

VAGINAL PROGESTERONE DECREASES PRETERM BIRTH AND NEONATAL MORBIDITY AND MORTALITY IN WOMEN WITH A TWIN GESTATION AND A SHORT CERVIX: AN UPDATED META-ANALYSIS OF INDIVIDUAL PATIENT DATA

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Short title: vaginal progesterone decreases preterm birth in women with a twin gestation and a short cervix

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ABSTRACT

OBJECTIVE To assess the efficacy of vaginal progesterone for the prevention of preterm birth and neonatal morbidity and mortality in asymptomatic women with a twin gestation and a sonographic short cervix (cervical length ≤ 25 mm) in the midtrimester.

METHODS Updated systematic review and meta-analysis of individual patient data from randomized controlled trials comparing vaginal progesterone with placebo/no treatment in women with a twin gestation and a midtrimester sonographic cervical length ≤ 25 mm. MEDLINE, EMBASE, POPLINE, CINAHL, and LILACS (all from inception to December 20, 2016), bibliographies, the Cochrane Central Register of Controlled Trials, Research Registers of ongoing trials, Google scholar, and conference proceedings were searched. The primary outcome measure was preterm birth < 33 weeks of gestation. Two reviewers independently selected studies, assessed the risk of bias, and extracted the data. Pooled relative risks (RRs) with 95% confidence intervals (CIs) were calculated.

RESULTS Individual patient data were available from 303 women (159 assigned to vaginal progesterone, 144 assigned to placebo/no treatment) and their 606 infants from six randomized controlled trials. One study including women with a CL between 20-25 mm provided 74% of total sample size of the IPD meta-analysis. Vaginal progesterone, compared with placebo/no treatment, was associated with a statistically significant reduction in the risk of preterm birth < 33 weeks of gestation

(31.4% vs. 43.1%; RR 0.69, 95% CI 0.51-0.93; moderate-quality evidence). Moreover, vaginal progesterone administration was associated with a significant decrease in the risk of preterm birth <35, <34, <32 and <30 weeks of gestation (RRs from 0.47 to 0.83), neonatal death (RR 0.53, 95% CI 0.35-0.81), respiratory distress syndrome (RR 0.70, 95% CI 0.56-0.89), composite neonatal morbidity and mortality (RR 0.61, 95% CI 0.34-0.98), use of mechanical ventilation (RR 0.54, 95% CI 0.36-0.81), and birthweight <1500 g (RR 0.53, 95% CI 0.35-0.80) (all moderate-quality evidence). There were no significant differences in neurodevelopmental outcomes at 4-5 years of age between the vaginal progesterone and placebo groups.

CONCLUSION The administration of vaginal progesterone to asymptomatic women with a twin gestation and a sonographic short cervix in the midtrimester reduces the risk of preterm birth occurring at <30 to <35 gestational weeks, neonatal mortality and some measures of neonatal morbidity, without any demonstrable deleterious effects on childhood neurodevelopment.

INTRODUCTION

Twin births have become more prevalent in developed countries over the last decades.¹⁻³ In 2014, the twin birth rate in the United States was 33.9 per 1000 live births, the highest rate ever recorded.⁴ Twin gestations are at increased risk of maternal, perinatal, and infant morbidity and mortality, as well as long-term neurodevelopmental disability.⁵⁻¹³ Moreover, twin gestations also have a significant impact on health care costs and quality of life for both parents and children.^{7,14,15}

Preterm birth is the most important factor determining neonatal morbidity and mortality among twins. The risk of preterm birth <37 and <32 weeks of gestation is 8- to 9-fold higher in twin than in singleton gestations.⁴ Several interventions have been proposed to reduce the rate of preterm birth in twin gestations such as bed rest,¹⁶ prophylactic tocolysis,¹⁷ nutritional advice,¹⁸ 17 α -hydroxyprogesterone caproate,¹⁹ vaginal progesterone,¹⁹ cerclage,²⁰ and cervical pessary.^{21,22} Unfortunately, these interventions have not been shown to reduce the risk of preterm birth in unselected twin gestations.

