Long-term AZT Exposure Alters the Metabolic Capacity of Cultured Human Lymphoblastoid Cells

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The antiretroviral efficacy of 3′-azido-3′-deoxythymidine (AZT) is dependent upon intracellular mono-, di-, and triphosphorylation and incorporation into DNA in place of thymidine. Thymidine kinase 1 (TK-1) catalyzes the first step of this pathway. MOLT-3, human lymphoblastoid cells, were exposed to AZT continuously for 14 passages (P1–P14) and cultured for an additional 14 passages (P15–P28) without AZT. Progressive and irreversible depletion of the enzymatically active form of the TK-1 24-kDa monomer with loss of active protein was demonstrated during P1–P5 of AZT exposure. From P15 to P28, both the 24- and the 48-kDa forms of TK-1 were undetectable and a tetrameric 96-kDa form was present. AZT-DNA incorporation was observed during P1–P5 of AZT exposure. When MOLT-3 cells were grown in AZT-free media (P15–P29), there was a reduction in AZT-DNA incorporation and MN formation; however, TK-1 depletion and the persistence of S-phase delay were unchanged. These data suggest that in addition to known mutagenic mechanisms, cells may become resistant to AZT partially through inactivation of TK-1 and through modulation of cell cycle components.

Key Words: nucleoside analog; antiretrovirals; thymidine kinase.

The genotoxicity of the DNA chain–terminating nucleoside reverse transcriptase inhibitors (NRTIs) has been amply demonstrated (International Agency for Research on Cancer, 2000), but specific underlying mechanisms other than DNA incorporation and mutagenesis are unclear. Since clinical NRTI therapy is administered long term and because human bone marrow is a target for NRTI toxicity (Richman et al., 1987; Yarchoan et al., 1989), human lymphoblastoid (MOLT-3) cells were used here to study the genotoxic mechanisms of long-term NRTI exposure. It is well established that 3′-azido-3′-deoxythymidine (AZT) acts as an HIV-1 reverse transcriptase inhibitor (Mitsuoka et al., 1985) by DNA incorporation that is required for chain termination and by blocking the nucleotide-binding site of the HIV-1 reverse transcriptase (Furman et al., 1986).

Thymidine kinase 1 (TK-1) catalyzes the monophosphorylation of AZT in the pathway of activation to DNA incorporation and is considered to be the rate-limiting step in NRTI metabolism. In cultured human cells, AZT resistance has been reported to be correlated with decreases in TK-1 activity (Avramis et al., 1993; Han et al., 2004) and hypermethylylation of the TK1 gene (Nyce et al., 1993; Wu et al., 1995). TK-1 exists intracellularly as an active monomeric form (24 kDa), an inactive dimeric form (48 kDa), an inactive tetrameric (96 kDa) form (Sherley and Kelly, 1988). The triphosphorylated AZT becomes incorporated into virus and host DNA in place of thymidine, inducing chain termination that results in DNA fragmentation and the formation of micronuclei (MN) (Olivero, 2007). Induction of MN occurring as a result of exposure to AZT has been reported in many different studies using various cell cultures and animal species. For example, increased levels of MN were reported in CD4 lymphocytes (Stern et al., 1994) and human lymphocytes and Chinese hamster ovary cells (Gonzalez and Larripa, 1994), HeLa cells (Arana and Jagetia, 2001), and human H9 cells (Agarwal and Olivero, 1997) all as a result of exposure to AZT. In addition, an increase in frequency of erythrocyte MN was observed in mouse pups exposed to AZT plus the NRTI didanosine (ddI) during gestation and after birth (Bishop et al., 2004; Dertinger et al., 1996; Witte et al., 2004) and in TK−/− and TK−/− mice (Dobrovolsky et al., 2005). The formation of AZT-DNA incorporation and AZT-induced MN suggests that chronic AZT exposure may induce mutations and functional
alterations in DNA replication. Increase in the frequency of complete Tk gene deletions following exposures to AZT and ddi have been reported (Meng et al., 2002) in vitro and in vivo (Von Tungeln et al., 2004). Furthermore, mutational analysis suggests that AZT induces a unique pattern of mutations in the Tk gene of mice and that the major mechanisms of AZT-induced mutagenesis involve deletion and recombination (Mittelstaedt et al., 2004).

