Presence of *Toxoplasma gondii* tissue cysts in human semen: Toxoplasmosis as a potential sexually transmissible infection

Wen Han Tong, Jana Hlaváčová, Samira Abdulai-Saiku, Šárka Kaňková, Jaroslav Flegr, Ajai Vyas

*School of Biological Sciences, Nanyang Technological University, 60 Nanyang Drive, 637551, Singapore*

*Department of Philosophy and History of Science, Faculty of Science, Charles University, Viničná 7, CZ – 128 44 Prague, Czechia*

**Article Info**

**Objectives:** *Toxoplasma gondii* is a widely prevalent protozoan parasite in human populations. This parasite is thought to be primarily transmitted through undercooked meat and contamination by cat feces. Here, we seek to determine if *Toxoplasma gondii* cysts can be found within human semen.

**Methods:** We used a mixture of histological and immunofluorescence stains to visualize *Toxoplasma gondii* cysts in thin smears of human semen. Further, we probed for presence of bradyzoite-specific mRNA transcription using in-situ hybridization.

**Results:** We visualized *Toxoplasma gondii* cysts in ejaculates of immune-competent and latently infected human volunteers. We confirmed the encystment by probing transcription of a bradyzoite-specific gene in these structures. These observations extend previous observations of the parasite in semen of several non-human host species, including rats, dogs, and sheep.

**Conclusions:** *Toxoplasma gondii* infection is a clinically significant infection, in view of its high prevalence, its purported role in neuropsychiatric disorders such as schizophrenia, as well as in the more serious form of congenital toxoplasmosis. Our demonstration of intact *Toxoplasma gondii* cysts in the ejaculate supports the possibility of sexual transmission of the parasite and provides an impetus for further investigations.

© 2022 The British Infection Association. Published by Elsevier Ltd. All rights reserved.

**Summary**

Introduction

*Toxoplasma gondii* (thereafter *Toxoplasma*) is a protozoan and intracellular parasite with a two-stage life. Its range of intermediate hosts includes a wide variety of homeothermic animals, including livestock, humans, and human commensals. Sexual reproduction or gametogenesis is confined to the gut of members of the cat family Felidae. *Toxoplasma* is one of the most prevalent human parasitic infections, with epidemiological studies suggesting a prevalence rate of ~30–40% worldwide. Prevalence is estimated to be ~11% in the USA and 17% in Singapore, with some geographical locations reporting a prevalence rate as high as 70%.1–5

*Toxoplasma* differentiates into fast-dividing tachyzoites during the acute phase of the infection, disseminating in a wide range of tissue types. As the host body mounts an immune response and inflammation, *Toxoplasma* migrates to immune-privileged sites like the brain and eyes. Here it differentiates into bradyzoites housed within tissue cysts, characterized by slow division and quiescent metabolism.1 These tissue cysts go through periodic recrudescence and cycles of immune response and re-encystment. The serological presence of anti-*Toxoplasma* IgG antibodies is often used as a proxy for the presence of latent *Toxoplasma* infection. Immune-competent humans typically do not display overt clinical symptoms.6 Therefore, toxoplasmosis is commonly thought of as a latent and clinically asymptomatic infection with lifelong persistence. However, this perspective is undergoing revision at present. Several reports of retrospective and prospective studies associate *Toxoplasma* infection status with an increased risk of developing neuropsychiatric disorders including bipolar disorder, addiction, obsessive compulsive disorder and schizophrenia.7–13

Humans are primarily infected through exposure to cat feces or by consuming undercooked meat containing parasitic cysts.14 Vertical transmission from acutely infected mothers to the developing fetus and organ transplants are other possible modes of *Toxoplasma* transmission in humans.15 Interestingly, there is growing evidence to non-human hosts that *Toxoplasma* can also be sexually transmissible. For example, *Toxoplasma* breaches the blood-testes barrier in rats and encysts within epididymis during latent infection.16,17 *Toxoplasma* in this host species transfers to the females through ejaculate and establishes chronic infection in both females and males, with documented vertical transmission.
females and resultant offspring. Aside from rats, viable parasites have been isolated from the semen of asymptomatic seropositive dogs\(^{18}\) and goats.\(^{19}\) Subsequently, female dogs and goats inseminated with this infected semen test positive for anti-Toxoplasma antibodies. Serocconversion is also achieved when female sheep are inseminated with Toxoplasma-spiked semen.\(^{20}\) These strands of evidence in non-human species suggest that Toxoplasma can antagonistically cross the blood-testes barrier, an uncommon ability amongst pathogens due to the highly impervious nature of this tissue barrier.

