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Protection against Vaccinia Virus Challenge by CD8 Memory T Cells Resolved by Molecular Mimicry

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Live vaccinia virus (VV) vaccination has been highly successful in eradicating smallpox. However, the mechanisms of immunity involved in mediating this protective effect are still poorly understood, and the roles of CD8 T-cell responses in primary and secondary VV infections are not clearly identified. By applying the concept of molecular mimicry to identify potential CD8 T-cell epitopes that stimulate cross-reactive T cells specific to lymphocytic choriomeningitis virus (LCMV) and VV, we identified after screening only 115 peptides two VV-specific immunogenic epitopes that mediated protective immunity against VV. An immunodominant epitope, VV-e7r192, did not generate cross-reactive T-cell responses to LCMV, and a subdominant epitope, VV-a11r189, did generate cross-reactive responses to LCMV. Infection with VV induced strong epitope-specific responses which were stable into long-term memory and peaked at the time virus was cleared, consistent with CD8 T cells assisting in the control of VV. Two different approaches, direct adoptive transfer of VV-e7r-specific CD8 T cells and prior immunization with a VV-e7r-expressing ubiquitinated minigene, demonstrated that memory CD8 T cells alone could play a significant role in protective immunity against VV. These studies suggest that exploiting cross-reactive responses between viruses may be a useful tool to complement existing technology in predicting immunogenic epitopes to large viruses, such as VV, leading to a better understanding of the role CD8 T cells play during these viral infections.

Prior immunity to unrelated pathogens can sometimes significantly alter the course and outcome of unrelated virus infections in mice, and this can be either beneficial or detrimental to the host (43, 47, 57, 58). For instance, immunity to lymphocytic choriomeningitis virus (LCMV) can protect mice from a lethal dose of vaccinia virus (VV), and these mice may show altered T-cell-mediated immunopathology resulting in bronchiolitis obliterans or acute fatty necrosis (8, 46). Heterologous immunity may play a role in the variability observed in human disease outcome, from subclinical to lethal, in individuals with the same infection or vaccination.

In 1980, the World Health Organization announced that smallpox (variola) had been eradicated by vaccination with VV; however, smallpox is now considered a potential bioterrorist agent (23). The use of modified VV for mixed-modality vaccines to boost immune responses that have been primed with other agents, such as DNA vaccines or adenovirus vectors expressing viral epitopes, is now being advocated (1). Concerns have arisen about the safety of VV as a vaccine. Adverse events, such as fulminant disseminated vaccinia, have occurred in immunodeficient individuals, while many significant immunity-mediated conditions, such as severe dermatological diseases (erythema multiforme and erythema nodosum), arthritis, pericarditis, myocarditis, and encephalitis, have occurred in healthy individuals following VV vaccination against smallpox (7, 19, 32, 36). Heterologous immunity may be playing a role in mediating some of this immunopathology, especially in adults, who would have large complex memory pools following a lifetime of infections. Thus, studies on heterologous immunity in the mouse model may be highly relevant to human disease, since vaccination with VV has again become prevalent.

Despite the great success of this live vaccine, the precise mechanisms of immunity associated with protection are still poorly understood. There has been evidence for both cellular immunity and humoral immunity playing a role. The level of serum-neutralizing antibody has been correlated with protective immunity. However, the observation that T-cell-deficient individuals had serious and at times fatal infections following VV immunization while agammaglobulinemic children did not suffer such complications suggests that cellular immunity plays an important role in clearing the infection (18, 36, 37, 39). It has been shown that both arms of the immune response are complementary in mediating protection against ectromelia virus (mouse pox), a poxvirus closely related to VV. CD8 T-cell responses were essential for clearing ectromelia virus early in a primary infection, while antibodies were important later in ectromelia virus infection (15, 25). The importance of CD8 T-cell responses during both primary and secondary VV infections is less clear. Previous studies using VV infection of mice have had equivocal results concerning the role of CD8 T cells, although most studies are consistent with CD8 T cells playing at least a supportive role in protective immunity (3, 13, 49, 50, 59). Ongoing research is focusing on developing new and safer vaccines (14). In order to achieve this, it is important to have a better understanding of the impact of T-cell responses on VV infection, including their roles both in protective immunity and in mediating immunopathology.

VV infection of LCMV-immune mice leads to the activation...
of T cells specific to many different LCMV epitopes. LCMV-NP\textsubscript{205}–specific CD8 T cells expanded most frequently, presumably due to cross-reactivity with VV (27). Interestingly, LCMV-NP\textsubscript{205}–specific CD8 T-cell responses are also cross-reactive with a similar epitope encoded by Pichinde virus (PV), PV-NP\textsubscript{205}. Six of eight amino acids are common between the LCMV and PV epitopes in sites important for interaction with the T-cell receptor (TCR) (5). CD8 T-cell responses may be cross-reactive with different antigens (5, 10, 27, 56, 58), and a common model for cross-reactivity is molecular mimicry, in which two peptides have sequence similarity at the sites of TCR recognition (16, 17). We used the concept of molecular mimicry as a premise to identify potential cross-reactive epitopes between LCMV and VV. This has led to the identification of two VV-specific CD8 T-cell epitopes in mice. Both of these epitopes induced highly effective VV-specific acute protection of two VV-specific CD8 T-cell epitopes in mice. Both the other epitope, VV-e7r\textsubscript{130}, which is described by using the concept of molecular mimicry. We here identify a matrix of cross-reactive responses involving five different epitopes and three different viruses (M. Cornberg, S. C. Clute, F. M. Saccoccio, S. K. Kim, Y. N. Naumov, M. A. Brehm, R. M. Welsh, and L. K. Selin, submitted for publication).

