



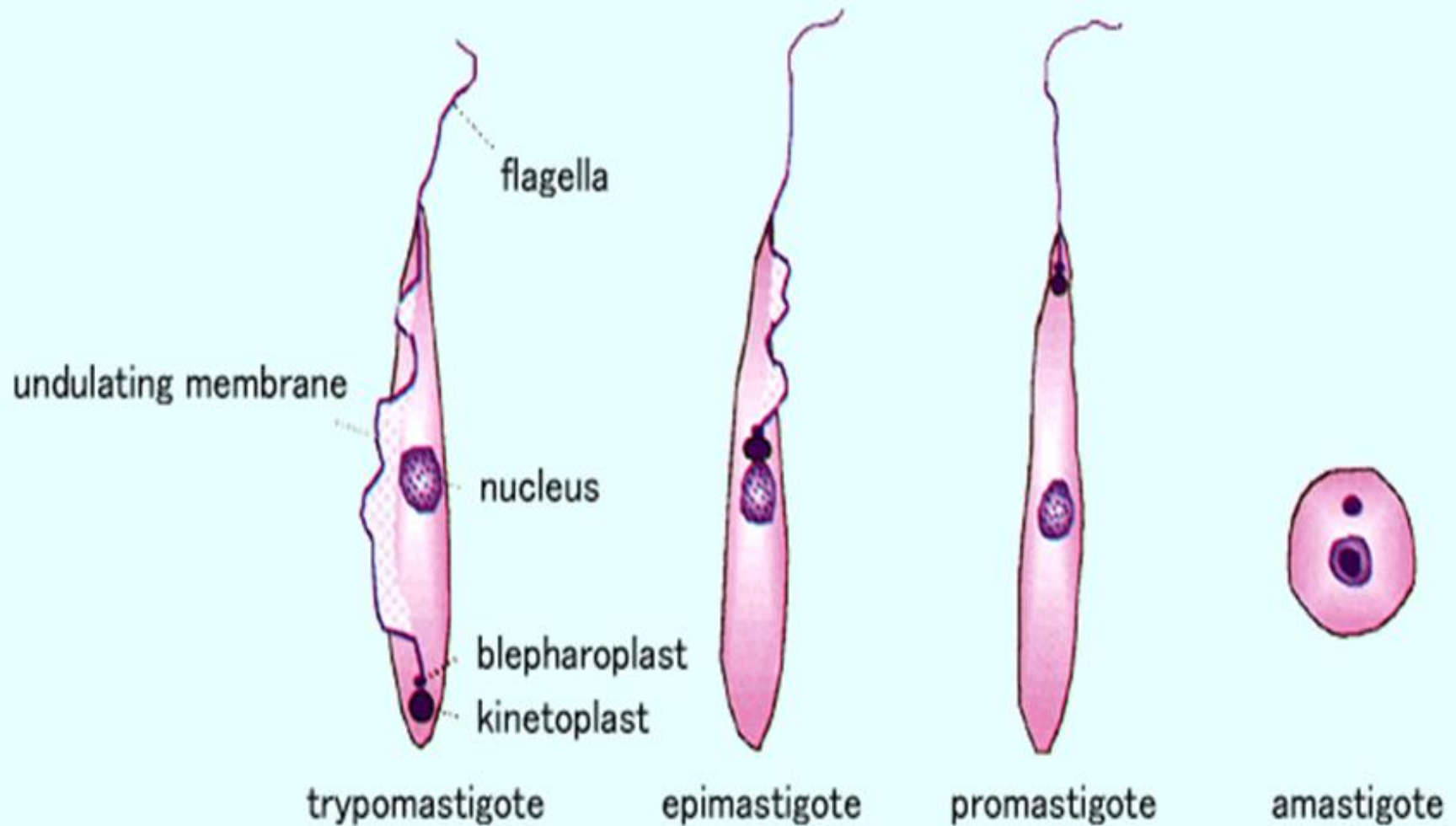
Haemoflagellates

Leishmania

Dr MONA BADR

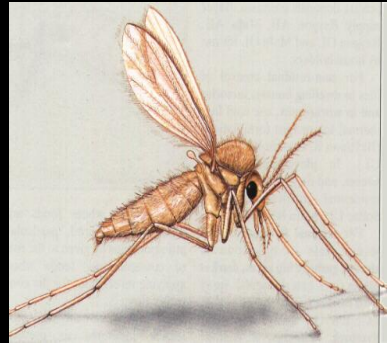


Different stages of Haemoflagellates



Lishmaniasis

- Lishmaniasis is a disease caused by parasites name Lishmania, it is spread by the bite of certain type of SANDFLIES.



