Immortalization of CD4+ and CD8+ T Lymphocytes by Human T-Cell Leukemia Virus Type 1 Tax Mutants Expressed in a Functional Molecular Clone

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The human T-cell leukemia virus type 1 (HTLV-1) transcriptional trans-activator Tax has been demonstrated to have transforming activity in multiple cell culture and transgenic-mouse models. In addition to activating transcription from the viral long terminal repeat (LTR) through the cyclic AMP response element (CREB/ATF) family of transcription factors, Tax activates the expression of multiple cellular promoters through the NF-κB pathway of transcriptional activation. The Tax mutants M22 and M47 have previously been demonstrated to selectively abrogate the ability of Tax to activate transcription through the NF-κB or CREB/ATF pathway, respectively. These mutations were introduced in the tax gene of the ACH functional molecular clone of HTLV-1, and virus produced from the mutant ACH clones was examined for the ability to replicate and immortalize primary human lymphocytes. While virus derived from the clone containing the M47 mutation retained the ability to immortalize T lymphocytes, the M22 mutant lost the ability to immortalize infected cells. These results indicate that activation of the CREB/ATF pathway by Tax is dispensable for the immortalization of T cells by HTLV-1, whereas activation of the NF-κB pathway may be critical.

The human T-cell leukemia virus type 1 (HTLV-1) infects and immortalizes T lymphocytes in vitro and is the causative agent of adult T-cell leukemia/lymphoma and HTLV-1-associated myelopathy/tropical spastic paraparesis in vivo (16, 22, 53, 57). The HTLV-1 genome encodes a 40-kDa transcriptional trans-activator known as Tax, which has been demonstrated to have transforming activity (2, 17, 43, 63, 70, 78). In addition to activating transcription from the viral long terminal repeat (LTR) (10, 13, 27, 66), Tax activates the expression of a number of cellular genes, many of which either encode proteins involved in the regulation of cellular proliferation (i.e., interleukin 2 [IL-2] [68], IL-2 receptor α chain [3, 25, 40, 68], and proliferating cell nuclear antigen [PCNA] [58]) or are proto-oncogenes (c-fos [14] and c-sis [75]). Furthermore, Tax alters the activity of a number of cell cycle regulators, including cyclin D (49), the mitotic checkpoint regulator MAD1 (30), the cyclin-dependent kinases (Cdk) Cdk4 and Cdk6 (65), the Cdk inhibitor p16INK4a (41, 72), and the tumor suppressor p53 (46, 55, 56). Thus, it is likely that Tax disregulates the cell cycle through many different mechanisms, leading to the eventual immortalization and transformation of the infected cell.

Tax activates transcription through a number of different transcription factors, including cyclic AMP response element (CREB/ATF) (4, 7, 8, 74, 80), NF-κB (3, 15, 24, 32, 44), p65 serum response factor (SRF) (14, 71), Sp1 (75), and NGFI-A/Egr-1 (75). Tax activates transcription through the CREB/ATF pathway by at least two distinct mechanisms. First, Tax binds CREB1 and increases the affinity of CREB1 for the three 21-bp repeats in the HTLV-1 LTR, which contain nonconsensus CREB response elements and are involved in Tax-activated and basal expression of the LTR (1, 5, 39, 80, 81). Second, Tax interacts with the CREB transcriptional coactivators CREB-binding protein (CBP) and p300, thereby recruiting them to the CREB–Tax–21-bp repeat complex (6, 20, 37). Likewise, Tax has been shown to interact with various members of the NF-κB family and their inhibitors, including p50, p100, and p105 (23, 47, 64). Recent evidence suggests that Tax also increases NF-κB activity by increasing the activity of mitogen-activated protein/extracellular signal-related kinase kinase 1 (MEKK1) and NF-κB-inducing kinase (NIK), which phosphorylate and activate the IκB kinases IKKα and IKKβ (15, 79). The IKK kinases then phosphorylate the NF-κB inhibitors IκBα and IκBβ, leading to IκB degradation and nuclear translocation of the active NF-κB subunits (44, 45).

