General Information
- *Toxoplasma gondii* is a protozoal parasite capable of infecting any warm-blooded animal, including humans.
- Wild and domestic cats are the only known definitive hosts of *Toxoplasma*; they can develop both systemic and patent intestinal infection. All other animals and humans serve as intermediate hosts in which the parasite may cause only systemic infection, which typically results in the formation of tissue cysts.
- In all species, *Toxoplasma* infection is usually subclinical, although it may occasionally cause mild, non-specific signs. Infection may have much more serious consequences in immunocompromised or pregnant animals and people.
- The major modes of transmission include consumption of undercooked meat containing *Toxoplasma* cysts, fecal-oral transfer of *Toxoplasma* oocysts from cat feces (either directly or in contaminated food, water or soil), and vertical transmission from mother to fetus if primary infection occurs during pregnancy.
- The risk of contracting *Toxoplasma* infection from cleaning the litter box of a house cat is actually very small, especially if a few simple precautions such as appropriate hand washing are observed.

Prevalence of *Toxoplasma*
*Toxoplasma* is one of the most widespread zoonotic pathogens in the world. In most animals and people, primary infection results in a detectable antibody titre for the life of the host, therefore seroprevalence (i.e. previous exposure to the parasite but not necessarily clinical disease) increases with age.

Humans
- Because toxoplasmosis is not a reportable disease in Ontario and most of North America, it is difficult to estimate the prevalence of infection in animals or people. An average 15 000 cases of clinical toxoplasmosis are reported annually in the USA, but it has been estimated that the actual number of cases is likely closer to 225 000. It is estimated that 50% of these cases occur due to foodborne transmission of *Toxoplasma*.
- Seroprevalence in the general population in persons over 12 years of age was estimated to be 22.5% in one North American study performed from 1988-1994. Worldwide, seroprevalence ranges from 0-100% depending on country, geographic area and even ethnic group.
- Between 40 to 400 children born in Canada each year are infected with *Toxoplasma* before birth.

Animals
- The prevalence of oocyst shedding in cats is very low (0-1%), even though serological studies indicate that at least 15-40% of cats have been infected, depending on how the cats are fed and whether they go outdoors.
- Disease and exposure are more common in cats and other pets that go outdoors, hunt, or are fed raw meat.

Zoonotic Risk & Other Risk Factors
- The risk of transmission of *Toxoplasma* from a household cat can be easily controlled by means of simple infectious disease control procedures (see Infection Control below). In one study from Norway, cleaning a cat litter box was found to be a strong risk factor for exposure to *Toxoplasma*. However, the same study, and another European study, showed no association between *Toxoplasma* exposure and living with or near a cat.
- Contamination of water sources and soil with the feces of wild or domestic cats is more difficult to control, and can lead to infection following ingestion of oocysts on unwashed, uncooked vegetables or in contaminated water. Cockroaches and coprophagic flies may also serve as mechanical vectors of *Toxoplasma*, resulting in contamination of food, water or utensils with infectious oocysts.
- Contact with contaminated soil or sand, such as in a garden or a sandbox, is also be associated with *Toxoplasma* infection.
- Consumption of undercooked meat is one of the principle risk factors for *Toxoplasma* infection, and people who handle raw meat (such as abattoir workers) may also be more commonly exposed to the parasite.
- The importance of these various risk factors likely varies considerably between ethnic groups due to differences in cultural habits regarding exposure to undercooked meat, soil and cats.

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Life Cycle of *Toxoplasma*

Unsporulated oocysts of *Toxoplasma* are passed in the feces of acutely infected cats. The oocysts usually sporulate in 1-5 days forming two sporocysts, each containing four infective sporozoites. When the oocysts are ingested by any warm-blooded host, the sporozoites excyst, invade intestinal cells and begin to divide asexually to produce tachyzoites. These then migrate throughout the body, invading tissue cells and multiplying until the cells rupture. Eventually the tachyzoites encyst, becoming bradyzoites, within cells of the central nervous system, muscle, and sometimes other organs. The cysts typically persist until death of the host without causing clinical signs. If the host is eaten by another animal, the bradyzoites excyst in the intestine and the process is repeated, forming new tissue cysts.