Interestingly, the other epitope, VV-e7r\textsubscript{130}, is described in more detail here, did not appear to activate cross-reactive LCMV-specific memory T-cell responses, even though it was identified by using the concept of molecular mimicry. We here demonstrate that, by use of VV-e7r\textsubscript{130}–ubiquitinated minigenec immunization or adoptive transfer of VV-e7r\textsubscript{130}–specific CD8 T-cell lines, memory CD8 T cells can play a significant role in mediating protective immunity to VV during a secondary infection.

MATERIALS AND METHODS

Mice. C57BL/6 (B6; H-2\textsuperscript{b}) male mice were purchased from The Jackson Laboratory (Bar Harbor, ME), and B6.S/J-ptprca (L5.1) congenic male mice were purchased from Tacsonic Farms (Germantown, NY). Mice were used at 2 to 8 months of age. All mice were maintained under specific-pathogen-free conditions in the University of Massachusetts Medical School, Department of Animal Medicine.

Cell lines. ATCC Vero cells, used in plaque assays, were cultured in minimum essential medium (MEM; Invitrogen, Carlsbad, CA). MC57G cells, which are an H-2\textsuperscript{a}-expressing melanomatous-derived fibroblast cell line from B6 mice, were used as stimulators in intracellular cytokine staining (ICS) assays or as targets for \textsuperscript{51}Cr release cytotoxicity assays and were maintained in MEM. MC57G cells were infected with VV at a multiplicity of infection of 10 PFU/cell and incubated for 1 h at 37°C. The TAP-2-deficient B6-derived T-lymphoma cell line, RMA-S, kindly provided by Hans-Gustaf Ljunggren (Karolinska Institute, Stockholm, Sweden), was grown in RPMI. RMA-S cells were pulsed with 1 \textmu M peptide for 1 h at 37°C and then were used as targets in \textsuperscript{51}Cr release cytotoxicity assays. As stimulators for CD8 T-cell lines, RMA-S cells were incubated with 1 \textmu M peptide for 1 h at 37°C and then irradiated (3,000 rads). All cell lines were supplemented with 100 U/mL penicillin G, 100 \textmu g/mL streptomycin sulfate, 2 mM l-glutamine, 10 mM HEPES, and 10% heat-inactivated (50°C, 30 min) fetal bovine serum (FBS; Sigma Chemical Co., St. Louis, MO).

Viruses. The Western Reserve (WR) strain of VV, a DNA virus in the orthopoxvirus family, was propagated in L929 cells (45). LCMV (Armstrong strain), an RNA virus in the Old World arenavirus family, was propagated in BHK-21 baby hamster kidney cells (45). The mouse adapted influenza virus A/PR/8/34 (H1N1), an RNA virus in the orthomyxovirus family, was grown in the allantoic fluid of 10-day-old embryonated chicken eggs (SPAFA, Preston, CT) (9).

Infection protocols. To assess virus infections, mice were infected intraperitoneally (i.p.) with 5 \times 10\textsuperscript{4} PFU of LCMV or 10\textsuperscript{4} PFU VV and methoxyflurane (Metofane)-anesthetized mice were challenged intranasally with 70 PFU of influenza A virus. To control for culture contaminants, VV stocks were purified through a sucrose gradient and diluted in Hanks’ balanced salt solution (HBSS; Invitrogen, Carlsbad, CA) and LCMV was diluted more than 40-fold in HBSS. Mice were considered immune 6 weeks after infection or later. Control naive mice were either left uninoculated or inoculated with tissue culture media or HBSS. The control mice were always age matched to mice of the experimental group and housed exactly under the same pathogen-free conditions.

Virus titration. VV titers in each of the organs (fat pads, testes, spleens, lungs, kidneys, brains, salivary glands, hearts, and livers) were determined by plaque assays on ATCC Vero cells with the use of a 10% homogenate of tissue taken from individual mice, as described elsewhere (46).

Identification and screening for potential cross-reactive VV epitopes. In order to identify potential VV epitopes cross-reactive with the H-2\textsuperscript{k}-restricted LCMV-NP\textsubscript{205} epitope (YTVKYPNL) (55), we searched the VV sequence for 8-mers which maintained the H-2\textsuperscript{k}-binding motif. Two H-2\textsuperscript{k}-restricted epitopes were identified, one in the VV protein e7r (positions 130 to 137 (STLNFNNL), and the second in the VV protein a11r, positions 88 to 95 (CLSNLQFNL)). For controls, the influenza virus epitope NP\textsubscript{436}\textsubscript{D} (ASNEENMETL) (52) and the LCMV epitope NP\textsubscript{396}\textsubscript{D} (FQPQNGTML) (20) were used.

Cell surface and tetramer staining by flow cytometry. Single cell suspensions were prepared from spleenocytes, peritoneal exudates (PECs), or peripheral blood. Erythrocytes were lysed with 0.84% NH\textsubscript{4}Cl solution. Cell suspensions were incubated in fluorescence-activated cell sorter buffer (phosphate-buffered saline [PBS] containing 2% FBS and 0.2% sodium azide) with 1 \textmu M tetramer (Clone 53-6.7) and fluorescein isothiocyanate–anti-mouse CD44 (Clone IM7). For control, the influenza virus epitope NP\textsubscript{436}\textsubscript{D} (ASNEENMETL) (52) and the LCMV epitope NP\textsubscript{396}D (FQPQNGTML) (20) were used.

