Replication-defective virus vaccine-induced protection of mice from genital herpes simplex virus 2 requires CD4 T cells

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Abstract

Replication-defective herpes simplex virus 2 (HSV-2), used as an immunization strategy, protects against HSV-2 challenge in animal models. The roles of replication-defective virus-induced T cell subsets in control of HSV-2 infection have not been established. Mice lacking B cells (μMT) were immunized, depleted of CD4 or CD8 T cells, and then challenged intravaginally with HSV-2 to elucidate T cell subset contributions in the absence of virus-specific antibody. Immunized, CD4-depleted μMT mice developed severe infection of the genital tract and nervous system. In contrast, depletion of CD8 T cells from μMT mice did not attenuate protection. Immunized wild-type mice depleted of CD4 T cells also developed more severe HSV-2 infection than mice from which CD8 T cells were depleted. Thus, immunization with replication-defective virus induces T cell responses that effectively control HSV-2 infection in the absence of HSV-immune antibody, and CD4 T cells play the predominant role in this protective effect.

Keywords

HSV-2; vaginal; B cell-deficient; CD4 T cells; immunization

Introduction

Herpes simplex virus 2 (HSV-2) is a sexually transmitted virus that causes most genital ulcerative disease. The virus currently infects 17% of adults in the United States and up to 75% of the adult population worldwide (Obasi, Mosha et al., 1999; Kamali, Nunn et al., 1999; Xu, Lee et al., 2007). Lesions caused by HSV-2 facilitate infection by HIV (Wald & Link, 2002; Freeman, Weiss et al., 2006), and HSV-2 shed in the genital tract of pregnant women can be transmitted to babies during birth, often with devastating consequences (Kimberlin & Whitley, 2005). No vaccine against HSV-2 has been licensed for use despite decades spent developing inactivated or glycoprotein-based vaccines. An adjuvanted glycoprotein vaccine provides limited protection, but only to seronegative women (Stanberry, Spruance et al., 2002). Development of alternative vaccination strategies is therefore warranted, and increased understanding of immune protective mechanisms that underlie their success in animal models may help further improve vaccine design.
Studies examining mechanisms of immune protection of the mouse vaginal tract and sacral sensory ganglia from HSV infection have often utilized intravaginal (i.vag.) inoculation of mice with attenuated HSV-2 as a method of generating immunity (McDermott, Smiley et al., 1984; Milligan & Bernstein, 1995b; Parr, Kepple et al., 1994; Harandi, Svennerholm et al., 2001). In these mice, genital secretions contain virus-specific secretory IgA and IgG (Milligan & Bernstein, 1995b; Parr, Bozzola et al., 1998), and circulating virus-specific IgG is also stimulated (McDermott, Brais et al., 1990; Parr & Parr, 1998a). Transfer of serum from HSV-immune mice to non-immune mice does not reduce HSV replication in the vaginal mucosa but does protect the nervous system and prevent a lethal outcome (McKendall, Klassen et al., 1979; Openshaw, Asher et al., 1979; McDermott, Brais, & Evelegh, 1990; Eis-Hubinger, Schmidt et al., 1993; Schneweis, Brado et al., 1988). B-cell-deficient (μMT) mice inoculated i.vag. with attenuated HSV-2 do not control challenge virus replication in the vaginal mucosa as readily as wild-type mice, but fare better than unimmunized mice (Dudley, Bourne et al., 2000; Parr & Parr, 2000; Harandi, Svennerholm et al., 2001). These observations indicate first, that the initial inoculation with attenuated HSV-2 stimulates T cell responses with some protective efficacy, and second, that HSV-specific antibody does influence virus replication in the mucosa. Interestingly, HSV-immune serum fully reconstitutes protection of the vaginal epithelium when transferred to immunized μMT mice (Dudley, Bourne, & Milligan, 2000), suggesting that early control of HSV-2 replication in the genital mucosa mediated by HSV-specific serum antibody requires the presence of immune T cells. Collectively, these studies indicate antibody generated by immunization with attenuated HSV-2 can ameliorate subsequent vaginal infection, but suggest a critical contribution of cell-mediated immunity to protection against HSV-2 challenge.

