Transmission of *Toxoplasma gondii* from Infected Dendritic Cells to Natural Killer Cells

Catrine M. Persson,¹‡ Henrik Lambert,¹,² Polya P. Vutova,¹,² Isabel Dellacasa-Lindberg,¹,² Joanna Nederby,¹ Hideo Yagita,³ Hans-Gustaf Ljunggren,¹ Alf Grandien,¹ Antonio Barragan,¹,²* and Benedict J. Chambers¹*

Center for Infectious Medicine, Department of Medicine, Karolinska Institutet, Karolinska University Hospital Huddinge, 141 86 Stockholm, Sweden; Department of Parasitology, Mycology and Environmental Microbiology, Swedish Institute for Infectious Disease Control, 171 82 Solna, Sweden; and Juntendo University School of Medicine, Tokyo 113-8421, Japan³

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The obligate intracellular parasite *Toxoplasma gondii* can actively infect any nucleated cell type, including cells from the immune system. In the present study, we observed that a large number of natural killer (NK) cells were infected by *T. gondii* early after intraperitoneal inoculation of parasites into C57BL/6 mice. Interestingly, one mechanism of NK cell infection involved NK cell-mediated targeting of infected dendritic cells (DC). Perforin-dependent killing of infected DC led to active egress of infectious parasites that rapidly infected adjacent effector NK cells. Infected NK cells were not efficiently targeted by other NK cells. These results suggest that rapid transfer of *T. gondii* from infected DC to effector NK cells may contribute to the parasite's sequestration and shielding from immune recognition shortly after infection.

*Toxoplasma gondii* causes chronic infections in up to one-third of the human population and in animals (22, 31). In healthy individuals, primary *T. gondii* infection causes relatively mild symptoms, whereas in the immunocompromised patient or in the developing fetus, life-threatening manifestations lead to severe neurological and ocular damage (11, 28, 37). Following oral infection, *T. gondii* parasites typically pass across restrictive biological barriers and rapidly disseminate (13). In this process, *T. gondii* actively infects a great variety of cell types, including epithelial cells and blood leukocytes (12, 21). In infected cells, the parasites establish nonfusigenic parasitophorous vacuoles, where they can replicate (27, 32, 38).

Natural killer (NK) cells and dendritic cells (DC) are two important cell types of the innate immune system. DC-NK cell interactions are important not only in host defense but also for the development of adaptive immune responses (5, 9). The activation of DC by pathogens leads to cytokine secretion, which activates NK cells, which, in turn, via cytokines or by direct cell-cell contact, may determine the adaptive immune responses that follow (9, 29). DC are sensitive to NK cell-mediated lysis in vitro and can be eliminated by NK cells in vivo (4, 6, 17, 19, 33, 43). Viral or bacterial infection of DC can reduce their sensitivity to NK cell-mediated lysis by increasing the expression of classical and nonclassical major histocompatibility complex class I molecules on the cell surface (14, 35, 43).

DC and NK cells play critical roles in innate immunity during acute *Toxoplasma* infection, being early sources of interleukin-12 (IL-12) and gamma interferon (IFN-γ), respectively (16, 20, 24, 34, 40). It has recently been suggested that infected DC, and possibly other leukocytes, can act as Trojan horses, potentiating the dissemination of the parasite from the point of infection to distal parts (8, 26). In the early phase of infection with *T. gondii*, NK cell recruitment to the site of infection is mediated by CCR5-binding chemokines (24). IFN-γ production by NK cells, induced by IL-12 from infected DC or macrophages, has been suggested to be the primary contribution of NK cells to the host defense against *T. gondii* (18, 25, 39). It can also drive cytotoxic CD8⁺ T-cell immunity to *T. gondii* even in the absence of CD4⁺ T cells (7). NK cells can also kill *T. gondii*-infected target cells (42), and perforin has been demonstrated to be important in protecting mice in the chronic stage of infection (10). In the present study, we investigated NK cell interactions with *T. gondii*-infected DC and, surprisingly, demonstrated how this interaction leads to *T. gondii* infection of NK cells.

