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1 Luteal activity following follicular drainage of subordinate follicles for twin pregnancy
2 prevention in bi-ovular dairy cows

3

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16

17 Short title: Follicular drainage in bi-ovular cows

18

19 Abstract

20

21 Twin pregnancy is undesirable in dairy cattle. This study examines luteal activity
22 following ultrasound-guided puncture and drainage of the smaller pre-ovulatory follicle
23 at timed AI in cows with a pre-ovulatory follicle in each ovary. Luteal activity was
24 determined through Doppler ultrasonography and plasma progesterone (P4)
25 concentrations. The effects of GnRH treatment on Day 7 post-AI on subsequent luteal

26 activity were also assessed. Two study groups were established: a control group of 29
27 cows and a follicular drainage (FD) group of 28 cows. After drainage, all cows
28 developed a corpus luteum (CL) in the drained ovary. On Day 21 post-AI, drainage
29 induced CL and fellow CL were similar in terms of size and vascularization. According
30 to a GLM repeated measures analysis of variance ($P < 0.001$), non-treated drained cows
31 had lower P4 concentrations on Day 21 post-AI than non-treated non-drained cows,
32 whereas GnRH treated cows, both drained and non-drained, showed the highest P4
33 concentrations at this time point. Twin pregnancy was recorded in 3 of the 8 pregnant
34 control cows, whereas no twins were observed in the FD group. Our results indicate that
35 luteal structures following follicular drainage were functional. As for the presence of an
36 additional CL, this could suggest a reduced risk of pregnancy loss. In addition, luteal
37 activity was significantly increased following GnRH treatment on Day 7 post-AI in
38 drained cows.

39 Keywords: Double ovulation, Follicular co-dominance, Color Doppler ultrasonography,
40 Dephereline. Post-AI GnRH

41

42 Introduction

43

44 Although infrequent in monotocous mammalian species, natural multiple births involve
45 in most cases the simultaneous formation of two or more ovulatory follicles on either
46 the same ovary or on both ovaries. The birth of twins was probably a basis for the early
47 development of concepts of human fertility (López-Gatius and Hunter, 2018a) and is
48 very welcome for the economy of beef cattle breeding (Echternkamp et al., 2004;
49 Hashiyada, 2017). However, twin pregnancy is not desirable in dairy cattle and its
50 economic burden in the range of \$97 to \$225 is dependent on the type of twin

51 pregnancy (unilateral vs. bilateral), parity, and the days in milk when the twin
52 pregnancy occurs (Mur-Novales et al., 2017). In fact, twin pregnancy is the main non-
53 infectious factor that compromises pregnancy maintenance during the first trimester of
54 gestation and is logically related to subsequent twinning (López-Gatius and García-
55 Ispuerto, 2010; López-Gatius et al., 2017). While spontaneous embryo reduction rates of
56 11.2% to 28.4% have been reported occurring at around 28–40 days of gestation
57 (López-Gatius and Hunter, 2005; Silva-del-Río et al., 2009; López-Gatius et al., 2010),
58 twin pregnancy losses can exceed 50%, especially during the warm period of the year in
59 some countries (López-Gatius et al., 2004; Andreu-Vázquez et al., 2011). Apart from
60 pregnancy loss, the reproductive performance and productive lifespan of a cow
61 delivering twins are greatly reduced (Beerepoot et al., 1992; Bicalho et al., 2007;
62 Andreu-Vázquez et al., 2012b). Twin pregnancy rates can exceed 18% in some herds
63 (Andreu-Vázquez et al., 2012a), and two strategies have been tested to mitigate their
64 negative effects: hormone therapy or artificial embryo reduction (López-Gatius et al.,
65 2017; López-Gatius and Hunter, 2017b). In effect, GnRH treatment at the time of
66 pregnancy diagnosis increases pregnancy survival and is accompanied by an increase in
67 the twin reduction rate (García-Ispuerto and López-Gatius, 2018).

68

69 Other proposed approaches to reduce twinning in cows on Days 28–41 of gestation
70 consist of manual rupture of the amniotic vesicle and transvaginal ultrasound-guided
71 aspiration of allanto-amniotic fluid or intra-luteal instillation of PGF2 α in one of the
72 two corpora lutea (López-Gatius, 2005; Andreu-Vázquez et al., 2012c; López-Gatius
73 and Hunter, 2016). However, although inducing twin reduction avoids the negative
74 effects of twinning, these interventions may increase the risk of pregnancy loss.

