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Individual or group antenatal education for childbirth or parenthood, or both (Review)

Gagnon AJ, Sandall J

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[Intervention Review]

Individual or group antenatal education for childbirth or parenthood, or both

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ABSTRACT

Background

Structured antenatal education programs for childbirth or parenthood, or both, are commonly recommended for pregnant women and their partners by healthcare professionals in many parts of the world. Such programs are usually offered to groups but may be offered to individuals.

Objectives

To assess the effects of this education on knowledge acquisition, anxiety, sense of control, pain, labour and birth support, breastfeeding, infant-care abilities, and psychological and social adjustment.

Search methods

We searched the Cochrane Pregnancy and Childbirth Group's Trials Register (April 2006), CINAHL (1982 to April 2006), ERIC (1984 to April 2006), EMBASE (1980 to April 2006) and PsycINFO (1988 to April 2006). We handsearched the Journal of Psychosomatic Research from 1956 to April 2006 and reviewed the reference lists of retrieved studies. We updated the search of the Cochrane Pregnancy and Childbirth Group's Trials Register on 7 July 2011 and added the results to the awaiting classification section of the review.

Selection criteria

Randomized controlled trials of any structured educational program provided during pregnancy by an educator to either parent that included information related to pregnancy, birth or parenthood. The educational interventions could have been provided on an individual or group basis. Educational interventions directed exclusively to either increasing breastfeeding success, knowledge of and coping skills concerning postpartum depression, improving maternal psycho-social health including anxiety, depression and self-esteem or reducing smoking were excluded.

Data collection and analysis

Both authors assessed trial quality and extracted data from published reports.

Main results

Nine trials, involving 2284 women, were included. Thirty-seven studies were excluded. Educational interventions were the focus of eight of the studies (combined n = 1009). Details of the randomization procedure, allocation concealment, and/or participant accrual or loss for these trials were not reported. No consistent results were found. Sample sizes were very small to moderate, ranging from 10 to 318. No data were reported concerning anxiety, breastfeeding success, or general social support. Knowledge acquisition, sense of control, factors



related to infant-care competencies, and some labour and birth outcomes were measured. The largest of the included studies (n = 1275) examined an educational and social support intervention to increase vaginal birth after caesarean section. This high-quality study showed similar rates of vaginal birth after caesarean section in 'verbal' and 'document' groups (relative risk 1.08, 95% confidence interval 0.97 to 1.21).

Authors' conclusions

The effects of general antenatal education for childbirth or parenthood, or both, remain largely unknown. Individualized prenatal education directed toward avoidance of a repeat caesarean birth does not increase the rate of vaginal birth after caesarean section.

[Note: the 58 citations in the awaiting classification section may alter the conclusions of the review once assessed.]

PLAIN LANGUAGE SUMMARY

Individual or group antenatal education for childbirth or parenthood, or both

Benefits of antenatal education for childbirth, and the best educational approaches to use, remain unclear.

Antenatal education aims to help prospective parents prepare for childbirth and parenthood. Prospective parents often look to antenatal education to provide important information on issues such as decision making about and during labour, skills for labour, pain relief, infant and postnatal care, breastfeeding and parenting skills. There are many varied ways of providing this antenatal education and some may be more effective than others. The review found nine trials involving 2284 women. Interventions varied greatly and no consistent outcomes were measured. The review of trials found a lack of high-quality evidence from trials and so the effects of antenatal education remain largely unknown. Further research is required to ensure that effective ways of helping health professionals support pregnant women and their partners in preparing for birth and parenting are investigated so that the resources used meet the needs of parents and their newborn infants.



BACKGROUND

Extent of implementation of antenatal education

Antenatal education classes, including 'childbirth education programs', 'prenatal classes', and antenatal groups are attended by an important percentage of pregnant women worldwide. Reports can be found of programs based in the US, the UK, Canada, Mexico, Brazil, Finland, Germany, Australia, Japan, and China, to name a few. Many maternity providers, including public health departments, hospitals, private agencies and charities, and some obstetricians' and midwives' practices, are reported to provide antenatal education. In other parts of the world, antenatal preparation is still less of a formality and knowledge of the birth experience and care of children is passed from mothers to daughters or from traditional birth attendants to those in their care. Living arrangements in which several generations share common space also lend themselves to active participation in the experience of birth and consequently there may be no defined need for structured antenatal education. The existence of structured education in preparation for childbirth and parenthood has come about as traditional methods of information sharing have declined. Typically these programs have not been based on the expressed needs of attendees, but rather on the messages that the educators themselves believed they should impart.

Goals and content of antenatal education

Antenatal education programs often have a range of aims, such as to: influence health behaviour; build women's confidence in their ability to give birth; prepare women and their partners for childbirth; prepare for parenthood; develop social support networks; promote confident parents; and contribute to reducing perinatal morbidity and mortality. Antenatal education thus comprises a range of educational and supportive measures that help parents and prospective parents to understand their own social, emotional, psychological, and physical needs during pregnancy, labour, and parenthood.

Theoretical approaches to antenatal education

Common approaches which were identified in the early literature on this topic were 'natural childbirth' (Dick-Read 1933) and psychoprophylaxis (Lamaze 1958) methods. The similarity of these two techniques is their emphasis on a healthy pregnancy, physical fitness, education on the physiology of normal birth, elimination of fear during labour, use of relaxation and breathing techniques, and continuous support by a familiar person. These two approaches have experienced relative degrees of popularity in North America and Europe since their inception. Tenets from these approaches still form the basis of many childbirth education programs today (Lothian 2005). Within the last two decades, an approach called 'Active Birth' has enjoyed and continues to enjoy some popularity (Balaskas 1992), and latterly 'hypnobirth', which teaches selfhypnosis, relaxation, and breathing techniques (Mongan 2005). The essence of these approaches lies in helping mothers to identify and develop their own bodily resources for birth. It is meant to empower women through the birth experience. The choice of these and other approaches have largely been thought to be reflective of the nature of the agency sponsoring the educational program.

Reasons for participating in antenatal education

Regardless of the theoretical perspective, the question arises as to whether what is taught in classes meets the needs of attendees (Nolan 1997a; Nolan 1997b). In a survey of those planning to attend independent (private) childbirth education classes in Montreal, women reported that their main reason for attending childbirth classes was to reduce their anxiety about labour and birth (Gagnon 1995). In that same survey, men stated their primary reasons for attending antenatal classes as being to satisfy the wishes of their partners and to learn about infant care. A UK-based survey found that women would like information on physical and psychological changes during pregnancy, fetal development, what will happen during labour and childbirth, their options during labour and childbirth and how to care for themselves during this time, possible complications, and how to care for the baby after birth (Sullivan 1993).

Delivery of antenatal education

In addition to differences in theoretical approaches, there is huge variation in the provision of antenatal education, with variation in the underlying aims and the way classes are delivered. Programs range from intensive one-day classes, 'bench' classes (offered while waiting to be seen by a healthcare provider), to several classes over several weeks. There is variation in whether programs are offered individually or in groups; venues include teachers' homes, community centres, hospitals and clinics; and classes or groups may be for pregnant women only or for women and their partners or birth partners.

Teaching-learning methods include self-learning programs, didactic presentations, videos, group discussions, and programs based solely on adult learning principles in which mothers identify their own learning needs and develop an individualized learning program from there. The content of classes varies considerably and is likely to be affected by the underlying aims and the way classes are delivered, as well as the skills, experience, and motivation of the teacher. These factors may impact on the effectiveness of antenatal education programs. Because of these wide-ranging goals, the effectiveness of antenatal education has been broken down by looking at specific goals: either looking at the effects on the experience of childbirth, or on adjustment to parenthood. Few studies have observed how women use preparation methods, but women reported difficulty in using them in a hospital setting (Spiby 1999).

Lay- versus professionally-offered antenatal education

The majority of studies did not explore women's (and partners') expectations and views in depth. There are surprisingly few indepth explorations of antenatal preparation and of lay-run groups, and the majority of research is of professionally-run classes that may be directed at explaining and justifying polices, resulting in increased compliance. Lay-run classes that are structured to include the lay perspective and the interests of women have had less attention.

Social capital in antenatal education

Very few studies have investigated the impact of building support networks for women. Yet there is a substantial body of evidence that the development of what is termed 'social capital' has a major influence on health and wellbeing. Social capital refers to



features of social organisation, such as civic participation, norms of reciprocity, and trust in others, which facilitate co-operation for mutual benefit. Thus, social capital is the advantage gained by an individual or group/community of individuals, a result of being part of a social network (Aldridge 2002).

High-risk participants of antenatal education

The vast majority of literature on antenatal preparation describes courses offered to typical attendees, well-educated women in the middle-to-upper socio-economic strata. Many care providers feel that this advantaged group will do well regardless of which, if any, antenatal education program they attend. Consequently, evaluating educational program effects in this population may not be of interest. Rather, evaluating the effects of educational programs for medically or socially high-risk mothers or fathers may be of greater interest to clinicians and important in terms of public health. Descriptions of educational programs from the fathers' perspectives and for special populations are fewer in number and show differences in the educational needs of these parents-tobe. Studies in the UK and Australia have examined the views and experiences of a group of new fathers concerning the role classes played in their transition to fatherhood, and fathers' sense of exclusion from such processes (Barclay 1996; Smith 1999).

Birth experience attenuation of antenatal education

The notion that the effects of antenatal education might be attenuated by what occurs at the time of birth is a common theme mentioned by childbirth educators and mothers. Encouraging a woman's assertiveness to express her birth wishes and concerns might lead to disappointment at birth. This notion points to the need to determine the independent effects of childbirth education. If their effects are attenuated, should the resources currently directed towards antenatal education be re-directed toward programs of education for caregivers at the time of birth, or elsewhere?

Summary

In summary, evidence for the best method to deliver antenatal education is lacking. There is a great deal of variation in antenatal education programs, which must be considered in conducting a systematic review. This review will therefore include reports of programs developed from various theoretical perspectives, with a variety of objectives, content, and teaching-learning methods employed, offered to a variety of populations. Due to the nature of antenatal education, the review must be broad rather than specific in trying to determine if there is any effect of any educator-provided antenatal educational intervention in any population.

OBJECTIVES

The objectives of this review are based on stated women's and men's reports of their reasons for attending classes. The primary objective is to assess the effects of structured antenatal education delivered by an educator to an individual or group on:

(a) knowledge acquisition;

(b) anxiety;

(c) maternal sense of control/active decision-making during labour and birth;

(d) labour pain;

(e) use of medications to reduce pain;

(f) partner involvement at birth;
(g) breastfeeding success;
(h) infant care abilities;
(i) social support (defined as general emotional and homemaking/ parenting support); and
(j) psychological and social adjustment to parenthood.

A secondary objective is to assess the effects of structured antenatal education on obstetrical interventions.

METHODS

Criteria for considering studies for this review

Types of studies

We considered studies if they met the following inclusion criteria: controlled trial evaluating structured forms of antenatal education classes provided to individual parent(s)-to-be or groups of future parents; random allocation to treatment and control groups; violations of allocated management insufficient to materially affect outcomes; loss to follow up insufficient to materially affect the comparison; and data available in a form suitable for analysis.

Types of participants

One or both expectant parents.

Types of interventions

Any structured (organized) educational program, offered to individuals or groups by an educator, related to the birth of an infant including preparation for childbirth, child care, and adjustment of the parents associated with parenthood.

Types of outcome measures

Factors that could be affected by antenatal education for either childbirth or parenthood including:

- 1. knowledge acquisition;
- 2. anxiety;
- 3. maternal sense of control/active decision-making, self-confidence;
- 4. labour pain;
- 5. use of medications to reduce pain;
- 6. partner involvement at birth;
- 7. breastfeeding success;
- 8. infant care abilities;
- 9. general social support;
- 10.psychological and social adjustment to parenthood;
- 11.obstetrical interventions.

Search methods for identification of studies

Electronic searches

We searched the Cochrane Pregnancy and Childbirth Group's Trials Register by contacting the Trials Search Co-ordinator (April 2006). We updated this search on 7 July 2011 and added the results to Studies awaiting classification.

The Cochrane Pregnancy and Childbirth Group's Trials Register is maintained by the Trials Search Co-ordinator and contains trials identified from:



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

Details of the search strategies for CENTRAL, MEDLINE and EMBASE, the list of handsearched journals and conference proceedings, and the list of journals reviewed via the current awareness service can be found in the 'Specialized Register' section within the editorial information about the Cochrane Pregnancy and Childbirth Group.

Trials identified through the searching activities described above are each assigned to a review topic (or topics). The Trials Search Coordinator searches the register for each review using the topic list rather than keywords.

In addition, we searched CINAHL (1982 to April 2006) (Appendix 1), EMBASE (1980 to April 2006) (Appendix 2), ERIC (1984 to April 2006) (Appendix 3 and PsycINFO (1988 to April 2006) (Appendix 4).

Searching other resources

We also handsearched the Journal of Psychosomatic Research beginning in 1956 up to April 2006 and reviewed the reference lists of retrieved studies.

We did not apply any language restrictions.

Data collection and analysis

The full text of all studies identified from all sources were obtained. Inclusion criteria were applied to the texts and their methodological quality was assessed.

