |
| Range of motion, articular (MeSH) | Joint Range of Motion<br>Joint Flexibility<br>Flexibility, Joint<br>Range of Motion<br>Passive Range of Motion |
| Postural Balance (MeSH) | Musculoskeletal Equilibrium |

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| | | | |
|---|---|---|---|
| | | | Equilibrium, Musculoskeletal<br>Postural Equilibrium<br>Equilibrium, Postural<br>Balance, Postural |
| | | Walk Test<br>(MeSH) | Test, Walk<br>Tests, Walk<br>Walk Tests<br>6-Minute Walk Test<br>6 Minute Walk Test<br>6-Minute Walk Tests<br>Test, 6-Minute Walk<br>Tests, 6-Minute Walk<br>Walk Test, 6-Minute<br>Walk Tests, 6-Minute<br>Incremental Shuttle Walk Test<br>Endurance Shuttle Walk Test |
| | | Cardiorespiratory fitness (MeSH) | Fitness, Cardiorespiratory |
| **Total items found** | **Without filters:** 1657 | | |
| | **With Humans filter:** 1135 | | |
| | **With Full Text filter:** 1388 | | |
| | **With Humans & Full Text Filter:** 930 | | |

The search recruited articles published until 15.01.21: no starting date limit was set for the search.

MeSH (Medical Subject Headings) is the National Library of Medicine controlled vocabulary thesaurus used for indexing articles for PubMed

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**Online Supplemental Material 2, Table S3.** Search strategy used and number of articles found in **Web of Science**

| ***Search Strategy*** | |
|---|---|
| TS=("Pregnant women" OR "pregnancy" OR "pregnan*") AND (("Physical Conditioning" OR "Physical fitness" OR "Exercise Test*" OR "Arm Ergometry Test*" OR "Bicycle Ergometry Test*" OR "Step Test*" OR "Treadmill Test*" OR "Physical Fitness Test*" OR "Cardiopulmonary Exercise Test*" OR "Fitness Tracker*" OR "Physical Fitness Tracker*" OR "Activity Tracker*" OR "Personal Fitness Tracker*") OR ("Muscle Strength" OR "Muscular Fitness" OR "Muscle strength dynamometer*") OR ("Joint Range of motion" OR "Joint flexibility" OR "Flexibility" OR "Range of motion" OR "Passive Range of Motion") OR ("Postural Balance" OR "Musculoskeletal Equilibrium" OR "Equilibrium" OR "Postural Equilibrium") OR ("Walk Test*" OR "6-Minute Walk Test*" OR "Incremental Shuttle Walk Test*" OR "Endurance Shuttle Walk Test") OR ("Cardiorrespiratory Fitness" OR "Cardiovascular Fitness OR "Aerobic Fitness" OR "Aerobic Capacity" OR "Maximal Oxygen Consumption" OR "V02max") OR ("Agility" OR "running speed" OR "aerobic fitness" )) | |
| **Total items found** | 1687 |

The search recruited articles published until 15.01.21 no starting date limit was set for the search.

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**Online Supplemental Material 3. Comprehensive description of the three quality assessment scores used in the present systematic review.**

The first quality score [24], was used to evaluate the quality of the articles that assessed validity. This list included three items based on sample size, description of the article population and statistical analysis to assess validity of each article. The validity quality score ranged from 0 to 6 (Table-S3). A score of 0-2 defined a very low-quality article; a score of 3-4 defined a low-quality article; and a score of 5-6 defined a high-quality article.

The second quality score [25] was employed to rate the studies that measured reliability (ESM 4 – Table-S4). This ranking was formed by four items based on description of the participants, the time interval, the results and appropriateness of statistical analyses. Each item in both, was rated from 0 (the lowest quality) to 2 (the highest quality). The reliability quality score ranged from 0 to 8 (ESM 4– Table-S4). A score of 0-1 defined a very low-quality article; a score of 2-5 defined a low-quality article; and a score of 6-8 defined a high-quality article.

The third quality score (ESM 4– Table-S5) was created to evaluate those studies that assessed association of PF with health-related outcomes. We adapted a score previously used in the Effective Public Health Practice Project (EPHPP) [26] which has been used in similar reviews [27]. The health-related outcomes quality score ranged from 0 to 5 (ESM 4– Table S5). A score of 0-2 defined a very low-quality article, a score of 3-4 defined a low-quality article, and a score of 5 defined a high-quality score. Three quality scores were calculated by counting the number of positive items.

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**Online Supplemental Material 3 Table S4. Quality assessment criteria to evaluate validity studies.**

| Grading system parameter | Grade | Criterion |
|---|---|---|
| Number of study subjects | 0 | n < 10 |
| | 1 | n= 11-50 |
| | 2 | n>51 |
| Description of the study population regarding to age, sex, health status, fitness levels, etc | 0 | Less items than required for grade 1 |
| | 1 | At least age and week of gestation. |
| | 2 | Age, week of gestation, health status and fitness levels and more. |
| Statistical analysis included in the study | 0 | Those not included in grade 1 |
| | 1 | Error indexes or regression analysis |
| | 2 | ≥3 items of Bland-Altamn plot and or ANOVA for repeated measurements |

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**Electronic Supplementary Material 4 Table S5. Quality assessment criteria to evaluate reliability studies.**

| Grading system parameter | Grade | Criterion |
|---|---|---|
| Description of the participants | 0 | Less items than required for grade 1. |
| | 1 | At least age and week of gestation. |
| | 2 | Age, week of gestation, health status and fitness levels and more. |
| Description of the time interval | 0 | Interval unknown. |
| | 1 | Vague and imprecise information about interval. |
| | 2 | Precise and complete description about interval. |
| Description of the results | 0 | Less results presented than required for grade. |
| | 1 | Description of test-retest results or description of the differences. |
| | 2 | Description of test-retest results and description of the differences. |
| Appropriateness of statistic | 0 | Only coefficient of variation |
| | 1 | Everything between grades 0 and 2 (normally – but not always – correlation plus an additional statistic). |
| | 2 | At least paired statistics, ANOVA for repeated measures (or non-parametrical corresponding tests) or Bland-Altman method. |