75

76 To eliminate the risk of twin pregnancy without reducing fertility, the puncture and
77 drainage of subordinate follicles at the time of insemination has recently proved to be an
78 efficient procedure (López-Gatius and Hunter, 2018b). This technique overcomes the
79 risk of pregnancy loss related to the process of induced twin reduction and increases the
80 incidence of additional corpora lutea. In this latter study, all drained follicles developed
81 as a corpus luteum 7 days later (López-Gatius and Hunter, 2018b). Draining with no
82 suction supposedly spares a sufficient number of granulosa cells in the follicle which
83 could subsequently give rise to luteal tissue. If these additional luteal structures are
84 functional they may reduce the subsequent risk of pregnancy loss. In effect, a number of
85 corpora lutea exceeding the number of embryos has emerged as a strong factor
86 promoting maintenance of a pregnancy (López-Gatius, 2012).

87

88 The objective of the present study was to assess luteal activity based on Doppler
89 ultrasonography (Acosta et al., 2003; Matsui and Miyamoto, 2009) and plasma
90 progesterone (P4) concentrations following the puncture and drainage of subordinate
91 follicles at timed AI in bi-ovular cows. A second goal was to evaluate the effect of
92 GnRH treatment on Day 7 post-AI on subsequent luteal activity and fertility.

93

94 Materials and Methods

95

96 Experimental animals

97

98 All procedures were approved by the Ethics Committee on Animal Experimentation of
99 the University of Lleida (license number CEEA.06-01/12). The study population was a
100 commercial dairy herd of Holstein-Friesian lactating dairy cows in northeastern Spain.

101 During the study period (November 2017 to April 2018), the mean number of lactating
102 cows in the herd was 240, and mean annual milk production was 10,190 kg per cow. In
103 our geographical area, a clear negative influence of heat stress from May to September
104 on the reproductive performance of lactating dairy cows has been extensively described
105 (López-Gatius, 2003; García-Ispuerto et al., 2007). In effect, ovulation failure increases
106 dramatically under heat stress conditions (López-Gatius et al., 2005; López-Gatius and
107 Hunter, 2017a). Thus, to reduce the number of preovulatory follicles failing to ovulate,
108 this study was performed during the cool period of the year (November to April). Cows
109 were fed complete rations and milked twice daily. Only healthy cows free of detectable
110 reproductive disorders and free of clinical diseases during the study period (days -5 to
111 +28 from insemination) were included. Exclusion criteria were the following: mastitis,
112 lameness, digestive disorders and pathological abnormalities of the reproductive tract
113 detectable by ultrasonography. Cows were selected from groups synchronized for fixed-
114 time insemination (García-Ispuerto and López-Gatius, 2014). Cows were treated with a
115 controlled internal drug release insert (CIDR containing 1.38 g of P4; Zoetis, New
116 York, NY, USA) plus a GnRH agonist (dephereleline: 100 µg gonadorelin acetate [6-
117 DPhe] i.m; Gonavet Veyx, Ecuphar, Barcelona, Spain) upon CIDR insertion. The CIDR
118 was left in place for 5 d, and these animals were also given cloprostenol (500 µg i.m.;
119 PGF Veyx Forte, Ecuphar, Barcelona, Spain) on CIDR removal. Twenty-four h and 36
120 h later, the cows received a second cloprostenol dose and a second GnRH dose,
121 respectively, and were inseminated 50–56 h after CIDR removal. The GnRH agonist
122 used in the present study was selected for its high efficiency in improving luteal
123 function (García-Ispuerto et al., 2019). Cows with a 2.5–3.5 body condition score on a
124 scale of 1 to 5 (Edmondson et al., 1989) and producing >30 kg milk per day were
125 selected for follicular drainage at the time of insemination. A combination of

126 ultrasonography and manual rectal palpation was used to confirm a cow was in estrus
127 and ready for service (López-Gatius and Camón-Urgel, 1988; López-Gatius and Hunter,
128 2017a). Only cows with at least two follicular structures equal to or larger than 12mm in
129 diameter in the absence of a corpus luteum, and with the two largest follicles located
130 one on each ovary, were included in the study. Since double ovulation has been related
131 to the least possible size differences between the larger and smaller follicle irrespective
132 of the individual diameter of each follicle (López-Gatius et al., 2018), only cows with a
133 size difference under 2mm between the two co-dominant follicles were included in the
134 study.