Several additional lines of evidence directly support a prominent role for HSV-immune T cells in protecting against HSV re-infection. Treatment with cyclosporine A selectively depresses T cell function and results in more severe HSV infections and reactivations in humans and in animal models (Field & Gottsch, 1995; Schneweis, Brado et al., 1988). Transfer of HSV-immune T cells but not B cells to naïve mice reduces lethality after HSV-2 infection (McDermott, Goldsmith et al., 1989). In addition, depletion of T cells from mice previously inoculated i.vag. with attenuated HSV-2 completely abrogates resistance to vaginal HSV-2 challenge over the first 5 days post-infection (Dudley, Bourne, & Milligan, 2000; Parr & Parr, 1998b). The T cell subset principally responsible for protection remains at issue. Some studies using depletion of HSV-immune T cell subsets found that CD4 T cells play the more significant role in protection of the genital tract and prevent acute infection of the nervous system by HSV (Milligan, Bernstein et al., 1998b; Schmidt, Eis-Hubinger et al., 1993; Milligan & Bernstein, 1997; Kuklin, Daheshia et al., 1998). In addition, depletion of CD4 T cells from wild-type mice immunized intranasally with vaccinia virus expressing HSV glycoprotein B, or from μMT mice immunized intranasally with attenuated HSV-1, reduces protective efficacy against HSV-1 replication after vaginal challenge more than does CD8 T cell depletion (Kuklin, Daheshia et al., 1998). Another study similarly using T cell subset depletion from mice previously immunized with attenuated HSV-2 concluded that CD8 T cells provide protection of the genital epithelium from subsequent HSV-2 infection (Parr & Parr, 1998b); however, in this study the extent of vaginal infection was evaluated by histology and quantification of virus protein instead of virus replication. Adoptive transfer of HSV-immune CTL has also suggested virus-immune CD8 T cells are protective against reinfection (McDermott, Goldsmith et al., 1989; Schneweis, Brado et al., 1988). Collectively, these studies indicate an important role for HSV-immune T cells in protection of the genital tract and nervous system, but the role of individual T cell subsets may depend on such variables as the immunizing agent or evaluation methods.

Although i.vag. administration of attenuated HSV-2 has been exceedingly useful in illuminating specific immune mechanisms protecting the genital tract against re-infection, this
method of immunization causes extensive infection of vaginal tissue and sub-lethal disease even at low doses (McDermott, Smiley et al., 1984) and so is not tenable as a vaccination strategy. Replication-defective virus offers increased safety over attenuated, replication-competent virus, making it a potentially viable approach to vaccination. Furthermore, the broad spectrum of immunogenic viral proteins expressed by replication-defective virus may elicit immunity of greater magnitude and sophistication than a glycoprotein subunit vaccine. Replication-defective HSV-2, administered subcutaneously (s.c.) as an immunizing agent, greatly reduces the severity of subsequent genital HSV-2 infection in mice and guinea pigs (Da Costa, Morrison et al., 2001; Morrison, Da Costa et al., 1998; Da Costa, Bourne et al., 1997). We previously determined that HSV-specific antibody generated by s.c. immunization of WT mice with an ICP8−, replication-defective HSV-2 strain protects the nervous system against infection and signs of disease and also reduces infection of the vaginal mucosa if immune T cells are present (Morrison, Zhu, & Thebeau, 2001). However, the role of T cells independent of antibody, and the T cell subsets stimulated by replication-defective virus that mediate protection have not been elucidated. We therefore depleted T cell subsets from mice previously immunized with replication-defective HSV-2, and examined infection and disease of the genital mucosa and nervous system that occurs in their absence after HSV-2 challenge.

Results

The protective roles of replication-defective virus-immune, CD4 and CD8 T cell subsets would ideally be examined in a system that did not include the potentially confounding presence of immune serum antibody, such as in mice lacking B cells. Concerns linger, however, about the integrity of T cell responses in mice genetically deficient in B cells, with some studies noting differences (Dudley, Bourne, & Milligan, 2000; Homann, Tishon et al., 1998; Deshpande, Zheng et al., 2000) and others finding normal responses (Deshpande, Zheng et al., 2000; Topham, Tripp et al., 1996; Asano & Ahmed, 1996; Morrison, Zhu, & Thebeau, 2001). We therefore studied the contributions of immune CD4 and CD8 T cells to protection in B cell-deficient (μMT) mice, bred onto a BALB/c background, and then repeated the studies in BALB/c (WT) mice to compare and confirm results in an otherwise intact host.

