

Genito-Urinary System

*Schistosomiasis –
Schistosoma hematobium*

- Schistosomiasis—S.-hematobium
- Candidiasis—Candida-albicans
- Gonorrhea
- Chlamydia trachomatis, Ureaplasma and Gardnerella
- Syphilis
- Trichomoniasis
- Ectoparasitic infections: pubic lice, Scabies
- HSV
- HPV
- HIV

Helminthes

- Helminths are invertebrates characterized by elongated, flat or round bodies.
- Platyhelminthes (flat)
 - **Trematodes (flukes)**
 - Cestodes (tapeworms)
- Nematelminthes (circular)
 - Nematodes (Roundworms)

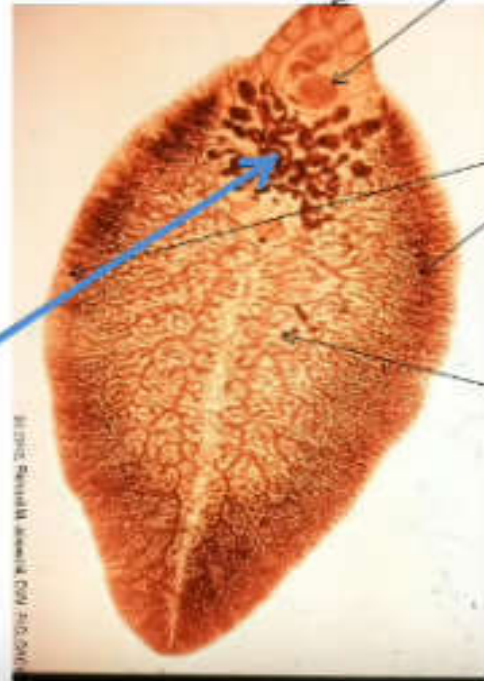


Trematodes



**Flat
Uterus**

Trematodes



Suckers

**Digestive
system**

Testes

Intermediate host : snail



حلازونة ياما الحلازونة خبينى
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Flukes, or trematodes

- Leaf-shaped, and vary in length from a few millimetres to 8cm.
- Excluding **blood flukes**, trematodes are **hermaphroditic**
 - having both male and female reproductive organs.
 - Both self-fertilisation and cross-fertilisation occur.
- **Blood flukes (schistosomes)** are the only bisexual flukes that infect humans
 - Within the definitive (human) host, male and female worms inhabit the lumen of blood vessels and are found in close physical association.



Common flukes (trematodes)

Organ inhabited

Fluke

Lung

Paragonimus westermani

Intestine

Fasciolopsis buski, *Heterophyes heterophyes*, *Mettgonimus yokagawi*

Liver

Clonorchis sinensis, *Opithorchis* species, *Fasciola hepatica*

Blood

Schistosoma mansoni, *Shistosoma haematobium*, *Schistosoma japonicum*



SCHISTOSOMIASIS

(BLOOD FLUKE INFECTION)

Parasitology

- The schistosomes are a group of closely related flukes that inhabit the portal vascular system of a number of animals.
- Of the five species known to infect humans, three, *S. mansoni*, *S. haematobium*, and *S. japonicum*, are of primary importance



- The 1 to 2 cm male possesses a deep ventral groove, or “schist.”
- Within this gynecophoral canal it carries the longer, more slender female in lifelong copulatory embrace.



- Each pair deposits 300 (*S mansoni*, *S haematobium*) to 3000 (*S japonicum*) eggs daily for the remainder of its 4- to 35-year life span.

- After mating of the adult worms in the portal vein, the conjoined couple use their suckers to ascend the mesenteric vessels against the flow of blood.
- *S. japonicum* enters the superior mesenteric vein,
 - the venous radicals of the small intestine and ascending colon;
- *S. mansoni* and *S. haematobium* are directed to the inferior mesenteric system.

- ***S. haematobium*** passes through the hemorrhoidal plexus to the systemic venous system, ultimately coming to rest in the venous plexus of the bladder and other pelvic organs.