A short cervix, traditionally defined as a transvaginal sonographic cervical length (CL) ≤ 25 mm in the midtrimester of pregnancy, is an important risk factor for spontaneous preterm birth and has emerged as one of the strongest and most consistent predictors of preterm birth in asymptomatic women with singleton²³⁻²⁹ or twin gestations.³⁰⁻⁴³ Currently, there is compelling evidence that vaginal progesterone administration to asymptomatic women with a singleton gestation and a sonographic short cervix decreases the risk of preterm birth and neonatal morbidity and mortality.⁴⁴⁻⁴⁶ The efficacy of vaginal progesterone in women with a twin gestation and a short cervix has been less studied.

A meta-analysis of individual patient data (IPD) published in 2012 reported on the efficacy of vaginal progesterone in preventing preterm birth and neonatal morbidity and mortality in asymptomatic women with a twin gestation and a CL ≤ 25 mm in the midtrimester.⁴⁷ A total of 52 women (104 infants) from three randomized controlled trials (RCTs) were included in the study. The use of vaginal progesterone was associated with a significant 44% reduction in the risk of composite neonatal morbidity and mortality (relative risk [RR], 0.56; 95% confidence interval [CI], 0.30-0.97) and a 30% non-significant reduction in the risk of preterm birth <33 weeks of gestation (RR, 0.70; 95% CI, 0.34-1.44). Since that time, additional RCTs evaluating the use of vaginal progesterone in twin gestations have been published; therefore, reassessment of the efficacy of this intervention in women with a twin gestation and a short cervix is justified.

The objective of this study was to update the previous IPD meta-analysis on the efficacy of vaginal progesterone in asymptomatic women with a twin gestation and a sonographic CL ≤ 25 mm at midtrimester for the prevention of preterm birth and neonatal morbidity and mortality.

METHODS

The study was conducted according to a prospectively prepared protocol and reported in accordance with the Preferred Reporting Items for Systematic reviews and Meta-Analyses (PRISMA) statement.⁴⁸ The review was registered with PROSPERO, number CRD42016039682.

Data sources and searches

We searched MEDLINE, EMBASE, POPLINE, CINAHL, and LILACS (all from inception to December 20, 2016), the Cochrane Central Register of Controlled Trials, and Research Registers of ongoing trials using a combination of keywords and text words related to *progesterone*, *preterm birth*, *randomized controlled trial*, and *twin gestation*. Google scholar, proceedings of congresses on obstetrics, maternal-fetal medicine, and ultrasound in obstetrics, reference lists of identified studies, previously published systematic reviews, and review articles were also searched. Experts in the field were contacted to identify further studies. No language restrictions were applied.

Study selection

RCTs in which asymptomatic women with a twin gestation and a sonographic short cervix (CL \leq 25 mm) in the midtrimester were randomly allocated to receive vaginal progesterone or placebo/no treatment for the prevention of preterm birth and/or adverse perinatal outcomes were eligible for inclusion in the review. Trials were included if the primary aim of the study was to prevent preterm birth in women with a twin gestation and a short cervix, or to prevent preterm birth in women with an unselected twin gestation but for whom outcomes were available in those with a pre-randomization CL \leq 25 mm. We excluded quasi-randomized trials, trials that evaluated vaginal progesterone in women with preterm labor, arrested preterm labor (as maintenance tocolysis), premature rupture of membranes, or second trimester bleeding, trials that assessed vaginal progesterone in the first trimester only to prevent miscarriage, and studies that did not report clinical outcomes. Studies published as abstracts alone were excluded if additional information on methodological issues and results could not be obtained.

All of the potentially relevant studies were retrieved and reviewed independently by two authors to determine inclusion. Disagreements were resolved by discussion among the reviewers.

Data collection

The corresponding author of each eligible trial was contacted and asked to provide anonymized data (without identifiers) about baseline characteristics and outcomes for every randomly assigned patient as well as data on study characteristics and

details of interventions and cointerventions. All initial communications with authors were based on a template explaining the study and the data required. Data provided by the investigators were merged into a master database specifically constructed for the review. Data were checked for missing information, errors, and inconsistencies by cross-referencing the publications of the original trials. Quality and integrity of the randomization processes were assessed by reviewing the chronological randomization sequence and pattern of assignment, as well as the balance of baseline characteristics across treatment groups. Inconsistencies or missing data were discussed with the authors and corrections were made when deemed necessary.

