

collect from the same animals in two different physiological stages: 67 ± 7 days of pregnancy and 30 days after weaning. Discriminant analysis with PCA/MDR algorithm coupled to Mahalanobis distance (MD), were used to identify pregnancy through spectra similarity, assuming three or more as dissimilar spectra. Spectra of pregnant or non-pregnant ewes from one breed were not consistently similar to other breed in the same reproductive stage, with a range of 33 to 89% of prediction accuracy in the spectral match. Greater breed effect was observed for pregnant ones, probably reflecting prolificacy differences. Spectra from pregnant animals were different from non-pregnant ones and vice versa (98 to 100% of prediction accuracy), regardless of the breed, suggesting they may be used to classify animals based on pregnancy. To minimize breed effect, all spectra were pooled to create two databases (pregnant and non-pregnant). From these data sets, a randomized subset of 42 fecal spectra, not used to equation calibration, were selected and used for validation of procedure. Both pregnant and non-pregnant discriminant equations were able to identify and classify the animals according to pregnancy with 100% of accuracy, confirming the potential of fecal NIRS to identify pregnancy in livestock. However, a statistical significant difference was observed for diet CP and organic matter digestibility (OMD) between pregnant and non-pregnant ewes (T-test, $P < 0,001$), reflecting changes in pasture quality during data collection. Thus, the potential effect of season on pregnancy-related spectral differences remains to be investigated and also a larger database need be evaluated before this technique is recommended for early pregnancy diagnosis in sheep.

Key Words: discriminant analysis, gestation, NIRS

688 Ability to culture of cells from postmortem goat skin tissues stored at room temperature for different time intervals. M. Singh* and X. Ma, *Fort Valley State University, Fort Valley, GA.*

Animal cloning technology has renewed the interest in postmortem tissue storage, since these tissues can be used to reintroduce the lost genes back into the breeding pool in animal agriculture, preserve the genetic diversity, and revive endangered species. Several studies have demonstrated that cell survival decreases with increasing postmortem tissue storage. However, the limits of time interval within which live cells can be recovered from dead animals is not adequately studied. Cell viability and their potential to in vitro culture ensure nuclear integrity, a requirement for successful cloning of animals. To test the postmortem storage limits of animal tissues, 2–3 mm² skin pieces (n = 70) from ears of 3 breeds of goats (n = 7) were cultured after 0, 2, 4 and 6 d of postmortem storage at 24°C. After 10 d of culture, outgrowth of fibroblast-like cells (>10 cells) around the explants was scored. All the explants irrespective of breed displayed outgrowth of cells on the dish containing fresh tissues (d 0). However, the number of explants exhibiting outgrowth reduced with increasing time interval. Only 53.8% explants displayed outgrowth after 2 d of tissue storage. The number of explants displaying outgrowth was much smaller after 4 d (16.7%) and 6 d (13.3%) of storage. In general, the number of outgrowing cells per explant, on a given day, also decreased with increasing postmortem storage time interval. To test the differences between cell cultures, obtained from postmortem fresh and stored tissues, secondary cultures were established from one of the goats exhibiting outgrowth of cells after 6 d of tissue storage. Comparison of both the cell lines revealed similar cell morphology and growth curves, and had doubling times of 23.0 h and 22.6 h, respectively. These results suggest that live cells can be recovered from skin tissues of goats and perhaps other animals even after 6 d of their death with comparable growth profiles.

Key Words: fibroblasts, postmortem tissue storage, goat skin