A number of mutations in Tax have been described which selectively abrogate the ability of Tax to upregulate transcription through the CREB/ATF and NF-κB transcriptional activation pathways (67, 69, 78). Two of these mutations, termed M22 and M47, have been extensively characterized and were chosen for examination in this study (69). The M22 mutation is a double-amino-acid substitution of an alanine and serine for the threonine and leucine amino acids at positions 130 and 131, respectively. The M22 mutant is defective for CREB/ATF activation, but the mutation has only a minimal effect on CREB/ATF activity (69). Conversely, the M47 mutation is a substitution of arginine and serine for the two leucine amino acid residues at positions 319 and 320, and the mutant is defective for CREB/ATF activation while retaining the ability to activate NF-κB (69). In addition to M22 and M47, other Tax point mutations have been described which also selectively abrogate CREB/ATF (C29S, H52Q, L296G, and L320G) or NF-κB (C23S, S258A, and G148V) activation (67, 78).

The effects of these mutations on the ability of Tax to immortalize or transform various cell types has been examined, including established rat fibroblast cell lines and primary hu-
man peripheral blood mononuclear cells (PBMC) (2, 62, 70, 78). However, the results of these studies have been conflicting, and no overall consensus has been reached as to whether Tax activation of the CREB/ATF or NF-kB pathway is critical for cellular immortalization or transformation. In this study, we examined the effects of these mutations on the ability of an infectious molecular clone of HTLV-1 to immortalize primary human PBMC. These results indicate that while CREB/ATF activation is dispensable for cellular immortalization, NF-kB activation appears to be important. Furthermore, activation of CREB/ATF by Tax may be important in the preferential immortalization of CD4+ T cells by HTLV-1.

**MATERIALS AND METHODS**

**Construction of ACH Tax mutants.** Construction of the ACH molecular clone of HTLV-1 has been previously described (34). Tax expression clones containing wild-type Tax (pcTax), the M22 mutation, and the M47 mutation were generous gifts from Warner C. Greene (61, 69). The Tax expression clones were digested with AccI and Smal (New England Biolabs, Beverly, Mass.), and a 961-bp fragment encoding amino acids 18 to 336 of Tax was cloned into an AccI-Smal-digested ACH subclone (ACH2Kml), which consists of nucleotides 6121 to 11813 of the ACH plasmid. A 2,344-bp Nol-EcoRI fragment from ACH2Kml was then cloned into an Nol-EcoRI-digested ACH to generate ACH.pcTax, ACHM22, and ACH.M47.

**Viral particle production assay.** Human 29T kidney fibroblasts seeded to 30% confluence in 10-cm-diameter dishes were transfected with 10 µg of ACH, pcTax, ACHM22, ACH.M47, or pBlueScript KS (Stratagene, La Jolla, Calif.) (empty vector) with 30 µl of Lipofectamine (Life Technologies, Gaithersburg, Md.) in OPTI-MEM I (Life Technologies). After transfection, the cells were cultured in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal calf serum (FCS), 2 mM L-glutamine, 50 µM of penicillin/ml, and 50 U of streptomycin/ml. Cell culture supernatants were collected at various time points posttransfection, and virus particle production was monitored by p19 enzyme-linked immunosorbent assay (ELISA) (Cellular Products, Buffalo, N.Y.) according to the manufacturer’s instructions.

**Transfection of PBMC.** Human PBMC were purified from healthy donors by Ficoll-Paque (Pharmacia, San Diego, Calif.) centrifugation and activated for 72 h with 10 µg of phytohemagglutinin-P (PHA) (Sigma, St. Louis, Mo.)/ml and 50 U of recombinant human IL-2/ml in RPMI 1640 medium supplemented with 10% FCS, 2 mM L-glutamine, 50 µg of penicillin/ml, and 50 U of streptomycin/ml. Cells were electroporated with 25 µg of ACH plasmid at 250 V, 1,800 µF capacitance, and 720Ω resistance with an ECM 680 electroporator (BTX, San Diego, Calif.). The cells were then cultured in RPMI 1640 medium supplemented with 5 µg of PHA/ml and 50 U of IL-2/ml for 6 weeks, after which they were cultured in the same medium without PHA. At various time points, cell culture supernatants were collected for p19 determination, and relative cellular viability was assessed by MTT conversion assays on 100 µl aliquots of cells as described by Hansen et al. (19).