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**Electronic Supplementary Material 4 Table S6. Quality assessment criteria to evaluate health-related outcomes studies.**

| Grading system parameter | Grade | Criterion |
|---|---|---|
| Description of the study sample regarding to number of participants, age, sex, health status, fitness levels, etc | 0 | n ≤ 25 and including less item than required for grade 1. |
| | 1 | N ≥26 and at least age and gestational week. |
| Adequate assessment and report of physical fitness test. | 0 | Items for grade 1 are not included within the article. |
| | 1 | Validity and/or reliability reported of test and detailed description of testing protocol. |
| Adequate assessment of health-related outcomes | 0 | Items for grade 1 are not included within the article. |
| | 1 | Validity or reliability of the outcome measure reported and/or measurement procedure adequately described. |
| Adequate adjustment of confounders | 0 | No adjustment was done. |
| | 1 | Adjustment of confounders such as age and sex were done. |
| Description of both number and reasons to withdrawal and dropout. | 0 | No description included. |
| | 1 | Description included. |

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**Electronic Supplementary Material 5 Table S7. Overview of studies included in the systematic review and description of physical fitness tests.**

| **Reference (authors, year)** | **Sample Size (n)** | **Gestation Weeks (SD) or range in weeks** | **Mean age (SD), or range, in years** | **Fitness Test and Short Description** |
|---|---|---|---|---|
| | | ***Cardiorespiratory Fitness*** | | |
| ***Cycle-ergometer protocol*** | | | | |
| Pomerance et al., (1974)[1] | 54 | 17.5-27 | 35-37 | Ad hoc, steady-state test at 60 rpm at 450, 600 and 300 kpm. |
| Erkkola, (1976)[2] | 120 | (2 weeks before term) | 20-26 | 1) Ad hoc, incremental submaximal test at 150, 300 and 450 kpm/min. 2) Arstila ECG test. |
| Morton et al, (1985)[3] | 23 | 40.15 (1.5) | 28.5 (2.1) | Ad hoc, steady-state test at 40 to 50 rpm and at 300 kpm . $min^{-1}$ for 6 min. |
| Veille et al., (1985)[4] | 17 | 35 (2) | 31 (1) | Ad hoc, incremental submaximal test at 50 and 60 rpm at 50W for 10-15 min to 70% HR max (no formula). |
| Jovanovic et al., (1985)[5] | 6 | 37.1 (0.9) | 28.5 (1.7) | Ad hoc, incremental submaximal self-administered test to 50% $VO_2$ max or exertion equivalent to usual training. |
| Wong & McKenzie, (1987)[6] | 20 | 3 time-points, (10-14; 22-24; 34-36) | 29.13 | Ad hoc, incremental submaximal test at 50 rpm at 25, 50, 75 and 100 W for 5-6 min to 150 bpm. |
| Kulpa et al., (1987)[7] | 141 | First trimester | 18-34 | Bruce protocol to 75% of HR max. |

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| | | | | |
|---|---|---|---|---|
| Carpenter et al., (1988)[8] | 45 | 29 (3.7) | 25.2 (3) | Ad hoc, incremental test. 2 phases: a) *submaximal*: at 0, 30 and 60W for 6 min. b) *maximal*: at 60W to volitional fatigue. |
| Moore et al.,, (1988)[9] | 11 | 21.3 | 26.6 | Ad hoc, incremental submaximal test at free pace for 20 min to 60 to 75% HR max. (220-age). |
| Sady & Carpenter, (1988)[10] | 40 | 29.2 (3.9) | 25.9 (3.3) | 2 incremental tests: 1) *Submaximal test* at 0 W, 30 W and 60 W at 30%, 50% and 70% of $VO_2$ max. 2) *Maximal test* increasing 10 W every 2-min stage to volitional fatigue. |
| Artal et al., (1989)[11] | 37 | 29.8 (0.5) | 28.3 (1.8) | Ad hoc, incremental maximal test at 25, 50, 75W and increments of 25W every 2-min stage until exhaustion. |
| Hume et al., (1990)[12] | 30 | | 28 | Ad hoc, steady-state submaximal test at 60% $VO_2$ max for 20min. |
| Sady et al., (1990)[13] | 9 | 25.6 (3.0); | 29 (4.9) | Ad hoc, incremental test. 2 phases: a) *submaximal*: at 0, 30 and 60 W for 18 min. 6-min each stage. b) *maximal*: incremental continuous to volitional fatigue. |
| Field et al., (1991)[14] | 13 | 33 ± 2 | 30 (4) | Modified Balke protocol to 70% HR max (no formula) |
| Rafla & Beazely, (1991)[15] | 21 | 28-37 | - | Ad hoc, incremental submaximal test from 60 rpm to 70% HR max (220-age) |
| Bung et al., (1991)[16] | 1 | 3 time-points, (24, 28, 37) | 25 | Ad hoc, incremental submaximal test from 15 W to 150 bpm. |
| Young & Treadway, (1992)[17] | 5 | 33 (1) | 29 (1) | Ad hoc, steady-state submaximal test at 50% $VO_2$ max for 30 min. |
| Clapp et al., (1993)[18] | 120 | 16-39 | - | Ad hoc, steady-state submaximal test at 60% ± 3% $VO_2$ max for 30 min. |
| Lotgering et al., (1995)[19] | 33 | 3 time-points, 16.1 (1); 25 (0.7); 35 (0.6) | 30.9 (0.7) | Ad hoc, incremental submaximal test. After 3 min at 15W, to increase 10 W every 30 sec until peak aerobic power. |