Methodological quality was determined through assessment of the likely presence or absence of biases which might have affected the internal validity of the trials. These included: selection bias through assessment of allocation concealment, attrition bias through assessment of loss of study participants to analysis and protocol deviations, and performance bias through assessment of blinding of participants, caregivers, and outcome assessors.

Statistical analyses were based on intention-to-treat and were calculated using the Cochrane statistical package, Review Manager (RevMan 2003). Results are reported as weighted mean differences and 95% confidence interval (CI), and relative risk and 95% CI.

Because an insufficient number of acceptable trials was identified, the intended subgroup analyses could not be performed to examine the effects of: (a) specific class content (eg, childbirth, childcare), (b) specific childbirth education approaches (eg, natural, psychoprophylaxis, active birth), (c) specific teaching approaches (eg, didactic, experiential), or effects in (d) specific population groups (eg, low income, multiethnic).

RESULTS

Description of studies

See the tables of 'Characteristics of included studies' and 'Characteristics of excluded studies'. Nine trials, involving 2284 women, met the study inclusion criteria. Studies were conducted in the United States, Canada, and Iran. Trials directed toward women or couples included two focused on increasing fetal attachment, one on labour, baby care, counseling, and neuromuscular exercises, one on age-appropriate development, one on early parenthood, one on a combination of 16 interventions including education about pregnancy, peer support, and health behavior, and one on the predisposing, enabling, and reinforcing factors for attempting a vaginal birth after caesarean. Two trials were directed towards men including one focusing on sexuality, pregnancy and prenatal care, labour and delivery, and infant and child care, while the other focused only on newborn care and related paternal behavior. (Fiftyeight reports from an updated search on 7 July 2011 have been added to Studies awaiting classification.)

Risk of bias in included studies

The largest of the included studies (n = 1275), examining an individualized prenatal education and support program to increase vaginal birth after caesarean section, was of high quality. The remaining trials were of uncertain quality, since details of the randomization procedure, allocation concealment, and/or participant accrual/loss were not always reported.

Effects of interventions

We included nine trials, involving 2284 women. We excluded 37 studies. Eight of the included studies (combined n = 1009) tested more general educational interventions. Specific interventions, populations, and outcomes measured were different in each study. No data from these trials were reported on anxiety, breastfeeding success, or general social support. Knowledge acquisition, sense of control, factors related to infant-care competencies, and some labour and birth outcomes were measured.

Three of the studies showed beneficial effects on knowledge gained. In one, weighted mean group differences (WMD) of 1.62 (95% confidence interval (CI) 0.49 to 2.75) on a scale of 20 were found (n = 48; Corwin 1999); a second, focusing on fathers-to-be, showed a WMD of 9.55 (95% CI 1.25 to 17.85) on a 75-item scale (n = 28; Westney 1988); and a third study showed a range of being of 20% (relative risk (RR) 1.20, 95% CI 1.07 to 1.35) to over twice as likely (RR 2.22, 95% CI 1.27 to 3.90) of knowing about specific issues (Klerman 2001).

Those receiving prenatal education in conjunction with several other interventions were found to be nearly 50% more likely to perceive mastery for behaviour change (RR, 1.48, 95% CI 1.05 to 2.09). Length of labour was found to be shorter in those receiving prenatal education in one study (Mehdizadeh 2005); in active labour: WMD -1.10 hours, 95% CI -1.64 to -0.56; in the second stage of labour: WMD -8.70, 95% CI -12.60 to -4.80. Benefits to attachment were found in very small studies (WMD 52.6 attachment behaviors, 95% CI 21.82 to 83.38, n = 10, Carter-Jessop 1981; and WMD 9.70 attachment behaviors, 95% 0.15 to 19.25, n = 22, Davis 1987). Similar results were found for satisfaction with maternal role preparation (WMD 21.59 of a possible score ranging from 24 to 120, 95% CI 11.23 to 31.95, n = 16, Hamilton-Dodd 1989). The results of all these smaller studies are likely to be subject to significant bias since



reports of baseline characteristics by group are largely absent and when group differences were reported, were not addressed in terms of their potential effect on outcome assessment. No differences were found in any other outcomes measured. The ninth and largest of the included studies (n = 1275; Fraser 1997), examining an educational and social support intervention to increase vaginal birth after caesarean section, showed similar rates of vaginal birth after caesarean section in 'verbal' and 'document' groups (RR 1.08, 95% CI 0.97 to 1.21).

DISCUSSION

Although women and their partners in many countries are commonly referred to structured antenatal education programs, the benefits of these programs to participants and their newborn infants remain unclear.

With the exception of one large trial aimed at increasing the rate of vaginal birth after caesarean delivery, the trials were small to moderate and of uncertain methodological quality. The usual benefit of meta-analyses of increasing statistical power by combining small studies was not achieved, since each study was testing the effect of a different intervention on one or more different outcomes. Further, this variation did not permit proposed subgroup analyses to be conducted to examine the effects of specific class content, theoretical underpinnings, teaching approaches, or effects in specific population groups.

AUTHORS' CONCLUSIONS

Implications for practice

No recommendations for practice changes can be made at this time since there exists insufficient evidence to determine the effects of person-to-person antenatal education for childbirth or parenthood, or both. The widespread popularity of antenatal classes, particularly in some northern countries, testifies to the desire of many (particularly first-time) expectant parents or their caregivers for antenatal education.

Implications for research

The extensive amount of resources placed in attendance and provision of structured antenatal education programs suggest the need for a large, well-designed clinical trial to answer the question of effectiveness. Conducting such a trial in this field will be very challenging due to their popularity, particularly in some northern countries. In previous work, we found only 10% of women initially inquiring about classes to state their willingness to be randomized to either classes or a placebo arm (Gagnon 1995). Further, compliance to the assigned intervention may be difficult. Women not already planning to attend classes are unlikely to attend even if they are randomized to do so.

Difficulties in conducting a randomized controlled trial in this field can also be expected to arise when deciding just what and in whom to test antenatal education given the great variation in content, in approach, and in the populations to whom it is currently provided.

In conducting such a trial, one should ensure that outcomes to be measured include those defined as important by both the consumers of this education and those referring parents-to-be for education.

[Note: the 58 citations in the awaiting classification section may alter the conclusions of the review once assessed.]

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Methods	Randomization to experimental and placebo/usual care groups, procedure not described. Outcome as- sessor was masked to group allocation.
Participants	10 healthy, married, "white", nulliparous women, 21-34 years old, attended by private obstetricians in the US, 32-37 weeks' gestation who were attending childbirth education classes. Reportedly an equivalent number of breastfeeders and equivalent attachment in both groups (no data provided). No other baseline comparisons reported.

Carter-Jessop 1981 (Continued)	
Interventions	All participants attended childbirth education classes. Group A: attachment intervention (n = 5). Consisted of 2-3 weekly encounters with women (apparent- ly not in groups) by the same nurse. During these encounters the women were encouraged to: (1) feel for the fetus' parts and to check for the fetal position daily, (2) increase their awareness of fetal activi- ty and notice how they can affect that activity, and (3) rub, stroke, and gently massage their abdomens over the fetus. Group B: usual prenatal care (n = 5).
Outcomes	Maternal attachment measured by observing the frequency of attachment behaviors for 1 second every 10 seconds during a 10-minute period at day 2-4 postpartum using the 'postnatal attachment test' developed by the author using the results of Marshall Klaus' work. No reliability or validity testing was done on this 'test'. There were 9 attachment behaviors which could be observed; no range of possible frequencies for these behaviors was given. Site of postpartum data collections was not stated. No SDs were reported. Means: attachment group, 106.6, control group, 54. Estimates of the SD from the crude data as reported: 23.09 (attachment group) and 26.46 (usual care group).
Notes	No SDs were reported. Means: attachment group, 106.6, control group, 54. Estimates of

Risk of bias		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	B - Unclear

Methods	Randomization to experimental and placebo/usual care groups, procedure not described. In one sec- tion, authors state that those in classes were randomized, elsewhere it states that those who registered for classes by telephone were randomized. Masking to group allocation by provider or outcome asses- sor, or both, was not described.
Participants	48 adults (stated as 48 couples in the abstract) in expectant parent classes in the US. These were moth ers with or without a partner/support person. Inclusion criteria were not described. Criteria to exclude from the analysis were: non-English speaking, bedrest due to preterm labour, delivery prior to the in- tervention, missing 'significant' (not defined) portions of the class, attending refresher or VBAC classes Baseline data were presented including gender, making it unclear as to whether the data presented re fer to the women or women and partners. Their average ages were 32-33 years old, 83.3% (control (C) group) to 91.7% (experimental (E) group) were Caucasian, 67-71% held bachelors' degrees. The groups were not equivalent on female gender (E = 66.7%, C = 54.2%), income over \$60,000 USD (E = 41.7%, C = 20.8%), and multiparity (E = 8.3%, C = 20.8%).
Interventions	The number of classes, their duration, and rate of attendance were not reported for either group. Group A: 'expanded' or integrated childbirth education (n = 24 plus those subsequently excluded for not completing 'significant portions of the class'). Participants were taught information about age- appropriate development in the first year of life through demonstrating parallels between labour challenges and early parenting challenges. The goal was to give parents the ability to generalize their labour coping skills to postpartum coping. Group B: 'traditional' or usual prenatal education (n = 24 plus those subsequently excluded for not completing 'significant portions of the class'). Usual education was not described.
Outcomes	Parenting knowledge as measured by the Prenatal Parenting Scale, a 20-item true/false questionnaire developed by the author. No reliability or validity data were provided. No scoring information was provided.



Corwin 1999 (Continued)

Notes

Selection bias may have occurred given that at least some of the exclusion criteria appear to have been applied postrandomization. Analyses of effects, parent knowledge score means, did not address the issue of clinically important baseline differences in gender, income, and parity.

Risk of bias

Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	B - Unclear

Davis 1987 Methods Randomization to experimental and control groups. The participant and provider were not masked to group allocation; the outcome assessor was masked. Allocation concealment was not described. 22 17-32 year old, low-risk, predominantly "white" English-speaking pregnant women (32-37 weeks' Participants gestational age) in a US clinic. Baseline characteristics by group did not differ. Interventions The "intrauterine attachment" intervention consisted of individual instruction given to women in a "clinical setting at scheduled visits". It was based on the theoretical framework of Barnard and Rubin and included: (1) demonstrating fetal position assessment through abdomen palpation and drawing pictures of what was found 3 times per week; (2) exploring any maternal behavior that the woman felt as an effect on fetal activity; (3) rubbing, stroking, massaging the abdomen for fetal response and taking note. 3 classes were scheduled; duration of these classes was not reported. Placebo/usual care was not described. Outcomes Selected maternal attachment subscales: affectionate behavior, proximity maintaining, caretaking behavior, and mother's attention were reported based on observations at 2-4 days postpartum. Overall score obtained by totalling the frequency of occurrence of each behavior in each subscale for 20 seconds of each minute for 15 minutes of observation. No ranges were provided. Notes **Risk of bias** Bias Authors' judgement Support for judgement Allocation concealment Unclear risk D - Not used (selection bias)

Methods	Eligible consenting women from one of 12 hospitals (1 US and 11 Canadian) were randomized through a central telephone answering service within strata defined by hospital and women's motivation to at-
	tempt vaginal delivery. Masking of outcome assessor was not described.
Participants	Inclusion criteria: previous low transverse caesarean, gestational age < 28 weeks, planning to give birth in a participating hospital, receiving prenatal care from a physician in a participating hospital, knowl- edge of English or French sufficient to permit completion of a self-administered questionnaire. Exclusion criteria: previous vaginal birth after caesarean, classic caesarean section or myomectomy scar, known multiple gestation.



raser 1997 (Continued)	1301 women were randomized, 1275 completed the study. No group differences were seen on charac- teristics measured at baseline.	
Interventions	Experimental: 'verbal' group (n = 641): offered an individualized prenatal education and support pro- gram that was provided by a research nurse with expertise in prenatal instruction and a resource per- son with strong communication skills and personal experience of a VBAC. The approach taken in pro- viding the intervention was to consider the predisposing, enabling, and reinforcing factors for decidin to attempt a VBAC. Pain relief options for labour and methods for permanent contraception (for those considering tubal ligation at the time of the caesarean section) were discussed. Each of 2 contacts las ed approximately 1 hour. They usually occurred at 21 and 27 weeks' gestation. The first contact was provided to 98% of the group, the second was provided to 72% of those in the low motivation for VBA stratum and 79% of those in the high motivation stratum. Control: 'document' group (n = 634): received a pamphlet highlighting the benefits of VBAC rather tha an elective repeat caesarean section.	
Outcomes	Primary outcome: proportion of women achieving vaginal delivery. Other outcomes included: pro- portion attempting vaginal delivery, and maternal and neonatal morbidity. These data were obtained through medical record review. Women's sense of control over the birth experience was obtained as measured by the Birth Experience Rating Scale (shortened version of the Labour Agentry Scale, no reli- ability or validity data reported). This scale included 18 Likert-type items, each with 7 steps resulting in a maximum total score of 126.	
Notes	Sample sizes by group were not presented for the Birth Experience Rating Scale thus prohibiting its in- clusion in a meta-analysis. Additional exploratory results were presented on sub-groups defined by maternal motivation for VBAC	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	A - Adequate

Hamilton-Dodd 1989

Methods	Women were randomized to either the maternal preparation program or routine care. Randomization procedures were not described. Unclear whether outcome assessors were masked to group allocation.
Participants	22 women from a medical center in Coving, California, USA, were initially randomized. 6 women with- drew although no information is provided regarding when and from which group. Data were reported on 16 women who were successfully recruited and completed the study. Inclusion criteria were: par- ticipation approved by the physician, 18-35 years old (inclusive), absence of major complications dur- ing pregnancy, women planning to be the primary caretakers of their infants for the first 2 months post- birth, predominantly anglophone. Exclusion criteria: major complications during birth, < 38 weeks' or > 41 weeks' gestation, 5-minute Apgar < 7, birthweight < 2500 g, mental or physical anomalies identified at birth or during the project. Maternal age and caesarean birth rate (25%) were similar in both groups. Group differences were found in parity and infant gender.
Interventions	Attendance (or lack thereof) at general childbirth education classes was not reported for any partici- pants. Group A: maternal preparation program (n = 8). 4 individual sessions of 1 1/2 to 2 hours length were provided by 1 of 3 occupational therapists over a 3 to 7 week period beginning within 1 month prior to birth and finishing when the infant reached 2-3 weeks of age. Content included activities of daily living incorporating the maternal role, infant development, physiological changes in the new mother, and mother-infant relationship. In general, broad concepts were discussed rather than specific information. The occupational therapists were available for telephone inquiries. Group B: routine care (n = 8), not described.

