Toxoplasma gondii tachyzoites cross retinal endothelium assisted by intercellular adhesion molecule-1 in vitro

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Retinal infection is the most common clinical manifestation of toxoplasmosis. The route by which circulating Toxoplasma gondii tachyzoites cross the vascular endothelium to enter the human retina is unknown. Convincing studies using murine encephalitis models have strongly implicated leukocyte taxis as one pathway used by the parasite to access target organs. To establish whether tachyzoites might also interact directly with vascular endothelium, we populated a transwell system with human ocular endothelial cells. Human retinal endothelial monolayers permitted transmigration of tachyzoites of RH and three natural isolate strains. Antibody blockade of intercellular adhesion molecule-1 significantly reduced this migration, but did not impact tachyzoite movement across an endothelial monolayer derived from the choroid, which lies adjacent to the retina within the eye. In demonstrating that tachyzoites are capable of independent migration across human vascular endothelium in vitro, this study carries implications for the development of therapeutics aimed at preventing access of T. gondii to the retina.

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Toxoplasma gondii is an apicomplexan parasite that infects approximately two billion persons worldwide.1 The most common clinical manifestation of toxoplasmosis is an ocular infection involving the retina (that is, retinitis), with secondary involvement of the adjacent choroid in severe cases (that is, retinochoroiditis). Ocular toxoplasmosis typically affects healthy adults, and causes blindness in 24% of affected eyes.2 In immunocompromised patients, particularly those who suffer from acquired immunodeficiency syndrome, infection with the parasite may result in a fatal encephalitis.1

After entering the human host, T. gondii spreads to the central nervous system via the bloodstream in tachyzoite form. Two independent groups have convincingly demonstrated that infected monocytes and/or dendritic cells carry tachyzoites across the blood-brain barrier in a murine model of toxoplastic encephalitis.3,4 However, as stated by Tardieux and Ménard,5 a ‘central question’ relates to ‘whether tachyzoites interact with endothelial barriers’. The human retinal vasculature is characterized by a dense capillary network,6 and circulating leukocytes move slowly through retinal capillaries because of their wider diameter, stiff viscoelastic properties and tendency to adhere to the endothelium.7 If lysis of a tachyzoite-infected leukocyte occurred during trans-capillary passage, the potential would exist for a direct interaction between tachyzoite and retinal vascular endothelium. Both free and intracellular tachyzoites circulate in the blood of individuals infected with T. gondii.8

Using a conventional transwell migration system populated with human ocular vascular endothelial cells, we investigated the possibility that, in addition to ‘taxiing’ in a leukocyte, T. gondii tachyzoites might be capable of accessing the human retina by migrating unassisted across the retinal vascular endothelium.

RESULTS
We observed migration of live T. gondii tachyzoites through a simulated human retinal vascular endothelium in a transwell system over a 4-h period (Figure 1a). Smaller numbers of heat-killed tachyzoites, loaded into upper chambers of control transwells, were also recovered from lower chambers at this time. However, there was no significant difference (P>0.05) in permeability to high molecular weight dextran for endothelial monolayers incubated with live, heat-killed or no tachyzoites, and permeability under these conditions remained significantly less (<0.001) than permeability of wells containing membranes coated with collagen alone (Figure 1b),...
suggesting that the endothelial monolayer remained intact for the duration of the experiment. Universal expression of CD144 (VE-cadherin) by CD31-positive retinal endothelial cells after extended confluent culture (Figure 1c), also was consistent with the formation of intercellular junctions across the monolayers. To address the possibility that transendothelial movement was peculiar to RH strain tachyzoites, we performed the same assay, but separately substituted one of three different natural parasite isolates for the clonal strain. The same result was obtained in this series of experiments (Figure 1d). To investigate a possible role for the cell adhesion molecule, intercellular adhesion molecule (ICAM)-1, in migration of *T. gondii* tachyzoites across human retinal vascular endothelium, we performed endothelial transmigration assays after pre-incubating endothelial monolayers with anti-human ICAM-1 antibody or control immunoglobulin G1 (IgG1). ICAM-1 blockade significantly reduced tachyzoite movement across the endothelium by ∼50% (Figure 2a, *P* < 0.001). In contrast, specific antibody blockade of the related cell adhesion molecule, vascular cell adhesion molecule (VCAM)-1, did not significantly impact tachyzoite migration (Figure 2b, *P* > 0.05). To determine whether ICAM-1-mediated endothelial transmigration within the eye was specific to retina, we performed the same experiment, but seeded the transwell membrane with endothelial cells that were isolated from choroid, in place of the retina. Choroid is the ocular tissue that encircles the retina. Consistent with specificity of the retinal endothelial interaction, ICAM-1 antibody blockade did not impact tachyzoite migration across simulated human choroidal vascular endothelium (Figure 2c, *P* > 0.05). When studies were repeated, we observed variation in absolute numbers of migrated parasites, but relationships between experimental conditions remained consistent.