**PCR amplification of Tax.** Genomic DNA was purified from ACH.pcTax- and ACH.M47-immortalized cell lines by the Wizard Prep genomic DNA purification kit (Promega, Madison, Wis.) according to the manufacturer’s instructions. PCR (95°C for 1 min; 60°C for 2 min; 72°C for 3 min; 35 cycles) was performed on 200 ng of genomic DNA with the following primers: 5′-CGGAA TTCATGCCCCACATCCGGGTTTGG-3′ and 5′-CCGGAATACTTCCATGGCCCACTTCCCAGGGTTTGG-3′, which amplifies the entire Tax open reading frame (ORF). The PCR product was then digested with BglII to assess the presence of the M47 mutation. Alternatively, the PCR product was digested with AccI and Smal and cloned into AccI-Smal-digested Tax expression plasmid pEX (kindly provided by O. John Semmes). For activity determination, 5 µg of each Tax expression clone was cotransfected with 1 µg of HTLV LTR-luciferase in 29T cells with Lipofectamine. The cells were harvested 24 h posttransfection, and luciferase activity was measured in 30 µg of whole-cell extract as described above. Transfection efficiency was normalized by measurement of CAT activity in 3 µg of cell extracts. CAT activity was determined by phase extraction of butyryl chloromethylphenyl with xylene (35).

**Infection of PBMC and microtiter infectivity assay.** For the viral replication assay, 5 × 10⁶ PCTAX-2 and M47-3 cells were lethally gamma irradiated (6,000 rads) and cocultured with 5 × 10⁶ PBMC activated with PHA and IL-2. The cells were cultured in RPMI supplemented with 50 U of IL-2/ml, and supernatants were collected at various time points and assayed for virus particles by p19 ELISA. The microtiter infectivity assays were performed as previously described by Persaud et al. (54). In brief, 10⁶ PHAHL-activated PBMC were cocultured with either 10⁵, 10³, or 10⁰ lethally irradiated (6,000 rads) PCTAX-2, PCTAX-3, M47-2, or M47-3 cells in replicates of 20 at each dilution in 96-well microtiter plates. The plates were cultured in RPMI with 50 U of IL-2/ml were split 1:2 approximately once per week. At 8 to 10 weeks postcoculture, individual wells were examined microscopically for the presence of viable cells.

**FACS analysis of immortalized cell lines.** One million ACH.pcTax- or ACH.M47-immortalized cells were washed in fluorescence-activated cell sorter (FACS) staining buffer (1× phosphate-buffered saline, 0.1% FCS, 0.05% sodium azide) and incubated in the same buffer for 30 min on ice with mouse anti-human CD3-phycocerythrin (PE), anti-CD25-PE, anti-CD4-fluorescein isothiocyanate, or anti CD8-PE (Pharmingen). The stained cells were washed twice in the staining buffer, fixed in 1% paraformaldehyde-phosphate-buffered saline, and analyzed on a FACSCalibur flow cytometer (Becton-Dickinson, San Jose, Calif.).

**RESULTS**

**Construction and activity of ACH Tax mutants.** A molecular clone of HTLV-1 known as ACH, which produces infectious viral particles capable of immortalizing primary human lymphocytes and establishing productive infections in rabbits, has been previously described (11, 34). The ACH Tax mutant clones were constructed as described in Materials and Methods and are diagrammed in Fig. 1. The resulting ACH Tax mutant clones contained the tax gene from HTLV-1 strain C91/PL between nucleotides 7346 and 8307, which differs from the CH strain in the ACH clone by 5 amino acid codons. As the first 17
and the last 7 amino acids are identical in the CH and C91/PL strains, the resulting tax gene in the ACH clones is identical to C91/PL tax. Since the C91/PL and CH strains differ by 5 amino acids, we also constructed a positive control clone which contained wild-type C91/PL tax (ACH.pcTax). The CREB/ATF- and NF-κB-activating activities of the M22 and M47 mutants have been previously examined in a number of cell lines and in a variety of expression systems (2, 69). However, we wanted to confirm that these mutants had the expected activities with respect to CREB and NF-κB activation when expressed from the ACH clone. The ACH.pcTax, ACH.M22, and ACH.M47 clones were cotransfected with either an HTLV LTR-luciferase reporter construct or human immunodeficiency virus (HIV) LTR-CAT reporter construct in human 293T kidney fibroblasts, and luciferase or CAT activity was measured (Table 1). As expected, the ACH.M47 clone expressed a Tax protein which failed to transactivate the HTLV LTR-luciferase construct, which is dependent on CREB, but had wild-type activity for the HIV LTR-CAT reporter plasmid, which is activated by NF-κB. Conversely, the ACH.M22 clone expressed a Tax protein which activated the HTLV LTR to near-wild-type levels but did not activate the HIV LTR. Therefore, when expressed from the ACH clone, the M22 and M47 Tax mutants have the originally described activity with respect to CREB and NF-κB activation.

