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## **CRYOPRESERVATION OF GOAT SPERMS COLLECTED FROM DIFFERENT REGIONS OF THE EPIDIDYMIS**

**K. U. E. Perera, H. B. S. Ariyaratne**

*Department of Basic Veterinary Sciences,  
Faculty of Veterinary Medicine and Animal Science, University of Peradeniya*

Cryopreservation of epididymal sperms in domestic and non-domestic animals has a number of important practical applications such as preservation of valuable genetic material from non-breedable animals.

In the present study, we assessed the motility, morphology and viability of sperms from the head, body and tail regions of 11 epididymides obtained from goats from the Kandy abattoir soon after slaughter. Eosin and nigrosin staining was used to assess the changes of morphology and viability.

Statistically significant differences were observed in the motility and viability of sperms in the head, body and tail of the epididymis ( $P < 0.05$ ) before and after cryopreservation. Before cryopreservation, the highest motility and viability were detected in sperms collected from the tail of the epididymis and the mean values ( $\pm$  standard error) observed were  $75.45 \pm 5.22\%$  and  $78.18 \pm 5.65\%$  for motility and viability respectively. These values reduced to  $39.09 \pm 8.31\%$  and  $37.72 \pm 4.96\%$  after freezing. The lowest motility and viability were observed in sperm of the head of the epididymis, which were  $5.27 \pm 2.72\%$ , and  $47.27 \pm 10.52\%$  respectively and reduced to  $1.00 \pm 0.89\%$  and  $24.18 \pm 6.90\%$  after freezing. The motility and the viability of sperms from the body of the epididymis were found to be  $35.7 \pm 31.39\%$  and  $58.3 \pm 16.48\%$  and reduced to  $14.8 \pm 13.7\%$  and  $31.5 \pm 8.31\%$  respectively due to freezing. Cryopreservation significantly increased ( $P < 0.05$ ) numbers of head and tail abnormalities, such as small heads, acrosomal defects, coiled tails and bent tails, in sperms collected from all three parts of the epididymis. A significant difference ( $P < 0.05$ ) was detected in the proportion of sperms having proximal cytoplasmic droplets among the head, body and tail regions of the epididymis, whereas no such difference ( $P > 0.05$ ) was observed for the distal cytoplasmic droplet in sperms collected from the different regions before cryopreservation. Similar trends were observed in sperms collected from these regions after freezing. Analysis of variance revealed a significant interaction ( $P < 0.05$ ) between cryopreservation and the region of the epididymis for the parameters of motility, viability and acrosomal defects.

From these observations, it was concluded that the cryopreservation was most successful when the sperms were obtained from the tail of the epididymis.