Immunization with Live and Dead *Chlamydia muridarum* Induces Different Levels of Protective Immunity in a Murine Genital Tract Model: Correlation with MHC Class II Peptide Presentation and Multifunctional Th1 Cells

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Immunization with Live and Dead Chlamydia muridarum Induces Different Levels of Protective Immunity in a Murine Genital Tract Model: Correlation with MHC Class II Peptide Presentation and Multifunctional Th1 Cells

Hong Yu,* Karuna P. Karunakaran,* Isabelle Kelly,** Caixia Shen,* Xiaozhou Jiang,* Leonard J. Foster,† and Robert C. Brunham*  

Mice that were intranasally vaccinated with live or dead Chlamydia muridarum with or without CpG-containing oligodeoxynucleotide 1862 elicited widely disparate levels of protective immunity to genital tract challenge. We found that the frequency of multifunctional T cells coexpressing IFN-γ and TNF-α with or without IL-2 induced by live C. muridarum most accurately correlated with the pattern of protection against C. muridarum genital tract infection, suggesting that IFN-γ–producing CD4+ T cells that highly coexpress TNF-α may be the optimal effector cells for protective immunity. We also used an immunoproteomic approach to analyze MHC class II-bound peptides eluted from dendritic cells (DCs) that were pulsed with live or dead C. muridarum elementary bodies (EBs). We found that DCs pulsed with live EBs presented 45 MHC class II C. muridarum peptides mapping to 13 proteins. In contrast, DCs pulsed with dead EBs presented only six MHC class II C. muridarum peptides mapping to three proteins. Only two epitopes were shared in common between the live and dead EB-pulsed groups. This study provides insights into the role of Ag presentation and cytokine secretion patterns of CD4+ T effector cells that correlate with protective immunity elicited by live and dead C. muridarum. These insights should prove useful for improving vaccine design for Chlamydia trachomatis. The Journal of Immunology, 2011, 186: 3615–3621.
peptides presented by MHC molecules may differ when DCs are pulsed with dead and live EBs. The immune correlates for protection against *C. trachomatis* infection are known to be IFN-γ-mediated Th1 immune responses (20–22). However, other infection model systems have shown that using IFN-γ as a single immune correlate may not be sufficient to screen for protective immunity. Interestingly, this has been also observed for *C. muridarum* following adoptive transfer of CD4+ T cell clones (23). Flow cytometry allows assessment of individual cytokines simultaneously on a single-cell basis. Recent reports describing various disease targets in different animal models have demonstrated a correlation between protection and high-quality T cells that coexpress multiple cytokines (24–26). One of the effector cytokines shown in these studies mediating control of intracellular infection is TNF-α. TNF-α in combination with IFN-γ can synergize to mediate killing of pathogens (27). IL-2 has no direct effector function but strongly enhances the expansion of effector T cells. Therefore, it is of interest to use a cytokine set of no direct effector function but strongly enhances the expansion of dead EBs.

**Materials and Methods**

**Chlamydia**

*C. muridarum* mouse pneumonitis strain Nigg was grown in HeLa 229 cells in Eagle’s MEM (Invitrogen) supplemented with 10% FCS. Elementary bodies were purified by discontinuous density gradient centrifugation as previously described (28). Purified EBs were aliquoted and stored at –80°C in sucrose-phosphate-glutamic acid buffer and thawed immediately before use. The infectivity and the number of inclusion-forming units (IFU) of purified EBs were determined by immunostaining using anti-EB mouse polyclonal Ab followed by biotinylated anti-mouse IgG (Jackson ImmunoResearch Laboratories) and a diaminobenzidine substrate (Vector Laboratories) (29). Dead EBs were prepared either by heating to 56˚C for 30 min or by exposure to ultraviolet light from a G15T8 ultraviolet lamp (D. William Fuller, Chicago, IL) at a distance of 5 cm for 45 min at room temperature as previously described (8). Viability was tested on HeLa 229 cells to ensure complete inactivation of the heat- or ultraviolet-treated EBs. The IFU for both live and dead EBs were calculated from the titers determined on original stocks as described above.