**TABLE 1. Activity of Tax mutants and summary of immortalization assays**

<table>
<thead>
<tr>
<th>Clone</th>
<th>HTLV Activity (%)</th>
<th>HIV Activity (%)</th>
<th>Immortalized Cultures/No. of Transfections (No. of donor PBMC)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACH.wt</td>
<td>ND</td>
<td>ND</td>
<td>4/5 (5)</td>
</tr>
<tr>
<td>ACH.pcTax</td>
<td>100</td>
<td>100</td>
<td>5/6 (4)</td>
</tr>
<tr>
<td>ACH.M22</td>
<td>80</td>
<td>7</td>
<td>0/6 (4)</td>
</tr>
<tr>
<td>ACH.M47</td>
<td>2</td>
<td>138</td>
<td>5/5 (4)</td>
</tr>
<tr>
<td>Empty vector</td>
<td>0</td>
<td>0</td>
<td>0/8 (7)</td>
</tr>
</tbody>
</table>

Luciferase or CAT activity above empty vector cotransfection normalized to 100% for the wild-type ACH.pcTax clone. A 100% activity corresponds to an approximately 20-fold increase in luciferase activity for HTLV LTR-luciferase and a 2-fold increase in CAT activity for HIV LTR-CAT. ND, not done.

**Viral particle production.** To assess the abilities of the ACH.pcTax, ACH.M22, and ACH.M47 clones to produce viral particles, 293T cells were transfected with 10 μg of each clone and viral particle production was assayed by p19 antigen ELISA approximately every 24 h posttransfection (Fig. 2). The ACH.pcTax and ACH.M22 clones were both capable of producing relatively high levels of viral particles, with up to 15,000 pg of p19 antigen/ml being detected in the culture supernatants at 90 h posttransfection. Therefore, the M22 mutation in the ACH clone does not appear to affect viral particle production, as would be expected from its ability to activate the HTLV LTR-luciferase construct efficiently. This result rules out any major defect in the Rex protein, a splicing regulator required for efficient gag expression (26, 27), whose ORF overlaps that of Tax and in which the M22 mutation also changes two amino acids (Prp[149]Cys[150] to Leu[149]Ala[150]). However, the ACH.M47 clone produced much lower amounts of virus in this transient transfection assay, with approximately 400 pg of p19/ml being detected by 90 h posttransfection. This result was expected due to the defect in LTR activation by the M47 mutant.

**Immortalization of primary human lymphocytes.** To test the ability of the ACH Tax mutant clones to induce immortalization, 20 million Ficoll-Paque-purified human PBMC were activated with 10 μg of PHA/ml and 50 U of IL-2/ml for 72 h and transfected by electroporation with the ACH.wt, ACH.pcTax, ACH.M47, and ACH.M22 plasmids. Cellular viability was monitored by MTT conversion assays. An example of one experiment, in which two separate transfections were done with ACH.pcTax and ACH.M47 and three transfections were done with ACH.M22, is shown in Fig. 3. As expected, the cells transfected with the ACH.wt and ACH.pcTax clones continued to proliferate indefinitely while the mock-transfected cells proliferated transiently and died by approximately 60 days posttransfection. Interestingly, like the mock-transfected cells, the ACH.M22-transfected cells also ceased to proliferate. Surprisingly, the ACH.M47-transfected cells also became immortalized, despite the fact that virus produced by this clone is not predicted to replicate as efficiently as ACH.wt- or ACH.pcTax-derived virus. PBMC from multiple donors were likewise transfected and monitored, and the results are summarized in Table 1. These results indicate that while CREB/ATF activation activity appears to be dispensable for immortalization of primary PBMC, NF-κB activation by Tax may be important for cellular immortalization.