Outcome measures

The primary outcome measure was preterm birth <33 weeks of gestation. Secondary outcome measures included preterm birth <37, <36, <35, <34, <32, <30 and <28 weeks of gestation; spontaneous preterm birth <33 and <34 weeks of gestation; respiratory distress syndrome (RDS); necrotizing enterocolitis; intraventricular hemorrhage; proven neonatal sepsis; retinopathy of prematurity; fetal death; neonatal death; perinatal death; a composite outcome of neonatal morbidity and mortality (defined as the occurrence of any of the following events: RDS, intraventricular hemorrhage, necrotizing enterocolitis, proven neonatal sepsis, or neonatal death); birthweight <1500 and <2500 g; admission to the neonatal intensive care unit (NICU); use of mechanical ventilation; and long-term

neurodevelopmental outcomes (suspected or diagnosed developmental delay, cerebral palsy, intellectual disabilities, vision impairment, hearing loss, cognitive and behavioral impairments, and motor, communication and learning disorders at any age in childhood).

Assessment of risk of bias

The risk of bias in each included trial was assessed independently by two authors using the criteria outlined in the Cochrane Handbook for Systematic Reviews of Interventions.⁴⁹ This tool assesses seven domains related to risk of bias (random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, selective reporting, and other bias) and categorizes studies by low, unclear, or high risk of bias in each domain. Disagreements in risk of bias assessment were resolved through consensus.

Statistical analysis

We included all randomized women and their fetuses/infants and performed all analyses on an intention-to-treat basis. For outcomes dealing with gestational age at birth, the unit of analysis was the pregnancy, whereas for perinatal outcomes, the unit of analysis was the fetus/neonate. Data from individual patients were combined in a 2-stage approach in which outcomes were analyzed in the original trial and then summary statistics were generated using standard summary data meta-analysis techniques to give an overall measure of effect (pooled RR with 95%

CI).⁵⁰ Heterogeneity of the results among studies was tested with the quantity I^2 .⁵¹ We pooled results from individual studies using a fixed-effect model if substantial statistical heterogeneity was not present (<50%). If I^2 values were $\geq 50\%$, a random effects model was used to pool data across studies. For adverse perinatal outcomes, we estimated pooled RRs using analytical methods that assumed independence between neonates. However, to avoid incorrect conclusions due to the non-independence of newborns from twin gestations, we also used a generalized linear model with generalized estimating equations to estimate parameters while controlling for cluster correlations.⁵²⁻⁵⁴ The number needed to treat (NNT) for benefit or harm with the 95% CI was calculated for outcomes for which there was a statistically significant reduction or increase in risk difference based on control event rates in the trials.⁵⁵

Subgroup analyses were performed to evaluate the effect of vaginal progesterone by CL (<10, 10-20, and 21-25 mm), daily dose of vaginal progesterone (100, 200, and 400 mg), and obstetrical history (no previous spontaneous preterm birth <37 weeks and at least 1 previous spontaneous preterm birth <37 weeks). A test for interaction between the treatment and subgroups was performed to examine whether treatment effects differed among subgroups.⁵⁶⁻⁵⁸ An interaction P -value ≥ 0.05 was considered to indicate that the effect of treatment did not differ significantly among subgroups. We planned to carry out sensitivity analyses to explore the effect of trial quality assessed by

allocation concealment and random sequence generation (considering selection bias) and blinding (considering performance and detection biases), with studies rated as "high risk of bias" or "unclear risk of bias" for these domains being excluded from the analyses in order to assess whether this made any difference to the overall result. Subgroup and sensitivity analyses were only performed for the primary outcome of preterm birth <33 weeks of gestation and for the secondary outcome of neonatal death. We also planned to explore potential sources of heterogeneity and to assess publication and related biases if at least 10 studies were included in a meta-analysis, but these analyses were not undertaken due to the limited number of trials included in the review.

