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Fetal fibronectin testing for reducing the risk of preterm birth.

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Fetal fibronectin testing for reducing the risk of preterm birth.

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[Intervention Review]

Fetal fibronectin testing for reducing the risk of preterm birth

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ABSTRACT

Background

Fetal fibronectin (FFN) is an extracellular matrix glycoprotein localized at the maternal-fetal interface of the amniotic membranes, between chorion and decidua, where it is concentrated in this area between decidua and trophoblast. In normal conditions, FFN is found at very low levels in cervicovaginal secretions. Levels greater than or equal to 50 ng/mL at or after 22 weeks have been associated with an increased risk of spontaneous preterm birth. In fact, FFN is one of the best predictors of preterm birth in all populations studied so far, and can help in selecting which women are at significant risk for preterm birth. This is an update of a review first published in 2008.

Objectives

To assess the effectiveness of management based on knowledge of FFN testing results for preventing preterm birth.

Search methods

For this update, we searched Cochrane Pregnancy and Childbirth's Trials Register (7 September 2018), ClinicalTrials.gov, the WHO International Clinical Trials Registry Platform (ICTRP) (7 September 2018), and reference lists of retrieved studies.

Selection criteria

Randomized controlled trials of pregnant women screened with FFN for risk of preterm birth. Studies included are based exclusively on knowledge of FFN results versus no such knowledge, and we have excluded studies including women with only positive or only negative FFN results.

Data collection and analysis

Two review authors independently assessed trials for inclusion and risk of bias, extracted data, and checked them for accuracy. The quality of the evidence was assessed using the GRADE approach.

Main results

We identified 16 trials, of which six were eligible for inclusion. The six included studies randomized 546 women with singleton gestations and threatened preterm labor (PTL) at 23 0/7 to 34 6/7 weeks. A total of 277 women were randomized to knowledge and 269 to no knowledge of FFN. No trials were identified on asymptomatic women or multiple gestations.

The risk of bias of included studies was mixed. For selected important outcomes, preterm birth before 37, 34, and 32 weeks, and maternal hospitalization, we graded the quality of the evidence and created a 'Summary of findings' table. For these outcomes, the evidence was graded as mainly low quality due to the imprecision of effect estimates.

Management based on knowledge of FFN results may reduce preterm birth before 37 weeks (21.6%) versus controls without such knowledge (29.2%) (risk ratio (RR) 0.72, 95% confidence interval (CI) 0.52 to 1.01; 4 trials; 357 women; low-quality evidence). However, management based on knowledge of FFN results may make little or no difference to preterm birth before 34 (RR 1.09, 95% CI 0.54 to 2.18; 4 trials; 357 women; low-quality evidence) or maternal hospitalization (RR 1.06, 95% CI 0.79 to 1.43; 5 trials; 441 women; low-quality evidence). The evidence for preterm birth before 32 weeks is uncertain because the quality was found to be very low (average RR 0.79, 95% CI 0.16 to 3.96; 4 trials; 357 women; very low-quality evidence).

For all other outcomes, for which there were available data (preterm birth less than 28 weeks; gestational age at delivery (weeks); birthweight less than 2500 g; perinatal death; tocolysis; steroids for fetal lung maturity; time to evaluate; respiratory distress syndrome; neonatal intensive care unit (NICU) admission; and NICU days), knowledge of FFN results may make little or no difference to the outcomes.

Authors' conclusions

The evidence from this review suggests that management based on knowledge of FFN results may reduce preterm birth before 37 weeks. However, our confidence in this result is limited as the evidence was found to be of low quality. Effects on other substantive outcomes are uncertain due to serious concerns in study design, inconsistency, and imprecision of effect estimates. No trials were identified on asymptomatic women, or multiple gestations.

Future studies are needed that include specific populations (e.g. singleton gestations with symptoms of preterm labor), a study group managed with a protocol based on the FFN results, and that report not only maternal but also important perinatal outcomes. Cost-effectiveness analyses are also needed.

PLAIN LANGUAGE SUMMARY

Fetal fibronectin testing for reducing the risk of preterm birth

What is the issue?

To assess the effectiveness of management of pregnant women based on a knowledge of fetal fibronectin test results for preventing preterm birth, compared with not having that knowledge. Fetal fibronectin (FFN) acts as a 'glue' between the pregnancy and the uterus. Normally very low levels of FFN can be found in secretions of the vagina and cervix. Raised levels at or after 22 weeks have been associated with an increased risk of spontaneous preterm birth.

Why is this important?

Preterm birth before 37 weeks is the main cause of sickness and death for newborn infants. Most women who give birth preterm have preterm labor symptoms such as contractions but many of the women with symptoms go on to deliver at term (37 weeks or more). Fetal fibronectin (FFN) is a test that can identify the women with symptoms of preterm labor who are most at risk for preterm birth. The level of FFN is measured in secretions from the vagina or cervix.

What evidence did we find?

We found six randomised controlled studies involving 546 women who were pregnant with one baby and were showing signs of preterm labor at between 23 to 34 weeks' gestation. We graded the following evidence as mainly low quality because of the low number of women in the studies and a wide variation in findings. We found that the number of births before 37 weeks may be slightly reduced when women and their doctors know the results of the FFN test (21.6% versus 29.2%; 4 trials; 357 women). However, knowledge of FFN results may make little or no difference for the other outcomes with available data, including: maternal hospitalization (5 trials; 441 women); use of uterine relaxants (tocolysis) to try to prevent labor; earlier preterm births; women's gestational age at delivery; babies with a birthweight less than 2500 g; newborn deaths; the number of babies with respiratory distress syndrome; giving steroids to mature the unborn babies' lungs; and number of days in a neonatal intensive care unit (NICU).

What does this mean?

This review of six studies did not find enough evidence to say whether or not the FFN test should be used in the management of women showing signs of preterm labor. A screening test such as FFN can only be considered effective if interventions based on the screening results, such as giving drugs to relax the uterus, reduce the number of preterm births. Further research should be encouraged.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON *[Explanation]*

FFN knowledge versus no knowledge in reducing the risk of preterm birth						
Patient or population: women with singleton pregnancies and threatened preterm labor (PTL) between 23 to 35 weeks Setting: hospital settings in United Kingdom and United States Intervention: FFN knowledge Comparison: no knowledge						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	No. of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with no knowledge	Risk with FFN knowledge				
Preterm birth < 37 weeks	Study population		RR 0.72 (0.52 to 1.01)	434 (5 RCTs)	⊕⊕○○ LOW ¹²	
	292 per 1.000	211 per 1.000 (152 to 295)				
Preterm birth < 34 weeks	Study population		RR 1.09 (0.54 to 2.18)	357 (4 RCTs)	⊕⊕○○ LOW ¹³	
	79 per 1.000	86 per 1.000 (43 to 172)				
Preterm birth < 32 weeks	Study population		Average RR 0.79 (0.16 to 3.96)	357 (4 RCTs)	⊕○○○ VERY LOW ⁴⁵	
	56 per 1.000	36 per 1.000 (14 to 98)				
Maternal hospitalization	Study population		RR 1.06 (0.79 to 1.43)	441 (5 RCTs)	⊕⊕○○ LOW ¹³	
	268 per 1.000	284 per 1.000 (211 to 383)				

* **The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: confidence interval; **RR:** risk ratio; **OR:** odds ratio;

GRADE Working Group grades of evidence

High quality: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect

Very low quality: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect

¹ We downgraded (1) level for serious limitations in study design due to some studies having unclear risk of bias for several domains and one study being at high risk of detection bias

² We downgraded (1) level for serious imprecision due to wide CI just crossing the line of no effect

³ We downgraded (1) level for serious imprecision due to wide CI crossing the line of no effect

⁴ We downgraded (2) levels for very serious imprecision due to wide CI crossing the line of no effect and a small number of events

⁵ We downgraded (1) level for serious inconsistency due to evidence of statistical heterogeneity $I^2 = 46\%$

BACKGROUND

Description of the condition

Preterm birth is defined by the World Health Organization (WHO) as birth between 20 and 36 6/7 weeks. Its incidence is about 5% to 12% in most low-, middle-, and high-income countries. This incidence is increasing in many countries, including low-income countries.

Preterm birth is the main cause of neonatal morbidity and mortality in most countries, especially in high- and middle-income countries. In the USA, 75% of perinatal mortality occurs in preterm babies; more than two-thirds of perinatal mortality (60% of total) occurs in infants born at less than 32 weeks. Mortality and morbidity are inversely associated with gestational age at birth. Morbidities include respiratory distress syndrome, bronchopulmonary dysplasia, intraventricular haemorrhage, necrotizing enterocolitis, sepsis, retinopathy, etc. All members of a family in which a preterm birth occurs suffer greatly, in several aspects, including medically, socially, psychologically, and financially (Berghella 2016).

Description of the intervention

Fetal fibronectin is an extracellular matrix glycoprotein. Fetal fibronectin in biologic fluids is produced by amniocytes and by cytotrophoblast. It is present throughout gestation in all pregnancies. It is not subject to genetic polymorphism. There are very high levels in amniotic fluid (100 µg/mL) in the second trimester, and 30 µg/mL at term. It is localized at the maternal-fetal interface of the amniotic membranes, between chorion and decidua, where it is concentrated in this area between decidua and trophoblast. Here it acts as a 'glue' between the pregnancy and the uterus. Concentration of fetal fibronectin protein found in blood is 1/5th that found in amniotic fluid; it is not present in urine. In normal conditions, this glycoprotein remains in this area between chorion and decidua, and very low levels are found in cervicovaginal secretions after 22 weeks (less than 50 ng/mL). Levels above this value (greater than or equal to 50 ng/mL) at or after 22 weeks in the cervicovaginal secretions collected by a swab have been associated with an increased risk of spontaneous preterm birth (Berghella 2016).

The fetal fibronectin test assesses risk of preterm birth by measuring amount of fetal fibronectin in cervicovaginal secretions. In fact, fetal fibronectin is one of the best predictors of preterm birth in all populations studied so far, including low- and high-risk women without preterm labor, twins, and women in preterm labor (Leitich 1999). The overall sensitivity and specificity are 56% and 84% for preterm before 37 weeks, respectively, but vary according to gestational age at collection, population studied, prevalence of preterm

birth, single versus multiple screening, etc. (Leitich 1999). Its positive predictive value varies from about 9% to 46% depending on the incidence of preterm labor in the population studies (Leitich 1999). Even at 13 to 22 weeks, higher (using 90th percentile) fetal fibronectin levels are associated with a two- to three-fold increase risk in subsequent spontaneous preterm labor.

How the intervention might work

The majority of women presenting with symptoms of preterm labor, such as contractions, back pain, and increase in discharge, do not deliver preterm. Fetal fibronectin has been shown to predict which women would deliver preterm among those presenting with symptoms of preterm labor (threatened preterm labor). By predicting better which women to target interventions such as tocolysis on, fetal fibronectin screening of women with threatened preterm labor could decrease the incidence of preterm birth.