- The disease can present in three main ways :
 - 1-Cutaneous
 - 2-Mucocutaneous
 - 3-Viseral (Kala-azar) most serious form of the disease can be fatal if not treated.

Leishmania Parasites and Diseases

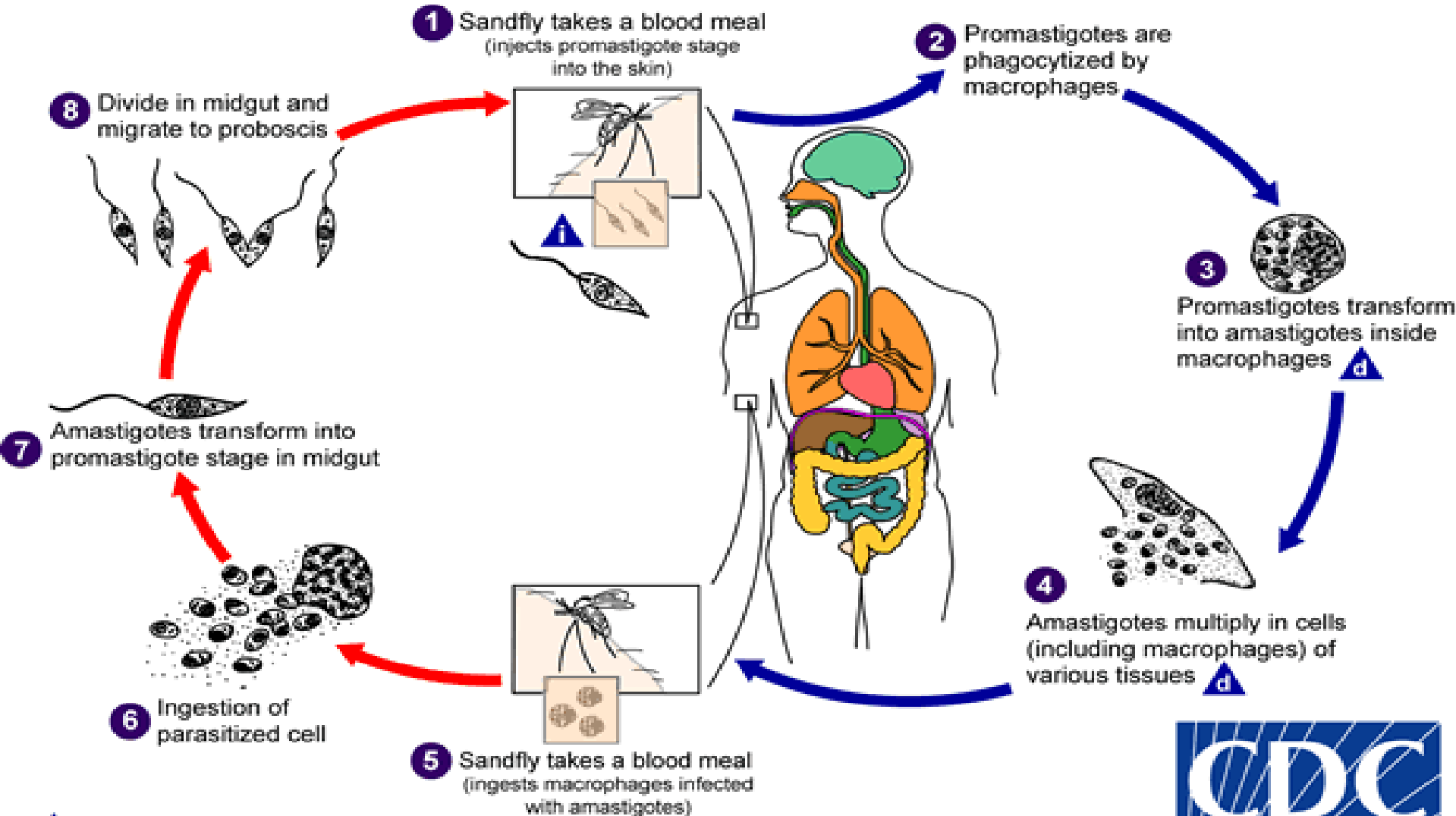
SPECIES	Disease
<i>Leishmania tropica</i> * <i>Leishmania major</i> * <i>Leishmania aethiopica</i> <i>Leishmania mexicana</i>	Cutaneous leishmaniasis
<i>Leishmania braziliensis</i>	Mucocutaneous leishmaniasis
<i>Leishmania donovani</i> * <i>Leishmania infantum</i> * <i>Leishmania chagasi</i>	Visceral leishmaniasis

