What is the importance of zoonotic trichomonads for human health?

Julia M. Maritz¹, Kirkwood M. Land², Jane M. Carlton¹, and Robert P. Hirt³

¹Center for Genomics and Systems Biology, Department of Biology, New York University, New York, NY 10003, USA
²Department of Biological Sciences, University of the Pacific, Stockton, CA 95211, USA
³Institute for Cell and Molecular Biosciences, Faculty of Medical Sciences, Newcastle University, Newcastle upon Tyne, NE2 4HH, UK

Trichomonads are common parasites of many vertebrate and invertebrate species, with four species classically recognized as human parasites: Dientamoeba fragilis, Pentatrichomonas hominis, Trichomonas vaginalis, and Trichomonas tenax. The latter two species are considered human-specific; by contrast, D. fragilis and P. hominis have been isolated from domestic and farm mammals, demonstrating a wide host range and potential zoonotic origin. Several new studies have highlighted the zoonotic dimension of trichomonads. First, species typically known to infect birds and domestic mammals have been identified in human clinical samples. Second, several phylogenetic analyses have identified animal-derived trichomonads as close sister taxa of the two human-specific species. It is our opinion, therefore, that these observations prompt further investigation into the importance of zoonotic trichomonads for human health.

The trichomonad lineage in phylum Parabasalia

Trichomonads are anaerobic, flagellated protists belonging to the large and diverse groups Trichomonadida and Tri-trichomonadida of phylum Parabasalia [1]. They are characterized by the presence of three to five anterior flagella, hydrogenosomes – hydrogen-producing organelles corresponding to anaerobic versions of mitochondria [2], a parabasal body (a large Golgi), and a complex cytoskeleton. A few species have been isolated from environmental samples and may represent free-living species; however, the majority of species form symbiotic interactions (see Glossary) with various animal hosts. Among the parasitic trichomonads, several species inhabit the oral, digestive, and urogenital tracts of invertebrate and vertebrate hosts, including livestock, pets, and humans.

Historically, phylum Parabasalia was divided into two groups based on morphological characteristics; however, the recent inclusion of molecular data recovered six groups: Trichomonadida, Tritrichomonadida, Hypotrichomonadida, Cristomonadida, Spirotrichonympha, and Trichonympha [3]. The Trichomonadida, Tritrichomonadida, and Hypotrichomonadida are of primary concern to parasitologists; however, the evolutionary relationships within and between these groups are unclear [4]. Several molecular phylogenies have attempted to resolve these evolutionary relationships using phylogenetic markers such as ribosomal RNA (rRNA) and protein coding genes (Figure 1), which give inconsistent phylogenies [4,5].

Four species of trichomonad are considered human parasites: Trichomonas vaginalis (found in the urogenital tract) [6], Trichomonas tenax (localized to the oral cavity) [7], and Pentatrichomonas hominis and Dientamoeba fragilis (located in the digestive tract) [8,9]. Only one species has well-established pathogenic potential: T. vaginalis, the cause of the most prevalent non-viral sexually transmitted infection in humans, trichomoniais [10]. Only T. vaginalis and T. tenax are considered human-specific, with the former characterized by the richest, although still limited, epidemiology data [11], but very little is known about the latter. P. hominis and D. fragilis can cause gastrointestinal symptoms in some patients, such as abdominal pain and diarrhea [8,12]. D. fragilis has also been proposed as a potential causative agent of irritable bowel syndrome (IBS) [13,14], but debate surrounds its pathogenicity, infection route, and epidemiology [15]. In addition, several trichomonad species are of veterinary importance, such as the avian pathogens Trichomonas gallinae, Tetratrichomonas gallinarum, and Histomonas meleagridis [16–19], and Trichomonas foetus, the causative agent of a venereal disease in cattle [20]. This extensive host range, along with the isolation of D. fragilis [21] and P. hominis [22] from various animal hosts, suggests that certain species of trichomonads may exhibit the characteristics of zoonoses. Although the question of zoonotic trichomonad has been considered for some years (e.g., [23,24]), recent results from several different sources have highlighted this potential. Here we summarize the clinical and phylogenetic studies that suggest a zoonotic potential for trichomonads, discuss their implications for human health, and the next steps required for investigation into their epidemiology, pathology, and evolution.