Why it is important to do this review

Preterm birth remains one of the main problems in obstetrics, given its association with perinatal morbidity and mortality, and high costs to care for these sick neonates. Half of preterm birth is preceded by preterm labor. The management of women with threatened preterm labor is controversial, with great difficulty in identifying the subgroup (about 30%) of women presenting with this condition who indeed deliver preterm. Being able to show that screening with fetal fibronectin decreases preterm birth would contribute to potentially save and/or ameliorate millions of lives of neonates worldwide every year. On the contrary, should this study show no benefit from fetal fibronectin screening, this screening, currently used in many units around the world, could be stopped, resulting in significant savings in time and money (Berghella 2016). This is an update of a review first published in 2008.

OBJECTIVES

To assess the effectiveness of management based on knowledge of FFN testing results for preventing preterm birth.

METHODS

Criteria for considering studies for this review

Types of studies

Published and unpublished randomized and quasi-randomized controlled trials. Cluster-randomized trials were also eligible for inclusion, though none were identified for this update. Abstracts were eligible for inclusion if sufficient information was provided to judge the quality and potential for bias of these trials.

Types of participants

Pregnant women between the gestational ages of 22 and 34 weeks screened with fetal fibronectin for risk of preterm birth.

Types of interventions

A screening test such as fetal fibronectin can only be considered effective if interventions based on fetal fibronectin screening results reduce the outcome of preterm birth. Interventions based on fetal fibronectin screening results can also be classified as:

1. interventions based on knowledge of fetal fibronectin results (e.g. fetal fibronectin is collected on all women, but women are randomized so that in 50% of them the result is available to them and the managing obstetrician, while in 50% the fetal fibronectin is blind to them and the managing obstetrician; or fetal fibronectin screening is done only on half of the women);
2. interventions based on positive fetal fibronectin;
3. interventions based on negative fetal fibronectin.

This review focuses exclusively on (1), i.e. interventions based on knowledge of fetal fibronectin results.

Types of outcome measures

Primary outcomes

- (1) Preterm birth (less than 37 weeks)

Secondary outcomes

- (2) Preterm birth less than 34 weeks
- (3) Preterm birth less than 32 weeks
- (4) Preterm birth less than 28 weeks
- (5) Gestational age at delivery
- (6) Birthweight less than 2500 g
- (7) Perinatal death (fetal death and neonatal death)
- (8) Maternal hospitalization
- (9) Tocolysis
- (10) Steroids for fetal maturity
- (11) Time to evaluate (time from arrival to hospital for evaluation of preterm labor to decision regarding admission, discharge, or extended monitoring)
- (12) Respiratory distress syndrome
- (13) Intraventricular haemorrhage
- (14) Necrotizing enterocolitis
- (15) Sepsis
- (16) Neonatal intensive care unit (NICU) admission

(17) NICU days

(18) Maternal well-being (e.g. stress level, etc.)

(19) Economic analysis (cost-effectiveness, cost utility)

We will report outcomes 1, 2, 3, 4, 5, 8, 9, 10, 11, 18, and 19 with 'mothers' as the denominator. We will report outcomes 6, 7, 12, 13, 14, 15, 16, and 17 with 'fetuses/neonates' as the denominator.

Search methods for identification of studies

The following methods section of this review is based on a standard template used by Cochrane Pregnancy and Childbirth.

Electronic searches

For this update, we searched Cochrane Pregnancy and Childbirth's Trials Register by contacting their Information Specialist (7 September 2018).

The Register is a database containing over 24,000 reports of controlled trials in the field of pregnancy and childbirth. It represents over 30 years of searching. For full current search methods used to populate Pregnancy and Childbirth's Trials Register including the detailed search strategies for CENTRAL, MEDLINE, Embase and CINAHL; the list of handsearched journals and conference proceedings; and the list of journals reviewed via the current awareness service; please follow this [link](#).

Briefly, Cochrane Pregnancy and Childbirth's Trials Register is maintained by their Information Specialist and contains trials identified from:

1. monthly searches of the Cochrane Central Register of Controlled Trials (CENTRAL);
2. weekly searches of MEDLINE (Ovid);
3. weekly searches of Embase (Ovid);
4. monthly searches of CINAHL (EBSCO);
5. handsearches of 30 journals and the proceedings of major conferences;
6. weekly current awareness alerts for a further 44 journals plus monthly BioMed Central email alerts.

Search results are screened by two people and the full text of all relevant trial reports identified through the searching activities described above is reviewed. Based on the intervention described, each trial report is assigned a number that corresponds to a specific Pregnancy and Childbirth review topic (or topics), and is then added to the Register. The Information Specialist searches the Register for each review using this topic number rather than keywords. This results in a more specific search set that has been fully accounted for in the relevant review sections ([Included studies](#); [Excluded studies](#); [Ongoing studies](#)).

In addition, we searched [ClinicalTrials.gov](#) and the WHO International Clinical Trials Registry Platform (ICTRP) for unpublished, planned and ongoing trial reports (7 September 2018) using the search methods detailed in [Appendix 1](#).

Searching other resources

We reviewed the reference list of all articles, in particular trials and review articles. We contacted all researchers of included trials to provide actual databases and any pertinent further information. We contacted experts in the field for additional and ongoing trials. We did not apply any language or date restrictions.

Data collection and analysis

For methods used in the previous version of this review, see [Berghella 2008](#).

For this update, the following methods were used for assessing the two trials that were identified as a result of the updated search.

Selection of studies

The two review authors independently assessed for inclusion all the potential studies identified as a result of the search strategy. We resolved any disagreement through discussion. Vincenzo Berghella is a co-author on one of the excluded trials ([Ness 2007](#)) and was not involved in the eligibility assessment for this trial as this was assessed by Gabriele Saccone.

Data extraction and management

We designed a form to extract data. For eligible studies, the two authors extracted the data using the agreed form. We resolved discrepancies through discussion. Data were entered into Review Manager software ([RevMan 2014](#)) and checked for accuracy. When information regarding any of the above was unclear, we planned to contact authors of the original reports to provide further details.

Assessment of risk of bias in included studies

The two authors independently assessed risk of bias for each study using the criteria outlined in the *Cochrane Handbook for Systematic Reviews of Interventions* ([Higgins 2011](#)). Any disagreement was resolved by discussion or by involving a third assessor.

(1) Random sequence generation (checking for possible selection bias)

We described for each included study the method used to generate the allocation sequence in sufficient detail to allow an assessment of whether it should produce comparable groups.

We assessed the method as:

- low risk of bias (any truly random process, e.g. random number table; computer random number generator);
- high risk of bias (any non-random process, e.g. odd or even date of birth; hospital or clinic record number);
- unclear risk of bias.

(2) Allocation concealment (checking for possible selection bias)

We described for each included study the method used to conceal allocation to interventions prior to assignment and assessed whether intervention allocation could have been foreseen in advance of, or during recruitment, or changed after assignment.

We assessed the methods as:

- low risk of bias (e.g. telephone or central randomization; consecutively numbered sealed opaque envelopes);
- high risk of bias (open random allocation; unsealed or non-opaque envelopes, alternation; date of birth);
- unclear risk of bias.

(3.1) Blinding of participants and personnel (checking for possible performance bias)

We described for each included study the methods used, if any, to blind study participants and personnel from knowledge of which intervention a participant received. We considered that studies were at low risk of bias if they were blinded, or if we judged that the lack of blinding was unlikely to affect results. We assessed blinding separately for different outcomes or classes of outcomes.

We assessed the methods as:

- low, high or unclear risk of bias for participants;
- low, high or unclear risk of bias for personnel.

(3.2) Blinding of outcome assessment (checking for possible detection bias)

We described for each included study the methods used, if any, to blind outcome assessors from knowledge of which intervention a participant received. We assessed blinding separately for different outcomes or classes of outcomes.

We assessed methods used to blind outcome assessment as:

- low, high or unclear risk of bias.

(4) Incomplete outcome data (checking for possible attrition bias due to the amount, nature and handling of incomplete outcome data)

We described for each included study, and for each outcome or class of outcomes, the completeness of data including attrition and exclusions from the analysis. We stated whether attrition and exclusions were reported and the numbers included in the analysis at each stage (compared with the total randomized participants), reasons for attrition or exclusion where reported, and whether missing data were balanced across groups or were related to outcomes. Where sufficient information was reported, or could be supplied by the trial authors, we planned to re-include missing data in the analyses which we undertook.

We assessed methods as:

- low risk of bias (e.g. no missing outcome data; missing outcome data balanced across groups);

- high risk of bias (e.g. numbers or reasons for missing data imbalanced across groups; 'as treated' analysis done with substantial departure of intervention received from that assigned at randomization);
- unclear risk of bias.

(5) Selective reporting (checking for reporting bias)

We described for each included study how we investigated the possibility of selective outcome reporting bias and what we found. We assessed the methods as:

- low risk of bias (where it is clear that all of the study's prespecified outcomes and all expected outcomes of interest to the review have been reported);
- high risk of bias (where not all the study's prespecified outcomes have been reported; one or more reported primary outcomes were not prespecified; outcomes of interest are reported incompletely and so cannot be used; study fails to include results of a key outcome that would have been expected to have been reported);
- unclear risk of bias.

(6) Other bias (checking for bias due to problems not covered by (1) to (5) above)

We described for each included study any important concerns we had about other possible sources of bias.

(7) Overall risk of bias

We made explicit judgements about whether studies were at high risk of bias, according to the criteria given in the *Handbook* (Higgins 2011). With reference to (1) to (6) above, we planned to assess the likely magnitude and direction of the bias and whether we considered it is likely to impact on the findings.

Assessment of the quality of the evidence using the GRADE approach

For this update, the quality of the evidence was assessed using the GRADE approach as outlined in the [GRADE handbook](#) in order to assess the quality of the body of evidence relating to the following outcomes for the main comparison (FFN knowledge versus no knowledge):

1. Preterm birth < 37 weeks
2. Preterm birth < 34 weeks
3. Preterm birth < 32 weeks
4. Maternal hospitalization

GRADEpro Guideline Development Tool was used to import data from Review Manager 5.3 (RevMan 2014) in order to create a 'Summary of findings' table. A summary of the intervention effect and a measure of quality for each of the above outcomes was produced using the GRADE approach. The GRADE approach uses five considerations (study limitations, consistency of effect,

imprecision, indirectness, and publication bias) to assess the quality of the body of evidence for each outcome. The evidence can be downgraded from 'high quality' by one level for serious (or by two levels for very serious) limitations, depending on assessments for risk of bias, indirectness of evidence, serious inconsistency, imprecision of effect estimates, or potential publication bias.

Measures of treatment effect

Dichotomous data

For dichotomous data, we presented results as summary risk ratio with 95% confidence intervals.

Continuous data

We used the mean difference if outcomes were measured in the same way between trials. In future updates, if necessary, we will use the standardised mean difference to combine trials that measured the same outcome, but use different methods.

Unit of analysis issues

Cluster-randomized trials

We planned to include cluster-randomized trials in the analyses along with individually-randomized trials, however all included studies were individually-randomized trials.

Cross-over trials

We planned to exclude cross-over trials. However, no cross-over trials were found during the search process.

Dealing with missing data

For included studies, levels of attrition were noted. In future updates, if more eligible studies are included, the impact of including studies with high levels of missing data in the overall assessment of treatment effect will be explored by using sensitivity analysis.

