Standard Operating Procedures (SOPs) for diagnosis of *Fasciola* spp. and Paramphistomidae eggs in cattle using the Sedimentation technique

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1. GOAL

These SOPs provide instructions on the laboratory activities, i.e. the preparation and copromicroscopic analysis of cattle faecal samples for the diagnosis of *Fasciola* spp. and Paramphistomidae eggs using the Sedimentation technique, in the framework of the PREPARE4VBD project.

2. EQUIPMENTS, MATERIALS AND REAGENTS

- Gloves
- Balance
- Becker
- Conical becker
- Cylinder
- Gauze or wire mesh (i.e., metal sieve having an aperture of 250 μm)
- Petri plate or glass slide and coverslide
- Conventional optical microscope
- A hand tally counter

3. METHODOLOGY AND PROCEDURES

3.1 Sedimentation Technique

- 1. Homogenize the sample thoroughly.
- 2. Weight 10 g of faeces in a becker, add 90 ml of tap water, homogenize and filter the faecal suspension using a double layer of gauze or a wire mesh having an aperture of 250 μ m in a conical becker with a capacity of 500 ml.
- 3. Add tap water up to 250 ml.
- 4. Wait for the mixture allowed to sit for 4 minutes.
- 5. Discard the supernatant.
- 6. Refill the conical becker with tap water up to 250 ml.
- 7. Repeat steps 4 to 6 for three times more.
- 8. Discard the last time the supernatant and transfer 10-15 ml of sediment in a Petri plate or add some drops of sediment on a glass slide, add a coverslip and examine under the microscope.
- 9. Look for eggs using a 10x objective and count the eggs.

3.2 Cleaning Sedimentation kit

You can wash all the plastic materials after use before the next assay.

References

Foreyet W.J. Veterinary Parasitology. Fifth Edition, 2001, Iowa State Press, pp. 145.

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Appendix 1. Faecal sample analysis sheet - Sedimentazion Technique

Farm ID:		
City:	Province:	
GPS coordinates: Latitude (E/W):	Longitude (N/S): Altitud	le:
Name of Analyst: Date of Analyzes:		
Cattle ID	Fasciola spp.	Paramphistomidae
	Number of eggs	Number of eggs
1		
2		
3		
4		
5		
6		
7		
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