New evidence supports the zoonotic potential of trichomonads

Human trichomonad infections are not body site-specific
The four trichomonad species recognized as human parasites were initially thought to be site-specific [25] (Table 1). However, various clinical studies have shown that they can...
Transmission:
Parasitic Symbiosis: Dysbiosis:
Disease incidence: the number of new disease cases that occur in a population for a given time period (typically per year).
Dysbiosis: an imbalance of the microbiota (the microbial populations at a particular body site of an animal host) that leads, or predisposes, the host to disease conditions.
Emerging infectious disease: outbreaks of previously unknown diseases or known diseases that show an increase in incidence, expansion in geographic range, or spread to a new population. Emerging infections can be caused by previously unknown or undetected infectious agents, newly evolved strains, environmental changes, and changes in human demography.
Intermediate host: a host in or on which a pathogen spends a part of its life, usually a transition period, but does not reach sexual maturity.
Mutualism (mutualist): a form of symbiosis between two organisms in which both benefit from the relationship. Some gut parabasalids from termites are thought to represent mutualists.
Opportunistic: a potential pathogen that typically does not cause disease in a healthy host, but can in particular situations cause disease, for example, owing to the compromised immune system (e.g., attributable to AIDS, chemotherapy, or malnutrition) of the host. Lung trichomonads represent probable opportunistic infections – see main text.
Parasitic (parasite): a non-mutual symbiosis between two species where one, the parasite, benefits at the expense of the other, the host. Parasites typically do not kill their hosts but exploit them for resources necessary for their survival. Obligate parasites cannot complete their life cycles and reproduce without a suitable host.
Pathogenic (pathogen): a broad term that refers to the ability of an organism to cause disease. It is typically used to describe an infectious agent or microorganism, such as a bacterium, protist, virus, etc., that causes disease in its host. Some pathogens, for example, parasites Acanthamoeba spp. and Naegleria fowleri and fungi Aspergillus spp. are free-living species thriving in the environment and occasionally infect humans, often in an opportunistic manner in compromised hosts.
Pathogenicity: the ability of a pathogen to overcome host defences and cause disease.
Re-emerging infectious disease: the reappearance of a historically known infectious disease after a significant decline in incidence. Acquired resistance of pathogens to antimicrobial medications is an important factor in the re-emergence of many diseases. Examples include West Nile virus, cholera, MRSA (methicillin-resistant Staphylococcus aureus).
Reservoir: the habitat or host that harbors an infectious agent, where it can live, grow, and multiply. Reservoirs can include humans, animals, and serve as a source of potentially disease outbreaks.
Symbiosis: a close and often long-term relationship between two or more different biological species. These relationships can be obligate or facultative, mutualistic, commensalistic, or parasitic.
Transmission: the passing of an infectious agent from one host to another host. Infect transmission routes include: physical contact, contact with a contaminated environment or surface, airborne transmission, and fecal–oral transmission. Indirect transmission routes involve another organism such as an insect vector or intermediate host.
Vector: an organism that carries and transmits a pathogen from an infected individual to another individual.
Virulence: a property of a pathogen, such as specific structural elements or biochemical compounds commonly called virulence factors that cause a reduction in host fitness or damage to the host. It is now recognized that virulence is multifactorial and involves characteristics of both the pathogen and its host, which influence the outcome of their interaction and hence the observed virulence (e.g., an opportunistic pathogen in immunocompromised patients).
Zoonosis: an infectious organism, such as a bacterium, virus, parasite, or fungus, transmissible between wildlife or domesticated animals and humans. Examples include: (i) the Lyme disease bacterium Borrelia transmitted to humans by ticks from a natural reservoir in rodents; (ii) the malaria parasite Plasmodium knowlesi transmitted by Anopheles vectors that causes malaria in monkeys and humans; and (iii) Cryptosporidium parvum, a parasite found in cats, dogs, and farmed animals and transmitted as a cyst in contaminated water, food, or through the fecal–oral route. Zoonoses are the leading cause of emerging infectious diseases worldwide, responsible for devastating disease outbreaks, mortality, and serious socioeconomic consequences.
Zoonotic potential: the potential for infectious diseases of wildlife or domestic animals to be transmitted to humans.

