

# Early Control of Highly Pathogenic Simian Immunodeficiency Virus/Human Immunodeficiency Virus Chimeric Virus Infections in Rhesus Monkeys Usually Results in Long-Lasting Asymptomatic Clinical Outcomes

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**In contrast to simian immunodeficiency viruses (SIVs), which induce immunodeficiency over a 1- to 2-year period, highly pathogenic simian-human immunodeficiency viruses (SHIVs) cause an irreversible and systemic depletion of CD4<sup>+</sup> T lymphocytes in macaque monkeys within weeks of inoculation. Nonetheless, the seemingly more aggressive SHIVs have proven to be easier to control by the same vaccine regimens which fail to contain SIV. Because early events during *in vivo* infections may determine both the pathogenic consequences of the challenge virus and its sensitivity to interventions that prevent disease, we have evaluated the effects of inoculum size and a potent antiretroviral drug on the development of disease in monkeys infected with SHIV<sub>DH12R</sub>. The results obtained show that in a majority of inoculated animals, suppression of SHIV replication during the first 2 weeks of infection, which prevents complete loss of CD4<sup>+</sup> T cells, leads to very low to undetectable postpeak viremia and an asymptomatic clinical course for periods up to 4 years.**

During the past few years, pathogenic simian-human immunodeficiency viruses (SHIVs) have largely supplanted simian immunodeficiency viruses (SIVs) as the primate lentivirus of choice for the challenge of macaque monkeys in vaccine experiments (1, 4, 6, 35, 40, 49). This has occurred for two principal reasons: (i) SHIVs bear the human immunodeficiency virus type 1 (HIV-1) envelope glycoprotein, thereby permitting an assessment of anti-HIV-1 neutralizing antibody (NAb) induction, and (ii) SHIVs cause an unusually rapid, irreversible, and systemic elimination of CD4<sup>+</sup> T lymphocytes within 3 to 4 weeks of inoculation (17, 19, 33). Although the latter pathogenic phenotype permits an early assessment of vaccine efficacy against disease, it is profoundly different from the clinical course commonly associated with SIV and HIV-1 infections, which are characterized by more-moderate depletions of CD4<sup>+</sup> T cells and the development of clinical immunodeficiency over a much longer time frame (1 to 2 years and 10 years, respectively) (8, 21, 28, 31). Despite their seemingly more aggressive pathogenicity *in vivo*, SHIVs have proven to be easier to control by the same vaccination regimens that fail to protect rhesus monkeys from challenges with pathogenic SIV strains such as SIV<sub>mac239</sub> and SIV<sub>E660</sub> (15, 32). Because

these discrepancies in vaccine sensitivity might reflect fundamental differences in the mechanisms underlying the diseases induced by SIV and SHIVs, we have examined how a directed intervention (administration of a potent reverse transcriptase [RT] inhibitor) during the first 2 weeks of the acute infection or the conditions of initiating the primary infection by varying the inoculum size might modulate the natural history of pathogenic SHIV infections over a 2- to 4-year observation period. The results obtained have been compared with those previously reported for SIV.

In the present study, we used uncloned SHIV<sub>DH12R</sub> stock (13, 17) and found that the complete and irreversible depletion of CD4<sup>+</sup> T cells in infected rhesus monkeys could be abolished, following a single 4-week course of anti-retroviral therapy (using 9-[2-(*R*)-(phosphonomethoxy) propyl]adenine [PMPA]) initiated on day 5 postinfection. Three of four animals remain alive and healthy, with low to undetectable levels of plasma viral RNA, 1.5 to 3 years after cessation of treatment. Furthermore, one of two monkeys receiving similar 4-week PMPA regimens beginning on day 14 postinoculation (when plasma viral RNA levels were quite high [ $5.1 \times 10^7$  RNA copies/ml] and the number of CD4<sup>+</sup> T lymphocytes had started to decline) was also asymptomatic for more than 3 years postinfection. Similar benign long-term clinical outcomes were observed in several naïve monkeys inoculated with very low doses (1 to 25 tissue culture infective doses [TCID<sub>50</sub>]) of SHIV<sub>DH12R</sub> even though they had experienced high levels ( $10^7$  to  $10^8$  RNA copies/ml) of peak viremia between weeks 2 and 3. Thus, blunting of the acute SHIV<sub>DH12R</sub> infection, which prevents the complete elimination of CD4<sup>+</sup> T cells, resulted in an

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uneventful and durably maintained (2 to 4 years) clinical course in the majority of inoculated rhesus monkeys.

#### MATERIALS AND METHODS

**Animal experiments.** Rhesus macaques (*Macaca mulatta*) were maintained in accordance with the guidelines of the Committee on Care and Use of Laboratory Animals (*Guide for the Care and Use of Laboratory Animals*, NIH 85-23, Department of Health and Human Services, 1985) and were housed in a biosafety level 2 facility; biosafety level 3 practices were followed. Animals were anesthetized with intramuscular injections of ketamine hydrochloride (Ketaject; Phoenix Pharmaceutical, Inc. St. Joseph, Mo.) and acepromazine maleate (Fermenta Animal Health Co. Kansas City, Mo.) during phlebotomies or SHIV inoculations.

**Virus and plasma viral RNA measurements.** The emergence, characterization, and preparation of the highly pathogenic SHIV<sub>DH12R</sub> stock have been described previously (13, 16, 17). In the experiments described, the indicated amounts of SHIV<sub>DH12R</sub> were inoculated into the saphenous veins of anesthetized rhesus monkeys. Plasma viral RNA levels were determined by real-time PCR (ABI Prism 7700 sequence detection system; Applied Biosystems, Foster City, Calif.) utilizing reverse-transcribed viral RNA from macaque plasma samples as templates as previously described (13).

**Lymphocyte immunophenotyping.** EDTA-treated blood samples were stained with fluorochrome-conjugated monoclonal antibodies (MAbs) (CD3-fluorescein isothiocyanate [Serotec Ltd., Oxford, England, or BD Biosciences Pharmingen, San Diego, Calif.], CD4-allophycocyanin, CD8-peridinin chlorophyll protein, and CD20-phycoerythrin [BD Biosciences Immunocytometry Systems, San Diego, Calif.]) and analyzed by flow cytometry (FACSort; BD Biosciences Immunocytometry Systems) as previously described (39).

**PMPA treatment of SHIV<sub>DH12R</sub>-infected macaques.** PMPA was generously provided by Norbert Bischofberger, Gilead Sciences, Inc. (Foster City, Calif.). An aqueous solution (60 mg/ml) of PMPA was prepared after adjusting the pH to 7.0 with NaOH and filtration through a membrane filter (pore size, 0.22  $\mu$ m) as previously described (44). PMPA was administered intramuscularly (30 mg/kg of body weight) every 24 h for 4 weeks starting on day 5 (for monkeys N056, N077, BC74, and BI36) or on day 14 (for monkeys 96N130 and 96N135) postinfection with  $10^3$  TCID<sub>50</sub> of the SHIV<sub>DH12R</sub> stock.

**Virus isolation from SHIV<sub>DH12R</sub> chronically infected animals.** Peripheral blood mononuclear cells (PBMC) ( $10^7$  cells) from infected monkeys (or a mixture of PBMC, lymph node, and bone marrow cells [ $\sim 10^7$  cells total] from macaque RhH520) were resuspended in 10 ml of RPMI 1640 medium (Cambrex Bio Science, Walkersville, Md.) supplemented with 10% fetal bovine serum (Hy-Clone, Logan, Utah) and 20 U of recombinant human interleukin-2 (IL-2) (Roche Diagnostics Corporation, Indianapolis, Ind.)/ml. These specimens were cocultivated with  $10^7$  naive monkey PBMC which had been previously stimulated with concanavalin A (Amersham Biosciences Corp., Piscataway, N.J.) (25  $\mu$ g/ml) for 24 h and were maintained an additional 48 h in the presence of IL-2. All of the medium was replaced daily during 14 days of cocultivation, while the cell concentration of  $2 \times 10^6$  cells/ml was maintained. Virus production was assessed by RT assays of the daily culture supernatant samples, and the specimen with the highest level of RT activity was used as stock virus after titration in MT-4 cells (39).