Recently, epidemiological and therefore indirect evidence of male-to-female sexual transmission in humans has been published. A study of sexual partners showed that seropositivity in men increases the risk of infection in women.\(^{21}\) There are strong indices that oral sex could be an important source of infection in heterosexual women and homosexual men.\(^{22}\)

Against this backdrop, we examined the possibility that live Toxoplasma parasites can be ejaculated along with semen in immunocompetent human subjects by examining the semen of seropositive immunocompetent men for the presence of the tissue cysts.

**Materials and methods**

**Sample collection**

Human volunteers were recruited from the Center for Assisted Reproduction, Department of Obstetrics and Gynaecology of the First Faculty of Medicine of Charles University and General University Hospital in Prague, Czechia. The goal and experimental procedures of this study were explained to all participants. Volunteers signed informed consent before the participation. The mean age of the participants was 36 years (SD = 5.14; range: 26–54 years). Ethical approvals were obtained from institutional review boards in General University Hospital in Prague (#384/16; 92/17) and Faculty of Science, Charles University (#2015/29).

Venous blood and semen samples were collected under aseptic conditions. Serum was separated from whole blood via centrifugation (2500 rpm for 10 min) and stored at –20°C before analysis. The full set of samples comprised 723 men, of which 172 were found to contain anti-Toxoplasma IgG antibodies (23.8%). Pairs of semen and matched serum samples were shipped to Singapore on dry ice after obliterating volunteer identity. These shipments contained a subset of the collected samples and were artificially enriched for seropositive participants. Numbers utilized in the staining experiments are listed in successive paragraphs. The serological status was anonymized upon shipping. All procedures conducted in Nanyang Technological University were reviewed and approved by the institutional review board (#IRB-2017–11–021).

**Serology**

Serum was examined for the presence of anti-Toxoplasma IgG antibodies using commercial ELISA kits (Abcam). In line with the manufacturer’s recommendation, the amount of anti-Toxoplasma IgG antibody above ≥ 35 U/ml was considered seropositive, and values < 30 U/ml were categorized as seronegative. Values between 30 to 35 U/ml were classified as equivocal and inconclusive.

**Giemsa stain**

Semen samples from fifty individuals that were sero-positive for anti-Toxoplasma gondii IgG but sero-negative for IgM were used for Giemsa stain for preliminary visualization of the tissue cysts. 50µl of semen sample was pipetted onto ten Superfrosted glass slides (Fisher Scientific) after thawing. The semen sample was smeared across the glass slides and allowed to air-dry completely for an hour. Semen smear slides were stored at 4 °C prior to usage.

Giemsa stain was carried out as previously described. In brief, semen-smeared slides were fixed for 10 min in 100% methanol at room temperature and then completely air-dried. The slides were then stained by Giemsa stain (Sigma-Aldrich) for 45 min at room temperature. The working Giemsa stain was prepared at a 1:5 dilution in phosphate-buffered saline and passed through a 0.45 µm syringe filter. After staining, the slides were rinsed with deionized water and air-dried completely. Slides were coverslipped with a mountant and visualized in a microscope (Zeiss Live Cell observer light microscope, 40X objective lens with a 1.2X digital magnification; total magnification = 400X). A semen sample was determined to be positive if >1 of the ten slides exhibited visual evidence of the cyst-like structures.

**Fluorescence immunohistochemistry**

In order to specifically visualize cyst wall of Toxoplasma gondii cysts, semen samples from seropositive and seronegative individuals were probed for lectin Dolichos biflorus agglutinin binding. Samples from 66 seropositive subjects and 17 seronegative controls were used. None of the subjects included in the determination were positive for IgM antibodies. Experimenters were blind to serostatus during the staining.

Fluorescence immunohistochemistry was carried out as previously described. The human semen smear slides (5µl volume) were fixed in 4% paraformaldehyde (wt/vol. in buffered saline) for 30 min at room temperature. Slides were then permeabilized in buffered saline containing 0.25% (v/v) Triton X-100 for 20 min. Slides were blocked with bovine serum albumin for 1 h at room temperature (Sigma-Aldrich, 1% v/v). Smear slides were then incubated with rhodamine-labeled Dolichos biflorus agglutinin (Vector Laboratories, dilution 1:100), a lectin that binds to carbohydrates found in the Toxoplasma tissue cyst wall. Slides were then repeatedly washed in buffered saline and air-dried. Slides were subsequently coverslipped with a mountant containing DAPI to stain nuclear boundaries (Life Technologies). Tissue cysts were visualized with a laser scanning confocal microscope (Carl Zeiss LSM 710), using a 40X objective lens with a 1.2X digital magnification (total magnification = 400X).

