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CD4+ T Cells Are Not Required for the Induction of Dengue Virus-Specific CD8+ T Cell or Antibody Responses but Contribute to Protection after Vaccination

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The contribution of T cells to the host response to dengue virus (DENV) infection is not well understood. We previously demonstrated a protective role for CD8+ T cells during primary DENV infection using a mouse-passaged DENV strain and IFN-α/βR−/− C57BL/6 mice, which are susceptible to DENV infection. In this study, we examine the role of CD4+ T cells during primary DENV infection. Four I-Ak-restricted epitopes derived from three of the nonstructural DENV proteins were identified. CD4+ T cells expanded and were activated after DENV infection, with peak activation occurring on day 7. The DENV-specific CD4+ T cells expressed intracellular IFN-γ, TNF, IL-2, and CD40L, and killed peptide-pulsed target cells in vivo. Surprisingly, depletion of CD4+ T cells before DENV infection had no effect on viral loads. Consistent with this observation, CD4+ T cell depletion did not affect the DENV-specific IgG or IgM Ab titers or their neutralizing activity, or the DENV-specific CD8+ T cell response. However, immunization with the CD4+ T cell epitopes before infection resulted in significantly lower viral loads. Thus, we conclude that whereas CD4+ T cells are not required for controlling primary DENV infection, their induction by immunization can contribute to viral clearance. These findings suggest inducing anti-DENV CD4+ T cell responses by vaccination may be beneficial. The Journal of Immunology, 2010, 185: 5405–5416.
CD4⁺ T cells can also induce chemokine production that attracts CD8⁺ T cells to sites of infection (19). However, the requirement for CD4⁺ T cell help for Ab and CD8⁺ T cell responses is not absolute, and may be specific to the pathogen and/or experimental system. For instance, it has been shown that CSR can occur in the absence of CD4⁺ T cells (20), and the primary CD8⁺ T cell response is CD4⁺-independent under inflammatory conditions (17).

Despite the known importance of CD4⁺ T cells in the host response to pathogens, to our knowledge no study has yet examined the role of CD4⁺ T cells during primary DENV infection, and no CD4⁺ T cell epitopes have been identified from DENV-infected mice. In this study, we sought to define the contribution of CD4⁺ T cells to the host response to primary DENV2 infection using IFN-α/βR⁻/⁻ mice. Infection with DENV2 resulted in CD4⁺ T cell expansion and activation. To study DENV-specific CD4⁺ T cells, a predictive algorithm was used to identify MHC class II (I-A²)-binding peptides from the entire proteome of DENV2, which is ~3390 aa and encodes three structural (core, envelope, and membrane), and seven nonstructural (NS; NS1, NS2A, NS2B, NS3, NS4A, NS4B, NS5) proteins. Four CD4⁺ T cell epitopes from the NS2B, NS3, and NS4B proteins were identified, one of which contains an immunodominant CD8⁺ T cell epitope that we identified previously (16).

DENV2-specific CD4⁺ T cells were of a Th1 phenotype, with intracellular expression of IFN-γ, TNF, IL-2, and CD40L, and could mediate in vivo cytotoxicity. However, depletion of CD4⁺ T cells did not have a significant effect on viral clearance, and CD4⁺ T cells were not required for the induction of the DENV2-specific Ab or CD8⁺ T cell responses. Immunization with dominant CD4⁺ T cell epitopes led to enhanced viral clearance, demonstrating that CD4⁺ T cells can contribute to the anti-DENV2 immune response, and supporting the development of a DENV vaccine that induces a robust CD4⁺ T cell response.

Materials and Methods

Mice and infections

C57BL/6 (H-2b) mice were obtained from The Jackson Laboratory (Bar Harbor, ME) and subsequently bred at the animal facility at La Jolla Institute for Allergy and Immunology. IFN-α/βR⁻/⁻ mice on the C57BL/6 background were obtained from W. Yokoyama (Washington University, St. Louis, MO) via C. Ware (La Jolla Institute for Allergy and Immunology). B6.SJL mice were purchased from Taconic Farms (Germantown, NY). Mice were between 5 and 8 wk of age. Mice were infected i.v. in the lateral tail vein or retro-orbitally with 200 μl DENV2 strain, S221, in 5% FBS/PBS. Blood was obtained from anesthetized mice by retro-orbital puncture. All mouse experiments were approved by the Animal Care Committee at La Jolla Institute for Allergy and Immunology.

