EFFECT OF MONENSIN ON REPRODUCTIVE PERFORMANCE AND MILK PRODUCTION IN LACTATING DAIRY COWS

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ABSTRACT

Thirty-one lactating Friesian dairy cows were used to determine the effect of monensin supplement on blood components, reproductive performance, body condition score (BCS) and milk production. Animals were divided into three groups as follows: Group 1 (n = 9) was considered as control group without monensin supplementation; group 2 (n = 12) was received 300 mg monensin/head/day for 40 days before calving and continued to day 60 postpartum and the cows in group 3 (n = 10) were received the same dose of monensin from calving till day 60 postpartum. Monensin had no significant effect on blood metabolites. BCS at calving was significantly (P < 0.05) increased in group 2 fed monensin compared with the other groups, but, the increase in BCS at conception was significant (P < 0.05) in both groups fed monensin compared with that of control group. Supplementation with monensin in group 2 significantly (P < 0.05) reduced the interval from calving to both first estrus and first AI and to conception as well as the length of the first estrus cycle (19.67 d). A higher pregnancy rate was recorded in group 2 (66.7%) and group 3 (50.0%) compared with that in control group (22.2%). Moreover, monensin significantly increased milk production by about 2.16 kg/d (7.2%) in group 2 but had no significant effect on milk composition. It can be concluded that monensin can be used as a feed supplement for the lactating dairy cows for 40d before calving till day 60 postpartum to improve their reproductive performance and to increase milk production without any marked changes in milk fat and other milk components.

INTRODUCTION

After parturition, energy balance is one of the common factors affecting the interval from parturition to first estrus and conception at the time of breeding in dairy cattle (Butler and Smith, 1989; Canfield and Butler, 1990). During early and peak lactation, dairy cows are unable to consume enough feed to meet the energy balance resulting in a state of negative energy balance (NEB) causing a poor reproductive efficiency (Randel, 1990). In dairy cows experiencing NEB postpartum, there is a deficiency in the release of LH rather than FSH (Beam and Butler, 1999). Thus, resumption of normal episodic LH release necessary for preovulatory follicular development has been proposed as the key event in the return to ovarian cyclicity of cows in NEB (Canfield and Butler, 1990, 1991). Early resumption of ovulatory cycles has also been shown to enhance conception rate to first insemination (Lucy et al., 1992).

Monensin sodium, an ionosphore antibiotic produced by Streptomyces cinnamonensis, selectively modifies the ruminal flora and improves the digestive efficiency in beef and dairy cattle (Schelling, 1984). Monensin supplementation lead to increase ruminal propionate (Hegazy, 1997 and Van Der Werf et al., 1998), and blood glucose concentration (Duffield et al., 1998; Green et al., 1999), and reduced ruminal degradation of dietary protein (Zinn and Borques, 1993). Reduced blood glucose concentration have been associated with reduced reproductive performance (Butler and Smith, 1989). Therefore, monensin may improve reproductive performance by reducing the ruminal degradation of dietary protein and increasing blood glucose. Earlier studies suggest that monensin may have some effects on pituitary and ovarian function (Randel and Rhodes, 1980 and Randel et al., 1982) in beef and dairy heifers and in cows (Orluna and Carlson, 1985) and shorten the interval from calving to the return to first observed estrus in dairy cows (Abe et al., 1994).

In lactating cows treated with monensin, it was reported that production of milk and milk protein may be increased by reduced methanogenesis, increased propionate, and increased blood glucose (Van der Werf et al., 1998; Phipps et al., 2000). However, milk fat has been shown to be depressed (Phipps et al., 2000; Duffield et al., 2003) or unchanged (Lean et al., 1994 and Duffield et al., 1999)

with monensin treatment. The objectives of this study were to investigate whether treatment with monensin improved reproductive performance, BCS and milk production in Friesian dairy cows.

MATERIALS AND METHODS

Animals and management:

Thirty-one multiparous Friesian cows were used in this study. They were selected based on their expected calving date from the herd located at Sakha Experimental Station, Ministry of Agriculture. The trial began in July 2003 and ended in April 2004. All animals were fed ration to meet their maintenance and milk production requirements according to the standard allowances recommended by N.R.C. (1989).