**Effect of coreceptor-specific inhibitors on SIV and SHIV replication.** TAK-779 and AMD3100 were obtained from the AIDS Research and Reference Reagent Program, Division of AIDS, National Institute of Allergy and Infectious Diseases, National Institutes of Health. AD101 was provided by Bahighe Baroudy (Schering Plough Research Institute, Bloomfield, N.J.). Chemokine coreceptor usage was determined as described previously (51), with minor modifications. Briefly, uninfected rhesus PBMC were prepared as described above. On day 3,  $5 \times 10^4$  cells were dispensed in 96-well round-bottom plates. Various concentrations (0, 0.05, 0.1, 0.5, 1, 5, and 10  $\mu$ M) of small-molecule CCR5- or CXCR4-specific inhibitors (AMD3100 [11], TAK-779 [3], and AD101 [41]) were added to duplicate wells and incubated for 1 h at 37°C. After this incubation, each test virus was spinoculated (30) at  $1,200 \times g$  for 1 h at a multiplicity of infection of 0.1. On day 5 postinfection, virus replication was assessed by RT assays of the culture supernatants.

#### RESULTS

**SHIV<sub>DH12R</sub>-induced disease is rapid, irreversible, and complete.** SIV<sub>mac</sub>/SIV<sub>sm</sub> infection of rhesus macaques typically causes a gradual decline of CD4<sup>+</sup> T cells in the peripheral blood and the induction of immunodeficiency over a 1- to 2-year period (21, 31). As is the case for HIV-1, the development

of disease by SIV does not require the complete elimination of the CD4<sup>+</sup> T-lymphocyte subset. In contrast, highly pathogenic SHIVs, including SHIV<sub>DH12R</sub>, cause a rapid, systemic, and complete depletion of CD4<sup>+</sup> T cells in rhesus macaques within 3 to 4 weeks of virus inoculation and death from immunodeficiency during the ensuing 3 to 7 months (17, 19, 33). As shown in Fig. 1, nine animals inoculated intravenously with moderate to high (500 to 5,000 TCID<sub>50</sub>) levels of SHIV<sub>DH12R</sub> experienced the characteristic CD4<sup>+</sup> T-cell loss within several weeks (Fig. 1A) and were euthanized 15 to 30 weeks postinfection due to uncontrollable diarrhea, marked weight loss, or the onset of opportunistic infections. Plasma viral RNA levels in SHIV<sub>DH12R</sub>-infected rhesus macaques typically reached  $10^7$  to  $10^8$  copies/ml at 2 to 3 weeks postinoculation, coinciding with the rapid loss of CD4<sup>+</sup> T lymphocytes. After declining 20- to 400-fold from the initial peak of viremia, the plasma viral loads gradually increased to the  $10^7$  RNA copies/ml level. Of the 28 monkeys inoculated with 500 TCID<sub>50</sub> or more of SHIV<sub>DH12R</sub>, 26 exhibited the pattern shown in Fig. 1. The other two monkeys were the only recipients of SHIV<sub>DH12R</sub> (5,000 TCID<sub>50</sub>) from the same thawed vial of stock virus, and both experienced a delayed and transient depletion of their CD4<sup>+</sup> T cells. Each has remained asymptomatic for more than 3 years. We presently have no explanation for the unusual course of infection in these two monkeys except that they were the only animals inoculated with virus from the same vial of SHIV<sub>DH12R</sub>.

**SHIV<sub>DH12R</sub> induces disease in an inoculum size-dependent manner.** Unlike the immunodeficiency caused by SIV<sub>mac</sub>/SIV<sub>sm</sub>, the results that Endo et al. previously reported showed that the disease induced by SHIV<sub>DH12R</sub> (characterized by rapid, systemic, and irreversible loss of CD4<sup>+</sup> T cells) is dose dependent (13). As shown in Fig. 2 (upper panel inset), all five animals inoculated intravenously with 25 TCID<sub>50</sub> or less of SHIV<sub>DH12R</sub> experienced significant depletions of CD4<sup>+</sup> T lymphocytes during the first 4 weeks of infection. However, in contrast to macaques inoculated with larger amounts of SHIV<sub>DH12R</sub> (500 TCID<sub>50</sub> or more), the CD4<sup>+</sup> T-cell levels did not fall to baseline values in any of these animals but stabilized at the 3 to 12% range over the first 10 weeks of infection. Each of these macaques still generated peak plasma virus loads of  $10^7$  to  $10^8$  RNA copies/ml within 2 to 3 weeks of infection. The plasma viral RNA loads subsequently fell in all five of the macaques but, starting at week 10, began to rise in one monkey (H521) and were accompanied by a further decline of its CD4<sup>+</sup> T cells to less than 1% at week 16. Macaque H521 was subsequently euthanized at week 24 because of anorexia and marked weight loss.

The remaining four monkeys inoculated with low doses of SHIV<sub>DH12R</sub> were alive and clinically asymptomatic at 4 years after their initial exposure to virus. Two distinct patterns of circulating CD4<sup>+</sup> T cells were observed. For two of these chronically infected monkeys (6071 and 6074), the CD4<sup>+</sup> T-lymphocyte numbers have remained depressed (10 to 14%) for nearly 4 years whereas, for animals 6049 and H520, they have climbed back to near preinoculation levels (Fig. 2, upper panel). The recovery or continued maintenance of CD4<sup>+</sup> T-cell numbers following their initial loss is clearly different from the results observed with animals inoculated with 500 TCID<sub>50</sub> or