For all outcomes, analyses were carried out, as far as possible, on an intention-to-treat basis, i.e. we attempted to include all participants randomized to each group in the analyses. The denominator for each outcome in each trial was the number randomized minus any participants whose outcomes were known to be missing.

Assessment of heterogeneity

We assessed statistical heterogeneity in each meta-analysis using the Tau², I² and Chi² statistics. We regarded heterogeneity as substantial if I² was greater than 30% and either Tau² was greater than zero, or there was a low P value (less than 0.10) in the Chi² test for heterogeneity. If we identified substantial heterogeneity (I² above 30%), we planned to explore it by prespecified subgroup analysis.

Assessment of reporting biases

In future updates, if there are 10 or more studies in the meta-analysis, we will investigate reporting biases (such as publication bias) using funnel plots. We will assess funnel plot asymmetry visually. If asymmetry is suggested by a visual assessment, we will perform exploratory analyses to investigate it.

Data synthesis

We carried out statistical analysis using the Review Manager software (RevMan 2014). We used fixed-effect meta-analysis for combining data where it was reasonable to assume that studies were estimating the same underlying treatment effect: i.e. where trials were examining the same intervention, and the trials' populations and methods were judged sufficiently similar.

If there was clinical heterogeneity sufficient to expect that the underlying treatment effects differed between trials, or if substantial statistical heterogeneity was detected, we used random-effects meta-analysis to produce an overall summary if an average treatment effect across trials was considered clinically meaningful. The random-effects summary will be treated as the average range of possible treatment effects and we will discuss the clinical implications of treatment effects differing between trials. If the average treatment effect is not clinically meaningful, we will not combine trials. If we used random-effects analyses, the results were presented as the average treatment effect with 95% confidence intervals, and the estimates of Tau² and I².

Subgroup analysis and investigation of heterogeneity

Had we identified substantial heterogeneity, we planned to investigate it using subgroup analyses and to consider whether an overall summary was meaningful, and if it was, to use a random-effects analysis to produce it.

We planned to restrict subgroup analyses to primary outcomes for the following subgroups:

1. Symptomatic women with threatened preterm labor versus asymptomatic women
2. Women with multiple gestations versus women with singleton gestations

We also planned to assess subgroup differences by interaction tests available within RevMan (RevMan 2014) and to report the results of subgroup analyses quoting the Chi² statistic and P value, and the interaction test I² value.

Given that all included trials enrolled only singleton gestations with threatened preterm labor, no subgroup analyses were performed.

Sensitivity analysis

We carried out sensitivity analyses for the primary outcome to explore the impact of trial quality assessed by concealment of allocation, high attrition rates, or both. Poor-quality studies were excluded from the analyses in order to assess whether this made any difference to the overall result.

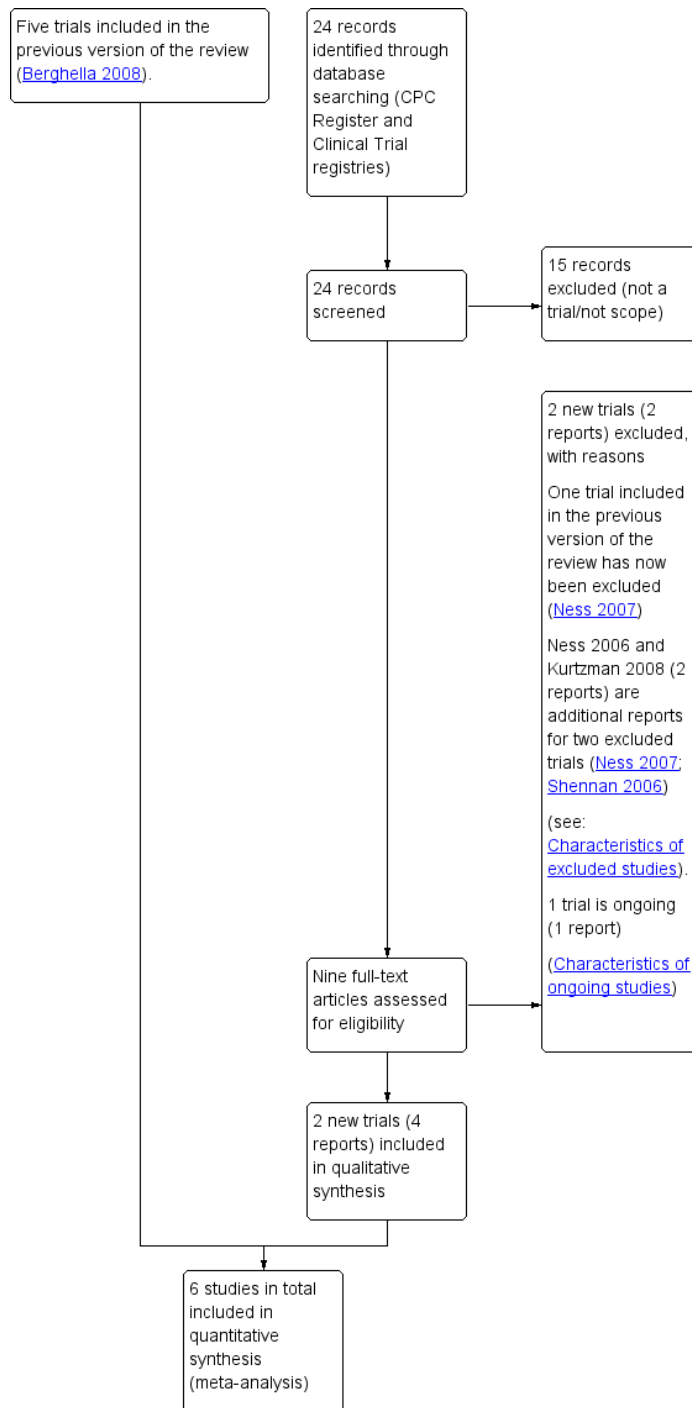
RESULTS

Description of studies

Results of the search

See: [Figure 1](#)

Figure 1. Study flow diagram.



For this update, we identified nine trial reports to assess. We included two new trials (Dutta 2011; Lee 2013) (four reports). We excluded two new trials (Gallot 2018; Osório 2010) (two reports) and added one trial report to ongoing (NCT01431885) (one report). Two reports were additional reports for two excluded trials (Ness 2007; Shennan 2006). One trial previously included has been excluded in this update (Ness 2007).

All women in the included studies had singleton gestations with threatened preterm labor (PTL) between 22 and 34 weeks. There were no trials comparing knowledge to no knowledge of fetal fibronectin (FFN) in asymptomatic women or women with multiple gestations. We identified no quasi-randomized trials.

Included studies

The six included studies randomized 546 women, of which 277 were randomized to knowledge and 269 to no knowledge of FFN.

Settings

All trials were based in hospital settings. One trial took place in the United Kingdom (Dutta 2011); all the other trials took place in the United States. Dutta 2011 and Plaut 2003 were multicenter and the remaining four trials were based in single hospitals.

Participants

All trials included women with singleton gestations and threatened PTL from around 23 to around 35 weeks' gestation. All the trials excluded multiple pregnancies apart from Plaut 2003. We requested patient-level databases from all authors, and obtained them from Grobman 2004, Lee 2013, Lowe 2004, and Nguyen 2002. Dutta 2011 and Plaut 2003 also replied to our query, but had no additional data to what they had already published. Our analysis was based only on singleton gestations. We avoided repeated entries of the same patients, which was possible since we had patient-level data for the one trial where this occurred (Plaut 2003).

Interventions and comparisons

Of 546 women, 277 were randomized to knowledge of FFN and 269 to no knowledge of FFN. Intervention groups had the FFN

following signs of PTL. Time taken to obtain results was not reported in most trials. In Lowe 2004, it took less than one hour, and Plaut 2003 took between one and two hours. Dutta 2011 and Nguyen 2002 did not perform FFN on women in the control group. The other trials performed FFN on participants in the control group but did not disclose the results. Only Lee 2013 and Nguyen 2002 had a protocol for positive FFN tests.

Funding

Two trials were supported by affiliated universities (Dutta 2011; Lowe 2004), and two trials did not disclose their funding sources (Lee 2013; Nguyen 2002). Two were sponsored by Adeza Biomedical which produced the FFN tests used in the trials (Grobman 2004; Plaut 2003).

Declarations of interest

Dutta 2011 and Lee 2013 declared that no authors had any conflicts of interest. None of the other trials reported whether or not there were conflicts of interest.

Excluded studies

We excluded seven of the 11 excluded studies because they only included women with positive (six trials) (Andrews 2001; Bisits 2004; Goldenberg 2001; Hauth 2001; ISRCTN43735180; Shennan 2006) or only with negative (one trial) (Elliott 2005) FFN results. One excluded study (Kalchbrenner 1999) compared speculum versus digital collection of FFN, with no management reported based on FFN results. Another study was excluded because it used different diagnostic tests, and only partly used FFN (Gallot 2018) and one study was published only as an abstract with insufficient information provided to judge the quality and risk of bias (Osório 2010). Another excluded study used mainly transvaginal ultrasound cervical length, and not FFN, in the algorithm studied (Ness 2007).

Risk of bias in included studies

See Figure 2; Figure 3.

Figure 2. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

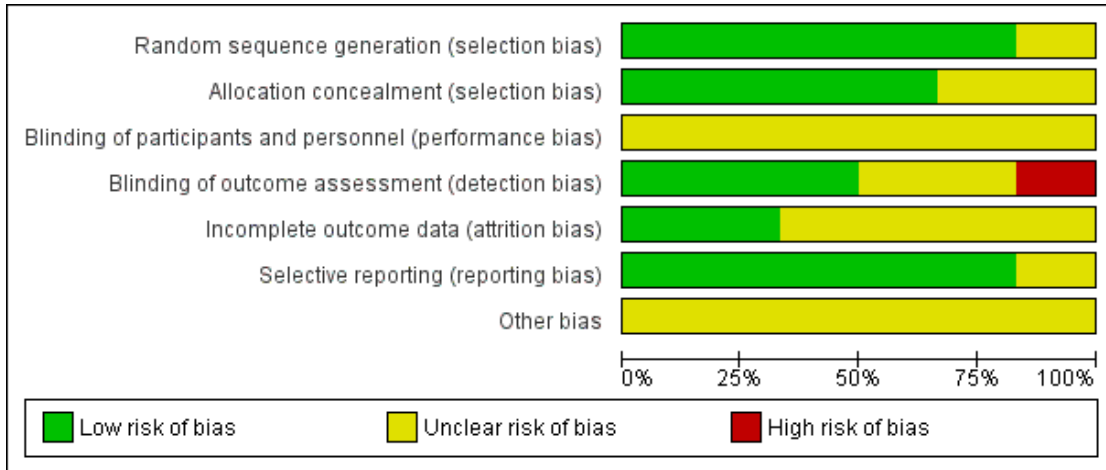


Figure 3. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Dutta 2011	+	?	?	?	?	+	?
Grobman 2004	+	+	?	+	+	+	?
Lee 2013	+	+	?	+	+	+	?
Lowe 2004	+	+	?	-	?	+	?
Nguyen 2002	?	?	?	?	?	?	?
Plaut 2003	+	+	?	+	?	+	?

