Effect of PGI-2 on uterine activity in vivo in non-pregnant ovariectomized goats (Capra hircus)

R. G. Cooke and A. M. Homeida*

University of Liverpool Veterinary Field Station, Leahurst, Chester High Road, Neston, South Wirral, Cheshire L64 7TE, UK; and *Faculty of Veterinary Science, University of Khartoum, Khartoum, Sudan

Summary. Jugular administration of 200 μg PGI-2 salt significantly reduced spontaneous uterine activity in ovariectomized, oestrogen-primed goats; the effect was acute and persisted for about 3 h. Peripheral plasma concentrations of 6-keto-PGF-1α, the stable metabolite of PGI-2, decreased to 50% of initial values after 30 min; but at the start of uterine recovery were in excess of 2 ng.ml⁻¹. Uterine reactivity to both oxytocin and PGF-2α after PGI-2 administration was unaffected.

Keywords: prostacyclin; uterus; motility; goat; PGF-2α; oxytocin

Introduction

Prostaglandins (PG) of the E and F series are generally considered to be the key prostanoids involved in the modulation of uterine activity. However, a role for prostacyclin (PGI-2), a potent vasodilator and inhibitor of platelet aggregation (Moncada et al., 1976), has to be defined. PGI-2 is present in the myometrium and endometrium of non-pregnant and pregnant animals including rats, guinea-pigs, sheep (Jones et al., 1977; Williams et al., 1978), goats (Cooke & Homeida, 1987) and women (Omini et al., 1979). In-vitro experiments have produced conflicting results: PGI-2 may either inhibit (Omini et al., 1979) or stimulate and then inhibit the pregnant human myometrium (Wikland et al., 1983); myometrial activity in the rat, however, is stimulated (Williams et al., 1979). Ly & Challis (1982) showed that PGI-2 inhibited both the electrical and mechanical activity of the non-pregnant sheep uterus in vivo, but responsiveness to PGF-2α and oxytocin were unaffected.

The aims of this study were to examine in ovariectomized goats the effect of PGI-2 on spontaneous uterine activity, and also the effects of PGI-2 on uterine reactivity to both PGF-2α and oxytocin.

Materials and Methods

Animals and treatments. Four mature female goats of mixed breeding but similar age (3–4 years) and weight (40–45 kg) were used. The animals, anaesthetized with sodium thiopentone intravenously and maintained with fluothane and oxygen, were ovariectomized through a left-flank incision. A soft polyethylene, air-filled balloon (27 × 5 mm; Portex Ltd, Hyde, Kent, UK) attached to a more rigid 40-cm length of cannula was inserted into the uterus via a small incision in the uterine wall; intrauterine pressure was monitored by a Bell and Howell pressure transducer and recorded on a Devices MX-2 machine. Details of the techniques have been described previously (Jones & Knifton, 1975). Following a post-surgical recovery period of 10 days all 4 animals were treated daily with 60 μg oestradiol benzoate (Intervet, Cambridge, UK) intramuscularly (i.m.) for 7 days.

On the 6th day of oestradiol administration 200 μg PGI-2 sodium salt (Upjohn Co., Kalamazoo, MI, USA) in 1 ml sterile saline (0.9% w/v NaCl) were injected into the jugular vein of each goat using 23-gauge needles. Uterine activity was recorded for 30 min before and for 2-5 h after PGI-2 administration. Jugular venous blood was collected by direct venepuncture into chilled heparinized tubes immediately before treatment and then at 10-min intervals for 120 min. Plasma was separated, snap-frozen in CO₂-acetone and stored at −20°C; samples were later analysed for the
Fig. 1. Intrauterine pressure in an ovariectomized goat before and after administration of 200 µg PGI-2 salt.

Fig. 2. Mean (± s.e.m.) uterine activity (●) and jugular venous plasma concentrations of 6-keto-PGF-1α (○) in 4 ovariectomized goats before and after administration of 200 µg PGI-2 salt at time 0.

6-oxo-PGF-1α, the stable metabolite of PGI-2. Before administration of PGI-2, 1 ml saline had been injected to assess whether the drug vehicle produced any change in uterine motility.

After a period of recovery (4 h) when uterine activity returned to that observed in the pre-injection phase, dose–response relationships were determined to doses of oxytocin (Universal Biologicals, Cambridge, UK) and PGF-2α (Lutalyse: Upjohn Co.); both were suitably diluted in saline so that each was contained in a maximum of 1 ml, and were injected into the jugular vein. To minimize possible time-related effects doses of oxytocin and PGF-2α were administered in random order; each was separated by a 15-min interval. The same sequence of injections was then repeated 20 min after administration of 200 µg PGI-2 as described above.
Table 1. Mean (± s.e.m.) uterine responses (Montevideo units) in 4 ovariectomized, oestradiol-treated goats to i.v. oxytocin and PGF-2α before and after treatment with 200 μg PGF-2α.

