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<tr>
<td>Authors(s)</td>
<td>Beltman, Marijke Eileen; Lonergan, Patrick; Diskin, M. G.; Roche, J. F.; Crowe, Mark</td>
</tr>
<tr>
<td>Publication date</td>
<td>2009-04-15</td>
</tr>
<tr>
<td>Publication information</td>
<td>Theriogenology, 71 (7): 1173-1179</td>
</tr>
<tr>
<td>Publisher</td>
<td>Elsevier</td>
</tr>
<tr>
<td>Item record/more information</td>
<td><a href="http://hdl.handle.net/10197/4663">http://hdl.handle.net/10197/4663</a></td>
</tr>
<tr>
<td>Publisher's statement</td>
<td>This is the author's version of a work that was accepted for publication in Theriogenology. Changes resulting from the publishing process, such as peer review, editing, corrections, structural formatting, and other quality control mechanisms may not be reflected in this document. Changes may have been made to this work since it was submitted for publication. A definitive version was subsequently published in Theriogenology (VOL 71, ISSUE 7, (2009)) DOI: <a href="http://dx.doi.org/10.1016/j.theriogenology.2008.12.014">http://dx.doi.org/10.1016/j.theriogenology.2008.12.014</a></td>
</tr>
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<td>Publisher's version (DOI)</td>
<td>10.1016/j.theriogenology.2008.12.014</td>
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Effect of progesterone supplementation in the first week post conception on embryo survival in beef heifers

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Abstract

Progesterone is essential for establishment and maintenance of pregnancy in mammals. The objective of this study was to examine the effect of elevating progesterone during the different physiological stages of early embryo development on embryo survival. Estrus was synchronized in cross-bred beef heifers (n=197, ~2-6 years old) and they were inseminated 12-18 h after estrus onset (=Day 0). Inseminated heifers were randomly assigned to 1 of 3 treatments: (1) Control, n=69; (2) progesterone supplementation using a Controlled Internal Drug Release Device (CIDR) from Day 3 to 6.5, n=64; or (3) progesterone supplementation using a CIDR from Day 4.5 to 8, n=64. Body condition (BCS) and locomotion scores (scale of 1-5) were recorded for all animals. Animals with a locomotion score ≥ 4 (very lame) were excluded. Embryo survival rate was determined at slaughter on Day 25. Conceptus length and weight were recorded and the corpus luteum (CL) of all pregnant animals was dissected and weighed. Supplementation with exogenous progesterone increased (P<0.05) peripheral progesterone concentrations, but did not affect embryo survival rate compared with controls. Mean CL weight, conceptus length and conceptus weight were not different between treatments. There was a positive relationship (P<0.04) between the increase in progesterone concentrations from Days 3 to 6.5 and embryo survival rate in treated heifers and a similar trend existed between the increase from Days 4.5 to 8 (P<0.06). There was also a positive relationship (P < 0.05) between the progesterone concentration on Day 6.5 and the embryo survival rate in treated heifers. A direct correlation was seen between locomotion score and embryo survival rate, with higher (P<0.05) early embryo survival rates in heifers with a lower locomotion score. In conclusion, supplementation with progesterone at different stages of early embryo development increased peripheral progesterone concentration and resulted in a positive association between changes in progesterone concentration during the early luteal phase and embryo survival rate. Supplementation with progesterone had no effect on either CL weight or conceptus size in pregnant animals. Lameness had a significant negative effect on early embryo survival.

Keywords: Cattle, progesterone, embryo survival, locomotion score
1. Introduction

Progesterone is an essential hormone for the establishment and maintenance of pregnancy in mammals [1]. In heifers, only 55% of single inseminations lead to a successful full term pregnancy despite a fertilization rate of 90-95% [2]. A major portion of this conception loss occurs between Days 8 and 16 of pregnancy [3] with relatively little late embryonic or early fetal mortality. Recent data from high producing dairy cows suggests that the percentage of early embryonic loss in such animals is even larger than in heifers [2]. The peripheral concentration of progesterone in early pregnancy has a significant effect on the survival of the embryo/conceptus [4]. Low concentrations of progesterone in the initial days following conception (Day 3-8) are associated with smaller embryos on Day 16 of pregnancy which may produce insufficient interferon-τ to suppress the luteolytic secretion of prostaglandin F2α [5]. Similarly, a delay in the post-ovulatory increase in progesterone has been associated with a decreased pregnancy rate in both cows [6,7] and heifers [2].