Quality of evidence

We used the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach, as outlined in the GRADE Handbook,⁵⁹ to assess the quality of evidence for primary and secondary outcome measures. We considered evidence from randomized controlled trials as high quality but downgraded the evidence one level for serious (or two levels for very serious) limitations based upon the following: design (risk of bias), consistency across studies, directness of the evidence, precision of estimates, and presence of publication bias. The GRADEpro Guideline Development Tool⁶⁰ was used to import data from Review Manager in order to create a 'Summary of findings' table to

report the quality of the evidence. The GRADE approach results in an assessment of the quality of a body of evidence in one of four grades:

1. High: We are very confident that the true effect lies close to that of the estimate of the effect.
2. Moderate: 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.
3. Low: Our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.
4. Very low: We have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect.

We performed all statistical analyses using Review Manager (RevMan; version 5.3.5; The Nordic Cochrane Centre, Copenhagen, Denmark) and SAS version 9.2 (SAS Institute, Cary, NC) software.

Informed consent was provided by the patients upon enrollment in each of the original trials. In the present study, the data were not used for any purposes other than those of the original trial, and no new data were collected. Therefore, informed consent specifically for this project was not considered necessary. This study was exempted for review by the IRB Administration Office of Wayne State University.

RESULTS

Selection, characteristics, and risk of bias of studies

A total of 213 records were identified, nine of which were retrieved for full-text review. Three of these studies, which evaluated vaginal progesterone in unselected twin gestations^{61,62} or pregnancies conceived by *in-vitro* fertilization or intracytoplasmic sperm injection,⁶³ were excluded because CL was not measured or collected before randomization or there were no data for women with a CL ≤ 25 mm at randomization (Figure 1). Six studies, including a total of 303 women (606 fetuses/infants) with a CL ≤ 25 mm, met the inclusion criteria,⁶⁴⁻⁶⁹ 159 women were assigned to vaginal progesterone and 144 to placebo/no treatment. Minimal differences were noted in baseline maternal characteristics between the vaginal progesterone and placebo/no treatment groups (Table 1).

The individual characteristics of the studies included in the meta-analysis are shown in Table 2. Five studies were double-blind, placebo-controlled trials.⁶⁴⁻⁶⁸ The remaining study compared vaginal progesterone with no treatment.⁶⁹ Three studies were performed in low/middle-income countries,^{65,68,69} two in high-income countries,^{66,67} and one in both low/middle- and high-income countries.⁶⁴ Two trials were specifically designed to evaluate the use of vaginal progesterone in women with a twin gestation and a sonographic short cervix (CL ≤ 15 mm⁶⁴ and CL between 20-25 mm⁶⁹). The remaining four studies tested the effect of vaginal progesterone in women with unselected twin gestations and their authors provided data relevant to women with a CL ≤ 25 mm before randomization.⁶⁵⁻⁶⁸ The trial that

assessed vaginal progesterone in women with a CL between 20-25 mm⁶⁹ provided data for 224 mothers and their 448 infants. The other five studies provided data for 79 women and 158 infants.

Three studies used vaginal progesterone 200 mg/day (capsule,⁶⁴ pessary,⁶⁶ or ovule,⁶⁸), one used vaginal progesterone suppositories 100 mg/day,⁶⁵ one used vaginal progesterone suppositories 400 mg/day,⁶⁹ and the remaining study used vaginal progesterone suppositories 200 or 400 mg/day.⁶⁷ Treatment was started between 20-24 weeks of gestation in five trials,^{64-67,69} and between 18-21 weeks of gestation in the remaining trial.⁶⁸ Five studies reported that participants received study medication from the time of enrollment until ~34 weeks of gestation,⁶⁴⁻⁶⁸ and one from enrollment until 37 weeks of gestation.⁶⁹ Two trials included only women with a dichorionic twin gestation.^{67,69} Major fetal abnormalities, cerclage in place or planned, allergy to progesterone, and hepatic dysfunction were reported as exclusion criteria in most studies. The primary outcome measure was preterm birth <34 weeks for two trials,^{66,69} <37 weeks for two trials,^{65,67} spontaneous preterm birth <34 weeks for one trial,⁶⁴ and mean gestational age at delivery for the remaining study.⁶⁸ The study by El-refaie et al⁶⁹ did not collect data for some neonatal morbidities such as necrotizing enterocolitis, intraventricular hemorrhage, proven neonatal sepsis and retinopathy of prematurity.

