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## Brief communication

# Extra-amniotic prostaglandin- $E_2$ for termination in the second and early third trimesters

## A.A. Rouzi<sup>a,\*</sup>, H. Hawasawi<sup>b</sup>, H. Abduljabbar<sup>a</sup>

<sup>a</sup>Departments of Obstetrics and Gynecology, King Abdulaziz University Hospital, Jeddah, Saudi Arabia <sup>b</sup>King Fahd Armed Forces Hospital, Jeddah, Saudi Arabia

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Termination of pregnancy in the second and early third trimesters has always been a common challenge in obstetrical practice. In the United States dilation and evacuation in the second trimester is preferred [1], while in the United Kingdom prostaglandin analogues are favored [2]. Between November 1995 and June 1998, 43 consecutive pregnant women underwent termination of pathological pregnancy in second and early third trimesters according to a protocol using extra-amniotic prostaglandin- $E_2$  (EAPGE<sub>2</sub>) in the Department of Obstetrics and Gynecology at King Fahd Armed Forces Hospital, Jeddah, Saudi Arabia. The protocol was approved by the institution review board. Consent was obtained from all women. Ultrasonography was used routinely to confirm the gestational age. On the day of termination, a 16-French Foley's catheter with a 30-ml balloon was positioned in the internal cervical os

to reach the extra-amniotic space. The rate of infusion of Prostin<sup>TM</sup> (dinoprostone, UPJOHN, Puurs, Belgium) was 1 ml/h initially. This was increased by 1 ml/h every 2 h until a maximum of 3 ml/h was reached or satisfactory uterine response was accomplished. If abortion did not occur after 12 h or if the catheter fell out before achieving satisfactory response, 40 U of oxytocin in 500 ml of normal saline was started at a rate of 30 mU/min.

The indications for termination of pregnancy are listed in Table 1. The gestational age (mean  $\pm$  S.D., range) at the time of termination was  $21.2 \pm 4.97$ , 13-31 weeks. Extra-amniotic PGE<sub>2</sub> was associated with abortion in all women. The mean induction-to-abortion interval was  $8.1 \pm 7.8$ , 1-26 h. Nineteen women needed supplemental oxytocin. Thirteen women underwent surgical evacuation of the uterus after delivery of the fetus. The mean blood loss (clinically estimated) was  $145.8 \pm 120$  ml. There was no major complication related to the use of EAPGE<sub>2</sub>. However,

<sup>\*</sup>Corresponding author. Tel.: +966-26772027; fax: +966-25372502.

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Table 1 Indications for termination of pregnancy

Indication	Number
Intrauterine fetal death	35
Trisomy 13, 18	2
Lethal fetal anomalies	4
Partial mole	1
Premature rupture of membranes remote from term	1

one woman underwent laparotomy because of perforation of the uterus at the time of evacuation of a retained placenta. Repair of the uterus was done. Fifteen women were given antibiotics (nine women prophylactically and six women for febrile morbidity).

Previous studies have confirmed the safety and effectiveness of EAPGE<sub>2</sub> gel [3,4]. The induction-to-abortion interval with EAPGE<sub>2</sub> gel ranged from 14 to 16.8 h. In our study EAPGE<sub>2</sub> was used as an infusion. This approach was safe, effective, and associated with shorter induction-to-abortion interval. Randomized clinical trials comparing

 $EAPGE_2$  gel with  $EAPGE_2$  infusion are required to confirm our findings.

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