Cytotoxicity assays. Standard in vitro chromium (\textsuperscript{51}Cr) release assays were performed to measure antiviral cytotoxic-T-lymphocyte (CTL) activity as described previously (45). The in vivo cytotoxicity assay was performed according to recently published techniques (2, 28), with splenocytes from B6 or congenic L5.1 mice used as target cells. After the lysis of red blood cells, splenocytes were divided into two or four populations. One population was pulsed with 1 \textmu M ovalbumin amino acids 257 to 264 (SIINFEKL) as a control, and other populations were pulsed with 1 \textmu M of the indicated VV peptides for 60 min at 37°C. Each population was labeled with a different concentration (2 \textmu M or 0.4 \textmu M) for the two-population experiment; 2 \textmu M, 0.67 \textmu M, 0.33 \textmu M, or 0.11 \textmu M for the
four-population experiment) of carboxyfluorescein diacetate-succinimidy ester (CFSE; Molecular Probes, Eugene, OR). After CFSE labeling, equal amounts of cells were mixed together, washed, and resuspended in PBS. A total of $2 \times 10^7$ cells were injected intravenously into each recipient mouse. Specific in vivo cytotoxicity was determined by harvesting splenocytes from recipient mice 16 h after intravenous cell transfer. CFSE-labeled target populations were quantitated by flow cytometry. When splenocytes from LYS.1 congenic mice were used as target cells, splenocytes from recipients were contained with a PE-conjugated LYS.1-specific monoclonal antibody (CD45.1, clone A20; BD Pharmingen). By gating on LYS.1-specific cells, it was possible to include four target populations per sample. Uninfected, LCMV- and VV-infected C57BL/6 mice were used as recipients. The amount of specific in vivo killing was calculated, as described elsewhere (2), as follows:

$$100 - \left( \frac{\text{[(% peptide pulsed in infected/% unpulsed in infected)/(% peptide pulsed in uninfected/% unpulsed in uninfected) \times 100]}{}} \right).$$

In vitro expansion of antigen-specific CTLs. Splenocytes ($10^7$) from VV- or LCMV-immune mice were cocultured with peptide-pulsed RMA-S cells ($10^6$) in RPMI supplemented with 100 U/ml penicillin G, 100 μg/ml streptomycin sulfate, 2 mM l-glutamine, 10 mM HEPES, 1 mM sodium pyruvate, 0.1 mM MEM-nonessential amino acids, 0.05 mM β-mercaptoethanol, and 10% FBS for 4 to 5 days at 37°C at 5% CO2. The IL-2 culture supplement BD T-Stim (BD Biosciences, San Diego, CA) was added after 4 to 5 days of culture. Peptide stimulation was repeated every 4 to 5 days. After 20 to 25 days of stimulation (four or five stimulations), T-cell lines were analyzed.

Adoptive transfer of antigen-specific T cells into mice. T-cell lines were restimulated without peptide stimulation for 7 days. Thereafter, live cells were separated with Lympholyte (Cedarlane, Hornby, Canada). Cells were suspended in PBS at $10^7$ cells/ml and incubated with 2 μM CFSE for 15 min at 37°C. After incubation, cells were washed twice with HBSS, and $10^6$ cells were injected i.p. into recipient mice. Mice were infected with 10 PFU VV i.p. on the same day. Three to four days after infection, mice were sacrificed. Fat pads and/or testes were analyzed for VV titers. PECs were analyzed for immune responses and division of transferred T cells. Surface staining was performed as described above, with fluorochrome-labeled antibodies peridinin chlorophyll protein–anti-mouse anti-CD45.2 (LY5.2, clone 104) and APC–anti-mouse CD8α (clone 53-6.7). Transferred donor cells were identified as positive for CD45.2 and CD8 when congenic mice were used.

VV-e7r130-specific minigene DNA immunization. Ubiquitinated DNA vaccine constructs expressing VV-e7r130 epitope were prepared as described elsewhere (42). The genes for VV-e7r130 were cloned into the F3Ub expression vector kindly provided by Lindsay Whitton (Scripps Institute, San Diego, CA). Clean lipopoly saccharide-free plasmid DNA was produced by using plasmid Giga kit (QIAGEN, Valencia, CA), according to manufacturer’s protocol. For immunization, the quadriceps muscles of B6 mice were injected with 100 μg (50 μg for each muscle) of DNA construct formulated in sterile 0.9% NaCl. The control mice were immunized with 100 μg of F3Ub vector or LCMV-NP396-specific minigenome or were not immunized. Mice received three DNA immunizations (100 μg each) separated by two weeks. Two weeks after the last immunization, mice were infected with 10^6 PFU VV. Six days after VV infection, mice were sacrificed. Splenocytes and PECs were analyzed by intracellular cytokine staining and by in vitro 51Cr release assays. Fat pads were evaluated for virus titers by plaque assays.

Statistical analysis. Descriptive statistics are expressed as mean values ± standard errors of the mean. Comparisons between groups were performed with the Student t test (two tailed).