Allocation

Risk of selection bias was considered low in four of the six included trials. These four trials used computer-generated random sequence generation and sealed opaque envelopes to conceal allocation (Grobman 2004; Lee 2013; Lowe 2004; Plaut 2003). One study (Dutta 2011) used telephonic random number generation but did not describe allocation concealment. The sixth study (Nguyen 2002) did not describe random sequence generation or allocation concealment (only the abstract of this study has been published).

Blinding

Women and staff were aware of the intervention they were allocated to due to its nature but it is unclear if this affected results. Laboratory personnel who performed the FFN test were blinded to women's characteristics and outcomes in Grobman 2004, Lee 2013, and Plaut 2003. One trial was unblinded (Lowe 2004). It was unclear in the remaining trials whether outcome assessors were blinded.

Incomplete outcome data

Two studies were assessed as having low risk of attrition bias because they reported no loss to follow-up (Grobman 2004; Lee 2013). In the remaining four trials, there was insufficient detail in the reports to permit a judgement of low or high risk and so these trials were assessed as being at unclear risk (Dutta 2011; Lowe 2004; Nguyen 2002; Plaut 2003). See [Characteristics of included studies](#) for further details.

Selective reporting

Risk of reporting bias was considered low in five of the six included studies; no deviations from original protocols were noted. The sixth study (Nguyen 2002) did not describe reporting details (only the abstract of this study has been published).

Other potential sources of bias

The included studies did not have enough information to adequately assess the presence of other forms of bias and, as such, were deemed to have an unclear risk of bias. Nguyen 2002 was published only as abstract. Two studies (Grobman 2004; Plaut 2003) were sponsored by Adeza Biomedical which manufactures the FFN tests, however the results from these studies did not seem to be skewed favourably toward the FFN tests.

Effects of interventions

See: [Summary of findings for the main comparison Main comparison \(FFN knowledge versus no knowledge\)](#)

See: [Summary of findings for the main comparison](#) the comparison (i.e. FFN knowledge versus no knowledge) and for selected important outcomes, we graded the quality of the evidence and created 'Summary of findings' tables. For all these outcomes, the evidence was graded as mainly low quality due to the imprecision of effect estimates, limitations in study design, and inconsistency of results.

We included six trials (involving 546 women) in this review. All women included had singleton gestations and threatened PTL.

Fetal fibronectin knowledge versus no knowledge

Primary outcome

Preterm birth before 37 weeks appeared to favour management based on knowledge of FFN results (20.7%) compared to controls without such knowledge (29.2%) though the confidence intervals (CIs) just crossed the line of no effect so we could not be certain of this effect (risk ratio (RR) 0.72, 95% CI 0.52 to 1.01; 5 trials; 434 women; low-quality evidence; [Analysis 1.1](#)). Three out of five trials had RRs less than one, with no heterogeneity.

We performed a sensitivity analysis for the primary outcome, by removing Nguyen 2002 which was assessed to be at unclear risk of allocation concealment. Removing this data from the analysis indicated that knowledge of FFN may result in fewer preterm births before 37 weeks (19.4% compared to 29.9%) with CI below the line of no effect (RR 0.67, 95% CI 0.46 to 0.97; 4 trials; 357 women; [Analysis 2.1](#)).

Secondary outcomes

Effects on other secondary outcomes are uncertain due to serious concerns in study design, inconsistency, and imprecision of effect estimates:

- preterm birth less than 34 weeks (RR 1.09, 95% CI 0.54 to 2.18; 4 trials; 357 women; low-quality evidence; [Analysis 1.2](#))
- preterm birth less than 32 weeks (average RR 0.79, 95% CI 0.16 to 3.96; 4 trials; 357 women; $\text{Tau}^2 = 0.93$; $I^2 = 46\%$; very-low quality evidence; [Analysis 1.3](#))
- preterm birth less than 28 weeks (RR 0.63, 95% CI 0.15 to 2.59; 4 trials; 357 women; [Analysis 1.4](#))
- gestational age at delivery (weeks) (mean difference (MD) 0.14, 95% CI -0.36 to 0.63; 5 trials; 456 neonates; [Analysis 1.5](#))

- birthweight less than 2500 g (no events in either group; one trial; 68 neonates; [Analysis 1.6](#))
- perinatal death (RR 2.21, 95% CI 0.09 to 52.27; 2 trials; 164 neonates; [Analysis 1.7](#))
- maternal hospitalization (RR 1.06, 95% CI 0.79 to 1.43; 5 trials; 441 women; low-quality evidence; [Analysis 1.8](#))
- tocolysis (RR 0.97, 95% CI 0.75 to 1.24; 6 trials; 531 women; [Analysis 1.9](#))
- steroids for fetal lung maturity (RR 1.04, 95% CI 0.79 to 1.38; 5 trials; 441 neonates; [Analysis 1.10](#))
- time to evaluate (time from arrival to hospital for evaluation of PTL to decision regarding admission, discharge, or extended monitoring (hours)) (MD 0.55, 95% CI -0.39 to 1.50; random-effects; 6 trials; 528 women; $\text{Tau}^2 = 0.55$; $I^2 = 52\%$; [Analysis 1.11](#))
 - respiratory distress syndrome (RR 0.91, 95% CI 0.06 to 14.06; 2 trials; 148 neonates; [Analysis 1.12](#))
 - NICU admission (RR 2.36, 95% CI 0.92 to 6.07; 2 trials; 124 neonates; [Analysis 1.13](#))
 - economic analysis (hospitalization charges): [Nguyen 2002](#) reported data regarding hospitalization charges and found that management based on FFN test required higher hospitalization charges (US dollars) (MD 153.00, 95% CI 24.01 to 281.99; [Analysis 1.14](#)).

Intraventricular haemorrhage, necrotizing enterocolitis, sepsis, NICU days, and maternal well-being were not reported in any trial.

DISCUSSION

Summary of main results

Management based on knowledge of FFN results may reduce preterm birth before 37 weeks (21.6%) versus controls without such knowledge (29.2%). However, management based on knowledge of FFN results may make little or no difference to preterm birth before 34 weeks or maternal hospitalization. The evidence for preterm birth before 32 weeks is uncertain because the quality was found to be very low.

For all other outcomes, for which there were available data (preterm birth less than 28 weeks; gestational age at delivery (weeks); birthweight less than 2500 g; perinatal death; tocolysis; steroids for fetal lung maturity; time to evaluate; respiratory distress syndrome; neonatal intensive care unit (NICU) admission; and NICU days), knowledge of FFN results may make little or no difference to the outcomes.

Therefore, further research is necessary before fetal fibronectin testing can be routinely recommended. There are no randomized trials comparing knowledge versus no knowledge of fetal fibronectin

in asymptomatic pregnant women, or in women with multiple gestations.

Overall completeness and applicability of evidence

We attempted to be as inclusive as possible in the search strategy. Nonetheless, the studies reported are predominantly from the US, which may limit the external validity of these results. The evidence from the included studies is also based on the cutoff for the fFN bedside test, also known as Quickchack fFN (50 ng/L). It should be noted, that the use of quantitative fFN which has since become available, may affect any future conclusions of this review.

It is still unclear which interventions are most beneficial once fetal fibronectin results are known. The Ness positive study ([Ness 2007](#)) reported a detailed protocol of management based on cervical length mainly (and fetal fibronectin results for a small subgroup), and could be replicated in further research trials. Our review did not include by design assessment of effectiveness of interventions based on positive fetal fibronectin testing, or negative fetal fibronectin testing. We identified no trials on women without signs or symptoms of labor.

Quality of the evidence

For the main comparison (fetal fibronectin knowledge versus no knowledge) and for most important outcomes (preterm birth less than 37 weeks; preterm birth less than 34 weeks; preterm birth less than 32 weeks; maternal hospitalization), we graded the quality of evidence using the GRADE approach. For most of these outcomes, the evidence was graded as low-quality mainly due to the imprecision of effect estimates and limitations in study design. We graded preterm birth less than 32 weeks as very low-quality due to evidence of inconsistency in results, in addition to imprecision of effect estimates and limitations in study design ([Summary of findings for the main comparison](#)).

Potential biases in the review process

We acknowledge that our analysis of preterm delivery is flawed as it was analyzed as four separate outcomes (< 37, < 34, < 32, < 28 weeks); and clearly these outcomes are not independent. A much better analysis would be to use the time to birth for each participant, but that would require individual patient data (IPD). This approach is not feasible without the availability of IPD. Vincenzo Berghella is a co-author on one of the excluded trials ([Ness 2007](#)) and was not involved in the eligibility assessment for this trial as this was assessed by Gabriele Saccone.

Agreements and disagreements with other studies or reviews

The findings of this review largely agree with our prior Cochrane Review and other reviews on this topic, with now additional studies and larger sample size (Berghella 2008; Berghella 2016).

AUTHORS' CONCLUSIONS

Implications for practice

The evidence from this review suggests that management based on knowledge of FFN results may reduce preterm birth before 37 weeks. However, our confidence in this result is limited as the evidence was found to be of low-quality. Effects on other substantive outcomes are uncertain due to serious concerns in study design, inconsistency, and imprecision of effect estimates. No trials were identified on asymptomatic women, or multiple gestations. Currently, the only management protocol for screening of women with threatened preterm labor shown by randomized trial data to decrease preterm birth has been that based mainly on transvaginal ultrasound cervical length, with fetal fibronectin only in women with cervical length 20 mm to 29 mm (Berghella 2016; Ness 2007).

Implications for research

Given that the evidence suggests that there may be a reduction of

around 28% in the primary outcome, further research could be encouraged, minimizing attrition bias, to better understand whether and under what circumstances the predictive characteristics of the fetal fibronectin test can be translated into better clinical management. Future studies could include specific populations (e.g. singleton gestations with symptoms of preterm labor), a study group managed with a protocol based on the fetal fibronectin results, and report not only maternal but also significant perinatal outcomes. Cost-effectiveness analyses are also needed.