Three ACH.pcTax-immortalized cell lines and three ACH.M47-immortalized cell lines were chosen for further analysis and will be referred to as PCTAX-1, PCTAX-2, PCTAX-3, M47-1, M47-2, and M47-3. The PCTAX-1 and M47-1 cell lines were derived from PBMC from the same donor, whereas the PCTAX-2, PCTAX-3, M47-2, and M47-3 cell lines were derived from PBMC of a different donor. These cell lines were examined for cell surface marker expression by FACS analysis, and all were determined to be activated T cells. Two of the ACH.pcTax-immortalized cell lines were CD4+ CD8−, while the third was CD4− CD8+. In contrast, two of the ACH.M47 cell lines were CD4− CD8+, while one expressed both CD4 and CD8. To assess the pattern of proviral integra-
immortalizing activity. However, the M22 mutant fails to immortalize the immortalized cells. Southern blot analysis was performed on BamHI-digested genomic DNA with a probe derived from an 8.5-kb SacI fragment of the HTLV-1 genome. This analysis revealed both clonal and oligoclonal patterns of proviral integration in the immortalized cells and a lack of pBluescript vector sequences from the ACH clone flanking the 3’ LTR (data not shown). Thus, integration of the HTLV-1 provirus occurred through the viral replication cycle and not through random integration of the transfected plasmid.

Conservation of M47 mutation and lack of Tax activity in M47-immortalized cell lines. One potential explanation for the ability of the ACH.M47 mutant to immortalize primary PBMC despite being impaired for replication is that a reversion may have occurred in the tax gene which restored the ability of the M47 mutant Tax protein to activate the CREB/ATF pathway. To assess this possibility, the tax ORF was PCR amplified from proviral genomic DNA isolated from the three ACH.pcTax and three ACH.M47-immortalized cultures. The PCR product was then digested with BglII, which is diagnostic for the presence of the M47 mutation. Whereas the three PCR products from the ACH.pcTax-immortalized cells were not cleaved with BglII, all three of the PCR products derived from the ACH.M47-immortalized cells were cut (Fig. 5A). This analysis rules out the possibility of a change at any of the 6 nucleotides which comprise the M47 mutation, as any change would have resulted in the loss of the BglII site. Additionally, the PCR products were cloned and sequenced and confirmed for the presence of the M47 mutation (data not shown).

This analysis, however, does not rule out the possibility of the presence of a mutation elsewhere in the Tax protein which is somehow suppressing or compensating for the 2-amino-acid M47 mutation. To examine this possibility, the AccI-XmaI fragment containing the coding region for Tax amino acids 18 to 336 from the above-described PCR products were cloned into a CMV promoter-driven Tax expression clone (pIEX; provided by O. J. Semmes) and cotransfected with the HTLV LTR-luciferase reporter construct in 293T cells (Fig. 5B). Positive and negative control plasmids were likewise constructed by PCR amplifying and cloning the tax ORFs from the original ACH.pcTax and ACH.M47 clones. As expected, the control expression plasmids derived from the original ACH.pcTax and ACH.M47 clones had the expected activity with respect to HTLV LTR activation, with the pTax clone activating the LTR approximately 30-fold and the M47 clone having <10% of the activity of the wild-type expression clone. Furthermore, the expression clones derived from the PCR-amplified proviral tax genes from the PCTAX-1, -2, and -3 and the M47-1, -2, and -3 cell lines also had the expected activity for activation of the HTLV LTR. Therefore, a mutation had not occurred elsewhere in the tax gene which had restored the ability of Tax to activate the CREB/ATF pathway.

Although the above analysis rules out the possibility of a reversion in Tax, it does not address the possibility of a cellular mutation or alteration in cellular environment which results in an increase of CREB/ATF activity in the ACH.M47-immortalized cells. This could potentially be due to an increased level of CREB expression or CREB phosphorylation or an enhanced CREB-CBP-p300 interaction. Alternatively, the cells immortalized with the ACH.M47 mutant may be the result of an inadvertent selection for cellular clones which express abnormally high levels of Tax, thus compensating for the lack of CREB activation by the M47 mutant. To assess this possibility, the three ACH.pcTax- and ACH.M47-immortalized cell lines were directly transfected by electroporation with the HTLV LTR-luciferase reporter construct. The cells were additionally cotransfected with a CMV-CAT reporter to normalize for transfection efficiency. The cell lines immortalized with the ACH.pcTax clone all had higher HTLV-1 LTR activity than the ACH.M47-immortalized cell lines, which expressed levels of luciferase similar to those of PHA-IL-2-stimulated PBMC (Fig. 6A). Furthermore, Western immunoblot analysis indicated that while Tax expression varied among the cell lines, expression of Tax in the M47 cell lines was not consistently upregulated and was similar to the levels in the PCTAX cell lines (Fig. 6B). Finally, the PCTAX and M47 cell lines were examined for activated NF-kB activity by electrophoretic mobility shift assay analysis with a probe containing the NF-kB binding site from the IL-2Rα promoter. As expected, all PCTAX- and M47-immortalized cell lines had increased nuclear NF-kB compared to activated PBMC, indicating the expression of a functional Tax protein (data not shown). Therefore, these results indicate that the ACH.M47-immortalized cells do not have abnormally altered CREB/ATF activity or Tax expression which suppresses or otherwise compensates for the M47 mutation.