* Endemic in Saudi Arabia

The life cycle of *Leishmania*

Sandfly Stages

Human Stages



i = Infective Stage

d = Diagnostic Stage

Clinical types of cutaneous leishmaniasis



■ *Leishmania major*: Zoonotic cutaneous leishmaniasis: wet lesions with severe reaction

■ *Leishmania tropica*: (human to human) cutaneous leishmaniasis: Dry lesions with minimal ulceration

■ Oriental sore (most common) classical self-limited ulcer



CUTANEOUS LISHMANIASIS THE COMMON TYPE

This starts as a painless papule on exposed parts of the body ,generally the face.

The lesion ulcerates after a few months producing an ulcer with an indurate margin.

1-

1-dry-type-lesion

In some cases the ulcer remains dry and heals readily (Lishmania Tropica , Oriental sore

2- wet-type-lesion

In some other cases the ulcer may spread with an inflammatory zone around , these known to heal slowly (Lishmania major)





UNCOMMON TYPES OF CUTANEOUS

LISHMANIASIS

- Diffuse cutaneous leishmaniasis (DCL):

Caused by *L. aethiopica*, diffuse nodular non-ulcerating lesions, seen in a part of Africa, people with low immunity to *Leishmania* antigens. Diffuse cutaneous (DCL), and consists of nodules and a thickening of the skin, generally without any ulceration, it needs numerous parasite.



- Leishmaniasis recidiva (lupoid leishmaniasis):

Severe immunological reaction to *leishmania* antigen leading to persistent dry skin lesions, few parasites.



Mucocutaneous Leishmaniasis

Leishmania braziliensis

The lesion starts as a pustular swelling in the mouth or on the nostrils. The lesion may become ulcerative after many months and then extend into the naso-pharyngeal mucous membrane.

Secondary infection is very common with destruction of the nasal cartilage and the facial bone. |



cutaneous & muco-cutaneous leishmaniasis

Diagnosis:

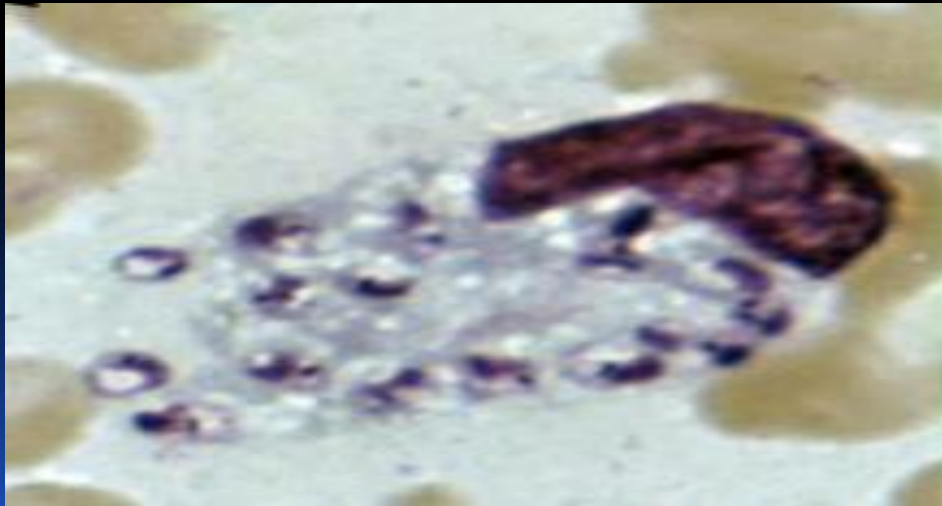
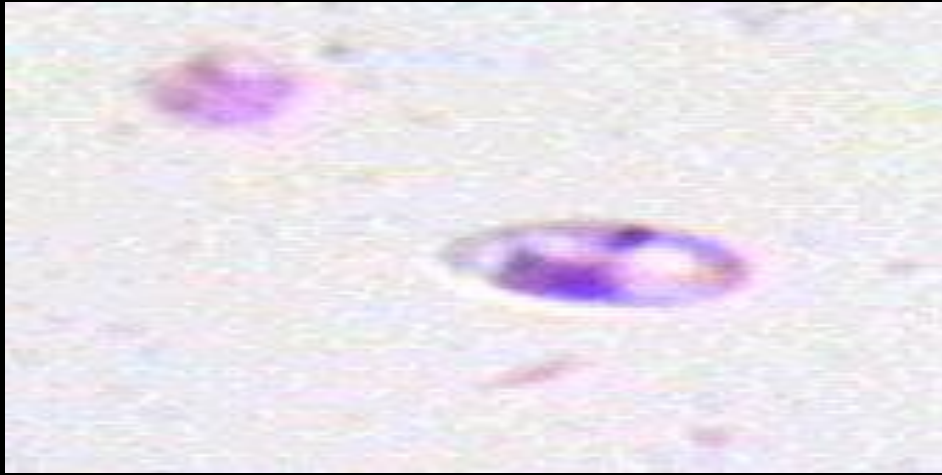
The parasite can be isolated from the margin of the ulcer.

Smear: Giemsa stain – microscopy for LD bodies (amastigotes) in tissue macrophages.

- **Biopsy:** microscopy for LD bodies or culture in **NNN** medium for promastigotes.