Experimental design:

Cows entered the trial at 50 days before the predicted calving date. Upon entering the trial, cows were paired according to body condition score (BCS) and calving date before being allocated randomly into three treatment groups. The first group $(G_1; n = 9)$ was left without monensin supplementation and considered as control. The second group $(G_2; n = 12)$ was given 300 mg monensin sodium premix per day per cow (Elancoban 200, Elanco Animal Health, division Eli Lilly Export S.A., Geneva, Switzerland) for 40 days before expected calving date and continued to d 60 post-calving. The third group $(G_3, n = 10)$ was given 300 mg monensin/d per cow from the day of calving until d 60 post-calving.

Data collected:

All cows were scored for BCS biweekly by the same operator at the afternoon milking using a scale from 1 to 5 with quarter-point increment (1 = thin to 5 = obese; Edmondson et al., 1989). Calf birth weight was also recorded for each cow. All reproductive events were recorded including the number of days from calving to the first observed estrus, first service and conception as well as number of services per pregnancy.

Pregnancy rate (PR) was defined for treated and control groups as number of cows that were pregnant divided by the total number of cows of each group. First service PR was defined for each group as the number of cows that were pregnant to their first service

divided by the total number of cows for which a first service date were recorded. Cows were machine milked twice daily and milk yield was recorded. Milk samples were collected biweekly from the successive a.m and p.m. milkings and analyzed for the percentage of milk fat, protein, lactose, and SNF.

Blood samples were collected biweekly with anticoagulant and then centrifuged at 1200 x g for 20 min. Plasma was decanted and stored frozen at -20°C until glucose, albumin, cholesterol and triglycerids analysis. All cows were observed for any sign of diseases until the end of treatment.

Statistical analysis:

Data were statistically analysed using (Chi-square test) General Linear Models Procedure adapted by SPSS (1997) for User's Guide.

RESULTS AND DISCUSSION

Results presented in Table (1) indicated that supplementation of dairy cow with monensin had insignificant effect on the plasma concentration of albumin, glucose, cholesterol and triglycerides. These results are consistent with that reported by Sauer et al. (1989) and Hayes et al. (1996). However, other studies have reported an increase in plasma glucose concentration (Abe et al., 1994; Duffield et al., 1998 and Green et al., 1999). In addition, O'Kelly and Spiers (1990) found that ruminal bacteria in steers supplemented with monensin had an increase in lipid content with an associated increase in the circulating level of cholesterol.

Table (1): Changes in some plasma metabolites in lactating dairy cows supplemented with monensin during pre and/or post-calving.

Parameters Treatment PRE (G₂) Control (G₁) POST (G₃) 46.45 ± 3.4 46.2 + 3.946.35 + 3.4Albumin g/L Glucose mmol/L 3.14 ± 0.59 3.21 ± 0.6 3.20 ± 0.6 Cholesterol mmol/L 5.13 ± 1.1 5.30 + 1.25.35 + 1.2Triglycerids (TG) mmol/L 0.11 0.11 0.12

Postpartum reproductive parameters for cows supplemented with monensin are presented in Table 2. The results showed a significant (P < 0.05) reduction in postpartum interval to both first estrus and first AI in cows fed monensin pre-calving (46.5; 58.5 d) compared to that of cows in control group (63.4; 73.0 d). However, the interval in group 3 which started monensin treatment post-calving (54.9; 66.7 d) was intermediate and did not significantly different than that of the other two groups. These results are in agreement with those of Tallam et al. (2003) and Heuer et al. (2001) who reported that the interval from calving to first estrus or ovulation occurred earlier in cows fed monensin than control cows. Additionally, Hegazy (1997) also reported that the interval from calving to first estrus was significantly decreased by 26 days in buffalo cows supplemented with monensin pre-calving than in the control group.