**Mice**

Female C57BL/6 mice (5 to 6 wk old) were purchased from Charles River Canada and housed under pathogen-free conditions at the Animal Facility of the Jack Bell Research Center. All experiments were performed in strict accordance with University of British Columbia guidelines for animal care and use.

**Immunization**

Three groups of mice were immunized with 1) 1500 IFU live EBs intranasally once, 2) 5 × 10^7 IFU dead EBs plus 20 μg CptG-ODN 1826 (5′-TCCATGACGTCTCAGGTT-3′) phosphorothiolate modified; Integrated DNA Technologies) intranasally three times at 2-wk intervals, and 3) 5 × 10^7 IFU dead EBs intranasally three times at 2-wk intervals. A nonvaccinated group was included as a naive control.

**Genital tract infection challenge and C. muridarum quantification**

Five weeks after the live EB immunization and 1 wk after the last death EB immunization, mice were injected s.c. with 2.5 mg medroxyprogesterone acetate (Depo-Provera; Pharmacia and Upjohn). One week after medroxyprogesterone acetate treatment, mice were challenged intravaginally with 1500 IFU *C. muridarum*. Cervicovaginal washes were taken at day 5 postinfection and were stored at –80°C for *Chlamydia* titration on HeLa cells as described previously (29).

**Multiparameter flow cytometry**

Six weeks after the live EB immunization and 2 wk after the last death EB immunization or 1 wk after live EB challenge, mice from specified groups were sacrificed and cells harvested from spleen were stimulated with killed EBs (5 × 10^5 IFU/ml) in complete RPMI 1640 for 4 h at 37°C. Brefeldin A was added at a final concentration of 1 μg/ml, and cells were incubated for an additional 12 h before intracellular cytokine staining. Cells were surface stained for CD3, CD4, and CD8 as well as with the viability dye, red fluorescent reactive dye (L23102; Molecular Probes), followed by staining for IFN-γ, TNF-α, and IL-2 by using a BD Cytoperm Plus Fixation/Permeabilization (BD Pharmingen) kit. Finally, the cells were resuspended in a 4% formaldehyde solution. All Abs and all reagents for intracellular cytokine staining were purchased from BD Pharmingen except where noted. We collected 200,000 live splenocytes per sample by using a FACSARia flow cytometer (BD Biosciences) and analyzed the data by using FlowJo software (Tree Star).

**DC pulsing with live or dead EBs**

DCs were generated as previously described (30). Briefly, bone marrow cells were isolated from the femurs of C57BL/6 mice and cultured in Falcon petri dishes at 4 × 10^5 cells in 50 ml DC medium. DC medium was IMDM supplemented with 10% FCS, 0.5 mM 2-ME, 4 mM l-glutamine, 50 μg/ml gentamicin, and 5% of culture supernatant of murine GM-CSF–transfected plasmacytoma X63-Ag8 and 5% of culture supernatant of murine IL-4–transfected plasmacytoma X63-Ag8, which contained 10 ng/ml GM-CSF and 10 ng/ml IL-4, respectively. The above two cell lines were kindly provided by Dr. F. Melchers (Basilea Institute, Switzerland). At day 3, half of culture supernatants were removed and fresh DC medium was added. At day 5, nonadherent cells (purity of >50% CD11c+) were transferred to new dishes and cultured at 25 × 10^5 cells in 50 ml DC medium containing 25 × 10^5 IFU live EBs or dead EBs, respectively, at 37°C in 5% CO2 for 12 h. The cells pulsed with live EBs or dead EBs were then harvested and stored in –80°C.