Viral replication and microtiter infectivity assay. Despite being impaired for replication in the transient transfection assay (Fig. 2) and producing lower amounts of virus early in the transfection of PBMC (data not shown), the ACH.M47-immortalized PBMC produce virus at levels similar to ACH.pcTax-immortalized cells. We examined whether or not the virus being shed from the ACH.M47-immortalized cells was as infectious for activated PBMC as wild-type virus. One ACH.pcTax-immortalized cell line (PCTAX-2) and one ACH.M47-immortalized cell line (M47-3) were chosen that expressed very similar amounts of viral particles (approximately 500 pg/ml/10⁶ cells/24 h). Furthermore, these two cell lines were both derived from the same donor, and both expressed CD4 on their surfaces (although M47-3 also expressed CD8). Therefore, PCTAX-2 and M47-3 are similar in many respects, except for the presence of wild-type or M47 mutant Tax in the inte-
grated provirus. Five hundred thousand cells were lethally irradiated (6,000 rads) and cocultured with $5 \times 10^6$ PHA-IL-2-activated PBMC, and viral replication was monitored by p19 ELISA on cell culture supernatants. As expected, the virus from the ACH.M47-immortalized cells replicated less efficiently than the wild-type virus due to the inability of M47 Tax to efficiently transactivate the LTR (Fig. 7A).

To further examine the replication and immortalizing activity of the ACH.M47 virus, a quantitative microtiter infectivity-immortalization assay was employed as described by Persaud et al. (54). In this assay, $10^4$ PBMC were cocultured with 10-fold dilutions ($10^3$, $10^2$, 10) of irradiated PCTAX-2, PCTAX-3, M47-2, or M47-3 cells in replicates of 20 in 96-well microtiter plates. The cultures were examined at 8 to 10 weeks postcoculture, and the percentage of cultures which became immortalized at each dilution of infected cells was quantitated (Fig. 7B). Despite being less able to replicate efficiently, the virus from the M47-2 and M47-3 cell lines infected and im-

**FIG. 4.** Cell surface phenotype of immortalized cell lines. ACH.pcTax- and ACH.M47-immortalized cells were stained with anti-human CD3, CD25, CD4, and CD8 and analyzed by flow cytometry. Although all cell lines express CD3 and CD25, expression of CD4 and CD8 is variable, with a tendency for CD4 expression by ACH.pcTax-transfected cells and CD8 expression by cells immortalized with the ACH.M47 clone. The open histogram on each plot corresponds to unstained cells, while the shaded histogram corresponds to cells stained with the indicated antibody.
mortalized the uninfected PBMC at each dilution of infected cells at a rate similar to that of the PCTAX-2- and PCTAX-3-derived virus. Thus, there does not seem to be a quantitative difference in the immortalizing activity of the M47 mutant Tax.

Cell surface phenotype of immortalized cell lines. Analysis of the cells immortalized by transfection of PBMC by the ACH.pcTax and ACH.M47 proviral clones revealed that while all cell lines were activated T cells, expression of CD4 and CD8 varied, with a slight bias for CD4 expression by ACH.pcTax-immortalized cells and for CD8 expression by ACH.M47-immortalized cells. To examine this phenomenon further, 12 ACH.pcTax and 12 ACH.M47 cell lines produced during the course of the microtiter infectivity assay were examined for cell surface marker expression (Table 2). As was observed for the ACH-transfected PBMC, the immortalized cell lines derived from infection with the M47-2- or M47-3-derived virus were more likely to be either CD8$^+$ T cells or mixed cultures of CD4$^+$ and CD8$^+$ cells. Likewise, the cultures infected with the PCTAX-2- or PCTAX-3-derived virus were more likely to be CD4$^+$ T lymphocytes. This suggests that activation of the CREB/ATF pathway by Tax may be important for the preferential immortalization of CD4$^+$ cells compared to that of CD8$^+$ cells by HTLV-1, despite the fact that the virus can infect both types of cells.