On the contrary, Lean et al. (1994) reported that there was no effect of monensin on the time of first estrus in lactating dairy cows despite the increased of glucose concentration. The present study indicated that, Monensin had only a significant $(P \le 0.05)$ effect on the length of first estrus cycle for cows in G₂ as compared to that in control one (Table 2). This finding is consistent with that of Tallam et al. (2003) who reported that supplementation with monensin resulted in a shorter duration of first estrus cycle than control cows. The shorter postpartum interval to first estrus in cows fed monensin suggests earlier resumption of normal ovarian activity. The reduction of this interval may be due to the metabolic effects of monensin through various metabolic signals which may have a positive influence on hypothalamic-pituitary function resulting in accelerated ovarian recovery postpartum (Randel and Rhodes, 1980; Randel et al., 1982 and Tallam et al., 2003). One of the effect of monensin feeding is the increased propionate: acetate production in the rumen (Hegazy, 1997). Infusion propionate into rumen has shown to enhance the secretion of LH in response to GnRH challenge in beef heifers (Rutter et al., 1983). Therefore, the increases in ruminal propionate by monensin supplementation would have positive effects on LH release. The timing of monensin treatment may also explain the occurrence of earlier first estrus in the cows in group 2 by modifying the pituitary-ovarian axis before calving (Tallam et al., 2003).

Table (2): Postpartum reproductive parameters of cows in response

to monensin feeding.

Parameters	Treatment		
	Control (G _i)	PRE(G ₂)	POST(G ₃)
No. cows	9	12	10
Interval from calving to first estrus, d	63.4 ± 5.6 °	46.5 ± 2.46 b	54.9 ± 4.04 ab
Interval from calving to first Al, d	73.0 ± 6.09 °	58.5 ± 2.1 b	66.7 ± 4.15 ab
Length of first estrus cycle, d	21.33 ± 0.44		20.38 ± 0.46 ab
Interval from calving to conception, d	113.78 ± 5.19 °	83.83 <u>+</u> 8.49 ^b	93.8 ± 8.85 ab
Service/pregnancy (n)	2 ± 0.00	1.5 ± 0.19	1.4 ± 0.24
Pregnancy/1st service %	1/9 (11.11)*	4/12 (33.33) b	3/10 (30.0) b
Pregnancy rate %	2/9 (22.2)	8/12 (66.7)°	5/10 (50.0) b

values within each row carrying different superscripts differ (P < 0.05)

The postpartum interval from calving to conception was significantly (P < 0.05) decreased in cows supplemented with monensin pre-calving (83.83d) than in the control group (113.78 d). This interval in cows given monensin post-calving (93.8 d) was intermediate and did not significantly differ than that in the other two groups. These findings disagree with that previously reported by Duffield et al. (1999) and Heuer et al. (2001) but are close to that of Beckett et al. (1998) in dairy cows and Hegazy (1997) in buffalo cows. A higher pregnancy rates/first service (33.33 and 30%; P < 0.05) were recorded for cows fed monensin (G₂ and G₃) than control cows (11.11%). These results confirm an earlier report that improving energy balance by monensin supplementation increases the conception rate at first service (Senatore et al., 1996). The present finding is also consistent with that of Haves et al. (1996) who reported that monensin might improve reproductive performance of cows if administration was started early enough to remove the negative effect of reduced feed intake. On the contrary, the present result disagrees with that of Duffield et al. (1999) who reported that treatment with monensin had no significant effect on any measure of reproductive performance. Moreover, Beckett et al. (1998) found that monensin supplementation did not alter the pregnancy or first service pregnancy rates. Pregnancy rates (Table 2) were higher in group 2 (66.7%; P < 0.01) and group 3 (50%; P < 0.05) fed monensin as compared with that in control group (22.2%).

The higher percentage of fertility in the cows supplemented with monensin may be due to monensin altered the follicular population, the size of second follicle on either side of ovaries and increased the number

be value within each row carrying different superscripts differ (P < 0.05)

values within each row carrying different superscripts differ (P < 0.01).

of follicular waves than control one (Reed and Whisnant, 2001). On the other hand, the lowest pregnancy rate shown in control group may be explained by the findings of Kendriek et al. (1999) that low energy intake was associated with poor quality of oocytes collected between day 30 and 100 of lactation causing low fertility.