**Identification of MHC class II-bound peptides**

We acquired 6 × 10^4 DCs pulsed with live EBs or dead EBs. The immunoproteomic approach to identify MHC class II-bound peptides from pulsed DCs involved multiple steps as previously described (18). Briefly, the pulsed DCs were lysed and MHC class II (I-Aβ) molecules were purified using allele-specific anti-MHC mAb affinity columns. MHC class II molecules bound to the affinity column were then eluted and the MHC-bound peptides were separated from MHC molecules by acetic acid treatment and ultrafiltration through a 5-kDa cutoff membrane to remove high molecular mass material. The purified MHC-bound peptides were analyzed qualitatively using an LTQ Orbitrap XL (Thermo Scientific) online coupled to a nanoflow HPLC using a nanospray ionization source. The mass spectrometer was set to fragment the five most intense multiply charged ions per cycle. Fragment spectra were extracted using DTASuperCharge (http://msquant.sourceforge.net) and searched using the Mascot algorithm against a database comprised of the protein sequences from *C. muridarum*.

**Statistical analysis**

Data were analyzed with the aid of the GraphPad Prism software program. The Kruskal–Wallis test was performed to analyze data for *C. muridarum* sheddings from multiple groups, and the Mann–Whitney U test was used to compare medians between pairs. Comparison of cytokine productions and mean fluorescence intensity (MFI) as determined by multiparameter flow cytometry between two groups were analyzed using two-tailed t test. A p value <0.05 was considered significant. Data are presented as means ± SEM.

**Results**

**Live and dead C. muridarum induce different levels of protection against C. muridarum genital tract infection**

We compared live to dead EBs with or without CptG-ODN 1862 using intranasal routes of vaccination. Six weeks after live EB or 2 wk after final dead EB immunization, we compared *Chlamydia* inclusion titers in cervicovaginal washes at day 6 following genital tract challenge with live EBs to measure protection. As shown in Fig. 1, excellent protection was observed in mice given live EBs intranasally. Compared to the nonvaccinated group, mice intranasally vaccinated with dead EBs formulated with CptG exhibited a partial reduction in *C. muridarum* titers in the vaginal wash (p < 0.01). Vaccination with dead EBs alone did not reduce...
cervicovaginal shedding, demonstrating that vaccination with dead EBs alone failed to induce any measurable protective immunity against *C. muridarum* infection.

**Functional characteristics of distinct populations of Th1 responses after live or dead EB immunization**

To define a correlation between characteristics of Th1 responses and protection, we used multiparameter flow cytometry and assessed Ag-specific IFN-γ, TNF-α, and IL-2–producing CD4+ T cells in mice following immunization before and after *C. muridarum* challenge. As shown in Fig. 2, among mice intranasally immunized with live EBs, a seven-color flow cytometry panel was used to simultaneously analyze multiple cytokines at a single cell level. A typical gating tree hierarchically identifies functional populations of CD4+ T cells based on the fluorescence staining of live cells, CD3+ cells, and IFN-γ-, TNF-α-, and IL-2–producing CD4+ T cells (Fig. 2A). CD4+ cells that express each cytokine (IFN-γ, TNF-α, and IL-2) are entered into a Boolean gating analysis that separately identifies the seven populations that express each possible combination (Fig. 2B).

To assess the characteristics of the Th1 responses in vaccinated mice, the total Ag-specific CD4+ T cell cytokine responses comprising IFN-γ, TNF-α, or IL-2 from splenocytes were measured in each of the vaccine groups, 6 wk after live EB or 2 wk after final dead EB immunization. Overall live EB intranasal immunization showed the highest number of IFN-γ-, TNF-α-, and IL-2–secreting cells per 10^6 splenocytes (Fig. 3A). The number per 10^6 splenocytes of IFN-γ/TNF-α double-positive CD4+ T cells was strikingly elevated in the live EB intranasally immunized group (Fig. 3B). The group immunized with dead EBs predominantly produced single-positive IFN-γ-producing CD4+ T cells (Fig. 3B). The nonvaccinated group (naive mice) showed very low background cytokine levels, indicating that cytokine-producing cells detected in the experimental system are *C. muridarum*-specific. Collectively, the data show that the total number of IFN-γ–producing CD4+ T cells is not strictly predictive of vaccine-elicted protection. Rather, protective efficiency is better correlated with cytokine coexpression profile of the CD4+ T cells.

