Parasitological examination of stool

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Caprology or Scatology ?

Introduction

- While isolation of the infecting agent and detection ofspecific antibodies are the major methods in bacteriology and virology, they are of much less importance in parasitology than morphological identification of the parasite by microscopy.
- Due to complex antigenic structure and extensive cross-reactions, serological diagnosis is of limited value in parasitic infections.

- Morphological diagnosis of parasites consists of two steps:
- (1) detection of the parasite or its parts in clinical samples (2) its identification.
- I. Detection depends on collection of the appropriatesamples and the ir examination by suitabletechniques.
- 2. Identification requires adequate skill and expertise in recognizing the parasite in its various stages and its differentiation from morphologically similar artefacts.

Collection of Fresh Stool Specimen

- collected in a suitable,clean, wide mouthed container like a plastic container with a lightfitting lid, waxed cardboard box, or match box.
- The specimen should not be contaminated with water, urine, or disinfectants.

Liquid stools :examined or preserved within 30 minutes of passage
Soft stools:within 1 hour of passage
Formed stool:with in 24 hours of passage.
Normally passed stools are preferable,

- although samples obtained after purgative (sodium sulfate) or high saline enema may also be used.
- Examination of fresh specimens is necessary for observing motility of protozoan parasites.

Gross Examination:

- consistency
- color,
- Odor
- pH
- presence of blood, pus, Mucus.
- Parasites: tapeworm proglottids, roundworm, pinworm.

Microscopic Examination

- microscope should be equipped with a micrometereyepiece, as it is often essential to measure the size of parasites.
- For example, the differentiation between cysts of the pathogenic *Entamoeba histolytica* and thenonpathogenic *E. hartmanni* is based entirely on their sizes.

Microscopy should also include contributory findings

- Charcot-Leyden crystals
- Pus cells(WBCs)
- Red blood cells (RBCs)
- Macrophages.

Methods:1Wet Mounts

- Wet saline mounts: Trophozoites of E.histolytica, Balantidium coli and Giardia lamblia.
- Eggsof helminths
- Rhabditiform larvae of Strongyloides stercoralis
- Iodine staining(Lugol's iodine):cysts, as it prominently stains the glycogen vacuoles and nuclei. Protozoan cyst stained with iodine show yellow gold ,cytoplasm, brown glycogen material and pale retractile nuclei.

Wet Preparation



2:Permanent Stained Smears

Iron-hematoxylin stain:

Trlchrome stain

Modified Ziehl-Neelsen (acid-fast) stain

Concentration Methods

They can be classified as

- Floatation :feces are suspended in a solution of high specific gravity, so that parasitic eggs and cysts float up and get concentrated at the surface
- Sedimentation: feces are suspended in a solution with low specific gravity, so that the eggs and cysts get sedimented at the bottom, either spontaneously or by centrifugation.

Floatation Methods

Saturated salt solution technique

Zinc sulfate centrifugal floatation

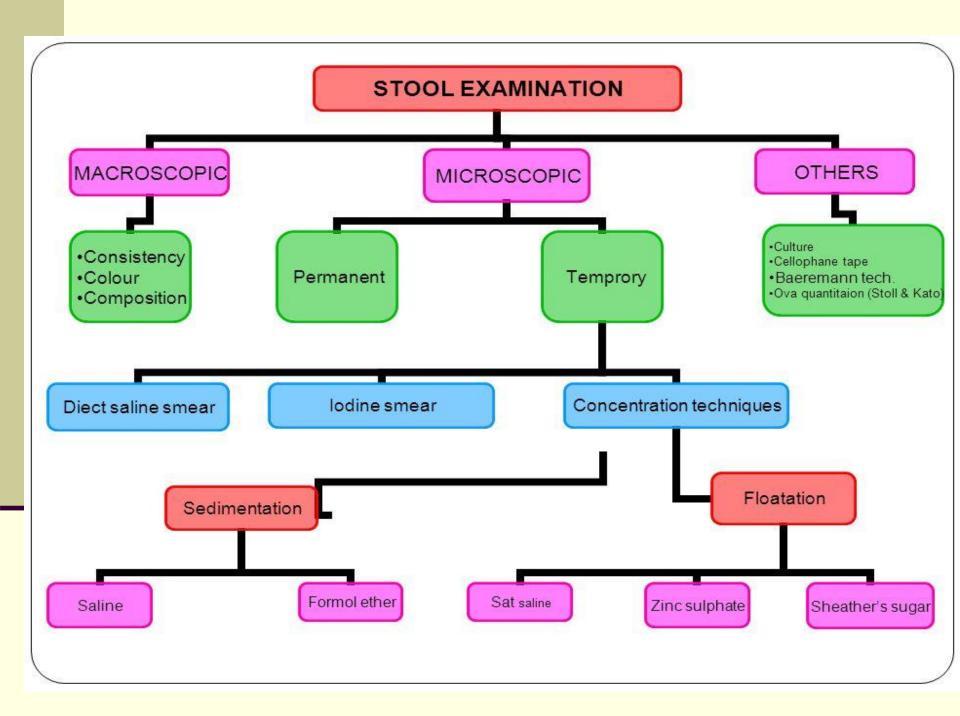
- Applicable for Fertilised egg of roundworm, hookworms and whipworm,
- Not applicable for eggs of tapeworms, unfertilized egg of Ascaris lumbricoides, eggs of trematodes and protozoan cysts

