

## **INHIBIN IN SERUM AND FETAL FLUID OF SHE- CAMEL THROUGHOUT PREGNANCY**

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### **SUMMARY**

Blood samples were collected from 40 pregnant she- camels slaughtered at Cairo abattoir. The animals were divided into three groups according to the stage of pregnancy, eraly stage up to 5 months (15 animals), middle stage from 5-9 months (13 animals) and late stage of pregnancy above 9 months (12 animals). Amniotic fluid was collected from each animal individually. Inhibin was measured in serum and amniotic fluid using indirect method. The results showed that inhibin level increaseed with the advancement of pregnancy in both serum and amniotic fluid. Moreover, the level of inhibin in amniotic fluid was higher than its corresponding serum level during the different stages of pregnancy.

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### **INTRODUCTION**

During the last decades inhibin received great attention where several studies were designed to study the relationship between inhibin and gonadotropin regulation in most species (Dejong, 1979, Franchimont et al., 1980), Franchimont et al., 1980 .Famworth et al., 1988 and Burger, 1993). Also, several studies were performed to determine its site of formation in both sexes (De jong and Sharp, 1976, Tsonis et al., 1983 and Dekretser et al., 1996). Moreover, other studies aimed to clarify its chemical nature (Robertson et al., 1985, Miyamoto et al., 1986 and Knight et al., 1989).

Recently the study of inhibin during pregnancy has some attention especially in humans (France et al., 1996, Lambert et al., 1996, Nambo et al., 1996 and Miller et al. 1997). However, such

studies in domestic animals seem to be scarce specially in she - camels. So this work aimed to give basic information about inhibin in serum and fetal fluids in different stages of pregnancy in the she-camel.

## MATERIAL AND METHODS

Blood samples were collected randomly from 100 labelled she- camel slaughtered at Cairo abattoir. After evisceration blood samples of non pregnant animals were discarded and those of pregnant animals were divided into three groups according the stage of pregnancy, early (15 animals), middle (13 animals) and late stage of pregnancy (12 animals). Amniotic fluid was collected from each animal of the three groups individually using separate, clean and dry syringes.

Stage of pregnancy was determined according to the method of Crown Rump Length (C.R.M) as follow (C.R.M <  $18.44 \pm 1.31$  cm) < 5 months for early pregnancy, (C.R.M <  $18.44 \pm 1.31$  to  $43.87 \pm 0.909$  cm) 5 to < 9 months for middle pregnancy

and (C.R.M,  $43.87 \pm 0.909$  to  $81.75 \pm 0.563$  cm) 9 to < 13 months for late pregnancy (Al-Agawany and Gad, 1991). Sera and fetal fluid were kept at -20c till inhibin assay. Sera and fetal fluid were subjected to steroid extraction according to the method of Welschen et al. (1977). Inhibin potency of the steroid -free samples was estimated by indirect method throughout comparing the percentage of FSH suppression in the ovariectomized rats as a result of injection of these samples with those obtained from the log - dose response curve of different doses of standard porcine inhibin (Ali, 1998). FSH was estimated using direct ELISA technique (Voller et al., 1979). Statistical analysis was performed according to Snedecor (1971).

## RESULTS

Data presented in table (1) showed that the higher level of inhibin potency was found during late stage of pregnancy in serum and amniotic fluid. Moreover, inhibin potency was significantly higher in amniotic fluid than its corresponding serum level throughout the different stages of pregnancy.

**Table (1):** Level of inhibin potency (unit/ml) in serum and amniotic fluid of she-camel throughout pregnancy.

Stage of pregnancy	Serum	Amniotic fluid
Early stage (up to 5 months)	0.57 ± 0.02 <sup>aA</sup>	0.94 ± 0.02 <sup>abA</sup>
Middle stage (5-9 months)	0.64 ± 0.03 <sup>bA</sup>	1.17 ± 0.04 <sup>aA</sup>
Late stage (above 9 months)	0.77 ± 0.05 <sup>abA</sup>	1.28 ± 0.08 <sup>bA</sup>

± Standard error.

- Values within the same column having the same small letter are significantly different at least at (P<0.05).
- Values within the same row having the same capital letters are significantly different at (P<0.001)

## DISCUSSION

In the present investigation, it was found that inhibin level in serum of pregnant she-camel increase gradually with the advancement of pregnancy. Such result comes in accordance with previous studies in other species including man (Rieley et al., 1996; Nambo et al., 1996 Noble et al., 1997. Miller et al., 1997 and Wallace et al., 1997).

It is recorded that corpus luteum is considered as a source of inhibin in some species (Davis et al., 1986; Burger et al., 1996 and Bird et al., 1997). Fetal gonads have been proved to be also a source of inhibin in most species if animals (Nambo et al., 1996 and Miller et al., 1997). Moreover, the placenta in women is also referred to as an inhibin source (Mayo et al., 1996 and Petraglia, 1997). Based on these result the increased inhibin level in serum of pregnant she

camel is anticipated and may be due to all these factors or some of them.

The present results also showed that inhibin level in the amniotic fluid in pregnant she camel increased gradually with the increase in pregnancy duration. This finding comes in agreement with the results of previous studies in other species (Riley et al., 1996; Nambo et al., 1996 and Miller et al., 1997). It can be suggested that such increase in inhibin level in the amniotic fluid is due to the inhibin secretory activity of foetal gonads (Aria et al., 1997; Nambo et al., 1996 and Miller et al., 1997).

Further investigation are required to declare whether the placenta in she-camel possesses the potentiality of inhibin secretory activity. Moreover, the obtained results clarify that the levels of inhibin of amniotic fluid is higher than its corresponding levels of serum at different

stages of pregnancy. This results comes also, in accordance with previous studies (Nambo et al., 1996; Miller et al., 1997 and Wallace et al., 1997) and it again supports the speculation that fetal gonads in camel have inhibin secretory activity.

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