135

136 Follicular draining and experimental design

137

138 Cows were assigned in chronological order of estrus synchronization to a control (n=30)
139 or follicular drainage (FD) (n=30) group (Fig. 1). Follicular puncture and drainage of
140 the subordinate follicle in the FD group was performed by ovum pick-up procedures as
141 previously described (López-Gatius and Hunter, 2018b). Briefly, a portable B-mode
142 ultrasound scanner (E.I. Medical IBEX LITE; E.I. Medical Imaging, Loveland CO,
143 USA) equipped with a convex 5–10 MHz (E.I. Medical IBEX MC8.0 10-6
144 Microconvex; E.I. Medical Imaging, Loveland CO, USA) transvaginal transducer was
145 used for draining through a sterile 19G 25-mm long needle located in the tip of the
146 transducer's metal guide. The vulva and the perineal region of the cow and the metal
147 guide were washed in an iodine solution and the transducer probe was coated with a
148 sterile preservative. Then, the metal guide containing the needle was introduced into the
149 dorsal vaginal fornix, which was to the left or right of the cervix depending on the side
150 of the subordinate follicles. Next, the ovary was positioned transrectally against the tip

151 of the transducer probe so that the follicle was separated only by the vaginal wall. The
152 vaginal wall was then pierced in a cranial direction through the fornix with the needle
153 and introduced into the follicular antrum (López-Gatius and Hunter, 2018b). Follicular
154 contents flowed rapidly through the metal guide with no suction. In the FD group, the
155 largest follicle was considered the dominant follicle and was not drained. Cows showed
156 no signs of discomfort during intra-follicular puncture. Cows were inseminated
157 immediately after follicular drainage with thawed semen from a single ejaculate. In 14
158 drained and 14 nondrained cows, depherenline treatment was randomly given on Day 7
159 post-AI. Ovulation, determined in the FD group by the presence of a corpus luteum in
160 the dominant follicle, and in the control group by the presence of one or two corpora
161 lutea, was assessed 7 days post-AI. Corpus luteum size was recorded in ovulating
162 ovaries and the luteal structure as a corpus luteum in drained ovaries 7 days post-AI
163 taken as the mean of two measurements approximating the greatest length and width. In
164 the case of cavity CL, the mean value of the luteal wall was also recorded. Pregnancy
165 diagnosis was performed by ultrasound at 28 days post-AI (Fig. 1). All gynecological
166 examinations and inseminations were performed by the same operator. Luteal activity
167 was evaluated through plasma P4 concentrations determined on Days 7 and 21 post-AI
168 and through Doppler-ultrasonography on Day 21 post-AI.

169

170 Luteal activity measurements

171

172 Blood samples were collected on Days 7 and 21 post-AI from the coccygeal vein into
173 two heparinized vacuum tubes (BD Vacutainer™; Becton-Dickenson and Company,
174 Plymouth, UK), centrifuged within 20 min (10 min, 1600 g) and the plasma stored at
175 -20°C until analysis.

176 A commercial enzyme-linked immunosorbent assay (ELISA) kit was used to determine
177 plasma progesterone concentrations (Ridgeway Scientific, Alvington, Gloucestershire,
178 UK). The sensitivity of the assay was 0.15 ng / ml. Samples were tested in duplicate,
179 and all samples were analyzed in a single assay (intra-assay coefficient of variation,
180 6%).