Here, we pursued mechanistic studies exploring consequences of prolonged AZT exposures using high concentrations of AZT that have been shown to induce genotoxicity and minimum cell death and compared those to therapeutic doses. Additionally, we have explored the persistence of AZT-induced effects once the exposure was terminated. To this end, we cultured MOLT-3 cells continuously in the presence of AZT for 14 passages (P14), followed by an additional 15 passages in the absence of the drug. We explored cell cycle parameters by flow cytometry and DNA damage by a radioimmunoassay (RIA) able to detect AZT incorporated into DNA and by documenting MN formation. Additionally, we investigated the role of chronic AZT exposure as a modulator of TK-1 oligomerization and functional capacity.

MATERIALS AND METHODS

Cell culture and treatment. The human lymphoblastoid cell line MOLT-3 (ATCC, Manassas, VA) was cultured using Rosman-Park-Memorial-Institute (RPMI) 1640 medium supplemented with 10% fetal bovine serum (ATCC). AZT (Sigma-Aldrich Co., St Louis, MO) exposures were performed in 75-cm² plastic culture flasks. AZT was dissolved in PBS (pH 7.2) (Biosource, Rockville, MD), and the final concentration was calculated from absorbance at 266 nm with a molar extinction coefficient of 11,500. Triplicate experiments were performed to assess the incorporation of 0, 5, 10, 50, 100, 200, and 800µM AZT into MOLT-3 cell DNA, for up to 14 passages. In a separate study, five different batches of MOLT-3 cells were exposed continuously to 800µM AZT for up to P14, and for three of those five experiments, AZT was removed at P14 and cells were grown for an additional 15 passages (to P29) in the absence of drug. Finally, in a repeat experiment, MOLT-3 cells were exposed to 10µM AZT, a human plasma level concentration, for up to 14 passages. MOLT-3 cells, on three separate occasions to 0 and 800µM AZT, were collected at P1, P4, P5, and P14. Cells from all the treatments and unexposed cells were centrifuged and pellets were fixed with 70% alcohol to obtain suspensions of 8 x 10⁵ cells/ml media. Ten slides/passage/experiment were obtained by centrifugation of 50 µl of the cell suspension/slide (~4000 cells) using a cytospin (Shandon Cytospin 2; Thermo Fisher Scientific Inc., Waltham, MA). Cells were stained with a 5% solution of Giemsa stain in PBS (pH 7.4) for 5 min. MN were scored.

Cytotoxicity analysis and time to subculture. MOLT-3 cells, from five independent trials, were collected 24 h after exposure to either 0 or 800µM AZT. For each experiment, cells were counted twice in a Coulter Counter (Coulter Electronics Limited, UK), and cell counts from drug-exposed flasks were compared with those from the unexposed flasks. Survival in exposed cells was expressed as percentage of survival in unexposed cells. In addition to counting, survival was established by trypan blue exclusion, with numbers of viable cells compared in exposed and unexposed cells. A separate observation was made in 15 independent trials to establish time to subculture. The ability of the cells to proliferate was also evaluated based on color changes in the culture medium indicated by phenol red. Typically, a yellowish color is observed when a critical amount of cells is present, point at which the cells were subcultured.

Preparation of DNA and measurement of AZT incorporated into DNA by RIA. DNA from unexposed cells and AZT-exposed cells at P1, P4, and P14 was obtained by nonorganic isolation (Chemicon International, Temecula, CA). The incorporation of AZT into MOLT-3-DNA was determined in cells exposed to 0, 5, 10, 50, 100, 200, and 800µM AZT by an AZT-RIA (Olivero et al., 1994). Briefly, a rabbit polyclonal anti-AZT antibody (Sigma-Aldrich Co.), which also recognizes AZT in DNA (Olivero et al., 1994), was reconstituted, diluted 1:7500, and incubated with MOLT-3-DNA for 90 min at 37°C. An aliquot (100 µl) containing ~20,000 cpm of [³²P]AZT tracer (16 Ci/mmol; Moravek Biochemicals Inc., Mountain View, CA) was added to each tube together with 100 µl of the secondary antibody, goat anti-rabbit immunoglobulin G (IgG; Sigma-Aldrich Co.), and reconstituted in 12 ml of 2-amino-2-hydroxyethyl-propane-1,3-diol (Tris) buffer (pH 8.00). The mixture was incubated for 25 min at 4°C, centrifuged at 1942 x g for 15 min at 4°C, and the resulting supernatant was decanted. The pellets were dissolved in 200 µl 0.1M NaOH, transferred to scintillation vials containing 6 ml Ecolite (MP, Irvine, CA), and counted in a liquid scintillation counter. The amount of standard AZT, added to 3 µg of unexposed carrier MOLT-3-DNA, required to inhibit antibody binding by 50% was 2.86 ± 0.9 (average ± SD, n = 12) pmol AZT/tube. The lower limit of detection was 4.2 molecules of AZT/10⁶ nucleotides.