Circulating progesterone concentrations can be increased by either exogenous supplementation [8-11] or ovulation of a dominant follicle, using GnRH or hCG for example, to form an accessory corpus luteum (CL) [12-14]. The earliest day on which a dominant follicle is large enough to consistently respond to GnRH or hCG is Day 5 [15]. The accessory CL created by treatment on Day 5 leads to increased plasma progesterone concentrations from Day 8 which reach a maximum on Days 16-17 [16]. However, this increase in progesterone may be too late to influence embryo survival [11,17]. A further study that involved the administration of GnRH during the second wave dominant follicle did not have a consistent positive effect on pregnancy rate [14].

Direct exogenous progesterone supplementation allows more control over the time of administration and results in an immediate increase in progesterone concentrations. This approach may therefore be a more effective method to use to increase circulating progesterone concentrations. This can be achieved by injection of progesterone or the insertion of a progesterone-releasing intravaginal device. Several studies have
described the effect of exogenous progesterone supplementation on development of the embryo. One of the earliest such studies was that of Garrett et al [18] who reported a dramatic advancement in conceptus development by Day 14 following administration of 100 mg progesterone on Days 1 to 4 of the cycle. Mann and Lamming [11] did a meta analysis of all studies that supplemented progesterone during early pregnancy and examined the effect of day of initiation of progesterone supplementation on the subsequent pregnancy rates. They concluded that progesterone supplementation early in pregnancy leads to an overall increase in pregnancy rate of 5%; however, supplementation after Day 6 had no beneficial effect. The same study found that when the initial pregnancy rate was low (<50%) there was a large improvement in pregnancy rate, but when the pregnancy rate was higher (>50%) there was no additional benefit.

More recent studies have shown that insertion of an intravaginal device between Days 2 and 7 increased the survival of the conceptus by 10-15% [6,11]. However, one study found that supplementation commencing within the first 2 days after ovulation and continuing for 7 days suppressed fertility [19] indicating possible interference with the development of the endogenous CL during this period. A recent study reported an increase in pregnancy rate from 35% to 48% when progesterone was supplemented to dairy cows with a used CIDR from Day 3.5 up until Day 10 [20]. This increase in pregnancy rate was ascribed to the embryotrophic effects of progesterone, especially in cows with low concentrations of progesterone at the time of supplementation. The consensus of most studies is that the critical period for progesterone in supporting increased early embryo development rates is from Day 3 to 7. In support of this, a recent study from our group has demonstrated that elevating progesterone concentrations from Day 3 to 7 post conception significantly increased the length of the conceptus on Day 13 and Day 16, despite having no phenotypic effect on development at pre-hatching (Day 5 and Day 7) stages [21].

The objective of this study was to examine the effect of timing of progesterone supplementation relative to key stages of embryo development (morula, blastocyst and elongation) within the first week of pregnancy on embryo survival rate in cattle. The relationship between progesterone concentration, body condition score, degree of lameness and embryo survival rate was also analysed.
2. Materials and Methods

2.1. Animals and treatments

Cross-bred beef heifers (n=197), approximately 2 years old and weighing 487 ± 3.4 kg were used. All animals were housed in a slatted floor facility in a commercial feedlot under the same management and had ad-lib access to a total mixed ration designed to achieve an average liveweight gain of 1.26 kg/heifer/day. The body condition and locomotion score of all heifers were recorded at one time point during the trial. Locomotion was scored as described by Spreacher et al [22], where a score of 1 represented normal locomotion, 2 represented mild lameness, 3 moderate lameness, 4 lame and 5 severely lame. Heifers that showed an arched back while both standing and walking and were either non or partially weight bearing on the affected foot (locomotion score 4 and 5) were excluded from the experiment. A total of 12 animals were assigned a locomotion score 4 and 5, representing 6% of the total number of animals. Body condition was scored on a 5-point scale with increments of 0.25 as described by Edmonson et al [23].

All heifers were synchronized using two injections of a prostaglandin F$_{2\alpha}$ analogue (PG, Prosolvin, Intervet Ireland Ltd.) given 9 days apart. Heifers were checked for signs of estrus 4 times daily commencing 36 h after the second PG injection and only those recorded in standing estrus (=Day 0, n=197) within the period of 36-60 hours after the 2nd PG injection were inseminated 12-18 h later with frozen-thawed semen from a single ejaculate of a fertile bull. Over all, 59% of synchronized animals were observed in standing estrus in this time period.

Following insemination, heifers were randomly assigned to 1 of 3 treatment groups: (1) control, n=69, (2) supplemented with progesterone by means of a Controlled Internal Drug Release device (CIDR 1.9g, Pfizer UK) from Day 3 until Day 6.5 (n=64), corresponding to the transition of the embryo from the 8-cell to the morula stage of development, or (3) supplemented with progesterone from Day 4.5 until Day 8 (n=64) corresponding to the transition of the embryo from the 16-cell to the
blastocyst stage of development. A time line for the treatments and blood sampling is shown in Figure 1.