Materials and Methods

Animals. C57BL/6 (B6), B6.recombination activating gene-1 (B6.RAG1)⁻/⁻, and B6.perforin (B6.pfp)⁻/⁻ mice (6 to 10 weeks old) were housed under standard conditions at the Department of Microbiology, Tumor and Cell Biology at the Karolinska Institutet and at the Karolinska University Hospital Huddinge, Stockholm, Sweden. All procedures were performed in conformance with both institutional and national guidelines.

Antibodies. Anti-FAS-L and anti-TRAIL monoclonal antibodies (mAbs) (41) were purified from cell culture media. Anti-FAS-L and anti-TRAIL Mabs were injected intraperitoneally (i.p.) at 500 μg/mouse 24 h prior to inoculation of parasites or adoptive transfer of parasite-infected DC. All labeled antibodies used for flow cytometry were obtained from Becton Dickinson (San Diego, CA).

Parasites and infection. Green fluorescent protein (GFP)-expressing type I RH-LDM (1) and type II PTG-GFP565T (31) *T. gondii* tachyzoites were main-
tained by serial 2-day passage in human foreskin fibroblast monolayers. Human foreskin fibroblasts were propagated in Dulbecco’s modified Eagle’s medium (Invitrogen, Paisley, United Kingdom) with 10% fetal calf serum (BioWhittaker, Verviers, Belgium), 20 μg/ml gentamicin, 2 mM glutamine, and 0.01 M HEPES (Invitrogen).

For infection of DC or NK cells in vitro, cells were harvested and incubated with freshly egressed GFP-expressing T. gondii tachyzoites at the indicated multiplicities of infection (MOI) for 16 to 24 h unless stated otherwise. Infection rates were assessed by flow cytometry by counting GFP cells. To inhibit parasite replication, 50 μM pyrimethamine (Sigma-Aldrich, Steinheim, Germany) was added to the cultures of GFP-expressing T. gondii tachyzoites and DC for the 16 to 24 h (30). Replication of parasites was assessed by flow cytometry and epifluorescence microscopy. Heat-killed parasites was generated as previously described (26).

Preparation of bone marrow-derived DC. Bone marrow-derived DC were generated as described previously (19). Briefly, bone marrow-derived cells were cultured in Dulbecco’s modified Eagle’s medium (Invitrogen) containing 10 ng/ml recombinant granulocyte-macrophage colony-stimulating factor (BioSource, Brussels, Belgium). The cells were harvested after 6 days and replated overnight. DC were further purified with anti-CD11c MAb-coated beads (Milteny Biotec, Bergisch Gladbach, Germany).

NK cell preparation. DX5+ cells from spleens of B6, B6.RAG1−/−, and B6.pfp−/− mice were purified by using the MACS separation system (Miltenyi Biotec) according to the manufacturer’s guidelines. Purified cells were resuspended in complete αMEM medium (10 mM HEPES, 2 × 10−3 M 2-mercaptoethanol, 10% fetal calf serum, 100 U/ml penicillin, 100 U/ml streptomycin) and cultured in 1,000 U recombinant IL-2 (Biosource)/ml for 6 days.

Cytotoxicity assays. Target cells (DC or NK cells) were incubated for 1 h in the presence of 51Cr (Amersham, Oxford, United Kingdom) and then washed thoroughly in phosphate-buffered saline (PBS). After 4 h of effector and target cell coinoculation, cell culture supernatants were taken from these wells and analyzed by using a gamma radiation counter (Wallac Oy, Turku, Finland). Specific lysis was calculated according to the following formula: % specific lysis = [(experimental release − spontaneous release)/(maximum release − spontaneous release)] × 100.