Sugar floatation technique:

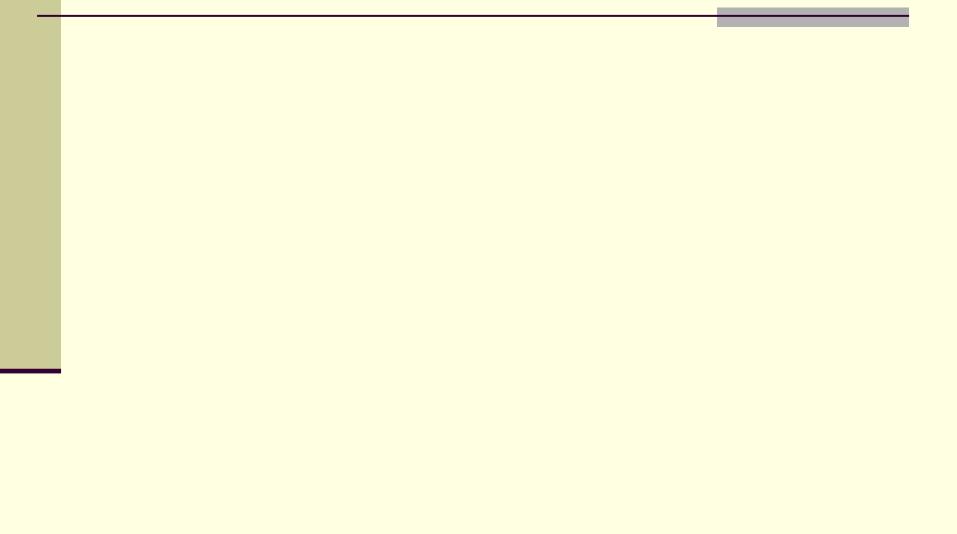
- Sheather's sugar floatation technique is recommended
- for the detection of cryptosporidia infection

Sedimentation Methods

- Formal-ether sedimentation technique
- Baermann concentration method
- The method is useful for all helminth eggs and protozoan cysts.