2.2. Blood sampling

All heifers were blood sampled via jugular venepuncture for subsequent measurement of progesterone at the time of CIDR insertion, 1 day post insertion and at CIDR removal. Control animals were sampled at the same times as those in both treatment groups (n=6 samples per animal). Blood samples were stored at ambient temperature for a maximum of 6 h and at 4°C for a further 18 h. Serum was decanted after centrifugation for 20 min at 1600 x g and stored at -20°C until subsequent analysis.

2.3. Early embryo survival rate

Embryo survival was determined at slaughter on Day 25. Following slaughter, the uterus was opened longitudinally and, where present, the conceptus was removed. Conceptus length was measured with an optical calipers and the weight was recorded. The CL of all pregnant animals were dissected out of the ovaries and weighed to establish whether or not there was an effect of progesterone supplementation on the development of the CL of pregnancy.

2.4. Progesterone assay

Progesterone was measured using a validated time-resolved fluorescent immunoassay (FIA) with an AutoDELFIA™ Progesterone kit (Perkin Elmer, Wallac Oy, Turku, Finland). The inter-assay coefficients of variation (CV) were 6.5%, 6.5% and 8.6% for low, medium and high quality controls serum pools, respectively. The intra-assay CV were 6.4%, 6.4% and 8.5% for high, medium and low quality control sera, respectively. The sensitivity of the assay was 0.01 ng/ml.

2.5. Statistical analyses

Embryo survival rate was compared using Fishers Exact Chi-square analysis as described by Snedecor and Cochran [24]. Changes in serum progesterone
concentration between pre- and post-insertion (or control equivalent) samples were calculated for individual animals in order to evaluate the incremental increase in progesterone. This increment and the concentrations of progesterone on individual days were compared by ANOVA with Bonferroni and one way t-test using SPSS for Windows. The relationships between body condition score, locomotion score and embryo survival rates were analysed using logistic regression with terms for both linear and quadratic effects included in the model where appropriate.

Within treatments the relationship between concentrations of progesterone on Day 3.0, 6.5 and changes between Days 3 and 6.5 (Treatment 1), Days 4.5, 8.0 and changes between Days 4.5 and 8 (Treatment 2) and embryo survival rate were evaluated using logistic regression in SAS 9. Progesterone was initially included as a linear term in the model. If significant the model was re-run with progesterone included as a quadratic term. Terms used were treatment, breed, progesterone concentration and pregnancy status.

3. Results

3.1. Progesterone

Treatment with a CIDR on either Day 3 or 4.5 increased the mean progesterone concentration by 1.12 ng/ml (P<0.01) on the day after CIDR insertion (Day 4 and 5.5, respectively) compared with control animals (Table 1). The incremental increase in progesterone from pre- to post-insertion (or equivalent) samples was higher in CIDR-treated heifers versus controls (P<0.01).

Within treatment 1, there was a significant relationship (P<0.04) between the change in serum progesterone concentration from Day 3 to 6.5 and early embryo survival rate. A similar relationship (P<0.06) existed from Day 4.5 to 8 within treatment 2. The relationship between serum progesterone concentration on Day 6.5 and early embryo survival rate in treated heifers was also significant (P<0.05), while there was no significant relationship between progesterone concentrations and early embryo survival rate on all other days (P>0.4). The relationship between change in serum concentration of progesterone and embryo survival rate is shown in Figure 2.
3.2. Embryo survival and conceptus size

The mean embryo survival rate over all animals was 41.6% and was not affected by treatment. Similarly, the weight and length of the conceptus was not affected by treatment (Table 2).

3.3. Corpus luteum weight

Treatment with exogenous progesterone did not affect the weight of luteal tissue compared with control animals, irrespective of timing of treatment (Table 2). Weight of the CL was greater (P=0.053) in heifers receiving progesterone between Days 3 and 6.5 compared with those supplemented between Days 4.5 and 8 (Table 2).

3.4. Body Condition Score and Locomotion Score

Of the inseminated heifers, 51% showed normal stance and locomotion and therefore had a locomotion score of 1. Forty-four percent of the heifers had a locomotion score 2; these animals showed an arched back while walking, while their gait remained normal. These animals were regarded as mildly or sub-clinically lame. The remaining 5% scored 3 on locomotion, displaying an arched back when both standing and walking. These animals had a gait that was shorter in stride and were regarded as moderately lame. There was a negative association (P<0.02) between locomotion score and embryo survival rate, which was independent of serum concentrations of progesterone with higher early embryo survival rates in heifers that had low locomotion scores (Figure 3).