The risk of bias in each included study is summarized in Figure 2. All studies had adequate generation of allocation sequence and concealment of allocation,

and appeared to be free of selective outcome reporting and other sources of bias. Five studies were considered to be at low risk for selection, performance, detection, attrition, and reporting biases.⁶⁴⁻⁶⁸ The study by El-refaie et al⁶⁹ had high risk of performance and detection biases because patients, clinical staff and outcome assessors were not blinded to the allocated interventions. In addition, this trial was judged to be at unclear risk of attrition bias because the number of losses to follow-up was not balanced across study groups (7.2% in the vaginal progesterone group and 13.6% in the “no treatment” group).

Effect of vaginal progesterone on preterm birth

Women allocated to receive vaginal progesterone had a significantly lower risk of preterm birth <33 weeks of gestation (31.4% vs. 43.1%; RR 0.69, 95% CI 0.51–0.93; $P = 0.01$; $I^2 = 0\%$; six studies, 303 women; moderate-quality evidence) compared to those allocated to placebo/no treatment (Figure 3). In addition, vaginal progesterone was associated with a significant reduction in the risk of preterm birth <35 weeks (RR 0.83, 95% CI 0.69-0.99; moderate-quality evidence), <34 weeks (RR 0.71, 95% CI 0.56-0.91; moderate-quality evidence), <32 weeks (RR 0.51, 95% CI 0.34-0.77; moderate-quality evidence), <30 weeks (RR 0.47, 95% CI 0.25-0.86; moderate-quality evidence), and spontaneous preterm birth <33 weeks (RR 0.67, 95% CI 0.48-0.93; moderate-quality evidence) and <34 weeks (RR 0.71, 95% CI 0.54-0.93; moderate-quality evidence) (Table 3). The number of patients needed to treat with vaginal progesterone to prevent one case of preterm

birth occurring at <30 to <35 gestational weeks varied from 6-12. There were no significant differences between the study groups in the risk of preterm birth <37 weeks (moderate-quality evidence), <36 weeks (moderate-quality evidence), and <28 weeks (low-quality evidence).

Effect of vaginal progesterone on adverse perinatal outcomes

Infants whose mothers received vaginal progesterone had a significantly lower risk of neonatal death (RR 0.53, 95% CI 0.35-0.81; moderate-quality evidence), perinatal death (RR 0.58, 95% CI 0.39-0.84; moderate-quality evidence), RDS (RR 0.70, 95% CI 0.56-0.89; moderate-quality evidence), composite neonatal morbidity and mortality (RR 0.61, 95% CI 0.34-0.98; moderate-quality evidence), birth weight <1500 g (RR 0.53, 95% CI 0.35-0.80; moderate-quality evidence), and use of mechanical ventilation (RR 0.54, 95% CI 0.36-0.81; moderate-quality evidence) (Table 4). The NNT to prevent one case of these adverse perinatal outcomes varied from 6-8. There was no evidence of an effect of vaginal progesterone on necrotizing enterocolitis (low-quality evidence), intraventricular hemorrhage (low-quality evidence), proven neonatal sepsis (low-quality evidence), retinopathy of prematurity (low-quality evidence), fetal death (very low-quality evidence), birth weight <2500 g (moderate-quality evidence), and admission to NICU (moderate-quality evidence).