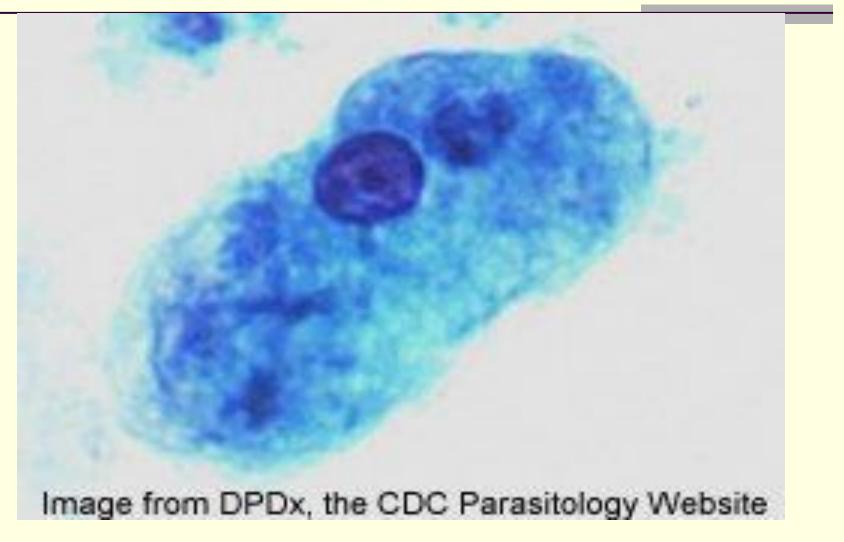
Identify: Iodine Preparation



Identify:Saline preparation



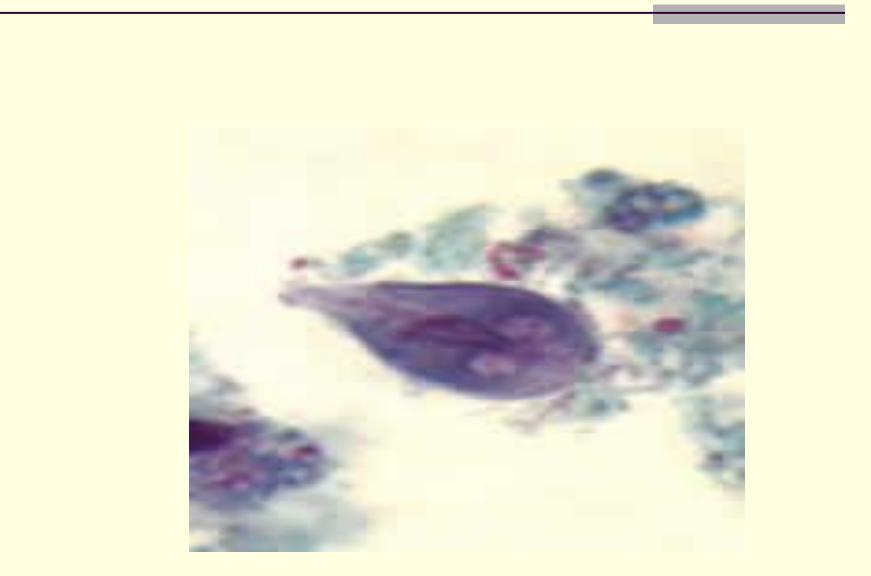








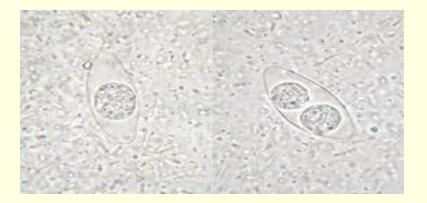
Trichrome stain

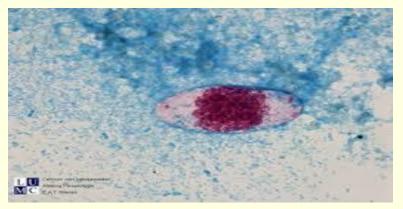


Modified ZN Stain

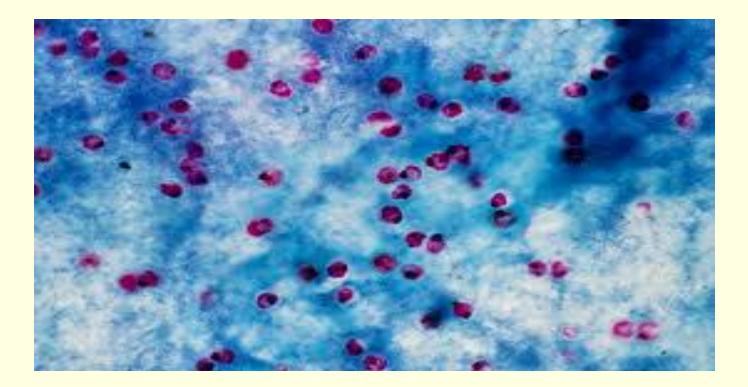


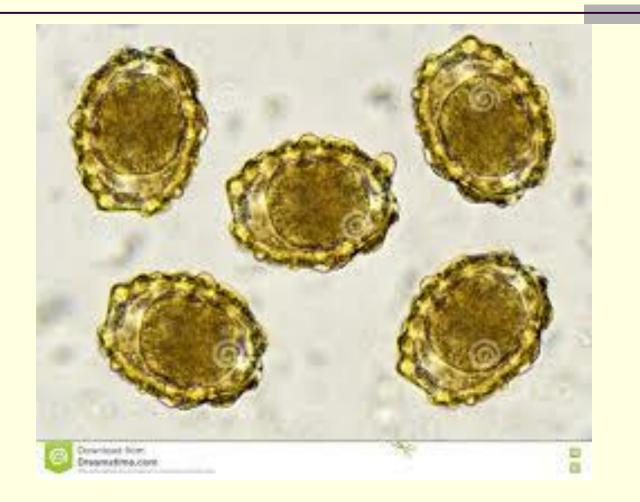
Modified ZN Stain





Modified ZN Stain



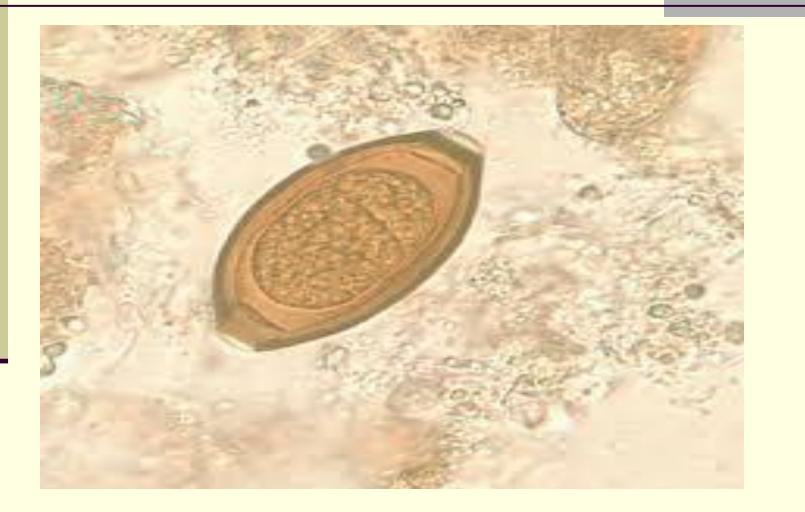
























Qoestion ?

- Example of Bile stain Eggs & Non Bile stain Eggs
- Eggs float in sss
- Eggs Sink in sss

Thank You