μMT or WT mice were sham-immunized or immunized s.c. with replication-defective HSV-2 to generate vaccine-specific immunity. Prior to i.vag. challenge with virulent HSV-2, CD4 or CD8 T cells were depleted in vivo using rat monoclonal antibodies specific for either subset. Control depletions consisted of injecting normal rat IgG. Depletions were continued every 4 days through day 7 post-challenge. μMT mice immunized with replication-defective HSV-2 and then control-depleted were able to restrict challenge virus replication in the genital mucosa (Fig. 1A). Titers were not reduced until 3 days post-challenge, however, consistent with previous observations (Morrison, Zhu, & Thebeau, 2001). Immunized μMT mice depleted of CD8 T cells also readily controlled replication in the genital mucosa by 3 days post-challenge (Fig. 1A). Titers were not significantly different than those observed in immunized control-depleted animals. In contrast, immunized CD4-depleted mice showed prolonged replication in the genital mucosa, with elevated titers at 3 to 5 days post-challenge that were indistinguishable from those of unimmunized mice (Fig. 1A). In WT mice, CD8 depletion had only a modest effect on the capacity of the immune response to limit virus infection. Slightly higher titers in CD8-depleted than control-depleted mice were observed only on days 2 and 3 post-challenge (Fig 1B). Overall, CD8- and control-depleted WT mice curtailed replication more effectively at early times post-challenge than their μMT counterparts. In contrast, CD4-depleted WT mice did not control replication of challenge virus in the genital mucosa (Fig. 1B), and virus titers resembled those seen in CD4-depleted μMT mice. Thus, replication-defective virus-immune, CD4 T cells have the principal role in limiting replication in the genital tract.
Signs of genital inflammation in μMT mice depleted of CD4 T cells were as severe as unimmunized mice and were markedly worse than CD8-depleted or control-depleted μMT mice (Fig. 2A). Correspondingly, immunized μMT mice depleted of CD4 T cells prior to challenge lost significant weight, whereas the overall health of CD8-depleted mice was less severely affected by the challenge virus infection (data not shown). In contrast, WT mice showed a clear difference between CD4-depleted and unimmunized groups with only about half of the former developing lesions (Fig. 2B). WT mice depleted of CD8 T cells, like their μMT counterparts, showed only mild signs of genital inflammation. Control-depleted WT mice remained completely protected (Fig. 2B). HSV-2 causes a more severe infection in the mouse model than in humans, with signs of illness extending to the nervous system in non-immune mice. Consequently, hind-limb paralysis developed in 90% of CD4-depleted or unimmunized μMT mice but in only 30% of the CD8-depleted and in none of the control-depleted mice (Table 1). Hind-limb paralysis developed in fewer CD4-depleted WT mice than control-depleted mice, and those paralyzed developed paralysis approximately 1 day later (Table 1). Not surprisingly, the CD4-depleted μMT mice died as rapidly as unimmunized controls, whereas immunized, CD8-depleted μMT mice rarely succumbed to infection (Fig. 3A). Although not all CD4-depleted WT mice developed genital lesions and paralysis, nearly all of the mice eventually succumbed to infection (Fig. 3B). The lethality of the infection in CD4-depleted mice precluded study of latency. Together, these results reveal a major contribution of virus-immune CD4 T cells to protection of the genital tract and nervous system from HSV-2-induced disease, but scant evidence of a CD8 T cell contribution.

