

## PROSTAGLANDIN-INDUCED LUTEOLYSIS IN THE RED DEER HIND

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The effectiveness of the prostaglandin F<sub>2α</sub> analogue, cloprostenol, to induce luteolysis in the red deer hind was investigated by monitoring plasma progesterone concentrations at 4 or 6 h intervals for up to 72 h. The effects of age, dose, ovulation rate and stage of the oestrous cycle were assessed in 2 experiments conducted in successive breeding seasons.

In Experiment 1, 20 pubertal and 20 adult hinds were synchronised for oestrus with intravaginal progesterone (CIDR, Carter Holt Harvey). Fourteen days after progesterone withdrawal (approximately 12 days after the expected time of oestrus; stags were absent), 0, 100, 250 or 500 µg of cloprostenol (*Estrumate*, Coopers) was administered i.m. and the hinds bled by jugular venepuncture. One hind had low progesterone (>0.5 ng/ml) at the time of treatment. All 3 doses were equally effective in lowering progesterone concentrations in both groups of hinds. By 12 h after administration concentrations averaged 0.70-0.93 ng/ml in the 3 treated groups compared with 1.89 ng/ml in the untreated hinds and at 24 h averaged 0.39-0.70 ng/ml and 2.13 ng/ml respectively.

In Experiment 2, adult hinds (n=28) were treated to induce oestrous synchronisation with either mono-ovulation (CIDR or CIDR + PMSG, 200 iu *Folligon*, Intervet) or superovulation (CIDR + PMSG + FSH, 0.5 units *Ovagen*, Immuno-Chemical Products or CIDR + PMSG + FSH + GnRH, 500 µg *Fertagyl*, Intervet). Mean (and range) ovulation rates determined at laparoscopy were CIDR 1.0 (1); CIDR + PMSG 1.29 (1-2); CIDR + PMSG + FSH 4.14 (0-11); CIDR + PMSG + FSH + GnRH 12.29 (6-27). The hinds were then allocated to receive 750 µg of cloprostenol i.m. either 11 or 15 days after progesterone withdrawal (approximately 9 or 13 days after the expected time of oestrus; stags were present). Three hinds were anovulatory or had low plasma progesterone at the time of cloprostenol treatment. In the remaining hinds, plasma progesterone concentrations were reduced within 6 h of cloprostenol and continued to decline thereafter (Table 1).

Table 1. Mean (± sem) ovulation rates and progesterone concentrations.

|                             |       | Mono-ovulation |            | Superovulation |             |
|-----------------------------|-------|----------------|------------|----------------|-------------|
|                             |       | d 11           | d 15       | d 11           | d 15        |
| Ovulation rate              | mean  | 1              | 1          | 9.8 ± 3.68     | 8.1 ± 2.20  |
|                             | range | -              | -          | 2 - 27         | 2 - 18      |
| Plasma progesterone (ng/ml) | 0 h   | 1.50 ± .16     | 2.30 ± .30 | 10.7 ± 3.92    | 10.8 ± 2.31 |
|                             | 12 h  | 0.94 ± .17     | 0.77 ± .11 | 6.39 ± 2.55    | 2.79 ± 0.55 |
|                             | 24 h  | 0.32 ± .02     | 0.37 ± .03 | 1.95 ± 0.88    | 1.10 ± 0.13 |

Cloprostenol was effective in inducing complete luteolysis (<1.0 ng/ml progesterone for at least 24 h) in all 13 hinds treated on d 15 and in 10/12 hinds treated on d 11. The two hinds (1 mono-ovulation, 1 superovulation) in which luteolysis appeared incomplete were characterised by short-lived (12-24 h) reductions in progesterone concentrations.

These results indicate that exogenous prostaglandin F<sub>2α</sub> is luteolytic in both young and adult red deer hinds and in hinds with multiple corpora lutea; however its effectiveness may be incomplete during the earlier stages of luteal function.