TABLE 1. Numbers of T. gondii-infected NK cells and T cells from the peritonea of infected mice

<table>
<thead>
<tr>
<th>Exptl conditions</th>
<th>No. of cells</th>
<th>Infected/uninfected cell ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Infected</td>
<td>Uninfected</td>
</tr>
<tr>
<td>Free T. gondii</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LDM NK cells b</td>
<td>1,425</td>
<td>29,055</td>
</tr>
<tr>
<td>T cells</td>
<td>1,197</td>
<td>181,224</td>
</tr>
<tr>
<td>PTG NK cells b</td>
<td>669</td>
<td>29,324</td>
</tr>
<tr>
<td>T cells</td>
<td>729</td>
<td>80,349</td>
</tr>
<tr>
<td>T. gondii-infected DC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LDM NK cells b</td>
<td>2,069</td>
<td>33,323</td>
</tr>
<tr>
<td>T cells</td>
<td>1,534</td>
<td>151,221</td>
</tr>
<tr>
<td>PTG NK cells b</td>
<td>2,419</td>
<td>21,197</td>
</tr>
<tr>
<td>T cells</td>
<td>1,558</td>
<td>60,099</td>
</tr>
</tbody>
</table>

a Total number of cells from six mice.

b Chi-square analysis comparing the total number of infected and uninfected NK cells with the total number of infected and T cells: P < 0.001.
prior to incubation with NK cells. For flow cytometry, cells were labeled with anti-NK1.1 and anti-CD11c MAbs. Dead cells were gated away by using propidium iodide.

**Ex vivo microscopy of infected lymphocytes.** For visualization of in vivo *Toxoplasma*-infected NK and T cells, DX5⁺ NK cells and CD3⁺ T cells were sorted from the peritoneal cavity with the MACS separation system and then seeded on glass coverslips coated with poly-L-lysine (Sigma-Aldrich). After 30 min at 37°C, the cells were washed once with BRB80 buffer [80 mM piperazine-N,N’-bis(2-ethanesulfonic acid) (PIPES), pH 6.9; 1 mM MgCl₂; 1 mM EGTA] and then fixed with 0.3% glutaraldehyde (TAAB Laboratories, Berkshire, United Kingdom) in BRB80 for 10 min at room temperature. Next, the cells were permeabilized with 0.1% Triton X-100 in PBS (PBST; Sigma, Steinheim, Germany) for 5 min at room temperature. Following a brief wash with PBS, pH 7.4, the coverslips were treated with 1 mg/ml sodium borohydride (Merck, Hohenbrunn, Germany) in PBS three times for 5 min each. The coverslips were then washed twice with PBST and incubated with phalloidin-Alexa 594 (Invitrogen, Carlsbad, CA) in PBST. Twenty minutes later, the coverslips were mounted with Vector Shield with 4',6'-diamidino-2-phenylindole (DAPI; Vector Laboratories, Burlingame, CA). Images were taken with a Leica DMRB microscope equipped with a QImaging Q20780 camera and processed with OpenLab software.

**Real-time confocal microscopy.** Consequences of NK cell interaction with infected DC were visualized with a spinning-disk confocal setup (Ultraview LCI-3 Tandem Scanning Unit; Perkin-Elmer, United Kingdom) on an Axiovert 200 M (Carl Zeiss, Germany) connected to a charge-coupled device camera (OrcaER; Hamamatsu, Japan). Cells were placed in a minichamber system (POCmini; LaCon, Germany) with a heating stage. Image acquisition and analysis of motility were performed with Openlab software (version 4.0.2) and Velocity software (Improvision Inc., United States).

**Statistical analysis.** Statistical analyses were performed with Prism Graph Version 4 (GraphPad Software Inc., La Jolla, CA).

**RESULTS**

**Infection of NK cells in mice inoculated i.p. with *T. gondii.*** Freshly egressed type I GFP⁺ RH-LDM *T. gondii* tachyzoites

