

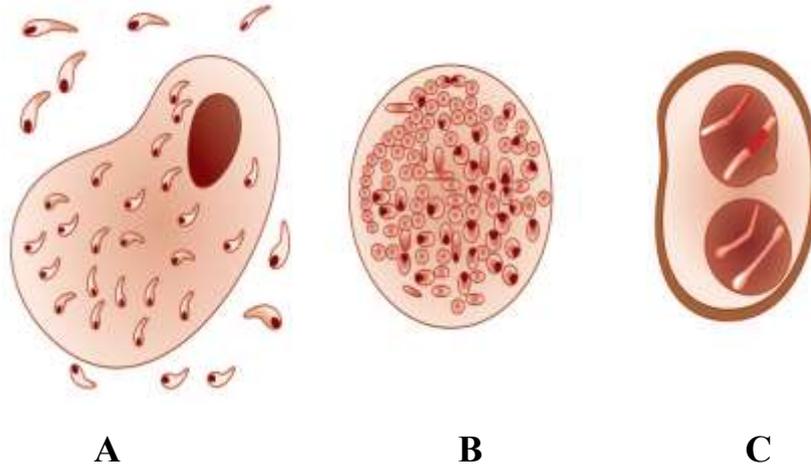


***Toxoplasma gondii* (Extra-intestinal coccidian)**

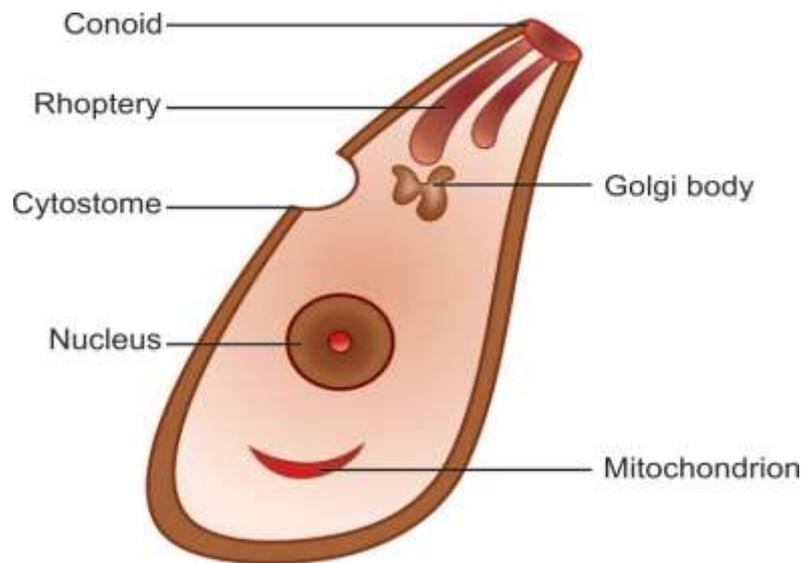
Morphology

T. gondii occurs in 3 forms

- **Trophozoite**
- **Tissue cyst**
- **Oocyst.**
- The trophozoite and tissue cyst represent stages in asexual multiplication (**schizogony**), while the oocyst is formed by sexual reproduction (**gametogony or sporogony**).
- All 3 forms occur in domestic cats and other felines, which are the definitive hosts and support both schizogony and gametogony.
- Only the asexual forms, trophozoites and tissue cysts are present in other animals, including humans and birds, which are the intermediate hosts.
- All the 3 forms are infectious to man. Trophozoites (Tachyzoites).
The trophozoite is crescent shaped, with one end pointed and the other end rounded.
- It measures 3–7 μm in length. The nucleus is ovoid and is situated at the blunt end نهاية حادة of the parasite.
- Electron microscopy reveals an **apical complex** at the pointed end.



Toxoplasma gondii: **A.** Smear from peritoneal fluid of infected mouse, showing crescentic tachyzoites: extracellular trophozoites and intracellular form within macrophage; **B.** Thick walled tissue cyst containing rounded forms bradyzoites; **C.** Oocyst containing 2 sporocysts with sporozoites inside



***Toxoplasma gondii*: Trophozoite (tachyzoite), fine structure seen by electron microscopy**

- The trophozoite stains well with Giemsa stain, the cytoplasm appearing **azure blue** and the nucleus, **red**.
- The actively multiplying trophozoite is seen intracellularly in various tissues during early acute phase of infection. Extracellular trophozoites can also be seen in impression smears.
- It can invade any nucleated cell and replicate within cytoplasmic vacuoles by a process called **endogony (internal budding)**, wherein 2 daughter trophozoites are formed, each surrounded by a membrane, while still within the parent cell. When the host cell becomes distended with the parasite, it disintegrates, releasing the trophozoites that infect other cells.
- During acute infection, the proliferating trophozoites within host cell may appear rounded and enclosed by host cell membrane. This is, called **pseudocyst** or **colony** and can be differentiated from tissue cysts by staining reactions.
- The rapidly proliferating trophozoites in acute infection are called **tachyzoites**
- The trophozoites are susceptible to drying, freeze thawing, and gastric digestion.