Each cytokine-producing cells in this analysis encompasses four distinct populations. For IFN-γ, it includes IFN-γ single-positive, IFN-γ/TNF-α double-positive, IFN-γ/IL-2 double-positive, and IFN-γ/TNF-α/IL-2 triple-positive CD4+ T cells. The relative percentage of the four distinct populations can help define the quality of the Th1 responses. Differences in the quality of IFN-γ, TNF-α, and IL-2 responses between vaccine groups is represented pictorially by pie charts (Fig. 4A, 4C, 4E). By quantifying the fraction of the total IFN-γ responses, we observed that the combination of IFN-γ/TNF-α/IL-2 triple- and IFN-γ/TNF-α double-positive CD4+ T cells encompassed 77, 46, 6, and 0% of the total IFN-γ–producing cells in live EBs, dead EBs plus CpG, and dead EBs alone and naive groups, respectively (Fig. 4A). The MFI for IFN-γ, TNF-α, and IL-2 is an additional measure of cytokine production on a single cell basis (Fig. 4B, 4D, 4F). The MFI for IFN-γ and TNF-α–double-positive CD4+ T cells from the live EB vaccine group was significantly higher than that from the other vaccine groups (Fig. 4B). Analysis of TNF-α–producing CD4+ cells also showed a correlation between the number of multifunctional Th1 cells simultaneously secreting IFN-γ, as well as TNF-α and the degree of protection among the vaccine groups (Fig. 4C). Even more impressively, the MFI for TNF-α of IFN-γ/TNF-α double-positive CD4+ T cells from the live EB vaccine groups is ∼5-fold higher as compared with dead EB alone group (Fig. 4D). For IL-2–producing CD4+ cells, the fraction of IFN-γ/TNF-α/IL-2–double-positive CD4+ T cells was associated with protection (Fig. 4E). There was no difference in the correlation of MFI for IL-2 with protection (Fig. 4F).

**IFN-γ/TNF-α/IL-2 triple- and IFN-γ/TNF-α double-positive CD4+ cells correlate with protection against *C. muridarum* infection**

We next assessed the magnitude and quality of Th1 responses among the different vaccine-primed groups 7 d after the intranasal *C. muridarum* challenge. There was little difference between the live EB and dead EB plus CpG vaccine groups in the three cytokine-producing CD4+ T cells apart from the higher total number of TNF-α–producing CD4+ T cells observed in the live EB group (Fig. 5A). Interestingly, nonvaccine-primed mice also developed similar numbers of IFN-γ–, TNF-α–, and IL-2–producing CD4+ T cells at day 7 after *C. muridarum* challenge when compared with mice that were vaccine primed with live EBs or dead EBs plus CpG. Vaccine priming with dead EBs alone induced significantly lower numbers of IFN-γ and IL-2 compared with the other vaccine groups (Fig. 5A). Of importance, >90% of IFN-γ–producing CD4+ T cells in the live EB vaccine group simultaneously expressed TNF-α with or without IL-2, whereas most IFN-γ–producing CD4+ T cells in the nonprotected groups (dead EBs alone and naive control) did not coexpress TNF-α (Fig. 5B). Moreover, the MFI for TNF-α of IFN-γ/TNF-α/IL-2 triple- and IFN-γ/TNF-α double-positive CD4+ T cells in the live EB vaccine groups was much higher than that in the nonprotected groups (Fig. 5C). The data demonstrated that IFN-γ–producing CD4+ T cells that highly coexpress TNF-α were the most reliable correlate of immunity against *C. muridarum* infection and were predominantly induced by live but not dead EB immunization.