The uniform paralysis observed in CD4-depleted μMT mice likely resulted from direct infection of the spinal cord and associated ganglia, but inflammation in the spinal cord could also result in CNS dysfunction (Bishop & Hill, 1991). To distinguish between these possibilities, the peripheral and central nervous systems of a cohort of immunized, T cell-depleted μMT mice were dissected at 7 days post-challenge, when signs of paralysis were developing. CD4-depleted and unimmunized cohort animals showed high titers of virus in the spinal cord, brainstem and brain (Fig. 4). In contrast, immunized mice that were CD8-depleted or given control Ig had low titers of virus (Fig. 4). Thus, immune control of acute HSV-2 infection of the nervous system also largely depends on the presence of CD4 T cells, and paralysis in these mice is likely due to vigorous replication of challenge virus in neurons. Together, these results strongly support a critical role for CD4 T cells induced by replication-defective virus in protecting the genital tract and nervous system from the deleterious effects of challenge virus infection. Possible supporting roles for HSV-immune antibody and CD8 T cells are discussed.

**Discussion**

Depletion of T cell subsets from WT or μMT mice vaccinated with replication-defective HSV-2 has revealed a decisive role for CD4 T cells in reducing HSV-2 replication in the genital tract, signs of genital and neurologic disease, and acute infection of the nervous system. Furthermore, the critical effector function of CD4 T cells is not the provision of help for generating immune serum antibody because the protective effect of CD4 T cells is still observed in mice lacking B cells. Because immunization of μMT mice with attenuated HSV (Milligan, Bernstein, & Bourne, 1998; Kuklin, Daheshia et al., 1998), vector-expressed HSV glycoprotein (Kuklin, Daheshia et al., 1998), or replication-defective HSV-2 (this study) have yielded similar results, we can surmise as a general phenomenon that CD4 T cells mediate significant protection of the mouse genital mucosa and nervous system independently of HSV-immune antibody. Nonetheless, CD8 T cells play an important role elsewhere in the body, curtailing HSV-1 infection or reactivation after cutaneous or ocular inoculation (van Lint, Ayers et al., 2004; Banerjee, Biswas et al., 2005; Liu et al., 2000).
T cell subset depletion from μMT mice previously immunized with replication-defective virus allowed us to discern T cell roles in protection in the absence of contributions from HSV-specific antibody. We also performed the same depletions in immunized WT mice, uniquely allowing us to corroborate the results obtained with μMT mice in a system that was not potentially prejudiced by lack of B cells, a population with antigen presenting and effector capacities. This comparison yielded very similar conclusions about the preeminent role of CD4 T cells in protection. Differences we did observe between the WT and μMT mouse models were modest: First, replication-defective virus-immune WT mice that received anti-CD8 or control antibody restrained early HSV-2 replication in the genital tract slightly better than the analogous μMT mice. Second, all immunized μMT mice were slightly more susceptible to lethal HSV-2 infection than immunized WT mice regardless of the subsequent depletion protocol. Third and most conspicuous, immunized WT mice depleted of CD4 T cells developed less severe genital disease and neurologic signs on average than did unimmunized mice, whereas in μMT mice, CD4-depleted and unimmunized groups were indistinguishable.

One interpretation of the less severe genital disease in CD4-depleted WT mice is that immune CD8 T cells assist in protection of WT but not μMT mice. In support of this possibility, CD8-depleted WT mice showed some genital inflammation, unlike mice treated with control Ig. Nonetheless, inflammatory signs were equally mild in CD8-depleted μMT and WT mice, arguing against this possibility. Alternatively, immune serum antibody (still present in the immunized, CD4-depleted WT mice) assists in protection against genital and neurological disease. This interpretation is consistent with our prior observations (Morrison, Zhu, & Thebeau, 2001) and those of Dudley et al. (Dudley, Bourne, & Milligan, 2000) that passively transferred immune serum antibody 1) significantly reduces the incidence and severity of genital and neurological disease after HSV challenge of immunized μMT mice, and 2) helps reduce early challenge virus replication in the genital mucosa, but only in the presence of virus-immune T cells. Our results here suggest that the CD4 T cell subset is the critical component working with antibody to reduce mucosal replication of challenge virus: lower replication over the first 2 days post-challenge was observed only in immunized WT mice that still contained CD4 T cells (Fig. 1B). Even so, we cannot formally rule out an additional protective effect of CD8 T cells, particularly in view of the slight inflammation observed in the genital mucosa of CD8-depleted WT mice compared with control-depleted mice.