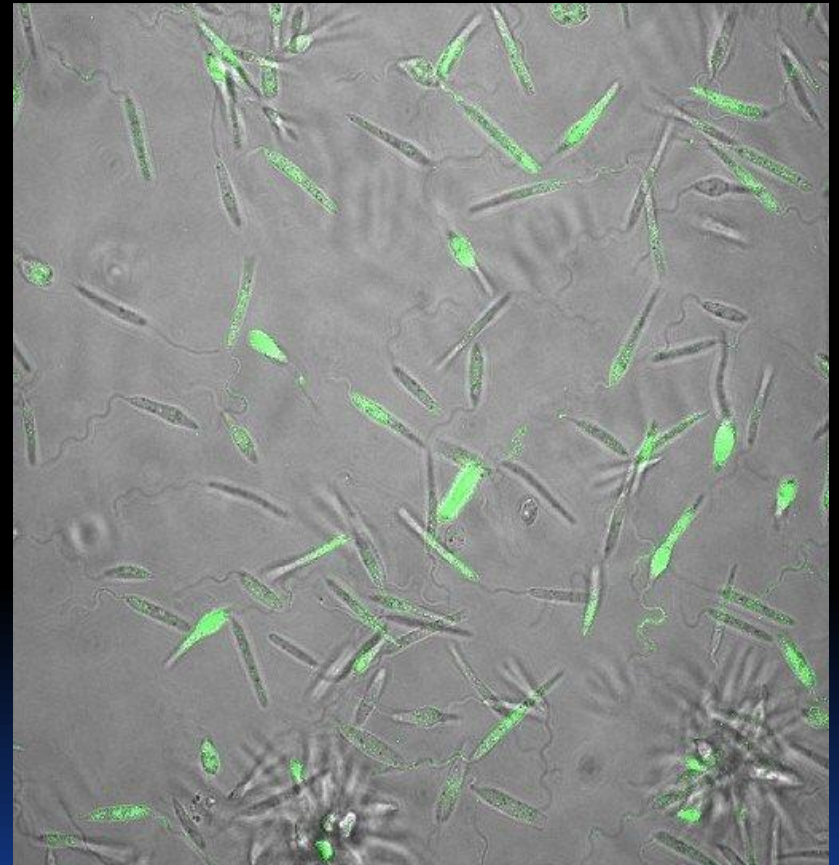
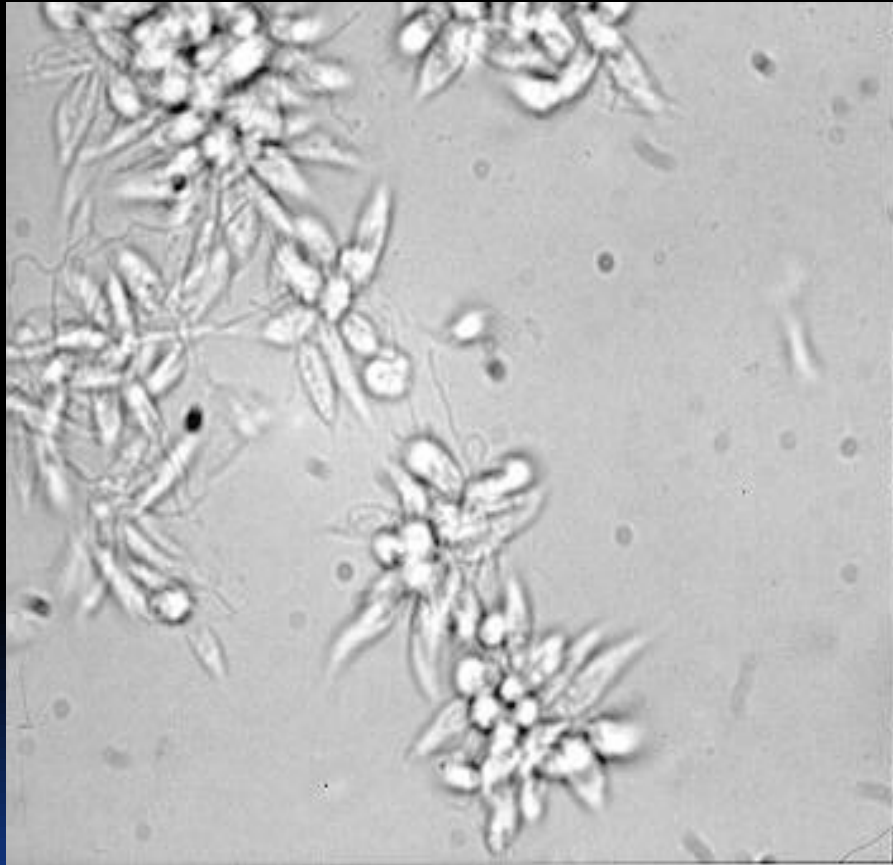
polymerase chain reaction (PCR) tests are available for the detection of *Leishmania* DNA