Hamilton-Dodd 1989 (Continued)

Outcomes	Maternal competence (measured by Ainsworth's 'System for Rating Maternal Care Behaviors'), adapta- tion to the maternal role (measured by Lederman's 'Postpartum Self Evaluation Questionnaire' and by Sheehan's 'Prenatal and Postnatal Questionnaire'), and satisfaction with the maternal role preparation provided with obstetrical care (measured by the Satisfaction with a Maternal Preparation Program Re- ceived in Conjunction with Obstetric Care Questionnaire). The System for Rating Maternal Care Behav- iors consists of 7 general maternal care sub-scales divided into a total of 22 behaviors rated on a scale of 1 to 9 with higher scores indicating greater competence. The Postpartum Self-Evaluation Question- naire is made up of 8 subscales containing 81 questions rated from 'very much so' (1) to 'not at all' (4). A ninth subscale was created by combining the last two. The Prenatal and Postnatal Questionnaire is a 10-item Likert-type Scale. The Satisfaction with a Maternal Preparation Program Received is a 24-item Likert Scale . Total scores ranged from 24 to 120 with higher scores indicating greater satisfaction.
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Notes

Analyses of effects did not address the issue of clinically important baseline differences in parity and infant gender.

Risk of bias

Bias Authors' judgement		Support for judgement				
Allocation concealment (selection bias)	Unclear risk	B - Unclear				

Klerman 2001

Bias	Authors' judgement Support for judgement
Risk of bias	
Notes	
Outcomes	Perceived quality of prenatal care, average time with nurse, average time at clinic, same nurse report- ed as providing care each visit, nurse reported as "very helpful", knowledge of risk factors, self-report of behavioral change, mean birthweight/age, low birthweight, mean gestational age, preterm birth, IU- GR, low APGAR score, NICU, congenital anomalies, mean maternal weight gain, and caesarean deliver- ies. No measurement details were provided for questions posed to women.
Interventions	The experimental group (n = 318) received augmented care, including 16 interventions of which group education was only one. 40-minute discussion groups and individual instruction were provided in a clinical setting. Written materials were distributed. Content focus was on pregnancy, peer social support, and health behavior education to minimize risk. Other supports for the experimental group (n = 318) included educationally-oriented peer groups, additional appointments and extended time with clinicians. The control group (n = 301) received "usual care" including prenatal classes that were at different times and locations than clinic appointments.
Participants	Included were: pregnant, African-American women eligible for medicaid and seeking prenatal care from Jefferson County, AL, USA from 1994-1996, scoring greater than 10 on a risk assessment scale, less than 26 weeks' gestational age, 16 years old, having had no major medical complications. Women hav- ing experienced alcoholism, high blood pressure, substance abuse, asthma, cancer, diabetes, epilep- sy, sickle cell disease, or HIV/AIDS were excluded. Most participants were single. Groups were similar or baseline characteristics. 656 women enrolled in the trial, outcome data at birth were available for 619, postpartum interview data (year 2) were available for 118.
Methods	Placebo/usual care controlled. It is unclear whether the participant, provider, and outcome assessor were masked.



Unclear risk

Klerman 2001 (Continued)

Allocation concealment (selection bias)

D - Not used

Mehdizadeh 2005							
Methods	Women were randomized (no details given).						
Participants	200 participants were recruited from Hedayat Hospital in Tehran, were under 35 years olds, and at 20 weeks' gestational age. The report states no statistically significant differences in demographic infor- mation by group (no data provided).						
Interventions		Women attended a total of 7 90-minute classes which included 30 minutes of instruction on baby care, stages of labour, and diet, 30 minutes of counseling, and 30 minutes of neuromuscular exercises.					
Outcomes	Back or pelvic pain, length of labour, vaginal delivery, oxytocin, episiotomy, birthweight, increased dai- ly activity, headache, and sleep disturbance. No measurement details were provided for questions posed to women.						
Notes							
Risk of bias							
Bias	Authors' judgement	Support for judgement					
Allocation concealment (selection bias)	Unclear risk	D - Not used					

Pfannenstiel 1991	
Methods	Randomized to 6 groups, 3 experimental and 3 placebo/usual care comparison groups. Randomization procedures were not described. Outcome assessors were masked to allocation.
Participants	67 men were recruited. Inclusion criterion: the mother had to agree to explain the study to the father. Of the 67 participating fathers, 22 of their partners were considered to be high medical risk. Their aver- age age was 22.5 years, they had an average of 11 years of education, held unskilled and semi-skilled positions, 70% were Caucasian, 88% were acquainted with their partners for over 1 year; 46% were married. There were reported to be no group differences at baseline on maternal or paternal age, ma- ternal or paternal education, socioeconomic status (Hollingshead), or race.
Interventions	3 experimental groups: high-risk pregnancy intervention with pre- and postmeasures (n = 11), low- risk pregnancy intervention with pre- and postmeasures (n = 11), low-risk pregnancy intervention with postmeasures only (n = 12). All 3 experimental groups received the same intervention: 2 1-1/2 hour classes were presented to fathers through the hospital clinic, 57% attended the group sessions, 43% at- tended tutorials. The content included newborn care, normal newborn behaviour, paternal self-image, attitude towards the infant, pregnancy, parent-infant interactions, normal child development. The educational ap- proach used was didactic and modeling. The background of the educators is not described. 3 control groups (n = 11 in each subgroup): content not described.
Outcomes	Paternal sensitivity and empathetic behaviour (as measured by the Assessment of Father-Infant Sensi- tivity scale) scored from 10 minute videotapes of infant feeding. The Assessment of Father-Infant Sen- sitivity Scale incorporates 12 items, each item is rated on a 5-point scale behaviorally defined from fail- ure in behavioral empathy (scores 1,2) to relative failure alternating with sensitive responses (score 3) to sensitive/empathic behaviors (score 4,5). In addition, it includes ratings for 6 father-plus-infant items



Pfannenstiel 1991 (Continued)

rated on a 5-point scale with higher scores indicative of more sensitive and empathic behavior. Interrater reliability ranged from .72 to .85.

Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement
	Judicio Judgement	oupport for Jungement

Westney 1988

Methods	Randomization to experimental and placebo/usual care group, procedure not described. Masking to group allocation by provider or outcome assessor, or both, not described.						
Participants	28 black, unmarried, 15-18 year old prospective fathers recruited from public schools, adolescent pre- natal clinics, recreation centers, other expectant fathers. Mean age did not differ by group. Reported not to differ on baseline knowledge.						
Interventions	Experimental: 4 2-hour weekly prenatal classes addressing: human sexuality, pregnancy and prenatal care, labour and delivery, infant and child care. All were presented by a female registered nurse-spe- cialist in maternal-child care. Teaching approaches included lectures, audiovisual aids, and group dis- cussions of concerns. Control: no pregnancy-related education program.						
Outcomes	Knowledge of human sexuality, pregnancy and prenatal care, labour and delivery, and infant and child care, measured using a 75-item questionnaire developed for this study. One form of this questionnaire was administered at baseline, another at completion of the classes for the experimental group and at 4 weeks postbaseline for the control group. No reliability or validity data were provided for this questionnaire. The range of possible scores was not given. It appears that the higher the score, the greater the knowledge.						
Notes	The knowledge score means and standards deviations used in this review were taken from Table 2 of the article. Unfortunately, the labelling of that table (4 rows of data, each only partially labelled) pro- hibits the reader from being fully certain that the scores used in this review are correct.						
Risk of bias							
Bias	Authors' judgement	Support for judgement					
Allocation concealment (selection bias)	Unclear risk	B - Unclear					

IUGR: intrauterine growth retardation NICU: neonatal intensive care unit VBAC: vaginal birth after caesarean section

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion								
Astbury 1980	90 primigravidae women, half attending childbirth education classes, half not, were randomly as- signed once in hospital in labour to either standard care, music, or information groups. Effects on anxiety were assessed; no standard deviations were reported thereby prohibiting inclusion of these data in the current review. No other outcomes were reported.								
Beck 1980	Participants were randomized to standard Lamaze preparation (n = 37) or Lamaze plus group de- sensitization to a hierarchy of labour and delivery-related scenes (n = 30). These groups were com- pared to each other and to self-selected controls. Only those participants who attended at least 4 of the 8 sessions were included in the data analyses, prohibiting an intention-to-treat analysis. Out- come data are not usable as presented, in that only correlations and F statistics are provided, and means and standard deviations cannot be calculated.								
Bjornson 1997	The aim of this study was to determine whether a video about infant immunization could inform parents as well as an oral presentation. There was no control condition permitting an assessment of the effects of education, the focus of the current review.								
Calabro 1996	The aim of this study was to determine whether health education materials in both English and Spanish were more effective when written at a lower rather than a higher reading level. The focus of the education material was to discourage alcohol use during pregnancy. There was no control condition permitting an assessment of the effects of education, the focus of the current review.								
Carson 1984	69 women were randomized to one of three groups: relaxation (in which relaxation techniques for labour were taught); abdominal palpation (in which identification of the fetal position and mas- sage were taught); and control. The objective of the study was to determine if abdominal palpation of the fetus would increase maternal attachment behavior. Published data include the mean num- ber of attachment behaviors observed (relaxation, 159, palpation, 156, control 142); no standard deviations are reported thereby prohibiting inclusion of these data in the current review. Caesare- an birth rate and breastfeeding data are only reported for the 3 groups combined.								
Chang 1987	Not a randomized controlled trial. This was a controlled experiment in which 1 of 4 childbirth class- es was provided with a pamphlet on seatbelt use, another received the pamphlet and 15-minute instruction, and the remaining 2 served as controls. There was no random assignment of individu- als or groups to an intervention.								
Cogan 1982	Childbirth education teachers, rather than parents, were randomly assigned to an experimental group which received communication skills training.								
D'Andrea 1994	Inner city African-American pregnant adolescents were randomized to receive either individual counseling and educational and community outreach-referral services (n = 32) or usual care (n = 32). Program effects on social networks and self-concept were assessed; no standard deviations were reported thereby prohibiting inclusion of these data in the current review. No other effects were assessed.								
Dalzell 1965	Not a randomized controlled trial. Each new participant was successively assigned to either prena- tal classes, home visits, individual counseling, or a control group.								
De Nuncio 2000	352 Latinas from two clinics in San Diego County were recruited and randomized to experimental and placebo/control care groups at the 34-36 week prenatal visit. The experimental intervention included immunization education; the control intervention was education to prevent SIDS. Group sizes were not reported, thereby prohibiting inclusion of these data in the current review.								
Durham 1986	Unit of analysis error and outcome data are not usable as presented. Published data are present- ed as the number of times medication was used to reduce pain summed for the entire group rather than for the individuals within the group (i.e., not analyzed based on the unit of randomization). Group means and standard deviations cannot be calculated.								
Escott 2005	The aim of this study was "to compare the use and effects of enhanced pre-existing coping strate- gies with the use and effects of coping strategies usually taught in National Health Service (NHS)								