Determination of BCS for cows in the three treatment groups (Table 3, Fig. 1) indicated that the values are similar at the start of treatments, thereafter, feeding monensin was significantly (P < 0.05) increased the BCS to reach 4.45 at calving for the cows supplemented with monensin pre-calving as compared with 4.14 and 4.18 for control cows supplemented with monensin post-calving, respectively. BCS was also significantly (P < 0.05) higher at conception for cows in the second group (3.65) and third group (3.45) which were treated with monensin compared with that of cows in control group (3.22). This finding confirms our suggestion that monensin improve energy utilization (Hegazy, 1997).

Table (3): Effect of monensin supplementation on BCS of the experimental cows and their calf birth weight

Parameters	Treatment			
	Control (G ₁)	PRE(G ₂)	POST(G ₃)	
BCS at:				
Initial	3.81 ± 0.15^{a}	3.85 ± 0.05^{a}	3.80 ± 0.12^{a}	
Calving	4.14 ± 0.1^{8}	4.45 ±0.05 ^b	4.18 ± 0.07^a	
Conception	3.22 ± 0.08^{a}	3.65 ± 0.06^{b}	3.45 ± 0.07^{b}	
Calf birth weight (kg)	31.89 ± 0.7	34.58 ± 1.12	32.00 ± 1.26	

^{a, b} values within each row carrying different superscripts differ (P < 0.05)

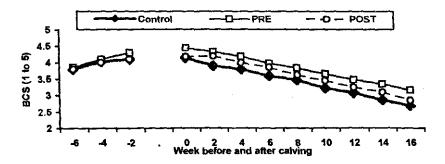


Fig. (1): Changes in BCS of lactating dairy cows during and after the treatment with monensin.

In contrast to the present results, Tallam et al. (2003) reported that monensin did not result in improvement in late prepartum body condition or at calving. However, the present results are in agreement with that of Hegazy (1997). Thereby, BCS at calving and postpartum nutrition influenced the percentage of cows with luteal activity or the occurrence of luteal activity (Vizcarra et al., 1998) and consequently affect the pregnancy rates and postpartum interval to estrus in cows (Richards et al., 1986). In agreement with the results of the previous studies (Duffield e al., 1998 and Wagner et al., 1999) monensin feeding reduced the loss of BCS after calving with increasing weeks of lactation than control cows (Fig. 1). It is previously noticed that the loss of body condition had negative impacts on the success of breeding (Duffield et al., 1999) and strongly related to reproductive performance (Pryce et al., 2001). Therefore, we suggest that BCS scored once biweekly can be used for either management and in a breeding program as an indirect selection criterion for fertility. Monensin supplementation during precalving period (G₂) resulted in a slight increase in birth weight of calves born than that of the other two groups (Table 3). These results were close to that of Hegazy (1997) and Vallimont et al. (2001).

Supplementation with monensin increased milk yield by about 2.16 kg/d (7.2%) in group 2 and 1.26 kg/d (4.49%) in group 3 compared to group 1 (control), but this increase was significant (P < 0.05) only in group 2 (Table 4). In agreement with the current results, monensin increased milk production in lactating dairy cows and magnitude of this increase was 7% (Van Der Werf et al., 1998) and 6.5% (Ruiz et al., 2001). In contrast Tallam et al. (2003) reported that monensin had no effect on milk yield.

Table (4): Milk yield and composition in the first 120 d of lactation for

cows treated with monensin pre or post-calving.

Items	Treatment		
	Control (G ₁)	PRE (G ₂)	POST (G ₃)
No. cows	9	12	10
Milk yield kg/d	13.4 ± 0.51°	15.56 ± 0.58 ^b	14.66 ± 0.53^{ab}
Fat %	3.15 ± 0.46	3.16 ± 0.36	3.13 ± 0.4
Protein %	2.68 ± 0.42	2.61 ± 0.33	2.5 ± 0.35
Lactose %	4.28 ± 0.53	4.32 ± 0.42	4.32 ± 0.46
SNF %	7.54 ± 0.7	4.54 ± 0.56	7.48 ± 0.6
SCC, cells/ml*1000	228.4 ± 3.7	230.5 ± 3.1	229.8 ± 3.3