ACKNOWLEDGEMENTS

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* *Indicates the major publication for the study*

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Dutta 2011

Methods	Randomized controlled trial
Participants	24 0/7-34 6/7 weeks singletons presenting to the hospital for the primary reason of uterine activity. Twins excluded. Number of participants = 93 (100 enrolled, but 7 excluded because they did not meet the inclusion criteria of which 4 were twins)
Interventions	FFN testing (44) versus no testing (44) Time FFN results available: unknown Protocol for FFN knowledge group: no
Outcomes	Primary: hospital admission
Notes	Unpublished data were not obtainable. Setting: 2 hospitals, Scotland Dates: December 2007 - March 2009 Conflict of interest: none Funding source: the trial is registered with the University of Edinburgh which is a charitable body, registered in Scotland, with registration number SC005336

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Telephonic randomization
Allocation concealment (selection bias)	Unclear risk	Not described
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not described
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not described
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	It is reported that those that did not follow the protocol were excluded from the study. No study flow diagram. In methods reports that 100 patients were enrolled, and 93 were available for analysis - 44 in the control group and 49 in the test group - but then denominator results differ - sometimes both groups reported as 44 and then in other places reports 49

Dutta 2011 (Continued)

Selective reporting (reporting bias)	Low risk	The trial was registered and there were no deviations from the original protocol in the final publication
Other bias	Unclear risk	Insufficient reporting to determine presence of other form of bias

Grobman 2004

Methods	Randomized controlled trial
Participants	24-34 week singletons with > 6 contractions/hour. Twins excluded Number of participants: 100 (50/50: knowledge/no knowledge)
Interventions	FFN knowledge or not Time FFN results available: unknown Protocol for FFN knowledge group: no
Outcomes	Primary: total costs
Notes	Intention-to-treat; only singletons; no protocol Positive FFN test in each group: 5 (10%) in knowledge, 3 (6%) in no knowledge group Setting: hospital, United States Dates: "12 month period" dates unclear Conflict of interest: not mentioned Funding source: supported by a grant from Adeza Biomedical, Sunnyvale, Calif

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated
Allocation concealment (selection bias)	Low risk	Computer-generated, opaque envelopes
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not reported
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Laboratory personnel who performed the FFN test were blinded to women's characteristics and outcomes
Incomplete outcome data (attrition bias) All outcomes	Low risk	No loss to follow-up. One woman randomized to the knowledge group - her results were not available because of problems with the machine - however it is reported that outcomes for this patient were analysed in the group to which she was randomized, suggesting they adhered

Grobman 2004 (Continued)

		to intention to treat analysis
Selective reporting (reporting bias)	Low risk	The trial was registered and there were no deviations from the original protocol in the final publication
Other bias	Unclear risk	Insufficient reporting to determine presence of other form of bias

Lee 2013

Methods	Randomized controlled trial
Participants	24 0/7-33 6/7 weeks singletons with symptoms suggesting PTL Number of participants: 76 (44/32: knowledge/no knowledge)
Interventions	FFN knowledge versus no FFN knowledge Time FFN results available: unknown Protocol for FFN knowledge group: yes
Outcomes	Time required to evaluate participants in triage in Labor and Delivery
Notes	Setting: medical centre, United States Dates: September 2006 - December 2010 Conflict of interest: none Funding source: not mentioned

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated
Allocation concealment (selection bias)	Low risk	Computer-generated
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not reported
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Laboratory personnel who performed the FFN test were blinded to women's characteristics and outcomes
Incomplete outcome data (attrition bias) All outcomes	Low risk	No loss to follow-up
Selective reporting (reporting bias)	Low risk	The trial was registered and there were no deviations from the original protocol in the final publication

Lee 2013 (Continued)

Other bias	Unclear risk	Insufficient reporting to determine presence of other form of bias
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Lowe 2004

Methods	Randomized controlled trial
Participants	23-34 week singletons/twins with contractions +/- cervical change Number of participants: 97 [89 singletons/8 twins] (46/51: knowledge/no knowledge)
Interventions	FFN knowledge or not Time FFN results available: < 1 hour Protocol for FFN knowledge group: no
Outcomes	Primary: length of stay
Notes	13/110 excluded post-randomization: unclear whether intention-to-treat; included twins; no protocol Positive FFN test in each group: 11 (24%) in knowledge group; FFN test not performed in no knowledge group Setting: hospital, United States Dates: August 2000 - May 2002 Conflict of interest: not mentioned Funding source: supported by Process Improvement Grant, University of Iowa

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated
Allocation concealment (selection bias)	Low risk	Computer-generated, opaque envelopes, blocks of 10
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Unblinded intervention
Blinding of outcome assessment (detection bias) All outcomes	High risk	Unblinded intervention
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	110 were enrolled into the study, but 13 were not included (reasons were documented - but did not present characteristics of these women or make it clear which group they had been randomized to) and so 97 women were available for the analysis. It is not clear exactly how many were initially randomised

Lowe 2004 (Continued)

Selective reporting (reporting bias)	Low risk	No deviation from the original protocol
Other bias	Unclear risk	Insufficient reporting to determine presence of other form of bias

Nguyen 2002

Methods	Randomized controlled trial
Participants	24-35 weeks (unclear if twins included) with symptoms of PTL Number of participants: 77 (42/35: knowledge/no knowledge)
Interventions	FFN testing done (and knowledge available) versus not done (so knowledge not available) Time FFN results available: unknown Protocol for FFN knowledge group: yes
Outcomes	Primary: cost-effectiveness
Notes	Abstract only; unclear if intention-to-treat, twins included, protocol, etc Positive FFN test in each group: not available in knowledge group; FFN not performed in no knowledge group Setting: hospital, United States Dates: not specified Conflict of interest: not mentioned Funding source: not mentioned

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No description available (only abstract published)
Allocation concealment (selection bias)	Unclear risk	No description available (only abstract published)
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	No description available (only abstract published)
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No description available (only abstract published)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Only abstract available - all enrolled appear to have been randomized
Selective reporting (reporting bias)	Unclear risk	No protocol available

Other bias	Unclear risk	No description available (only abstract published)
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Plaut 2003

Methods	Randomized controlled trial
Participants	24-34 6/7 week singletons/twins with 'symptoms of PTL' Number of participants: 114 (unclear from text) randomized [96 singleton, 12 twins] (analysed: 51/57: knowledge/no knowledge)
Interventions	FFN knowledge or not Time FFN results available: 1-2 hours Protocol for FFN knowledge group: no
Outcomes	Primary: transport rates (not reported)
Notes	6/114 excluded post-randomization: not intention-to-treat - 6 patients were excluded; 8 participants entered in study twice; stopped trial prematurely; sponsored by Adeza; included twins; no protocol Positive FFN test in each group: 6 (12%) in knowledge, 6 (10.5%) in no knowledge group Setting: 4 community hospitals, United States Dates: September 2000 - December 2001 Conflict of interest: not mentioned Funding source: supported by Adeza Biomedical, Sunnyvale, Calif

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated
Allocation concealment (selection bias)	Low risk	Computerized opaque envelopes
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not reported
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Laboratory personnel who performed the FFN test were blinded to women's characteristics and outcomes
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Consent obtained from 114 patients, 6 patients were excluded (reasons given), this left 108 swabs from 100 different patients - 8 patients were entered into the study twice. Results are presented for 108 swabs. Not clear number originally randomized into each group

Plaut 2003 (Continued)

Selective reporting (reporting bias)	Low risk	No deviation from the original protocol
Other bias	Unclear risk	Insufficient reporting to determine presence of other form of bias

FFN: fetal fibronectin

ITT: intention to treat

PTL: preterm labor

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Andrews 2001	Only women with positive FFN were included.
Bisits 2004	Only women with positive FFN were included. They were randomized to tocolysis or not
Elliott 2005	Only women with negative FFN were included.
Gallot 2018	Registration protocol of a trial using different diagnostic tests, and only in part on FFN
Goldenberg 2001	Only women with positive FFN were included (secondary analysis of NICHD BV/TV study)
Hauth 2001	Only women with positive FFN were included.
ISRCTN43735180	Only women with positive FFN were included.
Kalchbrenner 1999	This was a trial comparing speculum versus digital collection of FFN. There were no data regarding the results of the FFN test, and no intervention was reported based on FFN results
Ness 2007	The algorithm used was based mostly on cervical length results, and only in part (for cervical lengths 20-29 mm) on FFN
Osório 2010	Published only as abstract with no data available on methods or the outcomes of interest
Shennan 2006	Only women with positive FFN were included.

FFN: fetal fibronectin

Characteristics of ongoing studies *[ordered by study ID]*

[NCT01431885](#)

Trial name or title	2 methods of diagnosing preterm labor
Methods	Randomized controlled trial
Participants	Symptomatic women with more than 6 uterine contractions per hour
Interventions	Symptomatic preterm labor patients will be randomized to diagnosis of preterm labor by serial digital examination versus an algorithm incorporating transvaginal ultrasound measurement of cervical length and vaginal fetal fibronectin
Outcomes	Preterm birth < 37 weeks
Starting date	2011
Contact information	Conrad Chao; 001-559-499-6548; cchao@fresno.ucsf.edu
Notes	

DATA AND ANALYSES

Comparison 1. FFN knowledge versus no knowledge

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Preterm birth < 37 weeks	5	434	Risk Ratio (M-H, Fixed, 95% CI)	0.72 [0.52, 1.01]
2 Preterm birth < 34 weeks	4	357	Risk Ratio (M-H, Fixed, 95% CI)	1.09 [0.54, 2.18]
3 Preterm birth < 32 weeks	4	357	Risk Ratio (M-H, Random, 95% CI)	0.79 [0.16, 3.96]
4 Preterm birth < 28 weeks	4	357	Risk Ratio (M-H, Fixed, 95% CI)	0.63 [0.15, 2.59]
5 Gestational age at delivery	5	456	Mean Difference (IV, Fixed, 95% CI)	0.14 [-0.36, 0.63]
6 Birthweight < 2500 g	1	68	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
7 Perinatal death (fetal death and neonatal death)	2	164	Risk Ratio (M-H, Fixed, 95% CI)	2.21 [0.09, 52.27]
8 Maternal hospitalization	5	441	Risk Ratio (M-H, Fixed, 95% CI)	1.06 [0.79, 1.43]
9 Tocolysis	6	531	Risk Ratio (M-H, Fixed, 95% CI)	0.97 [0.75, 1.24]
10 Steroids for fetal lung maturity	5	441	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.79, 1.38]
11 Time to evaluate	6	528	Mean Difference (IV, Random, 95% CI)	0.55 [-0.39, 1.50]
12 Respiratory distress syndrome	2	148	Risk Ratio (M-H, Fixed, 95% CI)	0.91 [0.06, 14.06]
13 Neonatal intensive care unit admission	2	124	Risk Ratio (M-H, Fixed, 95% CI)	2.36 [0.92, 6.07]
14 Economic analysis (hospitalization charges)	1	77	Mean Difference (IV, Fixed, 95% CI)	153.0 [24.01, 281.99]

Comparison 2. FFN knowledge versus no knowledge: sensitivity analysis

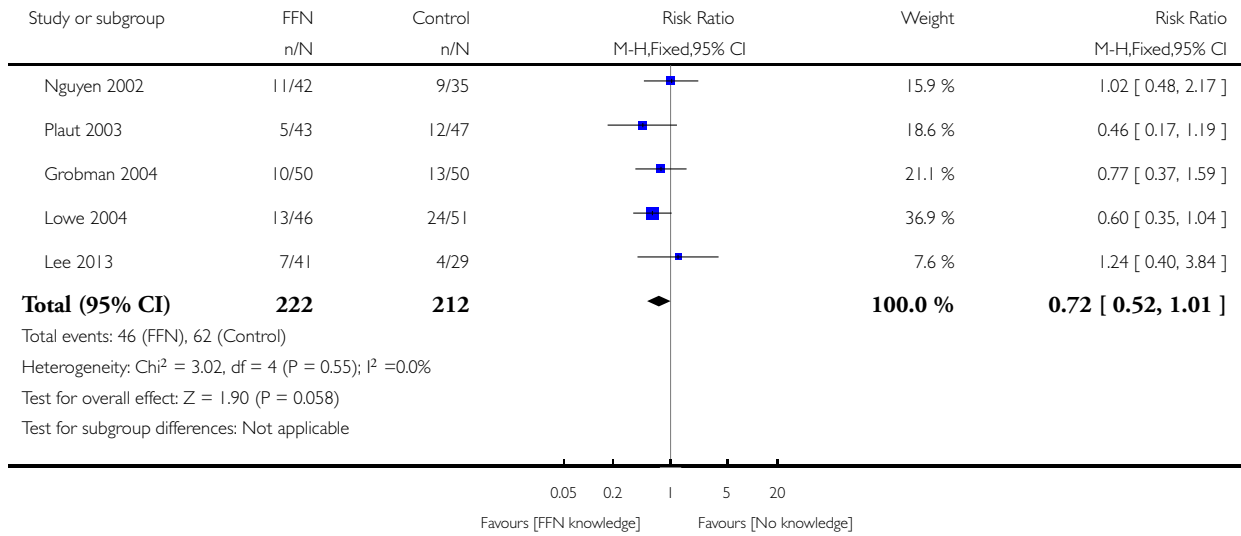
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Preterm birth < 37 weeks	4	357	Risk Ratio (M-H, Fixed, 95% CI)	0.67 [0.46, 0.97]