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| | | | | |
|---|---|---|---|---|
| Artal et al., (1995)[20] | 7 | 33.86±1.46 | 24.9 (2.18) | Ad hoc, incremental submaximal test. After 5 min per stage at 25, 50 and 75W, to increase 25W every 2 min to volitional fatigue. |
| O'Neill, (1996)[21] | 11 | 35.8 (1.1) | 30.3 (3.3) | 1) Ad hoc, steady-state test at 62.5 W for 15 min. 2) Ad hoc, steady-state test at 87.5 W for 15 min. 3) Ad hoc, steady-state test at 62.5 W for 30 min. |
| Soultanakis et al., (1996)[22] | 20 | 27.1 (1.3) | 31.4 (1.5) | 1) Incremental maximal with modified Balke protocol, increasing 25 W every 2-min at 60 rpm to $VO_2$max 2) Ad hoc, steady-state submaximal test during 1 hour at 50%-60% $VO_2$max at 60 rpm. |
| Manders et al., (1997)[23] | 12 | 29-32 | 20-36 | Ad hoc, incremental maximal test. After 5-min per stage at 50W, to increase 25 W/min to volitional fatigue. |
| Kemp et al., (1997)[24] | 23 | 33 (1) | - | Ad hoc, incremental maximal test at 20 W for 4 min. Then, increasing 20 W/min until exhaustion. |
| McGrath et al., (1999)[25] | 41 | 3 time-points: 17.45 (0.45); 26.5 (0.2) and 37.15 (0.15) | 29.4 (0.85) | Ad hoc, steady-state test with three 6-min stages and exercise brief (<5-min) between them. 1) 20 W to 110 bpm, 2) 45 W to 130 bpm 3) 70 W to 150 bpm. |
| Brenner et al., (1999)[26] | 20 | 27.0 (1.0) and 37.0 (1.0) | 29 (3.35) | Ad hoc incremental submaximal test for 3 min without resistance, then, increased 30 W/min to 170 bpm or RPE of 18. |
| MacPhail et al., (2000)[27] | 23 | 32 (4) | 20-40 | Idem Kemp et al., (1997) |
| Heenan et al., (2001)[28] | 28 | 34.7 (0.4) | 30.8 (1.5 | Idem Kemp et al., (1997) |
| Kennelly et al., (2002)[29] | 22 | 32.1 (1.4) | 25.9 (4.9) | Ad hoc incremental maximal test. After 2-min at 30 W, increasing 10 W/min at 50-60 rpm to achieve AT. |
| Heenan & Wolfe, (2003)[30] | 22 | 37.0 (0.2) | 29 (1.1) | 1) Ad hoc, incremental submaximal test at 20 W for 4 min. Then, to increase 20 W/min until 170 bpm. 2) Ad hoc, incremental ramp test from 0 W increasing work rate in 30-sec periods to 70 or 110% of VT. |

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| | | | | |
|---|---|---|---|---|
| Wolfe et al., (2003)[31] | 18 | 3 time-points: 19.2 (0.8) 27.8 (0.3) 37.0 (0.3) | 28.3 (0.25) | Ad hoc incremental submaximal test for 3 min of no resistance. Then, to increase 30 W/min to 170 bpm or RPE of 18. |
| Lindqvist et al., (2003)[32] | 14 | 5 time-points: 8, 15, 22, 29 and 36. | 29 (5) | Ad hoc incremental submaximal test for 2 min of no resistances. Then, to increase 20 W every 2 min to HR max or pulse oximetry below 95%. |
| Lynch et al., (2003)[33] | 23 | 16, 20, 24, 28, 32, 36 | 28.7(4) | Ad hoc incremental submaximal test at 60 rpm no resistance. Then, to increase 0.5 or 1 kP during two 3-min stages to 130 ± 5 bpm and 1 stage more to 145 ± 5 beats/min. |
| Heenan et al., (2003)[34] | 39 | 37.0 (0.2) | 28.5 (1.4) | 1) Ad hoc incremental submaximal test at 20 W for 4 min. Then, to increase 20 W/min until 170 bpm. 2) Ad hoc, incremental ramp test from 0 W increasing work rate in 30-sec period. (70 or 110% of VT) |
| Pirhonen et al., (2003)[35] | 14 | 5 time-points: (8, 15, 22, 29, 36) | 29.2 (4.6) | Ad hoc incremental submaximal test at 0 W and 20 W for 2 min. Then, to increase at 40 W and thereafter 30 W/min to 85% HR max (220-age) or pulse oximetry below 95%. |
| Kardel, (2005)[36] | 41 | 17, 30, 36 | 27.7 (1.95) | Ad hoc, incremental maximal test for 3-min stages at 50 W, 100 W and 150 W. After a rest, (no longer than 3-min) work maximally (200-280 W) for the first 30 seconds of 3-min stages. |
| McAuley et al., (2005)[37] | 14 | 17.05 (2.05) | 29.9 (0.85) | Ad hoc incremental submaximal and maximal test for 4 min at 20 W at 60-80 rpm. Then, to increase 20 W/min to 170 bpm or volitional fatigue. |
| Weissgerber et al., (2006)[38] | 11 | 7 - 22 | 25-40 | Ad hoc incremental submaximal test at 20 W for 4 min. Then, increasing 5 W/min until volitional fatigue or 170 bpm. |
| Jensen et al., (2007)[39] | 22 | 3 time-points: 19.7 (1.2), 28.2 (0.3), 36.3 (0.3) | 30.9 (0.9) | Idem test 1 of Heenan & Wolfe (2003). |
| Jensen et al., (2008)[40] | 15 | 34-38 | 30.6 (1.0) | Ad hoc incremental maximal test from 6-min resting period. After 25 W/2 min at cadence of 60 and 70 rpm to the point of volitional fatigue. |