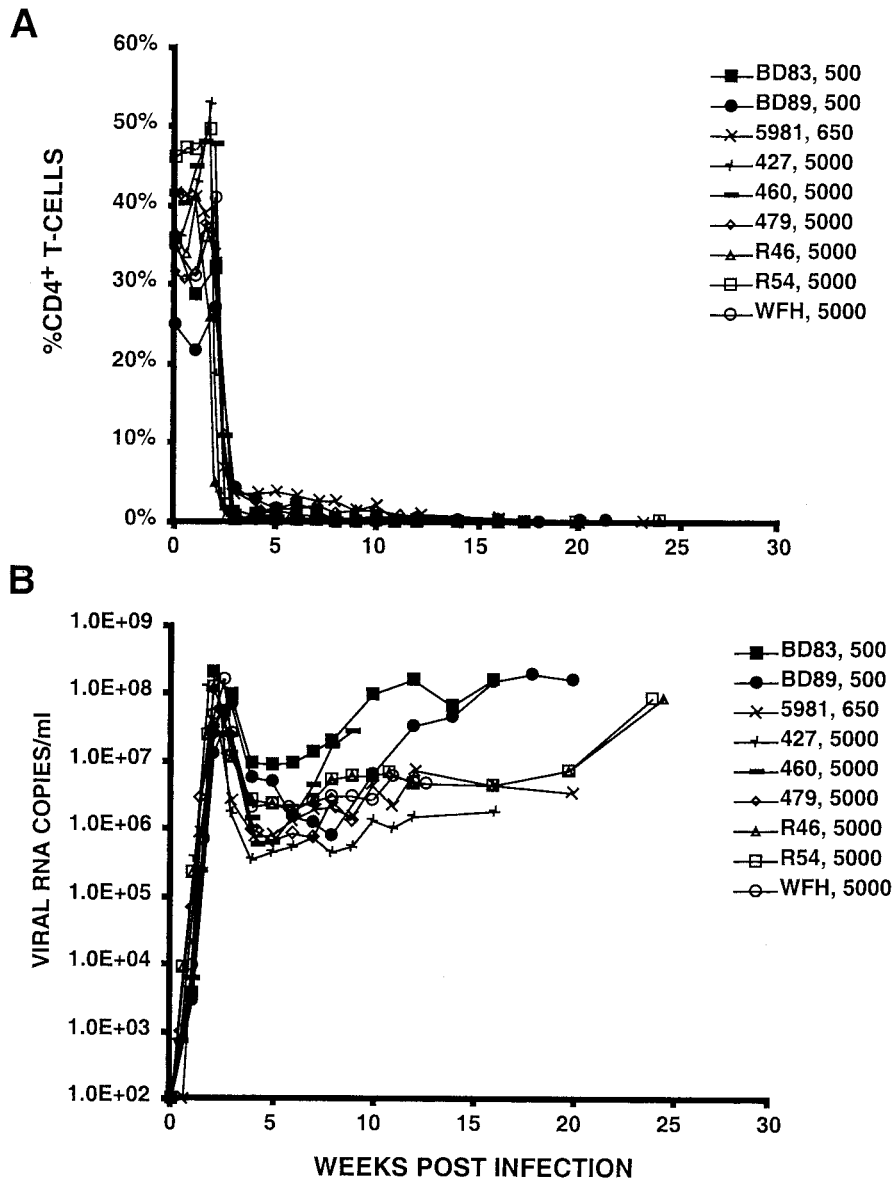


FIG. 1. Peripheral blood CD4<sup>+</sup> T-cell profiles (A) and plasma viral RNA loads (B) of SHIV<sub>DH12R</sub>-infected monkeys. Each animal was inoculated intravenously with the indicated amount (500, 650, or 5,000 TCID<sub>50</sub>) of SHIV<sub>DH12R</sub>. Peripheral blood CD4<sup>+</sup> T-cell numbers and plasma viral RNA levels were measured at the indicated times.

more of SHIV<sub>DH12R</sub>, whose depleted CD4<sup>+</sup> T-lymphocyte levels never increased over baseline levels (Fig. 1A).

It is worth noting that all of the macaques inoculated with 25 TCID<sub>50</sub> or less of SHIV<sub>DH12R</sub> generated peak plasma viral RNA levels comparable to those of the recipients of 500 TCID<sub>50</sub> or more of SHIV<sub>DH12R</sub> (compare Fig. 2, lower panel inset, with Fig. 1). Following the initial peak, the plasma viral RNA levels eventually became undetectable in three of the monkeys (6049, 6074, and H520), with only sporadic bursts of viremia that never exceeded 2 × 10<sup>3</sup> RNA copies/ml. The plasma viral RNA loads stabilized in the range of 10<sup>4</sup> to 10<sup>5</sup> RNA copies/ml in the fourth chronically infected macaque (6071).

Animals experiencing the complete and irreversible depletion of CD4<sup>+</sup> T cells following inoculation of SHIV<sub>DH12R</sub> are

unable to mount humoral immune responses (13). In contrast, rhesus monkeys infected with the nonpathogenic SHIV<sub>DH12</sub>, which carries the *env* gene from the HIV-1<sub>DH12</sub> primary isolate, typically experience a very modest and short-lived loss of CD4<sup>+</sup> T lymphocytes and generate robust titers of anti-virus NAbs (25). This is illustrated in Table 1 for macaque TPP, which was originally inoculated intravenously with a large SHIV<sub>DH12</sub> inoculum (1.6 × 10<sup>6</sup> TCID<sub>50</sub>) yet experienced only a transient decline of its CD4<sup>+</sup> T lymphocytes from 41% (pre-challenge) to 20% at week 2; by week 5 postinfection, its percent CD4<sup>+</sup> T-cell level had climbed to 35%, returning to preinoculation values by week 20. This animal produced detectable NAbs by week 3 postinoculation and titers greater than 1:1,000 by week 10, using an assay that measured complete neutralization of virus (38). The levels of NAbs in the

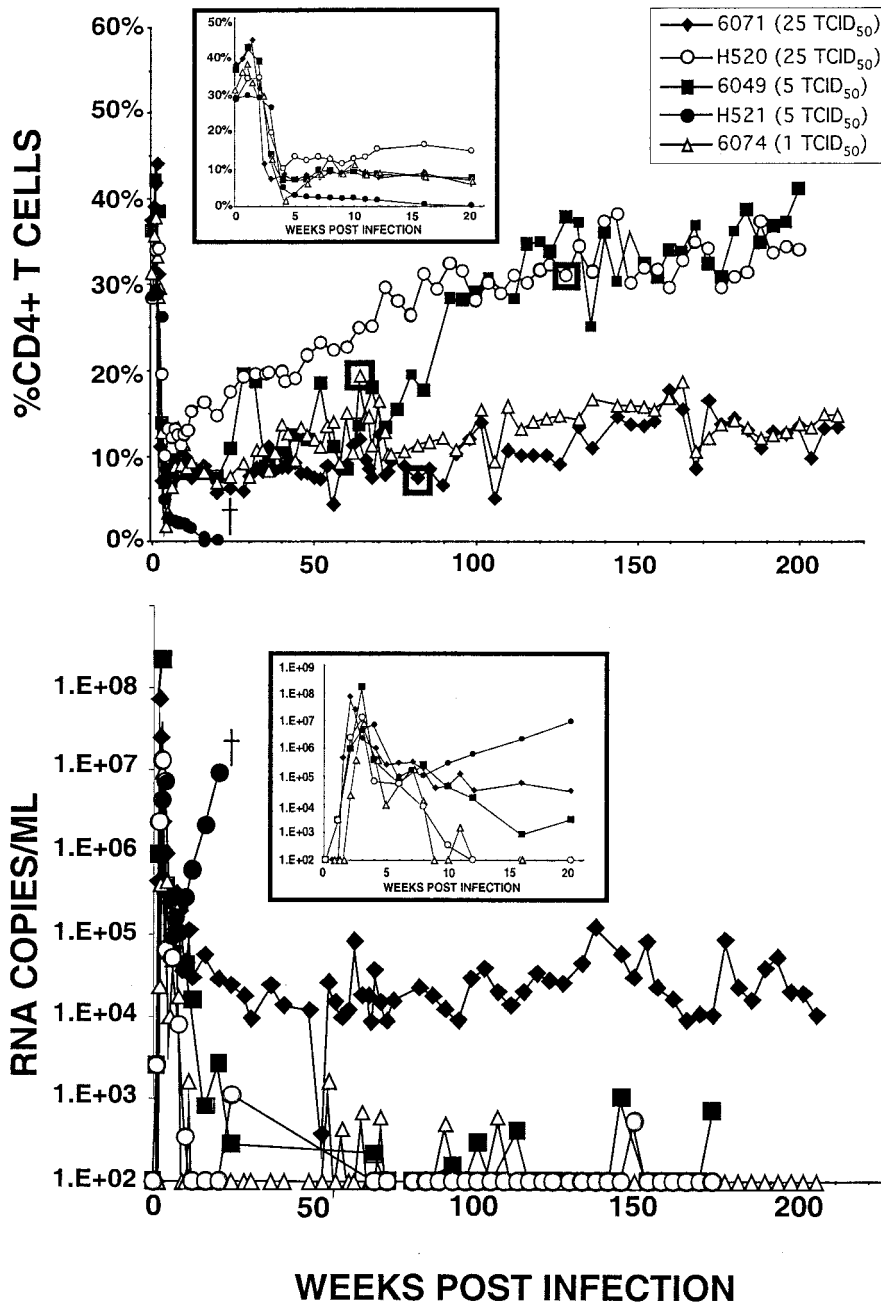


FIG. 2. Long-term effects of small intravenous SHIV<sub>DH12R</sub> inocula on disease development and control of viremia. Five macaques were inoculated intravenously with 1 to 25 TCID<sub>50</sub> of SHIV<sub>DH12R</sub>. The insets in each panel highlight percent CD4<sup>+</sup> T-cell levels (upper panel) and plasma viral RNA concentrations (lower panel) during the first 20 weeks of infection. The boxed data points in the upper panel indicate the times when virus was recovered from animal 6074 (week 64), animal 6071 (week 75), and animal H520 (week 137) by cocultivation of PBMC from each animal with PBMC from a naïve macaque.