Also found in atypical locations. For example, T. tenax, a commensal of the human mouth found in patients with poor oral hygiene [7], has been identified by microscopic and molecular methods in the upper and lower respiratory tracts [26,27]. One possibility that could account from this ‘aberrant’ location is inhalation of the parasite from the oral cavity into the respiratory tract. However, in some cases where T. tenax was identified in the respiratory tract, no parasites were found in the mouth [26]. Other human trichomonad species have also been identified in the respiratory tract including the sexually transmitted species T. vaginalis [28,29] and the gut parasite P. hominis [30], which suggests that these species too can proliferate outside their usual body sites.

At least five species of trichomonad, including P. hominis, T. tenax, T. vaginalis, T. foetus, and T. gallinarum, have been identified in the human respiratory tract and as causative agents of pulmonary trichomoniasis (Table 1). They have been found in up to 60% of patients with Pneumocystis pneumonia (PCP) and in up to 30% of patients with acute respiratory distress syndrome (ARDS) [31]. Because trichomonads are microaerophilic it is unlikely that they initiate and cause these diseases themselves, but may represent secondary and opportunistic infections that could exacerbate symptoms and prolong illness [23]. These trichomonad respiratory infections seem to depend upon: (i) the presence of bacteria on which to feed and (ii) local anaerobic conditions caused by PCP or ARDS-associated infections [32] but not necessarily upon immunosuppression, because drugs against PCP consistently cure patients of pulmonary trichomoniasis and, in one study, treated ARDS patients were not found to be immunocompromised [25]. Thus, the presence of an increasing number of distinct trichomonads in a broader range of clinical samples from patients with diverse diseases, such as AIDS, rheumatoid arthritis, prostate cancer, pulmonary infections (empyema and pneumonia in addition to PCP and ARDS), and digestive conditions such as diarrhea and IBS [33–35], is becoming increasingly apparent. Indeed, the frequency of pulmonary trichomoniasis infections may be higher than reported because transformation of parasites from the motile, pear-shaped stage to the ameboid stage renders microscopic identification in clinical samples difficult [31], highlighting the importance of molecular data to identify such infections [25].

Non-human species of trichomonad have been isolated from clinical samples
Trichomonads were thought to have strict host specificity [25]; however, trichomonad parasites not previously reported as infecting humans have recently been found in human clinical samples (Table 1). For example, parasites belonging to the genus Tritrichomonas can be isolated from the reproductive tract of cattle (Tritrichomonas foetus), the nasal mucosa and intestine of pigs (Tritrichomonas suis), and the intestine of non-human primates (Tritrichomonas mobilensis) [20]. Another example is T. foetus, historically considered specific to cattle [25,36]. Nonetheless, experimental cross-infections of the parasites between pigs and cattle in addition to analysis of molecular data suggest that these three species should be considered...
strains of the same species [20,37]. In addition, several different genotypes of *T. foetus* have been identified as causing diarrhea in cats in ~12 countries [38,39] and have also been isolated from dogs with diarrhea [40]. Moreover, in several new clinical cases, *T. foetus* or *T. foetus*-like organisms have unexpectedly been identified in the lungs of human patients [41]. Such findings suggest that *T. foetus* is a zoonotic parasite capable of colonizing an extensive range of hosts and body sites.