**In situ hybridization**

In-situ hybridization was performed on semen smears to visualize messenger RNA known to be specifically transcribed from the Toxoplasma genome in bradyzoites within tissue cysts (BAG1 mRNA sequence; GenBank: X82213.1). RNAse probes against BAG1 mRNA were commercially obtained (ACDbio) using proprietary techniques. A total of 12 semen samples were examined, of which 10 were sero-positive for IgG and 2 were sero-negative controls. None of the samples originated from IgM positive individuals. Experimenters were blind to the serostatus. Glass-mounted semen smears were stained in accordance with supplier instruction, made available with the probes. A minus-probe negative control was included. Slides were scanned using the Zeiss Live Cell observer light microscope using a 40X objective lens with a 1.2X digital magnification (total magnification = 400X).

**Results**

**Histological visualization confirmed the presence of Toxoplasma cysts in the semen**

Initially, Giemsa stain was used to visualize Toxoplasma cysts in semen smears. Toxoplasma cysts can be easily identified in
this staining preparation, characterized by a heavily stained interior containing bradyzoites along with a lightly stained cyst wall (Fig. 1A and B). A subset of semen from fifty volunteers, previously determined to be seropositive for IgG antibodies, was used for this experiment.

Ten glass slides were prepared by smearing a total of 50 μl of the sample. The presence of Toxoplasma was apparent in 35 out of 50 cases when the first slide from each individual was stained and visualized containing five μl of semen. We then stained an additional slide for 15 cases that did not show the presence of the parasite in the first instance, noting the presence of Toxoplasma in further ten cases. Upon continuing this successive examination, all 50 seropositive cases contained Toxoplasma cysts in ≥ 1 slide (Fig. 1C).

We subsequently sought to confirm the identity of Toxoplasma cysts by staining the cyst wall with a lectin with a known affinity for wall carbohydrates. A rhodamine-labeled Dolichos biflorus agglutinin was used for this purpose. We only examined a total volume of 5 μl of semen during this experiment. Differential interference contrast showed evidence of fluorescent cyst wall encompassing spherical cyst-like structures in the semen smears (Fig. 2). Out of 66 distinct seropositive samples, 26 exhibited immunohistological evidence of Toxoplasma gondii cysts (~39%). In contrast only 2 out of 17 seronegative samples (~12%) exhibited presence of cyst-wall like structures.

We further probed a bradyzoite-specific mRNA to confirm that the intra-cyst structures indeed constituted Toxoplasma bradyzoites. In-situ hybridization confirmed the presence of BAG-1 mRNA inside the tissue cysts, specifically demonstrating the transcription event specific to Toxoplasma encystment (Fig. 3). Seronegative samples did not show staining specific to Bag1 mRNA. Similarly, minus–probe negative controls did not show evidence of the staining.

Discussion

In this report, for the first time, we demonstrate the presence of Toxoplasma tissue cysts within human semen. These observations support the notion that Toxoplasma is a sexually transmissible infection in humans, congruent to prior observations made in non-human intermediate hosts.

Although infections in the human population are mainly asymptomatic (but see22,26), toxoplasmosis remains a public health problem for some populations. Symptomatic congenital infection by Toxoplasma, for example, presents with a prevalence rate of 0.7 cases per 1000 live births in the USA.27 Moreover, the infection during the first trimester of pregnancy in expectant mothers who have not been previously infected by Toxoplasma carries a substantial risk of miscarriages or stillbirth.28 The existence of the venereal route of the Toxoplasma transmission could potentially explain the poor efficiency of the educational campaigns that strive to reduce congenital toxoplasmosis risk. A large fraction of Toxoplasma infections in pregnant women cannot be attributed to the known risk factors,29,30 and sexual transmission could explain part of these infections. The potential of sexual transmission, thus, can signify particular risk in terms of pregnancy outcome and is also of consequence to the operation of sperm banks.

All methods employed in the current manuscript show in situ presence of the parasite. We have not confirmed viability of these tissue cysts using bioassay or artificial insemination. Our observations show the presence of tissue cysts in the semen but do not prove if these cysts can successfully initiate and establish infection in sexual partners post-coitus. This is an important gap in evidence because it is possible that the parasite presence in the semen could be a dead end for the transmission. Female reproductive tract is lined with blood-tissue barriers that might prove non-conducive for the infection. Moreover, it is also presently unclear if sexual transmission, even if possible, forms a significant fraction amongst all possible routes of infection to justify epidemiological implications.

