# Lab #3 <a href="McMasters & Ruminant Parasite Ova">McMasters & Ruminant Parasite Ova</a>

#### What you should accomplish during Lab #3.

- 1. A description of the laboratory organization.
- 2. After an introduction, students will prepare and examine a McMasters Technique
- **3.** Be able to Identify parasite ova & oocysts commonly found in ruminant feces.

#### LABORATORY ORGANIZATION

Parasitology drawers with supplies.

#### LABORATORY SUPPLIES

Some supplies that are used during the lab are listed below. Please note the "disposable supplies" versus the supplies that need to be retained.

#### Non-Disposable Supplies

- 1. McMaster's counting chamber
- 2. McMaster's measuring vial
- 3. Plastic cups
- 4. Tea strainers
- 5. Microscope slide box
- 6. Sodium nitrate containers
- 7. Test tube racks
- 8. Plastic Transfer Pipettes ←(\*\*\*Do Not Dispose\*\*\*)
- 9. Flotation vials (brown) ←(\*\*\*Do Not Dispose\*\*\*)
- 10. Centrifuge tubes ←(\*\*\*Do Not Dispose\*\*\*)

#### <u>Disposable</u>

- 1. Wooden applicator sticks
- 2. Wooden tongue depressors
- 3. Glass slides & Cover slips

## **Technique**

#### McMaster's Quantitation Technique

1) Fill McMaster's Graduated Vial (clear vial with 2 lines) to the bottom line with flotation solution (= 26 mls).

- 2) Add feces, about 4 gm, until the fluid level rises to the top line.
- 3) Pour this mixture into a clean beaker and mix thoroughly.
- 4) Pour mixture through a strainer into a 2<sup>nd</sup> clean beaker.
- 5) Mix strained mixture by pouring mixture from beaker to beaker a few times.
- 6) Withdraw a small amount of the well-mixed suspension with a pipette and load this into one side of the McMaster's counting chamber.
- 7) Mix suspension again by pouring mixture from beaker to beaker a few times.
- 8) Again, withdraw a small amount of the well-mixed suspension with a pipette and load the second side of the McMaster's counting chamber.
- 9) Wait 1 minute for eggs to rise to the top of the chamber.
- 10) Focus on the lines of the McMaster's chamber with 4X, then examine the chamber with 10X. (Scan for ova).
- 11) Examine the entire ruled area, counting all the eggs within the ruled areas
- 12) Add the total egg from each side of the chamber.
- 13) Multiply the sum of the 2 chambers by 25 to determine the eggs per gram (epg).
- Note: The McMaster's Quantitation Techniques is mainly for the quantitation of <u>Strongyle-type ova</u>, thus only strongyle-type eggs should be counted. However, a general idea (i.e. none, few, many...) of the number of other nematode ova, cestode ova & coccidian oocysts should be noted.
- Note: The McMaster's Chambers can NOT be examined with the 40X or 100X objectives.

## **Exercises**

- 1. Fecal Worm Egg Count & Fecal Floatation.
  - a) Perform a McMasters on the provided goat feces.
    - i. Count the number of strongyle-type eggs in each grid.
  - b) Determine and record the resulting FWEC.

(Grid A: + Grid B: ) 
$$X 25 = epg$$

- c) Perform a Fecal Floatation with the strained fecal suspension.
- 2. Examine specimens of parasite diagnostic stages presented on the overhead monitors.

Oocysts: Eimeria.

Ova: Strongyle-type, Nematodirus, Strongyloides, Trichuris, Moniezia, Fasciola.

Utilize this lab and the Parasitology website to learn these diagnostic stages as one will be responsible for identifying these on the Lab Practical.

# Lab #3 <u>Laboratory Discussion</u> Complete and turn in as directed.

1. The FWEC: epg.	
2. List the Parasite ova / oocysts that you found i goat feces.	n the