Study	Reason for exclusion								
	antenatal education on women's experience of pain and emotions during labour". There was no control condition permitting an assessment of the effects of education, the focus of the current review.								
Ferland 1981	There was not random assignment to groups; "2 of the classes constituted the experimental group (n = 19); the 2 remaining classes, the control group (n = 22)".								
Fischer 1972	Not a randomized controlled trial. Primiparous women were assigned to classes or a control condi- tion based on their expected dates of delivery.								
Ford 2002	282 14-20 year old primarily single women were recruited from 5 different urban clinics in the US. The intervention was mastery modeling and peer support. A group of 6-8 pregnant girls were paired with other pregnant teens with similar due dates. The intervention took place in a clinic setting. Social cognition theory provided the theoretical basis for the classes, which introduced modeling and rehearsing skills to achieve the following: 1) increased knowledge about pregnancy, 2) preparation for childbirth, 3) working with the healthcare system, 4) assessing adolescent and infant health, and 5) preventing unplanned pregnancy. Normal first labour, birth, and exercise were included. Intervention and control groups received written material. The control group received individual instruction. The distribution of age and "race" differed by group, bringing into question the randomization procedure. Group sizes were not reported, thereby prohibiting inclusion of these data in the current review.								
Glazier 1997	The interventions compared were written ones: a pamphlet on triple marker screening versus one on activities of daily living during pregnancy. No information sharing by persons (as in a class or one-to-one teaching) was tested.								
Goodson 1985	Not a randomized controlled trial. Alternate prenatal classes were assigned a 30-minute lecture on child passenger safety. There was no random assignment of individuals or groups to an interven- tion.								
Leitch 1999	Data presented is incompatible with Review Manager analysis methods.								
Linares 2006	255 women were randomized to the standard arm (a standard educational sheet about a particu- lar pregnancy-related topic) and 370 to the intervention arm (an educational sheet with gestation- al age-specific information, an explanation of the importance of next appointment, and a section of the woman's individual medical or pregnancy complications). Effects on missed appointments were assessed; no data by group were presented thereby prohibiting inclusion of these data in the current review.								
Manandhar 2004	This is a study of a community-based participatory intervention meant to reduce neonatal mortali- ty and is outside the realm of this review.								
McEnery 1986	Strong likelihood of selection bias and absence of an intention-to-treat analysis. 105 women ini- tially volunteered for the study. 69 appear to have been randomized (35 to the 'educated' group and 34 to the 'not educated' group) although the point in the process at which this was done is un- clear since reasons for exclusion include 'infant not examined at 1 year'. 19 women from the experi- mental group who did not attend more than 3 classes were moved to the control group for analyses leaving 16 in the 'educated' and 53 in the 'not educated' groups.								
Midmer 1995	Possible selection bias. Couples were reportedly randomized to receive either two classes on par- enting communication in addition to usual childbirth education (experimental group) or usual childbirth education (control group). The authors state that "each couple had an equal probabil- ity of being randomized to the experimental group or the control group". However, later in the text, the authors state "more [were] assigned to the experimental group to ensure adequate class size" (experimental group n = 41 couples, 58-62 individuals were analyzed; control group n = 29 couples, 42-45 individuals analyzed). The method of assignment to assure equal class size was not specified. Outcome data are not usable as presented. Published data include means without stan- dard deviations.								

Study	Reason for exclusion							
Nichols 1987	No outcome data are provided.							
O'Cathain 2002	This was a cluster-randomized controlled trial with a written intervention. Leaflets on promoting informed choice in women using maternity services were used. Information sharing by persons (as in a class or one-to-one relationship) was not tested. The intervention is beyond the scope of this review.							
Petrowski 1981	Strong likelihood of selection bias and attrition bias. 56 women were initially randomized, 20 were subsequently lost, leaving 36 across 4 treatment conditions. 4 were subsequently non-randomly added to obtain 4 groups of 10 each. Outcome data are not usable as presented. Published data include F statistics only; means and standard deviations can not be calculated.							
Polley 2002	This study focuses on limiting weight gain during pregnancy. The objective of it was to determine whether a stepped care behavioral intervention would decrease the percentage of women who gain more than the Institute of Medicine recommendation. The intervention is beyond the scope of the current review.							
Schachman 2004	US military wives were randomized to traditional childbirth education or "baby boot camp" (a childbirth-parenting preparation program based on a resilience paradigm), thereby prohibiting an assessment of effects, the focus of this review.							
Schonberger 2004	Study intervention is focused on reducing newborn exposure to allergens and smoke, not a focus of this review.							
Spence-Cagle 1984	All couples received regular Lamaze classes and were subsequently randomized to receive "added interpersonal needs content" to determine if the later might increase "love scores". No data were provided in the report.							
Springer 1996	Uses women attending childbirth classes as the population from which they drew their sample. In- dividuals were not randomized. This article does not evaluate antenatal education.							
St. James-Roberts	Women attendees of childbirth education classes were "randomly assigned to EMG and SCL biofeedback groups and to two equivalently sized control groups such that age, social class, and [anxiety] scores were counterbalanced across groups group sizes were 13 SCL, 13 SCL-control, 11 EMG, and 11 EMG-control". The effects of childbirth education itself, the focus of this review, were not assessed.							
Sullivan 1984	99 pregnant women were randomly assigned to receive either four self-directed programmed in- struction booklets with slides (experimental intervention) or no intervention (control). Since the intervention was not provided by an individual, the focus of the current review, it was excluded. Outcome data were also not presented in a format suitable for this review. Published data include mean group knowledge scores (experimental, 47.75; control, 27.57) without standard deviations. Degrees of freedom, mean square, f statistic and P values of an analysis of variance of a 3 x 2 factor- ial design were also reported.							
Timm 1979	Outcome data are not usable as presented and attrition bias may be present. Published data in- cludes mean birthweight adjusted for gestational age without standard deviations. No other means or standard deviations are provided or can be calculated. Of 146 women agreeing to participate, 118 completed the study; numbers were unevenly distributed by group (Group A n = 40, Group B n = 31, Group C n = 47), suggesting attrition bias.							
Turan 2001	Couples in Istanbul, Turkey were randomized to an antenatal education program in a couple's group, a women-only group, or a men-only group. They received group educational sessions, a booklet, and a telephone counseling service. Outcomes included men's knowledge regarding breastfeeding, preventative healthcare, and family planning as well as contraceptive use. The number of subjects analyzed at follow up was not reported.							

Study	Reason for exclusion
Washington 1983	Outcome data are not usable as presented and attrition bias may be present. Published data in- clude mean knowledge acquisition (experimental group, 13.2; control group, 10.3) without stan- dard deviations and standard deviations can not be calculated. Of 68 women randomized, data are presented on 40, 20 having attended classes and 20 not. The 28 lost were unaccounted. The author reports the mean difference as being statistically significant.
Zhang 1996	The experimental intervention is unclear; attempts to contact the trial author for further informa- tion have been unsuccessful.
Zimmermann-Tansella	The aim of this study was to determine the effects of Respiratory Autogenic Training versus tradi- tional psychoprophylaxis on relaxation and anxiety during the course, and pain and behaviour dur- ing birth. There was no control condition permitting an assessment of the effects of education, the focus of the current review.

EMG: electromyographic SCL: skin conductance level SIDS: sudden infant death syndrome

DATA AND ANALYSES

Comparison 1. Individual maternal education versus routine care

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Observed maternal competence	1	16	Mean Difference (IV, Fixed, 95% CI)	1.87 [-16.04, 19.78]
2 Adaptation to the maternal role (Le- derman tool)	1	16	Mean Difference (IV, Fixed, 95% CI)	12.0 [-4.83, 28.83]
3 Adaptation to the maternal role (Sheehan tool)	1	16	Mean Difference (IV, Fixed, 95% CI)	2.12 [-1.82, 6.06]
4 Satisfaction with maternal role preparation	1	16	Mean Difference (IV, Fixed, 95% CI)	21.59 [11.23, 31.95]

Analysis 1.1. Comparison 1 Individual maternal education versus routine care, Outcome 1 Observed maternal competence.

Study or subgroup	Treatment		Control			Mean Difference				Weight	Mean Difference
	Ν	N Mean(SD)		N Mean(SD)		Fixed, 95% CI					Fixed, 95% CI
Hamilton-Dodd 1989	8	157.9 (22.4)	8	156 (13)						100%	1.87[-16.04,19.78]
Total ***	8		8				•			100%	1.87[-16.04,19.78]
Heterogeneity: Not applicable											
Test for overall effect: Z=0.2(P=0.84)											
			Fa	vours control	-100	-50	0	50	100	Favours trea	itment



Analysis 1.2. Comparison 1 Individual maternal education versus routine care, Outcome 2 Adaptation to the maternal role (Lederman tool).

Study or subgroup	Tre	eatment	c	Control		Mean Difference				Weight M	lean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% CI				Fixed, 95% CI
Hamilton-Dodd 1989	8	126.4 (20.3)	8	114.4 (13.4)						100%	12[-4.83,28.83]
Total ***	8		8				-			100%	12[-4.83,28.83]
Heterogeneity: Not applicable											
Test for overall effect: Z=1.4(P=0.16)											
			Fa	vours control	-100	-50	0	50	100	Favours treatme	nt

Analysis 1.3. Comparison 1 Individual maternal education versus routine care, Outcome 3 Adaptation to the maternal role (Sheehan tool).

Study or subgroup	Tre	eatment	Control			Mean Difference			Weight	Mean Difference	
	Ν	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% C	I			Fixed, 95% CI
Hamilton-Dodd 1989	8	39.3 (4)	8	37.1 (4)						100%	2.12[-1.82,6.06]
Total ***	8		8							100%	2.12[-1.82,6.06]
Heterogeneity: Not applicable											
Test for overall effect: Z=1.05(P=0.29))										
			Fa	vours control	-10	-5	0	5	10	Favours treatm	ent

Analysis 1.4. Comparison 1 Individual maternal education versus routine care, Outcome 4 Satisfaction with maternal role preparation.

Study or subgroup	Tre	reatment Co		Control		Mea	an Difference		Weight	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	xed, 95% CI			Fixed, 95% CI
Hamilton-Dodd 1989	8	101.4 (10.8)	8	79.8 (10.3)					100%	21.59[11.23,31.95]
Total ***	8		8				•		100%	21.59[11.23,31.95]
Heterogeneity: Not applicable										
Test for overall effect: Z=4.09(P<0.0	0001)				1					
			Fa	vours control	-100	-50	0 50	100	Favours trea	atment

Comparison 2. Paternal education classes versus routine care

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Paternal sensitivity	1	66	Mean Difference (IV, Fixed, 95% CI)	0.38 [-0.04, 0.80]
2 Paternal plus infant sensitivity	1	66	Mean Difference (IV, Fixed, 95% CI)	0.65 [0.04, 1.26]

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
3 Paternal knowledge	1	28	Mean Difference (IV, Fixed, 95% CI)	9.55 [1.25, 17.85]

Analysis 2.1. Comparison 2 Paternal education classes versus routine care, Outcome 1 Paternal sensitivity.

Study or subgroup	Tre	Treatment		Control		Mean Difference		Weight		Mean Difference	
	N	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% C				Fixed, 95% CI
Pfannenstiel 1991	34	9.3 (0.9)	32	9 (0.9)			+			100%	0.38[-0.04,0.8]
Total ***	34		32				•			100%	0.38[-0.04,0.8]
Heterogeneity: Tau ² =0; Chi ² =0	0, df=0(P<0.0001	L); I ² =100%									
Test for overall effect: Z=1.76	(P=0.08)										
			Fa	vours control	-10	-5	0	5	10	Favours treatme	ont

Favours treatment Favours control

Analysis 2.2. Comparison 2 Paternal education classes versus routine care, Outcome 2 Paternal plus infant sensitivity.

Study or subgroup	Tre	eatment	Control			Mean Difference				Weight	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% C	I			Fixed, 95% CI
Pfannenstiel 1991	34	12.4 (1.2)	32	11.8 (1.3)			+			100%	0.65[0.04,1.26]
Total ***	34		32				•			100%	0.65[0.04,1.26]
Heterogeneity: Not applicable											
Test for overall effect: Z=2.1(P=0.04)						ī					
			Fa	vours control	-10	-5	0	5	10	Favours treatme	nt

Analysis 2.3. Comparison 2 Paternal education classes versus routine care, Outcome 3 Paternal knowledge.

Study or subgroup	Treatment		Control			Mean Difference			Weight	Mean Difference
	N Mean(SD)		N Mean(SD)		Fixed, 95% CI					Fixed, 95% CI
Westney 1988	15	35.9 (12.6)	13	26.4 (9.7)					100%	9.55[1.25,17.85]
Total ***	15		13				•		100%	9.55[1.25,17.85]
Heterogeneity: Not applicable										
Test for overall effect: Z=2.26(P=0.02)										
			Fa	vours control	-100	-50	0 50) 100	Favours treat	ment

Comparison 3. Childbirth education classes + maternal attachment preparation versus childbirth education classes alone

Outcome or subgroup title	No. of studies	No. of par- ticipants	Statistical method	Effect size
1 Frequency of maternal attachment behav- iors	1	10	Mean Difference (IV, Fixed, 95% CI)	52.60 [21.82, 83.38]

Analysis 3.1. Comparison 3 Childbirth education classes + maternal attachment preparation versus childbirth education classes alone, Outcome 1 Frequency of maternal attachment behaviors.