RESULTS

Cross-reactive peptide motif identifies H2-Kk-restricted VV-specific CD8 T-cell epitopes. Defining VV-specific epitopes has not been a priority until recent times, as smallpox had been essentially eradicated. Identifying epitopes for VV can be a daunting task, as VV is a very large DNA virus of the orthopoxvirus family, and expresses over 200 proteins, with the potential for thousands of CD8 T-cell epitopes (21). Human and mouse VV-specific CD8 T-cell epitopes have only very recently been identified (13, 33, 35, 51, 53). We took a unique approach to identifying VV-specific epitopes because our previous work had revealed selective expansion of LCMV epitope-specific T cells during VV infection, consistent with potential cross-reactive T-cell responses (8, 26, 27). In order to further examine cross-reactive responses between VV and LCMV, we had to define VV-specific epitopes. Since CD8 T cells specific for the LCMV-NP396 peptide often expanded during VV infection of LCMV-immune mice, this peptide was used as a template peptide sequence to identify potential cross-reactive epitopes with sequence similarity. By use of a general DNA/RNA and protein analysis software (DNASIS) and the criteria that the epitope should have an H-2Kk binding motif and greater than 30% sequence similarity to LCMV-NP396 (YTVKYKNPL), 115 8-mer peptides were identified in the VV genome. By screening these peptides by use of an intracellular IFN-γ assay with VV- or LCMV-specific CD8 T cells, we identified two VV-specific epitopes, e7r130 (STLNFNNL) and a11r198 (AIVNYANL) (Fig. 1A). The IFN-γ responses to VV-e7r130 ranged between 1% and 9.9% (mean ± standard error of the mean, 2.2% ± 0.5%; n = 22) of all CD8 T cells six days after VV infection, whereas the IFN-γ response to VV-a11r198 ranged between 0.1% and 0.75% (mean ± standard error of the mean, 0.23% ± 0.05%; n = 17). Six days after VV infection, the VV-e7r130 and VV-a11r198 IFN-γ responses accounted for approximately 10% and 1% of the anti-CD3 response, respectively. The VV-e7r130 and VV-a11r198 peptide-specific T-cell responses accounted for approximately 20% and 2%, respectively, of the total VV-specific response as determined by stimulation with the MHC class I-matched VV-infected MC57G cells. In vivo CTL assays using peptide-pulsed splenocytes from C57BL/6 mice as targets showed about 90% specific killing (mean, 87%; range, 83 to 92%; n = 6) for VV-e7r130 and 19% (range, 15 to 23%; n = 3) for VV-a11r198 (Fig. 1B). In vitro 51Cr release assays on syngeneic target cells (RMA-S) coated with peptides, although not as sensitive, showed a similar hierarchy (data not shown). These two newly identified VV epitopes were also found to have the same relative binding affinity for H-2Kk as LCMV-NP205; the epitope used to identify them (Fig. 1C). We consider VV-e7r130 an immunodominant H-2Kk-restricted epitope for VV, while VV-a11r198 is a subdominant epitope in primary VV infection.

When these peptides were tested in IFN-γ or CTL assays with splenocytes from mice infected with only LCMV 9 days earlier, VV-a11r198-induced significant IFN-γ production in up to 1% of all CD8 T cells (mean, 0.8%; range, 0.4 to 1%; n = 5) and in vivo killing was more than 80% (86%; range, 78 to 94%; n = 6), whereas VV-e7r130 failed to stimulate CD8 T cells to produce IFN-γ or demonstrate cytotoxic activity (Fig. 1A and B). Although VV-e7r130 was identified by its potential to be cross-reactive with LCMV, cells from LCMV-infected mice did not appear to recognize this epitope.

Newly identified VV CD8 T-cell epitopes e7r130 and a11r198 are maintained in the T-cell memory pool. To analyze if the VV epitope-specific immune response could be maintained in the CD8 T-cell memory pool, we analyzed VV-infected C57BL/6 mice during the acute phase of infection and more than six weeks after infection. VV-e7r130 and VV-a11r198 responses in the spleen peaked at 6 days after VV infection as measured by IFN-γ production upon in vitro peptide stimulation (for e7r, a mean ± standard deviation of 2.2% ± 0.5% and a range of 1 to 9.9% [n = 22]; for a11r, a mean ± standard deviation of 0.23% ± 0.05% and a range of 0.1% to 0.75% [n = 17]) (Fig. 1A and 2A and B). CD8 T-cell responses specific
to either epitope gradually declined over the first 30 days after VV infection and stabilized thereafter, as demonstrated by little change in their total number or frequencies between day 30 and 200 postinfection (Fig. 2A and B). Memory CD8 T cells specific for VV-e7r130 frequencies were stable at 0.4% (range, 0.2 to 0.6%; n = 11) and for VV-a11r198 at 0.05% ± 0.02% (range, 0.01 to 0.1%; n = 8) for more than 200 days postinfection (Fig. 2A and B). The responses against the VV epitopes were also detectable in peripheral sites, such as the peritoneal cavity, the initial site of virus entry, at slightly higher frequencies than in the spleen (Fig. 2A, row ii). Tetramer staining with H-2Kb-containing VV-e7r130 allowed direct visualization of antigen specific CD8 T cells without in vitro stimulation and showed results similar to those determined by intracellular cytokine assays (Fig. 2A, row iii).

Analyzing intracellular cytokine production for both IFN-γ and TNF-α during the acute and memory phases of VV infection demonstrated that VV-specific CD8 T cells could produce both cytokines in response to VV-e7r130 and VV-a11r198. The determined ratio ([IFN-γ + TNF-α] /[IFN-γ + TNF-α-]) was 0.5 at the peak of the VV-specific CD8 T-cell response (day 6) and greater than 2 after 15 days after VV infection. This is consistent with the previously demonstrated concept that mature memory CD8 T cells are capable of producing both cytokines (Fig. 2A, row iv) (48).