181

182 Both ovaries were examined by color Doppler ultrasonography on Day 21 post-AI
183 (Zonare Medical Systems Inc., USA equipped with a 7.5 MHz transducer) and surfaces
184 of corpora lutea scanned and digitalized in a video. Images of cross-sections of each CL
185 were also recorded. The diameter (mean of two measurements) and area were obtained
186 in the ultrasound image that visually represented the largest CL. In the case of CLs with
187 a cavity, the latter was measured and subtracted to obtain the area of luteal tissue. Three
188 cross-sectional images covering the largest areas of the respective CL were finally
189 recorded, and the most vascularized picture selected for each corpus luteum. All corpora
190 lutea were digitalized in B-mode, power and color Doppler mode. All images were
191 digitized in DICOM format. Fixed pre-installed Doppler system controls were used to
192 exclude variations in recordings. The color Doppler images were used to analyze
193 vascular areas in the luteal structures. The vascular area was calculated from each
194 selected CL picture using Image J® software (National Institutes of Health, Bethesda,
195 MD, USA). The percentage of vascularized CL ($\text{color Doppler area}/\text{total CL area} \times 100$)
196 was then determined for each evaluation. All ultrasound analyses were performed by the
197 same operator.

198

199 Data collection and statistical analyses

200 The following data were recorded in each animal: parturition and treatment dates; parity
201 (primiparous versus multiparous); milk production at AI (low producers <40 kg versus
202 high producers \geq 40 kg); days in milk at AI (DIM; <90 days postpartum versus \geq 90 days
203 postpartum); follicular size at AI (diameter of the follicles \geq 12 mm); follicular drainage
204 (control vs FD); treatment on Day 7 post-AI (non-treated vs GnRH group); CL size and
205 number of CL at 7 and 21 days post-AI; CL vascularization on Day 21 post-AI (four
206 classes: no vascularization; < 25%; between 25 and 50%; >50%); ovary in which
207 follicular or luteal structures were recorded (right versus left ovary); plasma
208 progesterone concentration on Days 7 and 21 post-AI; and conception rate after FTAI.
209 Conception rate was defined as the percentage of cows that became pregnant at FTAI
210 out of the total number of cows in the corresponding group.

211

212 The effects of the above-mentioned variables and those of plausible interactions on
213 plasma progesterone concentrations on Days 7 and 21 post-AI were assessed by general
214 linear model (GLM) repeated measures analysis of variance using the SPSS computer
215 package, version 11.5 (SPSS Inc., Chicago, IL, USA).

216

217 Differences in CL size on Day 21 post-AI in the control and FD groups and GnRH-
218 treated and non-treated cows were analyzed by the Student's test. Possible significant
219 effects of drainage, treatment and drainage-treatment interaction on the conception rate
220 were explored by Tukey-Kramer multiple comparison tests. Significance was set at $P <$
221 0.05. Values are expressed as the mean \pm the standard deviation (SD).

222

223 3. Results

224

225 Three cows with three corpora lutea at 7 days post-AI were withdrawn from the study.
226 The final study population was comprised of 57 cows: 29 in the control group and 28 in
227 the FD group. Ovulation failure (absence of a CL 7 days after AI) was registered in 2
228 control cows. The data collected in these two animals were eliminated from the P4
229 analyses but maintained for the conception rate analysis. Table 1 shows luteal dynamics
230 on Days 7 and 21 post-AI following drainage and GnRH treatment. Double ovulation
231 was recorded in 12 cows (41.4%) in the control group. After follicular drainage, all
232 cows showed a CL in the drained ovary, whereas the corresponding dominant follicle
233 failed to ovulate in 6 (21.4%) of the 28 FD cows. No pregnancies were produced in
234 these latter 6 cows. On Day 21 post-AI, luteal structures were not detected in 12 cows: 7
235 in the control group and 5 in the FD group. No pregnancies were recorded in these 12
236 cows and the P4 data obtained in these animals were removed from the analyses. Of the
237 28 cows receiving GnRH on Day 7 post-AI, 6 had a GnRH-induced CL on Day 21 post-
238 AI: 3 in the control and 3 in the FD group (Table 1). The GnRH induced CL was the
239 third CL in the FD group. Mean milk production at the time of treatment, DIM at AI
240 and number of lactations, were 44.5 ± 9.5 kg, 123.0 ± 70.1 days, 2.7 ± 1.6 lactations,
241 respectively (mean \pm SD). In the GLM repeated measures analysis of variance, parity,
242 milk production and days in milk at AI were found to have no effects on luteal patterns.
243 Luteal activity was assessed only in cows with at least one CL on Day 7 (n=55) or Day
244 21 (n=45) post-AI. The mean size of the CL on Day 7 post-AI resulting from ovulation
245 (16.2 ± 4 mm, ranging from 8 to 24 mm) was larger ($P < 0.03$, Student's test) than that
246 resulting from drained follicles (8.2 ± 3.5 mm, ranging from 5 to 15 mm). In 22 FD
247 cows that had both ovulating and drainage CL on Day 21 post-AI, no significant
248 differences in size and vascularization between both types of CL could be detected.
249 Similar CL vascularization patterns were also observed in the remaining cows with two