Tissue cyst

Tissue cysts are the resting form of the parasite.

- They are found during chronic stage of the infection and can be found in the brain (most common site), skeletal muscles, and various other organs.
- The cyst wall is eosinophilic and stains with silver, in contrast to the pseudocyst.
- With periodic acid Schiff (PAS) stain, the cyst wall stains weakly, and the parasites inside are stained deeply. The slowly multiplying parasites within the cyst are called **bradyzoites**.

- The cyst is round or oval, 10–20 μm in size and contains numerous bradyzoites. Cysts remain viable in tissue for several years.
- In immunologically normal hosts, the cysts remain silent, but in the immunodeficient subjects, they may get reactivated, leading to clinical disease.
- It is relatively resistant and when the raw or undercooked meat containing the cysts is eaten, infection occurs.
- The cyst wall is disrupted by peptic or tryptic digestion and the released parasites initiate infection by invading intestinal epithelial cells.
- They reach various tissues and organs through blood and lymphatic dissemination.
- Cysts are susceptible to desiccation, freezing, and thawing, and heat above 60°C.

Oocyst

Oocysts develop only in intestine of the definitive hosts .

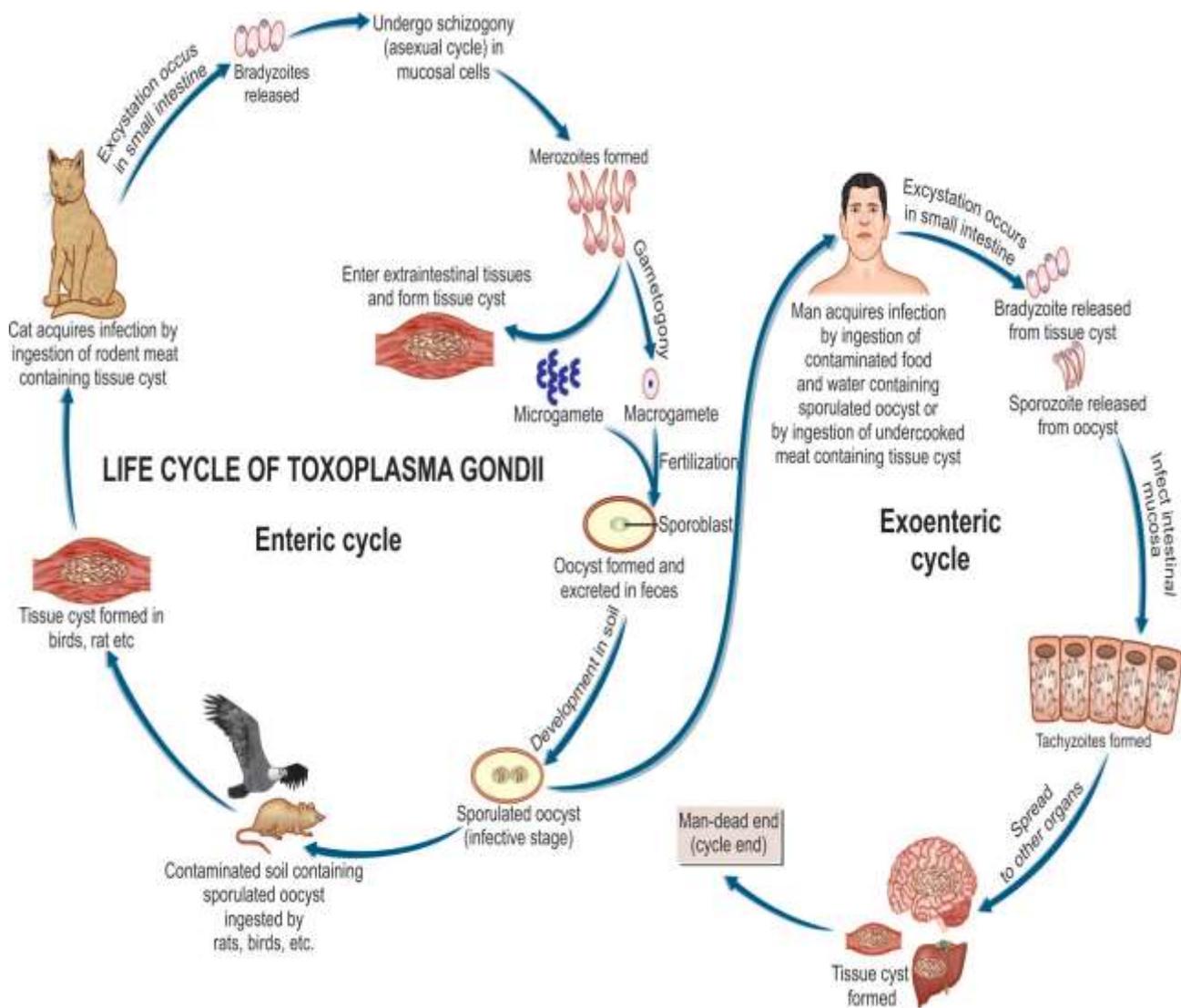
It is oval in shape and measures 10–12 μm in diameter. Each cyst is surrounded by a thick resistant wall.