The interpretation that HSV-immune, CD8 T cells play a very limited role in protecting either the genital tract or the nervous system from HSV-2 infection depends upon evidence that they are induced by replication-defective virus, and that they mount normal responses in the context of a B cell-deficient mouse. CD8 T cells specific for HSV-2 are found in the genital lymph nodes and vaginal epithelium of mice after i.vag. immunization with attenuated HSV-2 (Milligan & Bernstein, 1997; Milligan & Bernstein, 1995a) or intranasal immunization with adenovirus expressing HSV-1 gB (Gallichan & Rosenthal, 1996). Immunization of WT and μMT mice with replication-defective virus also induces CD8 T cell responses (Morrison, Zhu, & Thebeau, 2001), and in preliminary experiments we have found higher percentages of activated CD8 T cells in genital lymph nodes after immunization or challenge of μMT mice (data not shown).

In μMT mice, CD8 T cell recruitment to the vaginal mucosa 20 hr after HSV challenge is reportedly normal (Parr & Parr, 2000), as are CD8 cytolytic responses to HSV (Deshpande, Zheng et al., 2000), and CD4 and CD8 responses to other viruses (Topham, Tripp et al., 1996; Asano & Ahmed, 1996). We had previously shown that CD8 T cells of μMT mice elicited by replication-defective virus produce IFNγ in normal amounts in response to challenge (Morrison, Zhu, & Thebeau, 2001). In preliminary experiments to further analyze T cell subsets in μMT mice, we have also found no differences in expression of early activation marker CD69 at 3 d after infection, or of CD25 (IL-2 receptor alpha chain) at 4 and 7 d after infection.
compared with WT mice. However, CD40L expression on CD8 T cells from μMT mice appears discernibly reduced (data not shown). This deficit or delay in CD40L expression potentially could have functional consequences (Kemball, Lee et al., 2006). Thus we must cautiously interpret our data to include the possibility of a minor contribution of replication-defective virus-immune CD8 T cells to protection of the genital mucosa and nervous system from HSV-2 infection and disease, as seen in WT mice but not in μMT mice containing potentially impaired CD8 T cells.

The prominent role of virus-immune CD4 T cells in protection leads to speculation about the protective effector mechanism. A large proportion of CD4 T cells in the genital lymph nodes stain IFNγ-positive during acute and recall responses to HSV (Milligan & Bernstein, 1995a), CD4 T cells are the main producers of IFNγ (Milligan & Bernstein, 1995a; Milligan & Bernstein, 1997), and IFNγ is known to be a critical component of the protective response to genital infection with HSV-2 (Asano & Ahmed, 1996; Milligan, Bernstein, & Bourne, 1998; Parr & Parr, 1999). In addition, we (Morrison, Zhu, & Thebeau, 2001) and others (Milligan, Dudley-McClain et al., 2004; Kolaitis, Doymaz et al., 1990) have demonstrated CD4 T cell cytolytic activity in the draining lymph nodes and vaginal epithelium soon after i.vag. HSV-2 infection. Either cytokine or cytolytic effector functions represent a plausible mechanism of CD4 T cell-mediated defense, though further experiments will be necessary to establish this point. Regardless of mechanism, replication-defective virus, used in a mouse model to vaccinate against HSV-2 infection, stimulates a strong protective effect mediated through CD4 T cells functioning in concert with, but to a large extent independently of providing help for, HSV-specific antibody. These results support development of vaccines that express viral proteins containing immunodominant CD4 epitopes to generate strong protective immunity.

**Materials and Methods**

**Cells and viruses**

A partially purified preparation of the ICP8−, replication-defective HSV-2 strain 5BlacZ (Da Costa, Bourne, Stanberry, & Knipe, 1997) was prepared for immunizations by high speed centrifugation of clarified supernatants of infected, ICP8-complementing S2 cells (Gao & Knipe, 1989) as previously described (Morrison, Da Costa, & Knipe, 1998). Stocks of HSV-2 strain G-6, a plaque-purified isolate of strain G, were grown in Vero cells and prepared as previously described (Morrison & Knipe, 1996). Titer of virus stocks was determined by standard plaque assay on S2 or Vero cell monolayers (Knipe & Spang, 1982).