Body condition scores of all 197 heifers are presented in Table 3. Of all heifers, 56% had a body condition score of 4 or higher. There was a significant relationship (R²=0.02, P<0.05) between BCS and progesterone levels on Day 3, but there was no relationship between BCS and embryo survival.
4. Discussion

With the estimated percentage of early embryonic death in dairy cows increasing from 28% in 1980 to 43% in 2006 [2], a large body of current research is aimed at understanding the factors determining embryo survival in the early stages of pregnancy. The available evidence indicates that increased progesterone concentrations during the early stages of pregnancy are associated with an advancement of embryonic development leading to an increased level of interferon-τ production and an associated increase in pregnancy rate. Many studies on progesterone supplementation, in an attempt to improve pregnancy rate have been carried out, with variable results. Part of the variation in results is probably related to the fact that only certain animals, those with naturally low circulating progesterone, may exhibit beneficial effects following exogenous progesterone supplementation. In addition, the precise physiological stage of embryo development which most benefits from high progesterone is not known. Recent studies have demonstrated that cows with low progesterone at a time when the embryo is at the morula to blastocyst stage have smaller embryos that produce less interferon-τ on Days 13-16 [5]. It has also been shown that there is an increased probability of embryo survival in heifers and cows that have elevated progesterone on Days 4-6 [2]. Thus, it appears that a critical window for progesterone supplementation is between the morula and hatching blastocyst stages. The present study was conducted to elevate progesterone during these critical stages of embryo development. Beef heifers were used to exclude the confounding factors that can occur in dairy cows, such as lactation, negative energy balance and uterine contamination. The results show that elevation of progesterone does not increase the number of animals that become pregnant, but it was seen that there was a significant relationship between the change in serum progesterone concentrations from Day 3 to 6.5 and embryo survival.

There was no beneficial effect of progesterone supplementation on conceptus survival and size on Day 25. Using a similar supplementation protocol, Carter et al. [21] reported a difference in embryo size on Day 16 when progesterone was supplemented
from Day 3 onwards. These two observations would support the idea that larger
embryos on Days 13-15 may be more effective in suppressing the luteolytic
mechanism around the initiation of elongation due to an advancement in embryonic
growth, rather than the production of larger embryos per se and that by Day 25 this
difference has disappeared. This suggests that progesterone may be only crucial for
advancing embryonic growth to ensure a strong anti-luteolytic signal (maternal
recognition), but once maternal recognition has occurred developmental rate is less
relevant. The possibility that CIDR insertion somehow negatively affected the
development of the endogenous CL, as has been reported in previous studies [19] is
unlikely as the CL weight of control and supplemented groups did not differ in the
present study.

There was a relatively high incidence of subclinical lameness amongst the animals
used in this study, most likely due to the high energy diet that they were fed [25]. The
fact that the animals were housed on slats contributed to their possible discomfort
[26]. The amount of subclinical lameness was already evident in the low incidence of
estrus expression (59%) in the heifers (within 36-60 h after the second PGF2α
injection) used in the experiment. Estrus expression on concrete and slats is
compromised by the ground surface, since cattle do not like to mount on this surface
[27,28]. Early embryo survival across all animals was low, irrespective of treatment
and the lameness could have had a negative impact on the potential positive effects of
treatments. A normal pregnancy rate in heifers would be expected to be up to 60-65%
[2,29]. It is known that lameness has a negative effect on fertility and various authors
have shown that dairy cows that are lame have decreased fertility, starting with lower
estrus expression [27,30,31] and lower conception rate with more inseminations per
conception [32]. To our knowledge this is the first study that has related locomotion
score with early embryo survival rate; there was a clear negative relationship between
the two parameters. This supports the hypothesis that lameness has an adverse effect
on fertility. Indeed, Walker et al. [31,33] demonstrated that lameness decreased the
intensity of estrus expression and progesterone concentrations in dairy cattle.
Lameness can be seen as a chronic stressor as it is a painful long term condition.
Chronic stress and pain has often been associated with disturbances of the
hypothalamic-pituitary-adrenal axis, which can lead to disrupted patterns of pulsatile
GnRH release as well as deprivation of the ovarian structures of gonadotrophin
support resulting in low progesterone concentrations [33]. When cows are lame, the
chance that they ovulate is reduced compared with that of normal cows. Dobson [34]
demonstrated that cows that did not ovulate did not have an LH surge and that their
LH pulse frequency was lower than that of the ovulating cows. Therefore reduced
embryo survival may be negatively associated with lameness in cattle.