Study or subgroup	Tr	eatment	с	ontrol		Mean Difference			Weight	Mean Difference	
	Ν	Mean(SD)	Ν	Mean(SD)		Fixed, 95% CI					Fixed, 95% CI
Carter-Jessop 1981	5	106.6 (23.1)	5	54 (26.5)			-	-	_	100%	52.6[21.82,83.38]
Total ***	5		5						•	100%	52.6[21.82,83.38]
Heterogeneity: Not applicable											
Test for overall effect: Z=3.35(P=0)						1					
			Fa	vours control	-100	-50	0	50	100	Favours trea	itment

Comparison 4. Expanded childbirth education classes versus traditional childbirth education classes

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Parenting knowledge	1	48	Mean Difference (IV, Fixed, 95% CI)	1.62 [0.49, 2.75]

Analysis 4.1. Comparison 4 Expanded childbirth education classes versus traditional childbirth education classes, Outcome 1 Parenting knowledge.

Study or subgroup	Tre	Treatment		ontrol		Mean Difference			Weight	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	xed, 95% CI			Fixed, 95% CI
Corwin 1999	24	16.8 (1.7)	24	15.2 (2.3)					100%	1.62[0.49,2.75]
Total ***	24		24				•		100%	1.62[0.49,2.75]
Heterogeneity: Tau ² =0; Chi ² =0	, df=0(P<0.0001	.); I ² =100%								
Test for overall effect: Z=2.81(F	P=0)									
			Fa	vours control	-10	-5	0	5 10	Favours treatme	ent

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Vaginal birth after caesarean	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	1.08 [0.97, 1.21]
2 Attempted vaginal delivery	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.97, 1.12]
3 Scheduled caesarean	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	0.90 [0.74, 1.11]
4 Unsuccessful attempt at vaginal birth af- ter caesarean	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	0.96 [0.77, 1.19]
5 Urgent caesarean	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	0.88 [0.58, 1.33]
6 Uterine rupture or dehiscence	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	1.32 [0.46, 3.78]
7 Hysterectomy	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	0.20 [0.01, 4.11]
8 Maternal blood transfusion	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	0.66 [0.11, 3.93]
9 Perinatal death	1	1280	Risk Ratio (M-H, Fixed, 95% CI)	0.50 [0.09, 2.69]
10 Apgar score < 7 at 5 minutes	1	1280	Risk Ratio (M-H, Fixed, 95% CI)	0.66 [0.24, 1.84]
11 Neonatal intensive care unit admission	1	1280	Risk Ratio (M-H, Fixed, 95% CI)	1.00 [0.75, 1.34]
12 Instrumental delivery	1	1275	Risk Ratio (M-H, Fixed, 95% CI)	1.03 [0.80, 1.31]

Comparison 5. Individual vaginal birth after caesarean education/support versus routine care

Analysis 5.1. Comparison 5 Individual vaginal birth after caesarean education/ support versus routine care, Outcome 1 Vaginal birth after caesarean.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N			M-H, F	ixed, 9	5% CI				M-H, Fixed, 95% Cl
Fraser 1997	339/641	310/634				+				100%	1.08[0.97,1.21]
Total (95% CI)	641	634				•				100%	1.08[0.97,1.21]
Total events: 339 (Treatment), 310 (Co	ontrol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.42(P=0.15)					1						
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 5.2. Comparison 5 Individual vaginal birth after caesarean education/ support versus routine care, Outcome 2 Attempted vaginal delivery.

Study or subgroup	Treatment n/N	Control n/N	Risk Ratio M-H, Fixed, 95% Cl							Weight	Risk Ratio M-H, Fixed, 95% Cl
Fraser 1997	465/641	440/634				+				100%	1.05[0.97,1.12]
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



Study or subgroup	Treatment n/N	Control n/N			Ris M-H, Fi	sk Rat ixed, t				Weight	Risk Ratio M-H, Fixed, 95% Cl
Total (95% CI)	641	634				•				100%	1.05[0.97,1.12]
Total events: 465 (Treatment), 440 (Control)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.23(P=0.22	2)				1						
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 5.3. Comparison 5 Individual vaginal birth after caesarean education/support versus routine care, Outcome 3 Scheduled caesarean.

Study or subgroup	Treatment	Control			Risk Ra	io			Weight	Risk Ratio
	n/N	n/N		M-H	, Fixed,	95% CI				M-H, Fixed, 95% Cl
Fraser 1997	137/641	150/634			-+-				100%	0.9[0.74,1.11]
Total (95% CI)	641	634			•				100%	0.9[0.74,1.11]
Total events: 137 (Treatment),	150 (Control)									
Heterogeneity: Tau ² =0; Chi ² =0,	, df=0(P<0.0001); I ² =100%									
Test for overall effect: Z=0.98(F	9=0.33)									
	Fa	vours treatment	0.1 (0.2 0.5	1	2	5	10	Favours control	

Analysis 5.4. Comparison 5 Individual vaginal birth after caesarean education/support versus routine care, Outcome 4 Unsuccessful attempt at vaginal birth after caesarean.

Study or subgroup	Treatment	Control			Ri	sk Rat	io			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	5% CI				M-H, Fixed, 95% Cl
Fraser 1997	126/641	130/634				-+				100%	0.96[0.77,1.19]
Total (95% CI)	641	634				+				100%	0.96[0.77,1.19]
Total events: 126 (Treatment), 130 (C	Control)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.38(P=0.71)										
	Fa	avours treatment	0.1	0.2	0.5	1	2	5	10	Favours control	

Analysis 5.5. Comparison 5 Individual vaginal birth after caesarean education/support versus routine care, Outcome 5 Urgent caesarean.

Study or subgroup	Treatment	Control			Ri	sk Rat	io			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% Cl
Fraser 1997	39/641	44/634			_	+				100%	0.88[0.58,1.33]
Total (95% CI)	641	634				•				100%	0.88[0.58,1.33]
Total events: 39 (Treatment), 44	(Control)										
Heterogeneity: Tau ² =0; Chi ² =0, d	f=0(P<0.0001); I ² =100%										
Test for overall effect: Z=0.62(P=0	0.54)										
	Fa	vours treatment	0.1	0.2	0.5	1	2	5	10	Favours control	



Analysis 5.6. Comparison 5 Individual vaginal birth after caesarean education/ support versus routine care, Outcome 6 Uterine rupture or dehiscence.

Study or subgroup	Treatment	Control			Ri	sk Ra	tio			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed,	95% CI				M-H, Fixed, 95% CI
Fraser 1997	8/641	6/634						_		100%	1.32[0.46,3.78]
Total (95% CI)	641	634						-		100%	1.32[0.46,3.78]
Total events: 8 (Treatment), 6 (Control))										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.52(P=0.61)											
	Fa	avours treatment	0.1	0.2	0.5	1	2	5	10	Favours control	

Analysis 5.7. Comparison 5 Individual vaginal birth after caesarean education/support versus routine care, Outcome 7 Hysterectomy.

Study or subgroup	Treatment	Control	Risk Ratio			Weight	Risk Ratio		
	n/N	n/N		M-H, Fi	xed, 9	95% CI			M-H, Fixed, 95% CI
Fraser 1997	0/641	2/634				_		100%	0.2[0.01,4.11]
Total (95% CI)	641	634				-		100%	0.2[0.01,4.11]
Total events: 0 (Treatment), 2 (Control)								
Heterogeneity: Not applicable									
Test for overall effect: Z=1.05(P=0.3)									
	Fa	avours treatment	0.001	0.1	1	10	1000	Favours control	

Analysis 5.8. Comparison 5 Individual vaginal birth after caesarean education/ support versus routine care, Outcome 8 Maternal blood transfusion.

Study or subgroup	Treatment	Control			Ris	sk Rat	tio			Weight	Risk Ratio
	n/N	n/N			M-H, Fi	ixed,	95% CI				M-H, Fixed, 95% Cl
Fraser 1997	2/641	3/634			+			-		100%	0.66[0.11,3.93]
Total (95% CI)	641	634								100%	0.66[0.11,3.93]
Total events: 2 (Treatment), 3 (Control))										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.46(P=0.65)											
	Fa	avours treatment	0.1	0.2	0.5	1	2	5	10	Favours control	

Favours treatment Favours control

Analysis 5.9. Comparison 5 Individual vaginal birth after caesarean education/support versus routine care, Outcome 9 Perinatal death.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio
	n/N	n/N		M-H	l, Fixed, 95	% CI			M-H, Fixed, 95% CI
Fraser 1997	2/643	4/637				1		100%	0.5[0.09,2.69]
		Favours treatment	0.01	0.1	1	10	100	Favours control	



Study or subgroup	Treatment	Control		R	isk Ratio)		Weight	Risk Ratio
	n/N	n/N		М-Н, Г	ixed, 95	% CI			M-H, Fixed, 95% Cl
Total (95% CI)	643	637						100%	0.5[0.09,2.69]
Total events: 2 (Treatment), 4 (Control)	l.								
Heterogeneity: Not applicable									
Test for overall effect: Z=0.81(P=0.42)						1			
	Fa	vours treatment	0.01	0.1	1	10	100	Favours control	

Analysis 5.10. Comparison 5 Individual vaginal birth after caesarean education/ support versus routine care, Outcome 10 Apgar score < 7 at 5 minutes.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% Cl
Fraser 1997	6/643	9/637								100%	0.66[0.24,1.84]
Total (95% CI)	643	637								100%	0.66[0.24,1.84]
Total events: 6 (Treatment), 9 (Control))										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.79(P=0.43)											
	Fa	avours treatment	0.1	0.2	0.5	1	2	5	10	Favours control	

Analysis 5.11. Comparison 5 Individual vaginal birth after caesarean education/ support versus routine care, Outcome 11 Neonatal intensive care unit admission.

Study or subgroup	Treatment	Control			Risk Ratio					Weight	Risk	Ratio
	n/N	n/N			M-H, F	ixed, 9	5% CI				M-H, Fix	ed, 95% CI
Fraser 1997	82/643	81/637								100%		1[0.75,1.34]
Total (95% CI)	643	637				\blacklozenge				100%		1[0.75,1.34]
Total events: 82 (Treatment), 81 (Contr	rol)											
Heterogeneity: Not applicable												
Test for overall effect: Z=0.02(P=0.98)												
	Fa	avours treatment	0.1	0.2	0.5	1	2	5	10	Favours control		

Analysis 5.12. Comparison 5 Individual vaginal birth after caesarean education/support versus routine care, Outcome 12 Instrumental delivery.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N			М-Н, F	ixed, 9	95% CI				M-H, Fixed, 95% CI
Fraser 1997	109/641	105/634								100%	1.03[0.8,1.31]
Total (95% CI)	641	634				•				100%	1.03[0.8,1.31]
Total events: 109 (Treatment), 105 (C	ontrol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.21(P=0.83))										
	Fa	avours treatment	0.1	0.2	0.5	1	2	5	10	Favours control	

Comparison 6. Promotion of intrauterine attachment versus routine care

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Affectionate behavior (maternal attach- ment subscale)	1	22	Mean Difference (IV, Fixed, 95% CI)	9.70 [0.15, 19.25]
2 Proximity maintaining (maternal attach- ment subscale)	1	22	Mean Difference (IV, Fixed, 95% CI)	2.87 [-1.04, 6.78]
3 Mother's attachment (maternal attach- ment subscale)	1	22	Mean Difference (IV, Fixed, 95% CI)	-1.90 [-8.52, 4.72]

Analysis 6.1. Comparison 6 Promotion of intrauterine attachment versus routine care, Outcome 1 Affectionate behavior (maternal attachment subscale).

Study or subgroup	Tre	eatment	Control			Mean Difference				Weight	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	ixed, 95% CI				Fixed, 95% CI
Davis 1987	10	77.2 (8.6)	12	67.5 (14)						100%	9.7[0.15,19.25]
Total ***	10		12				•			100%	9.7[0.15,19.25]
Heterogeneity: Not applicable											
Test for overall effect: Z=1.99(P=0.05)											
			Favo	urs treatment	-100	-50	0	50	100	Favours contro	l

Analysis 6.2. Comparison 6 Promotion of intrauterine attachment versus routine care, Outcome 2 Proximity maintaining (maternal attachment subscale).

Study or subgroup	Treatment		c	ontrol	Mean Difference	Weight	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
Davis 1987	10	42.7 (4.7)	12	39.8 (4.6)		100%	2.87[-1.04,6.78]
Total ***	10		12			100%	2.87[-1.04,6.78]
Heterogeneity: Not applicable							
Test for overall effect: Z=1.44(P=0.15)				1		1	
			-	10	5 0 5	10 =	

Favours treatment -10 -5 0 5 ¹⁰ Favours control

Analysis 6.3. Comparison 6 Promotion of intrauterine attachment versus routine care, Outcome 3 Mother's attachment (maternal attachment subscale).