FIG. 2. NK cell-mediated lysis of DC is enhanced upon infection with *T. gondii* and is dependent on perforin and live-parasite infection. (A) Lysis of DC infected with tachyzoites (MOI of 3) by NK cells from B6 and B6.pfp⁻/⁻ mice versus the effector-to-target cell ratio. The results of three separate experiments are shown ± the standard error of the mean. (B) Lysis of DC infected with tachyzoites, tachyzoites pretreated with pyrimethamine, or heat-killed tachyzoites versus the effector-to-target cell ratio. The results of three separate experiments are shown ± the standard error of the mean. (C) Lysis of tachyzoite-infected B6 NK cells and DC by uninfected B6 NK cells versus the effector-to-target cell ratio. The results of three separate experiments are shown ± the standard error of the mean.
were injected into the peritoneal cavities of B6 mice. After 48 h, approximately 30% of the CD11b<sup>+</sup> myeloid cells were infected (data not shown and reference 26). Surprisingly, when gating on the lymphocyte population, a significant number of NK cells were also infected with GFP<sup>+</sup> T. gondii tachyzoites. Among the total lymphocytes, the relative number of NK cells infected was significantly greater than that of infected T cells (Fig. 1A and Table 1). Interestingly, similar results were obtained when GFP<sup>+</sup> RH-LDM T. gondii tachyzoite-infected DC were injected into the peritoneal cavities of B6 mice (Fig. 1B and Table 1). Results were similar with type II GFP<sup>+</sup> PTG/ME49 T. gondii tachyzoites, both when inoculated as free parasites and in DC (Table 1). Ex vivo examination of the infected NK cells showed proliferating intracellular tachyzoites (Fig. 1C). Some NK cells had multiple vacuoles with replicating parasites (Fig. 1D). These data demonstrate that not only can myeloid cells, including macrophages and DC, become infected but also lymphocytes, including NK cells, following i.p. injection of T. gondii tachyzoites in B6 mice. During the 72-h period that we examined, the number of T cells in the peritoneal cavity remained relatively constant, whereas the number of NK cells increased, as previously observed by Khan et al. for NK cell recruitment into the spleen and liver (24).

**FIG. 3.** NK cells become infected following lysis of infected DC in vitro. (A) Lymphokine-activated killer cell cultures containing one NK cell (left side) to three T cells (right side) following 2 h of culture with GFP<sup>+</sup> tachyzoite-infected DC (MOI of 1). Bar graphs demonstrate the difference between infected NK cells and T cells. Accumulated data from three experiments are shown (P < 0.01 [Student’s t test]). (B) Infection of B6 (left side) and B6.pfp<sup>−/−</sup> (right side) NK cells following 2 h of culture with DC infected with GFP<sup>+</sup> tachyzoites (MOI of 1). One representative of three experiments is shown. The NK cell/DC ratio was 3:1. Bar graphs demonstrate the difference between infected B6 and B6.pfp<sup>−/−</sup> NK cells. Accumulated data from three separate experiments are shown (P < 0.01 [Student’s t test]).
20% of the wild-type NK cells had become infected, as determined by GFP expression (Fig. 3B, left side). In contrast, only a small proportion of NK cells from B6.pfp−/− mice were infected under similar conditions (Fig. 3B, right side). Upon treatment with the parasite egress-promoting agent dithiothreitol, a significant increase in the transfer of parasites from DC to NK cells from B6.pfp−/− mice was observed. Under these conditions, NK cell infection levels were almost in line with those of NK cells from B6 mice (data not shown). Altogether, these data suggest that perforin-dependent lysis of infected DC by NK cells leads to the egress of tachyzoites that then infect surrounding NK cells. The transfer of parasites from infected DC to NK cells was further analyzed by live imaging. In the resulting films, motile NK cells physically interacted with infected DC (Fig. 4A to C). Interaction between the smaller NK cells and larger DC for approximately 5 min was followed by (D and E) lysis of the infected DC and rapid egress of parasites (indicated by an asterisk). (F) Shortly after the parasites' egress, surrounding NK cells became infected by the GFP-expressing parasites (white arrows). Scale bar = 25 μm.

DISCUSSION

NK cells are one important component of the immune system involved in the control of T. gondii infection (16, 20, 24, 40). In the present study, we demonstrate that T. gondii tachyzoites are rapidly transferred to NK cells during infection. We show here that NK cell-mediated killing of infected DC leads to rapid egress of viable parasites, which can then infect effector NK cells, possibly enabling them to resist immune elimination.