Earlier non-human studies have demonstrated that ejaculated Toxoplasma cysts can result in seroconversion in females as a direct result of intercourse.16,31,32 Such clear data remains absent in the case of humans, although several strands of evidence support that possibility. A recent study using a similar cohort of clients from the Center of Assisted Reproduction, for example, showed that Toxoplasma infection was 1.4 times more prevalent in women part-
Fig. 2. Representative confocal images depicting *Toxoplasma* cyst wall (red), stained with rhodamine-labeled *Dolichos biflorus* lectin, superimposed on a brightfield differential interference contrast image of the cyst. Scale bar = 10 μm. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.).

ners of the infected men at a prevalence rate of 25.6%, compared to partners of uninfected men at a rate of 18.2%. Concordance of serostatus between sexual partners cannot, by itself, provide strong support to the possibility of sexual transmission. Partners often share living environments and other risk factors like exposure to cats, soil, or cooking practices. Nonetheless, the increased risk of the infection due to living with infected partners appears specific to women and not men. This sexual dichotomy supports the idea of sexual transmission through ejaculate. Moreover, there is a positive association between the prevalence of *Toxoplasma* infection and the prevalence of other sexually transmitted diseases in humans, which further consolidates the argument that *Toxoplasma*
can be sexually transmitted.\textsuperscript{14} \textit{Toxoplasma} cysts are resistant to, and possibly even need to be activated by, gastric juices, with consumption of undercooked meat being a primary risk factor for the infection. A recent study raises the possibility that oral sex can be a risk factor for \textit{Toxoplasma} infection in heterosexual and homosexual partners.\textsuperscript{72}\ According to this, the seroprevalence of toxoplasmosis was increased in women and (non-significantly) also in men reporting to be engaged in oral sex (fellatio, not cunnilingus).\textsuperscript{29}

The blood-testes barrier consists of Sertoli cells that are present at the exterior surfaces of seminiferous tubules and form inter-cellular tight junctions.\textsuperscript{33,34} Two conductive but distinct microenvironments exist for the development of germ cells. The basal part of seminiferous tubules houses spermatogonial cell division, differentiation, and renewal of preleptotene.\textsuperscript{35} The basal part is separated from the adluminal or apical region by the blood-testes barrier, creating an immune-privileged environment for spermogenesis, meiosis, and spermiation.\textsuperscript{36,37} The blood-testes barrier is one of the most impervious barriers within the mammalian body. Most infections found in human seminiferous tubules, epididymis, and testes represent a retrograde invasion from the perineal canal.\textsuperscript{38,39} Relevant examples of such retrograde infections include \textit{Neisseria gonorrhoeae} or \textit{Chlamydia trachomatis}.\textsuperscript{38,39} Further example of an infection with retrograde ascent includes \textit{Pseudomonas aeruginosa}.\textsuperscript{40} In contrast, the anterograde passage of pathogens from the testicular lumen side is much infrequent in immunocompetent males. \textit{Toxoplasma} might provide an interesting exception to this rule. Congruent with the presence of the parasite within testes, \textit{Toxoplasma} increases testosterone production in rats\textsuperscript{41} and humans\textsuperscript{42} and facilitates testosterone-dependent phenotypes in males.\textsuperscript{43-45}

Observations in this study lay the groundwork for future experiments to determine if the presence of the parasite in human semen can be generalized to other population cohorts. Further experiments are also warranted to examine the possibility and relative importance of sexual transmission in the epidemiology of \textit{Toxoplasma} infection and the etiology of congenital toxoplasmosis.

\textbf{Statement of ethics}

All experimental procedures were reviewed and approved by Institutional review boards of University Hospital in Prague (#384/16; 92/17), Faculty of Science at Charles University (#2015/29), and Nanyang Technological University (#IRB-2017-11-021).

\textbf{Availability of data and materials}

Authors confirm that, should the manuscript be accepted, the data supporting the results will be archived in an appropriate public repository such as Dryad or Figshare, and the data DOI will be provided at the end of the article.

\textbf{Declaration of Competing Interest}

Authors declare an absence of any competing interests.

\textbf{Acknowledgements}

This work was financially supported by the Ministry of Education, Singapore (grant RG136/15) and Charles University Grant Agency (grant 262721). We would like to thank the medical staff from the Center for Assisted Reproduction, Department of Obstetrics and Gynaecology, First Faculty of Medicine, Charles University and General University Hospital for their assistance with sample collection.

\textbf{References}