Immunodominant VV-e7r130-specific CD8 T cells elicit effector function during the acute phase of VV infection. During the acute phase of VV infection, VV-e7r130-specific CD8 T-cell cytotoxic activity paralleled cytokine production. In vivo killing of VV-e7r130-labeled targets was 87% (range, 83 to 92%; n = 6) 6 days after VV infection and declined to 34% (range, 10 to 46%; n = 5) and 18% (range, 16 to 22%; n = 5) 8 and 15 days,

FIG. 1. VV- or LCMV-specific CD8 T cells recognize the subdominant VV-a11r198 epitope, while only VV-specific CD8 T cells recognize the dominant VV-e7r130 epitope. (A) Intracellular IFN-γ assay. Splenocytes from naïve or day 6 VV- or day 8 LCMV-infected C57BL/6 mice were stimulated as indicated in an ICS assay. The percentage of CD8 T cells producing IFN-γ is recorded in the upper left quadrant (gated on CD8 cells). Data are representative of five experiments (two to six mice/group). No stim, no stimulation. (B) In vivo cytotoxicity assay. CTL activity was analyzed in naïve or day 6 VV- or day 8 LCMV-infected C57BL/6 mice using CFSE-labeled targets coated with LCMV-NP205 (a), VV-a11r198 (b), VV-e7r130 (c), or control ovalbumin SIINFEKL (d) peptide. Data are representative of two experiments (three to five mice/group). (C) RMA-S stabilization assay. The newly identified VV epitopes stabilize H-2Kb on RMA-S cells in a standard stabilization assay. The mean fluorescence intensities (MFI) of the H-2Kb expression to different concentrations of the indicated peptides are shown. Data are representative of two similar experiments.
respectively, after VV infection (Fig. 2C). Staining for total CD8 T-cell granzyme B was consistent with these results (Fig. 2C). Interestingly, VV-e7r130-specific in vitro killing measured by 51Cr release assay was more than fourfold lower than in vivo killing (19% versus 87%) 5 days after VV infection, indicating a much higher sensitivity for the in vivo killing assay. The kinetics of VV-specific CD8 effector functions coincided with a decrease of VV titers measured in several organs. VV titers peaked at day 3 after infection and were significantly reduced 6 days postinfection. VV was cleared in the spleen and liver by day 8 postinfection but persisted slightly longer in fat pads and testes, clearing by 15 days postinfection (Fig. 2D). VV PFU could not be detected in any organ (testis, fat pads, spleen, liver, lung, brain, or salivary gland) 30 days postinfection, suggesting that the virus had been completely cleared from the host.

VV-specific CD8 T-cell lines generated from VV-immune mice respond to VV in vitro and in vivo. We next established VV-e7r- and -a11r-specific T-cell lines by stimulating splenocytes from VV-immune mice with the VV-specific peptides for 20 to 25 days in the presence of the IL-2. In 51Cr release cytotoxicity assays, both lines were able to lyse VV-infected
targets (Fig. 3A). In vitro ICS assays also demonstrated that the VV-e7r-specific and a11r-specific lines produce IFN-γ and TNF-α in response to VV-infected MC57G cells (Fig. 3B). However, stimulating with VV-infected MC57G cells in an ICS assay was not as sensitive a technique as peptide stimulation or the in vitro cytotoxicity 51Cr release assay using VV-infected or peptide-coated targets. Both CFSE-labeled VV epitope-specific cell lines upon adoptive transfer proliferated in vivo in the recipient mice upon VV infection, whereas control influenza A virus NP366-specific lines did not (Fig. 3C). These results demonstrate that these epitopes were processed and presented during VV infection.

**VV-e7r130-specific CD8 T cells reduce VV load.** To investigate a potential protective capacity of CD8 T-cell responses against VV infection, we used two different methods, either adoptive transfer of VV-e7r-specific CD8 T cells into the peritoneal cavity or immunization with ubiquitinated minigenes expressing VV-e7r130. To directly test if VV-specific CD8 T-cell responses could protect against VV infection, rested VV-e7r130-specific CD8 T-cell lines were adoptively transferred i.p. into recipient syngeneic naïve mice, which were then infected with VV. Control mice were injected with influenza virus NP366-specific CD8 T-cell lines, which were documented not to be cross-reactive with VV. Tetramer or ICS assays demonstrated that more than 95% of each CD8 T-cell line was specific for the peptide used to generate the line (Fig. 4A). Three days after adoptive transfer, the CFSE-labeled VV-e7r-specific CD8 T cells had gone through three to six rounds of division in response to the VV infection, while the influenza virus NP366-specific cells did not proliferate (Fig. 4B). There was a significant 99% reduction in VV titers 3 days after VV infection in the fat pads of the mice injected with the VV-e7r-specific line compared to the titers in the mice injected with the control influenza virus NP366-specific line (Fig. 4C). Similar results were observed in a second experiment, in which a PBS injection was compared to injection with the VV-e7r130 cell line (Fig. 4C). Our VV dose was not lethal, but control mice appeared to be more ill, as demonstrated by significantly greater weight loss (Fig. 4D), ruffled fur, shivering, and a decrease in