250 CL. The mean size of GnRH induced CL on Day 21 post-AI (11.1 ± 2.5 mm, ranging
251 from 6 to 20 mm) was significantly smaller ($P < 0.0001$, Student's test) than that of the
252 remaining CL (18.7 ± 6.2 mm, ranging from 8 to 36 mm). Vascularization of the
253 GnRH-induced CL was similar to that of CL partners.

254

255 Mean plasma P4 concentrations increased significantly ($P < 0.0001$, Student's test) from
256 Day 7 to 21 post-AI (2.3 ± 1.6 and 3.4 ± 3.1 ng/mL, respectively). General linear model
257 repeated measures of variance revealed that the treatment x drainage interaction had a
258 significant effect (between subject effect repeated measures ANOVA; $P < 0.001$) on
259 plasma P4 concentrations on Day 21 post-AI. Nontreated drained cows had lower P4
260 concentrations on Day 21 post-AI than those their non-treated non-drained partners,
261 whereas GnRHtreated cows, both drained and non-drained, showed the highest P4
262 concentrations on Day 21 post-AI (Fig. 2).

263

264 Over the course of the study, 19 cows (33.3%) of the 57 cows enrolled became
265 pregnant: 8 control and 11 FD cows. The conception rate was significantly higher ($P <$
266 0.05) in drained cows treated with GnRH on Day 7 post-AI than non-treated drained
267 cows. Twin pregnancy was recorded in 3 of the 8 pregnant control cows, whereas twins
268 were not registered in the FD group (Table 1). Corpus luteum vascularization on Day 21
269 post-AI was positively correlated with plasma P4 concentrations and pregnancy rate
270 (Table 2).

271

272 Discussion

273

274 In the present study, we examined bi-ovular lactating dairy cows with a pre-ovulatory
275 follicle in each ovary. Findings confirm our previous data indicating that puncture and
276 drainage of the smaller preovulatory follicle (subordinate) in these cows at timed
277 artificial insemination serves to avoid a risk of twin pregnancy (López-Gatius and
278 Hunter, 2018b). Further, while showing individual variation in size, induced luteal
279 structures present in the drained ovary in all cows subjected to this technique were
280 found to be functional. On Day 21 post-AI, these drainage-induced CL were similar in
281 size and vascularization to their counterparts of ovulatory origin. Moreover, luteal
282 activity was greatly increased following GnRH treatment on Day 7 post-AI.

283

284 In this setting, however, the high risk of ovulation failure of the dominant follicle
285 suggests that the drainage procedure could impair fertility. Accordingly, to improve the
286 chances of selecting double ovulating animals, only cows with a size difference smaller
287 than two mm between the two co-dominant follicles were included in this study (López-
288 Gatius et al., 2018). In addition, the experiment was performed during the cool period of
289 the year, when the risk of ovulation failure clearly decreases (López-Gatius et al., 2005).
290 Six of the 28 (21.4%) drained cows failed to ovulate. We nevertheless anticipate that
291 treatment with GnRH or hCG at the time of insemination will improve the chances of
292 ovulation for the dominant follicle. Further comprehensive studies are needed to
293 confirm this assertion.

294

295 A further critical point was the impaired functional condition of the luteal structures
296 observed on Day 21 post-AI after follicular puncture. Thus, our results suggest some
297 form of limitation with time after ovulation, seemingly associated with a reduced cell
298 proliferation and steroid secretion potential. Lowest mean plasma P4 concentrations on

299 Day 21 post-AI were recorded in the non-treated drained cows. However, in these
300 animals, treatment with GnRH on Day 7 post-AI significantly increased plasma P4
301 concentrations and the pregnancy rate, thus overcoming the possible detrimental effect
302 of follicular draining. In effect, we would expect a reduced risk of early fetal loss due to
303 this significant increase in plasma P4 concentrations in response to GnRH treatment
304 (López-Gatiús, 2012). Sub-optimal levels of circulating P4 have been extensively
305 related to such losses (Ayad et al., 2007; Karen et al., 2014; Gábor et al., 2016).
306 However, although the conception rate was significantly higher in drained cows treated
307 with GnRH on Day 7 post-AI than in non-treated drained cows, this finding on fertility
308 need to be interpreted with caution. A higher sample size in each group should be
309 required to safely draw any conclusion in future studies.