^P values within each row carrying different superscripts differ (P < 0.05)

The increase in milk yield (Table 4) from cows treated with monensin may be due to that monensin increased the supply of glucogenic precursors resulting from changes in pattern of rumen fermentation (Lean and Wade, 1997). It was reported that the addition of monensin to the diet may compared with an increase in concentrate components of the diet because both lead to an increase in ratio of propionic to acetic and butyric acid resulting in increased milk production (Van Der Werf et al., 1998). The negligible effects of monensin on milk composition are similar to that of a previous studies (Beckett et al., 1998; Vallimont et al., 2001) and disagree with that of others (Phipps et al., 2000 and Duffield et al., 2003).

In conclusion: the present study indicates positive effects of monensin use on dairy cows when monensin is fed for 40d before calving and continued 60 d post-calving. Monensin supplementation improved BCS, reproductive performance, fertility rate and increased milk yield in lactating dairy cows without any changes in milk components.

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تأثير الموننسين على الكفاءة التناسلية وإنتاج اللبن لأبقار اللبن الحلابة ' ابراهيم محمود عبد الرازق _ 'عبد السلام موسى متولى 'عبد الستار عبد العزيز شتا

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الملخص العربي

استخدم في هذه الدراسة ٣١ بقرة فريزيان لدراسة تأثير الموننسين على بعض مكونات الدم ، الكفاءة التناسلية ، درجة امتلاء الجسم (BCS) وإنتاج اللبن حيث تسم تقسيمها الى ثلاث مجموعات كالتالى: المجموعة الأولى (٩ بقسرات) وقد اعتبرت كمجموعة ضابطة غير معاملة. أما المجموعة الثانية (١٢ بقرة) فقد اعطيت ٥٠ مجم موننسين / أس/يوم لمدة ٤٠ يوم قبل الولادة المنتظرة استمرت حتى ٦٠ يوم بعد الولادة ، في حين أن المجموعة الثالثة (١٠ بقرات) قد اعطيت نفس جرعة الموننسين المستخدمة في المجموعة الثالثة وحتى ٦٠ يوم بعد الولادة.

أظهرت النتأنج أنه لم يكن للموننسين أى تأثير معنوى على مكونسات السدم المدروسة ، أدت الى ظهور تحسن معنوي فى السـ BCS المجموعة الثانية عند الولادة مقارنة بالمجاميع الاخرى بينما تحسن السـ BSC عند حدوث الحمل فسـى المجموعة الثانية و الثالثة مقارنة بالمجموعة الضابطة. إعطاء الموننسين أدى إلى انخفاض معنوى فى طول الفترة من الولادة وحتى أول شياع وأول تلقيحه وكذا حتى حدوث الحمسل ، كما أدى أيضا إلى انخفاض معنوى فى طول أول دورة شياع وذلك بالنسبه لأبقار المجموعة الثانية (١٩٠٨) وفى المجموعة الثالثة فى نسبة الحمل المتحصل عليها فى المجموعة الثانية (٢٠١٧) وفى المجموعة الثالثة فى نسبة الحمل المتحصل عليها فى المجموعة الثانية (٢٠١٧) فضلا عن ذلك قبد أدت المعاملة بالموننسين إلى زيادة كمية اللبن المنتجة زيادة معنوية بمقدار ٢٠١٦ كجم/يوم (٢٠٧٧) فى المجموعة الثانية مقارنة بالمجموعة الضابطة وذلك دون أى تغيير فى نسبة دهون اللبن أو مكونات اللبن المخرى. وعليه فإنه يمكن استخدام الموننسين كإضافة غذائيسة غذائيسة لابقار الفريزيان الحلابة خلال فترة الس ، ٤ يوم قبل الولادة وحتى ، ٦ يوم بعد الولادة لتحسين كفائتها التناسلية ومعدل الخصوبة بها وكذلك زيادة إنتاجيتها مسن اللبن دون حدوث أى تغيير فى نسبة دهون اللبن ومكونات اللبن الأخرى.