Cell culture and viral stocks

The hybridoma clones SFR3, GK1.5, and 2.43, which produce rat anti-human HLA-DR5, anti-mouse CD4, and anti-mouse CD8 IgG2b Ab, respectively, were from American Type Culture Collection (Manassas, VA), and were grown in protein-free hybridoma medium supplemented with penicillin, streptomycin, HEPES, Glutamax, and 2-ME (all from Invitrogen, Carlsbad, CA) at 37°C, 5% CO₂, C6/36, an Aedes albopictus mosquito cell line, was cultured in Leibovitz’s L-15 medium (Invitrogen, Carlsbad, CA) supplemented with 10% FBS, and were grown in protein-free hybridoma medium supplemented with penicillin, streptomycin, and HEPES at 28°C. Aedes albopictus mosquito cell line was cultured in Leibovitz’s L-15 medium (Invitrogen, Carlsbad, CA) supplemented with 10% FBS, and were grown in protein-free hybridoma medium supplemented with penicillin, streptomycin, and HEPES at 28°C.
with DENV2 (10^6 GE/well) in 50 μl 0.1 M NaHCO₃. The virus was UV inactivated, and plates were left overnight at 4°C. The plates were then washed to remove unbound virus using 0.05% (v/v) Tween 20 (Sigma-Aldrich) in PBS. After blocking with blocker casein blocking buffer (Thermo Scientific) for 1 h at room temperature, 1:3 serial dilutions of serum in a total volume of 100 μl were added to the wells. After 1.5 h, wells were washed and bound Ab was detected using HRP-conjugated goat anti-mouse IgG Fc portion or HRP-conjugated donkey anti-mouse IgM μ-chain (Jackson ImmunoResearch Laboratories, West Grove, PA) and tetramethylbenzidine (eBioscience).

Ab-virus neutralization assay
Serum was heat inactivated at 56°C for 30 min. Three-fold serial dilutions of serum were then incubated with 5 × 10^6 GE of DENV2 for 1 h at room temperature in a total volume of 100 μl PBS. Next, ~5 × 10^6 CD3/36 cells/well of a 24-well plate were infected with 100 μl of the virus-Ab mix for 1 h at 28°C. Cells were washed twice with 500 μl PBS, and then incubated at 28°C in 500 μl L-15 medium containing 5% FBS, penicillin, and streptomycin for 24 h. For each Ab dilution, the percentage of infected cells was determined by flow cytometry, as previously described (25), using 2H2-­­-biotin (IG2a anti-pM/M, DENV1–4 reactive) and streptavidin allophycocyanin (BioLegend). The percentage of infected cells was normalized to 100% (infection without serum).

CD8 in vivo cytotoxicity assay
IFN-α/βR−/− mice (recipients) were infected with 10^10 GE of DENV2. Some mice were depleted of CD4+ T cells before infection. Splenocytes (targets) were harvested from donor B6.SJL congenic mice (CD45.1) 7 d after. RBCs were lysed, and the target cells were pulsed with varying concentrations of a pool of four H-2d–restricted DENV2 peptides (Mₖ 467–475, N₅₂ 164–172, N₅₅ 267–275, N₅₅ 336–343) or DMSO for 1 h at 37°C. The cells were then washed and labeled with CFSE (Invitrogen) in PBS/1% BSA for 10 min at 37°C. Cells were labeled with 1 μM CFSE (CFSEith) or 100 nM CFSE (CFSEith/0) or left unlabeled. After washing, the cell populations were mixed and 5 × 10^5 cells from each population were injected i.v. into naive or infected recipient mice. After 4 h, the mice were sacrificed and splenocytes were stained with anti-CD45.1 allophycocyanin (eBioscience) and analyzed by flow cytometry, gating on CD45.1+ cells. The percentage of killing was calculated as follows: 100 − [%percentage DENV peptide pulsed in infected mice]/%percentage DMSO pulsed in infected mice]/%percentage DENV peptide pulsed in naive mice]/%percentage DMSO pulsed in naive mice) × 100.

CD4 in vivo cytotoxicity assay
IFN-α/βR−/− mice (recipients) were infected with 10^10 GE of DENV2. Some mice were depleted of CD4+ or CD8+ cells before infection. Splenocytes (targets) were harvested from donor B6.SJL congenic mice (CD45.1) 7 d after. RBCs were lysed, and the target cells were pulsed with 1.7 μg (=1 μM) each of N₅₂ 164–172, N₅₅ 267–275, and N₅₅ 336–343 (or DMSO) for 1 h at 37°C. The cells were then washed and labeled with CFSE in PBS/0.1% BSA for 10 min at 37°C. DENV2 peptide-pulsed cells were labeled with 1 μM CFSE (CFSEith) or DMSO-pulsed cells with 100 nM CFSE (CFSEith%). After washing, the two cell populations were mixed and 5 × 10^6 cells from each population were injected i.v. into naive or infected recipient mice. After 16 h, the mice were sacrificed, splenocytes were stained, and the percentage of killing was calculated as described for the CD8 in vivo cytotoxicity assay.