If an intermediate host is eaten by a cat, the bradyzoites invade the intestinal epithelial cells and undergo five stages of asexual reproduction (merogony) followed by formation of microgamonts and macrogamonts. The microgamonts divide to form flagellated microgametes, which then fertilize the macrogamonts. The fertilized macrogamont forms a wall and becomes an unsporulated oocyst approximately 10 μm x 12 μm in size, which is passed in the feces of the cat. When fed tissue cysts, 97% of cats infected for the first time will produce oocysts, usually within 3-10 days. They may shed for up to 20 days, but the majority of oocysts will be shed in just 1-2 days. Only 20% of cats fed oocysts will develop a patent infection, and the prepatent period may be 18 days or more. Contrary to previous beliefs, studies have shown that oocysts can be shed in low numbers by previously infected cats that are challenged again with the parasite or that become immunosuppressed due to disease or drug therapy.

Transmission of *Toxoplasma*

Carnivorous animals are often infected with *Toxoplasma* through ingestion of bradyzoites from tissue cysts in infected prey, as are people who eat undercooked meat, particularly that of pigs, sheep and goats. *Toxoplasma* cysts are less commonly found in poultry and rarely found in beef. The prevalence of *Toxoplasma* infection in commercial farm animals has decreased significantly with the advent of intensive management practices. Free-range poultry, swine and small ruminants, marsupials (e.g. kangaroos) and some wild game are more likely to harbour cysts.

Oocysts are only shed by cats. **Unsporulated oocysts in fresh feces are not infective**; they need appropriate oxygen, humidity and temperature to sporulate. Sporulated oocysts are the most environmentally resistant life stage of the parasite. Ingestion of as few as ten oocysts may infect an intermediate host, while ingestion of 100 or more oocysts can cause a patent infection in a cat, which may shed tens to hundreds of millions of oocysts. Sporulated oocysts from cat feces can survive passage through the intestine of a dog.

Tachyzoites are potentially infective, and may be found in the tissues of acutely infected animals, as well as the milk of sheep, goats, cows, and sometimes chicken eggs. However, tachyzoites are killed relatively easily by pasteurization, and uncommonly survive gastric digestion, although this may be more of a concern in infants who have lower concentrations of peptic enzymes. Any kind of cooking will kill tachyzoites in an egg. *Toxoplasma* can also be transmitted by organ transplants and blood transfusions, but this is uncommon.

**In utero transmission of *Toxoplasma*** occurs only if primary infection of the dam/mother occurs during pregnancy. Parasitemia then results in placentitis and infection of the fetus. This is more likely to occur in humans, sheep and goats, and sometimes mice, cats and dogs. Kittens can be infected transplacentally or through tachyzoites shed in the queen’s milk. Under normal circumstances, a female that has been exposed to *Toxoplasma* 4-6 months prior to pregnancy will develop sufficient immunity to protect herself and the fetus for the rest of her life. However, if the immune response is suppressed by drug therapy or disease such as HIV/AIDS in humans, both the mother and the fetus may become susceptible to infection again. In humans, the risk of the infection being passed on to the fetus increases from the first trimester (10-25%) to the third trimester (60-90%). However, the potential congenital defects are more severe with earlier infections.
Symptoms and Signs

Clinical signs of toxoplasmosis are caused by cellular destruction due to multiplying tachyzoites, which most commonly affect the brain, liver, lungs, skeletal muscle and eyes. Infection may be associated with other diseases such as distemper or ehrlichiosis in dogs, hemobartonellosis, feline immunodeficiency virus (FIV) infection in cats, or immunosuppressive therapy in any species.

Humans: Approximately 15% of cases of Toxoplasma infections are associated with clinical signs such as mild fever and lymphadenopathy. Signs may persist for 1 to 12 weeks; more severe disease is very rare in immunocompetent individuals. Of clinical cases, 0.2%-0.7% may develop ocular toxoplasmosis (retinitis), but this is more commonly associated with congenital infection. If more severe disease develops, signs may be related to encephalitis, hepatitis, myositis or pneumonia. Toxoplasma encephalitis develops at some time in approximately 40% of individuals with AIDS, and is fatal in 10-30% of these cases.

Approximately 10% of congenital Toxoplasma infections result in abortion or neonatal death. In 10-23% of congenital infections, signs are present at birth; these may include hydrocephalus, chorioretinitis, microcephaly and small size. Clinical signs are not apparent at first in 67-80% of cases. Ocular toxoplasmosis may occur in up to 1/3 of children that survive congenital infection.