Subgroup and sensitivity analyses

Subgroup analyses of the effect of vaginal progesterone on preterm birth <33 weeks and neonatal death are shown in Table 5. There was no evidence that women in any one of the prespecified subgroups benefit more or less from the use of vaginal progesterone than those in any other subgroup (all *P* for interaction ≥ 0.40). Nonetheless, vaginal progesterone was associated with a statistically significant reduction in the risk of preterm birth <33 weeks and neonatal death in women with a CL between 10-20 mm (RR 0.44, 95% CI 0.22-0.87 and 0.20, 95% 0.05-0.86, respectively) and women who used 400 mg of daily vaginal progesterone (RR 0.66, 95% CI 0.46-0.95 and 0.42, 95% 0.23-0.76, respectively). Moreover, vaginal progesterone significantly decreased the risk of neonatal death in women with a CL between 21-25 mm (RR 0.57, 95% 0.36-0.90) and women with no previous spontaneous preterm birth (RR 0.58, 95% 0.36-0.93).

When the sensitivity analysis was restricted to the five trials with adequate blinding of patients, clinical staff and outcome assessors,⁶⁴⁻⁶⁸ the effect of vaginal progesterone on the reduction in the risk of preterm birth <33 weeks and neonatal death was non-significant (RR 0.77, 95% CI 0.48-1.24 and 0.56, 95% CI 0.21-1.48, respectively). However, it should be noted that the sensitivity analyses did not change substantially the magnitude and direction of effect sizes obtained in the overall analyses. Sensitivity analyses based on allocation concealment and random sequence generation were not performed because there were no trials at unclear or high risk of bias for these domains.

Effect of vaginal progesterone on long-term neurodevelopmental outcomes

No study has reported the effects of vaginal progesterone on long-term neurodevelopmental outcomes in twin gestations with a short cervix. Thus far, two trials have reported the effects of prenatal exposure of vaginal progesterone on long-term neurodevelopmental outcomes in unselected twin gestations.^{70,71} In 2015, a follow-up study of one of the excluded trials⁶¹ reported that there was no significant difference in developmental delay (assessed using the Child Development Inventory tool) between twins exposed to either vaginal progesterone (42/140) or placebo (65/184) at a mean age of 55.5 months (odds ratio [OR] 0.87, 95% CI 0.46-1.63).⁷⁰ Recently, one of the studies included in the review⁶⁶ reported on the developmental performance of children exposed prenatally to vaginal progesterone (n=225) or placebo (n=212), at a mean age of 57 months.⁷¹ The developmental performance was evaluated by the parent-completed Ages and Stages Questionnaire (ASQ) screening tool. Overall, mean ASQ total scores were significantly higher in the vaginal progesterone-exposed group (269 ± 28) than in the placebo-exposed group (262 ± 31) ($P=0.03$), although there was no statistically significant difference in the risk of low ASQ score (<10th percentile) between the study groups (OR 0.47, 95% CI 0.21-1.06). A subgroup analysis showed that dichorionic twins that were exposed prenatally to vaginal progesterone had a significantly lower risk of having a low total ASQ score than those exposed to placebo (OR 0.34, 95% CI 0.14-0.86).

DISCUSSION

Principal findings

The main finding in this updated IPD meta-analysis is that the administration of vaginal progesterone to asymptomatic women with a twin gestation and a midtrimester sonographic short cervix significantly reduces the risk of preterm birth <33 weeks (primary outcome) by 31% and neonatal death by 47%. In addition, patients who received vaginal progesterone had a significantly decreased risk of preterm birth <35, <34, <32, and <30 weeks, spontaneous preterm birth <33 and <34 weeks, perinatal death, composite neonatal morbidity and mortality, RDS, birthweight <1500 g, and mechanical ventilation. Moreover, evidence from two trials that assessed vaginal progesterone in unselected twin gestations showed that there were no significant differences in the risk of neurodevelopmental disability at 4-5 years of age between children prenatally exposed to vaginal progesterone and those exposed to placebo.

Quality of the evidence

Evidence for most critical outcomes assessed with GRADE methodology was considered to be of moderate quality (Table 6). We downgraded the evidence from high-quality to moderate-quality because most of the pooled effect was provided by one study with moderate risk of bias. A judgment of moderate quality means that we have some confidence that our results approach the true impact of vaginal progesterone on preterm birth and adverse neonatal outcomes in twin gestations

with a short cervix; at the same time, we acknowledge that future trials may change these results.