DISCUSSION

The Tax protein of HTLV-1 has been demonstrated to have a number of activities which may all lead directly or indirectly...
PBMC were cocultured with 10^3, 10^2, or 10 gamma-irradiated PCTAX-2, microtiter infectivity-immortalization assay. Ten thousand PHA-IL-2-activated deviation of two replicate infections is indicated by the error bars. (B) Results of PBMC. Viral replication was determined by p19 antigen ELISA. The standard irradiated (6,000 rads) and cocultured with 5 x 10^3 viable cells, and the number of cultures which were immortalized at each dilution was determined.

The ability of Tax to upregulate the expression of a variety of genes involved in cellular proliferation or cell cycle control has also been proposed as contributing to the ability of Tax to transform cells. Among the genes demonstrated to be upregulated by Tax are c-fos and the genes for PCNA, platelet-derived growth factor (c-sis), IL-2, and IL-2Rα (3, 14, 25, 40, 58, 68, 75). Additionally, Tax has been shown to repress the transcription of p53 (76) and bax (9), which may lead to an inhibition of apoptosis in the infected cell.

Tax has been demonstrated to have transforming activity in a number of cell culture systems (2, 17, 43, 62, 63, 70, 78). However, the relative importance of the activation of the CREB/ATF and NF-κB pathways for transforming activity is controversial. Studies examining the transformation of established rodent cell lines have produced conflicting results. Smith and Greene examined the ability of the M22 and M47 mutations to transform Rat2 fibroblasts and determined that the CREB pathway is important for transformation (i.e., M47 was defective for cellular transformation, while M22 retained transforming activity) (70). However, subsequently Yamaoka et al. and Matsumoto et al. have demonstrated that the NF-κB pathway appears to be important for the transformation of Rat1 fibroblasts (43, 78). Studies utilizing viral transduction systems to evaluate the ability of Tax mutants to immortalize primary human PBMC have also produced varied results. Rosin et al. demonstrated that a Tax point mutation defective for NF-κB induction (S258A) retains the capability to immortalize primary lymphocytes when transduced by a herpesvirus saimiri vector (62). On the contrary, Akagi et al. demonstrated that the M22 Tax mutation fails to immortalize PBMC when transduced by a retroviral expression vector (2).

To cellular immortalization and transformation of the infected cell, Tax interacts with the mitotic checkpoint protein MAD1, and this interaction leads to the loss of MAD1 function (30). Tax also interacts with Cdk4 and Cdk6 and increases their activities in primary human T cells (65). Tax has also been shown to interact with and inactivate the Cdk inhibitor p16INK4a (41, 72). Additionally, Tax functionally inactivates p53 through an increase in phosphorylation, although the mechanistic details are still unclear (46, 55, 56). Finally, expression of Tax leads to DNA damage, which may be associated with the transcriptional repression of DNA polymerase-β (29) and/or increased PCNA expression (58).
primary PBMC, which may be the closest model of what occurs in vivo.

Our results indicate that the activation of the NF-κB pathway by Tax is critical for the immortalization of PBMC by HTLV-1. Disregulated NF-κB activity has been demonstrated to be transforming and associated with other human cancers. The IκB family member bel-3 is located at a site of chromosomal translocations in a specific type of B-cell leukemia (51). Likewise, the NF-κB family member p52 was identified in the cloning of a chromosomal translocation in a non-Hodgkin’s B-cell lymphoma (48). Additionally, the Epstein-Barr virus LMP-1 protein activates the NF-κB pathway by interaction with the tumor necrosis factor receptor (TNFR)-associated factors (TRAFs) and the TNFR-associated death domain protein (TRADD) through a pathway that involves NIK, IKKα, and IKKβ (18, 73). Furthermore, activation of NF-κB by LMP-1 through its interactions with the TRAFs and TRADD has been demonstrated to be important for its B-lymphocyte growth-transforming activity (12, 28).