Other examples of non-human trichomonads recently found to infect humans are members of the genus *Tetratrichomonas*, currently the largest genus in phylum Parabasalia. *Tetratrichomonas* species are found in the small intestine of a wide spectrum of invertebrate and vertebrate hosts, such as leeches, birds, and rodents [42]. Indeed, some species of tetratrichomonads are known to infect a wide range of unrelated hosts, such as *Tetratrichomonas prowazeki*, which has been found in species of amphibians and reptiles [42]. Another example, *Tetratrichomonas gallinarum*, is primarily thought of as an avian parasite of the digestive tract in domestic and wild birds [43], although its pathogenicity is not well established [17]. However, several recent studies have identified *Tetratrichomonas* strains isolated from human lungs or the human

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**Table 1. Trichomonad species identified in clinical studies***

<table>
<thead>
<tr>
<th>Species</th>
<th>Primary hosta, infection site</th>
<th>Host range: infection site</th>
<th>Diagnostic method used to detect in other hosts or sites</th>
<th>Clinical conditions</th>
<th>Refs</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Trichomonas vaginalis</em></td>
<td>Human; UGT</td>
<td>Humans; RT</td>
<td>PCR and sequencing of the ITS1–5.8S–ITS2 rRNA region</td>
<td>Trichomoniasis, pulmonary infections, AIDS</td>
<td>[10,27,28]</td>
</tr>
<tr>
<td><em>Trichomonas tenax</em></td>
<td>Human; DT and buccal cavity</td>
<td>Humans; RT</td>
<td>PCR and sequencing of the ITS1–5.8S–ITS2 rRNA region</td>
<td>Salivary trichomoniasis, pulmonary infections</td>
<td>[7,25,26]</td>
</tr>
<tr>
<td><em>Dientamoeba fragilis</em></td>
<td>Humans; DT</td>
<td>Humans and other mammals; DT</td>
<td>Fecal smears, PCR, and sequencing of the ITS1–5.8S–ITS2 rRNA region, 18S rRNA</td>
<td>Chronic diarrhea, IBS</td>
<td>[13,14,21]</td>
</tr>
<tr>
<td><em>Pentatrichomonas hominis</em></td>
<td>Not known; DT</td>
<td>Humans and other mammals; DT</td>
<td>Fecal smears, PCR, and sequencing of the ITS1–5.8S–ITS2 rRNA region</td>
<td>Diarrhea, pulmonary infections, rheumatoid arthritis</td>
<td>[8,30,33]</td>
</tr>
<tr>
<td><em>Tritrichomonas foetus</em></td>
<td>Bovine; UGT, DT</td>
<td>Humans and other mammals; RT</td>
<td>PCR and sequencing of the ITS1–5.8S–ITS2 rRNA region, EF-1α gene, and TR7/TR8 variable length region</td>
<td>Pulmonary infections, AIDS</td>
<td>[20,38,41]</td>
</tr>
<tr>
<td><em>Tetratrichomonas gallinarum</em></td>
<td>Birds; DT</td>
<td>Birds, humans; RT</td>
<td>PCR and sequencing of the ITS1–5.8S–ITS2 rRNA region</td>
<td>Pulmonary infections</td>
<td>[24,44]</td>
</tr>
<tr>
<td><em>Tetratrichomonas</em> sp.</td>
<td>Not known</td>
<td>Humans; RT</td>
<td>PCR and sequencing of the ITS1–5.8S–ITS2 rRNA region</td>
<td>Pulmonary infections</td>
<td>[24,43]</td>
</tr>
</tbody>
</table>

*Abbreviations: DT, digestive tract; UGT, urogenital tract; RT, respiratory tract; ITS, internal transcribed spacer; IBS, irritable bowel syndrome; PcP, *Pneumocystis* pneumonia; ARDS, acute respiratory distress syndrome.

*The primary host may not represent the true natural history of the species, which may have a broader host range.