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| | | | | |
|---|---|---|---|---|
| Kardel et al., (2009)[41] | 40 | 35-37 | 20-40 | Ad hoc incremental maximal test at 20 W for 2 min. Then, to increase (8-12 min) to ramp up 10% of the predicted maximal load. |
| Ong et al., (2009)[42] | 12 | 2 time-points: 18 and 28. | 30 (4) | Ad hoc, incremental submaximal test increasing 25 W/min to 75 % HR Max (220-age). |
| Thorell et al., (2010)[43] | 520 | 4 time-points: 10.9, 24.0, 29.7, 36.5. | 29.0 (4.4) | Ad hoc incremental submaximal test at 50 or 75 W (based on previous level) increasing 25 W/min to ≥125 bpm. |
| Rojas-Vega et al., (2011)[44] | 20 | 34±1.6 | 35.2 (3.6) | Ad hoc incremental submaximal test free of cadence and speed for 2 min. Then, to increase 25 W/ 2 min at 60 rpm to 150 bpm. |
| Thorell et al., (2015)[45] | 520 | 10.9 | 29.6 | Idem Thorell et al. (2010). |
| Kim et al., (2015)[46] | 32 | 13-35 | 24.8 (2.5) | Ad hoc, steady-state test with three 20-min phases:1) standing 2) pedalling at 50 W for 20 min 3) sitting. |
| Nakagaki et al., (2016)[47] | 20 | 25.1(6.3) | 33.7(4.2) | Ad hoc, incremental submaximal test at 50 rpm to 160 bpm or impossibility to maintain the pedalling rate. |
| Jedrzejko,et al., (2016)[48] | 22 | 37-41 | 24.4 (3.92) | Ad hoc, incremental submaximal test on supine cycle divided into three 4-min constant stages increasing from 25 W to 75 W. |

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| | | | | |
|---|---|---|---|---|
| Sussman et al., (2019)[49] | 23 | 2 time-points: 14-15 and 33-34 gw | 30 (3) | YMCA protocol. Incremental test on semirecumbent to 60-80% $HR_{Max}$ or RPE of 14 out 20. |
| Purdy et al., (2019)[50] | 63 | 4 groups: 10-12, 20-27, 30-37 | 30.5 (4.5) | Ad hoc, incremental maximal test on recumbent cycle at 25 W at 50 rpm for 5 min. Then, to increase 25 W/min at same speed to volitional fatigue. |
| Bilodeau et al., (2019)[51] | 58 | 3 time-points: 16.5 (1.0), 35.6 (0.9); 39.8 (1.1) gw | 30 (3.7) | Modified Bruce ramp protocol. |
| Matenchuk et al., (2019)[52] | 47 | 4 groups: nonpregnant; 1st trimester, 2nd trimester, 3rd trimester | | Ad hoc, incremental maximal test at 25 W at 50 rpm for 5 min. Then, to increase 25 W/min to volitional fatigue. |
| Correa et al., (2020)[53] | 48 | 2 time-points: 18; 36 gw. | | Ad hoc, incremental ramp submaximal test at 4 W for 4 min. Then, to increase 20 W/min until symptom limitation or $HR_{Max}$ (220-age). |
| Bijl et al., (2020)[54] | 40 | 11 (1) | | Ad hoc, incremental submaximal on an upright cycle ergometer for 3-min at 40rpm. Then, to increase at 60-70 rpm at 25 W followed by a rise of 5 Watt in every 12-s to 70% $HR_{max}$ (Tanaka formula). |
| ***Treadmill protocol*** | | | | |
| Sibley et al., (1981)[55] | 13 | 2 time-points: 21.9 (2.3); 33.9 (2.3) | 24.3 (1.4) | Balke protocol to 140 bpm. |
| Veille, (1985)[4] | 17 | 35 (2) | 31 (1) | Ad hoc, incremental submaximal walking test to 70% $HR_{Max}$ (no equation to calculate $HR_{max}$ shown). |
| Lewis et al., (1988)[56] | 28 | 2 time-points: 22 wg and 30 wg | 27.8 (3.3) | Modified Balke protocol. |
| Artal et al., (1989) [11] | 37 | 30.3 (1.9) | 25.9 (2.5) | Modified Balke protocol. |

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| Clapp, Little & Capeless, (1993)[18] | 120 | 16-39 | NR | 1) Ad hoc, steady-state test at 40% ± 3% $VO_2$ max for 30 min. 2) Idem at 60% ± 3% VO2max. |
| Winn et al., (1994)[57] | 12 | 26-36 | 32 (4) | Modified Bruce Protocol to 75% HR Max (220-age). |
| Marquez-Sterling et al., (2000)[58] | 15 | 19.1 (2.15) | 29.5 (3.1) | Ad hoc incremental test at 4 km/h and 0% grade for 2-min. Then, increasing 6 km/h and 2.5% every 2-min to 150 bpm. |
| Santos et al., (2005)[59] | 72 | 17.9 (3.6) | 27.3 (4.65) | Ad hoc, incremental ramp test from 2.4 km/h and 0% grade to AT. |
| Yeo et al., (2005)[60] | 9 | 19 (5) | 30 (3) | 2 Cornell Protocol (85%MHR; Karvonen formula) with 2 systems (VO2000 and CPX/D). |
| Mottola et al., (2006)[61] | 156 | 16-22 | 30.8 (3.7) | Modified Balke protocol with this equation $VO_2$ peak (predicted) = (0.055*peak HR) + (0.381* incline) + (5.541* speed (mph)) + (-0.090*BMI) -6.846 : incremental walking test at 3 mph for 5 min, 0% grade. Then, increase 2% every 2 min. Max inclination permitted 12% grade. Then, increasing speed 0.2 mph every 2-min to volitional fatigue. |
| Davenport, et al., (2008)[62] | 106 | 16-20 | 20-39 | Modified Balke protocol. Idem Mottola et al., 2006. |
| Oliveria et al., (2012)[63] | 187 | 3 time-points: 13, 20, 28. | 24.7 (5.5) | Modified Balke protocol. Idem Mottola et al. (2006). |
| Ruchat et al., (2012)[64] | 44 | 2 time-points: 16-20 and 34-36 | 30.8 (4.2) | Modified Balke protocol. Idem Mottola et al. (2006). |
| Szymanski, (2012)[65] | 45 | 30.4 (1) | 33.36 | Modified Balke protocol. Idem Mottola et al. (2006). |
| Salvesen et al., (2012)[66] | 6 | 25.5 | 32 | Ad hoc, incremental maximal test at 6% grade increasing speed in periods of 1km/h every 5-min to volitional fatigue. |