Quantitation of DENV burden in mice
Mice were euthanized by isoflurane inhalation, and blood was collected via cardiac puncture. Serum was separated from whole blood by centrifugation in serum separator tubes (Starstedt, Newton, NC). Small intestines were immediately placed into RNAlater (Qiagen, Valencia, CA) or Qiagen TissueLyser. Immediately after homogenization, all tissues were centrifuged (5 min, 4°C, 16,000 X g) to pellet debris, and RNA was isolated using the RNeasy mini kit (Qiagen). Serum RNA was isolated using the QIAamp viral RNA mini kit (Qiagen). After elution, viral RNA was stored at −80°C. Quantitative RT-PCR was performed according to a published protocol (26), except a MyQ single-color real-time PCR detection system (Bio-Rad, Hercules, CA) with One-Step quantitative RT-PCR kit (Quanta BioSciences, Gaithersburg, MD) was used. The DENV2 standard curve was generated with serial dilutions of a known concentration of DENV2 genomic RNA, which was in vitro transcribed (MAXScript kit; Ambion, Austin, TX) from a plasmid containing the cDNA template of S221, 3′ untranslated region. After transcription, DNA was digested with DNase I, and RNA was purified using the RNeasy mini kit and quantified by spectrophotometry. To control for RNA quality and quantity when measuring DENV in tissues, the level of 18S rRNA was measured using 18S primers described previously (27) in parallel real-time RT-PCR reactions. A relative 18S standard curve was made from total splenic RNA.

Peptide immunizations
IFN-α/βR−/− mice were immunized i.v. with 50 μg each of NS2B 108–122, NS3 198–212, and NS3 237–251 emulsified in CFA (Difco, Detroit, MI). After 11 d, mice were boosted with 50 μg peptide emulsified in IFA (Difco). Mock-immunized mice received PBS/DMSO emulsified in CFA or IFA. Mice were infected 13 d after the boost with 10^11 GE of DENV2 (some mice were depleted of CD4+ or CD8+ T cells 3 and 1 d before infection). Four days later, the mice were sacrificed and tissues were harvested, RNA was isolated, and DENV2 RNA levels were measured as described above.

Statistical analyses
Data were analyzed with Prism software version 5.0 (GraphPad Software, Inc., San Diego, CA). Statistical significance was determined using unpaired t test with Welch’s correction.

Results
CD4+ T cell activation and expansion following DENV2 infection
DENV2 (10^10 GE of S221) infection of IFN-α/βR−/− mice results in an acute infection, with viral replication peaking between 2 and 4 d postinfection (16). At this time the mice show signs of disease, including hunched posture and ruffled fur, and the virus is subsequently cleared from the serum by day 6. To determine the role of CD4+ T cells during the course of this primary DENV2 infection, we first examined the expansion of CD4+ T cells in the spleens of IFN-α/βR−/− mice 7 d postinfection with DENV2, and observed a 2-fold increase in CD4+ T cell numbers (Fig. 1A). The cells were activated, as measured by CD44 upregulation and CD62L downregulation on splenic CD4+ T cells (Fig. 1B) and on circulating blood CD4+ T cells, with the peak on day 7 postinfection (Fig. 1C). To study the CD4+ T cell response in the spleen in more detail, we performed immunohistochemistry on spleen sections obtained from naive mice and mice 3, 5, and 7 d after DENV2 infection. Sections were stained for CD4, CD8, and B220 to highlight B cell follicles, and F4/80 to show red pulp macrophages. As expected, in naive mice, we observed CD4+ and CD8+ T cells dispersed throughout the spleen, but preferentially in T cell areas, also known as the paracortical lymphoid sheath (PALS) (Fig. 1D). By day 3 after DENV2 infection, most of the CD4+ and CD8+ T cells had migrated to the PALS, with very few T cells observed in the red pulp. At day 5, the CD4+ cells were still concentrated in the PALS, at the border between the T cell area and B cell follicles, and also in the B cell follicles. At day 7 postinfection, the spleen had increased in size dramatically, and CD4+ T cells were found primarily in the PALS and B cell follicles. The localization of CD8+ T cells differed from the CD4+ T cells mainly in that at day 5 postinfection, many of the CD8+ T cells had left the T cell area and were found distributed throughout the red pulp and marginal zone. At day 7, the CD8+ T cells were observed in the PALS, marginal zone, and also the red pulp. These images illustrate the kinetics of the adaptive immune response to DENV2 in the spleen, and show CD4+ T cells in close proximity to both CD8+ T cells and B cells after DENV2 infection.

Tregs are a subset of CD4+ T cells that are characterized by the expression of the transcription factor, Foxp3 (28), and have been found to facilitate the early host response to HSV-2 (29) and help
control WNV infection (30). To determine whether DENV2 infection results in expansion of Tregs, we examined the number of CD4+Foxp3+ cells in the spleen 7 d postinfection, and found there was a decrease in the percentage of Tregs among total CD4+ cells, and no change in the number of Tregs, demonstrating that DENV2 infection does not lead to an expansion of Tregs in the spleen (Fig. 1).