Amastigotes of Leishmania



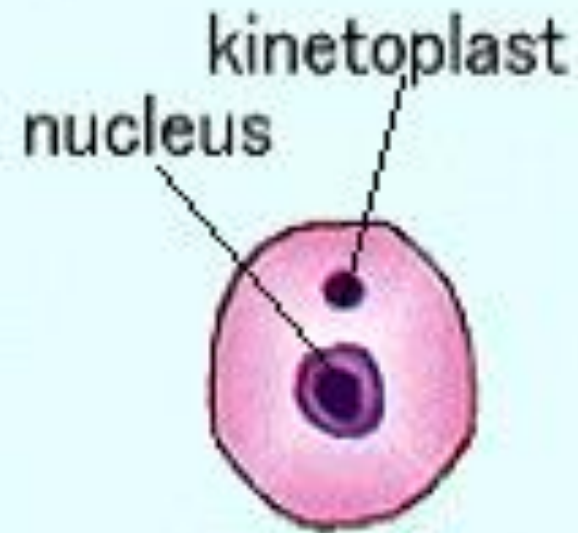
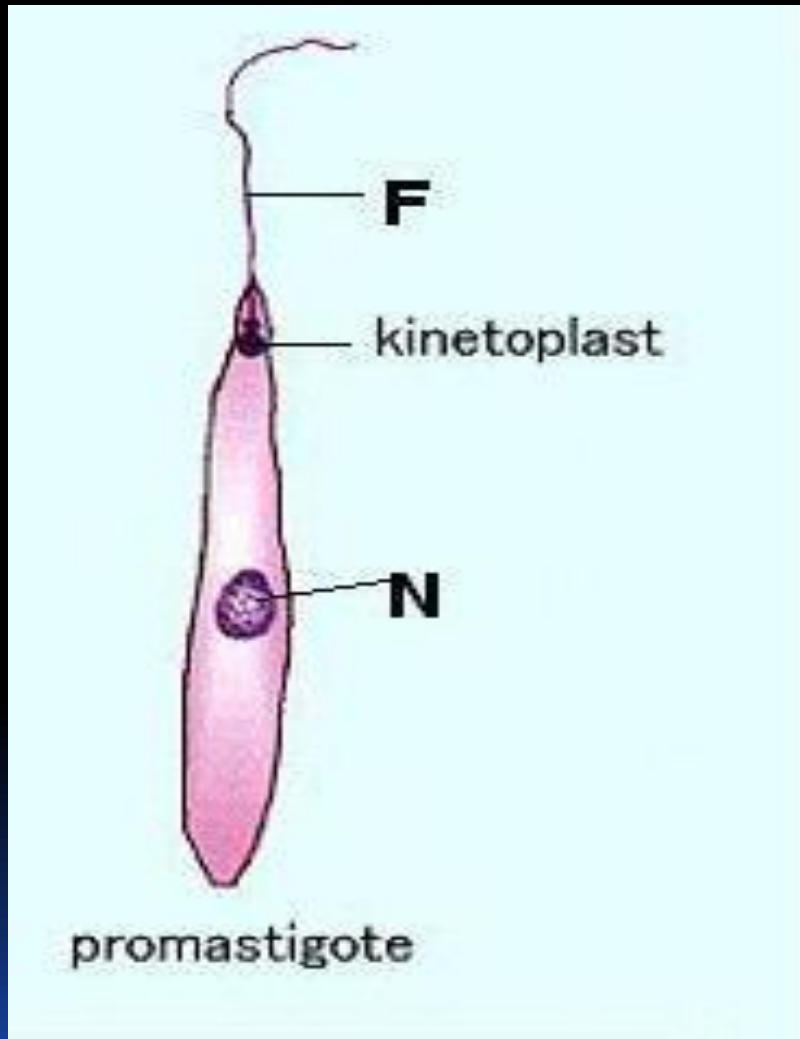
NNN medium





Promastigotes of Leishmania

Promastigotes of Leishmania



Amastigote of Leishmania

Treatment

- No treatment – self-healing lesions
- Medical:
 - **Pentavalent antimony (Pentostam),**
 - Antifungal drugs
 - +/- Antibiotics for secondary bacterial infection.
- Surgical:
 - Cryosurgery
 - Excision
 - Curettage

REFERENCE :WHO (2010) Control of leishmaniasis. Report of a meeting of an expert committee on the control of leishmaniasis.
http://whqlibdoc.who.int/trs/WHO_TRS_949_eng.pdf



Visceral leishmaniasis

- Is the most serious form, and is potentially fatal if untreated, There are geographical variations.
- The disease is called **kala-azar**
- ***Leishmania infantum* mainly affect children**
- ***Leishmania donovani* mainly affects adults**
- The incubation period is usually 4-10 months.
- The early symptoms are generally low grade fever with malaise and sweating and anemia.
- In later stages, the fever becomes intermittent and then liver enlargement, spleen enlargement or hepatosplenomegally because of the hyperplasia of the lymphoid –macrophage system and bone marrow.