310 To assess the luteal function of drainage-induced CL, we used Doppler ultrasonography.
311 This technique proved valuable especially when combined with plasma P4
312 concentrations, which are useful as an early predictor of pregnancy. Although early
313 pregnancy diagnosis was not an objective of the present study, our results reinforce
314 previous reports on early pregnancy prediction based on luteal activity on Days 14 to 21
315 post-AI in which CL luteum blood flow area was closely associated with plasma P4
316 concentrations and subsequent pregnancy, as extensively reported (Herzog et al., 2011;
317 Siqueira et al., 2013; Kanazawa et al., 2017).

318

319 As an overall conclusion, puncture and drainage of subordinate follicles at the time of
320 insemination eliminates the risk of twin pregnancy and increases the presence of
321 additional functional corpora lutea. GnRH treatment on Day 7 post-AI greatly improves
322 subsequent luteal function in cows undergoing follicular drainage. Results furthermore

323 suggest that GnRH or hCG given at the time of insemination could reduce the ovulation
324 failure rate of the dominant follicle following the puncture and drainage technique.

325

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327

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329

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331

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447 C.A.C., Viana, J.H.M., 2013. Color Doppler flow imaging for the early detection of
448 nonpregnant cattle at 20 days after timed artificial insemination. *J. Dairy Sci.* 96, 6461–
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- 450

451 Table 1. Luteal patterns and conception rate after drainage and treatment (n = 57).

Drainage	GnRH	2 CL D7	0 CL D21	Conception	Twins*
D0	D7	n (%)	n (%)	rate n (%)	n (%)
No	No	5/15 (33.3)	4/15 (26.7)	4/15 (26.7) ^{a,b}	1/4 (25)
	Yes	7/14 (50)	3/14 (21.4)	4/14 (28.6) ^{a,b}	2/4 (50)
Yes	No	10/14 (71.4)	3/14 (21.4)	3/14 (21.4) ^a	0/3 (0)
	Yes	12/14 (85.7)	2/14 (14.3)	8/14 (57.1) ^b	0/8 (0)

452 Values with different superscripts within columns denote significant differences

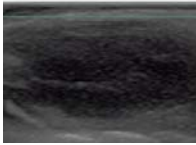
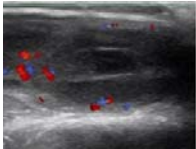
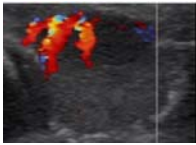
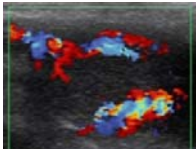
453 detected by the Tukey-Kramer test (P < 0.05).

454 *Values on pregnant cows

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457 Table 2. Sonograms on Day 21 post-AI for different levels of CL vascularization and
 458 their corresponding plasma P4 concentrations and conception rates (n = 45).

Vascularization	%	Mean P4 values \pm SD (ng/ml)	Conception rate n (%)	GnRH treated n (%)
	0	0.22 \pm 0.21 ^a	0/8 (0) ^a	4 (50)
	< 25	1.49 \pm 1.90 ^a	0/9 (0) ^a	4 (44)
	25-50	4.92 \pm 1.20 ^b	5/13 (38.5) ^{a,b}	7 (54)
	> 50	6.30 \pm 3.05 ^b	14/15 (93.3) ^b	9 (60)