*Pulmonary infections include PcP, ARDS-associated infections, pneumonia, and can lead to empyema.
oral cavity as *T. gallinarum* or *T. gallinarum*-like organisms [24,44,45]. Studies have also shown that genus *Tetratrichomonas* is much more diverse than previously thought and that *T. gallinarum* comprises at least three cryptic species with variable host specificity, some that represent human isolates [24,43]. Notably, experiments failed to transmit two *Tetratrichomonas* of human origin to birds, although the authors suggest this result could be explained by either adaptation of the *T. gallinarum*-like trichomonads to the human host or extensive in vitro culturing, so that infection of birds was no longer biologically achievable [24].

**Molecular phylogenies reveal close relationships between human and avian trichomonads**

Recent molecular phylogenetic analysis of trichomonads using rRNA and protein coding genes (e.g., *Rpb1*) has begun to answer important questions regarding trichomonad phylogeny. *Rpb1* is a ubiquitous eukaryotic gene coding for the largest subunit of RNA polymerase II and is present as a single copy in many eukaryotes. A recent analysis of *Rpb1* generated a fully resolved phylogeny of Trichomonadea, Tritrichomonadea, and Hypotrichomonadea, and species and isolates within these groups (Figure 2) [5]. Interestingly, the phylogeny recovered some avian isolates of *Trichomonas* spp. as sister taxa to *T. vaginalis*, and *T. tenax* as closely related to *T. gallinaceae*; these findings are consistent with previous phylogenies based upon rRNA and other protein coding genes [24,46–48]. The common ancestor to this complex is also related to the avian *T. gallinarum* (e.g., [42]). These new phylogenies complicate the inferred relationship between *T. vaginalis* and *T. tenax*, which on the basis of host specificity might be expected to be sister taxa [47]. In this scenario, an ancestor of both species infected humans and subsequently differentiated into two distinct species with different body site preferences. However, the
phylogenetic data suggest the zoonotic transfer of trichomonad parasites from humans to birds, and/or vice versa, on at least two occasions, or a combination of these events.

Other studies investigating outbreaks of avian trichomonosis in wild birds, primarily attributed to _T. gallinae_ infections, have also confirmed the close relationship between avian and human trichomonads [48]. Epidemic infections by _T. gallinae_ of passerine species in Europe have recently been described and associated with high mortality and population decline [18]. This dramatic case of disease emergence demonstrates the potential for a trichomonad to jump host species (columbiform to passerine) and spread rapidly through populations. The disease is thought to have been initiated and spread through transfer of the parasite via contaminated water and bird feeders, as well as through direct contact between passerines [17]. Furthermore, the recent isolation and preliminary characterization of _Trichomonas stableri_ associated with epidem- ic mortality in California Band-tail pigeons suggest that avian trichomonosis may be caused by pathogens other than _T. gallinae_ and _H. meleagridis_ [17,49]. Genetic analysis at multiple loci indicated _T. stableri_ to be more closely related to _T. vaginalis_ than to the bird-associated _T. gallinae_ [49] (Figure 2), further supporting the hypothesis that trichomonads are crossing host boundaries. Generating genetic markers and whole genome sequences of _T. stableri_ and other _T. vaginalis_-like isolates derived from cases of epidemic avian trichomoniasis will provide important insight into the evolution and origins of these pathogens. For example, does the _Trichomonas_ sp. isolate from a white-winged pigeon extremely closely related to isolates of _T. vaginalis_ represent a case of human to bird transfer (accession HM016231 in Figure 2)?