**DISCUSSION**

Elegant studies conducted in murine toxoplasmosis have shown clearly that *T. gondii* tachyzoites infect phagocytic leukocytes, which traffic parasites from the vascular tree to the brain. However, tachyzoites use a form of independent motility known as ‘gliding’ that is parasite actin-dependent. By gliding, they cross polarized epithelial and trophoblastic monolayers, and move through tissues, such as intestine, in which they navigate from the epithelium to submucosal vascular endothelium. These observations led us to hypothesize that *T. gondii* tachyzoites were also capable of crossing from the circulation to a target organ by direct interaction with the vascular endothelium. Recognizing that the most common clinical manifestation of toxoplasmosis was retinitis, we used a transwell system populated with human retinal endothelial cells to test our hypothesis. Our experiments demonstrated that tachyzoites were indeed capable of moving through a simulated human retinal endothelium.

We first evaluated migration of RH strain tachyzoites across a human retinal endothelial monolayer. Although RH is the most widely studied parasite strain, it has remained in culture for multiple decades, and, in addition to exhibiting heightened virulence in comparison with natural isolates of the same haplogroup, it has lost the ability to complete a full life cycle. Therefore, it was important
Figure 2 ICAM-1 blockade significantly inhibits T. gondii tachyzoite migration across simulated human retinal, but not choroidal, vascular endothelium. Graphs showing number of tachyzoites recovered from lower chambers of transwells divided by (a, b) human retinal endothelial cell monolayers or (c) human choroidal endothelial cell monolayers cultured on type 1 collagen, 4 h after upper chambers were loaded with 1 × 10³ live or heat-killed RH strain T. gondii tachyzoites (n = 6 wells, parasite viability = A, 55%; B, 52%; C, 95%). Monolayers were pre-incubated for 2 h with (a) mouse anti-human ICAM-1 IgG1 (α-ICAM-1) or (b) mouse anti-human VCAM-1 IgG1 (α-VCAM-1), or control mouse IgG1 (control). There was a significant reduction in tachyzoite migration across the retinal endothelial monolayer in the presence of anti-ICAM-1 antibody (***P<0.001), but not anti-VCAM-1 antibody (P>0.05) (Student’s t-test). Anti-ICAM-1 antibody did not reduce tachyzoite migration across the choroidal endothelial monolayer (P>0.05) (Student’s t-test). Each graph presents results that are representative of two independent experiments. In all graphs, bars represent mean and error bars represent s.e. of mean.

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**METHODS**

**Culture of T. gondii tachyzoites**

Per the classification of Khan et al.,¹³ clonal (RH: haplogroup 1), natural common (GT-1; haplogroup 1) and natural exotic (TgCatBR2 and GPHT: haplogroup 6) T. gondii isolates were used in these studies. Natural isolates were generously provided by L. David Sibley, PhD (Washington University, St Louis, MO, USA). Parasites were maintained in tachyzoite form by serial passage in confluent monolayers of human neonatal dermal fibroblasts (Cascade Biologics, Portland, OR, USA) in Dulbecco’s modified Eagle’s medium (Catalog number: 12100; Invitrogen-Gibco, Grand Island, NY, USA), supplemented with 44mM sodium bicarbonate and 1% heat-inactivated fetal bovine serum (FBS; Hyclone, Logan, UT, USA), at 37°C and 5% CO₂. Parasite viability was evaluated by plaque assay for every experiment; viability of 20% or greater was required, consistent with the previous description of the viability of a freshly egressed, high-passaged natural isolate from haplogroup 1.¹²

**Isolation and immortalization of human ocular endothelial cells**

Retinal and choroidal endothelial cells were obtained from non-diseased eyes of human cadaveric donors (Lions Vision Gift, Portland, OR, USA), adhering closely to previously published methods.¹⁴ In brief, retina or choroid were dissected from posterior eye cups and digested with graded solutions of ≤0.5 mg ml⁻¹ dispase (Invitrogen-Gibco) and ≤3 mg ml⁻¹ type II collagenase (Sigma-Aldrich, St Louis, MO, USA) in MCD-B-131 medium (Sigma-Aldrich) with 2% FBS. Endothelial cells were isolated from the digested...
tissue using magnetic Dynabeads (Dynal-Invitrogen, Oslo, Norway) coated with mouse monoclonal anti-human CD31 antibody (BD Pharmingen, San Diego, CA, USA), per the manufacturer’s instructions. Cells were cultured at 37°C and 5% CO₂ in MCDB-131 medium (Sigma-Aldrich) with 2–10% FBS and EGM-2 SingleQuots supplement, omitting gentamicin, hydrocortisone and FBS (Clonetics-Lonza, Walkerville, MD, USA).