As mentioned previously, the M22 Tax mutant has been demonstrated to lack NF-κB-activating activity in a variety of cell types, including Jurkat T cells (69) and primary human PBMC (2). Furthermore, it appears that M22 Tax fails to activate NF-κB by a defect in the activation of the IKK-activating kinases MEKK1 and NIK (15, 79). Although the effect of the M22 mutation on NF-κB activation is clear, other effects that the M22 mutation may have on other Tax functions is less well established. For example, Pise-Masison et al. have demonstrated that M22 mutant Tax is not able to functionally inactivate p53 as wild-type Tax is capable of doing (55). However, Mulloy et al. have shown that M22 retains the ability to inactivate p53 function (46). In addition, the M22 mutant Tax protein does not dimerize as efficiently as wild-type Tax (74), which may also influence various Tax activities, including HTLV LTR activation (1, 81). Therefore, we cannot rigorously exclude the possibility that the lack of immortalizing activity observed with the ACH.M22 clone may be due to a defect in Tax to interact with or activate or inactivate some additional factor. Examination of additional Tax mutants defective for NF-κB induction may help to conclusively define a role for NF-κB induction in HTLV-1-mediated cellular immortalization.

Our results also indicate that activation of the CREB/ATF pathway by Tax is not necessary for immortalization. This is somewhat surprising, as the ACH.M47-derived virus replicates to much lower levels than the wild-type ACH.p cxTax-derived virus, due to the defect in CREB activation. However, the ACH.M47-immortalized cells produce viral particles to levels similar to those of ACH.p cxTax-immortalized cells (data not shown). There are a number of possible explanations for this observation. The possibility exists that once the cells become immortalized, there are sufficient levels of CREB or other transcription factors to drive levels of viral gene expression that lead to relatively large amounts of viral particle production. Second, regulation of viral gene expression in the immortalized cell may be more complex than simple activation of the LTR by Tax, as a number of studies have identified suppressors of the LTR, possibly in the R and U5 regions (52, 77). In fact, expression of viral genes by the ACH clone appears to be restricted in lymphoid cell lines but not in primary PBMC (34). In addition, the fact that CREB activation is not necessary for immortalization may not be surprising, in that few cellular genes have been shown to be activated by Tax through the CREB/ATF pathway. However, though activation of the CREB/ATF pathway is dispensable for cellular immortalization (IL-2-dependent growth), our results do not rule out the possibility that Tax interaction with the CREB/ATF pathway is important for the emergence of fully transformed cells which proliferate independently of IL-2.

In vivo, CD4+ T cells represent the major infected cell type in asymptomatic individuals (59, 60), and the leukemic cells in patients with adult T-cell leukemia/lymphoma are CD4+ T cells in the majority of cases (21, 31, 33). In vitro, HTLV-1 can infect and immortalize CD4+ as well as CD8+ T cells, although immortalization and transformation of CD4+ cells is more common (42, 54). While the M47 mutant virus retains the ability to immortalize infected T cells, there appears to be a difference in the phenotype of the immortalized cells, as CD8+ cells become immortalized at a much higher frequency than that observed with the wild-type virus. Newbound et al. have demonstrated that the ability of Tax to activate the HTLV-1 LTR is greatly enhanced in CD4+ cells compared to CD8+ cells (50). Furthermore, it was proposed that this difference may account for the higher frequency of CD4+ HTLV-1 immortalized cells than CD8+ immortalized cells (50). This hypothesis is consistent with our findings. Since the virus with the M47 mutation does not efficiently transactivate the LTR, there would be no selective advantage for replication in CD4+ cells versus CD8+ cells. Therefore, one would expect an approximately equal frequency of CD4+ and CD8+ immortalized cells with the M47 mutant virus. Alternatively, activation of the CREB/ATF pathway may be important for increased proliferation and outgrowth of CD4+ cells, as was proposed by Akagi et al., who observed immortalization of CD4+ T cells by retrovirally transduced wild-type Tax and CD8+ cells by transduced M47 mutant Tax (2).

The results presented here support the possibility that NF-κB activation by Tax is important for the immortalization of primary T lymphocytes by HTLV-1. In addition, it appears that activation of the CREB/ATF pathway by Tax is dispensable for cellular immortalization. Examination of the immortalizing phenotypes of various Tax point mutants in a functional molecular clone of HTLV-1 likely mimics the infection of primary lymphocytes in vivo more closely than other cell culture models which have been used to study cellular transformation by HTLV-1. This system will be useful for the determination of immortalizing activity of additional Tax mutants defective for other activities of Tax which may contribute to the immortalization and transformation of infected cells, which will lead to a more complete understanding of Tax function and the pathogenesis of HTLV-1 infection.

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