- *Schistosoma japonicum*—veins of small intestines
- *Schistosoma mansoni*—portal veins of colon and rectum
- *Schistosoma haematobium*—veins of bladder and pelvic organs

S. haematobium

- On reaching the submucosal venules, the worms initiate oviposition.
 - Each pair deposits 300 eggs daily for the remainder of its 4- to 35-year life span.
- Ova lying immediately adjacent to the mucosal surface rupture into the lumen of the bladder and are passed to the outside in the urine.
- The eggs measure 60 by 140 μm and possess a terminal spine.

mansoni



haematobium



japonicum



Diagnostic Stage by Morphology

mansoni



-Lateral spine

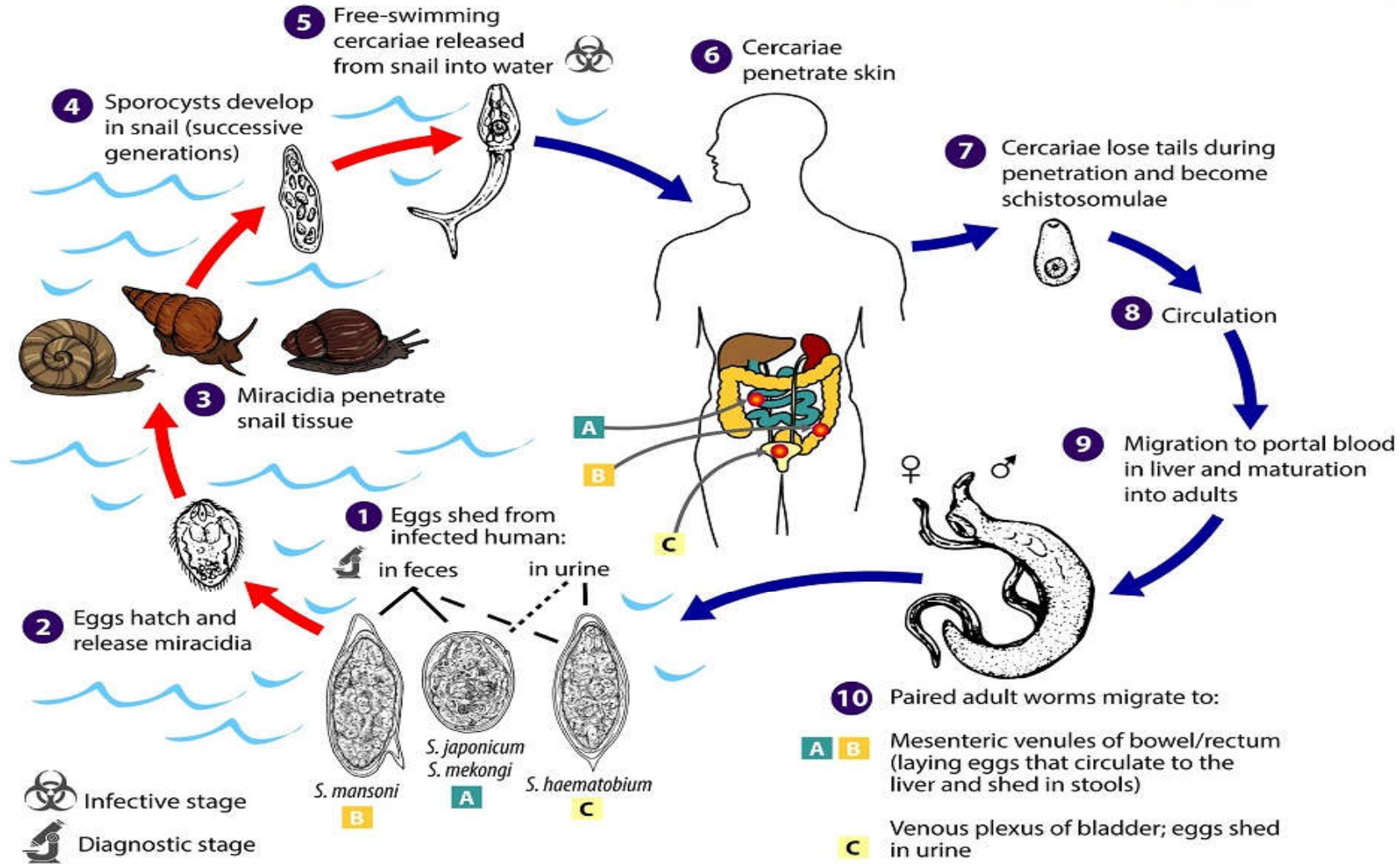
haematobium



-Central spine

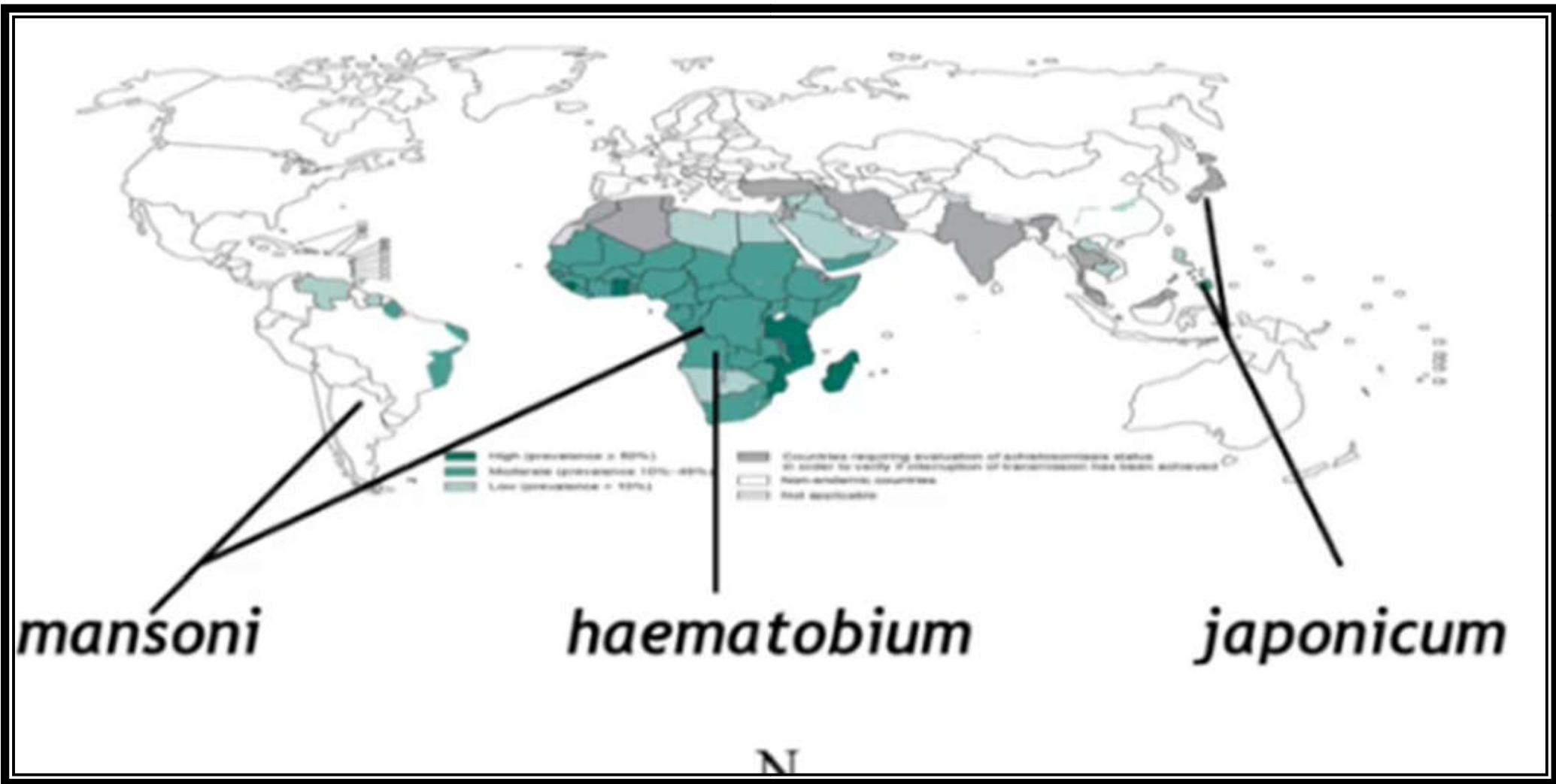
japonicum





Egg...miracidia.....snail.....cercariae.....skin...schistosomulae.....circulation

- Approximately 200 million people—almost 1 in 30 of all humans—are infected worldwide.
- Of these, roughly 200 000 will die annually.