The implications of zoonotic trichomonads for human health

According to current disease dynamic models, the zoonotic emergence of parasitic diseases in humans is typically associated with several characteristics, including broad host range, genetic variability, presence of genotypes better suited to the parasitism of humans, and modified pathogenic potential [50]. Emergent zoonoses are thought to appear through a number of different stages, for example, some develop as animal ‘parasitoses’ that are newly transmissible to humans, although the source of the disease remains the animal reservoir [50,51]. In other cases, parasites are able to cross the species barrier, modify their specificity, and become sustainably transmissible from human to human [50,52,53]. The evolution of these emergent parasitoses is not linear, and an explanation for such a complex process requires consideration of the multi-host ecology and complex dynamics of zoonotic infections [54]. Based on these models and the clinical and molecular evidence discussed previously, it may be that several trichomonads are at different stages of zoonotic emergence (Figure 3). These observations raise important questions regarding the implications of the potential widening pathological spectrum of trichomonads in humans and suggest that owing to links with other diseases these parasites may be of greater medical importance than previously thought.

Historically, trichomonads have not been considered as emerging infections because of their site- and host-specific occurrence. Nonetheless, the presence of trichomonads in a diverse array of clinical disorders suggests that they may exhibit a form of opportunism and multiply when local conditions are favorable. For example, the diseases in which trichomonads are found as co-infecting agents in respiratory infections are probably not limited to Pcp and ARDS-associated infections, but may include other pulmonary diseases such as cystic fibrosis [31,32]. Overall, the high prevalence of pulmonary diseases globally [55] combined with the higher burden of both lung conditions and zoonotic diseases among people in resource-limited settings [55,56] suggest that only the ‘tip of the iceberg’ of pulmonary trichomoniasis may currently be known. Digestive tract infections by trichomonads are also increasingly recognized as being common. Although the exact clinical profile of _D. fragilis_ is still poorly understood some consider this species to have pathogenic capabilities [12], and recent studies have associated the rise of IBS with a high prevalence (~40%) of _D. fragilis_ in Europe [35]. However, the pathogenicity of _D. fragilis_ has been questioned owing to the asymptomatic nature of many infections, and it is considered by some as a commensal of the intestinal flora [57]. Indeed, treatment of _D. fragilis_-infected children with metronidazole was not associated with better clinical outcomes [58].