**FIG. 3.** VV-e7r-specific and VV-a11r-specific lines generated from VV-immune mice respond to VV in vitro and in vivo. (A) In vitro 5-h 51Cr release cytotoxicity assays demonstrate specific lysis of peptide-coated RMA-S cells and VV-infected MC57G cells. Control targets included RMA-S cells, MC57G cells, and LCMV-infected MC57G cells. (B) In vitro ICS assays show that the e7r-specific and a11r-specific lines produce IFN-γ and TNF-α in response to VV-infected MC57G cells. As a control, these lines did not produce cytokines to LCMV-NP396 (data not shown). (C) Both VV-e7r-specific and VV-a11r-specific lines proliferated in response to VV in vivo. VV-e7r-specific CD8 T-cell line (VV-p10-3, generated from a VV-immune mouse) and a11r-specific CD8 T-cell line (L/P24-7, generated from an LCMV- and VV-immune mouse) proliferated in response to VV infection i.p. as assessed by loss of CFSE by day 4 after adoptive transfer of the cell line i.p. into syngeneic C57BL/6 mice. The PECs collected from these mice were used in an ICS assay and produced IFN-γ only to the VV peptide used originally to stimulate the line and did not respond to the alternate VV peptide or the control influenza virus NP366 peptide. The control NP366-specific line generated from an influenza virus-immune mouse did not proliferate in response to VV and did not produce IFN-γ in response to stimulation with VV peptides.
activity, than the mice treated by immunization with the VV-specific T-cell lines.

A second method utilized three intramuscular immunizations every 14 days with an ubiquitinated DNA minigene specific for the VV-e7r130 sequence to demonstrate the protective role of CD8 T-cell responses. The VV-e7r130-specific CD8 T-cell responses measured by intracellular IFN-γ (Fig. 5A and B) and in vitro cytotoxicity (Fig. 5C) assays were increased approximately three- to sixfold compared to nonimmunized mice or mice immunized with the F3Ub vector only. Six days after VV infection, the VV-e7r130-specific CD8 T-cell population was significantly higher, at 2.4 \( \times 10^6 \) total cells (range, 1.0 \( \times 10^6 \) to 5.1 \( \times 10^6 \); \( n = 5 \); \( P = 0.04 \), versus the F3Ub group) in VV-e7r130-immunized mice, than the T-cell populations in nonimmunized and F3Ub-immunized virus-immune mice, respectively, demonstrated antigen specificity in an ICS assay (representative of two similar experiments). (B) The VV-e7r-specific CD8 T-cell line, but not the IV-NP366-specific cell line, proliferated in response to VV infection (i.p.), as assessed by loss of CFSE day 3 after adoptive transfer (i.p.) of the cell line into syngeneic C57BL/6 mice. (C) Significant reduction in VV titers in two experiments on days 3 and 4 after VV infection in mice injected with the VV-e7r-specific (black circles) or control influenza virus NP366-specific (gray circles) (experiment 1) cell line or PBS (white circles; experiment 2) (# and *, \( P < 0.05 \); four or five mice/group). (D) Significant protection from weight loss day 4 after VV infection in mice injected with VV-e7r-specific cell line compared to protection from weight loss in mice given PBS (\( P < 0.003 \); five mice/group). Weight is expressed as a percentage of the original weight prior to infection.

FIG. 4. VV-e7r-specific CD8 T cells reduce VV titers in vivo. (A) VV-e7r-specific (VV-e7r130) or influenza virus NP366-specific (IV-NP366) CD8 T-cell lines generated from VV-immune or influenza virus-immune mice, respectively, demonstrated antigen specificity in an ICS assay (representative of two similar experiments). (B) The VV-e7r-specific CD8 T-cell line, but not the IV-NP366-specific cell line, proliferated in response to VV infection (i.p.), as assessed by loss of CFSE day 3 after adoptive transfer (i.p.) of the cell line into syngeneic C57BL/6 mice. (C) Significant reduction in VV titers in two experiments on days 3 and 4 after VV infection in mice injected with the VV-e7r-specific (black circles) or control influenza virus NP366-specific (gray circles) (experiment 1) cell line or PBS (white circles; experiment 2) (# and *, \( P < 0.05 \); four or five mice/group). (D) Significant protection from weight loss day 4 after VV infection in mice injected with VV-e7r-specific cell line compared to protection from weight loss in mice given PBS (\( P < 0.003 \); five mice/group). Weight is expressed as a percentage of the original weight prior to infection.

In a second experiment, we also observed a 90% reduction in VV titers (Fig. 5D), as well as significantly decreased weight loss (Fig. 5E) from VV-e7r minigene immunization compared with that from control LCMV-NP396 minigene immunization.