**Live EB-pulsed DCs present more *C. muridarum* MHC class II peptides compared with dead EB-pulsed DCs**

Because DCs are essential to induce *C. muridarum* cellular immunity via presentation of pathogen-specific Ags to naive T cells, we hypothesized that DCs exposed to live versus dead EBs may present distinct populations of peptides that may correlate with protection against *C. muridarum* infection. We used an immunoproteomic approach to identify *Chlamydia* MHC class II peptides eluted from DCs pulsed with live EBs or dead EBs. Eluted peptides were sequenced using mass spectrometry. We identified 45...
C. muridarum MHC class II-binding peptides from DCs pulsed with live EBs, but only 6 C. muridarum MHC class II-binding peptides from DCs pulsed with dead EBs (Table I). The 45 MHC class II-binding peptides identified in DCs pulsed with live EBs mapped to 13 distinct epitopes derived from 13 unique source proteins. The six MHC class II-binding peptides identified in DCs pulsed with dead EBs mapped to three distinct epitopes derived from three unique source proteins. The peptides derived from RplF and FabG were found in DCs pulsed with both live and dead EBs. There were 11 peptides, including PmpG and PmpE/F-2, found only in DCs pulsed with live EBs. Of note, our previous studies demonstrated that PmpG and PmpE/F-2 are immunodominant peptides/proteins that engender protective immunity against C. muridarum genital tract infection (19). We also compared the number of pathogen- and murine-derived MHC class II-binding peptides identified in DCs pulsed with live and dead EBs and found more peptides when the DCs were pulsed with live EBs. This was true for both C. muridarum and self-peptides and was more notable for C. muridarum-derived peptides than for self-peptides. The fold increase for pathogen-derived peptides from dead to live EB pulsing is 7.5 versus 2.2 for self-peptides (Table II). This suggests that DCs pulsed with live EBs present peptides via MHC more efficiently than do DCs pulsed with dead EBs, which in turn correlates with induction of better protective immunity by live EB immunization.

**Discussion**

Infection or immunization with viable intracellular pathogens typically induces immunological effects distinct from those elicited by inactivated organism (31). This also appears to be the case for Chlamydia (32, 33). For example vaccination of both humans and nonhuman primates with whole killed C. trachomatis resulted in limited protection with possible disease exacerbation upon re-infection (6, 7, 34). Strikingly, mice immunized with live C. muridarum developed complete protection (8, 35), whereas immunization with killed C. muridarum induced responses ranging from little to no protection (8), thereby providing a model system to investigate the underlying cellular and molecular basis for the failure of previous human and primate C. trachomatis vaccine trials. An understanding of differences in immune mechanisms...
elicited between live and dead organisms may serve to guide the future directions for *C. trachomatis* vaccine design.

There are two major findings reported from the present study. First, IFN-γ+–producing CD4+ T cells that highly coexpress TNF-α optimally correlated with protective immunity and were preferentially induced by live EB immunization. Second, exposure to live *C. muridarum* EBs induced DCs to present more pathogen-derived MHC class II peptides than did dead *C. muridarum* EBs.

CD4 Th1 effector cells clearly underpin *Chlamydia* immunity (13), and Igietseme et al. (23) originally demonstrated that adoptive transfer of a CD4 T cell clone specific to *C. muridarum*, which secreted both IFN-γ and TNF-α, resulted in resolution of chronic genital *C. muridarum* infection in nude mice. In contrast, a clone that produced IFN-γ alone was unable to resolve the infection in nude mice, even after 100 d, demonstrating that T cells that secrete multiple cytokines were substantially better at clearing infection than T cells with a limited cytokine profile. Building on these early promising results, we tested multiple cytokine secretion patterns at the single cell level using flow cytometry.