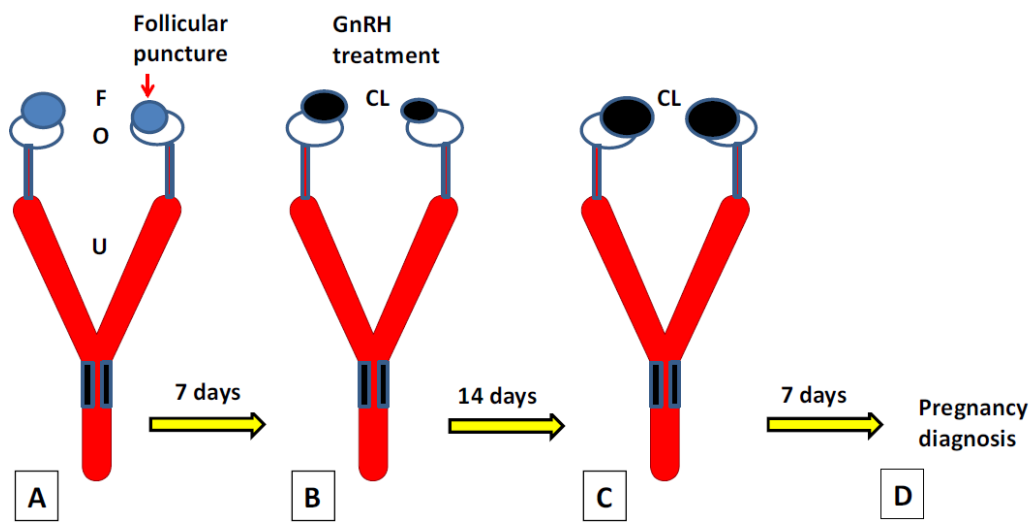
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460 Values with different superscripts within columns denote significant differences
 461 detected by the Tukey-Kramer test (P < 0.001).

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464 Fig. 1. Experimental design on cows with a preovulatory follicle in each ovary.
 465 Puncture and drainage of the smaller pre-ovulatory follicle was performed at timed AI
 466 in the drainage group (A). Ovulation and luteal activity were assessed on Day 7 post-AI.
 467 Luteal activity was determined through corpora lutea measurements and plasma
 468 progesterone determinations. A GnRH dose was administered at this time point in the
 469 treatment group of drained and non-drained cows (B). Luteal activity was determined
 470 through Doppler ultrasonography, corpora lutea measurements and plasma progesterone
 471 determinations on Day 21 post-AI (C). Pregnancy diagnosis was performed on Day 28
 472 post-AI (D). F: follicles; CL: corpora lutea; O: ovaries; U: uterus.

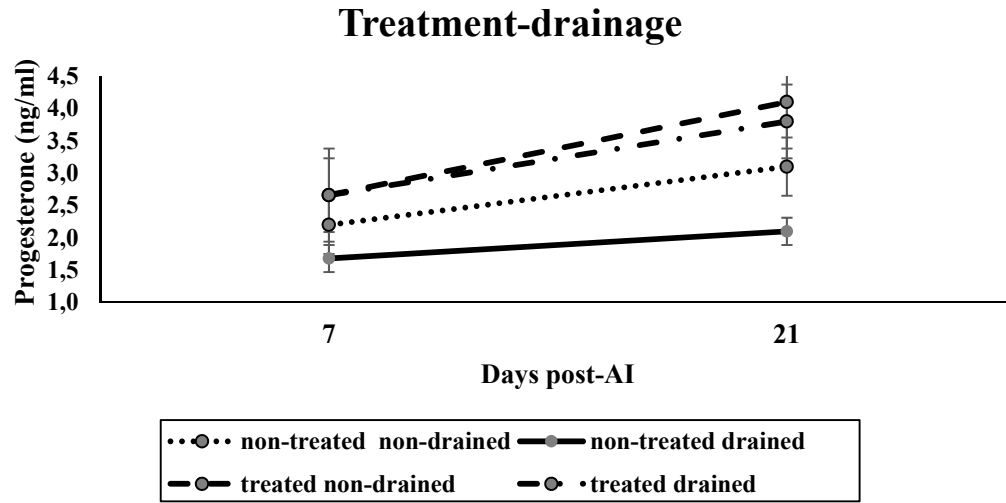


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475 Figure 2. Mean progesterone concentrations (ng/ml) recorded on days 7 and 21 post-AI
 476 after drainage and treatment (n = 45). Values differed significantly on Day 21 post-AI
 477 between non-treated drained and non-treated non-drained cows and between treated and
 478 non-treated cows (between subject effect repeated measures ANOVA; $P < 0.001$).

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