**DISCUSSION**

Through the study of T-cell cross-reactivity between two heterologous viruses, we found that sequence similarity to a known epitope may be a useful tool to help predict immunogenic epitopes to large viruses, such as VV. This led to the finding that VV infection induced potent CD8 T-cell responses that were well maintained into memory and that VV-specific CD8 T cells could play a significant role in mediating protective immunity to VV. Our initial observations of VV-induced proliferation of T cells specific to an LCMV epitope, NP205, in LCMV-immune mice indicated that this LCMV epitope-specific response potentially included cross-reactive T cells that could also recognize VV (8, 26, 27). Mechanisms for CD8 T-cell cross-reactivity can be manifold. Molecular mimicry, in which a different peptide retains sites that are necessary for interaction with the TCR, is one of several paths to cross-reactive T-cell responses (43), as in the case with the NP205 epitope of LCMV and PV (5). It had been difficult to identify VV-specific MHC-I-restricted epitopes, as VV encodes more than 200 proteins and theoretically thousands of potential epitopes (21). Basing our method on the concept of molecular mimicry, we scanned the VV genome for sequence similarities to LCMV-NP396 and identified two VV immunogenic epitopes after screening only 115 peptides. This would suggest that this technique adds another approach to identifying immunogenic epitopes and could complement more conventional methods using algorithms based on MHC binding motifs (H. G. Rammensee, J. Bachmann, and S. Stevanovic, SYFPEITHI, a database of MHC ligands and peptide motifs—epitope prediction [http://www.syfpeithi.de/Scripts/MHCServer.dll/EpitopePrediction.htm]). Paradoxically, we identified an immunodominant VV epitope, VV-e7r130, that did not generate cross-reactive T-cell responses with LCMV and a subdominant epitope, VV-a11r198, which did generate cross-reactive T-cell responses with LCMV. Further characterization of these responses in subsequent studies has confirmed these observations (Cornberg et al., submitted). Identification of these VV-specific epitopes led to a better understanding of the role CD8 T cells play during VV infection. This strategy may be useful for identifying new CD8 T-cell epitopes for large viruses such as VV. This technique is particularly useful for identifying epitopes that may be missed by the more conventional methods, such as VV-a11r198, VV-a11r198 was not identified in a recent report using algorithms to identify VV epitopes in C57BL/6 mice (35). Although the VV-a11r198 epitope is a weak epitope in naive C57BL/6 mice infected with VV because of its ability to activate cross-reactive LCMV-specific memory CD8 T cells, its frequency can be significantly increased in some LCMV-immune mice infected with VV based on the private specificity of the memory T-cell repertoire of each mouse (27). Both epitopes VV-e7r130 (STLNFNNL) and VV-a11r198 (AIVNYANL) were conserved among vaccinia virus strains (WR,
FIG. 5. VV-e7r-expressing ubiquitinated DNA minigene vaccine increased e7r-specific CD8 T cells and reduced VV load. Increased frequency (A) and total number (B) of the VV-e7r130 epitope-specific CD8 T cells day 6 after VV infection in mice immunized three times intramuscularly with e7r-expressing ubiquitinated DNA minigene as assessed by ICS assay. Data are from one representative mouse (five mice/group). The total number of VV-e7r-specific cells equals the frequency of antigen specific cells multiplied by the total number of viable cells. There was a greater number of VV-e7r130 epitope-specific cells in the VV-e7r130 minigene-immunized mice than in mice given either control, minigene vector or PBS (P values of <0.05 and <0.08, respectively; n = 5). Data are representative of two experiments. No stim, no stimulation. (C) Enhanced cytotoxicity as measured by 51Cr release assays on splenocytes from the VV-e7r130 minigene-immunized mice compared to controls given minigene vector or PBS. Targets included peptide-coated RMA-S cells or VV-infected MC57G cells (five mice/group). (D and E) Significant reduction in VV titers day 6 after VV infection in mice injected with VV-e7r-specific minigenes (■) compared to control-vector-immunized mice (■) (log PFU/ml, 2.6 ± 0.4 versus 3.7 ± 0.2; P < 0.05; seven and five mice, respectively) or control LCMV-NP396-specific minigene-immunized mice (○) (log PFU/ml, 2.7 ± 0.4 versus 3.7 ± 0.2; P < 0.07; eight and five mice, respectively). (E) Significant protection from weight loss during VV infection of VV-e7r130 minigene immunized mice (●) compared to control LCMV-NP396 minigene-immunized mice (○) (P < 0.0004; eight and five mice, respectively; paired Student’s t test). Weight is expressed as a percentage of the original weight prior to infection.
Ankara, and Copenhagen), other poxviruses (cowpox, monkeypox, camelpox, and ectromelia virus), and also variola virus. VV-e7r is a soluble myristylated late protein of unknown function, and it is not associated with the virus membrane (30). The 36- to 40-kDa putative VV protein a11r is one of the 49 gene products that are conserved in all sequenced poxviruses (54) and was found to be associated with the putative DNA-packaging protein A32L required for virion morphogenesis (34). Recently, Resch et al. showed that the a11r protein is also a late protein that seems to be required to form normal virion membranes (40).

Although VV-specific CTLs were first described in 1975 (29), the importance of CD8 T cells in poxvirus infections still remains somewhat controversial and remains an important question, since newer vaccines are being designed to include induction of CD8 T-cell responses. Earlier studies had mixed results. Studies with ectromelia virus (mouse pox), a poxvirus closely related to VV, in mice demonstrate a consistent and absolute role for CD8 T cells in the control of primary infection, although antibody responses have been found to be important later in infection (4, 6, 15, 25). Tscharko et al. have demonstrated that immunization with a newly identified immunodominant peptide provided significant protection against a secondary lethal ectromelia infection, suggesting that CD8 memory T-cell responses could contribute to protective immunity to ectromelia virus (53). However, studies with VV in mice have been less definitive in identifying a role for CD8 T cells in both primary and secondary infections (3, 13, 49, 50, 59).

Studies examining the role of CD8 T cells during primary infection with VV have suggested that CD8 T cells may not be essential but they do play a supportive role. In one study using high-dose VV, β2-microglobulin knockout mice (β2 m−/− mice) which lack CD8 T cells were able to recover from VV infection, suggesting that CD8 T-cell responses were not essential during the primary infection, but this study did not rule out that they could play a significant role. Recovery was described as the disappearance of skin lesions and weight gain after intradermal inoculation, and VV titers were not analyzed (50). In another study, Belyakov et al. (3) showed that CD8 T cells alone in the absence of an antibody response were not sufficient to protect against VV infection during a primary infection (analyzed by weight differences). However, the depletion of CD8 T cells in the absence of an antibody response prevented late recovery, suggesting that CD8 T cells could make the difference between survival and death (3). A recent study by Xu et al. also showed that CD8 T cells can contribute to protection against VV, but CD4 T cells and antibodies may play a more important role, at least in primary infection (59).