MN assay. In separate experiments, MOLT-3 cells were exposed to 0, 10, and 800µM AZT and collected at P1, P4, P5, and P14. MOLT-3 cells, exposed on three separate occasions to 0 and 800µM AZT, were collected at P1, P4, P5, and P14. Cells from all the treatments and unexposed cells were centrifuged and pellets were fixed with 70% alcohol to obtain suspensions of 8 x 10⁵ cells/ml media. Ten slides/passage/experiment were obtained by centrifugation of 50 µl of the cell suspension/slide (~4000 cells) using a cytospin (Shandon Cytospin 2; Thermo Fisher Scientific Inc., Waltham, MA). Cells were stained with a 5% solution of Giemsa stain in PBS (pH 7.4) for 5 min. MN were scored in 1000 MOLT-3 cells from 10 slides/passage.

Flow cytometric analysis of cell cycle parameters. MOLT-3 cells were exposed on three separate occasions to 800µM AZT during P1–P14 and then grown in AZT-free media for additional 10 passages. Cells were collected at every passage, pelleted, and washed with RPMI 1640 without serum before they were fixed in 1 ml 70% ice-cold ethanol, gently dropped while vortexing. Following an overnight fixation at 4°C, cells were pelleted by centrifugation and incubated with Ribonuclease A (Sigma-Aldrich Co.) at room temperature for 20 min. Propidium iodide (20–50 µg/ml) (Molecular Probes, Eugene, OR) was added to each cell suspension, and cells were kept in the dark at 4°C overnight. Cells were passed through a fluorescence-activated cell sorter (FACSCalibur; BD Biosciences, San Jose, CA) using the double discrimination module, and data were acquired using CellQuest (BD Biosciences) software. The cell cycle was modeled using ModFit software (Venty Software, Topsham, ME). Percentages of cells in G0–G1, S, and G2–M phases were calculated directly by the software.

Western blot for TK-1 protein analysis. Aliquots of unexposed and AZT-exposed MOLT-3 cells from each passage were lysed in radio-immune precipitation assay buffer (50mM Tris-HCl [pH 7.6], 150mM NaCl, 0.25% SDS, 1% Triton X-100, 1mM EDTA, and 0.5% Nonidet P-40 [NP-40]; Fluka Chemicals, Milwaukie, WI) with the addition of protease inhibitor tablets (Complete; Roche Diagnostics, Indianapolis, IN) for 30 min on ice, followed by sonication with an ultrasonic processor at a 20% amplitude with a 3-mm microtip (Sonicos VC 750; Sonicos and Materials Inc., Newtown, CT) for three pulses of 10 s each. Proteins were quantified by Bradford reaction (Bio-Rad, Hercules, CA). Samples were resolved on a 10% Bis-Tris vertical polyacrylamide gel (NuPage; Invitrogen) and then transferred to a polyvinylidene fluoride membrane (Immobilon-P; Millipore, Billerica, MA). Membranes were allowed to dry overnight and then were blocked with phosphate buffer saline 0.1% tween containing 5% nonfat dry milk. A TK-1 monoclonal antibody (QED Biosciences, San Diego, CA) was used to incubate the membrane overnight at 4°C. After incubation with an anti-mouse IgG–horseradish peroxidase (HRP)-conjugated secondary antibody (Novus Biologicals, Littleton, CA), the membrane was processed for chemiluminescence with an Enhanced Chemiluminescence...
Western Blotting Detection Kit (Amersham Biosciences, Buckinghamshire, UK). Controls for loading were carried out after stripping the membrane (Restore; Invitrogen) using a mouse anti-actin antibody (Chemicon International) followed by an anti-mouse IgG-HRP–conjugated secondary antibody (Novus Biologicals). Signal was revealed using electrochemiluminescence Western Blotting Detection Reagents as previously described (Olivero et al., 2008). Images were captured using a Lumi-imager (Roche Diagnostics).