**Antibody production**

Monoclonal antibodies specific for CD4 (GK1.5) or CD8 (53-6.72) were harvested from hybridoma culture supernatants of cells grown in Integra flasks (Integra Biosciences) according to the manufacturer’s recommendations. They were diluted to 1 mg/ml in sterile PBS for injection.

**Animals and immunizations**

BALB/c mice were purchased from the National Cancer Institute and were immunized at 6 wk of age. Mice bearing the μMT mutation (Igh6tm1Cgn) (Kitamura, Roes et al., 1991) were backcrossed 12 generations onto a BALB/c background in the Saint Louis University Department of Comparative Medicine. All animals were housed and treated in accordance with ALAC and Institutional guidelines. Mice were immunized s.c. in the hind flanks with 2x10^6 pfu of partially purified 5BlacZ in 20μl vol and rested for 4 wk before challenge.
Depletions and challenge

One week and one day prior to challenge, mice were injected s.c. in the neck ruff with 3 mg hydroxyprogesterone (Depoprovera, UpJohn) in 100 μl vol. Four days and one day prior to challenge mice received 0.3 mg monoclonal antibody in 0.3 ml i.p. Depletions were continued by antibody injections days 3 and 7 post-challenge. For challenge, mice were infected by i.vag. inoculation of 7.5 × 10^5 pfu G-6 in 5 μl vol. This represents 150 LD_{50} for BALB/c mice (Thebeau & Morrison, 2002). Challenge virus shed from the genital mucosa was quantified by swabbing vaginal vaults twice with calcium alginate swabs at 9 hr and days 1 through 5 post-infection. Duplicate swabs for each time point were frozen together in 1 ml PBS until plaque assays could be performed. Mice were monitored daily post-challenge for change in body weight, signs of disease and survival. Mice were weighed individually and mean change from initial body weight was calculated daily for each group. Disease scores were assigned in a blinded fashion based on the following scale: 0-no apparent signs of disease, 1-slight erythema and edema of the genitals, 2-prominent erythema and edema of the genitals, 3-severe erythema and edema with lesions on the genitals. Mean daily disease score ± SEM was calculated for each group. Virus replication in neural tissues was analyzed by dissection of brains, brainstems, and spinal cords from a cohort of mice 7 days after challenge. Tissues were stored frozen until use. For virus titer determination, the tissues were thawed and disrupted using a Mini-Bead Beater (BioSpec, Inc.) and diluted for standard plaque assay.

Flow cytometry

A cohort of mice was treated with monoclonal antibodies 3 days and 1 day prior to sacrifice. Splenocytes and genital lymph node cells were isolated and stained with anti-CD4 and anti-CD8 antibodies specific for different epitopes than the monoclonal antibodies used for depletions (Caltag). Flow cyt fluorometric analyses revealed that 95% of CD4^+ T cells were depleted and >98% of CD8^+ T cells from each tissue (data not shown).

Statistics

Significance of difference in virus or antibody titers on individual days was determined by Student’s t test. Proportion of mice with hind-limb paralysis or surviving infection was compared using the Fisher exact method. The Kruskal-Wallis non-parametric test was used to assess the significance of difference in disease scores on individual days post-challenge.

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Figure 1. Replication of HSV-2 in the genital mucosa of immunized depleted of CD4+ or CD8+ T cells
Groups of A) μMT or B) WT BALB/c mice were immunized s.c. with 2x10^6 pfu of HSV-2 5BlacZ and rested for 4 wk, or were left unimmunized. Immunized mice were then depleted of CD4+ or CD8+ T cells by injection of CD4 or CD8-specific monoclonal antibody, and challenged by i.vag. inoculation of 7.5x10^5 pfu HSV-2 G-6. A third group was injected with control antibody before challenge. Vaginal swab samples were collected at the indicated times post-challenge and titered by standard plaque assay. Data represent the geometric mean titers for 7 to 8 mice per group ± SEM. The experiment was repeated once. Data point at time 0 indicates the inoculum dose. *, P = 0.0138–0.0478; **, P = 0.001; ***, P = <0.0001–0.0003