Subgroup analyses

We evaluated several clinically important subgroups based on CL, daily dose of vaginal progesterone, and history of spontaneous preterm birth. Overall, subgroup analyses indicated that the beneficial effects of vaginal progesterone did not differ significantly across patient groups, as the interaction tests for subgroup differences were non-significant. Patients with a CL between 10-20 mm or those who received vaginal progesterone 400 mg/day seemed to have a larger-than-average reduction in the risk of preterm birth <33 weeks and neonatal death. However, analyses of categories such as CL <10 mm, dose of 100 or 200 mg of daily vaginal progesterone, and history of spontaneous preterm birth were based on small numbers of women, reflecting the pattern of recruitment to the original trials, in which most women had a CL between 10-25 mm, used vaginal progesterone 400 mg/day, and did not have a history of spontaneous preterm birth. As a result, our analysis was limited in its power to estimate effects within those groups of patients. Thus, although prespecified and clinically interesting, these subgroup analyses should be interpreted with caution.

Lack of long-term adverse neurodevelopmental outcomes in twins exposed to vaginal progesterone during pregnancy

Current evidence suggests that *in utero* exposure to vaginal progesterone, administered in twin gestations for the prevention of preterm birth, has no detrimental effects on long-term neurodevelopmental outcomes. A total of 761 surviving children who participated in two placebo-controlled trials of vaginal progesterone to prevent preterm birth in unselected twin gestations^{61,66} were evaluated at a mean age of ~56 months for neurodevelopmental outcomes.^{70,71} Both studies reported no significant differences in the risk of developmental delay⁷⁰ or suspected developmental delay⁷¹ between children whose mothers received vaginal progesterone and those whose mothers received placebo. It should be noted that vaginal progesterone had no effect on gestational age at delivery in both trials, which allowed the assessment of the direct effect of vaginal progesterone on childhood neurodevelopmental outcomes independent of any effect of vaginal progesterone on preterm birth. Interestingly, a subgroup analysis of one of these studies⁷⁰ found that dichorionic twins that were exposed prenatally to progesterone had a significantly reduced risk for a low total ASQ score, a higher total mean ASQ score, and higher mean ASQ scores in communication, gross motor skills, and personal/social skills in comparison with dichorionic twins that were exposed to placebo. These findings suggest a potential long-term benefit related to prenatal exposure of vaginal progesterone, which would not be surprising because there is some evidence indicating that progesterone could act as a neuroprotectant for brain disorders, mainly traumatic brain injury.⁷² Thereby, a direct beneficial effect of

vaginal progesterone on childhood neurodevelopment would be plausible. This issue deserves further investigation.

Lack of long-term adverse health outcomes in twins exposed to vaginal progesterone during pregnancy

With regard to the effects the prenatal exposure of vaginal progesterone on childhood health outcomes in twins, the follow-up study by McNamara et al⁷⁰ reported that there were no significant differences between vaginal progesterone-exposed and placebo-exposed twins with respect to death, congenital malformations, growth, health service utilization, and global health status at 3-6 years of age. The follow-up study by Vedel et al⁷¹ reported that the rates of diagnoses related to 10 organ systems, the median number of hospital admissions, and the median length of hospital stay did not differ significantly between the vaginal progesterone- and placebo-exposed twins up to 8 years of age. However, in subgroup analyses restricted to dichorionic twins and diagnoses made solely during hospital admission, the investigators found that diagnoses related to structural and functional abnormalities of the heart were significantly more frequent among children who were exposed prenatally to vaginal progesterone. Notwithstanding, these differences turned non-significant after Bonferroni adjustment for multiple comparisons. In conclusion, second- and third-trimester exposure to vaginal progesterone does not seem to have harmful effects on childhood health of twins.