To produce sufficient numbers for study, endothelial cells were immortalized by transduction with the murine recombinant amphotropic retrovirus, LXS16E6E7, which encodes for human papilloma virus E6 and E7 oncogenes, as well as a G418-resistance gene.18 LXS16E6E7 virus was kindly gifted by Denise A Galloway, PhD (Fred Hutchinson Cancer Institute, Seattle, WA, USA). Endothelial cells were cultured for 24h with LXS16E6E7 virus in supernatant harvested from PA317 packaging cell cultures. Percentage of immortalized cells was maximal by transduction in the presence of 5mgl⁻¹ hexadimethrine bromide (Sigma-Aldrich) (retinal endothelial cells), or by post-transduction exposure to 200μgml⁻¹ G418 antibiotic (Mediatech-Cellgro, Manassas, VA, USA) (choroidal endothelial cells). Following immortalization, cultures were purified by selection with anti-CD31 antibody-coated magnetic Dynabeads. Immortalized cells retained the endothelial phenotype as evidenced by: cobblestone morphology; expression of CD31 and von Willebrand factor; and capillary-like tube formation on provisional extracellular matrix (data not shown).

**Human ocular endothelial transmigration assay**

Polyethylene terephthalate transwell membranes (3μm pore-size, 0.3cm² diameter; BD Labware, Franklin Lakes, NJ, USA) were inserted into 24-well plates, coated with 30μg ml⁻¹ bovine type I collagen (Santa Cruz Biotechnology, Santa Cruz, CA, USA) for 2h at 37°C, and subsequently seeded with 30 000 endothelial cells suspended in MCDB-131 medium with 10% FBS and EGM-2 SingleQuots. Transwells were incubated at 37°C and 5% CO₂ for 4 to 5 days. One million freshly egressed live or heat-killed (that is, exposed to 55°C for 60min)19 tachyzoites, suspended in phenol red-free endothelial basal medium (Clonetics-Lonza) with 2.5% FBS and EGM-2 SingleQuots at 1/4 dilution, were loaded in upper chambers, and lower chambers were loaded with the same medium (n = 4–6 wells). After a 4h incubation at 37°C and 5% CO₂, parasites in lower chambers were counted by hemocytometer. In some experiments, upper chambers were pre-incubated for 2h with 10μg ml⁻¹ mouse anti-human ICAM-1 antibody (isotype IgG1; clone P2A4; Millipore, Temecula, CA, USA) or 30μg ml⁻¹ mouse anti-human VCAM-1 antibody (isotype IgG1; clone BBIG-V1; R&D Systems, Minneapolis, MN, USA), or control mouse IgG1 (clone 17117; R&D Systems) at the same concentration. Endothelial monolayer integrity was determined by permeability to 1mg ml⁻¹ Texas Red-conjugated 70 000MW dextran (Molecular Probes-Invitrogen, Eugene, OR, USA), per the method of Harhaj et al.20 For statistical analyses, the Student’s t-test was used to compare two groups and the ANOVA, with post-hoc comparisons. For all experiments, criteria for endothelial monolayer integrity were: (1) no significant difference (P ≥ 0.05) in dextran permeability of transwells containing endothelial monolayers and incubated with live versus heat-killed tachyzoites; (2) no significant increase (P ≥ 0.05) in permeability of transwells containing endothelial monolayers and incubated with live or heat-killed tachyzoites versus no tachyzoites; and (3) highly significant increase (P < 0.001) in permeability of transwells containing membranes coated with type I collagen alone versus wells containing endothelial monolayers and incubated with live, or heat-killed or no tachyzoites (n = 3–6 wells).

**Flow cytometric analysis of confluent human retinal endothelial cells**

Human retinal endothelial cells were plated for confluence in a 6-cm diameter dish. After 3 days, the cells were lifted with 0.05% trypsin, and surface-stained for 30min with V450-conjugated mouse anti-human CD31 antibody (isotype IgG1; clone WM59) and PE-conjugated anti-CD144 antibody (isotype IgG1; clone 55-7H1), or similarly labeled control mouse IgG1 (clone MOPC-21) (all obtained from BD Pharmingen) in phosphate buffered saline with 10% sodium azide and 10% FBS on ice. Subsequently, cells were washed with and suspended in the same buffer, and subjected to fluorescence activated cell sorting. Data were acquired on the BD LSR II flow cytometer (Becton-Dickinson, San Jose, CA, USA), and analyzed using FCS Express V3 (De Novo Software, Los Angeles, CA, USA).

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