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| Mottola et al., (2013)[67] | 40 | 35.7 (0.4) | 33.5 (0.7) | Ad hoc, steady-state test for 40-min, preceded by a 5-min warm-up increasing speed and inclination to 95% VT. |
| Bisson et al., (2013)[68] | 65 | 16 | 29.9 (4.5) | Modified Balke protocol. |
| LeMoyne et al., (2014)[69] | 67 | 1st trimester, 2nd trimester, 3rd trimester | 29.6 (5.5)<br>30.1 (3.1)<br>32.3 (3.7) | Ebbeling single-stage submaximal treadmill walking test. |
| Bisson et al., (2014)[70] | 61 | 16 (0.6) | 30.0 (4.5) | Modified Balke protocol. |
| Marshall et al., (2015)[71] | 51 | 3 time-points: 20, 32 | 29.2(5.3) | Ad hoc, incremental submaximal test at 0% grade and 3.21 km/h for 5-min. Then, two 5-min stages with speed and grades self-administered to moderate (brisk walk) and vigorous (jog/run) respectively. |
| Santos et al., (2016)[72] | 28 | 30.51 (3.3) | 26 (6.9) | Modified Balke protocol. |
| Hesse et al., (2018)[73] | 25 | 22.1 (1.4) | 30 (3.6) | Bruce protocol until volitional fatigue. |
| Baena-García et al., (2020)[74] | 127 | 16 | 32.9 (4.6) | Modified Bruce protocol until 85% $HR_{Max}$ |
| Dobson et al., (2020)[75] | 22 | 3 time-points: Early- (13–18 gw), mid- (24–28 gw) and late-pregnancy (34–37 gw). | 31.4 (3.7) | Submaximal incremental Walking Exercise Test (SWET) during 21-min on a treadmill. From 3.2 km/hr at 4 min at 2% grade, to increase 2% every 3 min over seven stages. |
| ***On track*** | | | | |
| Bung et al., (1991)[16] | 1 | 3 time-points(24, 28, 37) | 25 | Ad hoc, maximal test. 3 sprints of 200 m and one of 100 m on track |

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| Da Silva et al., (2010)[76] | 74 | 37 | 21.5 | 6-minute walk test. |
| Ramírez-Vélez et al., (2011)[77] | 64 | 2 time-points: 18.6 (3.4) and 16 weeks later. | 19.5 (2.3) | 6-minute walk test. |
| Hjorth et al., (2012)[78] | 304 | 25.0 (7.3) | 23.0 | Ad hoc, steady-state walking test for 250 m on ground level at their normal walking pace. |
| Price et al., (2012)[79] | 62 | 5 time-points: 12–14, 18–20, 24–26 and 30–32 | 29.05 | Ad hoc test walking or running as fast a as possible within comfort zone at a steady pace. Power = (weight x distance) / time. |
| Radzikowska et al., (2017)[80] | 45 | 3-7 | 24-36 | 6-minute walk test. |
| Oviedo-Caro et al., (2018)[81] | 134 | 20 | 32.5 (4.2) | 6-minute walk test. |
| Dennis et al., (2019)[82] | 300 | 37 (1.3) | 31 (4.2) | 6-minute walk test. |
| Amola et al., (2019)[83] | 34 | 3rd trimestre | 25.1 (7.5) | 6-minute walk test. |
| Birnbaumer et al., (2020)[84] | 39 | 26 (7) | 26 (3.4) | Ad hoc, incremental walking test on a 400 Walking speed was paced by audio every 10 m and started at 3 km/h. Then, to increase 0.5 km/h every 50 m to participants were unable to walk the given pacer speed. |
| ***Step Protocol*** | | | | |
| Dibblee & Graham (1983)[85] | 16 | 3 time-points: (the last month of each trimester) | 23-31 | Canadian Home Fitness Test. |
| Williams, Reilly et al. (1988)[86] | 16 (10 pregnant | First, second and third trimester. | 25.6 (3.6) | Ad hoc, incremental test at 115, 135, and 155 bpm for 5 min. |