- The oocystis formed by sexual reproduction (gametogony).
- Cats shed millions of oocysts per day in feces for about 2 weeks during the primary infection. The freshly passed oocyst is not infectious.
- They undergo sporulation in the soil with formation of 2 sporocysts, each containing 4 sporozoites. The sporulated oocyst is infective.
- Oocyst is very resistant to environmental conditions and can remain infective in soil for about a year.
- When the infective oocyst is ingested, it releases sporozoites in the intestine, which initiates infection.

Life Cycle

T. gondii completes its life cycle in 2 hosts.

- **Definitive host:** Cats and other felines, in which both sexual and asexual cycle takes place.
- **Intermediate hosts:** Man and other mammals, in which only the asexual cycle takes place.
- *T. gondii* has 2 types of life cycles:
 1. Enteric cycle
 2. Exoenteric cycle.



Life cycle of *Toxoplasma gondii*.

Enteric cycle

Enteric cycle occurs in cat and other definitive hosts.

- Both sexual reproduction (gametogony) and asexual reproduction (schizogony) occur within the mucosal epithelial cells of the small intestine of the cat.
- Cat acquires infection by ingestion of tissue cysts in the meat of rats and other animals or by ingestion of oocysts passed in its feces.
- The bradyzoites are released in the small intestine and they undergo asexual multiplication (schizogony) leading to formation of merozoites.
- Some merozoites enter extraintestinal tissues resulting in the formation of tissue cysts in other organs of the body.
- Other merozoites transform into male and female gametocytes and sexual cycle (gametogony) begins, with the formation of **microgamete** and **macrogamete**.

- A macrogamete is fertilized by motile microgamete resulting in the formation of an oocyst, which passes through maturation stages (**sporulation**) in the soil after being excreted from host through feces.
- A mature oocyst containing 8 sporozoites is the infective form which may be ingested by rats or other mammals to repeat the cycle

Exoenteric cycle

Exoenteric cycle occurs in humans, mice, rats, sheep, cattle, pigs and bird , which are the intermediate hosts. Humans acquire infection after:

- Eating uncooked or undercooked infected meat, particularly lamb and pork containing tissue cysts.
- Ingestion of mature oocysts through food, water, or fingers contaminated with cat feces directly or indirectly.
- Intrauterine infection from mother to fetus (**congenital toxoplasmosis**)
- Blood transfusion or transplantation from infected donors.
- Sporozoites from the oocysts and bradyzoites from the tissue cysts enter into the intestinal **mucosa** and multiply **asexually** and **tachyzoites** are formed (**endodyogeny**).
- Tachyzoites continue to multiply and spread locally by lymphatic system and blood.
- Some tachyzoites also spread to distant extraintestinal organs like brain, eye, liver, spleen, lung, and skeletal muscles and form **tissue cysts**. The slowly multiplying forms inside the tissue cysts are known as **bradyzoites**, which remain viable for years.
- The dormant bradyzoites inside the cyst may be reactivated in immune suppression causing renewed infection in the host.
- Human infection is a dead end for the parasite.
- ❖ Human toxoplasmosis is a zoonosis.
- ❖ The full natural cycle is maintained predominantly by cats and mice.
- ❖ Mice eat materials contaminated with oocysts shed in cats feces. Tissue cysts develop in mice.
- ❖ When such mice are eaten by cats, they get infected and again shed oocysts in feces.