Animals:

- In healthy, naïve cats, primary infection with Toxoplasma is typically either subclinical or may cause mild small-intestinal diarrhea for up to 10 days. In rare cases when cats develop significant clinical disease, signs may include fever, lethargy, anorexia, and others associated with pneumonia, hepatitis, myositis or encephalitis. Pancreatic, cardiac and ocular tissues are also commonly affected. Dermatitis and vomiting have also been reported. The onset of signs may be slow, or the disease may be rapidly fatal. Neurologic and ocular signs without systemic illness are more common with reactivated infections.
- Congenitally or lactationally infected kittens are often severely affected due to uncontrolled replication of tachyzoites and tissue damage. Kittens may be stillborn or die before weaning, typically with signs related to pneumonia, hepatitis or encephalitis. However, in some cases the only sign of disease may be chorioretinitis, and occasionally concurrent anterior uveitis.
- Clinical signs and tissue involvement are similar in dogs, but ocular lesions are much less common. Generalized toxoplasmosis typically occurs in dogs less than one year old. Clinical signs in older dogs are primarily associated with encephalitis and myositis, and appear very similar to Neospora caninum infection.

Clinical infection in sheep and goats is much more common than in dogs and cats, and is primarily associated with reproductive problems, including abortion and birth of weak young that are incoordinated and unable to feed themselves. The infection can also be common in birds, but it is rarely clinical.

Diagnosis

Once infected with Toxoplasma, people and animals usually develop a life-long protective antibody titre, unless the individual is severely immunocompromised and unable to mount or sustain an appropriate humoral immune response. The organism itself can be detected in tissues or body fluids using either PCR or bioassays in mice.

Infection in Humans: There are several different serological tests used to diagnose toxoplasmosis in humans that are intended to help differentiate between latent and acute infections. Serum IgM titers indicate recent infection, whereas serum IgG titers persist longer and therefore typically indicate previous infection. However, both types of antibodies are usually detectable within 1-2 weeks of infection. Some Toxoplasma IgM test kits have relatively high false-positive rates. Measurement of IgG avidity can also help “age” the antibody response. The modified latex agglutination test (MAT) detects IgG, but can help differentiate acute and chronic infections based on reactivity with acetone versus formalin-fixed antigen. It is considered extremely sensitive.

Serological screening of pregnant women is not generally recommended in the USA and Canada as it is in some European countries, because the risk of exposure to Toxoplasma is comparatively low. Diagnosis of in utero infection is most commonly accomplished by detecting Toxoplasma DNA in amniotic fluid using PCR.
Clinical Infection in Animals: Both leukopenia and leukocytosis may be observed in animals with clinical toxoplasmosis. Serum biochemistry abnormalities are related to the primary tissues involved (e.g. liver, muscle, pancreas). In acute infection, tachyzoites may be rarely seen in peripheral blood, cerebrospinal fluid (CSF), or lung or tracheal wash fluid, but are more commonly found in thoracic or abdominal effusions. Changes in the CSF of encephalitic cats are inconsistent, but may include increased protein and lymphocytic or mixed inflammatory cells.

A diagnosis of clinical toxoplasmosis is difficult to make antemortem in animals, and should be made based on a four-fold or greater change in antibody titre or a single high IgM titre, response to anti-Toxoplasma therapy, and exclusion of other differential diagnoses. Toxoplasma-specific antibody may also be detected in the aqueous humor or CSF of cats with ocular or neurologic signs, respectively, but the titre must be compared to that of a non-ocular or non-neurologic pathogen to determine if the antibody was produced locally.

Subclinical Infection in Cats: It is important to note that the development and persistence of IgM in infected cats is very inconsistent, and is not a reliable marker of acute Toxoplasma infection. Furthermore, some cats may not develop IgG titres until 4-6 weeks after infection, well after they have stopped shedding oocysts, and the titre may peak in as little as 2-3 weeks and remain high for years.

A cat that is IgG seropositive is unlikely to be shedding oocysts, and is unlikely to shed oocysts if it is exposed to the parasite. Nonetheless, exposure should be minimized as some seropositive cats may shed low numbers of oocysts if re-exposed. A cat that is seronegative is unlikely to be shedding oocysts, but is likely to develop a patent infection if it is exposed to Toxoplasma.

Oocyst Shedding in Cats: Cats typically only shed oocysts in their feces for 1-3 weeks following their first exposure to Toxoplasma, therefore fecal examination is usually unrewarding. If present, the oocysts are best detected using a centrifugal fecal floatation technique with Sheather’s sugar solution (specific gravity 1.26). The unsporulated oocysts have no distinct internal structures, and are approximately 10 μm in diameter (1/4 the size of Cystoisospora oocysts, and 1/8 the size of Toxocara cati eggs). They are indistinguishable from oocysts of some species of Hammondia and Besnoitia, which also occur in cats, but these oocysts should be considered Toxoplasma until proven otherwise.