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| | and 6 non-pregnant) | | | |
| Melzer et al., (2010)[87] | 44 | 38.27 | 31 (5.6) | Ad hoc, incremental test at 15-32.5 body lifts per minute (rate of change: 2.5 body lifts/ min$^2$). Mechanical power was calculated as: 9.81 m/s$^2$ x step height (m) x lift frequency (number of body weight lifts/ min) and expressed in J/min/kg |
| ***Muscular Fitness*** | | | | |
| Baker & Johnson (1994)[88] | 200 | NR | 28-32 | Hand Grip Sphygmomanometer Test: Pressing an inflated cuff of for 30-sec to MVCF over 3-min period. |
| Rogers & Tomilson (1998)[89] | 20 | NR | 5 times: 12, 18, 24, 30, 36 | Hand Grip Sphygmomanometer Test at 30% of MVCF for 2-min. |
| Feiner et al. (2000)[90] | 34 | 22-36 | 22-35 | Isometric Hand-Grip Test with dominant hand for 3 min at one-third of MVCF. |
| Gutke et al., (2008)[91] | 301 | 12-18 | 29 | 1) Maximal voluntary isometric hip extension test with a fixed sensor holding a sling around the thigh and pulling for 5 sec during 3 reps with 5-10-sec of rest. 2) Isometric back flexors endurance: Maintaining an abdominal crunch for a maximum of 120 sec. |
| Thorell et al., (2010)[43] | 520 | 1 time-points: 10.9 | 29.0 (4.4) | Sit-up test. Supine position with the knees at a 90º angle and the feet flat on the floor. 3 sets per 5 repetitions, without a rest or to stop when they were unable to perform of 15 repetitions of sit-ups. |
| O'Connor et al., (2011)[92] | 32 | 21-25 | 18-38 | Ad hoc 5 tests: 1) Seated leg press; 2) Leg curls; 3) Leg extension; (4) Lat pull; (5) Back extension. |

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| Hjorth et al., (2012)[78] | 304 | 25.0 (7.3) | 23.0 | Hand-Grip maximal strength test twice on dominant and non-dominant side alternatively. |
| Price et al. (2012)[79] | 62 | 5 time-points: 12–14, 18–20, 24–26 and 30–32 | 29.1 | Ad hoc test. Lifting a 7-kg medicine ball from the floor to waist height as many times possible for 1 min. |
| Bisson et al. (2013)[68] | 65 | 16 | 29.9 (4.5) | Hand-Grip maximal strength test twice on dominant and non-dominant side alternatively. Adjusting the handle of dynamometer. |
| Atay et al., (2015)[93] | 37 | 2 time-points: 20 and 32 | 29.6 (5.9) | Hand-Grip maximal strength test in a sitting position. |
| Petrov et al., (2015)[94] | 92 | 2 time-points: 13 and 35 | 30.7 (3.5) | Hand-Grip isometric peak strength. |
| Wickboldt (2015)[95] | 43 | 32 (4) | 37-42 | Hand-Grip maximal strength test during the uterine contraction. |
| Kalliokoski et al. (2016)[96] | 51 | NR | 28.3(6.4) | 1) Hand-Grip maximal strength test for 10 sec 3-times in each hand.<br>2) Ad hoc upper leg performance test through 3 movements: a) To rise once after a squat b) to stand on one leg for 30 sec 3) Trendelenburg's test. It was evaluated able or unable. |
| Ngaka et al. (2016)[97] | 50 | >37 | 28.8 (5.7) | Hand-Grip maximal strength test in a supine position. |
| Rodriguez-Díaz et al., (2017)[98] | 105 | 24-30 | 32.2 (4.7) | Hand-Grip maximal strength test for each hand. |

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| Zelazniewicz, (2018)[99] | 95 | 3 time-points (once in each trimester) | 29.6 (3.4) | Hand-Grip maximal strength test twice on dominant and non-dominant side alternatively. |
| Takeda et al., (2019)[100] | 21 | 22 and 23.25 gw. | 32 (3.3) | 1) Toe grip dynamometer 2) Hand-held dynamometer fixed to the legs of the chair with a belt not stretchable to assess quadriceps strength . |
| Baena-García et al., (2020)[74] | 156 | 16 | 32.9 (4.6) | 1) Hand-grip maximal strength twice on dominant and non-dominant side alternatively with 30 sec rest between them.<br>2) 30-sec Chair Stand Test |
| Yenisehir et al., (2020)[101] | 167 | Second and third trimester. | 28.4 (4.6) | 5 Times Sit to Stand test, 5 repetitions of sit-to-stand maneuver as fast as possible with fold arms across the chest. |
| | | ***Flexibility*** | | |
| Gilleard et al. (2002)[102] | 21 | 4 time-points: 18 or less, 24, 32, 38 | 21-40 | 3 tests measured with Expert Vision™ Motion Analysis System: 1) Seated and standing forward flexion 2) Seated and standing side-to-side flexion 3) Seated axial rotation |
| Marnach et al. (2003)[103] | 46 | 3 time-points: 8-12, 16-22, 34-36. | 28.8 (0.8) | Wrist flexion-extension and medial-lateral deviation using goniometer |
| Garshasbi et al. (2005)[104] | 212 | 17-22 | 26.4 (4.7) | Side bending test: Both sides. |
| rice et al., (2012)[79] | 62 | 5 time-points: 12–14, 18–20, 24–26 and 30–32. | 29.1 | Sit-and-reach test. |
| Lindgren et al. (2014)[105] | 200 | 3 time-points: 11, 24 and 36. | 28.4 (5.9) | Ad hoc machine to test passive abduction of the left fourth finger. |
| Atay et al., (2015)[93] | 37 | 2 time-points: 20 and 32, | 29.6 (5.9) | Back scratch test. |
| Rodriguez-Díaz et al., (2017)[98] | 105 | 24-30 | 32.2 (4.7) | Isquiosural flexibility test by goniometer. |