**Radiochemical enzyme–specific activity assay for TK-1.** Pellets from AZT-exposed cell suspensions (3 × 10^7 cells) collected at different passages in three separate experiments were resuspended in 1 ml of ice-cold extraction buffer (50mM Tris-HCl [pH 7.6], 2mM dithiothreitol [DTT], 1 protease inhibitor cocktail tablet [Roche Diagnostics], 20% glycerol, and 0.5% NP-40). Cell suspensions were frozen and thawed three times to improve the lysis process, and sonicated cell lysates were centrifuged for 10 min at 18,000 × g at 4°C. The supernatant was kept on ice and analyzed for protein concentration as described above using the Bradford reaction. Diethylaminoethyl cellulose filter discs (Whatman, Maidston, UK) were immersed in 10μM AZT-monophosphate (MP) for 10 min and allowed to dry completely. The reaction mixture (50mM Tris-HCl [pH 7.6], 2mM DTT, 5mM MgCl2, 5mM ATP, 10mM NaF, and 20μM [H]3-AZT; 12 Ci/mmol; Moravek Radiochemicals, Brea, CA) was used to incubate the protein extract at 37°C for 20 min. After that time, 20 μl of the suspension was spotted on the pretreated DEAE discs. The discs were allowed to air dry and washed three times with ammonium formate, transferred to liquid scintillation vials, and radioactivity was counted. TK-1 activity (counts per minute) was measured as directly proportional to the radioactivity obtained (Arner et al., 1992).

**Phosphorylation of [3H]-thymidine during cell growth.** Control cells and cells treated with 800μM AZT for 1, 45, and 59 passages were incubated for 30 min with [3H]-thymidine. Cells were harvested and lysed in 5% trichloroacetic acid. After 10 min on ice, the extracts were centrifuged at a 3-min centrifugation at 2000 × g at 4°C. A measured volume of the resultant supematant was neutralized by addition of AG-11A8 resin (0.5 g/ml extract), filtered, and the labeled deoxynucleoside/deoxynucleotide pools were analyzed by high pressure liquid chromatography as described (McKee et al., 2004).

**RESULTS**

**Cytotoxicity and Time to Subculture for MOLT-3 Cells Exposed Chronically to AZT**

MOLT-3 cells that were collected 24 h after exposure to 0, 10, or 800μM AZT exhibited a survival of 100, 81.9, and 69.2%, respectively. In separate experiments, time to subculture was assessed in MOLT-3 cells during the course of 14 passages of continuous exposure to 0, 10, or 800μM AZT (Fig. 1, open circles). In unexposed cells, passaging took place daily, and in AZT-exposed cells at P2–P5, a delay in the cell cycle was observed. However, Figure 1 shows that by five passages of AZT exposure, the rate of doubling was altered so that confluence was observed by 2-day intervals for P5–P14. Similar profile was observed for long-term MOLT-3 cells exposed to 10μM AZT.

During 14 passages of chronic exposure to 800μM AZT, the activity of TK-1, measured by the capacity to phosphorylate thymidine, revealed an abrupt decrease as early as P1, which lasted until P14 (Fig. 1). Insets in Figure 1 show Western blots of monomeric 24-kDa TK-1 from extracts obtained at the same passages for which time to confluence and activity were measured. The insets show unexposed cells and cells exposed to AZT for the passages indicated and were obtained from the

![FIG. 1](https://academic.oup.com/toxsci/article-abstract/115/1/109/1634250) Comparison of time to confluence (circles) and TK-1–specific activity (% of control [triangles]) for MOLT-3 cells exposed continuously to 800μM AZT. Time to confluence was determined visually, and TK-1 phosphorylation was obtained by radiolabeling assay (see “Materials and Methods” section). Insets show expression of monomeric 24-kDa TK-1 by Western blots, where the “ctrl” cells were unexposed and the exposed passages are indicated. For loading control, see Figure 5.
Western blot presented in Figure 5a. The reduction of TK-1 activity with continued culture time in the presence of AZT is clearly documented here.