Treatment of Toxoplasmosis Animals:
- Clindamycin (dogs 10-20 mg/kg, cats 10-12.5 mg/kg, PO or IM, q12h for 4 weeks) is the drug of choice for clinical toxoplasmosis in dogs and cats. This dose is higher than that on the product label. High-dose oral clindamycin therapy can cause anorexia, vomiting and diarrhea in dogs and cats. Improvement should be seen within 48 hours, but FIV-infected cats in particular may be more difficult to treat.
- Cats with ocular inflammation should also be treated with either topical (ideally) or systemic glucocorticoids.
- Pyrimethamine (dogs and cats 0.25-0.5 mg/kg, PO q12h for 4 weeks) and a sulfonamide (dogs and cats 30 mg/kg, PO q12h for 4 weeks) in combination are the second choice treatment. Cats must be monitored for signs of bone marrow suppression when this treatment regimen is used for more than 2 weeks. Newer macrolides (e.g. azithromycin, clarithromycin) may be effective against Toxoplasma, but are not licensed for use in animals. Doxycycline or minocycline may be used if there is concurrent infection with a different tetracycline-susceptible pathogen.
- Oocyst shedding in cats can be reduced by use of the therapeutic dose of clindamycin, or high-dose sulfonamide-pyrimethamine combinations (100 mg/kg-2.0 mg/kg, PO q24h for 1-2 weeks). Monensin (mixed in food as 0.02% w/w dry weight for 1-2 weeks) or toltrazuril (5-10 mg/kg PO q24h for 2 weeks) can also be used, if given within 1-2 days of exposure or administration of immunosuppressive therapy to the animal (which may cause recrudescence of infection). An oral vaccine to reduce oocyst shedding in cats has been developed, but is not currently commercially available.

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Infection Control

In the majority of studies, no direct association has been found between cat ownership and the risk of toxoplasmosis in people. Given the emotional benefits associated with owning a cat, and the minimal risk of transmission of Toxoplasma if appropriate hygiene is practiced, even pregnant or immunocompromised individuals do NOT need to give up their cats. Such individuals should avoid contact with cat feces and cat litter whenever possible by having someone else clean their cat’s litter box. If a cat is found to be shedding oocysts, it should be removed from the premises temporarily and treated to eliminate shedding. Because cats are usually meticulous groomers, it is unlikely that oocysts will be found on their fur, so regular handling is not a significant risk. **Individuals should always wash their hands thoroughly after contact with cat stool, litter or a litter box.**

Microwave cooking, salting and smoking do not consistently kill all infective Toxoplasma organisms. Freezing meat to −12°C for at least 24 hours will kill most Toxoplasma tissue cysts, but sporulated oocysts can survive at −20°C for up to 28 days. Washing kitchen utensils and surfaces that have come in contact with raw meat with soap and scalding hot water will kill any bradyzoites or tachyzoites present.

Oocysts take longer to sporulate under cooler conditions. At room temperature sporulation may occur within 1-5 days, but this can take 3 weeks at 11°C. Once sporulated, oocysts can survive even longer in the environment. They are resistant to most disinfectants, therefore immersion of litter boxes and other potentially contaminated instruments in boiling or scalding water is the preferred means of decontamination. Even sporulated oocysts are killed by heating to 55-60°C for 1-2 minutes. Cat feces should be disposed of daily to reduce the risk of transmission. Feces and dirty litter can be disposed of in a septic system if the litter is biodegradable, sealed tightly in a plastic bag and placed in the garbage, or incinerated. Backyard compost units do not produce sufficient heat to destroy oocysts and other pathogens potentially present in fecal material. Also keep cats out of sandboxes and other areas where children play that cats may be inclined to defecate.

Zoonotic Disease Risk

The zoonotic risk to the general population posed by Toxoplasma in cats is:

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Pregnant Women and Individuals with Compromised Immune Systems:

Education of individuals in these high risk groups about how to decrease the transmission of Toxoplasma is an important tool in the prevention of this disease. Recommendations for preventing toxoplasmosis include:

- Cook all meat to a minimum internal temperature of 67°C/153°F.
- Peel or thoroughly wash fruit and vegetables prior to consumption.
- Clean all surfaces and objects that come in contact with raw meat or unwashed fruit and vegetables.
- Avoid contact with cat litter and garden soil, otherwise wear gloves and wash hands thoroughly after.
- Avoid feeding raw meat to cats.
- Keep cats indoors so they do not become infected by eating small prey.

For these groups, the zoonotic risk posed by Toxoplasma in cats is likely:

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