<table>
<thead>
<tr>
<th>Oxytocin (mU)</th>
<th>Response</th>
<th>PGF-2α (μg)</th>
<th>Response</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Before</td>
<td>After</td>
<td>Before</td>
</tr>
<tr>
<td>0</td>
<td>290 ± 26*</td>
<td>28 ± 8</td>
<td>0</td>
</tr>
<tr>
<td>50</td>
<td>510 ± 41</td>
<td>552 ± 36</td>
<td>1</td>
</tr>
<tr>
<td>100</td>
<td>742 ± 61</td>
<td>726 ± 51</td>
<td>2</td>
</tr>
<tr>
<td>200</td>
<td>879 ± 71</td>
<td>1051 ± 80</td>
<td>4</td>
</tr>
</tbody>
</table>

*P < 0.001 compared to corresponding value after treatment.

Uterine responses were expressed in Montevideo units (Caldeyro-Barcia et al., 1957), i.e. the product of mean amplitude and the total number of contractions in a 10-min period.

Assay of 6-keto-PGF-1α. A specific radioimmunoassay for 6-keto-PGF-1α was used as described by Liggins et al. (1980), utilizing their antiserum (4V-208) at a dilution of 1:10 000; 6-keto [5,8,9,11,12,14,15(n)-3H]PGF-1α (5000 c.p.m./tube) was obtained from Amersham International (Bucks, UK). Standard curves were constructed using pure 6-keto-PGF-1α (Upjohn). The recovery of added tracer was 77 ± 15% (mean ± s.e.m., n = 9) and results were corrected for extraction losses; the recovery of standards (20, 50, 500, 1000 pg) added to goat plasma was 19-1 ± 1-9, 52-6 ± 7-8, 489 ± 15-7 and 893 ± 22-8 pg (mean ± s.e.m., n = 6). The non-specific binding was 8%, the sensitivity of the assay 10 pg/tube, and the intra- and inter-coefficients of variation were 6-2 and 12-6%, respectively (n = 12). Antiserum specificity has been reported previously (Liggins et al., 1980).

Data analysis. Data were analysed using two-way ANOVA and Duncan’s multiple range test, and Student’s t test for paired data.

Results

Injection of 1 ml saline vehicle did not have any effect on uterine motility in the 4 goats. Figure 1 shows a representative trace of the effects of PGF-2 (200 μg) on uterine motility in one ovariectomized goat during treatment with oestradiol benzoate. There was a rapid decline in activity within 10 min, with a peak effect after about 30 min. Activity returned slowly and was still significantly (P < 0.01) below pre-injection values after 2-0 h. The results for all 4 animals are presented in Fig. 2. Concentrations of 6-keto-PGF-1α decreased progressively to 50% of initial values after about 30 min; at the start of the recovery period concentrations were still greater than 2 ng.ml⁻¹. Table 1 shows the mean responses to various doses of intravenous oxytocin and PGF-2α before and after treatment with 200 μg PGF-2. Reactivity to both agents, at all dose levels, was not significantly (P < 0.05) affected, i.e. the dose–response curves for both oxytocin and PGF-2α were unaltered by PGF-2, even though spontaneous activity was significantly (P < 0.001) reduced.

Discussion

The results of this study show that PGF-2 inhibits spontaneous uterine motility in the goat; similar results have been reported for the sheep (Lye & Challis, 1982). Such an acute effect of PGF-2 may be of importance in rendering the uterus quiescent before implantation when progesterone concentrations are very low (Thorburn & Schneider, 1972; Irving et al., 1972). This is supported by several observations. While PGF-2 is present in the goat corpus luteum, highest concentrations of its metabolites, 6-keto-PGF-2α, have been found in the myometrium and endometrium, and concentrations increase significantly during early pregnancy (Cooke & Homeida, 1987); in this respect it may be beneficial to compare uterine activity, and responses to PGF-2, in non-pregnant and early pregnant goats. PGF-2 has also been shown to regulate uterine blood flow in the rat (Kennedy &
Zamecnik, 1978), to assist blastocyst implantation in women (Kelly, 1981), and be luteotrophic in the cow both in vitro and in vivo (Milvae & Hansel, 1980).

The prolonged effect of PGI-2 on the goat uterus was surprising considering its short half-life of 3 min in blood or biological buffers (Dusting et al., 1978). It is possible that 6-keto PGF-2α is responsible for the inhibitory effect, but the uterus recovered when plasma concentrations of the metabolite were in excess of 2 ng.ml⁻¹, and also it has been reported that 6-keto-PGF-2α has no effect on human myometrial activity in vitro (Omini et al., 1979). However, bearing in mind the possible species differences in uterine responses to PGI-2 referred to above, it would be useful to examine the effects of the metabolite on uterine motility in ruminants.

As observed in ovariectomized ewes (Lye & Challis, 1982) both oxytocin and PGF-2α were able to overcome the inhibitory effects of PGI-2, suggesting that prostacyclin has little effect on receptors to these hormones. This contrasts to the action of progesterone which blocks uterine responsiveness to both oxytocin and PGF-2α (R. G. Cooke & A. M. Homeida, unpublished observations), probably by inhibiting receptor formation (McCracken et al., 1984).

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References


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