CD4+ and CD8+ T cells are also important in the protection of the host from T. gondii infection (15, 16). However, we recently reported that T-cell-mediated cytotoxicity triggers rapid egress of parasites from their host cells in vitro and in vivo, an active process mediated by intracellular fluxes of Ca2+ induced by death signals from FAS-L, TRAIL, and perforin (36). Thus, primed but not naive CD8+ T cells could be infected by T. gondii upon interaction with infected cells (36). In the present study, we have focused on examining the first 24 to 72 h of infection, a time when the cytotoxic response is dominated by NK cells. This and the previous study raise new questions about the role of cell-mediated immunity in the establishment of acute and chronic Toxoplasma infections. From the pathogen's perspective, its rapid transfer from infected DC into NK cells is intriguing. It may be argued that
transmission of parasites from DC to NK cells contributes to the Toxoplasma parasite’s efficacy in establishing a primary infection while avoiding clearance as immune control mounts. Also, NK cells are not as well equipped to handle intracellular infections as antigen-presenting cells are, since NK cells do not possess intracellular killing pathways such as, e.g., nitric oxide. Since NK cells did not appear to target infected NK cells, NK cell infection may provide a reservoir in which the parasites proliferate. Thus, even though the rapid transfer of cell infection may provide a reservoir in which the parasites infections as antigen-presenting cells are, since NK cells do not possess intracellular killing pathways such as, e.g., nitric oxide. Since NK cells did not appear to target infected NK cells, NK cell infection may provide a reservoir in which the parasites proliferate. Thus, even though the rapid transfer of T. gondii from DC to NK cells may not mediate systemic dissemination per se, it may promote persistence of the parasite in a less hostile intracellular environment.

Additionally, NK cells are likely poorer at stimulating naïve T cells than are DC, since they lack high levels of the necessary costimulatory molecules (3). Therefore, T. gondii parasites may selectively recruit NK cells (24) and be strong activators of NK cells. This activation could lead to NK cell-mediated lysis of infected cells and production of IFN-γ that could eliminate the majority of the parasites. In the process, though, NK cells could become infected, thus creating a niche for the parasites. Therefore, parasites that have secluded themselves within NK cells could reach distant organs directly upon the migration of NK cells or indirectly upon the lysis of infected NK cells after parasite replication.

In terms of host defense, antibody responses to T. gondii may be more critical than generally appreciated in protecting the host by preventing cell-cell transmission of the parasite. In line with this hypothesis, B-cell-deficient mice survive past the early stage of infection by T. gondii but die 3 to 4 weeks postinfection (23). Therefore, the development of effective immunizations against the parasite may require the ability to evoke antibody-mediated responses to prevent chronic infection, since NK cell and T-cell cytotoxic responses may in fact aid the parasites’ survival, dissemination, and persistence.

This study demonstrates that T. gondii can use NK cells and potentially other lymphocytes to survive and multiply in the host. This may not be an isolated mechanism of immune evasion used by T. gondii. It has recently been shown that Neospora, a related apicomplexan parasite, also enhances the susceptibility of infected fibroblasts to NK cells (2) and can infect NK cells. However, we still need to determine if other pathogens similarly increase the NK cell sensitivity of targeted cells and, if so, provide an advantage for the persistence of infection. Two distinct possible mechanisms can be hypothesized by which pathogens can evade NK cell-mediated responses. Infections that promote the maturation of DC and NK cell resistance may use this strategy to bypass early elimination and thereby disseminate in the host. Additionally, pathogens that induce NK cell-activating ligands may take advantage of NK cell-mediated killing to continuously infect other cells and further the pathogen’s survival. In conclusion, the present data suggest a mechanism by which NK cells paradoxically may promote the dissemination of the parasite T. gondii.

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REFERENCES

ERRATUM

Transmission of *Toxoplasma gondii* from Infected Dendritic Cells to Natural Killer Cells


*Center for Infectious Medicine, Department of Medicine, Karolinska Institutet, Karolinska University Hospital Huddinge, 141 86 Stockholm, Sweden; Department of Parasitology, Mycology and Environmental Microbiology, Swedish Institute for Infectious Disease Control, 171 82 Solna, Sweden; and Juntendo University School of Medicine, Tokyo 113-8421, Japan*

Volume 77, no. 3, p. 970–976, 2009. Page 972: Figure 2C should appear as shown below, with the correct symbol labels.

![Graph showing the percentage of specific lysis](image-url)