four monkeys originally inoculated with 25 TCID<sub>50</sub> or less of the highly pathogenic SHIV<sub>DH12R</sub> were similarly determined. As shown in Table 2, all of these chronically infected animals generated NAb but at titers which were significantly lower than that measured in animal TPP. When the long-term profiles of NAb production in the four recipients of low-dose SHIV<sub>DH12R</sub> were analyzed, two distinct patterns (which paralleled those of the levels of CD4<sup>+</sup> T lymphocytes) became evident. Monkeys H520 and 6049, whose CD4<sup>+</sup> T-cell numbers

eventually returned to preinoculation levels, produced higher titers of NAb between weeks 30 and 50 postinfection than animals 6071 and 6074, which maintained persistently low levels of CD4<sup>+</sup> T cells (Table 2, Fig. 2, upper panel). Thus, although the SHIV<sub>DH12R</sub>-induced CD4<sup>+</sup> T-lymphocyte depletion was partial in the chronically infected monkeys, antibody production seemed to be significantly impaired.

The appearance of distinct long-term CD4<sup>+</sup> T-cell profiles in monkeys inoculated with low doses of SHIV<sub>DH12R</sub> suggested

TABLE 1. NAbS elicited against nonpathogenic SHIV<sub>DH12</sub><sup>a</sup>

Plasma sample taken at (wk)	Neutralizing titer against SHIV <sub>DH12</sub>
Preinfection	<1.7
2	<1.7
3	4
4	48
6	125
8	431
10	1,078
12	359

<sup>a</sup> NAb elicited in animal TPP following inoculation with  $1.6 \times 10^6$  TCID<sub>50</sub>.

the possible emergence in vivo of nonpathogenic variants that had been present in the original uncloned SHIV<sub>DH12R</sub> stock. This possibility was investigated by examining the pathogenic phenotype of the SHIVs recovered from three of the four chronically infected macaques between years 2 and 4 postinoculation from PBMC by cocultivation (Fig. 2). The viruses isolated from animals 6071 and 6074, both of which had persistently low CD4<sup>+</sup> T-cell levels, were inoculated intravenously (approximately  $10^5$  TCID<sub>50</sub>) into naïve monkeys WCH and WDC (Fig. 3). WCH and WDC rapidly and irreversibly lost CD4<sup>+</sup> T lymphocytes and died from immunodeficiency at weeks 35 and 36, respectively. This result indicates that chronically infected, but asymptomatic, monkeys 6071 and 6074 still harbored virus with a pathogenic phenotype indistinguishable from that of the original SHIV<sub>DH12R</sub>. In contrast, the SHIV recovered at week 137 from animal H520 induced a significant but incomplete decline in CD4<sup>+</sup> T lymphocyte levels following inoculation of macaque TXF (Fig. 3). By week 51, the level of CD4<sup>+</sup> T cells in animal TXF had risen to more than 13% and its plasma viremia level had fallen to  $4 \times 10^3$  RNA copies/ml.

Although one could argue that the dependence of disease development on inoculum size simply reflects the dilution of highly pathogenic variants present in the uncloned, genetically heterogeneous SHIV<sub>DH12R</sub> stock, the same phenomenon has been observed with the recently obtained molecular clone SHIV<sub>DH12R-CI-7</sub>, which consistently induces the prototypic rapid and complete CD4<sup>+</sup> T-cell depletion following the intravenous inoculation of 500 TCID<sub>50</sub> or more of virus (R. Sadjadpour, unpublished data).

Taken together, these results indicate that rhesus monkeys inoculated with small amounts of the highly pathogenic SHIV<sub>DH12R</sub> experienced a marked but incomplete loss of CD4<sup>+</sup> T cells and remained clinically asymptomatic over a nearly 4-year observation period. However, the capacity of these seemingly healthy animals (with depressed levels of CD4<sup>+</sup> T lymphocytes) to produce high titers of NAbS was impaired compared to that of infected animals with minimal or no loss of this T-cell subset. In one of the chronically infected macaques whose CD4<sup>+</sup> T-cell numbers had returned to preinoculation levels, the only virus recovered (at nearly 3 years postinfection) failed to induce the characteristic rapid and irreversible depletion of CD4<sup>+</sup> T cells. This attenuated SHIV variant, presumably present in the original uncloned SHIV<sub>DH12R</sub> inoculum, emerged following long-term replication in vivo.

**Administration of a single course of PMPA during the first weeks of SHIV<sub>DH12R</sub> infection can prevent virus-induced disease.**

The potent RT inhibitor PMPA has previously been shown to block both acute and chronic SIV infections in vivo (42, 43). When administered at 24 but not 72 h postinoculation with SIV<sub>smE660</sub>, a 28-day course of therapy prevented the establishment of infection (23). In an initial experiment to assess the efficacy of PMPA to contain highly pathogenic SHIVs, the initiation of treatment was deliberately delayed until 5 days following virus inoculation because of numerous reports showing that SHIVs can be contained by vaccine approaches that fail to control SIVs (1, 5, 6, 15, 32, 40). Four rhesus monkeys were inoculated intravenously with 1,000 TCID<sub>50</sub> of SHIV<sub>DH12R</sub> and received single daily intramuscular injections of PMPA (30 mg/kg of body weight) for 4 weeks beginning on day 5 postinoculation with virus. At the time of PMPA therapy initiation, SHIV<sub>DH12R</sub> had caused no loss of CD4<sup>+</sup> T lymphocytes, although  $\sim 10^4$  copies/ml of plasma viral RNA had already become detectable (Fig. 4, right inset in lower panel). Compared to the results for untreated acutely SHIV<sub>DH12R</sub>-infected monkeys (Fig. 1), which experienced a rapid depletion of their CD4<sup>+</sup> T cells and symptoms of immunodeficiency requiring euthanasia, the four recipients of PMPA experienced no appreciable loss of CD4<sup>+</sup> T lymphocytes during the first 20 weeks postinfection (Fig. 4, upper panel inset). Peak plasma virus loads (ranging from  $1.4 \times 10^4$  to  $8.8 \times 10^4$  RNA copies/ml)

TABLE 2. NAbS elicited against pathogenic SHIV<sub>DH12R</sub>

Plasma sample taken (wk)	Neutralizing titer against SHIV <sub>DH12R</sub> for animal:				
	6071 (25 TCID <sub>50</sub> )	H520 (25 TCID <sub>50</sub> )	H521 (5 TCID <sub>50</sub> )	6049 (5 TCID <sub>50</sub> )	6074 (1 TCID <sub>50</sub> )
Preinfection	<0.8	<3.1	<3.1	<3.1	<0.8
10	<0.8	16	<3.1	<3.1	<0.8
20	4	21	<1.6	7	11
30	12	21		63	13
40	7	44		76	8
50	5	34		ND	7
60	7	ND		ND	8
70	8	ND		ND	8
92	ND <sup>a</sup>	4		ND	ND
126	67	ND		ND	12
184	ND	12		ND	ND
212	7	ND		ND	12

<sup>a</sup>ND, not determined.