There are less data available for the role of CD8 T cells during secondary VV infection. One study by Snyder et al. (49) documented protection against lethal secondary VV challenge in HLA-A2 transgenic mice by vaccination with an MHC-I-restricted T-cell epitope, suggesting that CD8 memory T-cell responses were important in protection against VV. These mice received the VV peptide three times subcutaneously accompanied with 50 nmol of a hepatitis B virus core helper peptide, 5 μg of granulocyte-macrophage colony-stimulating factor (GM-CSF), 800 IU of IL-2, and incomplete Freund’s adjuvant. However, these mice with only a single CD8 epitope-specific memory response did not have complete protection, as some mice lost weight and some mice died. This did not occur in mice previously immunized with the whole virus (49). Another study by Daftary et al. (12) using HLA-A2 transgenic mice to identify human immunodeficiency virus (HIV)-specific CD8 T-cell epitopes, demonstrated that immunization with a Th-CTL fusion peptide against an HIV epitope led to enhanced clearance of recombinant VV expressing the HIV epitope but was highly dependent on coadministration of peptide with cytosine-phosphate-guanine DNA (12). However, in another study by Drexler et al., immunization with HLA-A2-restricted VV-specific CD8 peptide vaccines was not able to protect HLA-A2 transgenic mice against a secondary infection with WR VV infection (13). Interestingly, research into sequential heterologous virus infections, in which there are no cross-protective neutralizing antibodies, shows that cross-reactive LCMV-specific memory CD8 T cells mediate protective immunity early in VV infection, suggesting that CD8 T cells could play an important role upon secondary VV challenge (8, 46; Cornberg et al., submitted). Our studies would strongly support the contention that CD8 memory T cells could play an important role in protection against VV during secondary infections. VV-e7r130-specific ubiquitinated DNA immunization significantly enhanced epitope-specific CD8 memory T-cell responses in vivo and resulted in a 90% reduction of VV titers on secondary challenge. Another technique allowed a more direct test of the ability of effector type memory CD8 T cells to protect against VV in a peripheral site. Epitope-specific CD8 T cells were activated in vitro and then rested into a quiescent state before transfer into the intraperitoneal cavity prior to challenge with VV. Only 106 cells were required to reduce VV titers more than 95%. These data demonstrate a number of points. First, e7r130-specific CD8 T cells recognized VV-infected targets in vivo, supporting the concept that this epitope is processed and presented. Second, CD8 T cells provided protection against VV, especially when delivered at the site of the infection. This underlines the concept that peripheral effector memory T cells can contribute to the early clearance of a virus infection by their immediate response (8, 24, 31) and justifies the use of vaccines which induce CD8 T-cell-dependent immunity. Finally, T cells can encounter their antigen presented on APCs in the periphery at the site of the infection. It also suggests that adoptive transfer of in vitro generated antigen-specific CD8 T cells could be a potential therapeutic intervention against poxviruses as has been shown for other infectious diseases (41). This model using intraperitoneal infection of CD8 T cells also provides a useful tool to study the effect of in vitro generated CD8 T cells upon VV-infected mice or in other viral infection models. To our knowledge, this is the first report of immunization with an ubiquitinated minigene expressing a VV-specific CD8 epitope or with the direct transfer of a VV-specific CD8 T-cell line leading to protective immunity without added CD4 helper peptides or adjuvants, and this clearly demonstrates that memory CD8 T cells can mediate protection against VV.

These studies are an excellent baseline for a better understanding about VV-specific memory CD8 T-cell responses in the context of other heterologous virus infections. For instance, the VV epitope-specific CD8 T cells were maintained in the memory population with the same immune hierarchy.
even more than 200 days after VV challenge (Fig. 2A and B). However, this analysis is not comparable with the human scenario in which one is exposed to numerous antigens during a lifetime. Encountering new infections induces immune responses to the new pathogen and results in attrition of non-cross-reactive memory T cells specific to the previous antigen (44). Studies in humans have shown that there is a 10-fold decrease in VV-specific memory over a lifetime (22). Considering the number of infections which humans can encounter, this appears to be a minimal loss. However, when memory CD8 T cells are able to generate cross-reactive responses to new antigen, their high frequency and activation state give them an advantage over naïve T cells and can lead to a preferential expansion of the cross-reactive CD8 T cells, which can alter the hierarchy of T-cell responses (5, 11). It is possible that cross-reactive VV-specific T cells could preserve VV-specific memory and can explain the maintenance of VV-specific immune responses more than 40 years after smallpox vaccination with VV (22). It is possible that cross-reactive memory responses, which may be of lower avidity to the new heterologous virus, are not always as efficient at clearing the new pathogen as higher-avidity de novo non-cross-reactive responses. These lower-avidity responses to VV may be more prone to stimulating immunopathology especially if these responses are directed at late antigens like VV-a11r. Having these systems now in place, we are better able to address the role heterologous immunity and cross-reactivity play in mediating protective immunity and/or immunopathology during VV infection a model for smallpox vaccination.

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REFERENCES


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