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for CD4-depleted compared with control Ig-depleted mice. (For CD4-depleted compared with CD8-depleted mice: Fig. 1A, $P = 0.0478$ on day 2, and $P = <0.0001–0.0002$ on days 3 and 4; Fig. 1B, $P = 0.0016–0.0092$ on days 1 through 3, and $P <0.0001$ on day 4.)
Figure 2. Genital disease in immunized mice depleted of CD4⁺ or CD8⁺ T cells

Groups of A) µMT (n=9 to 11) or B) WT (n=7 to 10) mice treated as in Fig. 1 were examined over time post-challenge for signs of genital and neurologic disease. 0, no signs; 1, mild erythema and edema of the external genitalia; 2, moderate erythema and edema; 3, severe erythema and edema accompanied by lesions. Data were pooled from two independent experiments and represent the arithmetic mean ± SEM of all mice per group. *, P = 0.021; **, P = 0.002–0.006; ***, P <0.001 for CD4-depleted compared with control Ig-depleted mice. (For CD4-depleted compared with CD8-depleted mice: Fig. 2A, P = 0.021 on day 4, P = 0.002 on day 5, and P <0.001 on days 6 and 7; Fig. 2B, P = 0.015–0.026 for days 6 through 8.)
Figure 3. Survival of immunized mice depleted of CD4\(^+\) or CD8\(^+\) T cells

Groups of A) \(\mu\)MT (n=9 to 11) or B) WT (n=7 to 10) mice treated as in Fig. 1 were monitored over time post-challenge and sacrificed when moribund. *, \(P = 0.0147–0.0325\); **, \(P = 0.0055\); ***, \(P = <0.0001–0.0007\) for CD4-depleted compared with control Ig-depleted mice. (For CD4-depleted compared with CD8-depleted mice: Fig. 3A, \(P = 0.0325\) on day 8, and \(P = <0.0001–0.0007\) on days 9 through 14; Fig. 3B, \(P = 0.0047\) for day 11 through day 14.)
Figure 4. Replication of HSV-2 in the nervous system of immunized and T cell-depleted μMT mice
Groups of 6 to 8 5BlacZ-immunized μMT mice were depleted of CD4⁺ or CD8⁺ T cells prior to challenge as described in Figure 1. Mice were euthanized 7 days post-challenge. Neural tissues were dissected, homogenized and virus titer contained in them was determined by standard plaque assay. Values represent the geometric mean titers ± SEM. The experiment was repeated once. *** $P < 0.0001$ for CD4-depleted compared with control Ig-depleted mice. (For CD4-depleted compared with CD8-depleted mice: $P = 0.0006$ for spinal cord and $P < 0.0001$ for brainstem and brain.)
### Table 1
Percentage of mice developing hind-limb paralysis

<table>
<thead>
<tr>
<th>Mouse type</th>
<th>Depletion group</th>
<th>Percent developing paralysis</th>
<th>Mean day post-challenge ± SD</th>
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<tr>
<td>µMT</td>
<td>Control'</td>
<td>90.9</td>
<td>7.6±0.7</td>
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<tr>
<td></td>
<td>5B' + anti-CD4</td>
<td>88.9&lt;sup&gt;a&lt;/sup&gt;</td>
<td>7.6±1.3</td>
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<tr>
<td></td>
<td>5B' + anti-CD8</td>
<td>30.0</td>
<td>9.0±1.7</td>
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<tr>
<td></td>
<td>5B' + control Ig</td>
<td>0.0</td>
<td>--</td>
</tr>
<tr>
<td>BALB/c (WT)</td>
<td>Control'</td>
<td>85.7</td>
<td>7.5±0.8</td>
</tr>
<tr>
<td></td>
<td>5B' + anti-CD4</td>
<td>57.1&lt;sup&gt;b&lt;/sup&gt;</td>
<td>8.3±1.3</td>
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<tr>
<td></td>
<td>5B' + anti-CD8</td>
<td>0.0</td>
<td>--</td>
</tr>
<tr>
<td></td>
<td>5B' + control Ig</td>
<td>0.0</td>
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</tbody>
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<sup>a</sup>P = 0.0001 for anti-CD4 v. control Ig, and P = 0.0198 for anti-CD4 v. anti-CD8

<sup>b</sup>P = 0.0147 for anti-CD4 v. control Ig or anti-CD8