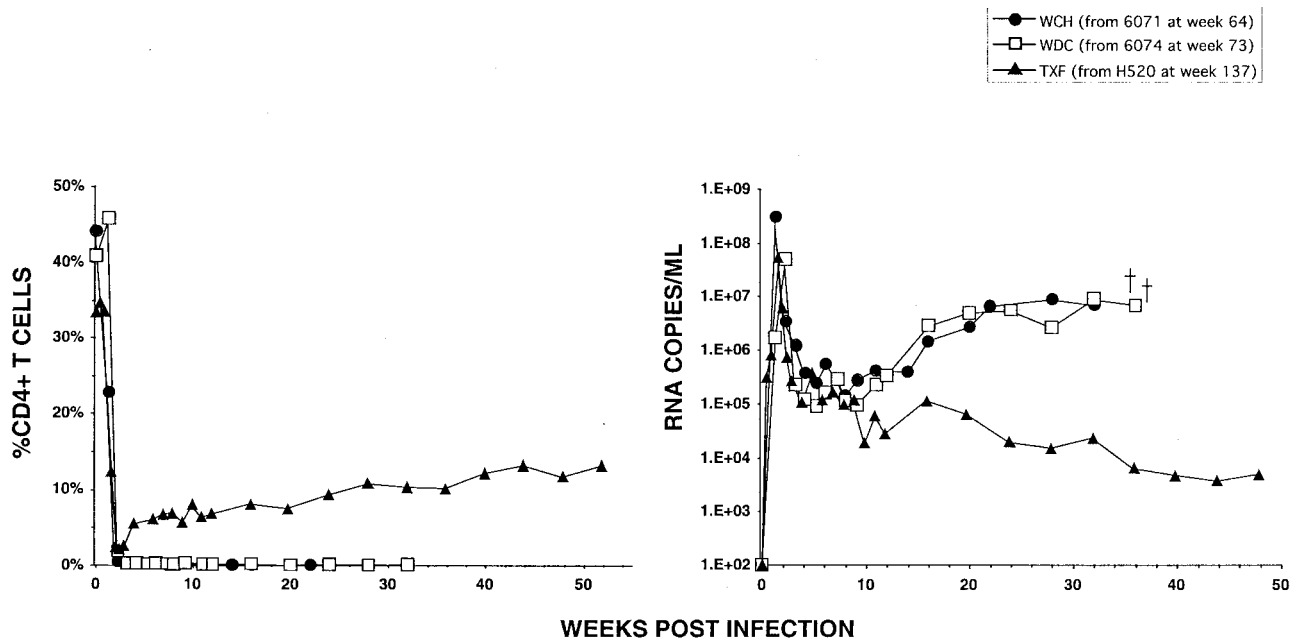


FIG. 3. Virus recovered from monkeys persistently infected with SHIV<sub>DH12R</sub> exhibited divergent pathogenic phenotypes. SHIVs isolated from animals 6074, 6071, and H520 (Fig. 2) were propagated *in vitro* and inoculated intravenously into monkeys WDC ( $7 \times 10^5$  TCID<sub>50</sub>), WCH ( $2.9 \times 10^4$  TCID<sub>50</sub>), and TXF ( $3.9 \times 10^5$  TCID<sub>50</sub>), respectively.

rapidly fell below the detection threshold ( $<200$  RNA copies/ml) during the period of PMPA administration and rebounded to somewhat higher levels following the cessation of treatment. In three of the four macaques (N077, BC74, and BI36), plasma viral RNA loads gradually declined to background levels by week 20 postinoculation (Fig. 4, lower panel). In the fourth animal (N056), the plasma viremia level fell approximately 10-fold between weeks 8 and 13 but, by week 20, had climbed to  $1.4 \times 10^5$  RNA copies/ml. The CD4<sup>+</sup> T-cell levels in this monkey also began to decline between weeks 20 and 40, reaching 10 cells/mm<sup>3</sup> on week 81, at which time the animal was euthanized because of anorexia, diarrhea, and marked weight loss. Nonetheless, three of the four rhesus monkeys receiving a single 4-week course of PMPA beginning on day 5 postinfection with SHIV<sub>DH12R</sub> maintained their CD4<sup>+</sup> T cells at preinoculation levels for nearly 3 years and readily controlled their virus loads during this period (Fig. 4). This clinical outcome is clearly different from that reported for SIV-infected monkeys in a study in which PMPA treatment was started more than 24 h postinoculation with virus (23).

In view of the success at maintaining CD4<sup>+</sup> T-lymphocyte levels and containing viremia (both acute and chronic) when PMPA treatment was started prior to the peak of virus production and before any loss of CD4<sup>+</sup> T cells was detectable, we next inoculated two rhesus monkeys (96N130 and 96N135) intravenously with 1,000 TCID<sub>50</sub> of SHIV<sub>DH12R</sub> but did not initiate daily PMPA administration until day 14 postinoculation with virus. At this point in the infection, the CD4<sup>+</sup> T-lymphocyte levels had begun to fall and the plasma virus loads ( $1.2 \times 10^8$  and  $5.2 \times 10^7$  RNA copies/ml [Fig. 5 insets]) were comparable to those measured for the group of untreated animals (Fig. 1). The subsequent 4-week course of PMPA therapy potently repressed the plasma viremia to background lev-

els in animal 96N135 and partially controlled virus loads in macaque 96N130. During the first 18 weeks of infection, both of the PMPA-treated monkeys experienced severe but partial depletions of their CD4<sup>+</sup> T cells, avoiding the irreversible and complete loss of this T-lymphocyte subset suffered by the untreated animals (compare Fig. 5 insets with Fig. 1). Although the CD4<sup>+</sup> T-cell levels stabilized at the 4 to 6% range in macaque 96N130, they declined even further after week 18 as plasma virus loads began to rise. Ultimately, this animal had to be euthanized at week 85 because of marked weight loss and intractable diarrhea (Fig. 5). This clinical course contrasted with that of monkey 96N135, which maintained excellent control of its plasma viremia over a 3-year observation period and whose CD4<sup>+</sup> T cells returned to their preinoculation levels.

To investigate the possible emergence of nonpathogenic variant virus in animal 96N135, whose CD4<sup>+</sup> T cells had returned to preinoculation levels, virus was isolated at weeks 30 and 46 from animals 96N130 and 96N135, respectively (Fig. 5), and used to inoculate (at  $10^4$  TCID<sub>50</sub>) two naïve macaques. As shown in Fig. 6, the virus recovered from animal 96N130, which had been unable to effectively control its viremia and ultimately died from immunodeficiency, caused the characteristic rapid and irreversible loss of CD4<sup>+</sup> T cells in monkey 3GD and death at week 36. In contrast, animal AM32, the recipient of virus isolated from healthy monkey 96N135, experienced only a modest and transient depletion of its CD4<sup>+</sup> T lymphocytes, which remained at near preinoculation levels for more than 2 years. Furthermore, the viremia in this monkey was rapidly suppressed and became undetectable by week 14 postinfection.

Taken together, these experiments indicate that antiretroviral therapy, when begun prior to peak viremia and before demonstrable depletion of CD4<sup>+</sup> T lymphocytes, was able to durably control virus replication and prevent disease induction