Because association is not evidence for causality, additional data are required to establish the pathogenicity of trichomonads in the digestive tract and ‘aberrant’ body sites such as the lungs. In this context, it is important to consider characteristics of both host and parasite with regard to the outcome of their interactions. For example, one extreme is represented by severely immune-compromised patients that are more susceptible to a wider range of microbial infections compared with immunocompetent hosts [59]. When studying the outcome of human–microbe interactions, a complex interplay among viruses, bacteria and archaea, microbial eukaryotes, and animal parasites influence the health status of the human host, with mucosal microbiota playing a key role influencing health and disease status [60,61]. Based on these considerations and examples, trichomonads may be more prevalent and have a wider pathological spectrum in humans than currently recognized, influencing human health through direct pathologies but also indirectly through dysbiosis of the mucosal microbiota and local inflammation, facilitating transmission of pathogens – a prime example being _T. vaginalis_ infection and bacterial vaginosis contributing to HIV transmission [61,62]. The potential influence of gut trichomonads to human health will also have to consider their potential impact on the gut microbiota, which might explain observations of the association between _D. fragilis_ and IBS through inducing gut dysbiosis [35]. Indeed, the ability of trichomonads to live on a variety of mucosal tissues may be the key to their wide host range and ability to develop infections at different body sites, as well as contribute directly or indirectly to pathologies. Once the capacity to thrive on vertebrate mucosal surfaces has developed, there may be less of a barrier to cross both species and mucosal sites. For example, in the case of
Figure 3. Speculative models of zoonoses caused by trichomonads. Trichomonads are listed on the right and colored according to primary hosts assigned historically in the literature. Unbroken lines represent known infections or transmission routes, and broken lines represent speculative infections or transmission routes for which data are lacking. The relationships are represented as follows: (blue box) trichomonads identified in wild bird species (e.g., green finch [16] and toucan [81]) in partially domesticated species (rock dove) and in fully domesticated species (chicken) circulate within these populations with variable host specificity [17] (blue unbroken circle with arrow). Two of the four avian trichomonads listed (Tetratrichomonas sp. and Tetratrichomonas gallinarum) have been identified in human lungs [24], and Trichomonas gallinae and Trichomonas stableri are also included owing to their close relationship to Trichomonas tenax and Trichomonas vaginalis [5,49]. [Red box] T. vaginalis and T. tenax are the two species considered human-specific, with known human-to-human infections (unbroken red circle). The close genetic relationship of the human and avian trichomonads (Figure 2) suggests either independent zoonotic acquisitions from avian sources (broken blue arrow) or transfer of the parasites from humans to birds through environmental contamination (broken red arrow). [Green box] Trichomonas foetus has been isolated from a variety of pets and farm animals, with the same strain known to infect cattle and pigs (unbroken green arrow) [29], but different genotypes infecting cattle and cats [29,31]; the origins of dog infections remain unclear [32]. Thus, there are at least two T. foetus genotypes capable of colonizing an extensive range of hosts, including humans [41] (broken green circle and arrow). The lack of precise epidemiological data is indicated by "?." [Purple box] Pentatrichomonas hominis has been isolated from a variety of pets and farm animals [22], but little is known about its infection route and epidemiology; the same strain could be circulating between all identified hosts (broken purple circle). [Orange box] Dientamoeba fragilis has been isolated from farm animals (pigs) and non-human primates (gorillas), with the same strain known to infect pigs and humans [21] (unbroken orange arrow). Recent evidence suggests that household pets do not play a role in transmission [82]; however, the origins remain unclear and multiple strains could be circulating in animal hosts (broken orange circle and arrow). Additionally, given recent prevalence and transmission data it seems unlikely that transmission from non-human hosts represents a significant proportion of infections. Contaminated surfaces and water [83], uncooked meat, or direct contact with pets and farm animals could lead to animal-to-human transmissions of trichomonads. Initial infections were presumably through the digestive tract (via oral ingestion) with further progression to the lungs for some (various) species or the urogenital tract (T. vaginalis).

T. foetus, this sexually transmitted species may represent a recent transfer from the digestive to the urogenital tract, with a capacity of the parasite to thrive in the gut in different species (e.g., pigs, cats, and dogs).

Investigating trichomonads as emerging infectious diseases and zoonoses

Trichomonads provide a unique system for the study of the origins and pathobiology of zoonotic and emerging infectious diseases. In addition, they have attracted interest as model systems for evolutionary biology and comparative genomics [63,64], and for biochemical, molecular, and cell biology investigations [2,65,66]. The T. vaginalis genome sequence published in 2007 was the first species of trichomonad to be sequenced [67], and others are currently underway including isolates of T. foetus, P. hominis, T. gallinae, and T. tenax. These sequences will enable comparative analysis of common and unique parasitic modes of life cycle, and possible adaptive mechanisms. For example, T. vaginalis and T. foetus appear to have evolved independently to colonize the urogenital tracts of different mammalian hosts [68]. In addition, T. foetus has been isolated from the digestive tract of cats and dogs, indicating that it is capable of colonizing a broad range of hosts and environments [69]. These two species of sexually transmitted trichomonad probably represent cases of convergent evolution and provide an opportunity to compare the derived similarities and the origins of these traits that coincide with a shared niche.