Study or subgroup	Tre	atment	Control			Me	an Differe	nce		Weight	Mean Difference	
	N	Mean(SD)	Ν	Mean(SD)	Fixed, 95% CI					Fixed, 95% CI		
Davis 1987	10	6.6 (7.6)	12	8.5 (8.3)						100%	-1.9[-8.52,4.72]	
			Favo	urs treatment	-10	-5	0	5	10	Favours contro	l	



Study or subgroup	Treatment		Control		Mean Difference					Weight	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	xed, 95% (CI			Fixed, 95% CI
Total ***	10		12							100%	-1.9[-8.52,4.72]
Heterogeneity: Not applicable											
Test for overall effect: Z=0.56(P=0.57)											
			Favou	rs treatment	-10	-5	0	5	10	Favours contro	l

Comparison 7. Augmented prenatal care versus routine care

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Prenatal care - very helpful	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.17 [1.05, 1.30]
2 Prenatal care - somewhat or not too help- ful	1	223	Risk Ratio (M-H, Fixed, 95% CI)	0.27 [0.11, 0.64]
3 Compared with last time, prenatal care rated - better	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.17 [1.05, 1.30]
4 Compared with last time, prenatal care rated - same	1	223	Risk Ratio (M-H, Fixed, 95% CI)	0.43 [0.25, 0.74]
5 Compared with last time, prenatal care rated - worse	1	223	Risk Ratio (M-H, Fixed, 95% CI)	0.10 [0.01, 0.77]
6 Average time reported with nurse per visit - less than 15 minutes	1	223	Risk Ratio (M-H, Fixed, 95% CI)	0.39 [0.24, 0.64]
7 Average reported time with nurse per visit - about 30 minutes	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.30 [1.02, 1.66]
8 Average time reported with nurse per visit - about 45 minutes	1	223	Risk Ratio (M-H, Fixed, 95% CI)	2.37 [0.96, 5.84]
9 Average time reported with nurse per visit - about 60 minutes	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.42 [0.48, 4.22]
10 Average reported time from entering to leaving clinic - about 1 hour	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.58, 1.89]
11 Average reported time from entering to leaving clinic - about 1.5-2.5 hours	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.48 [1.17, 1.86]
12 Average reported time from entering to leaving clinic - about 3 hours or more	1	223	Risk Ratio (M-H, Fixed, 95% CI)	0.37 [0.21, 0.67]
13 Same nurse reported as providing care each visit	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.13 [1.03, 1.24]
14 Nurse rated as "very helpful"	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.16 [1.03, 1.30]
15 Knowledge of risk factors - told she or her baby might be "at risk" or "have problems"	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.48 [1.03, 2.14]



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Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
16 Knowledge of risk factors - changed be- havior during pregnancy in response to in- formation about risks/problems	1	223	Risk Ratio (M-H, Fixed, 95% CI)	2.22 [1.27, 3.90]
17 Knowledge of risk factors - told how much weight to gain	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.20 [1.07, 1.35]
18 Self-report of behavioral change/status - regular vitamin-mineral supplementation	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.88, 1.16]
19 Self-report of behavioral change/status - smoking cessation (for smokers only)	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.40 [0.56, 3.48]
20 Self-report of behavioral change/status - perceived mastery	1	223	Risk Ratio (M-H, Fixed, 95% CI)	1.48 [1.05, 2.09]
21 Mean birthweight	1	607	Mean Difference (IV, Fixed, 95% CI)	44.0 [-50.51, 138.51]
22 Low birthweight (less than 2500 g)	1	607	Risk Ratio (M-H, Fixed, 95% CI)	1.48 [0.92, 2.39]
23 Mean gestational weight age, week	1	607	Mean Difference (IV, Fixed, 95% CI)	0.30 [-0.13, 0.73]
24 Preterm birth	1	607	Risk Ratio (M-H, Fixed, 95% CI)	0.77 [0.50, 1.18]
25 Intrauterine growth restriction	1	607	Risk Ratio (M-H, Fixed, 95% CI)	1.30 [0.82, 2.09]
26 Low Apgar score - at 1 minute	1	607	Risk Ratio (M-H, Fixed, 95% CI)	0.81 [0.53, 1.24]
27 Low Apgar score - at 5 minutes	1	607	Risk Ratio (M-H, Fixed, 95% CI)	1.59 [0.38, 6.58]
28 Neonatal intensive care unit stay	1	607	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.47, 1.09]
29 Congenital anomalies	1	607	Risk Ratio (M-H, Fixed, 95% CI)	0.68 [0.22, 2.12]
30 Mean maternal weight gain during preg- nancy, kg - if BMI less than 19.8 (at risk)	1	607	Mean Difference (IV, Fixed, 95% CI)	1.0 [-0.01, 2.01]
31 Mean maternal weight gain during preg- nancy, kg - if BMI is greater than or equal to 19.8 (not at risk)	1	607	Mean Difference (IV, Fixed, 95% CI)	0.10 [-1.16, 1.36]
32 Caesarean deliveries	1	607	Risk Ratio (M-H, Fixed, 95% CI)	0.80 [0.55, 1.17]

Analysis 7.1. Comparison 7 Augmented prenatal care versus routine care, Outcome 1 Prenatal care - very helpful.

Study or subgroup	Treatment	Control		Risk Ratio						Weight	Risk Ratio
	n/N	n/N			М-Н, F	ixed, 9	95% CI				M-H, Fixed, 95% Cl
Klerman 2001	110/118	84/105		<mark>+</mark>				1		100%	1.17[1.05,1.3]
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



Study or subgroup	Treatment	Treatment Control			Ri	sk Rat	tio			Weight	Risk Ratio	
	n/N	n/N			M-H, F	ixed,	95% CI				M-H, Fixed, 95% Cl	
Total (95% CI)	118	105				•				100%	1.17[1.05,1.3]	
Total events: 110 (Treatment),	84 (Control)											
Heterogeneity: Not applicable												
Test for overall effect: Z=2.79(P	=0.01)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment		

Analysis 7.2. Comparison 7 Augmented prenatal care versus routine care, Outcome 2 Prenatal care - somewhat or not too helpful.

Study or subgroup	Treatment Control Risk Ratio					Weight	Risk Ratio		
	n/N	n/N		М-Н, Р	ixed, 95%	6 CI			M-H, Fixed, 95% Cl
Klerman 2001	6/118	20/105			-			100%	0.27[0.11,0.64]
Total (95% CI)	118	105		-	-			100%	0.27[0.11,0.64]
Total events: 6 (Treatment), 20 (Control)								
Heterogeneity: Not applicable									
Test for overall effect: Z=2.96(P=0)									
		Favours control	0.01	0.1	1	10	100	Favours treatment	

Analysis 7.3. Comparison 7 Augmented prenatal care versus routine care, Outcome 3 Compared with last time, prenatal care rated - better.

Study or subgroup	Treatment	Control		Risk Ratio						Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	5% CI				M-H, Fixed, 95% CI
Klerman 2001	110/118	84/105				+				100%	1.17[1.05,1.3]
Total (95% CI)	118	105				•				100%	1.17[1.05,1.3]
Total events: 110 (Treatment), 84 (Co	ntrol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=2.79(P=0.01)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.4. Comparison 7 Augmented prenatal care versus routine care, Outcome 4 Compared with last time, prenatal care rated - same.

Study or subgroup	Treatment	Control	Risk Ratio				Weight	Risk Ratio		
	n/N	n/N		M-H, Fixe	d, 95% CI				M-H, Fixed, 95% Cl	
Klerman 2001	16/118	33/105	-		_			100%	0.43[0.25,0.74]	
Total (95% CI)	118	105	-					100%	0.43[0.25,0.74]	
Total events: 16 (Treatment), 33 (Con	itrol)									
Heterogeneity: Not applicable										
Test for overall effect: Z=3.07(P=0)										
		Favours control	0.1 0.2	0.5	1 2	5	10	Favours treatment		



Analysis 7.5. Comparison 7 Augmented prenatal care versus routine care, Outcome 5 Compared with last time, prenatal care rated - worse.

Study or subgroup	Treatment	Control		I	Risk Ratio			Weight	Risk Ratio M-H, Fixed, 95% Cl	
	n/N	n/N		м-н,	Fixed, 95%	% CI				
Klerman 2001	1/118	9/105		-				100%	0.1[0.01,0.77]	
Total (95% CI)	118	105			-			100%	0.1[0.01,0.77]	
Total events: 1 (Treatment), 9 (Control)									
Heterogeneity: Not applicable										
Test for overall effect: Z=2.21(P=0.03)										
		Favours control	0.01	0.1	1	10	100	Favours treatment		

Analysis 7.6. Comparison 7 Augmented prenatal care versus routine care, Outcome 6 Average time reported with nurse per visit - less than 15 minutes.

Treatment	Control	Risk Ratio	Weight	Risk Ratio M-H, Fixed, 95% Cl	
n/N	n/N	M-H, Fixed, 95% CI			
18/118	41/105		100%	0.39[0.24,0.64]	
118	105	•	100%	0.39[0.24,0.64]	
ol)					
			_		
	n/N 18/118	n/N n/N 18/118 41/105 118 105	n/N M-H, Fixed, 95% CI 18/118 41/105 118 105	n/N n/N M-H, Fixed, 95% Cl 18/118 41/105 100% 118 105 100%	

Favours control 0.1 0.2 0.5 1 2 5 10 Favours treatment

Analysis 7.7. Comparison 7 Augmented prenatal care versus routine care, Outcome 7 Average reported time with nurse per visit - about 30 minutes.

Study or subgroup	Treatment	Control		Risk Rat	io			Weight	Risk Ratio	
	n/N	n/N		M-H, Fixed, 9	95% CI				M-H, Fixed, 95% Cl	
Klerman 2001	73/118	50/105						100%	1.3[1.02,1.66]	
Total (95% CI)	118	105		-	•			100%	1.3[1.02,1.66]	
Total events: 73 (Treatment), 50 (Co	ontrol)									
Heterogeneity: Not applicable										
Test for overall effect: Z=2.09(P=0.0	4)									
		Envours control	0.1 0.2	0.5 1	2	5	10	Equation treatment		

Favours control 0.1 0.2 0.5 1 2 5 10 Favours treatment

Analysis 7.8. Comparison 7 Augmented prenatal care versus routine care, Outcome 8 Average time reported with nurse per visit - about 45 minutes.

Study or subgroup	Treatment	Control	Risk Ratio						Weight	Risk Ratio	
	n/N	n/N			М-Н, F	ixed,	95% CI				M-H, Fixed, 95% CI
Klerman 2001	16/118	6/105			1					100%	2.37[0.96,5.84]
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



Study or subgroup	Treatment	Control			Ri	sk Rat	tio			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% CI
Total (95% CI)	118	105								100%	2.37[0.96,5.84]
Total events: 16 (Treatment), 6 (Contro	ol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.88(P=0.06)				1							
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.9. Comparison 7 Augmented prenatal care versus routine care, Outcome 9 Average time reported with nurse per visit - about 60 minutes.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N			M-H, Fi	ixed,	95% CI				M-H, Fixed, 95% CI
Klerman 2001	8/118	5/105					+			100%	1.42[0.48,4.22]
Total (95% CI)	118	105						-		100%	1.42[0.48,4.22]
Total events: 8 (Treatment), 5 (Control	l)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.64(P=0.52)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.10. Comparison 7 Augmented prenatal care versus routine care, Outcome 10 Average reported time from entering to leaving clinic - about 1 hour.

Study or subgroup	Treatment	Control			Ri	sk Rat	io			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% CI
Klerman 2001	20/118	17/105								100%	1.05[0.58,1.89]
Total (95% CI)	118	105			-	\blacklozenge				100%	1.05[0.58,1.89]
Total events: 20 (Treatment), 17 (Cont	rol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.15(P=0.88)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.11. Comparison 7 Augmented prenatal care versus routine care, Outcome 11 Average reported time from entering to leaving clinic - about 1.5-2.5 hours.

Study or subgroup	Treatment	Control			Ri	sk Ra	tio			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed,	95% CI				M-H, Fixed, 95% Cl
Klerman 2001	83/118	50/105					+			100%	1.48[1.17,1.86]
Total (95% CI)	118	105					•			100%	1.48[1.17,1.86]
Total events: 83 (Treatment), 50 (Con	trol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=3.29(P=0)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



Analysis 7.12. Comparison 7 Augmented prenatal care versus routine care, Outcome 12 Average reported time from entering to leaving clinic - about 3 hours or more.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N		M-H, Fixed, 95% Cl							M-H, Fixed, 95% Cl
Klerman 2001	13/118	31/105								100%	0.37[0.21,0.67]
Total (95% CI)	118	105								100%	0.37[0.21,0.67]
Total events: 13 (Treatment), 31 (C	ontrol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=3.26(P=0)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.13. Comparison 7 Augmented prenatal care versus routine care, Outcome 13 Same nurse reported as providing care each visit.

Study or subgroup	Treatment	Control	Risk Ratio	Weight	Risk Ratio	
	n/N	n/N	M-H, Fixed, 95% CI		M-H, Fixed, 95% CI	
Klerman 2001	112/118	88/105	+	100%	1.13[1.03,1.24]	
Total (95% CI)	118	105	•	100%	1.13[1.03,1.24]	
Total events: 112 (Treatment), 88 (C	Control)					
Heterogeneity: Not applicable						
Test for overall effect: Z=2.6(P=0.01))			L .		

Favours control 0.1 0.2 0.5 1 2 5 10 Favours treatment

Analysis 7.14. Comparison 7 Augmented prenatal care versus routine care, Outcome 14 Nurse rated as "very helpful".