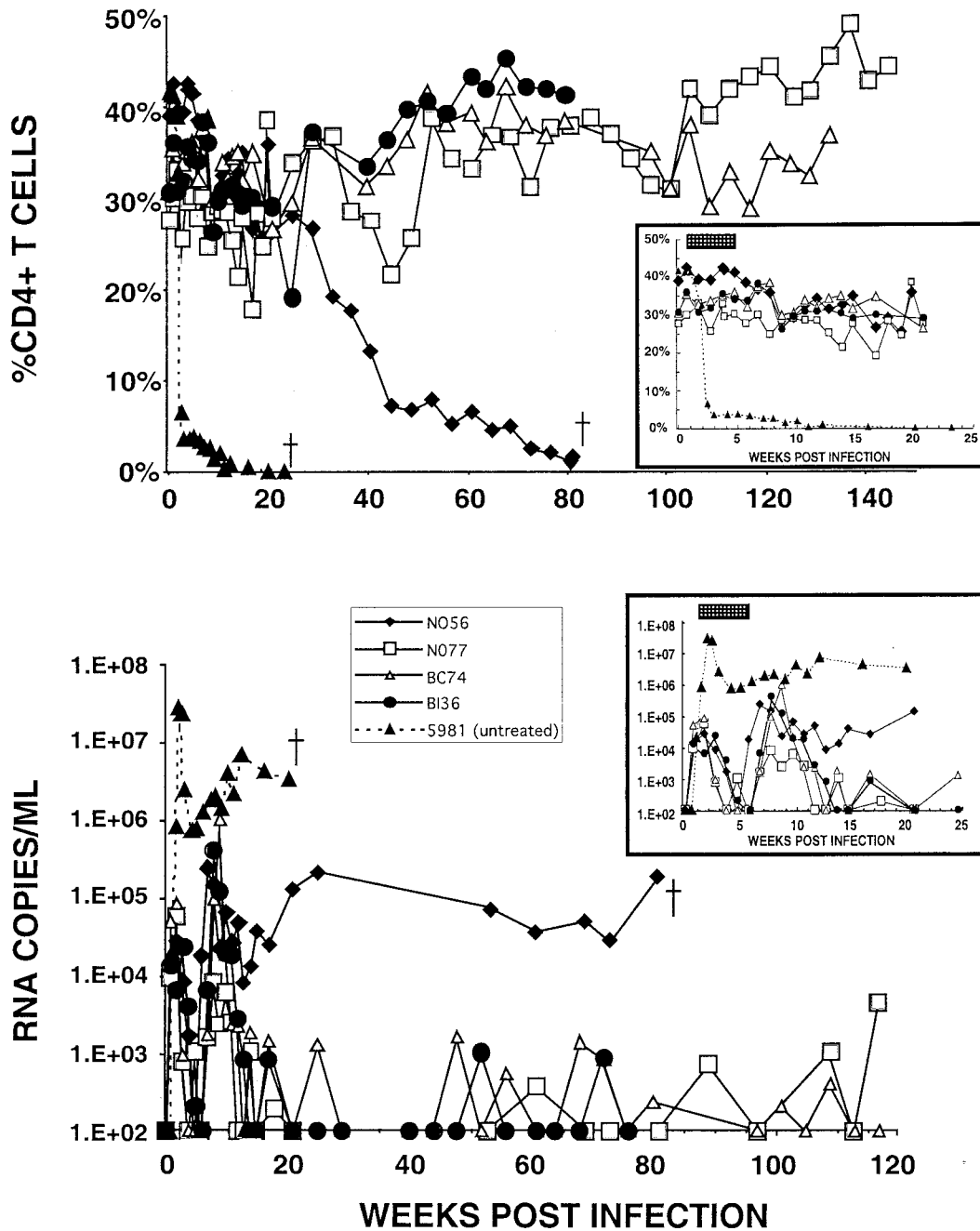


FIG. 4. A single 4-week course of PMPA treatment initiated on day 5 postinfection prevented irreversible CD4<sup>+</sup> T-cell depletion and durably controlled virus replication in three of four treated animals. Four rhesus monkeys (N056, N077, BC74, and BI36) were inoculated intravenously with 10<sup>3</sup> TCID<sub>50</sub> of the SHIV<sub>DH12R</sub> stock and treated for 4 weeks with PMPA (30 mg administered intramuscularly per day/kg of body weight) beginning on day 5 postinfection. An untreated monkey (5981) was inoculated intravenously with 650 TCID<sub>50</sub> of SHIV<sub>DH12R</sub>. The insets in each panel highlight percent CD4<sup>+</sup> T-lymphocyte (upper panel) and the plasma viral RNA (lower panel) levels during the initial 25 weeks of infection. The stippled bar in each inset indicates the 4-week period of PMPA administration.

in three of four monkeys inoculated with the highly pathogenic SHV<sub>DH12R</sub>. Starting PMPA administration at the time of peak viremia or after some loss of CD4<sup>+</sup> T lymphocytes had occurred was less effective but did block the characteristic SHIV-induced disease in one of two infected macaques. In the latter case, the healthy animal harbored virus exhibiting an attenuated pathogenic phenotype.

**SHIV<sub>DH12R</sub> utilized CXCR4, not CCR5, to enter monkey PBMC.** The distinctive patterns of disease induction and sensitivity to intervention by both antiviral agents and vaccines raised the possibility that SHIV<sub>DH12R</sub> and SIVs target different subsets of T lymphocytes during the initial phase(s) of in vivo infection. We therefore examined the chemokine coreceptor dependence of both for replication in rhesus monkey PBMC.