Incorporation of AZT into MOLT-3 Cell DNA

An RIA was used to determine the amount of AZT incorporated into DNA at different passages. Figure 2 shows AZT-DNA incorporation values for MOLT-3 cells exposed to 5, 10, 50, 100, 200, and 800µM AZT for 1, 5, and 14 passages. Incorporation of AZT was observed at all passages with all doses, with maximum incorporation, 246 molecules AZT/10^6 nucleotides, at P5 with the 200µM dose. For the 800µM AZT dose, the highest incorporation was also observed at P5 with 171 molecules AZT/10^6 nucleotides. Figure 2 shows no obvious dose response either with dose or with increasing passage number. In an additional experiment, MOLT-3 cells originally exposed to 800µM AZT for 14 passages were subsequently cultured in AZT-free media, for an additional 12 passages. No detectable AZT-DNA incorporation remained after these cells were cultured in the absence of AZT for 12 passages (data not shown).

Statistical Interpretation

Application of linear regression to the mean levels of AZT-DNA incorporation at P1 showed a positive trend with an R value of 0.737. However, the data did not pass the normality test (p = 0.03) due to the mean value at 10µM AZT. If the latter value for 10µM AZT is excluded, then the data for the remaining AZT concentrations pass a Normality Test, and a highly positive linear regression (R = 0.985) is observed at P1. One-way ANOVA was used in addition to the Student’s t-test to compare the data among groups on the MN experiments (Table 2), and the p value was of 0.0001.

DNA Damage Assessed by the MN Assay

The MN assay was performed as one indicator of AZT-induced DNA damage. The induction of MN in MOLT-3 cells exposed to 10µM AZT (Fig. 3) and 800µM AZT (Fig. 3) is shown for passages 0, 1, 4, 5, and 14. In Figure 3 statistically significant differences were observed for all the AZT-exposed passages compared to the unexposed controls (p values = 0.016, 0.0003, 0.0078, and 0.0022 for P1, P4, P5, and P14, respectively). Additionally, a comparison among the unexposed controls cultured in parallel for P1, P4, P5, and P14 showed no statistical significance (p = 0.084).

In cells exposed to 800µM AZT at P1, P4, P5, and P14, the MN (per 1000 cells examined) reached values of 20, 61, 38, and 35, respectively, a significant increase for passage 1 (p = 0.02) compared to the untreated control (P0), and highly significant increase for P4 (p = 0.0001), P5 (p = 0.0002), and P14 (p = 0.0001); p < 0.05 for each passage compared to P0. However at P27, after 13 passages of culturing in the absence of drug, MN values returned to control levels (8/1000 cells) (Table 2).

Cell Cycle Analysis of MOLT-3 Cells Exposed Chronically to 800µM AZT

Evaluation of changes in the MOLT-3 cell cycle profile was carried out by flow cytometry in cells exposed long term to AZT. Table 1 shows percentages of cells in G0-G1, S, and G2 phases determined in untreated (control) cells and in cells exposed to 800µM AZT at P1 and P14. Table 1 also shows values for cells at P24, which were exposed to AZT for 15 consecutive passages with no AZT, dark bar). Values are mean ± SD from three experiments. Comparison of MN induced by 10µM AZT frequency by Student’s t-test showed a significant increase for passage 1 (p = 0.016) compared to the untreated control (P0), P4 (p = 0.0003), P5 (p = 0.0078), and P14 (p = 0.0022). Values are mean ± SD from three experiments. Comparison of MN frequency induced by 800µM AZT by Student’s t-test showed a significant increase for passage 1 (p = 0.02) compared to the untreated control (P0), and highly significant increase for P4 (p = 0.0001), P5 (p = 0.0002), and P14 (p = 0.0001).
The oligomerization of TK-1, from monomer to dimer and tetramer, prompted us to investigate TK-1 phosphorylation–specific activity, and the data are shown in Figure 1 (solid triangles). An 80% decrease in TK-1 activity was observed in MOLT-3 cells at P1 in the presence of 800 μM AZT. During 13 consecutive passages, in which cells were cultured in the presence of 800 μM AZT, TK-1 activity remained near the assay limit of detection (Fig. 1, Table 2). A plot of the relationship between the phosphorylation activity of TK-1 and time to confluence is shown in Figure 1, with insets indicating TK-1 protein levels determined by Western blot at P1, P3, P5, and P12. The Western blot insets also show that the drop in expression of TK-1 protein in its inactive dimeric (48 kDa) and active monomeric (24 kDa) forms, for unexposed (P0) cells and cells exposed to 800 μM AZT from P1 to P14 as well as cells grown subsequently in AZT-free media (P17–P19). The 24-kDa band indicates the presence of active protein in cultures that have been either unexposed (control) or exposed to AZT for 24 h. Reduction in protein band intensity is apparent during subsequent passages 9–14 with AZT in the medium and passages 17–19 with no AZT. The inactive dimeric form, however, remains unchanged through P14 (Fig. 4).