Identification of DENV2 CD4+ T cell epitopes

To study the DENV2-specific CD4+ T cell response, we sought to identify MHC class II (I-$\alpha$)-restricted CD4+ T cell epitopes using a bioinformatics prediction method that has been previously used to map the CD4+ T cell response to mouse CMV (31). Briefly, the proteome of DENV2 was screened and 73 15-mer peptides predicted to bind I-$\alpha$ were identified. The peptides were tested by IFN-$\gamma$ ICS using splenocytes from DENV2-infected I-$\alpha$/βR−/− mice. Positive peptides (2× background) were then reordered as HPLC-purified (≥90%) and retested. Four positive peptides were identified, as follows: NS2B 108–122, NS3 198–212, NS3 237–251, and NS4B 96–110 (Fig. 2A, Table I). Similar to the DENV2-specific CD8+ T cell response (16), the epitopes identified in I-$\alpha$/βR−/− mice were also recognized by CD4+ T cells from DENV2-infected wild-type mice (Fig. 2B), and the magnitude of the CD4+ T cell response was higher in I-$\alpha$/βR−/− mice compared with wild-type mice, most likely due to increased viral replication. Notably, NS3 200–214 has been identified as a human HLA-DR15–restricted CD4+ T cell epitope (14, 32). It was also of interest that NS4B 96–110 contains a CD8+ T cell epitope.
epitope (NS4B99–107) that we had previously identified as the immunodominant epitope in both wild-type and IFN-α/βR^2/2^ C57BL/6 mice infected with DENV2 (16).

**Phenotype of DENV2-specific CD4^+^ T cells**

Multicolor flow cytometry was performed to study the phenotype of DENV2-specific CD4^+^ T cells. These cells produced IFN-γ, TNF, and IL-2 (Fig. 3). No intracellular IL-4, IL-5, IL-17, or IL-10 was detected (data not shown). The DENV2-specific CD4^+^ T cells also expressed CD40L, suggesting they are capable of activating CD40-expressing cells, which include dendritic cells (DCs) and B cells. The four DENV2-derived CD4^+^ T cell epitopes induced responses that differed in magnitude, but were similar in terms of phenotype. The most polyfunctional cells (those expressing IFN-γ, TNF, IL-2, and CD40L) also expressed the highest levels of the cytokines and CD40L. These results demonstrate that DENV2 infection elicits a virus-specific Th1 CD4^+^ T cell response in IFN-α/βR^−/−^ mice.

**Effects of CD4^+^ and/or CD8^+^ T cell depletions on DENV2 viral RNA levels**

To determine how CD4^+^ T cells contribute to controlling DENV2 infection, we depleted CD4^+^ T cells, CD8^+^ T cells, or both from IFN-α/βR^−/−^ mice and measured DENV2 RNA levels 5 d postinfection with 10^{10} GE of DENV2. We found no difference in viral RNA levels between control undepleted mice and CD4-depleted mice in the serum, kidney, small intestine, spleen, or brain (Fig. 4). As we observed previously (16), CD8-depleted mice had significantly higher viral loads than control mice. Depletion of both CD4^+^ and CD8^+^ T cells resulted in viral RNA levels that were significantly higher than those in control mice in all tissues examined, and equivalent to the viral RNA levels in CD8-depleted mice. These data show that CD4^+^ T cells are not required to control primary DENV2 infection in IFN-α/βR^−/−^ mice, and confirm an important role for CD8^+^ T cells in viral clearance.

**CD4^+^ T cells are not required for the anti-DENV2 Ab response**

Although CD4^+^ T cells were not required for controlling DENV2 infection, we wondered whether they made any contribution to the anti-DENV immune response, for example, by helping the B cell and/or CD8^+^ T cell responses. CSR, the process by which the Ig H chain C region is switched so the B cell expresses a new isotype of Ab, can be induced when CD40L-expressing CD4^+^ T cells engage

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**FIGURE 2.** Identification of DENV2-derived epitopes recognized by CD4^+^ T cells. A. Splenocytes were obtained from IFN-α/βR^−/−^ mice 7 d postinfection with 10^{10} GE of DENV2 and restimulated in vitro with DENV2-derived 15-mer peptides predicted to bind I-A^b^ cells. Cells were then stained for surface CD4 and intracellular IFN-γ and analyzed by flow cytometry. The four positive peptides identified are shown. In the dot plots, the percentage of CD4^+^ T cells producing IFN-γ is indicated. The responses of individual mice as well as the mean and SEM are also shown (n = 7–11). The response of unstimulated cells was subtracted from the response to each DENV2 peptide, and the net percentage and number of splenic CD4^+^ T cells producing IFN-γ are indicated. B. Splenocytes were obtained from wild-type C57BL/6 mice 7 d postinfection with 10^{10} GE of DENV2 and stimulated and stained as in A (n = 6).