Study or subgroup	Treatment	Control	Risk Ratio	Weight	Risk Ratio	
	n/N	n/N	M-H, Fixed, 95% CI		M-H, Fixed, 95% CI	
Klerman 2001	107/118	82/105	+	100%	1.16[1.03,1.3]	
Total (95% CI)	118	105	•	100%	1.16[1.03,1.3]	
Total events: 107 (Treatment),	, 82 (Control)					
Heterogeneity: Not applicable						
Test for overall effect: Z=2.51(F	P=0.01)					
		Faure as at a l	01 02 05 1 2 5	10		

Favours control 0.1 0.2 0.5 1 2 5 10 Favours treatment

Analysis 7.15. Comparison 7 Augmented prenatal care versus routine care, Outcome 15 Knowledge of risk factors - told she or her baby might be "at risk" or "have problems".

Study or subgroup	Treatment	Control		Ri	sk Rat	tio			Weight	Risk Ratio	
	n/N	n/N			М-Н, F	ixed,	95% CI				M-H, Fixed, 95% CI
Klerman 2001	50/118	30/105		<mark></mark>				1		100%	1.48[1.03,2.14]
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



itudy or subgroup Treatmen		Control			Ris	sk Rat	tio			Weight	Risk Ratio
	n/N n/		•		M-H, F	ixed,	95% CI				M-H, Fixed, 95% CI
Total (95% CI)	118	105					•			100%	1.48[1.03,2.14]
Total events: 50 (Treatment), 30 (Contro	ol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=2.1(P=0.04)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.16. Comparison 7 Augmented prenatal care versus routine care, Outcome 16 Knowledge of risk factors - changed behavior during pregnancy in response to information about risks/problems.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N		M-H, Fixed, 95% Cl							M-H, Fixed, 95% Cl
Klerman 2001	35/118	14/105					-	-		100%	2.22[1.27,3.9]
Total (95% CI)	118	105						-		100%	2.22[1.27,3.9]
Total events: 35 (Treatment), 14 (Cont	trol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=2.79(P=0.01)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.17. Comparison 7 Augmented prenatal care versus routine care, Outcome 17 Knowledge of risk factors - told how much weight to gain.

Study or subgroup	Treatment	Control		Risk Ratio						Weight	Risk Ratio
	n/N	n/N		M-H, Fixed, 95% Cl							M-H, Fixed, 95% Cl
Klerman 2001	108/118	80/105				+				100%	1.2[1.07,1.35]
Total (95% CI)	118	105				•				100%	1.2[1.07,1.35]
Total events: 108 (Treatment),	80 (Control)										
Heterogeneity: Not applicable						ĺ					
Test for overall effect: Z=2.99(F	?= 0)				1						
		Eavours control	0.1	0.2	0.5	1	2	5	10	Eavours treatment	

Favours control 0.1 0.2 0.5 1 2 5 10 Favours treatment

Analysis 7.18. Comparison 7 Augmented prenatal care versus routine care, Outcome 18 Self-report of behavioral change/status - regular vitamin-mineral supplementation.

Study or subgroup	Treatment	Control		Risk Ratio						Weight	Risk Ratio
	n/N	n/N		M-H, Fixed, 95% Cl							M-H, Fixed, 95% Cl
Klerman 2001	93/118	82/105				+				100%	1.01[0.88,1.16]
Total (95% CI)	118	105				•				100%	1.01[0.88,1.16]
Total events: 93 (Treatment), 82 (Con	trol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.13(P=0.9)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



Analysis 7.19. Comparison 7 Augmented prenatal care versus routine care, Outcome 19 Self-report of behavioral change/status - smoking cessation (for smokers only).

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N			M-H, F	ixed,	95% CI				M-H, Fixed, 95% Cl
Klerman 2001	11/118	7/105					I			100%	1.4[0.56,3.48]
Total (95% CI)	118	105			-					100%	1.4[0.56,3.48]
Total events: 11 (Treatment), 7 (Contro	ol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.72(P=0.47)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.20. Comparison 7 Augmented prenatal care versus routine care, Outcome 20 Self-report of behavioral change/status - perceived mastery.

Study or subgroup	Treatment	Control	Risk Ratio	Weight	Risk Ratio
	n/N	n/N	M-H, Fixed, 95% Cl		M-H, Fixed, 95% Cl
Klerman 2001	55/118	33/105		100%	1.48[1.05,2.09]
Total (95% CI)	118	105	•	100%	1.48[1.05,2.09]
Total events: 55 (Treatment), 33	(Control)				
Heterogeneity: Not applicable					
Test for overall effect: Z=2.26(P=0	0.02)			_1	
0 7 11	0.02)				

Favours control 0.1 0.2 0.5 1 2 5 10 Favours treatment

Analysis 7.21. Comparison 7 Augmented prenatal care versus routine care, Outcome 21 Mean birthweight.

Study or subgroup	Tre	eatment	Control		Mean Difference				Weight	Mean Difference	
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	ixed, 95% CI				Fixed, 95% CI
Klerman 2001	311	3076 (584)	296	3032 (603)						100%	44[-50.51,138.51]
Total ***	311		296				•			100%	44[-50.51,138.51]
Heterogeneity: Not applicable											
Test for overall effect: Z=0.91(P=0.36)											
			Fa	vours control	-1000	-500	0	500	1000	Favours trea	itment

Analysis 7.22. Comparison 7 Augmented prenatal care versus routine care, Outcome 22 Low birthweight (less than 2500 g).

Study or subgroup	Treatment	Control			Risk Ratio		Weight	Risk Ratio	
	n/N	n/N		M-H	, Fixed, 959	% CI			M-H, Fixed, 95% CI
Klerman 2001	39/311	25/296						100%	1.48[0.92,2.39]
Total (95% CI)	311	296			•			100%	1.48[0.92,2.39]
		Favours control	0.01	0.1	1	10	100	Favours treatment	



Study or subgroup	Treatment n/N	Control n/N	Risk Ratio M-H, Fixed, 95% Cl					Weight	Risk Ratio M-H, Fixed, 95% CI
Total events: 39 (Treatment),	25 (Control)								
Heterogeneity: Not applicable	2								
Test for overall effect: Z=1.63(P=0.1)								
		Favours control	0.01	0.1	1	10	100	Favours treatment	

Analysis 7.23. Comparison 7 Augmented prenatal care versus routine care, Outcome 23 Mean gestational weight age, week.

Study or subgroup	Tre	eatment	c	ontrol	Mean Difference			Weight M	lean Difference		
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	ixed, 95% (21			Fixed, 95% CI
Klerman 2001	311	39 (2.6)	296	38.7 (2.8)			+			100%	0.3[-0.13,0.73]
Total ***	311		296				•			100%	0.3[-0.13,0.73]
Heterogeneity: Not applicable											
Test for overall effect: Z=1.37(P=0.1	7)										
			Fa	vours control	-10	-5	0	5	10	Favours treatme	nt

Analysis 7.24. Comparison 7 Augmented prenatal care versus routine care, Outcome 24 Preterm birth.

Study or subgroup	Treatment	Control	Risk Ratio						Weight	Risk Ratio	
	n/N	n/N			M-H, Fi	ixed, 9	95% CI				M-H, Fixed, 95% CI
Klerman 2001	33/311	41/296			_					100%	0.77[0.5,1.18]
Total (95% CI)	311	296								100%	0.77[0.5,1.18]
Total events: 33 (Treatment), 41 (0	Control)										
Heterogeneity: Tau ² =0; Chi ² =0, df=	=0(P<0.0001); I ² =100%										
Test for overall effect: Z=1.22(P=0.	22)										
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.25. Comparison 7 Augmented prenatal care versus routine care, Outcome 25 Intrauterine growth restriction.

Study or subgroup	Treatment	Control		Risk Ratio					Weight	Risk Ratio	
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% Cl
Klerman 2001	37/311	27/296					-			100%	1.3[0.82,2.09]
Total (95% CI)	311	296								100%	1.3[0.82,2.09]
Total events: 37 (Treatment), 27 (Cont	rol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.11(P=0.27)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.26. Comparison 7 Augmented prenatal care versus routine care, Outcome 26 Low Apgar score - at 1 minute.

Study or subgroup	Treatment	Control		Risk Ratio						Weight	Risk Ratio
	n/N	n/N		M-H, Fixed, 95% CI					M-H, Fixed, 95% CI		
Klerman 2001	34/311	40/296								100%	0.81[0.53,1.24]
Total (95% CI)	311	296								100%	0.81[0.53,1.24]
Total events: 34 (Treatment), 40 (Cont	rol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.97(P=0.33)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.27. Comparison 7 Augmented prenatal care versus routine care, Outcome 27 Low Apgar score - at 5 minutes.

Study or subgroup	oup Treatment Control Risk Ratio						Weight	Risk Ratio			
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% CI
Klerman 2001	5/311	3/296					1			100%	1.59[0.38,6.58]
Total (95% CI)	311	296								100%	1.59[0.38,6.58]
Total events: 5 (Treatment), 3 (Cont	rol)										
Heterogeneity: Tau ² =0; Chi ² =0, df=0	(P<0.0001); I ² =100%										
Test for overall effect: Z=0.64(P=0.52	2)			1	1						
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.28. Comparison 7 Augmented prenatal care versus routine care, Outcome 28 Neonatal intensive care unit stay.

Study or subgroup	Treatment	Control		Risk Ratio				Weight	Risk Ratio		
	n/N	n/N			M-H, Fi	ixed, 9	95% CI				M-H, Fixed, 95% Cl
Klerman 2001	33/311	44/296			+					100%	0.71[0.47,1.09]
Total (95% CI)	311	296								100%	0.71[0.47,1.09]
Total events: 33 (Treatment), 44 (Cont	rol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.56(P=0.12)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.29. Comparison 7 Augmented prenatal care versus routine care, Outcome 29 Congenital anomalies.

Study or subgroup	Treatment	Control	ol Risk Ratio						Weight	Risk Ratio	
	n/N	n/N		M-H, Fixed, 95% Cl					M-H, Fixed, 95% Cl		
Klerman 2001	5/311	7/296			-					100%	0.68[0.22,2.12]
Total (95% CI)	311	296								100%	0.68[0.22,2.12]
Total events: 5 (Treatment), 7 (Control))										
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	



Study or subgroup	Treatment	Control	Risk Ratio M-H, Fixed, 95% Cl				Weight	Risk Ratio			
	n/N	n/N			M-H, F	ixed,	95% CI				M-H, Fixed, 95% CI
Heterogeneity: Not applicable											
Test for overall effect: Z=0.67(P=0.51)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 7.30. Comparison 7 Augmented prenatal care versus routine care, Outcome 30 Mean maternal weight gain during pregnancy, kg - if BMI less than 19.8 (at risk).

Study or subgroup	Tre	eatment	c	ontrol	l Mean Di		Mean Difference			Weight I	Mean Difference
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	xed, 95% CI				Fixed, 95% CI
Klerman 2001	311	15.3 (6.3)	296	14.3 (6.4)						100%	1[-0.01,2.01]
Total ***	311		296				•			100%	1[-0.01,2.01]
Heterogeneity: Not applicable											
Test for overall effect: Z=1.94(P=0.0	5)										
			Fa	vours control	-10	-5	0	5	10	Favours treatme	nt

Analysis 7.31. Comparison 7 Augmented prenatal care versus routine care, Outcome 31 Mean maternal weight gain during pregnancy, kg - if BMI is greater than or equal to 19.8 (not at risk).

Study or subgroup	Tre	eatment	с	ontrol	Mean Difference		ice		Weight	Mean Difference	
	Ν	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% (3			Fixed, 95% CI
Klerman 2001	311	12.9 (8.2)	296	12.8 (7.6)						100%	0.1[-1.16,1.36]
Total ***	311		296				•			100%	0.1[-1.16,1.36]
Heterogeneity: Not applicable											
Test for overall effect: Z=0.16(P=0.88	3)										
			Fa	vours control	-10	-5	0	5	10	Favours treatme	ent

Analysis 7.32. Comparison 7 Augmented prenatal care versus routine care, Outcome 32 Caesarean deliveries.