Lack of adverse maternal events

In our previous IPD meta-analysis,⁴⁷ in which all included studies used vaginal progesterone 90-200 mg/day, the rates of maternal adverse effects such as vaginal discharge and vaginal pruritus, and discontinuation of treatment because of adverse effects were similar between the vaginal progesterone and placebo groups. In 2013, the three-armed trial by Serra et al⁶⁷ comparing placebo with two different daily doses of vaginal progesterone (200 and 400 mg) reported a dose-dependent, non-significant trend towards a higher rate of intrahepatic cholestasis of pregnancy (0% in the placebo group, 1% in the 200 mg group, and 5% in the 400 mg group). Nonetheless, the larger study by El-refaie et al⁶⁹ reported that there was no significant difference in the rate of intrahepatic cholestasis of pregnancy between the group using 400 mg of daily vaginal progesterone (1%) and the no treatment group (0%). Moreover, this study found that the rates of vaginal pruritus, vaginal discharge, headache, skin rash, and gastrointestinal symptoms did not differ significantly between the study groups. Thus, it appears that a 400 mg daily dose of vaginal progesterone is not associated with an increased risk of adverse maternal effects as compared with a 200 mg daily dose of vaginal progesterone or placebo/no treatment.

Strengths and limitations

The main strengths of our meta-analysis include: (1) the use of patient-level data which offer several advantages over study-level analysis, including the ability to:

use more appropriate statistical methods not always feasible using study-level analysis, define outcome measures consistently across studies, investigate subgroups in which treatment may be either more or less effective, address questions that have not been satisfactorily resolved by individual trials, minimize publication and reporting biases, and to adjust for prognostic variables that may have confounded the original treatment comparisons; (2) the baseline balance in prognostic factors between the two study groups, which reduces the possibility of causing bias in the intervention effect estimates; (3) the absence of substantial heterogeneity in most of the meta-analyses performed. Indeed, all meta-analyses about the effect of vaginal progesterone on preterm birth had no observed heterogeneity ($I^2 = 0\%$), whereas the majority of meta-analyses regarding adverse perinatal outcomes had low heterogeneity or no heterogeneity; and (4) the sensitivity analyses restricted to trials at low risk of bias that were consistent with (and thus supportive of) the overall findings.

Some potential limitations must also be considered. First, only two trials were specifically designed to assess the efficacy of vaginal progesterone in women with a twin gestation and a sonographic short cervix. Second, 74% of the total sample size of the IPD meta-analysis was provided by one study⁶⁹ which included women with a CL between 20-25 mm and was not placebo-controlled. However, it should be highlighted that assessment and measurement of most outcomes included in our review are considered objective in nature, and thereby not likely to

be influenced by lack of blinding.⁴⁹ It is noteworthy that estimates of pooled RRs obtained after excluding this study were not significantly different than those obtained in the overall analyses. Moreover, the significant 39% reduction in the risk of composite neonatal morbidity and mortality associated to vaginal progesterone administration was obtained without including data from the study by El-refaie et al⁶⁹ in the meta-analysis. Third, the larger study⁶⁹ did not collect information about several neonatal morbidities such as necrotizing enterocolitis, intraventricular hemorrhage, proven neonatal sepsis, and retinopathy of prematurity. Finally, some subgroup analyses included a small number of patients, which limits the statistical power to estimate the effects within these subgroups.

Implications for practice and research

This updated IPD meta-analysis indicates that vaginal progesterone reduces the risk of preterm birth and neonatal morbidity and mortality in patients with a twin gestation and a sonographic short cervix, without any deleterious effects on childhood neurodevelopment. Although the results of our meta-analysis appear promising, further research is required before conclusive advice can be provided with regard to the benefits of using vaginal progesterone in women with a twin gestation and a short cervix. Evidence from this updated IPD meta-analysis and three ongoing RCTs comparing vaginal progesterone with placebo (NCT02697331, NCT02518594) or no treatment (NCT02329535) in ~750 women with a twin gestation and a sonographic short cervix will help to determine whether vaginal

progesterone can be recommended to these patients with the aim of preventing preterm birth and improving perinatal outcomes.

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FIGURE LEGENDS

Legend for Figure 1: Study selection process

Legend for Figure 2: Methodological quality of studies included in the systematic review

Legend for Figure 3: Forest plot of the effect of vaginal progesterone on the risk of preterm birth <33 weeks of gestation