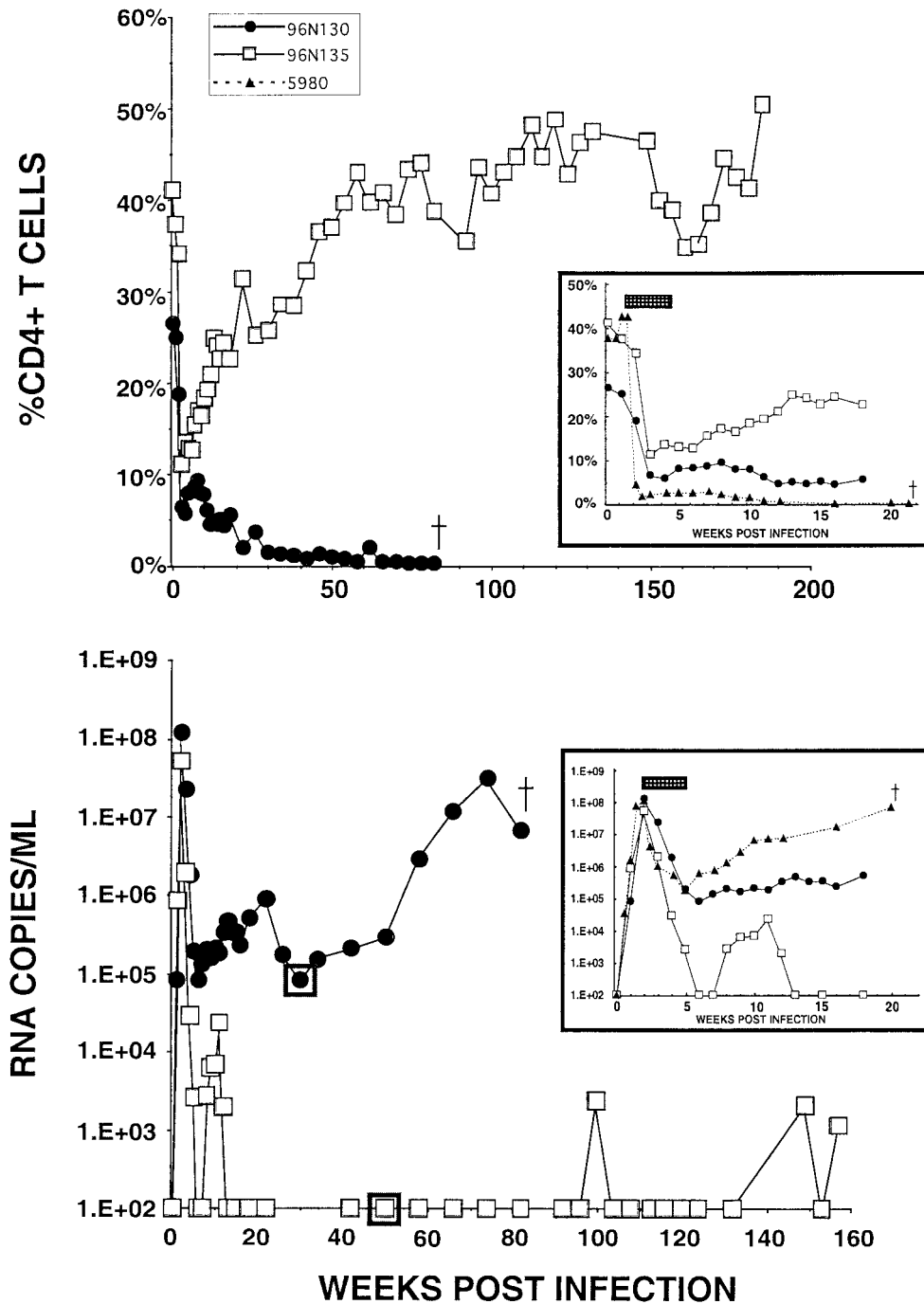


FIG. 5. Administration of PMPA to two rhesus monkeys beginning on day 14 postinfection. Two monkeys (96N130 and 96N135) were inoculated intravenously with  $10^3$  TCID<sub>50</sub> of the SHIV<sub>DH12R</sub> stock and treated for 4 weeks with PMPA (30 mg administered intramuscularly per day/kg of body weight) beginning on day 14 postinfection. An untreated monkey (5980) was inoculated intravenously with  $1.6 \times 10^4$  of SHIV<sub>DH12R</sub>. The insets in each panel highlight percent CD4<sup>+</sup> T-lymphocyte (upper panel) and the plasma viral RNA (lower panel) levels during the initial 20 weeks of infection. The stippled bar in each inset indicates the 4-week period of PMPA administration. The boxed data points in the lower panel indicate the times when virus was recovered from animal 96N130 (week 30) or 96N135 (week 46) by cocultivation of its PBMC with PBMC from a naïve macaque.

In these experiments, virus was spinoculated (30) onto PBMC in the presence of small-molecule coreceptor-targeted inhibitors specific for CCR5 or CXCR4. The production of progeny virus was measured as levels of RT activity released into the medium on day 5 postinfection. As shown in Fig. 7, infections of monkey PBMC by the T-cell-tropic SIV<sub>mac239</sub> and the mac-

rophage-tropic SIV<sub>mac316</sub> were both blocked by two different CCR5 inhibitors (TAK-779 and AD-101) but not by the CXCR4 inhibitor AMD3100. For SIV<sub>mac239</sub>, this result is in agreement with a previous report showing that CCR5 is the coreceptor used to infect macaque PBMC (51). The same assay was also performed with two different DH12-related SHIVs



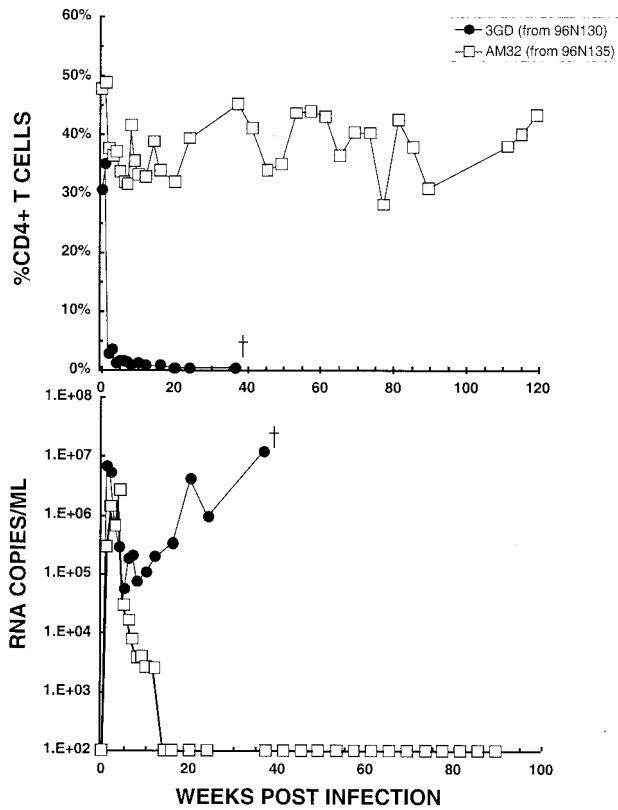


FIG. 6. Virus recovered from monkeys treated with PMPA starting on day 14 postinfection exhibited divergent pathogenic phenotypes in vivo. The SHIV isolates recovered from animal 96N130 at week 30 and animal 96N135 at week 46 (boxed data points in Fig. 5) were propagated in vitro, and  $10^4$  TCID<sub>50</sub> of each isolate was inoculated intravenously into macaques 3GD and AM32, respectively.

(SHIV<sub>DH12R</sub> and SHIV<sub>DH12R-CL-7</sub>, a recently obtained molecular clone that reproducibly induces a rapid, complete, and irreversible depletion of CD4<sup>+</sup> T cells in inoculated rhesus monkeys) (18) (R. Sadjadpour et al., unpublished data). For both SHIVs, the opposite outcome was observed: infection of PBMC was completely blocked by AMD 3100 and not by the CCR5 inhibitors. Some competition was observed with TAK-779 but only at the highest concentrations tested (5 to 10  $\mu$ M), perhaps reflecting nonspecific effects of this compound. These results indicate that both SIVs utilized CCR5 and SHIV<sub>DH12R</sub> used CXCR4 to enter and spread through cultured rhesus PBMC.

DISCUSSION

The results of this study show that (i) the immunodeficiency caused by highly pathogenic SHIV<sub>DH12R</sub> is dependent on inoculum size; (ii) SHIV-induced disease can be prevented by antiretroviral drug therapy initiated several days following exposure to virus; (iii) when the rapid and complete depletion of CD4<sup>+</sup> T cells induced by SHIV<sub>DH12R</sub> fails to occur, a majority of infected animals experience an asymptomatic clinical course; and (iv) highly pathogenic SHIV<sub>DH12R</sub> exclusively uses CXCR4 to enter macaque PBMC, whereas SIV<sub>mac239</sub> and SIV<sub>mac316</sub> utilize CCR5 to infect the same cells. When considered together with the strikingly different patterns of CD4<sup>+</sup> T-lymphocyte depletion and the temporal development of disease, these results suggest that the immunodeficiencies caused by SIV and highly pathogenic SHIVs are mechanistically dissimilar.

For SHIVs, clinical outcomes can be readily and durably modulated by therapeutic and vaccine interventions. In one of the longitudinal studies described in the present report, three of four macaques remained free of disease for 2 to 3 years following a single 28-day course of treatment (initiated on day

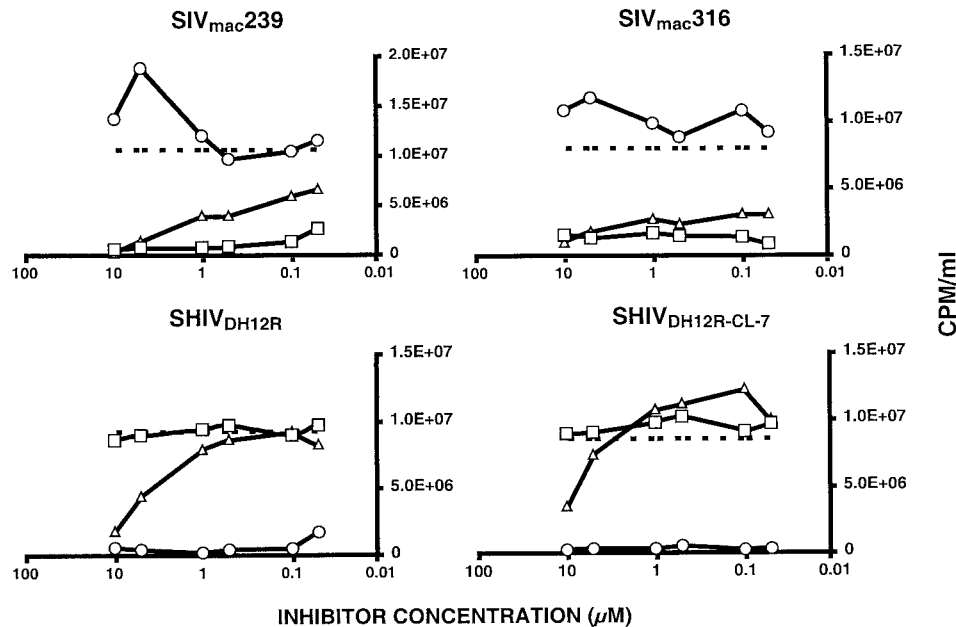


FIG. 7. Coreceptor usage by SHIVs and SIVs. SHIV<sub>DH12R</sub>, SHIV<sub>DH12R-CL-7</sub>, SIV<sub>mac239</sub>, and SIV<sub>mac316</sub> were spinoculated onto rhesus PBMC in the presence of the indicated amounts of small-molecule chemokine receptor inhibitors. RT activity released into the medium on day 5 postinfection was determined in the absence (dashed line) or presence of inhibitor. Lines with circles represent AMD 3100; lines with triangles represent TAK-779; lines with squares represent AD-101.