Species of trichomonad exhibit a range of genome sizes, from ~94 Mb for the P. hominis genome to ~177 Mb for the T. foetus genome [70]. The T. vaginalis genome sequence revealed the ~160-Mb genome to be a result of expanded transposable elements and protein coding gene families,
including those responsible for interaction of the parasite with its immediate environment [71–73]. It has been hypothesized that the large genome size may be a recent event that occurred when the most recent common ancestor of T. vaginalis underwent a population bottleneck during its transition from the digestive tract to the urogenital tract. Because genome size is positively associated with cell volume, an increase in genome size and concomitant increase in cell size might have increased its phagocytic potential as well as surface area for the interaction of the parasite with host cell tissue [67,73]. In this context, it will be particularly interesting to compare the genomes of T. vaginalis with its closely related isolates and species derived from birds (Figure 2) to gain detailed insight into correlations between genome evolution and how this might relate to parasite pathobiology in humans and birds. Lateral gene transfers from bacterial donors sometime in the evolutionary past have also importantly influenced the evolution of T. vaginalis protein coding genes [63,67]. For example, the parasite has gained an almost complete pathway for the degradation of complex glycans present in host mucosal secretions, factors which may contribute to its adaptive potential and pathogenicity [61,63]. Comparison of a broad range of trichomonad genomes will help test this hypothesis and may pinpoint gene families whose acquisition and/or expansions correlate with pathogenicity and facilitated or mediated transitions from: (i) an ancestral animal-only stage to human-inclusive infections or (ii) from the digestive to urogenital tracts.

Trichomonads also provide a unique system to study the features of a zoonotic lifestyle via comparative examination of molecular and cellular characteristics. For example, successful T. vaginalis infections are probably favored by virulence mechanisms such as cytadherence and phagocytosis [72,73]. The ~60 000 predicted protein coding genes of T. vaginalis [67] includes a plethora of candidate genes for surface molecules mediating interaction with host tissues and membrane trafficking and signaling, important processes involved in parasite pathobiology [72,73]. Cysteine proteases in particular have been identified as virulence factors central to the host–pathogen interface in T. vaginalis [61,72–75]. Transcriptomic studies have shown upregulation of some of these T. vaginalis virulence factors in response to contact with host cells in vitro [76] and have also documented their expression under in vitro growth conditions in T. foetus [69]. Similar to T. vaginalis, recent studies have shown the presence of cysteine proteases in the cell-free filtrate of T. gallinae and demonstrated their involvement in its in vitro cytopathogenic effects [77]. Mining other trichomonad genome data to identify important virulence proteins will improve our understanding of the molecular and cellular basis of infections and can be used to test hypotheses, such as whether zoonotic organisms show greater diversity in key virulence proteins underlying their capacity to parasitize a variety of host species and mucosal sites [78].

Concluding remarks and future perspectives
Although we have discussed several recent studies that provide strong evidence for the zoonotic origin and potential of trichomonads, regular and sustained zoonotic transmission of these microbes has yet to be definitively established. To improve our knowledge of the zoonotic origins of trichomonads, detailed investigations including systematic surveys of trichomonads in humans and animals will be required. Molecular methods have been instrumental in our understanding of the biology and complexity of trichomonads so far; however, more data and novel approaches are needed to resolve evolutionary relationships and to improve diagnostic tools. Wide sampling and whole genome sequencing of trichomonads, with subsequent comparative genomic investigations, will facilitate identifying the closest relatives of human trichomonad pathogens, providing a solid evolutionary framework for how these diseases have emerged and forming a basis for epidemiological studies across both animal (wild and domestic) and human hosts. Environmental studies such as the ‘Microbes, Sewage, Health and Disease’ metagenomics project in New York City (http://www.nyu.edu/about/news-publications/nyu-stories/video-mapping-nychs-metagenome.html) will provide important information on potential transmission routes and the patterns, nature, and occurrence of trichomonad infections in humans, animals, and birds; and establish the importance of trichomonads zoonotic transmissions – as has been established for species of Trypanosoma, Cryptosporidium, and Toxoplasma [79]. Furthermore, studies investigating the potential pathogenicity of these parasites in various mucosae (respiratory, digestive, and urogenital) are needed to determine the clinical significance and public health implications of trichomonads. Accumulation of these different types of data will advance our understanding of parasite biology and infection mechanisms, and provide approaches towards developing drug targets and vaccine candidates for species of this increasingly recognized medically and veterinary important lineage.

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