Study or subgroup	Treatment	Control		Risk Ratio						Weight	Risk Ratio
	n/N	n/N			М-Н, F	ixed,	95% CI				M-H, Fixed, 95% CI
Klerman 2001	43/311	51/296							100%	0.8[0.55,1.17]	
Total (95% CI)	311	296								100%	0.8[0.55,1.17]
Total events: 43 (Treatment), 51	(Control)										
Heterogeneity: Tau ² =0; Chi ² =0, c	df=0(P<0.0001); I ² =100%										
Test for overall effect: Z=1.16(P=	:0.25)										
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Comparison 8. Group maternal education versus routine care

Outcome or subgroup title	No. of studies	No. of partici- pants	Statistical method	Effect size
1 Back or pelvic pain (during labour)	1	200	Risk Ratio (M-H, Fixed, 95% CI)	0.70 [0.54, 0.90]
2 Length of labour (hours) - active phase	1	200	Mean Difference (IV, Fixed, 95% CI)	-1.10 [-1.64, -0.56]
3 Length of labour (minutes) - second stage	1	200	Mean Difference (IV, Fixed, 95% CI)	-8.7 [-12.60, -4.80]
4 Vaginal delivery	1	200	Risk Ratio (M-H, Fixed, 95% CI)	1.08 [1.00, 1.16]
5 Oxytocin	1	200	Risk Ratio (M-H, Fixed, 95% CI)	1.12 [0.93, 1.35]
6 Episiotomy	1	200	Risk Ratio (M-H, Fixed, 95% CI)	1.13 [0.85, 1.50]
7 Birthweight	1	200	Mean Difference (IV, Fixed, 95% CI)	14.0 [-55.11, 83.11]
8 Increased daily activity	1	200	Risk Ratio (M-H, Fixed, 95% CI)	1.28 [0.88, 1.85]
9 Headache	1	200	Risk Ratio (M-H, Fixed, 95% CI)	4.0 [1.16, 13.75]
10 Sleep disturbance	1	200	Risk Ratio (M-H, Fixed, 95% CI)	0.81 [0.64, 1.03]

Analysis 8.1. Comparison 8 Group maternal education versus routine care, Outcome 1 Back or pelvic pain (during labour).

Study or subgroup	Treatment	Control			Ri	sk Rat	tio			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed,	95% CI				M-H, Fixed, 95% CI
Mehdizadeh 2005	46/100	66/100			+					100%	0.7[0.54,0.9]
Total (95% CI)	100	100			-	▶				100%	0.7[0.54,0.9]
Total events: 46 (Treatment), 66 (Cont	rol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=2.78(P=0.01)					1						
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 8.2. Comparison 8 Group maternal education versus routine care, Outcome 2 Length of labour (hours) - active phase.

Study or subgroup	Tre	eatment	с	ontrol		Me	ean Differer	ice		Weight	Mean Difference
	N	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% C	3			Fixed, 95% CI
Mehdizadeh 2005	100	3.8 (1.4)	100	4.9 (2.4)			+			100%	-1.1[-1.64,-0.56]
Total ***	100		100				•			100%	-1.1[-1.64,-0.56]
Heterogeneity: Not applicable											
			Fa	vours control	-10	-5	0	5	10	Favours trea	tment



Study or subgroup	ıdy or subgroup Tre			Control		Me	an Differer	ice		Weight Mean Difference
	N	Mean(SD)	Ν	Mean(SD)		Fi	ixed, 95% (:1		Fixed, 95% CI
Test for overall effect: Z=3.96(H	P<0.0001)				_	1				
			F	avours control	-10	-5	0	5	10	Favours treatment

Analysis 8.3. Comparison 8 Group maternal education versus routine care, Outcome 3 Length of labour (minutes) - second stage.

Study or subgroup	Tre	eatment	c	ontrol		Me	ean Differend	e		Weight	Mean Difference
	N	Mean(SD)	Ν	Mean(SD)		F	ixed, 95% Cl				Fixed, 95% CI
Mehdizadeh 2005	100	17 (10.5)	100	25.7 (16.9)			+			100%	-8.7[-12.6,-4.8]
Total ***	100		100				•			100%	-8.7[-12.6,-4.8]
Heterogeneity: Not applicable											
Test for overall effect: Z=4.37(P	<0.0001)										
			Fa	vours control	-100	-50	0	50	100	Favours treatme	nt

-avours control -100 -50 0 50 100 Favours

Analysis 8.4. Comparison 8 Group maternal education versus routine care, Outcome 4 Vaginal delivery.

Study or subgroup	Treatment	Control			Risk Ratio			Weight	Risk Ratio
	n/N	n/N		M-H	, Fixed, 95%	6 CI			M-H, Fixed, 95% Cl
Mehdizadeh 2005	97/100	90/100			+			100%	1.08[1,1.16]
Total (95% CI)	100	100			۲			100%	1.08[1,1.16]
Total events: 97 (Treatment), 90 (Contr	ol)								
Heterogeneity: Not applicable									
Test for overall effect: Z=1.99(P=0.05)									
		Favours control	0.01	0.1	1	10	100	Favours treatment	

Analysis 8.5. Comparison 8 Group maternal education versus routine care, Outcome 5 Oxytocin.

Study or subgroup	Treatment	Control			Ri	sk Rat	io			Weight	Risk Ratio
	n/N	n/N			М-Н, F	ixed, 9	5% CI				M-H, Fixed, 95% Cl
Mehdizadeh 2005	73/100	65/100				+				100%	1.12[0.93,1.35]
Total (95% CI)	100	100				•				100%	1.12[0.93,1.35]
Total events: 73 (Treatment), 65 (Cont	trol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.22(P=0.22)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 8.6. Comparison 8 Group maternal education versus routine care, Outcome 6 Episiotomy.

Study or subgroup	Treatment	Control			Ri	sk Rati	io			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	5% CI				M-H, Fixed, 95% CI
Mehdizadeh 2005	52/100	46/100				-				100%	1.13[0.85,1.5]
Total (95% CI)	100	100				•				100%	1.13[0.85,1.5]
Total events: 52 (Treatment), 46 (Cont	trol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=0.85(P=0.4)					1						
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 8.7. Comparison 8 Group maternal education versus routine care, Outcome 7 Birthweight.

Study or subgroup	Tre	eatment	с	ontrol	Mean Difference			Weight I	Mean Difference		
	Ν	Mean(SD)	Ν	Mean(SD)		Fi	xed, 95% C	l			Fixed, 95% CI
Mehdizadeh 2005	100	3190 (284)	100	3176 (209)					_	100%	14[-55.11,83.11]
Total ***	100		100							100%	14[-55.11,83.11]
Heterogeneity: Not applicable											
Test for overall effect: Z=0.4(P=0.69)					1						
			Fa	vours control	-100	-50	0	50	100	Favours treatme	nt

Analysis 8.8. Comparison 8 Group maternal education versus routine care, Outcome 8 Increased daily activity.

Study or subgroup	Treatment	Control			Ri	sk Rati	io			Weight	Risk Ratio
	n/N	n/N			М-Н, F	ixed, 9	5% CI				M-H, Fixed, 95% CI
Mehdizadeh 2005	41/100	32/100				++	-			100%	1.28[0.88,1.85]
Total (95% CI)	100	100								100%	1.28[0.88,1.85]
Total events: 41 (Treatment), 32 (Contr	ol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.31(P=0.19)											
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

Analysis 8.9. Comparison 8 Group maternal education versus routine care, Outcome 9 Headache.

Study or subgroup	Treatment	Control			Risk Ratio			Weight	Risk Ratio
	n/N	n/N		M-H	H, Fixed, 95%	6 CI			M-H, Fixed, 95% Cl
Mehdizadeh 2005	12/100	3/100				+		100%	4[1.16,13.75]
Total (95% CI)	100	100						100%	4[1.16,13.75]
Total events: 12 (Treatment), 3 (Contro	ol)								
Heterogeneity: Not applicable									
Test for overall effect: Z=2.2(P=0.03)									
		Favours control	0.01	0.1	1	10	100	Favours treatment	

Analysis 8.10. Comparison 8 Group maternal education versus routine care, Outcome 10 Sleep disturbance.

Study or subgroup	Treatment	Control			Ri	sk Rat	io			Weight	Risk Ratio
	n/N	n/N			M-H, F	ixed, 9	95% CI				M-H, Fixed, 95% CI
Mehdizadeh 2005	52/100	64/100			-	+				100%	0.81[0.64,1.03]
Total (95% CI)	100	100			•					100%	0.81[0.64,1.03]
Total events: 52 (Treatment), 64 (Con	trol)										
Heterogeneity: Not applicable											
Test for overall effect: Z=1.7(P=0.09)					1						
		Favours control	0.1	0.2	0.5	1	2	5	10	Favours treatment	

APPENDICES

Appendix 1. Search terms for CINAHL

1982 to April 2006 (authors wrote and ran the search)

- 1. exp pregnancy/ or pregnancy.ti, ab, sh.
- 2. prenatal. ti, ab, sh.
- 3. antenatal. ti, ab, sh.
- 4. birth. ti, ab, sh. OR childbirth. ti, ab, sh.
- 5. early intervention (education)/ OR education/ OR education. ti, ab, sh.
- 6. preparation. ti, ab, sh.
- 7.5 OR 6
- 8. (mothers/ or motherhood. ti, ab, sh.) OR (father/ or fatherhood. ti, ab, sh.) OR (parent/ or parenthood)
- 9. 1 OR 2 OR 3 OR 4 OR 8

10.7 AND 9

MeSH terms were consulted and the appropriate terms were used in conjunction with natural language to create a broad search. Natural language was also used to describe concepts which may be termed differently in each database.

Appendix 2. Search terms for EMBASE

1980 to April 2006 (authors wrote and ran the search)

- 1. birth/ OR childbirth/ OR pregnancy.exp
- 2. prenatal/ OR antenatal.exp
- 3. 1 AND 2
- 4. 3 AND Education.exp
- 5. randomized controlled trial.exp AND 4
- 6. pain.exp/ OR knowledge acquisition.exp/ OR self-confidence.exp/ OR anxiety.exp/ OR decision-making.exp/ OR obstetrical intervention.exp/ OR partner involvement.exp/ OR breast-feeding.exp/ OR infant care.exp/ OR social support.exp
- 7. 6 AND randomized controlled trial.exp
- 8. 5 AND 7

Appropriate EMBASE keywords were selected to formulate a search strategy. Natural language was also employed to ensure a broad search. Terms were 'exploded' (exp) to access all related articles.

Appendix 3. Search terms for ERIC database

1984 to April 2006 (authors wrote and ran the search)

- 1. birth/ OR childbirth.ab, ti, hw, id OR pregnancy
- 2. (adult learning/ or didactic.ab, ti, hw, id) OR (class or course).ab, ti, hw, id. OR exp teaching methods/ OR exp educational research/





- 3. 1 AND 2
- 4. parenthood education/ OR preparation.ab, ti, hw, id OR childbirth.ab, ti, hw ADJ education.ab, ti, hw, id.
- 5. 3 OR 4
- 6. 5 NOT school ADJ curriculum

The thesaurus for the ERIC database was consulted and the appropriate terms were selected to formulate a search strategy. Terms that are followed by a forward slash are the controlled vocabulary for the ERIC thesaurus. Natural language was also employed to ensure a broad search. Childbirth education or preparation; teaching methods; and stress or anxiety reduction were the three concepts covered by this search.

Appendix 4. Search strategy for PsycINFO

1988 to April 2006 (authors wrote and ran the search)

- 1. exp birth/ OR childbirth.ti, ab, hw, id. OR pregnancy/ OR prenatal.ti, ab, hw, id OR antenatal.ti, ab, hw, id.
- 2. exp education/ or preparation.ti, ab, hw, id.
- 3. 1 and 2

Appropriate terms were selected from the PsycINFO thesaurus and the forward slash denotes this usage. Natural language was employed to help broaden the search. Terms were 'exploded' (exp) to access all related articles.

WHAT'S NEW

Date	Event	Description
7 July 2011	Amended	Search updated. Fifty-eight reports added to Studies awaiting classification.

HISTORY

Protocol first published: Issue 1, 1999 Review first published: Issue 4, 2000

Date	Event	Description
29 January 2009	Amended	The contact person for this review is now Jane Sandall.
3 September 2008	Amended	Converted to new review format.
17 May 2007	New citation required but conclusions have not changed	New author helped to prepare the update.
1 April 2006	New search has been performed	Search updated. Three new trials (Davis 1987; Klerman 2001; Mehdizadeh 2005) included and 15 new trials excluded (Astbury 1980; D'Andrea 1994; De Nuncio 2000; Escott 2005; Ford 2002; Leitch 1999; Linares 2006; Manandhar 2004; O'Cathain 2002; Pol- ley 2002; Schachman 2004; Schonberger 2004; Spence-Cagle 1984; Springer 1996; Turan 2001). The conclusions concerning the effects of antenatal education remain largely the same and are generally unknown.



CONTRIBUTIONS OF AUTHORS

For the update, Jane Sandall and Anita J Gagnon (AJG) collaborated on assessing the quality of and extracting the data for included studies. AJG conducted the quantitative analyses. AJG previously prepared the protocol and undertook the initial review, applying the inclusion criteria and extracting data for the included studies for the first version of this review.

DECLARATIONS OF INTEREST

None known.

SOURCES OF SUPPORT

Internal sources

- McGill University, Montreal, Canada.
- McGill University Health Centre Montreal, Canada.
- The SMBD Jewish General Hospital (for initial review), Montreal, Canada.
- King's College, London, UK.

External sources

- Le Fonds de la recherche en santé du Québec (FRSQ), Canada.
- La Fondation de recherche en sciences infirmières du Québec (FRESIQ)(for initial review), Canada.
- Groupe de recherche interuniversitaire en sciences infirmieres de Montreal (GRISIM), Canada.

INDEX TERMS

Medical Subject Headings (MeSH)

*Labor, Obstetric; *Parenting; Fathers [education]; Mothers [education]; Prenatal Care [methods]; Randomized Controlled Trials as Topic; Vaginal Birth after Cesarean

MeSH check words

Female; Humans; Pregnancy