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| Cherni et al., (2019)[106] | 17 | 3 occassions: first, second and third trimester | 36 (2) | 4 tests measured with optoelectronical system: 1) Extensometer of the metacarpophalangeal joint of the index. 2) Figertrip to floor test: from 20cm platform, to reach the floor with knees extended; 3) Sit-and-reach test adapted on delivery bed 4) Beighton score |
| Baena-García et al., (2020)[74] | 156 | 16 | 32.9 (4.6) | Back Scratch |
| | | ***Balance*** | | |
| ***Stabilometry – On force platform or pressures platform*** | | | | |
| Butler et al., (2006)[107] | 12 | 3 time-points: 11-14, 19-22, 36-39 | 32.9 (5.5) | Standing with eyes open and eyes closed for 30 sec each. 3 trials. 1 piece. Force Platform. |
| Ribas et al., (2007)[108] | 60 | 3 time-points: 1) Up to 12 week 2) 13-24 3) Upwards of 25 weeks | 23.3 (4.8) | Standing with bipedal support and eyes open for 5 sec. 2 pieces at 40 Hz. |
| Nagai et al., (2009)[109] | 43 | 30.3 (0.8) | 33 (0.65) | Standing with feet parallel, gazing a black 12-cm circle fixed at a 1.5 m distance with eyes open and eyes closed for 1 min each. 1 piece. |
| Oliveira et al., (2009)[110] | 20 | 3 time-points: 15.1 (1.8); 24.0 (2.4); 34.5 (2.5) | 28.7 (6.2) | Standing with 4 protocols at 50Hz and 2-min rest periods between them: 1) Eyes open with feet comfortably apart; 2) Eyes closed with feet comfortably apart; 3) Eyes open with feet together; 4) Eyes closed with feet together. 1 piece. |
| Karadag-Saygi et al., (2010)[111] | 35 | 33 (3) | 29.8 (4.5) | Standing for 60 sec. |
| Yu et al., (2013)[112] | 21 | NR | 30.2 (3.05) | Standing with heels on a line at 1.0 m from visual target with visual tasks and inspection tasks. |
| Ersal et al., (2014)[113] | 69 | 2 time-points: 20.9 (1.2) and 35.8 (1.5) | 28.3 (5.0) | Standing with feet hip-width apart and staring straight ahead on Equitest platform. |

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| 7Opala-Berdzik et al., (2014)[114] | 31 | 36.2 (1.2) | 28.2 (3.6) | Standing with arms at both sides and in a comfortable stance on a stable force platform with eyes open and eyes closed for 2 trials of 30-sec and 1-min rest between them. |
| Opala-Berdzik et al., (2015)[115] | 45 | 2 time-points: 13.1 (2.5) and 36.2 (1.2) | 28.2 (3.6) | Idem Opala-Berdzik et al., (2014) |
| Ozturk, (2016)[116] | 68 | 31.5 (4.73) | 30.3 (3.6) | Standing and arms extended in 6 different positions for 32-sec: 1) facingforward eyes open and eyes closed; 2) Eyes closed head rotated at 45º to the right 3) Idem 45º to the left; 4) Eyes closed, head tilted at 30º backward and 5) Idem 30º forward; 6) Standing on an unstable cushion, facing forward eyes open and eyes closed. 4 pieces. |
| Shibayama et al., (2016)[117] | 161 | 28-33 | 33.3 (4.7) | Standing and feet together for 30 sec on force platform. 1 piece. |
| Takeda et al., (2018)[118] | 100 | 2nd and 3rd trimester | 20-30 | Standing with the medial malleoli 100 mm apart for 10-sec. Then, moving forward, backward, right and left for 10-sec each. 2 pieces. |
| Moreira et al., (2017)[119] | 30 | 1st and 3rd trimester | 26.8 (5.1) | Standing with each foot positioned on each triaxial force plate (feet apart by ~20 cm) and arms along the body with eyes open focusing on a target located ~2 m in front and eyes closed for 3 trails of 60-sec each and 2-min rest. 2 pieces. |
| Opala-Berdzik et al., (2018)[120] | 70 | 10.8 (1.6) | 28.6 (4.4) | Standing with arms at both sides and in a comfortable stance on a stable force platform, with eyes open looking straight ahead at a wall 3m away for 2 trials of 30-sec and 1-min rest between them. |

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| Catena et al., (2019)[121] | 17 | 9 time-points: 16-20 gw, 36-40 gw and 1 time per month up to 7 months postpartum | 28.9 (4.0) | 2 trials: 1) quiet static in anatomical position for 10 s on a force plate; 2) Idem 1 on a back-board spanning two force plates. |
| Fontana et al., (2020)[122] | 24 | 23 (3) | 30 (6) | Standing barefoot two-legged stance with arms at both sides with eyes open at 2 m from a cross placed on a wall at eye level during 3 x 30s trials with 30 s rest intervals. The mean was retained on force platform. |
| Valerio et al., (2020)[123] | 40 | 30.8 (3.9) | 28 (2.5) | Standing barefoot with freestanding supports inside the platform and arms by their sides. And staring at a mark on the opposite wall. 3 trials with the eyes open and three trials with eyes closed, with 30 s rest intervals. |
| Takeda et al., (2019)[100] | 21 | 22 and 23.25 gw. | 32 (3.3) | Standing barefoot on 2 stabilometers. 3 trials: 1) 10-sec standing position; 2) 10-sec moving in the anterior position; 3) 10-sec moving in the posterior position. |
| ***Others*** | | | | |
| Atay et al., (2015)[93] | 37 | 2 time-points: 20 gw and 32 gw | 29.6 (5.9) | One-legged stand test. |
| | | ***Dynamic Balance*** | | |
| | | ***On platforms*** | | |
| Davies et al., (2002)[124] | 150 | Day of labour | 30.2 (5.8) | Balance Master Platform Tests: 1) Sit to Stand; 2) Walk Test, 3) Step and Quick Turn, 4) Step Up and Over. |