**TK-1 Enzyme–Specific Activity**

Cell lysates from consecutive passages of MOLT-3 cells exposed to 800 μM AZT were processed for Western blot analysis using an anti-TK-1 antibody. Figure 4 shows the reduction of TK-1 protein level with long-term culture of MOLT-3 cells in the presence of AZT. 

**Note.** *p Values < 0.05 (Student’s t-test). Values are mean ± SD for three experiments where AZT-exposed cells at P1, P14, and P24 were compared to the unexposed control.
The doubling time was accompanied by a loss of TK-1 monomeric protein (Fig. 1, P₁–P₅, and Fig. 5a).

**Phosphorylation of Thymidine during Cell Growth**

The data above suggested that in the presence of 800 μM AZT, thymidine tri-phosphate (TTP) for DNA replication was provided predominantly from the de novo rather than the salvage pathway. To further assess this point, control cells and cells continually exposed to 800 μM AZT for various passages were incubated with [³H]-thymidine for 30 min, and the conversion to [³H]-thymidine mono-phosphate, thymidine di-phosphate, and TTP was determined as described in “Materials and Methods” section. In the control cells, 60% of the label was phosphorylated, mostly to TTP (47%). After one passage in the presence of AZT, only a small amount of TTP (< 4% of total) was detected. At later passages, there was no detectable conversion of [³H]-thymidine to [³H]-TTP (data not shown).

Table 2 provides a summary of all the data presented in this paper. The doubling changes were similar to the percentage of cells in S-phase in that the cell cycle slowed immediately upon encountering the drug and subsequently (≥ P₅) shifted more toward normal. The TK-1-specific activity, in parallel with the TK-1 monomeric form, was completely depleted beyond P₅. However, the AZT-DNA incorporation and MN values were still high at P₁₄ in the absence of active TK-1 and were only decreased after AZT was removed from the media.

**DISCUSSION**

The antiretroviral nucleoside analog AZT becomes incorporated intracellularly into the nascent HIV-1 viral DNA via reverse transcriptase and into the host DNA via classical polymerases (Furman et al., 1986). Prior to this incorporation, the drug must be phosphorylated by TK-1, thymidylate kinase, and nucleoside diphosphate kinase (Furman et al., 1986). In this report, an inhibition of TK-1 protein expression and activity was found to correlate with length of long-term AZT exposure in human lymphoblastoid MOLT-3 cells (Table 2). Incorporation of AZT into DNA, and more general DNA damage revealed by MN, were observed during AZT exposure and decreased to background levels when AZT was removed from the medium. The lack of dose response seen with AZT incorporation into the DNA MOLT-3 cells could be explained by the fact that these cells are derived from a patient who had received prior multidrug chemotherapy and has unusually high terminal deoxynucleotidyl transferase activity.

Cells cultured for 14 passages with 800 μM AZT exhibited inhibition of TK-1 protein and phosphorylation activity along with cell cycle delay. Finally, when AZT was removed from the medium, active TK-1 did not recover and the cell cycle parameters did not return to normal, even though indicators of DNA damage, AZT incorporation, and MN returned to background levels. The dose used for most of the experiments presented here, 800 μM AZT, is the highest dose with low cytotoxicity observed in this cell line. This high dose was employed in order to reveal mechanisms of AZT-induced
genotoxicity. In addition, using 10μM AZT, a dose found in human plasma of HIV-1–infected patients, we also found significant DNA damage (Fig. 3).

In agreement with the previous findings that only the monomeric form of TK-1 performs active phosphorylation (Sherley and Kelly, 1988), we observed a TK-1 phosphorylating activity on the part of the 24-kDa form of TK-1 (Fig. 1). The activity and the monomer disappeared after five passages of exposure to 800μM AZT and were not restored in further passages. In addition, the expression of the inactive dimeric form (48 kDa) remained unchanged as time of exposure to AZT increased (Fig. 1). Moreover, after AZT removal, the monomeric form was undetectable, the dimeric form was dramatically reduced, and a tetrameric form of 96 kDa became evident (Fig. 4, right panel). A modest reduction was observed when cells were exposed long term to lower doses as observed in Figure 5b.