The heifers used for this experiment were finishing heifers which was apparent from
their high body condition scores. These animals were fed to gain an average of 1.26
kg/day and to finish quickly at the appropriate slaughter weight. Animals with an
excess in body condition have been known to have problems breeding [35]. In the
abattoir it was evident that all animals that were not pregnant were cyclic. It is a
possibility that heifers gaining weight at a high rate on an energy dense diet may have
higher steroid metabolism, similar to that seen in dairy cows [36], influencing the
possible positive effects of the supplementation of progesterone.

In conclusion, progesterone supplementation in heifers fed a high energy diet did not
improve early embryo survival but resulted in a positive association between change
in progesterone concentration during the early luteal phase and embryo survival rate.
However, moderate lameness had a negative effect on embryo survival on Day 25.
5. Acknowledgements

This work was supported by Science Foundation Ireland (Grant no.: ) and the first author received further support from the Research Development Fund from the School of Agriculture, Food Science and Veterinary Medicine, UCD, Dublin. The authors thank Pat Duffy and Mary Duane for their technical assistance, Penny Furney for help with progesterone assays and IAD Castlebellingham for access to the experimental animals.

6. References


Table 1: Mean (±s.e) serum progesterone concentration (ng/ml) in all heifers treated with a CIDR versus all control heifers. Jugular blood samples were of all heifers taken on day of insertion and one day post insertion.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>N</th>
<th>Pre-insertion</th>
<th>Post-insertion</th>
<th>Increment*</th>
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<tbody>
<tr>
<td>Control (D3-6.5)</td>
<td>69</td>
<td>0.184 ± 0.020(^a)</td>
<td>0.398 ± 0.042(^a)</td>
<td>0.221 ± 0.047(^a)</td>
</tr>
<tr>
<td>CIDR (D3-6.5)</td>
<td>64</td>
<td>0.189 ± 0.016(^a)</td>
<td>1.235 ± 0.071(^b)</td>
<td>1.073 ± 0.060(^b)</td>
</tr>
<tr>
<td>Control (D4.5-8)</td>
<td>69</td>
<td>0.469 ± 0.048(^a)</td>
<td>0.763 ± 0.095(^a)</td>
<td>0.294 ± 0.049(^a)</td>
</tr>
<tr>
<td>CIDR (D4.5-8)</td>
<td>64</td>
<td>0.420 ± 0.032(^a)</td>
<td>1.563 ± 0.086(^b)</td>
<td>1.166 ± 0.059(^b)</td>
</tr>
</tbody>
</table>

*Increment from pre- to post-treatment samples was calculated for individual animals to generate means and standard errors.

\(^a,b\)Within day of assay, means without common superscripts are different (P < 0.01).
Table 2: Embryo survival, conceptus length and weight and CL weight (±s.e) 25 days post conception in pregnant animals supplemented with progesterone from Day 3-6.5 or from Day 4.5-8 as well as the unsupplemented control animals.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>No. of Animals</th>
<th>No. from which embryos were recovered (%)</th>
<th>CL weight, g</th>
<th>Conceptus weight, g</th>
</tr>
</thead>
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<tr>
<td>Control</td>
<td>69</td>
<td>29 (42.0)</td>
<td>5.84 ± 0.19</td>
<td>0.49 ± 0.06</td>
</tr>
<tr>
<td>P4 Day 3 to 6.5</td>
<td>64</td>
<td>22 (34.4)</td>
<td>6.03 ± 0.15</td>
<td>0.55 ± 0.06</td>
</tr>
<tr>
<td>P4 Day 4.5 to 8</td>
<td>64</td>
<td>31 (48.4)</td>
<td>5.42 ± 0.16</td>
<td>0.55 ± 0.05</td>
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Table 3: Distribution of body condition scores amongst inseminated heifers (n=197).

<table>
<thead>
<tr>
<th>Body condition score</th>
<th>3.25</th>
<th>3.5</th>
<th>3.75</th>
<th>4</th>
<th>4.25</th>
<th>4.5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of animals</td>
<td>5</td>
<td>20</td>
<td>61</td>
<td>78</td>
<td>27</td>
<td>6</td>
</tr>
<tr>
<td>Number of pregnant animals</td>
<td>3</td>
<td>8</td>
<td>20</td>
<td>35</td>
<td>14</td>
<td>3</td>
</tr>
<tr>
<td>Percentage pregnant</td>
<td>60</td>
<td>40</td>
<td>33</td>
<td>45</td>
<td>52</td>
<td>50</td>
</tr>
</tbody>
</table>

Body condition was scored on a 5 point scale with steps of 0.25 as described by Edmonson et al [23]