When using multiparameter flow cytometry in other pathogen studies, multifunctional CD4+ T cells that cosecrete IFN-γ, TNF-α, and IL-2 have been reported to better correlate with protection than IFN-γ single-positive secretion in *Leishmania major* (24) and *Mycobacterium tuberculosis* (25). Olsen et al. (36) also recently reported a high number of Ag-specific CD4+ T cells coexpressing IFN-γ and TNF-α in mice intranasally immunized with live *C. muridarum*, and in the current study, we demonstrate that IFN-γ+–producing CD4+ T cells that highly coexpress TNF-α were optimally correlated with protective immunity to *C. muridarum* genital tract challenge.

Multiple mechanisms could contribute to the enhanced protection to *Chlamydia* infection as mediated by multifunctional CD4+ T cells. First, multiple cytokines could have direct and synergistic antimicrobial action. In vitro studies have demonstrated that IFN-γ and TNF-α individually and synergistically in combination restrict intracellular *Chlamydia* growth (37, 38). Both IFN-γ and TNF-α activate inducible NO synthase in murine cells (39), and Robinson et al. (40) demonstrated that TNF-α syn-

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**FIGURE 4.** The IFN-γ, TNF-α, and IL-2 MFI and fraction of Ag-specific triple-, double-, and single-positive cells for each vaccine group. EB-specific cytokine productions from splenocytes of vaccinated mice before challenge were analyzed by multiparameter flow cytometry. The fraction of IFN-γ (A), TNF-α (C), and IL-2 (E) EB-specific triple-, double-, and single-positive cells for each vaccine group. The MFI of IFN-γ (B), TNF-α (D), and IL-2 (F) EB-specific triple-, double-, and single-positive cells for each vaccine group. Three or four mice were in each group. Shown are results representative of two experiments. D-EB, dead EBs; L-EB, live EBs.

**FIGURE 5.** The magnitude and quality of CD4+ T cell responses in vaccinated mice after *C. muridarum* challenge. EB-specific cytokine productions from splenocytes of vaccinated mice after challenge were analyzed by multiparameter flow cytometry. A, The total number of IFN-γ+, TNF-α+, or IL-2+–producing CD4 T cells in 10^6 splenocytes. B, The fraction of IFN-γ EB-specific triple-, double-, and single-positive cells for each vaccine group. Four mice were in each group. C, The MFI of TNF-α EB-specific triple-, double-, and single-positive cells for each vaccine group. Shown are results representative of two experiments. D-EB, dead EBs; L-EB, live EBs.
ergistically enhanced IDO activity induced by IFN-γ at the level of transcription in human epithelial cells. Second, multifunctional CD4+ T cells may better correlate with protective immunity because they produce more cytokine on a per-cell basis, as recently shown in a study of Leishmania major (24). In our study, we also observed higher MFI values for IFN-γ and TNF-α among triple- and double-positive cells. Third, both IFN-γ and TNF-α upregulate MHC class II and ICAM-1 expression on human cervical and vaginal epithelial cells (41). Greater production of IFN-γ and TNF-α by multifunctional CD4+ T cells could thereby enhance recognition of Chlamydia-infected epithelial cells and T cell activation via TCR/peptide/MHC class II and LFA/ICAM-1 pathways. It has been reported that physical contact between T cells and epithelial cells via LFA-1–ICAM-1 interactions are critical for epithelial NO production to chlamydiacidal levels (42). Lastly, Jayarapu et al. (43) reported that CD4+ Th1 cells restrict epithelial Chlamydia growth by both inducible NO synthase-dependent and degranulation-dependent mechanisms, and multifunctional CD4+ T cells may be superior to single cytokine-producing cells in their degranulation potential. For instance, Kannanganat et al. (44) reported that CMV-specific CD4 T cells underwent degranulation with ~50% of IFN-γ/TNF-α/IL-2 triple producers and IFN-γ/TNF-α double producers degranulating following Ag stimulation, compared with <15% of IFN-γ single producers. These data suggest that multifunctional CD4 T cells that coexpress IFN-γ and TNF-α preferentially degranulate compared with single IFN-γ producers and thus may possess better killing potential. In aggregate, the data suggest that vaccination that induces IFN-γ and TNF-α coexpressing CD4 T cells may represent the optimal correlate of protective immunity to C. trachomatis infection by one or more of these mechanisms.