**Table I.** DENV2-derived CD4^+^ T cell epitopes

<table>
<thead>
<tr>
<th>Epitope</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>NS2B108–122</td>
<td>GLFPVSLPITAAAWY</td>
</tr>
<tr>
<td>NS3198–212</td>
<td>GKTTRKLPAIVRRAI</td>
</tr>
<tr>
<td>NS3237–251</td>
<td>GLAPIRYQTPAIRAEH</td>
</tr>
<tr>
<td>NS4B96–110</td>
<td>IGCSQVNPILTA</td>
</tr>
</tbody>
</table>

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CD40 on B cells (20). However, CSR can also occur in the absence of CD4+ T cell help. To determine whether the anti-DENV2 Ab response depends on CD4+ T cells, we first measured DENV2-specific IgM and IgG titers in the sera of control and CD4-depleted mice 7 d postinfection with 10¹⁰ GE of DENV2. As expected, we found no difference in IgM titers at day 7 between control and CD4-depleted mice (Fig. 5A). There was also no difference in IgG titers between control and CD4-depleted mice.

To measure the functionality of these DENV2-specific Abs, a flow cytometry-based neutralization assay was performed, in which C6/36 mosquito cells were infected with DENV2 in the presence of heat-inactivated sera obtained from control and CD4-depleted mice (Fig. 5A). There was also no difference in IgG titers between control and CD4-depleted mice. To measure the functionality of these DENV2-specific Abs, a flow cytometry-based neutralization assay was performed, in which C6/36 mosquito cells were infected with DENV2 in the presence of heat-inactivated sera obtained from control and CD4-depleted mice 7 d postinfection. The sera from control and CD4-depleted mice could neutralize DENV2 equally well (Fig. 5B). As reported previously (8), naive serum was able to prevent DENV infection of C6/36 cells, although not as efficiently as DENV-immune serum. We also looked for the presence of GC B cells, as the GC reaction is generally thought to be CD4+ T cell dependent (33). As expected, we found GC B cells were absent in the CD4-depleted mice (Fig. 5C). Based on the lack of GC B cells in the DENV2-infected CD4-depleted mice, we conclude that the early anti-DENV2 Ab response is CD4- and GC-independent.

**CD4+ T cells are not necessary for the primary DENV2-specific CD8+ T cell response**

Next, we assessed the role of CD4+ T cells in helping the CD8+ T cell response by examining the DENV2-specific CD8+ T cell response in control and CD4-depleted DENV2-infected mice. The numbers of splenic CD8+ T cells were equivalent in control and CD4-depleted mice (data not shown). IFN-γ ICS was performed...
using DENV2-derived H-2b–restricted immunodominant peptides that we identified previously (M60-67, NS2A8-15, and NS4B99-107) (16). Somewhat surprisingly, there was an increase in the number of DENV2-specific IFN-γ CD8+ T cells in CD4-depleted mice compared with control mice (Fig. 6A). To further characterize the phenotype of the CD8+ T cells generated in the absence of CD4+ T cells, we examined expression of TNF, IL-2, and CD107a (a marker for degranulation) in cells stimulated with NS4B99-107 (Fig. 6B, 6C). As also shown in Fig. 6A, the magnitude of the CD8+ T cell response was larger in the CD4-depleted mice, but the cytokine and CD107a expression profiles were comparable. Similar results were observed when cells were stimulated with M60-67 or NS2A8-15 (data not shown). Next, we examined the functionality of the DENV2-specific CD8+ T cells in an in vivo cytotoxicity assay, in which splenocytes were pulsed with a pool of four H-2b–restricted immunodominant peptides and CFSE labeled before injection into control or CD4-depleted DENV2-infected mice. CD8+ T cell-mediated cytotoxicity was very efficient; almost 100% killing was observed at peptide concentrations of 500 ng/ml (Fig. 6D). Therefore, the peptide concentrations were titrated down, and no difference in killing was observed between control and CD4-depleted mice at any concentration tested. These data reveal that the primary anti-DENV2 CD8+ T cell response, in terms of expansion, cytokine production, degranulation, and cytotoxicity, does not depend on CD4+ T cell help.