Pathogenicity and Clinical Features

The outcome of *Toxoplasma* infection depends on the immune status of the infected person.

- Active progression of infection is more likely in immunocompromised individuals. Toxoplasmosis has acquired great importance as one of the major fatal complications in acquired immunodeficiency syndrome (AIDS).
- Most human infections are asymptomatic.
- Clinical toxoplasmosis may be congenital or acquired.

Congenital toxoplasmosis

Congenital toxoplasmosis results when *T. gondii* is transmitted transplacentally from mother to fetus.

Parasite which can be transmitted from mother to fetus:

Toxoplasma gondii

Plasmodium spp.

Trypanosoma cruzi

- This occurs when the mother gets primary *Toxoplasma* infection, whether clinical or asymptomatic, during the pregnancy.
- The risk of fetal infection rises with progress of gestation; from 25%, when the mother acquires primary infection in first trimester to 65% in the third trimester. Conversely, the severity of fetal damage is highest, when infection is transmitted in early pregnancy.
- Mothers with chronic or latent *Toxoplasma* infection, acquired earlier, do not ordinarily infect their babies. But in some women with latent or chronic infection, the tissue cyst may be reactivated during pregnancy and liberate trophozoites, which may infect the fetus *in utero*
- Most infected newborns are asymptomatic at birth and may remain so throughout. Some develop clinical manifestations of toxoplasmosis weeks, months, and even years after birth.
- The manifestations of congenital toxoplasmosis include chorioretinitis التهاب المشيمة, cerebral calcifications التكتلسات الدماغية, convulsions تشنجات, strabismus التخلف العقلي, deafness الصم, blindness العمى, mental retardation التخلف العقلي, microcephaly صغر الرأس, and hydrocephalus استسقاء الرأس.
- A few children are born with manifestations of acute toxoplasmosis, which may include fever, jaundice, petechial rashes, microphthalmia, cataract, glaucoma, chorioretinitis, lymphadenopathy, hepatosplenomegaly, myocarditis, cerebral calcifications, and chorioretinitis.

Acquired Toxoplasmosis

- Infection acquired postnatally بعد الولادة is mostly asymptomatic.
- The most common manifestation of acute acquired toxoplasmosis is lymphadenopathy اعتلال العقد اللمفاوية; the cervical lymph nodes being most frequently affected.
- Fever, headache, myalgia الألم العضلي, and splenomegaly are often present. The illness may resemble mild flu and is self-limited, although the lymphadenopathy may persist.
- In some cases, there may be a typhus-like exanthema طفح يشبه التيفوس with pneumonitis التهاب رئوي, myocarditis التهاب عضلة القلب, and meningoencephalitis التهاب السحايا والدماغ, which may be fatal.

Ocular Toxoplasmosis

Another type of toxoplasmosis is ocular.

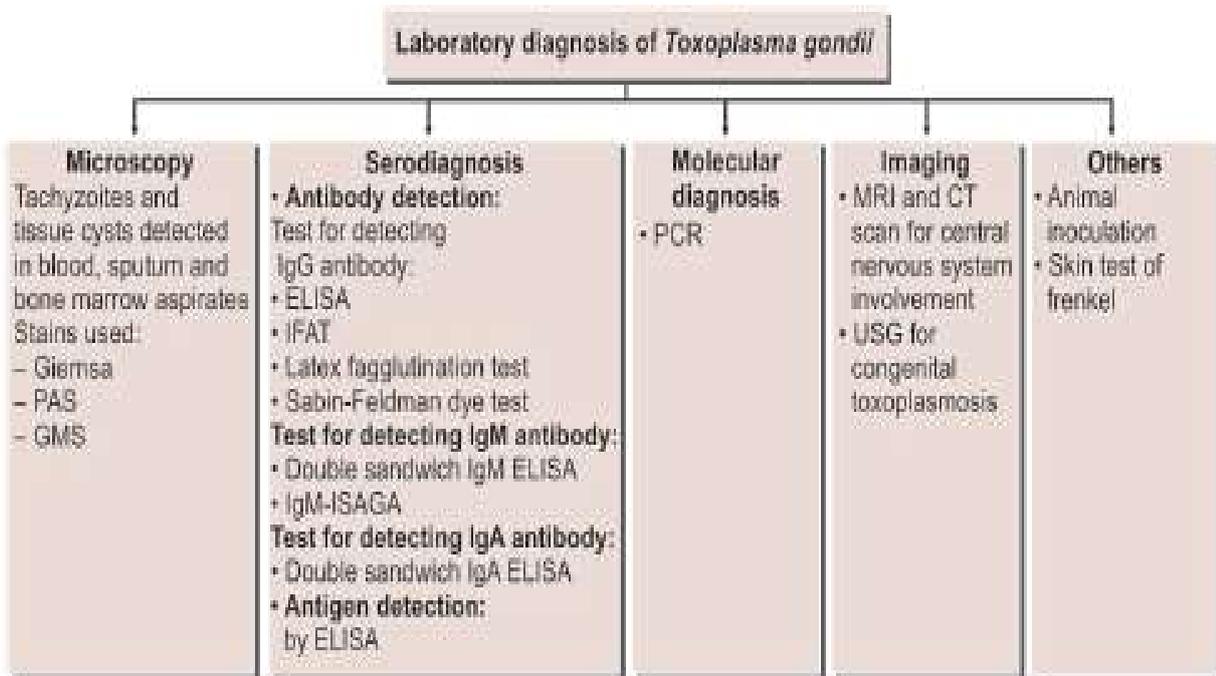
- It may present as uveitis, choroiditis, or chorioretinitis.
- Some cases may be so severe that they require enucleation. Toxoplasmosis *in* Immunocompromised Patients.
- Toxoplasmosis is most serious and often fatal in immunocompromised patients, particularly in AIDS, whether it may be due to reactivation of latent infection or new acquisition of infections.