Analysis 1.1. Comparison 1 FFN knowledge versus no knowledge, Outcome 1 Preterm birth < 37 weeks.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 1 Preterm birth < 37 weeks

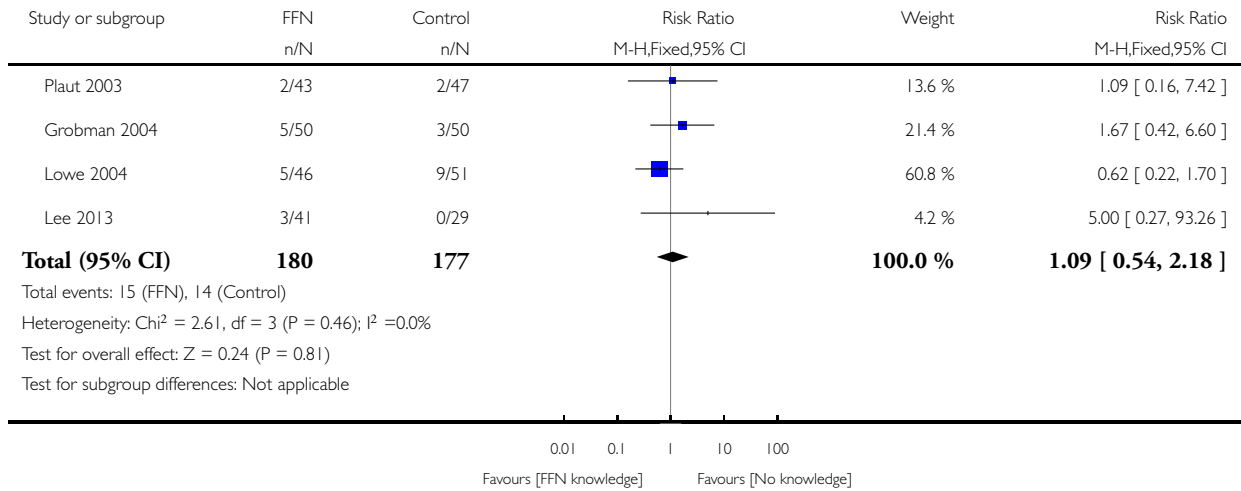


Analysis 1.2. Comparison 1 FFN knowledge versus no knowledge, Outcome 2 Preterm birth < 34 weeks.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 2 Preterm birth < 34 weeks

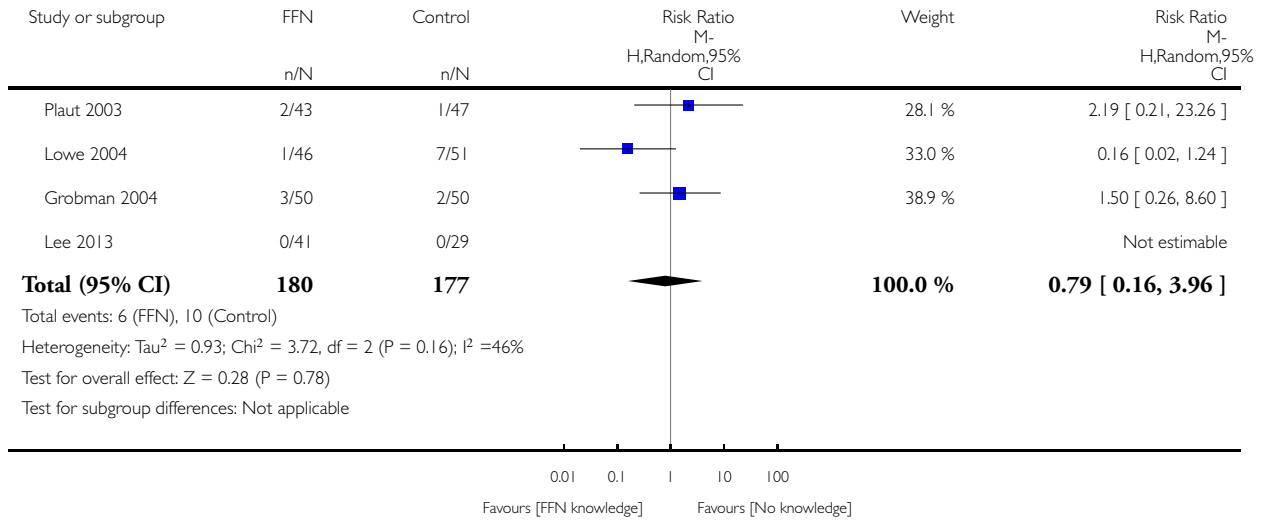


Analysis 1.3. Comparison 1 FFN knowledge versus no knowledge, Outcome 3 Preterm birth < 32 weeks.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 3 Preterm birth < 32 weeks

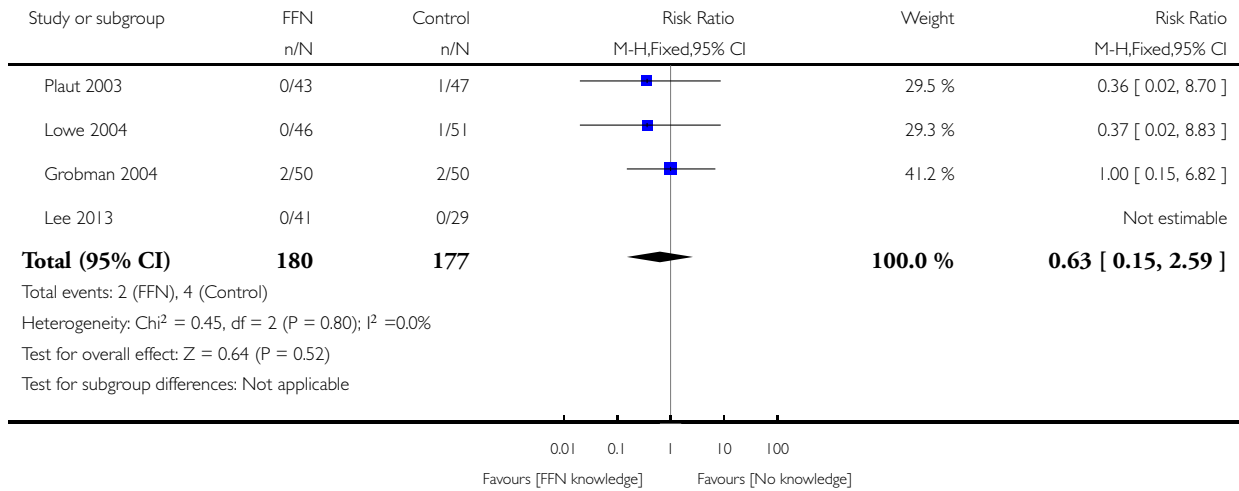


Analysis 1.4. Comparison 1 FFN knowledge versus no knowledge, Outcome 4 Preterm birth < 28 weeks.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 4 Preterm birth < 28 weeks

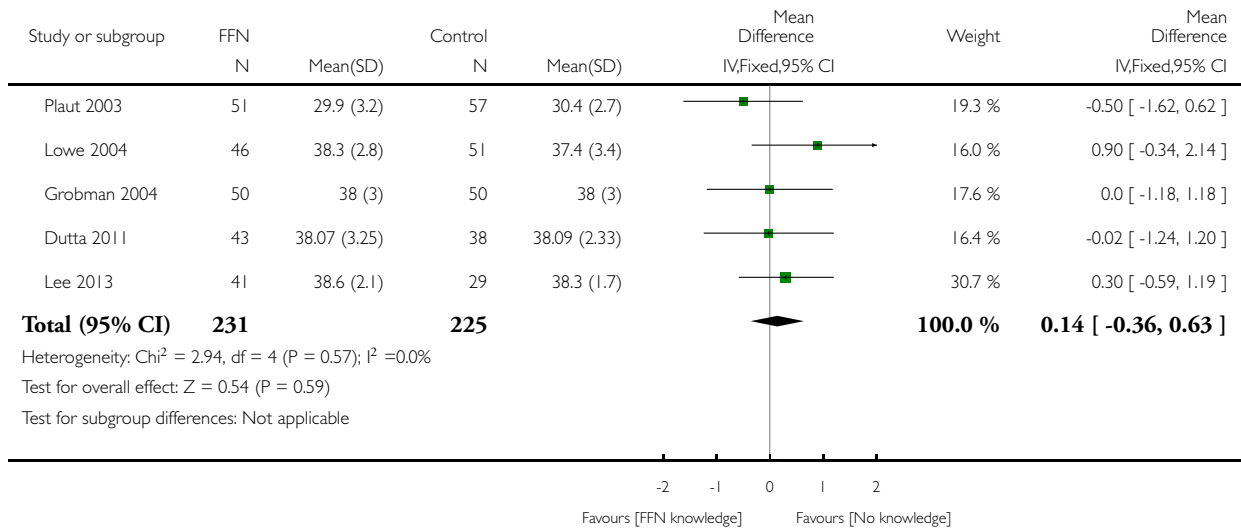


Analysis 1.5. Comparison 1 FFN knowledge versus no knowledge, Outcome 5 Gestational age at delivery.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 5 Gestational age at delivery

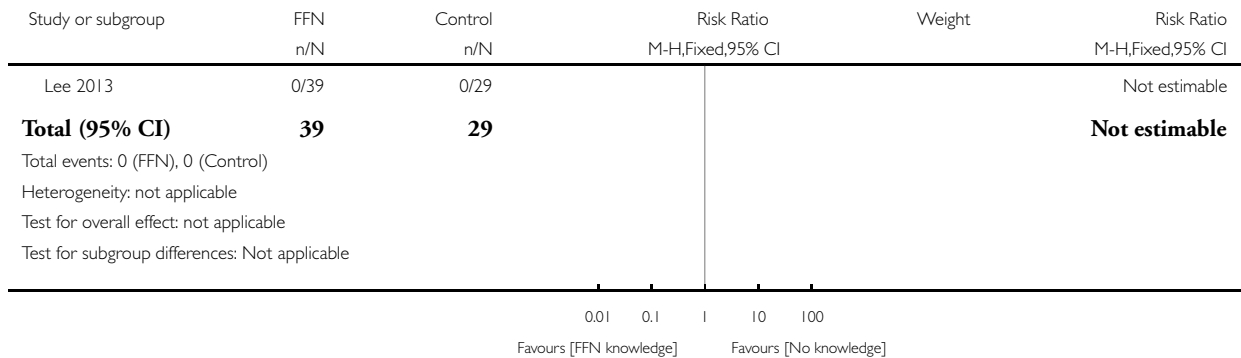


Analysis 1.6. Comparison 1 FFN knowledge versus no knowledge, Outcome 6 Birthweight < 2500 g.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 6 Birthweight < 2500 g