In vivo killing of I-Ab-restricted peptide-pulsed target cells in DENV2-infected mice

Although we found that the absence of CD4+ T cells had no effect on viral RNA levels on day 5 postinfection, it was possible that CD4+ T cells could still be contributing to the anti-DENV2 host response by killing infected cells. We therefore performed an in vivo cytotoxicity assay using splenocytes pulsed with the three peptides that contain only CD4+ T cell epitopes (NS2B108-122, NS3198-212, and NS3237-251) and not NS4B96-110, as we wanted to measure only CD4+ T cell-mediated killing. Approximately 30% killing of target cells was observed (Fig. 7). No cytolytic activity was observed when CD4+ T cells were depleted, whereas depletion of CD8+ T cells had no effect on killing, demonstrating that the cytotoxicity was CD4+ T cell mediated. Thus, DENV2-specific CD4+ T cells can exhibit in vivo cytolytic activity, although this effector function does not appear to significantly contribute to controlling primary DENV2 infection.

**Vaccination with DENV2 CD4+ T cell epitopes helps control viral load**

Having found that DENV2-specific CD4+ T cells can kill target cells, we next assessed whether immunization with CD4+ T cell epitopes could help control DENV2 infection. Mice were immunized with NS2B108-122, NS3198-212, and NS3237-251, before DENV2 infection, and we measured CD4+ T cell responses by ICS and viral RNA levels 4 d postinfection. Peptide immunization resulted in enhanced CD4+ T cell cytokine responses (data not shown), and significantly lower viral loads in the kidney and spleen (Fig. 8). The protective effect was mediated by CD4+ T cells, as CD4 depletion before infection abrogated the protective effect. Similarly, CD8 depletion resulted in no protection, demonstrating that protection after CD4+ T cell peptide immunization requires both CD4+ and CD8+ T cells. These data suggest that CD4+ T cells elicited by immunization protect by helping the CD8+ T cell response. Thus, although CD4+ T cells are not required for the primary CD8+ T cell or Ab response, and the...
absence of CD4+ T cells had no effect on viral RNA levels, vaccination with CD4+ T cell epitopes can reduce viral loads.

Discussion

Numerous studies have investigated the phenotype of DENV serotype cross-reactive T cells, which have been hypothesized to contribute to the pathogenesis of secondary heterologous infections, yet the actual contribution of T cells during DENV infection is unknown. The findings presented in this study, and in our previous study, reveal that CD8+ T cells play an important protective role in the response to primary DENV2 infection, whereas CD4+ T cells do not. CD4+ T cells expanded, were activated, and were located near CD8+ T cells and B cells in the spleen after DENV2 infection, yet they did not seem to affect the induction of the DENV2-specific CD8+ T cell or Ab responses. In fact, CD4+ T cell depletion had no effect on viral clearance. However, our data demonstrate that vaccination with CD4+ T cell epitopes prior to DENV infection can provide significant protection, supporting T cell peptide vaccination as a strategy for DENV immunization without the risk of ADE.

To the best of our knowledge, this is the first study to identify CD4+ T cell epitopes from DENV-infected mice. We found the DENV2-specific CD4+ T cells recognized epitopes from the NS2B, NS3, and NS4B proteins, and displayed a Th1 phenotype. CD4+ T cell epitopes have been identified in mice infected with other flaviviruses, including yellow fever virus, for which an I-A^b–restricted peptide from the E protein was identified (34), and WNV, for which six epitopes from the E and NS3 proteins were identified (35). DENV-derived epitopes recognized by human CD4+ T cells have been identified, primarily from NS proteins, including the highly conserved NS3 (10). One study identified numerous epitopes from the NS3_200-325 region, and alignment of consensus sequences for DENV1–4 revealed that this region is more conserved (78%) than NS3 as a whole (68%) (14), sug-
FIGURE 7. Cytotoxicity mediated by DENV2-specific CD4+ T cells. In vivo killing of DENV2-derived I-A\textsuperscript{b}-restricted peptide-pulsed cells. IFN-\alpha/\betaR\textsuperscript{-/-} mice (control, CD4-depleted, or CD8-depleted) infected 7 d previously with 10\textsuperscript{10} GE of DENV2 were injected i.v. with CFSE-labeled target cells pulsed with the three epitopes that contain only CD4\textsuperscript{+} T cell epitopes (NS2B\textsubscript{108-122}, NS3\textsubscript{198-212}, and NS\textsubscript{237-251} (n = 6 control, 3 CD4-depleted, and 3 CD8-depleted mice). After 16 h, splenocytes were harvested and analyzed by flow cytometry, and the percentage of killing was calculated.

