

Labour induction in women at term with unfavourable cervix

Submission date 16/08/2009	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 18/09/2009	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 18/09/2009	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Study information

Scientific Title
Sustained-release dinoprostone vaginal pessary with concurrent high-dose oxytocin infusion compared sustained-release dinoprostone vaginal pessary followed six hours later by high dose oxytocin infusion for labour induction in women at term with unfavourable cervix: a randomised controlled trial

Study objectives

Can we use concurrent oxytocin infusion with dinoprostone vaginal pessary for cervical ripening and labour induction?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Istanbul Bakirkoy Women and Children Hospitals Local Ethics Board approved on the 7th November 2008 (ref: 162)

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Labour induction

Interventions

Women who were assigned randomly to receive the sustained-released dinoprostone (Propess®, Vitalis, Turkey) with concurrent high-dose oxytocin (Group A) had a single dose placed high into the vaginal fornix. This sustained-released product releases dinoprostone at a low but steady rate (0.3 mg/h). It remained in the vagina for up to 12 hours, as recommended by the manufacturer. At the same time, oxytocin infusion at 4 milliunits/min was started for all participants. Oxytocin infusion was doubled every 30 minutes to a maximum of 40 milliunits/min or until four contractions in 10 minutes was achieved.

Women who were assigned randomly to receive the sustained-released dinoprostone followed six hours later by high-dose oxytocin (Group B) had a single dose placed high into the vaginal fornix. A standard high-dose of intravenous oxytocin was administered 6 hours after the insertion of the vaginal pessary. An initial dose of 4 mU/min was increased at 30 minute intervals by 4 mU/min to a maximum dose 40 mU/min or until four contractions in 10 minutes was achieved.

Intervention Type

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

Oxytocin, dinoprostone

Primary outcome(s)

Number (rate) of women who went to vaginally deliver within 24 hours of the initiation of the protocol.

Key secondary outcome(s)

1. Incidence of excess uterine activity (uterine hyperstimulation or uterine tachysystole)
2. Labour induction-to-delivery interval
3. Labour induction-to-active phase interval (defined as at least 6 uterine contractions per 20-minute intervals, with at least 70% effaced cervix and a cervical dilatation of greater than or equal to 4)
4. Total number of doses of dinoprostone pessary used
5. Meconium-stained liquor
6. Mode of delivery
7. Instrumental delivery rate
8. Maternal satisfaction score for the birth process obtained within 24 hours of delivery (a visual analog scale (VAS) with a range of 0 to 10, with higher score denoting greater satisfaction, was used to gauge maternal satisfaction)
9. Visual analogue scale pain score (ranged from 0 to 10, with 0 representing no pain to 10 representing unbearable pain)
10. Rates of maternal and neonatal complications:
 - 10.1. Maternal complications included:
 - 10.1.1. Incidence of maternal side effects (nausea, vomiting, diarrhoea, pyrexia)
 - 10.1.2. Postpartum haemorrhage (blood loss greater than 500)
 - 10.1.3. Third- or fourth-degree lacerations
 - 10.1.4. Intrapartum chorioamnionitis (defined as temperature greater than or equal to 38°C accompanied by maternal or fetal tachycardia [greater than 160 beats/min], uterine tenderness, malodorous amniotic fluid discharge, and/or maternal leukocytosis [white blood cell count greater than 15,000 cell/min³])
 - 10.1.5. Postpartum endometritis (defined as temperature greater than or equal to 38°C accompanied by uterine tenderness and/or purulent or foul-smelling lochia beyond the first 24 hours after delivery)
 - 10.2. Neonatal complications noted were:
 - 10.2.1. Apgar scores of less than 7 at 5 minutes
 - 10.2.2. Neonatal jaundice
 - 10.2.3. Rate of admission to the neonatal intensive care unit

Secondary analysis based on parity was also planned.

Completion date

01/02/2009

Eligibility

Key inclusion criteria

1. Single live fetus in cephalic presentation
2. Gestational age greater than or equal to 37 weeks as determined by the last menstrual period or by a first- or second-trimester ultrasound scan
3. Bishop score less than or equal to 4
4. Females aged 20 - 40 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

1. Any contraindication to vaginal delivery
2. Previous caesarean section
3. Multiple pregnancy
4. Estimated fetal weight greater than 4500 g
5. Breech presentation
6. Antepartum haemorrhage
7. Evidence of fetal distress

Date of first enrolment

01/10/2007

Date of final enrolment

01/02/2009

Locations**Countries of recruitment**

Türkiye

Study participating centre

Atakent Mah. Soyak Olypiakent Sitesi D10-57

Istanbul

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Sponsor information**Organisation**

Istanbul Bakirkoy Women and Children Hospital (Turkey)

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

Istanbul Bakirkoy Women and Children Hospital (Turkey) - Department of Obstetrics and Gynecology

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration