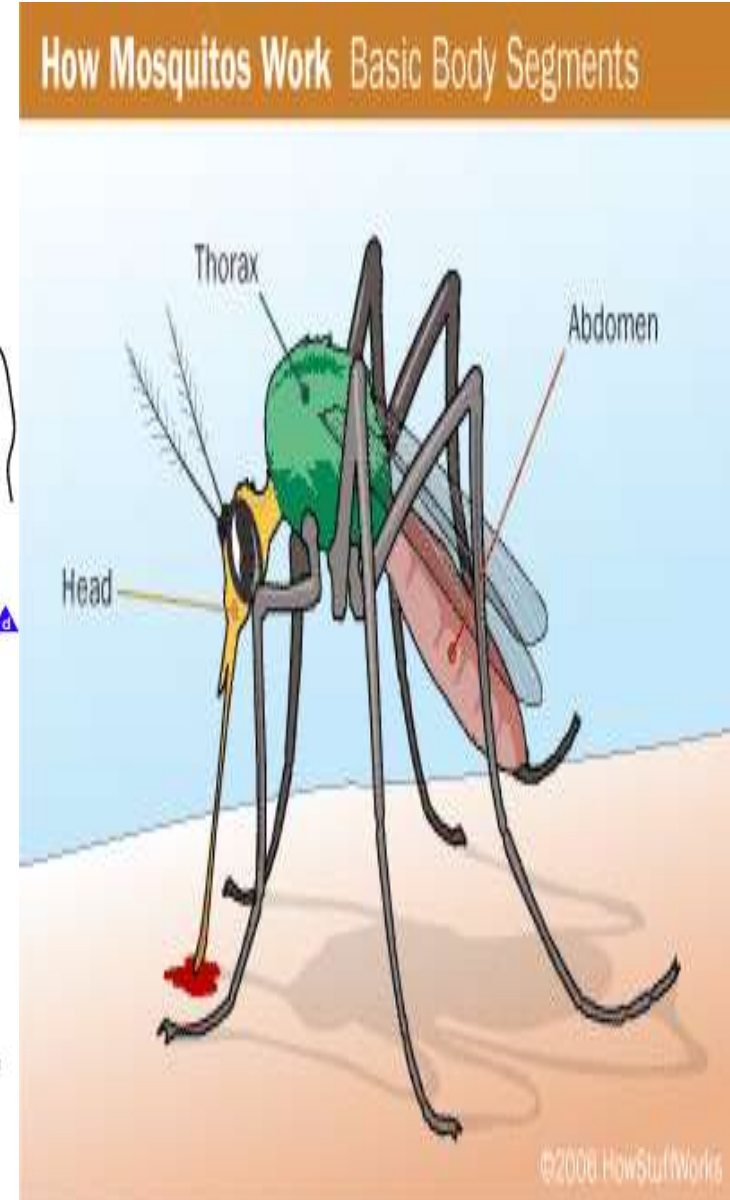
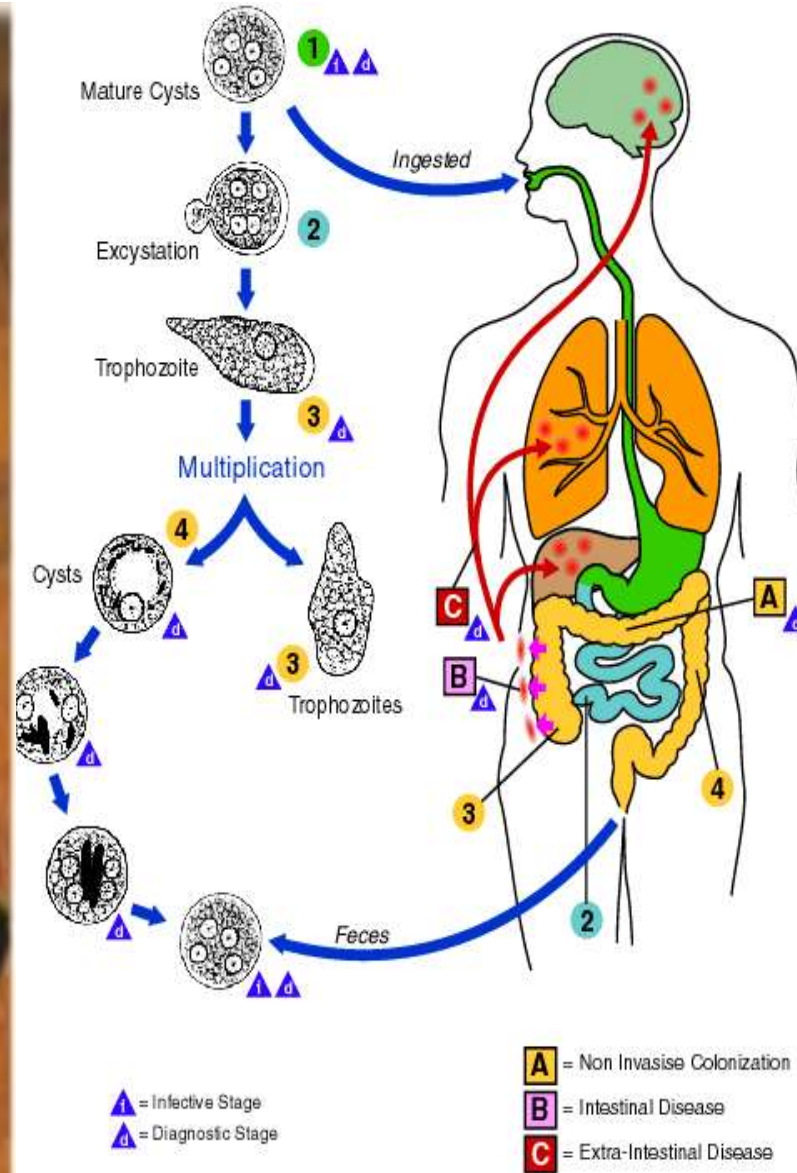


# MEDICAL PARASITOLOGY



# MEDICAL PARASITOLOGY

## INTRODUCTION :

- The *parasites* which infect man
- The *disease* they produce
- The *response* generated by them
- Various methods of *diagnosis*
- *Prevention.*

**PARASITE :** *An organism that entirely dependent on another organism ie its host, for all or part of its life cycle and metabolic requirements.*

### ***Microparasite & Macroparasite***

- On basis of their location – *Ectoparasite & Endoparasite*

**ECTOPARASITE** – lice – causes infestation.

**ENDOPARASITE** – all protozoan & helminthic – cause infection.

## **OBLIGATE PARASITE :**

- Cannot exist without a host eg. *Toxoplasma gondii*

## **FACULTATIVE PARASITE :**

- Live either a parasitic or free-living existence eg. *Naegleria fowleri*, *Acanthamoeba* spp., *B. mandrillaris*

**ACCIDENTAL PARASITE :** Attack an unusual host eg. *Echinococcus granulosus* in man

**HOST :** Harbours the parasite and provides the nourishment and shelter

**DEFINITIVE HOST :** sexual reproduction

**INTERMEDIATE HOST :** larval or asexual stages

**RESERVOIR HOST :** Harbours the parasite and serve as imp. source of inf. to other hosts

## **VECTOR :**

- Usually an insect, that transmits an infection from one host to another e.g. housefly

# HOST-PARASITE RELATIONSHIP

## SYMBIOSIS :

- An association, one cannot live without the help of the other

## COMMENSALISM :

- Only parasite gets benefit without causing any injury to the host

## PARASITISM :

- Parasite gets benefits from the host and host always suffers from some injury

# SOURCES OF INFECTION

## CONTAMINATED SOIL AND WATER :

- Eggs of *A.lumbricoids*, *Trichuris trichiura*
- Cysts of *E.histolytica*, *Giardia lamblia* etc.

## RAW OR UNDERCOOKED PORK :

- *T. solium*, *Trichinella spiralis*

## RAW OR UNDERCOOKED BEEF : *T. saginata*

## BLOOD-SUCKING INSECTS : *Plasmodium* spp., *Wuchereria bancrofti*, *Leishmania* spp. etc.



**HOUSEFLY** : *E. histolytica*

**DOG** : *Echinococcus granulosus*

**CAT** : *T. gondii*

**MAN** : *E. histolytica*, *Enterobius vermicularis* and  
*H. nana*

**AUTOINFECTION**: *E. vermicularis* & *S. stercoralis*

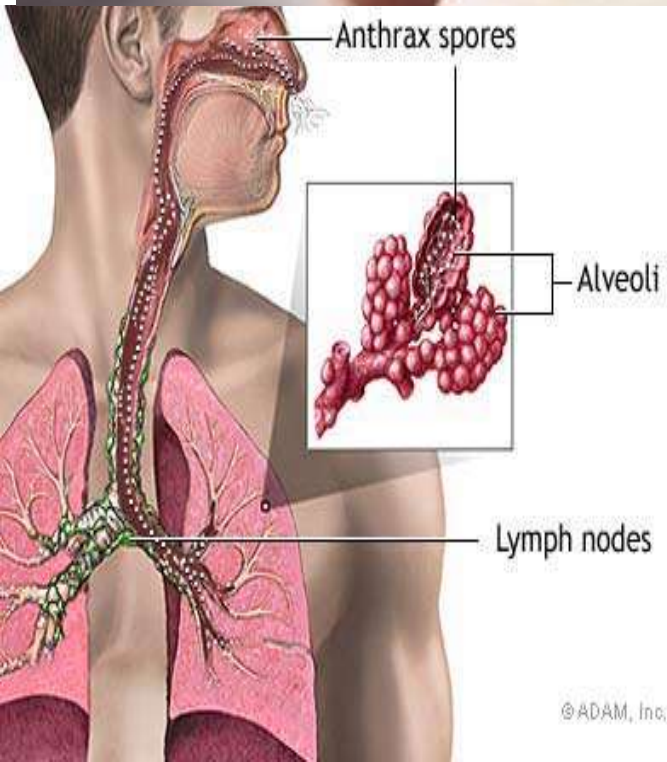
- Freshwater fishes , crab and crayfishes :  
*Diphyllobothrium latum*, *Paragonimus westermani*

# PORTAL OF ENTRY

**MOUTH** : E.histolytica, G.lambliia, B.coli, etc.

**INHALATION** : Eggs of E.vermicularis

**SKIN** : A.duodenale, S.stercoralis, Plasmodium spp., Leishmania spp., W.bancrofti etc.



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**CONGENITAL** : T.gondii, Plasmodium spp.

**SEXUAL CONTACT** : Trichomonas vaginalis etc.

**IATROGENIC INFECTION** : Malaria parasites  
may be transmitted by transfusion or by  
contaminated syringes and needles

The most common methods of transmission of HIV are:

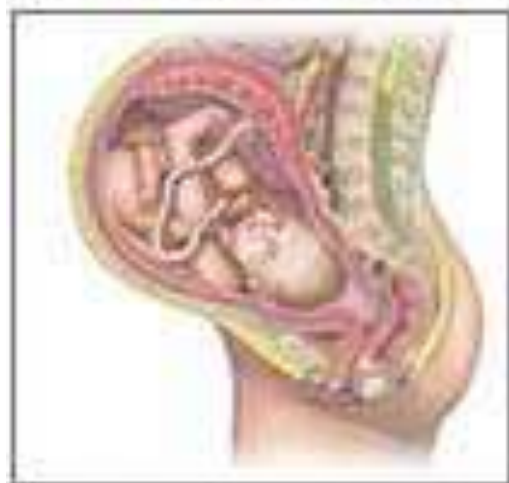


Unprotected sex with an infected partner



Sharing needles with infected person

Almost eliminated as risk factors for HIV transmission are:



Transmission from infected mother to fetus



Infection from blood products

# PATHOGENICITY

## TRAUMATIC DAMAGE :

- By entry of filariform larvae of *S.stercoralis* etc.
- By attachment of Hookworms to the intestinal wall
- Eggs of *S.haematobium* and *S.mansoni* in urinary bladder and intestinal canal
- Large worms - *A.lumbricoides* and *T.saginata* may produce intestinal obstruction

## **LYTIC NECROSIS –**

- *E.histolytica* secretes lytic enzymes which lysis tissues.
- *Plasmodium* spp., *Leishmania* spp. cause necrosis during their growth and multiplication

## **ALLERGIC MANIFESTATIONS :**

- By secretions and excretions of the growing larvae and the products liberated from dead parasites.

## **INFLAMMATORY REACTION :**

- Most of the parasites provoke cellular proliferation and infiltration at the site of their location.
- Cause eosinophilia
- Anaemia
- Black water fever in malaria
- Inflammation of L.I (E.histolytica)



## **NEOPLASIA :**

- Schistosoma haematobium can cause vesical carcinoma.

## **SECONDARY INFECTION :**

- The migrating larvae e.g. strongyloidiasis, ascariasis - may carry bacteria and viruses from intestine to the blood & tissue.

# IMMUNITY IN PARASITIC INFECTIONS

- Less efficient than bacterial and viral infections
- **CMI** – Cytotoxic T (Tc) cells, Natural killer (NK) cells, Activated macrophages.
- **AMI** – Antibody (produced by B-cells) mainly IgM, IgG, IgE.

# **PRESERVATION OF STOOL SPECIMENS :**

## **FORMALIN SOLUTION -**

- 10% formalin saline – 3:1
- (+) cysts, eggs and larvae
- (-) permanent stained smear, trophozoite, PCR

## POLYVINYL ALCOHOL (PVA) -

- Ethyl alcohol + HgCl<sub>2</sub> + GAA + Glycerine + PVA
- 3:1
- (+) cyst, trophozoites, trichrome staining
- (-) acid fast stain, safranin stain

## MERTHIONATE-IODINE- FORMALIN (MIF) SOLUTION -

- Sol. 1 – thiomersal + formaldehyde + glycerol
- Sol. 2 – Lugol's iodine
- **Stains and fixes** cysts, eggs, larvae without any need for further staining by wet mount
- Well preserved for 1 year or more

## SCHAUDINN'S SOLUTION -

- HgCl<sub>2</sub> + Ethyl alcohol + GAA + Glycerol
- 14:1
- It fixes and preserves the specimen for 1 year

# LABORATORY DIAGNOSIS

## DEMONSTRATION OF PARASITE :

### IN STOOL :

- Wet mount : Normal saline and Lugol's iodine (trophozoites, cysts, eggs)
- By concentration methods : Salt flotation or formal-ether con. method
- By Ziehl-Neelsen staining e.g. *Cryptosporidium parvum*, *Isospora belli*.

# EXAMINATION OF FAECES

## COLLECTION OF SPECIMEN :

- Normally passed stool
- No Barium enema specimen
- Three faecal samples x 3 days
- First two samples – During normal bowel movement
- 3<sup>rd</sup> sample – After magnesium sulphate purge
- Amount – Whole stool, series of stool samples, milligram amount



## **EXAMINATION OF STOOL SPECIMENS -**

- Liquid stool specimens – within 30 min
- Semiformed stool specimens – within 60 min
- Formed stool specimens – within 24 hrs

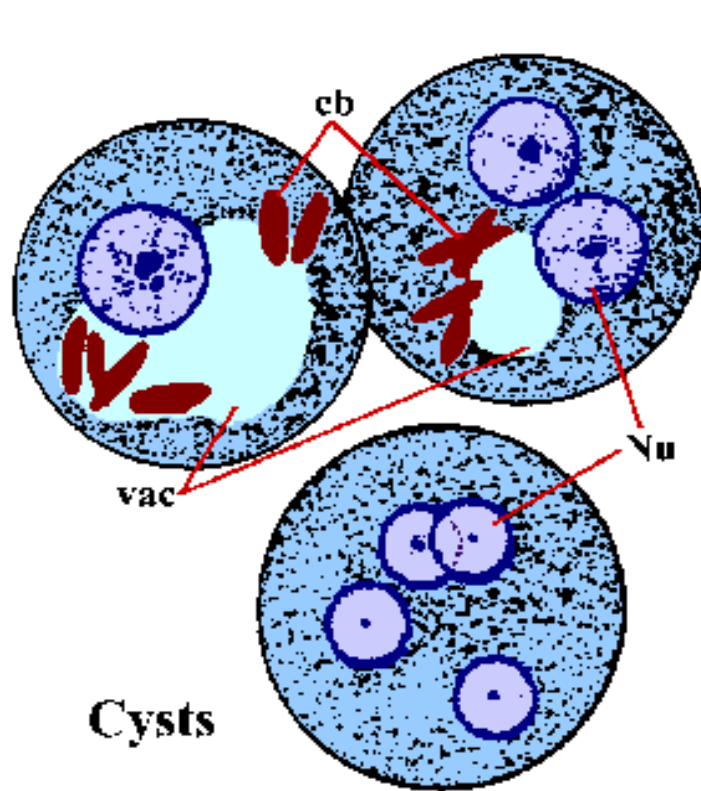
# METHODS OF EXAMINATION

## Macroscopic Examination -

- Consistency, Colour, Odour, blood or mucous
- Adult helminths – *A.lumbricoides*, *E.vermicularis*, segments of Tapeworms

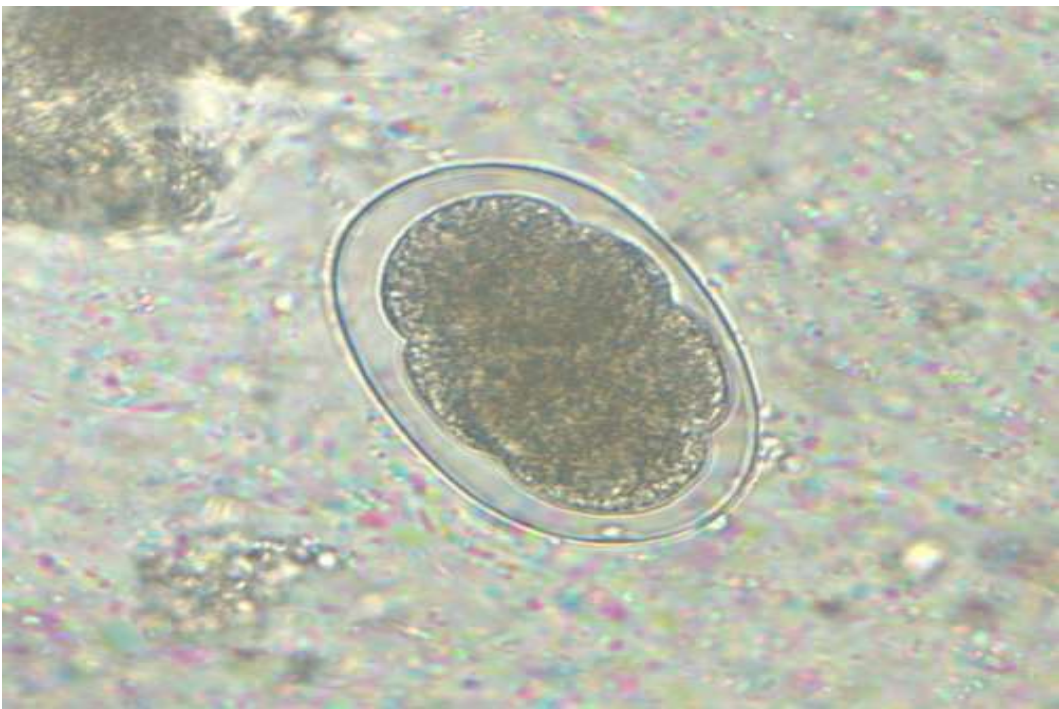
## Microscopic Examintion -

- Saline wet mount
- Iodine wet mount
- Stains – Iron-haematoxyline stain, Trichrome stain, Modified acid-fast stain

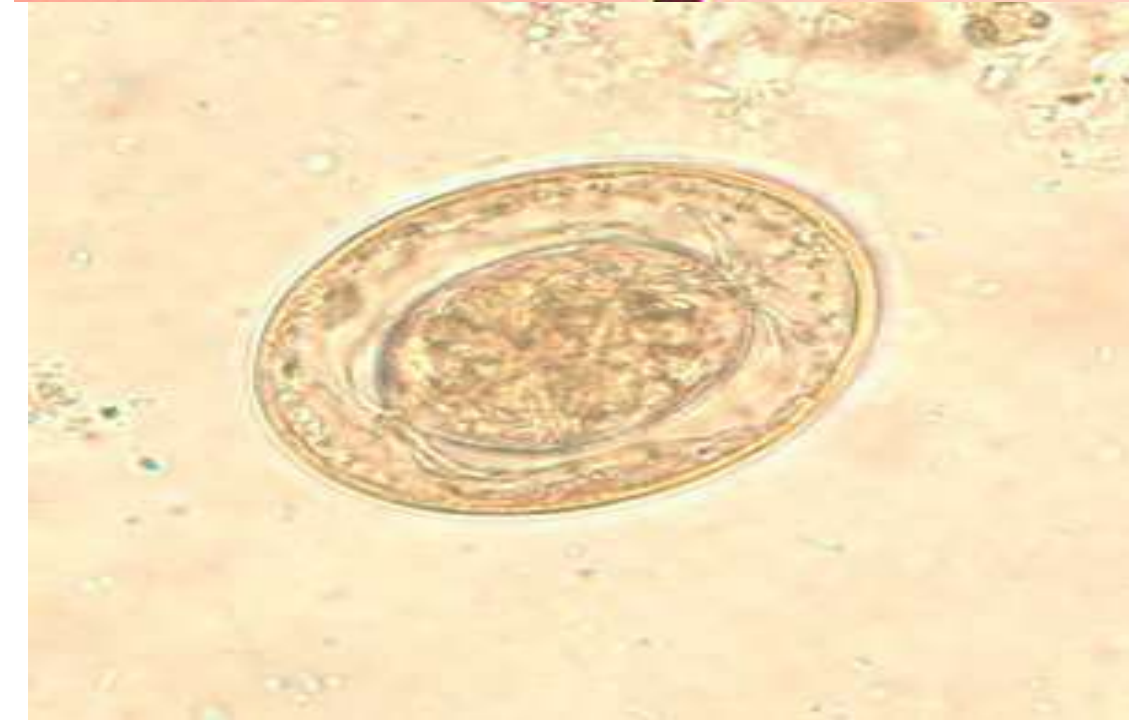




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# **CONCENTRATION METHODS :**

- Flootation Technique
- Sedimentation Technique

## **FLOATATION TECHNIQUE :**

### **SATURATED SALT FLOATATION TECHNIQUE:**

- All the helminthic eggs float except unfertilized eggs of *A.lumbricoides*, eggs of taenia and all intestinal flukes

### **ZINC SULPHATE (33%) CENTRIFUGAL FLOATATION TECHNIQUE –**

- Concentrates cysts of protozoa, eggs of nematodes and small tape worms

## **SEDIMENTATION TECHNIQUE :**

- Simple sedimentation
- Formalin-ether sedimentation



## FORMALIN-ETHER SEDIMENTATION -

- Stool + 10ml water ----> filtrate ----> centrifuge x 2000rpm x 2min ----> discard supernatant
- Sediments + 10ml saline ----> centrifuge ----> discard supernatant
- Sediments + 7ml formalin saline ----> stand for 10min or longer ----> +3ml ether ----> mix it
- Centrifuge x 2000rpm x 2min

**Four layers** --- ether --- debris --- formalin ---  
sediment

**Good method** – Hypertonic sol. rupture the cysts and eggs

## **QUANTIFICATION OF WORM BURDEN :**

- Direct smear egg count
- Stoll's method

### **Direct smear egg count -**

- 2mg of faeces in a drop of saline
- Examine under low power
- Count the no. of eggs and calculate it per gram

## STOLL'S METHOD -

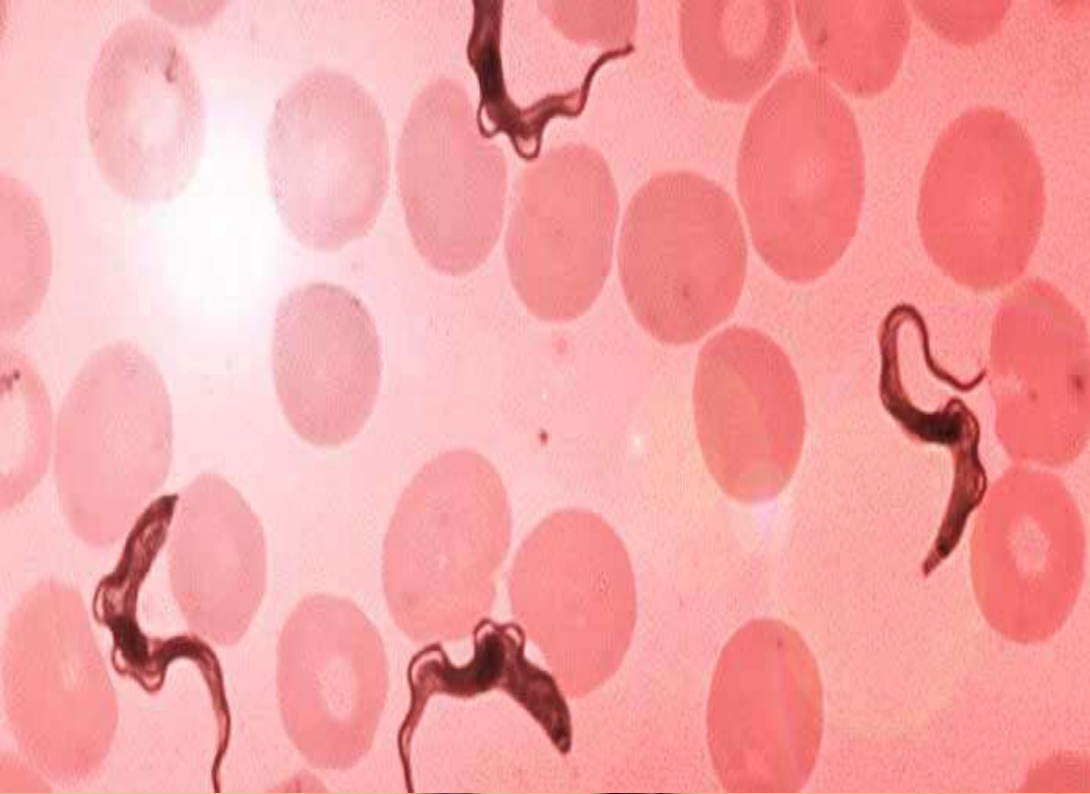
- 4gm of faeces + 56ml N/10 NaOH --- mix well
- Take 0.075ml on glass slide
- Count the eggs under low power ( a )
- $(a) \times 200 = \text{eggs / gm x 24 hr faecal sample}$

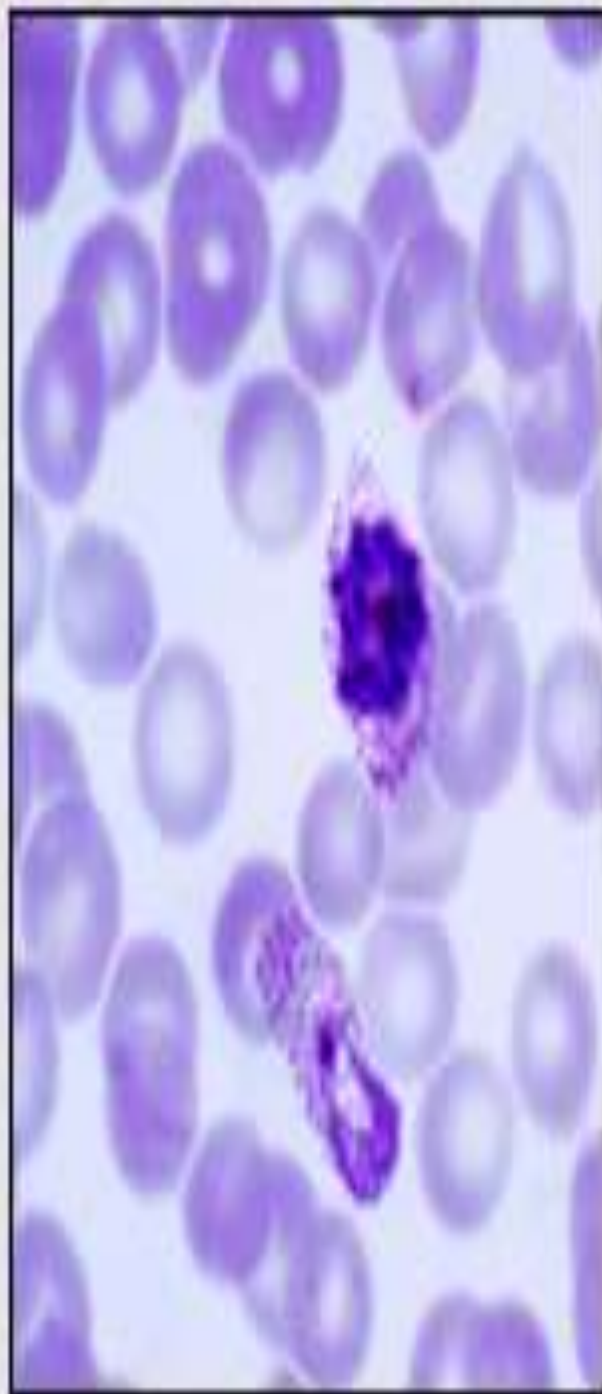
## **CORRECTION FACTOR (C.F)**

- Mushy-formed stool – **C.F – 1.5**
- Mushy stool – **C.F – 2**
- Mushy-diarrhoeic stool – **C.F – 3**
- Frankly diarrhoeic stool – **C.F – 4**
- Watery stool – **C.F - 5**

## In Blood :

- Wet mount – Trypanosomes and Microfilariae
- In Pbf – Thin & thick smear
- Staining – Leishman stain, Giemsa stain, Field stain, J.S.B stain
- Plasmodium spp., L.donovani, microfilariae of W.bancrofti





## **BLOOD CONCENTRATION METHODS :**

### **MICROHAEMATOCRIT CENTRIFUGATION –**

- For MP and trypanosome

### **TRIPLE CENTRIFUGATION –**

- 9ml blood + 1ml of 6% sod.citrate – centrifuge 100g x10min
- Supernatant -- centrifuge 250g x10min
- Supernatant -- centrifuge 700g x10min
- Sediments examined for trypanosomes



## **BUFFY COAT CONCENTRATION –**

- 5ml citrated blood
- For L.D, M.P, Trypanosomes

## **MEMBRANE FILTRATION -**

- For microfilariae in blood

## IN URINE :

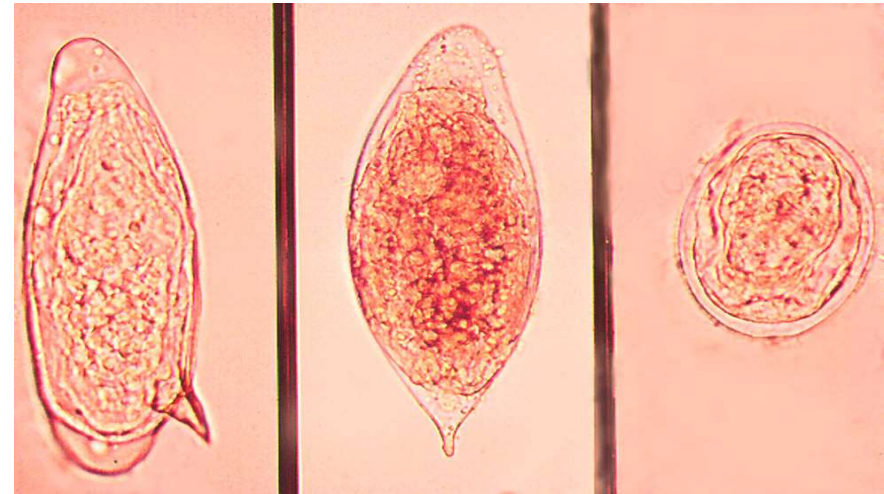
- Trophozoites of *T.vaginalis*, eggs of *S.haematobium*

## GENITAL SPECIMENS :

- Trophozoites of *T.vaginalis*

## CSF :

- Trophozoites of *N.fowleri*, *Acanthamoeba* spp., *B.mandrillaris*



## SPUTUM :

- Eggs of *Paragonimus westermani*, *E.histolytica*
- During migratory phase - larvae of *A.lumbricoides*, *Ancylostoma duodenale*, *Necator americanus*, *S.stercoralis*

## **TISSUE BIOPSY & ASPIRATION :**

- E.h from liver abscess
- G.lamblia from bile
- larvae of T.spiralis, T.solium in the muscle biopsy
- Scolices and brood capsules in the fluid aspirated from hydatid cyst

## **CULTURE :**

- E.h & G.l in stool
- Leishmania spp. in blood

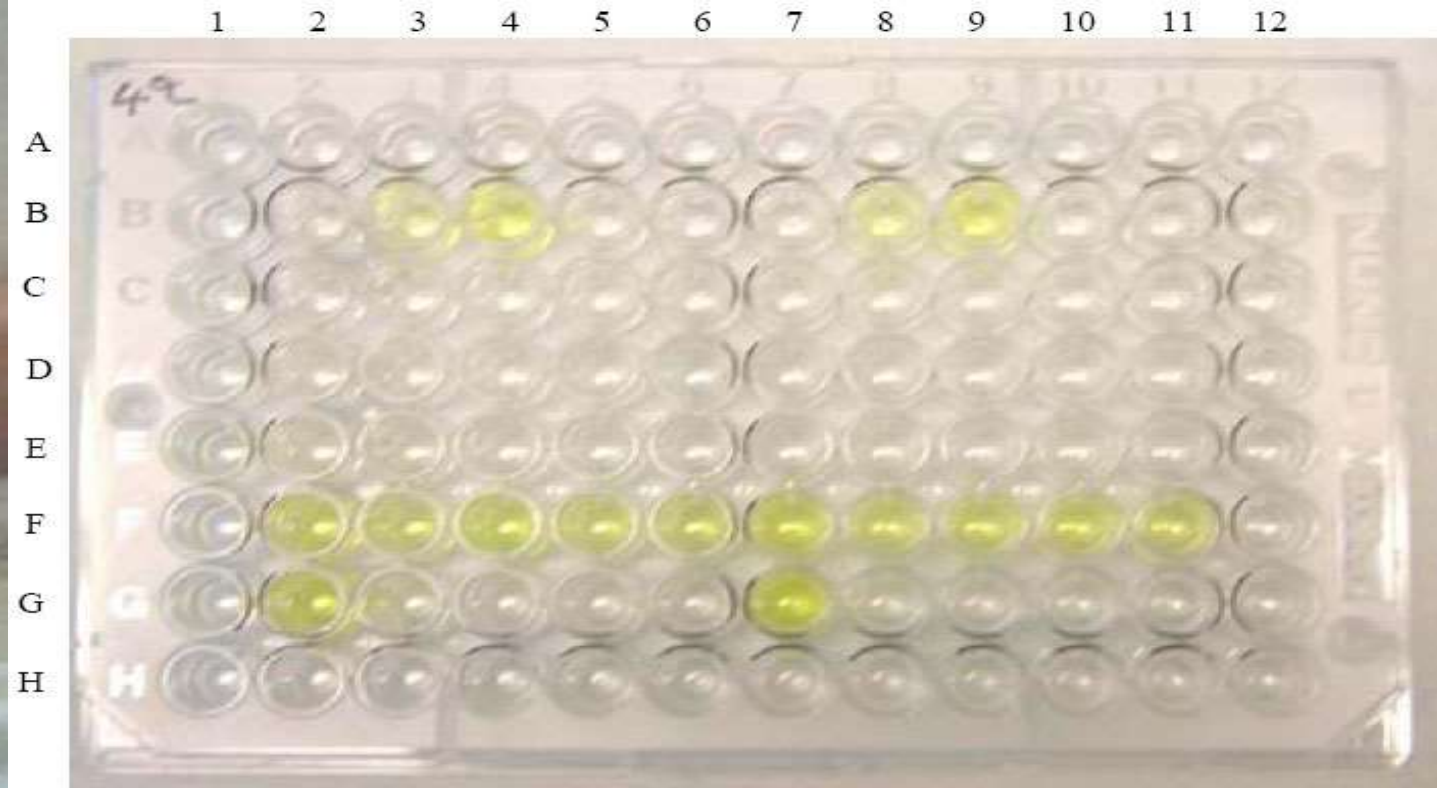
## **IMMUNODIAGNOSIS :**

**Skin Tests** – By Intradermal injection- Immediate & Delayed hypersensitivity reaction

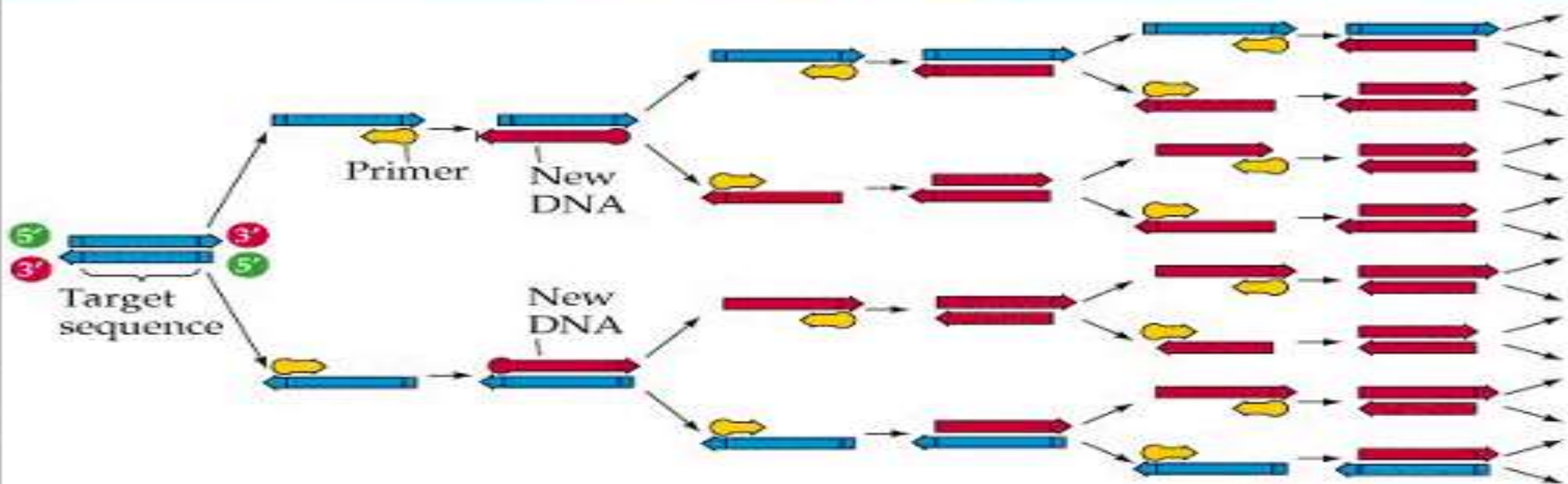
**Serological Tests** – Detection of antibodies or antigens by ELISA, RIA, Agglutination Tests, CFT, IHA etc.

## **MOLECULAR METHODS :**

DNA probes and Polymerase chain reaction (PCR)



### RESEARCH METHOD





**THANK YOU**