ANIMAL PROTOCOL REVIEW

Form 12. Adverse events and early ending of the study

Institutional Animal Care and Use Committee, Ross University, School of Veterinary Medicine

ADVERSE EVENTS (AEs) Note: All AEs must be reported within 48 hours.

Describe potential adverse events. (Possible local and/or systemic AEs)
How often will the animals be monitored for adverse events and by whom?
Apart from the Attending Veterinarian (AV) and the PI, to whom else will AEs be reported?

EARLY STUDY TERMINATION (for an individual animal)

Provide clinical endpoints for removing animals from the study. Describe the conditions, complications and criteria that would lead to euthanasia of the animal before the expected endpoint of the study.

All AEs most be reported by submitting a seperate Adverse Event Form to the AV and thus to the IACUC.

Note: All studies must have this form filled out to address possible adverse events and animal follow-up.