Hepatosplenomegaly in visceral leishmaniasis



Untreated disease can be fatal

After recovery it might produce a condition called post kala-azar dermal leishmaniasis (PKDL)



Visceral leishmaniasis

Diagnosis

(1) Parasitological diagnosis:

Bone marrow aspirate

Splenic aspirate

Lymph node

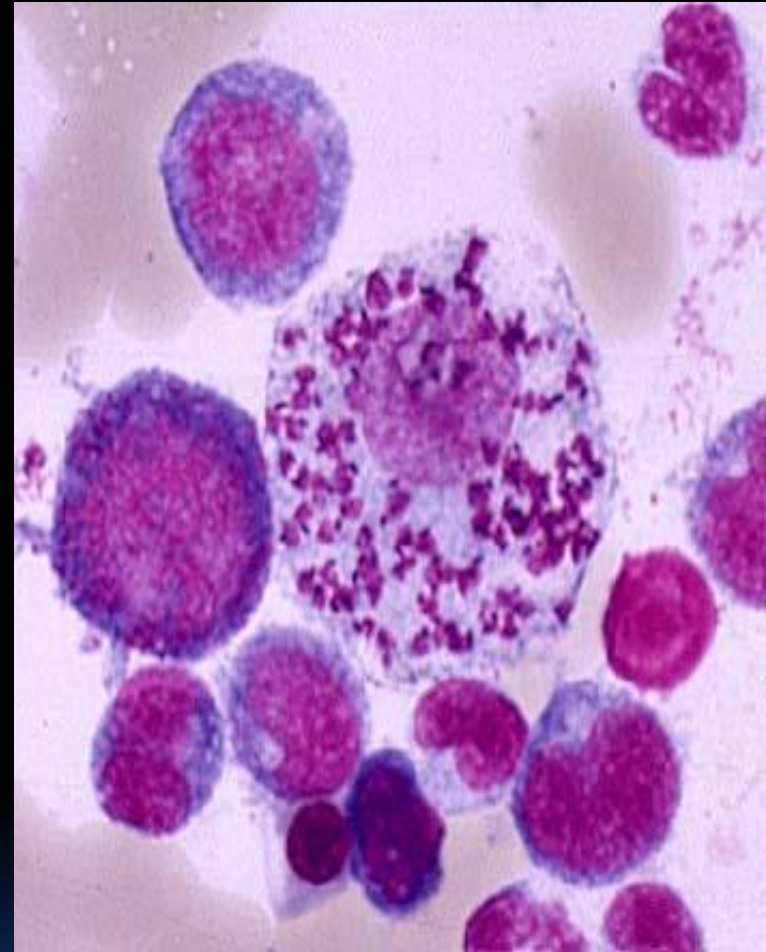
Liver biopsy

1. microscopy
2. culture in NNN medium

Diagnosis of Lishmaniasis

- Leishmaniasis is diagnosed in the hematology laboratory by direct visualization of the amastigotes (Leishman-Donovan bodies). Buffy-coat preparations of peripheral blood or aspirates from marrow, spleen, lymph nodes, or skin lesions should be spread on a slide to make a thin smear and stained with **Leishman or Giemsa stain** . Amastigotes are seen within blood macrophage and spleen monocytes or, less commonly, in circulating neutrophils .

Bone marrow aspiration



Bone marrow amastigotes

(2) Immunological Diagnosis:

- Specific serologic tests: Direct Agglutination Test (DAT), ELISA, IFAT
- Skin test (leishmanin test) for survey of populations and follow-up after treatment.

(3) *polymerase chain reaction* (PCR) tests are available for the detection of *Leishmania* DNA

Treatment of visceral leishmaniasis

- Recommended treatment varies in different endemic areas:
 - Pentavalent antimony- sodium stibogluconate (Pentostam)
 - Amphotericin B

Treatment of complications:

- Anaemia
- Bleeding
- Infections etc.

REFERENCE :WHO (2010) Control of leishmaniasis. Report of a meeting 571 of the WHO expert committee on the control of leishmaniasis. http://whqlibdoc.who.int/trs/WHO_TRS_949_eng.pdf