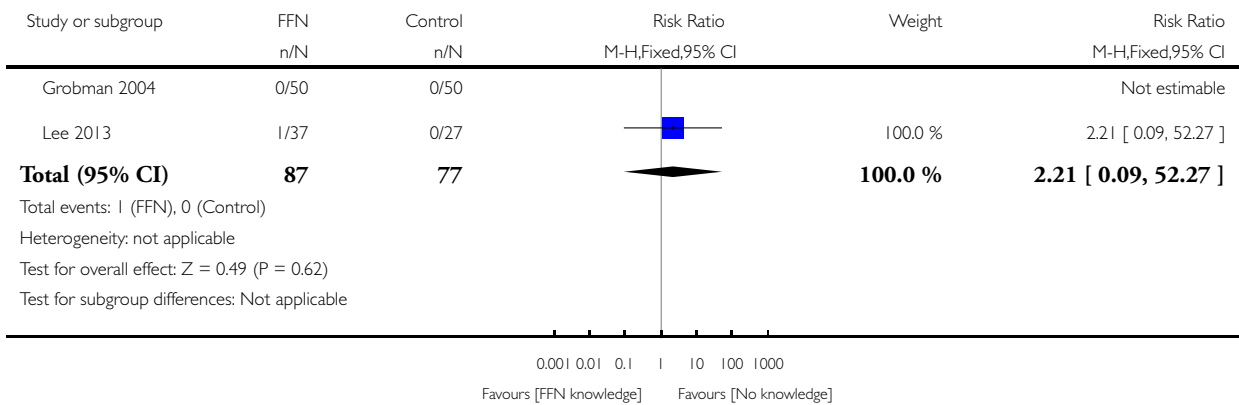


Analysis 1.7. Comparison 1 FFN knowledge versus no knowledge, Outcome 7 Perinatal death (fetal death and neonatal death).

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 7 Perinatal death (fetal death and neonatal death)

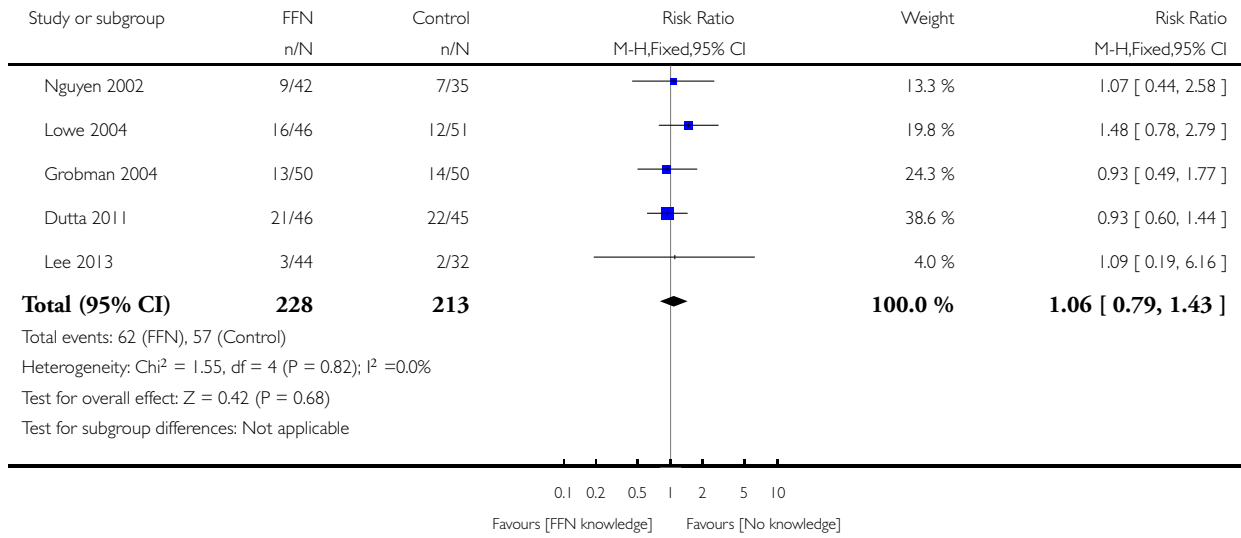


Analysis 1.8. Comparison 1 FFN knowledge versus no knowledge, Outcome 8 Maternal hospitalization.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 8 Maternal hospitalization

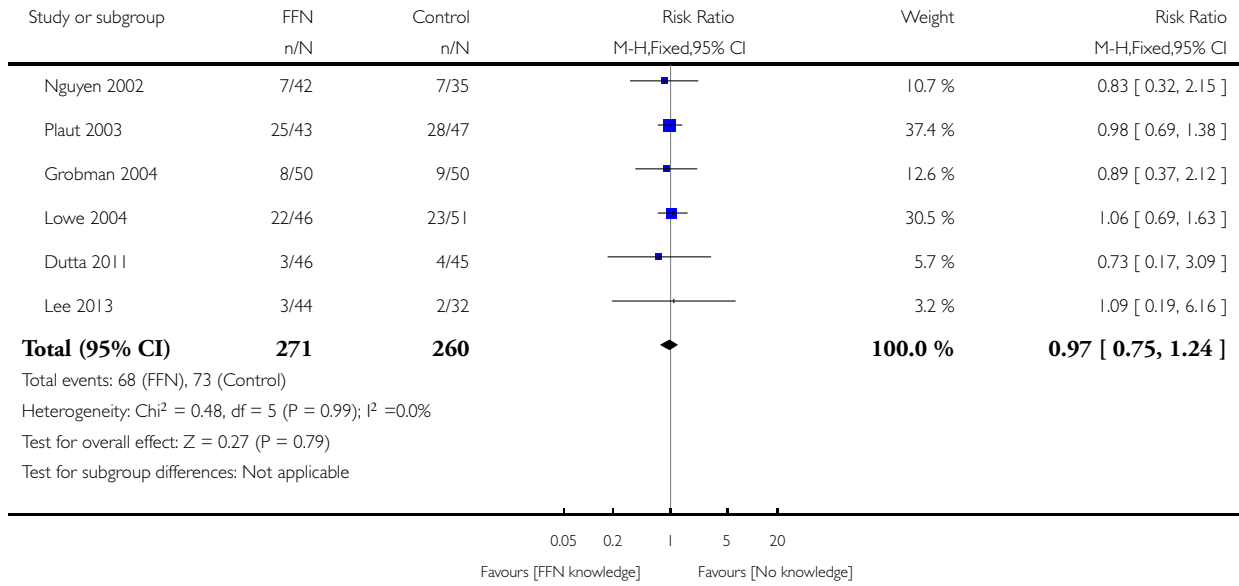


Analysis 1.9. Comparison 1 FFN knowledge versus no knowledge, Outcome 9 Tocolysis.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 9 Tocolysis

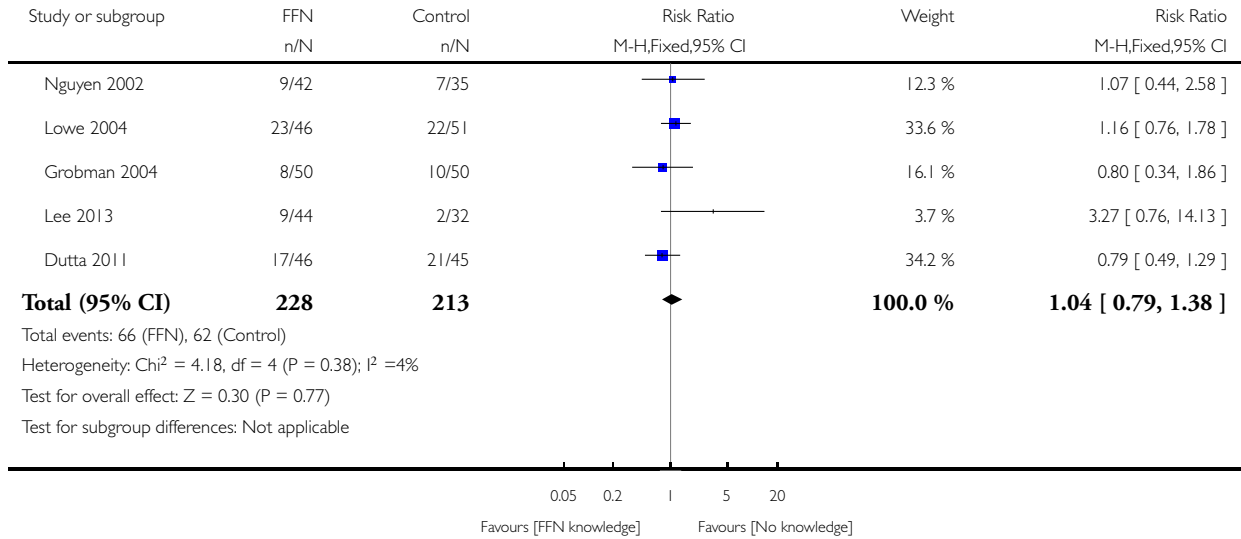


Analysis 1.10. Comparison 1 FFN knowledge versus no knowledge, Outcome 10 Steroids for fetal lung maturity.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 10 Steroids for fetal lung maturity

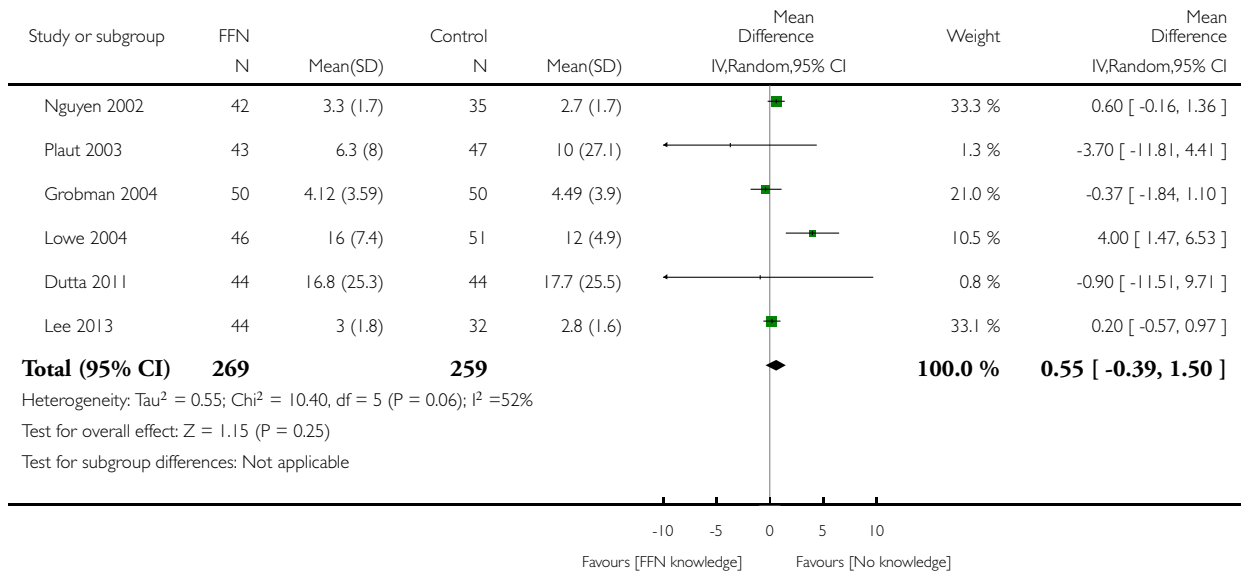


Analysis 1.11. Comparison 1 FFN knowledge versus no knowledge, Outcome 11 Time to evaluate.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 11 Time to evaluate