- Most infected patients carry fewer than 10 pairs of worms in the vascular system and, accordingly, lack clinical manifestations of disease.

- **PATHOGENESIS**
- There are three major clinicopathologic stages in schistosomiasis.
- The first stage is initiated by the penetration and migration of the schistosomula.
- The second or intermediate stage begins with oviposition and is associated with a complex of clinical manifestations.
- The third or chronic stage is characterized by granuloma formation and scarring around retained eggs.

- CLINICAL ASPECTS

- Early Stage (within 24 hours)

- Penetrating the skin,
 - immediate and delayed hypersensitivity to parasitic antigens
 - results in an intensely pruritic papular skin rash that increases in severity with repeated exposures to cercariae.

- As the viable schistosomula begin their migration to the liver, the rash disappears and the patient experiences fever, headache, and abdominal pain for 1 to 2 weeks.

□ Intermediate Stage (One to two months)

- an acute febrile illness that bears a striking resemblance to serum sickness.
- fever and chills,
- patients experience cough, urticaria, arthralgia, lymphadenopathy, splenomegaly, abdominal pain, and diarrhea.
- The onset of oviposition leads to a state of relative antigen excess, the formation of soluble immune complexes, and the deposition of these in the tissues of the host.

- *S japonicum* infection
 - clinical manifestations of encephalitis. Typically, leukocytosis, marked peripheral eosinophilia, and elevated levels of IgM, IgG, and IgE immunoglobulins are present..

Katayama Syndrome

❑ Chronic Stage

- Approximately one half of all deposited eggs reach the lumen of the bowel or bladder and are shed from the body.
- Those retained induce inflammation and scarring, initiating the final and most morbid phase of schistosomiasis.

- Soluble antigens excreted by the eggs stimulate the formation of T lymphocyte–mediated eosinophilic granulomas.
- Early in the infection, the inflammatory response is vigorous, producing lesions more than 100-fold larger than the inciting egg itself.

- Inflammatory and fibrotic reactions to retained eggs cause chronic disease,
- the severity of tissue damage is directly related to the total number of eggs retained.

- In *S. haematobium* infection,
 - the bladder mucosa becomes thickened, papillated, and ulcerated.
 - Hematuria and dysuria result; repeated hemorrhages produce anemia.
 - In severe infections the muscular layers of the bladder are involved, with loss of bladder capacity and contractibility.
 - Progressive obstruction leads to renal failure and uremia.
 - Bladder carcinoma is frequently seen.

- Other urogenital organs may also be involved, including the spermatic cord, testes, fallopian tubes, ovaries, and vagina.

- In *S mansoni* and *S japonicum* infections, the bowel mucosa is congested, thickened, and ulcerated. Patients experience abdominal pain, diarrhea, and blood in the stool.
- Eggs deposited in the larger intestinal veins may be carried by the portal blood flow back to the liver, where they lodge in the presinusoidal capillaries.

- The resulting inflammatory reaction leads to the development of periportal fibrosis and hepatic enlargement.

- Presinusoidal obstruction of blood flow can result in portal hypertension and serious manifestations of portal obstruction.

- Eggs that are carried around the liver in the portosystemic collateral vessels may lodge in the small pulmonary arterioles, where they produce interstitial scarring, pulmonary hypertension, and right ventricular failure.

- Immune complexes shunted to the systemic circulation may induce glomerulonephritis.
- Occasionally, eggs may be deposited in the central nervous system, where they may cause epilepsy or paraplegia.

- DIAGNOSIS

- Definitive diagnosis requires the recovery of the characteristic eggs in urine, or biopsy specimens.

- filtering the urine through a membrane filter.

- Cystoscopy with biopsy of the bladder mucosa may be required for the diagnosis of mild infection.
- Conventional serologic tests detect circulating antibodies with sensitivities exceeding 90% but cannot distinguish active from inactive infection.

- TREATMENT

- No specific therapy is available for treatment of schistosomal dermatitis.

- Antihistamines** and **corticosteroids** may be helpful in ameliorating their more severe manifestations.

- TREATMENT

- Several antihelmintic agents may be used.

- Praziquantel**, which is active against all three species of schistosomes, is the agent of choice, although there is increased resistance to this single-dose oral agent in mass therapy programs. (C/I pregnancy)

- PREVENTION