Recently, we used an immunoproteomic approach to directly identify C. muridarum T cell peptides presented by APCs. Because DCs are essential to induce C. muridarum immunity via presentation of Ags to naïve T cells, we reasoned that live and dead organisms may induce presentation of different peptides. Accordingly, we compared the MHC class II-bound C. muridarum peptides identified in DCs 12 h after pulsing with live and dead organisms. We found that DCs pulsed with live EBs presented 45 MHC class II C. muridarum peptides that mapped to 13 proteins, including RplF, PmpG, and PmpE/F-2. These three C. muridarum T cell Ags have been previously identified as immunodominant. Moreover, vaccination with individual RplF, PmpG, or PmpE/F-2 recombinant proteins in mice engendered significant resistance to challenge infection in both the lung and genital tract models (19). In our study, we compared six MHC class II C. muridarum peptides that mapped to three proteins and included only one (RplF) of the three immunodominant C. muridarum Ags. Among the 15 epitopes presented by DCs after 12 h pulsing with live or dead EBs, 5 (RplF, FabG, PmpG, PmpE/F-2, and TC0420) were previously identified when DCs were pulsed with live organisms for 24 h (18). Thus 10 new C. muridarum T cell epitopes (TC0518, FusA, TC0884, Tuf, TC0654, TC0190, Tsf, PmpE/F-1, AtPE; Table I) were identified in the current study.

Table II. Comparison of Chlamydia- and murine-derived MHC class II-bound peptides identified in DCs after 12 h pulsing with live and dead EBs

<table>
<thead>
<tr>
<th>Peptide Sequence (C. muridarum)</th>
<th>Peptide Sequence (Mouse)</th>
<th>Source Protein</th>
<th>Abbreviation</th>
</tr>
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<tbody>
<tr>
<td>VKGNFEVSPAAHIDPORP (13)</td>
<td>Riboosomal protein L6</td>
<td>RplF</td>
<td></td>
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<tr>
<td>SPQGNYAAAKAGIIFPS (3)</td>
<td>3-Oxoacyl-(acyl carrier) protein reductase</td>
<td>FabG</td>
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</tr>
<tr>
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<td>Polymorphic membrane protein G</td>
<td>PmpG</td>
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</tr>
<tr>
<td>GAFPHLAPSAANYINNGS (5)</td>
<td>Polymorphic membrane protein F</td>
<td>PmpE/F-2</td>
<td></td>
</tr>
<tr>
<td>GRDLNVTGPKQTDVGL (7)</td>
<td>Hypothetical protein TC0420</td>
<td>TC0420</td>
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<tr>
<td>EOTKIPIGTPIAVFSTEQGN (4)</td>
<td>Pyruvate dehydrogenase</td>
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<td>FusA</td>
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<td>SRALYOQPLAISEA (1)</td>
<td>Polymorphic membrane protein E</td>
<td>PmpE/F-1</td>
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*Numbers in parentheses denote total number of peptides of varying length but consisting of the same core residues.*
In conclusion, this study demonstrates that the cytokine secretion pattern and frequency of multifunctional T cells coexpressing IFN-γ, TNF-α, and/or IL-2 induced by live or dead Chlamydia muridarum vaccines correlated with the degree of protection against genital tract infection. Furthermore, DCs pulsed with live EBs presented more Chlamydia MHC class II peptides compared with dead EB-pulsed DCs. These findings provide mechanisms to explain the disparate protective immunity responses elicited by live and dead Chlamydia muridarum and should help guide future vaccine development for Chlamydia trachomatis.

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Disclosures

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