- In these patients, involvement of brain is most common.
- Clinical manifestation include encephalitis, altered mental state, seizures, cerebellar signs, meningismus, and neuropsychiatric manifestations.
- Besides central nervous system involvement, other organs involved are lungs, pancreas, gastrointestinal tract, eyes, heart, and liver.
- *Toxoplasma* pneumonia can be confused with pneumocystis pneumonia.

Host Immunity

Host defense against *Toxoplasma* infection involves both humoral (antibody-mediated) and cellular responses. Specific IgG antibody can lyse extracellular trophozoites, but activated T cells and natural killer cells appear to be more important in containing the infection and preventing clinical disease.

Laboratory Diagnosis



-Presence of only tissue cysts does not differentiate between active and chronic infection.

-The presence of cysts in placenta or tissues of newborn establishes congenital *Toxoplasma* infection.

Antibody detection

Diagnosis of acute infection with *T. gondii* can be made by detection of the simultaneous presence of IgM and IgG antibodies.

Tests for detecting IgG antibody include:

1. Enzyme-linked immunosorbent assay (ELISA)
2. Indirect fluorescent antibody test (IFAT)
3. Latex agglutination test
4. Sabin-Feldman dye test.

-Positive IgG titer ($>1:10$) can be detected as early as 2–3 weeks after infection. Peak level of antibody is observed in blood 4–8 weeks after infection.

-A positive IgM antibody titer indicates an early primary infection. The serum IgM titer can be measured by double sandwich IgM ELISA or IgM immunosorbent assay (IgM-ISAGA). Both assays are equally specific and sensitive. Negative IgM titer and positive IgG titer indicate distant infection.

-The double sandwich IgA-ELISA test is used for detecting congenital infection in newborns.

Antigen detection

Detection of antigen by ELISA indicates recent *Toxoplasma* infection.

-In AIDS and other immunocompromised patients, antigen detection is very useful.

-Detection of antigen in amniotic fluid is helpful to diagnose congenital toxoplasmosis.

Immunocompromised Patients

AIDS patients who are seropositive for *T. gondii* and have a CD4+ T lymphocyte count below $<100/\mu\text{L}$, should receive primary prophylaxis against *Toxoplasma* encephalitis.

*Trimethoprim-sulfamethoxazole is the drug of choice. If this drug cannot be tolerated by patients, dapsone-pyrimethamine is the recommended alternative drug of choice.

* Prophylaxis against toxoplasma encephalitis should be discontinued in patients who have responded to antiretroviral therapy (ART) and whose CD4+ T lymphocyte count has been above $200/\mu\text{L}$ for 3 months.

Notes

- ❖ Individuals at risk, particularly pregnant women, children, and immunocompromised persons should avoid contact with cat and its feces.
- ❖ Proper cooking of meal.
- ❖ Proper washing of hands and washing of vegetables and fruits before eating.
- ❖ Blood or blood products from seropositive persons should not be given and screening for *T. gondii* antibody should be done in all blood banks.

Control

It is difficult to control toxoplasmosis because of wide range of animal reservoirs. Currently, there is no effective vaccine available for humans. A genetically engineered vaccine is under development for use in cats.