FIGURE 8. Peptide immunization with CD4\textsuperscript{+} T cell epitopes results in enhanced DENV2 clearance. IFN-\alpha/\betaR\textsuperscript{-/-} mice were immunized s.c. with 50 \mu g each of the three DENV peptides that contain only CD4\textsuperscript{+} T cell epitopes (NS2B\textsubscript{108-122}, NS3\textsubscript{198-212}, NS\textsubscript{237-251}) in CFA, or mock immunized with DMSO in CFA. Mice were boosted 11 d later with peptide in CFA, then challenged with 10\textsuperscript{10} GE of DENV2 13 d later, and sacrificed 4 d postinfection. Separate groups of peptide-immunized mice were depleted of CD4\textsuperscript{+} or CD8\textsuperscript{+} T cells prior to infection. DENV2 RNA levels in the tissues were quantified by real-time RT-PCR and are expressed as DENV2 U normalized to 18S RNA. Each symbol represents one mouse, and the bar represents the geometric mean. \(p < 0.05; \ast \ast p < 0.01\).

gusting that the region contains good candidates for a DENV T cell epitope-based vaccine. Interestingly, one of the NS3-derived epitopes we identified has also been described as a human CD4\textsuperscript{+} T cell epitope. This species cross-reactive NS3 peptide may bind human HLAs promiscuously, making it a good potential vaccine candidate. Another intriguing finding was that one of the CD4\textsuperscript{+} T cell epitopes identified in this study contained the most immunodominant of the CD8\textsuperscript{+} T cell epitopes we had identified previously. Overlapping epitopes have also been found in lymphocytic choriomeningitis virus (LCMV) (36–38). The significance of overlapping epitopes is unknown, but is most likely related to homology between MHC class I and MHC class II, and may be associated with proteasomal processing. Overlapping epitopes may turn out to be common once the complete CD4\textsuperscript{+} and CD8\textsuperscript{+} T cell responses to other pathogens are mapped.

CD4\textsuperscript{+} T cells are classically defined as helper cells, as they help B cell and CD8\textsuperscript{+} T cell responses. However, inflammatory stimuli can override the need for CD4\textsuperscript{+} T cell help, and therefore, the responses to many acute infections are CD4-independent (17). DENV2 replicates to high levels in IFN-\alpha/\betaR\textsuperscript{-/-} mice, the mice appear hunched and ruffled at the time of peak viremia, and they have intestinal inflammation (T. Prestwood, unpublished observation). DENV2 replication depends on the virus and experimental system. The lack of GC B cells in CD4-depleted mice shows that the DENV2-specific IgG response at day 7 is CD4-independent. The goal of our study was to determine how CD4-independent CD8\textsuperscript{+} T cell responses have been demonstrated for vesicular stomatitis virus (42), rotavirus (43), and poliovirus (44). In addition, EBV (via latent membrane protein 1) can induce CD40-independent CSR (45), and mice deficient for CD40 or CD4\textsuperscript{+} T cells are able to mount an influenza-specific IgG response that is protective (46).

Similar to the studies mentioned above, our results demonstrate that the DENV2-specific IgG response at day 7 is CD4-independent. The lack of GC B cells in CD4-depleted mice shows that our CD4 depletions have a functional effect, and indicate anti-DENV IgG is being produced by extrafollicular B cells. It is possible that the absence of CD4\textsuperscript{+} T cells would have an effect on DENV2-specific Ab titers and/or neutralizing activity at later time points; however, the goal of our study was to determine how CD4\textsuperscript{+} T cells contribute to clearance of primary DENV2 infection, and we clearly show that the early anti-DENV2 Ab response is CD4-independent. The mechanisms by which T-independent Ab responses are induced are beginning to be elucidated. B lymphocyte stimulator, also known as B cell-activating factor, and a proliferation-inducing ligand are induced on DCs after activation with IFN-\alpha, IFN-\gamma, LPS, or CD40L, and mediate CD40-independent CSR in human B cells (47). B cell-activating factor and a proliferation-inducing ligand also induce CSR in mouse B cells in the absence of CD40 (48). T-independent CSR can also be induced via TLR signaling. For example, T-independent CSR to human papillomavirus-like particles occurs via TLR4 and MyD88 in mouse B cells (49), and TLR3 or TLR9 signaling can induce CSR in human B cells (50, 51). The pathways involved in mediating the CD4-independent CSR in DENV2-infected mice remain to be determined.

Like pathogen-specific Ab responses, the primary CD8\textsuperscript{+} T cell responses to many acute infections are also CD4-independent. CD4-independent CD8\textsuperscript{+} T cell responses have been demonstrated for Listeria monocytogenes (52, 53), LCMV (54), and influenza (55). Recently, a mechanism for how DCs can activate CD8\textsuperscript{+} T cells in the absence of CD4\textsuperscript{+} T cell help has been described. DCs activated with TLR3 or TLR9 agonists, or by influenza infection, upregulated CD40L and activated naïve CD8\textsuperscript{+} T cells in the absence of CD4\textsuperscript{+} T cells (56). In accordance with the studies
mentioned above, we found that the primary CD8+ T cell response to DENV2 did not depend on CD4+ T cells. In fact, we observed an enhanced DENV2-specific CD8+ T cell response in CD4-deficient mice compared with control mice at day 7, which has also been reported for influenza (55)- and WNV (40)-specific CD8+ T cell responses. In our study, this could be due to the depletion of Tregs, or an increased availability of cytokines (e.g., IL-2) in mice lacking CD4+ T cells. This enhanced CD8+ T cell response may explain why CD4-depleted mice have no differences in viral titers despite the fact that DENV2-specific CD4+ T cells demonstrate in vivo cytotoxicity.