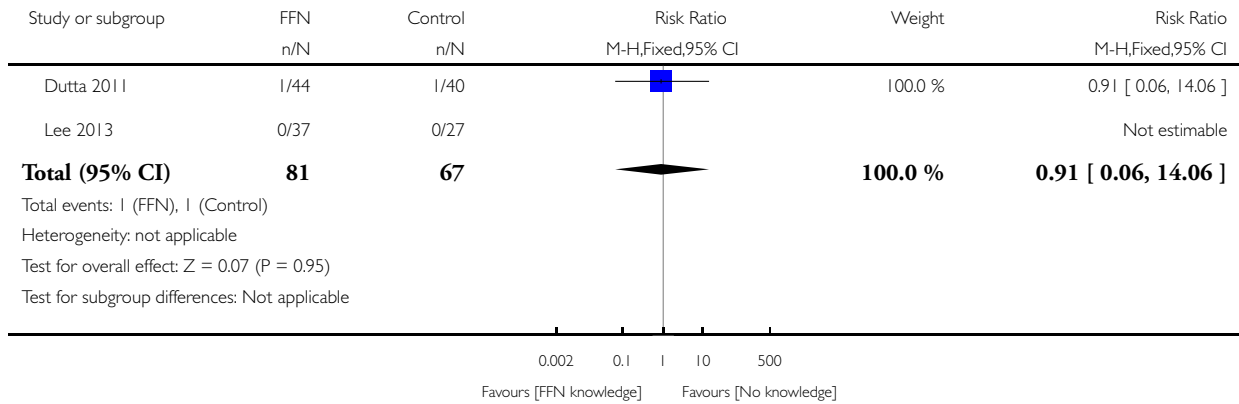


Analysis 1.12. Comparison 1 FFN knowledge versus no knowledge, Outcome 12 Respiratory distress syndrome.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 12 Respiratory distress syndrome

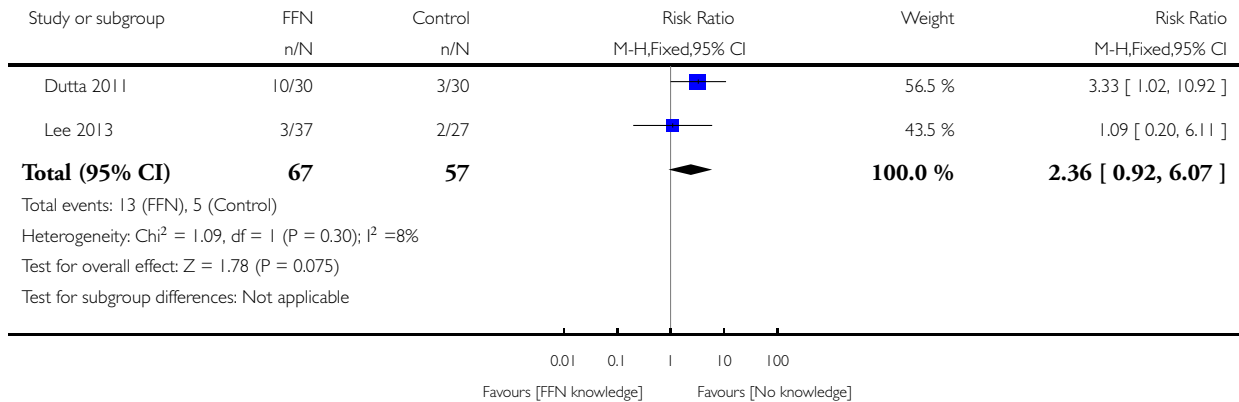


Analysis 1.13. Comparison 1 FFN knowledge versus no knowledge, Outcome 13 Neonatal intensive care unit admission.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 13 Neonatal intensive care unit admission

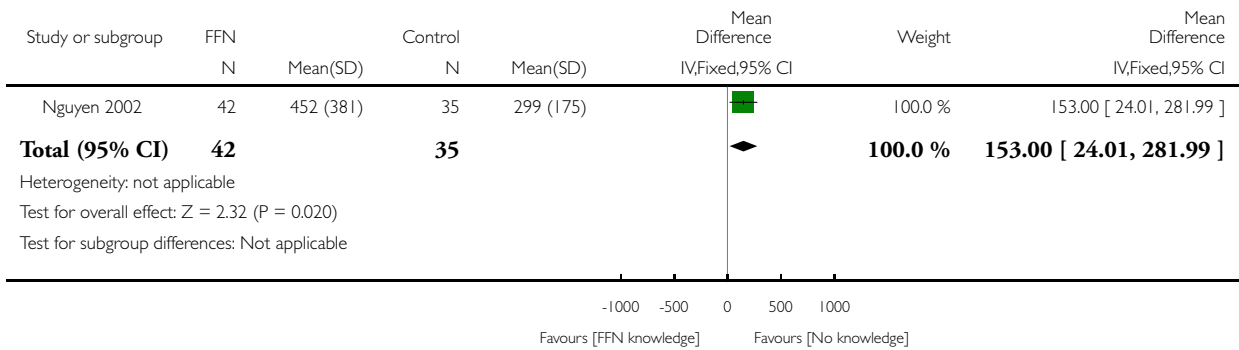


Analysis 1.14. Comparison 1 FFN knowledge versus no knowledge, Outcome 14 Economic analysis (hospitalization charges).

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 1 FFN knowledge versus no knowledge

Outcome: 14 Economic analysis (hospitalization charges)

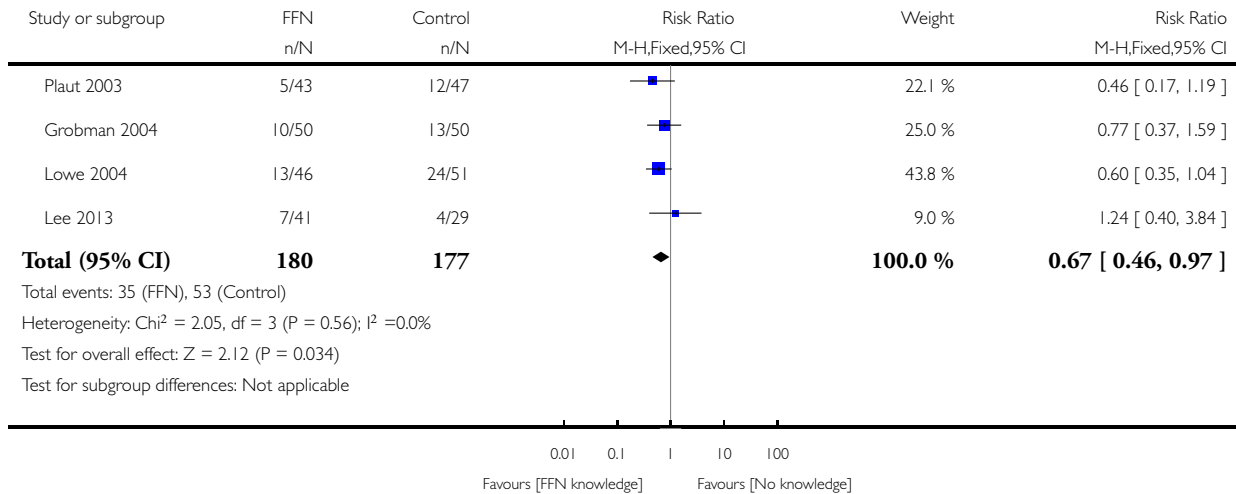


Analysis 2.1. Comparison 2 FFN knowledge versus no knowledge: sensitivity analysis, Outcome 1 Preterm birth < 37 weeks.

Review: Fetal fibronectin testing for reducing the risk of preterm birth

Comparison: 2 FFN knowledge versus no knowledge: sensitivity analysis

Outcome: 1 Preterm birth < 37 weeks



APPENDICES

Appendix I. Search methods for ICTRP, ClinicalTrials.gov and additional MEDLINE search

ICTRP

fetal AND fibronectin

ffn

ClinicalTrials.gov

fibronectin | Interventional Studies | pregnancy

fibronectin | Interventional Studies | preterm

Interventional Studies | preterm | Fetal fibronectin

WHAT'S NEW

Date	Event	Description
7 September 2018	New search has been performed	Search updated and new trials added. A 'Summary of findings' table has been incorporated
7 September 2018	New citation required but conclusions have not changed	Two new studies have been added (Dutta 2011 ; Lee 2013), but the conclusions have not changed.

HISTORY

Protocol first published: Issue 4, 2007

Review first published: Issue 4, 2008

Date	Event	Description
16 December 2007	Amended	Converted to new review format.

CONTRIBUTIONS OF AUTHORS

Vincenzo Berghella (VB), Gabriele Saccone:

1. helped with the initial idea of a fetal fibronectin review update;
2. met several times regarding all aspects of the development of the protocol and review;
3. contributed to the writing and the editing of the protocol and the review;
4. helped to revise and respond to the feedback received on the first draft of the protocol and the review.

VB prepared the first draft and finalised the revised draft of the protocol and the review in response to feedback.

VB is the guarantor of the review.

DECLARATIONS OF INTEREST

Vincenzo Berghella: is a co-author on one of the excluded trials ([Ness 2007](#)). VB was not involved in the eligibility assessment for this trial as this was assessed by Gabriele Saccone.

Gabriele Saccone: none known.

SOURCES OF SUPPORT

Internal sources

- None, Other.

External sources

- No sources of support supplied

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

A 'Summary of findings' table has been incorporated in this update and we added in a search of [ClinicalTrials.gov](https://clinicaltrials.gov) and the WHO International Clinical Trials Registry Platform ([ICTRP](https://www.who.int/ictcp)) for unpublished, planned, and ongoing trial reports.

In the original review, we also ran a separate search of MEDLINE (January 1966 to December 2007) using the search strategy detailed below:

MEDLINE

#1 exp Obstetric labor, premature/

#2 Fibronectins/

#3 #1 and #2

#4 fetal adj3 fibronectin

#5 #3 or #4

INDEX TERMS

Medical Subject Headings (MeSH)

Biomarkers [analysis]; Fetus; Fibronectins [*analysis]; Premature Birth [*prevention & control]; Randomized Controlled Trials as Topic

MeSH check words

Female; Humans; Pregnancy