5 postinoculation with virus) with a nonnucleoside RT inhibitor. Similarly, long-term monitoring of several animals exposed to 25 TCID<sub>50</sub> or less of SHIV<sub>DH12R</sub> indicated that they remained disease-free, with low to undetectable plasma virus loads, during a 4-year observation period. The simplest model consistent with all of the data is that a threshold level of systemic virus production which causes massive and systemic killing of all CD4<sup>+</sup> T-lymphocyte subsets in lymphoid tissues must be reached within the first week of a SHIV<sub>DH12R</sub> infection. A delay (caused by, e.g., anti-retroviral drugs, vaccines, or a low level of inoculum) of only a few days in achieving this threshold total body virus burden effectively abolishes the irreversible loss of CD4<sup>+</sup> T cells, presumably by allowing time for an immune response to develop and control virus replication (22). If the elimination of CD4<sup>+</sup> T lymphocytes is not complete, systemic viral loads will be contained and a majority of infected animals will remain asymptomatic for extended periods of time. While the prevention of SHIV-induced disease by PMPA treatment initiated on day 5 can be attributed to the direct suppression of virus replication by a potent RT inhibitor, the attenuating effect of low inoculum size is less understandable. Infections initiated with small amounts of pathogenic SHIVs are likely to be intrinsically slower because they require several additional replication cycles to catch up to those started with high virus inputs. The correspondingly slower kinetics of virus-induced cell killing during the critical first week of infection may therefore preserve sufficient numbers of CD4<sup>+</sup> T-cell activity to control virus replication.

The clinical features of SIV-induced immunodeficiency are quite different. First, although SIV can cause reductions in peripheral blood CD4<sup>+</sup> T-cell levels during acute infections, the depletions observed are relatively modest and never approach the complete and systemic loss observed in SHIV-infected monkeys (20, 36). More importantly, the clinical symptoms of immunodeficiency in SIV-infected animals do not require the total elimination of CD4<sup>+</sup> T lymphocytes but can become apparent at levels of 200 to 300 cells/ $\mu$ l or higher (10, 24). Second, the induction of AIDS in SIV-infected macaques does not appear to be dose dependent. Animals inoculated with 1:50- or 1:1,000-fold dilutions of a SIV<sub>mac251</sub> stock died from immunodeficiency within similar time frames (an average of 218 or 275 days, respectively) (9). In a separate study, several pig-tailed macaques infected with SIV<sub>smE660</sub> (0.3 to 30,000 50% monkey infective doses) or SIV<sub>sm543</sub> (0.6 to 600 50% monkey infective doses) developed disease in 5 to 25 or 3 to 19 months, respectively, independent of inoculum size (14). Finally, anti-retroviral agents appear to be less effective in the SIV-macaque system. When the most widely employed and effective agent, PMPA, was used and treatment was started at 24 h postinoculation with SIV<sub>smE660</sub>, plasma viremia became undetectable during and following the cessation of therapy (23). However, when PMPA administration was delayed until day 3 of the SIV infection, plasma virus loads increased significantly after treatment was stopped, reaching the levels measured in control monkeys. Although no information pertaining to final clinical outcomes was provided in that study, the results reported contrast with those observed for the SHIV-infected-PMPA-treated animals, three of four of which remain disease-free, with undetectable levels of plasma viral RNA, even

though the single course of PMPA treatment was delayed until 5 days postinoculation with virus.

The results obtained with inhibitors that target specific chemokine receptors may be key in explaining the different disease phenotypes elicited by SHIVs and SIVs: SHIVs use CXCR4 for entry of rhesus monkey PBMC, while SIVs use CCR5. These chemokine receptors are differentially expressed on leukocyte subsets and in individual body compartments. In monkey and human PBMC, for example, 80 to 95% of CD4<sup>+</sup> T cells express CXCR4 while only 5 to 10% produce detectable CCR5 (2, 46, 50) (Y. Nishimura, unpublished observation). A similar pattern of chemokine receptor expression has been reported for CD4<sup>+</sup> T lymphocytes in lymph nodes and spleen (46, 47). The vast majority of these CXCR4<sup>+</sup> CD4<sup>+</sup> cells share surface molecule expression patterns with naive CD4<sup>+</sup> T lymphocytes (29). In contrast, a large fraction of CD4<sup>+</sup> T cells in other organs, including the gut-associated lymphoid tissue, express CCR5 and memory lymphocyte markers (37, 46); many of these tissue-infiltrating memory CD4<sup>+</sup> T cells are also activated (26, 48). Rhesus monkeys inoculated with SIV<sub>mac</sub> typically experience a profound and selective loss of CD4<sup>+</sup> CCR5<sup>+</sup> memory T lymphocytes in the gut-associated lymphoid tissue during the first few weeks of infection and only modest depletions of CD4<sup>+</sup> T cells in the blood and peripheral lymphoid organs, the bulk of which are CXCR4<sup>+</sup> CCR5<sup>-</sup> naive cells (12, 45, 46, 48). In contrast, the expression of CXCR4 on virtually all naive and a significant fraction of memory CD4<sup>+</sup> T cells (29) (Y. Nishimura, unpublished observation) makes them prime targets of highly pathogenic SHIVs and could explain the unrelenting and rapid pattern of their depletion in SHIV, but not, SIV infections.

Thus, instead of causing the irreversible elimination of CD4<sup>+</sup> T cells in the blood and lymphoid tissues, SIV selectively and systemically depletes the CD4<sup>+</sup> CCR5 memory subset in infected monkeys. The loss of these cells results in a reduced capacity to produce gamma interferon, tumor necrosis factor alpha, and IL-2, following in vitro stimulation with phorbol myristate acetate plus ionomycin, within 2 to 3 weeks postinfection (27). This memory CD4<sup>+</sup> T-cell dysfunction persists for several months and worsens in those animals developing clinical disease.

When considered in the context of HIV-1 infections of humans, it is tempting to speculate that most infected individuals are exposed to HIV-1 inocula containing both syncytium-inducing CXCR4- and non-syncytium-inducing CCR5-utilizing variants. The X4 component may be responsible for the modest depletion of CD4<sup>+</sup> T cells that has been reported to accompany most acute HIV-1 infections (36). However, as CCR5-using variants become predominant, these CXCR4-utilizing HIV-1 strains may be readily suppressed and rapidly become undetectable in recently infected individuals (34, 52). Such a scenario is consistent with a report showing that syncytium-inducing variants of HIV-1 (presumably X4 virus), which were recovered during primary infections of two individuals following accidental parenteral virus transmissions, were subsequently supplanted in both patients by non-syncytium-inducing (presumably R5) virus (7). The slowly developing immune dysfunction (associated with chronic HIV-1 infection of several years' duration) could eventually permit archival CXCR4-utilizing virus to emerge in some infected individuals

and contribute to the accelerated decline of CD4<sup>+</sup> T cells during the late stages of disease.

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