Although CD4+ T cells did not play an important role in helping the Ab or CD8+ T cell responses, we found DENV2-specific CD4+ T cells could kill peptide-pulsed target cells in vivo. CD4+ T cells specific for other pathogens, including HIV (57) and influenza (58), demonstrate in vitro cytotoxicity. In vivo cytotoxicity assays have been used to show CD4+ T cell-mediated killing following infection with LCMV (59) and WNV (35). DENV-specific cytolytic human CD4+ T cell clones (60, 61) and a mouse (H-2d) CD4+ T cell clone (62) have been reported. Whether CD4+ T cells actually kill infected cells during DENV infection remains to be determined, but is possible, as MHC class II-expressing macrophages are targets of DENV infection (8). Based on the fact that CD4 depletion did not have a significant effect on viral clearance, it is unlikely that CD4+ T cell-mediated killing plays a major role in the anti-DENV2 response in this model.

An obvious caveat to using the IFN-α/βR−/− mice is that type I IFNs are known to help T cell and B cell responses through their actions on DCs, and can act directly on T cells (63). Type I IFNs were found to contribute to the expansion of CD4+ T cells following infection with LCMV, but not L. monocytogenes (64). Type I IFNs can induce the development of Th1 IFN-γ responses in human CD4+ T cells, but cannot substitute for IL-12 in promoting Th1 responses in mouse CD4+ T cells (65). Following Listeria infection, IL-12 synergized with type I IFN to induce IFN-γ production by CD4+ T cells (66). Although DENV does not replicate to detectable levels in wild-type mice, examining the CD4+ T cell response in these mice revealed that the same epitopes were recognized as in the IFN-α/βR−/− mice, but the magnitude of the epitope-specific response was greater in the IFN-α/βR−/− mice. This suggests that the high levels of viral replication in the IFN-α/βR−/− mice are sufficient to drive a DENV2-specific CD4+ IFN-γ response. Our results demonstrate a DENV2-specific CD4+ T cell response, including Th1-type cytokine production and cytotoxicity, in the absence of IFN-α/βR signaling, yet this response is not required for clearance of infection. It is, however, possible that CD4+ T cells contribute to protection during DENV infection of hosts with intact IFN responses.

We have previously shown that immunization with CD8+ T cell epitopes resulted in enhanced viral clearance (16), and the results presented in this study show that immunization with CD4+ T epitopes is also protective. Peptide immunization with CD4+ T cell epitopes has also been found to protect against WNV (35) and HSV-1 (67). Our results have significant implications for DENV vaccine development. Although a DENV vaccine is a global public health priority, designing a vaccine has been challenging, as it needs to induce protection against all four serotypes. The DENV vaccine candidates in development, some of which are in phase II trials, focus on eliciting an Ab response. The challenge is to induce and maintain a neutralizing Ab response against all four serotypes, as it is becoming increasingly clear that non-neutralizing Ab (or subneutralizing quantities of Ab) can actually enhance dengue disease (8, 9). A different approach would be a peptide vaccine that induces cell-mediated immunity, including both CD4+ and CD8+ T cell responses, which would not be able to prevent infection, but could reduce viral loads and disease severity, and would eliminate the risk of ADE. Such a vaccine should target highly conserved regions of the proteome, for example, NS3, NS4B, and/or NS5, and ideally include epitopes conserved across all four serotypes. A vaccine containing only peptides from these particular NS proteins would also preclude the induction of any Ab against epitopes on the virion, which could enhance infection, or Ab against NS1, which could potentially contribute to pathogenesis (68). Our peptide vaccination was given along with CFA, which is commonly used in mice to induce Th1 responses (69), which was the type of response observed after natural DENV infection. CFA is not a vaccine adjuvant approved for human use, and thus, any peptide vaccine developed against DENV will have to be formulated with an adjuvant that is approved for human use and promotes a Th1 response. Numerous vaccine adjuvants are currently under development and can be tested in our experimental system in the future.

Although our results indicate CD4+ T cells do not make a significant contribution to controlling primary DENV2 infection, the characterization of the primary CD4+ T cell response and epitope identification will allow us to determine the role of CD4+ T cells during secondary homologous and heterologous infections. CD4+ T cells are often dispensable for the primary CD8+ T cell response to infection, but have been shown to be required for the maintenance of memory CD8+ T cell responses after acute infection (70). Finally, our findings support a DENV vaccine strategy that induces CD4+ T cell, in addition to CD8+ T cell, responses.

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