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| Karadag-Saygi et al., (2010)[111] | 35 | 33 (3) | 29.75 (4.5) | Walking barefoot 4 m. |
| McCrory et al, (2010)[125] | 81 | 2 time-points: 20.9 (1.2) and 35.8 (1.5) | 28 (5.7) | The Motor Control Test protocol with translational perturbations. Equitest posture platform. |
| Branco et al., (2013)[126] | 22 | 27 (1.3) | 32.5 (2.6) | Walking barefoot for 10 m between 2 points in a straight line at a natural and comfortable speed for 3 min. |
| Cakmak et al., (2014)[127] | 41 | 6-12 | 26.5 (4.7) | Standing with knee flexed, arms placed across the chest and glare fixed ahead with open eyes on a movable platform provides up to 20º of surface tilt in a 360º range of motion for 3 trails of 20 sec each. |
| Inanir et al., (2014)[128] | 110 | 3 groups: 1st trimester, 2nd trimester and 3rd trimester. | 24.7 (5.2) | Idem to Cakmak et al., (2014) |
| ***3-D Camera motion capture system*** | | | | |
| Wu et al., (2004)[129] | 25 | 27 | 33.1 | Walking on a treadmill at different velocities (incrementing 0.11 m/s, from 0.17 up to 1.72 m/s; for 3 min at each level). |
| Forczeck et al., (2012)[130] | 13 | NR | 29.2 (3.5) | Walking barefoot at a self-selected speed across the room during 15 gait cycles.10 |
| Takeda et al., (2012)[131] | 16 | 24.85 (1.95) | 35 (1.4) | Stand-to-sit motion assessing the time taken to sit down; the leg joint moment; the antero-posterior and vertical floor reaction forces; and the range of motion of the lower limbs and trunk. |
| Gottschall et al., (2013)[132] | 13 | 2 time-points: 20 and 32 | 31.3 (4.5) | Walking along 25 m on a custom-built portable apparatus composed of a 2.4 m ramp inclined at 15° continuous with a 4.8 m plateau. |
| McCrory et al., (2014)[133] | 69 | 28.35 (1.35) | 28.0 (5.7) | Walking along the 8-m runway. |

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| Krkeljas, (2018)[134] | 35 | 3 time-points: 9-12 gw; 20-22 gw and 28-32 gw. | 27 (6.1) | Walking on a straight line, at a self-selected pace along the 15-m walkway. |
| Catena et al., (2019)[135] | 15 | 5 time-points: 16-20; 20-24, 24-28; 28-32; 32-36 gw. | 29.3 (3.7) | 60-second trial of semi-continuous stand-to-sit motion. 54 reflective markers were adhered to body land. |
| Catena et al., (2019)[136] | 15 | 7 time-points: 12-16; 16-20; 20-24, 24-28; 28-32; 32-36; 36-40 gw | 28.1 (4.3) | Walking on a treadmill for 60 seconds at a self-selected comfortable speed. |
| Forczek et al., (2019)[137] | 30 | 3 time-points: 12, 25, 36 gw. | 30.3 (3.4) | Walking barefoot at a self-selected speed during 12m intervals. 10 gait cycles. |
| Forczek et al., (2019)[138] | 14 | 2 time-points: pre-pregnancy; 1st trimester | 20-40 | Walking barefoot across room at a self-selected during 50 m with 1 min rest intervals. 10 gait cycles. |
| Catena et al., (2020)[139] | 23 | 5 time-points: 18, 22, 26, 30, 34 gw. | | Walking on a treadmill for 60 seconds at a self-selected comfortable speed. |
| Gimunova et al., (2020)[140] | 41 | 4 time-points: 14, 28, 37 gw | 30.5 (4.1) | Walking barefoot along a 6-meter walkway at a self-selected. |
| McCrory et al., (2020)[141] | 95 | 2 time-points: 2nd and 3rd trimester | 28.4 (5.5) | Walking along the 8m laboratory runway until walking speed stabilized. |
| Rothwell et al., (2020)[142] | 17 | 2 time-points: 16-20; 36-40 gw | 22-37 | Walking on a treadmill for 60 seconds at a self-selected comfortable speed. |
| Forczek et al., (2019)[143] | 36 | 3 time-points: 12; 25; 36 gw | 30.3 (3.4) | Walking across the room 50 m with 1-min rest interval. 10 gait cycles. |
| ***Others*** | | | | |
| Sawa et al., (2015)[144] | 27 | 2 groups: early pregnancy (≤27 gw) or late pregnancy (≥27 gw) | 30.9 (4.2) | Walking at self-pace speed along a 15-m smooth, horizontal corridor. It was recorded with 2 wireless motion-recording-sensor units and one piezo-resistive triaxial accelerometer. |

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| Błaszczyk et al., (2016)[145] | 28 | 1st trimestrer and 3rd trimester | 28.2 (3.4) | Walking along 10-m long walkway (back and forth 10 times) at self-space speed. It was recorded by custom made; self-adhesive copper foil electrodes attached to the soles of their shoes. |
| | | | ***Speed*** | |
| Evensen et al., (2015)[146] | 17 | 28.7 (7.4) | 31.1(2.3) | Ten-metres Timed walk Test (10mTWT) |
| Evensen et al., (2016)[147] | 18 | 28.9 (7.3) | 31.4 (2.7) | 10mTWT |
| | | | ***Multicomponent*** | |
| Evensen et al., (2015)[146] | 17 | 28.7 (7.4) | 31.1(2.3) | Timed Up and Go Test (TUG) |
| Evensen et al., (2016)[147] | 18 | 28.9 (7.3) | 31.4 (2.7) | TUG |
| Christensen et al., (2019)[148] | 74 | 23 | 31.2 (3.7) | TUG |

Ad hoc: test designed specifically for that study; NR: Not reported; PFS: Physical Fitness Score; kpm: kilopoundimeter; min: minutes; sec: seconds; $HR_{Max}$: Heart Rate Maximum; $VO_2$ max: oxygen consumption maximum; RPE: rate of perceived exertion; AT: anaerobic threshold; ICC: intraclass correlation coefficient; MVCF: maximal voluntary contraction force; HGS: hand-grip strength; m: meters; mm: millimeters; FP: force platform; PP: pressure platform.

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