Using a recombinant TK-1 model, others support the concept of a catalytically effective, ATP inducible tetrameric form with higher activity than the dimeric form (Munch-Petersen, 2009). Inhibition of TK-1 activity by AZT exposure has been reported in T-lymphocytic cells (Avramis et al., 1993) and attributed to hypermethylation of the TK-1 promoter (Wu et al., 1995). Inhibition of the TK-1 enzyme has been reported to occur in HIV-1–infected patients (Antonelli et al., 1996; Jacobsson et al., 1995, 1998; Turriziani and Antonelli, 2004) as well. Antonelli et al. (1996) reported that 10 patients exposed to therapeutic doses of AZT for 6 months exhibited a significant reduction in the phosphorylation efficiency of the enzyme TK-1.

In H9 cells, the DNA methylation inhibitor 5-azacytidine reversed AZT-induced downregulation of TK-1, again suggesting that TK-1 hypermethylation may account for the decreased TK-1 gene expression (Groschel et al., 2002; Hoever et al., 2003). However, those reports do not discuss the form of the enzyme inhibited or the changes in oligomerization taking place after treatment.

Our data suggest that AZT induces theimerization of the 24-kDa form (Fig. 4). Western blot analysis of long-term MOLT-3 cells exposed to 800μM AZT revealed the presence of a 24-kDa form and a 48-kDa form at earlier passages similar to the forms found in untreated cells. However, when exposure reached five passages, the 24-kDa form became undetectable and the 48-kDa form remained present. Furthermore, the only form present after removal of the drug for three to five passages was the 96-kDa tetramer (Fig. 4).

Interestingly, although TK-1 is considered to be the rate-limiting step in AZT incorporation into DNA, we measured incorporation at P14, several passages after both the TK-1 activity and the monomer protein were undetectable. Three possible scenarios could explain this dichotomy. First, although undetectable by Western blots, low levels of TK-1 protein could have been sufficient to phosphorylate AZT that was been incorporated into DNA. Second, incorporation taken place at earlier passages, when an active form of TK-1 was present, could have remained in the DNA without being removed by DNA repair mechanisms and could have been present in later passages, perhaps in those cells with impaired or reduced cell division capacity. Finally, the nuclear encoded mitochondrial pyrimidine deoxynucleoside salvage enzyme TK-2, involved in pyrimidine nucleoside phosphorylation, could have been responsible for AZT phosphorylation in the absence of TK-1. Evidence of a phosphorylation activity for that enzyme has been reviewed by Eriksson et al. (2002), indicating that analogs, such as AZT, arabinofuranosylthymine, 3’-fluoro-2’,3’-deoxythymidine, and ribothymidine, could be phosphorylated but with relatively low efficiency. Evidence of this activity has been reported elsewhere utilizing transgenic mice that individually expressed pathogenetic point mutants of human TK-2 (Kohler et al., 2008). TK-2 is constitutively expressed along the cell cycle and often is virtually the only thymidine kinase that is physiologically active in nonproliferating and resting cells (Perez-Perez et al., 2008). To rule out nucleotide pool imbalances, experiments were performed employing equimolar concentrations of thymidine to study analysis of cell cycle and other end points. Results (Yu et al., 2009) indicated that the doses of NRTIs used had no effect on the nucleotide pool.

It does not appear at the level of transcription, that salvage pathways have been activated, since the expression of the genes involved in this pathway, determined by microarray at pathways have been activated, since the expression of the genes involved in this pathway, determined by microarray at

In summary, this study showed that indicators of DNA damage, AZT-DNA incorporation, MN, and S-phase arrest occur in MOLT-3 cells exposed to AZT continuously for 15 passages, and only the S-phase arrest is not completely reversible in cells subsequently grown in AZT-free media. The study also showed that continuous AZT exposure resulted in an early (P3) complete downregulation of the TK-1 24-kDa protein accompanied by a loss in the ability of TK-1 to phosphorylate AZT. The losses were not reversible at later passages after AZT was removed from the media. The data
showing AZT-induced DNA damage in the absence of active TK-1 were unexpected and requires explanation. The role of chronic AZT exposure as a modulator of TK-1 oligomerization and functional capacity could be relevant to AZT resistance in patients treated long term.

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**REFERENCES**


Munch-Petersen, B. (2009). Reversible tetramerization of human TK1 to the high catalytic efficient form is induced by pyrophosphate, in addition to tripolyphosphates, or high enzyme concentration. *FEBS J.* 276, 571–580.


