Abacavir, an anti–HIV-1 drug, targets TDP1-deficient adult T cell leukemia

Kohei Tada,1 Masayuki Kobayashi,1,* Yoko Takiuchi,1 Fumie Iwai,1 Takashi Sakamoto,1 Kayoko Nagata,1 Masanobu Shinohara,1 Katsuhiko Io,1 Kotaro Shirakawa,1 Masakatsu Hishizawa,1 Keisuke Shindo,1 Norimitsu Kadowaki,1 Kouji Hirota,2 Junpei Yamamoto,3 Shigenori Iwai,3 Hiroyuki Sasanuma,4 Shunichi Takeda,4 Akifumi Takaori-Kondo1*

Adult T cell leukemia (ATL) is an aggressive T cell malignancy caused by human T cell leukemia virus type 1 (HTLV-1) and has a poor prognosis. We analyzed the cytotoxic effects of various nucleoside analog reverse transcriptase inhibitors (NRTIs) for HIV-1 on ATL cells and found that abacavir potently and selectively kills ATL cells. Although NRTIs have minimal genotoxicities on host cells, the therapeutic concentration of abacavir induced numerous DNA double-strand breaks (DSBs) in the chromosomal DNA of ATL cells. DSBs persisted over time in ATL cells but not in other cell lines, suggesting impaired DNA repair. We found that the reduced expression of tyrosyl-DNA phosphodiesterase 1 (TDP1), a repair enzyme, is attributable to the cytotoxic effect of abacavir on ATL cells. We also showed that TDP1 removes abacavir from DNA ends in vitro. These results suggest a model in which ATL cells with reduced TDP1 expression are unable to excise abacavir incorporated into genomic DNA, leading to irreparable DSBs. On the basis of the above mechanism, we propose abacavir as a promising chemotherapeutic agent for ATL.

INTRODUCTION

Adult T cell leukemia (ATL) is caused by human T cell leukemia virus type 1 (HTLV-1) (1–3). Because of resistance to most cytotoxic drugs, patients suffering from ATL have an extremely poor prognosis with conventional chemotherapy [3-year overall survival rate of 24% (4, 5)]. Even allogeneic hematopoietic stem cell transplantation only increases the 3-year overall survival rate to a modest 33% (6). The combination of azidothymidine (AZT, also called zidovudine) and interferon-α (IFN-α), first proposed by Gill et al. (7) and Hermine et al. (8), is currently considered the most effective therapy, with an overall 5-year survival rate that reaches 46% (9). Some speculate that this combination therapy represses ATL cells by activating the immune response against HTLV-1–infected ATL cells as well as by interfering with the productive replication of HTLV-1 (10). However, the molecular mechanism underlying the cytotoxic effect of AZT against ATL remains elusive.

AZT is a first-generation nucleoside analog reverse transcriptase inhibitor (NRTI) (11) and has been widely used for HIV-1 patients. NRTI is efficiently incorporated into newly synthesizing viral DNA by HIV-1 reverse transcriptase, but not by replicative DNA polymerases in host cells. Efficient incorporation by reverse transcriptase is attributable to a lack of proofreading activity as well as to less stringent discrimination (12). Incorporated NRTIs prevent further extension of DNA synthesis, thereby inhibiting the productive replication of HIV-1. NRTIs approved for the treatment of HIV-1 are believed to have a minimal genotoxic effect on host cells.

Tyrosyl-DNA phosphodiesterase 1 (TDP1) was initially identified as a key enzyme for eliminating covalently bound topoisoamerase I (TopI) from cleaved DNA ends (13). A mutation in the TDP1 gene causes spinocerebellar ataxia with axonal neuropathy (SCAN1). A defect in TDP1 causes hypersensitivity to camptothecin (CPT11), a chemotherapeutic TopI poison, which stabilizes the covalent binding of TopI to cleaved DNA ends and kills cancer cells by inducing double-strand breaks (DSBs) during DNA replication, as shown in both SCAN1 patient–derived cell lines and knockout mouse models (14, 15).

Here, we found that a therapeutic concentration of abacavir (ABC), an NRTI, induces chromosomal DSBs and thereby kills ATL cells but not non–HTLV-1–infected cells. ABC is widely used as a key drug for the treatment of HIV-1 infection because it is well tolerated by >90% of patients with few adverse effects and has a desirable bioavailability (16). Once ABC is incorporated into the cells, it is phosphorylated in a unique stepwise anabolism to be converted to the triphosphated guanine analog carbovir (CBV) (17). The therapeutic concentration (16, 18, 19) of ABC causes a marked increase in the number of DSBs. We here demonstrate that this pronounced accumulation of DSBs caused by ABC results primarily from the down-regulation of TDP1 in ATL cells. We thus propose ABC as a novel and potent therapeutic agent for ATL.

RESULTS

ABC selectively kills HTLV-1–infected cells, including ATL cells

We first examined the cytotoxic effects of six NRTIs (ddI, d4T, 3TC, AZT, TDF, and ABC), all of which are current therapeutic agents for HIV-1, on ATL cell lines [ED-40515(−), ED-40515(+), SYK-11L(+), and ATL-43T] and on HTLV-1–infected cell lines (MT-2 and Hut-102). AZT was toxic to the ED-40515(−), SYK-11L(+), ATL-43T, and MT-2 cell lines but not to the others, whereas ddI, d4T, 3TC, and TDF showed poor cytotoxicity (Fig. 1A). Strikingly, the clinically relevant concentration of ABC was highly toxic to all ATL- and HTLV-1–infected cell lines but not to non–HTLV-1–infected cell lines (Jurkat, H9, and SU-DHL-6) (Fig. 1, A and B, fig. S1, and table S1). We confirmed the cytotoxicity of ABC to MT-2 cells by cell counting assays as well as MTS assay (fig. S2) because the mitochondrial toxicity of the NRTIs (20) could have affected the MTS assay. Because IFN-α is combined with AZT to treat ATL patients (7–9), we also tested...
the synergistic effect of IFN-α with AZT and ABC. IFN-α did not enhance the lethality of either AZT or ABC in either MT-2 or ED-40515(−) cells (Fig. 1C), suggesting that the cytotoxic effect of IFN-α in vivo may not be by a direct action (10). We therefore conclude that ABC potently and selectively kills HTLV-1–infected cells, including ATL cells, in vitro.

We next examined the effect of ABC on cell cycle and apoptosis in ATL cells. ABC induced S/G2-phase arrest and apoptosis in ED-40515(−) cells, but not in Jurkat cells (Fig. 2, A to C, and fig. S3). This finding suggests that ABC may cause DNA damage by prematurely terminating the replication of host chromosomal DNA, thereby activating the

**Fig. 1.** ABC selectively kills HTLV-1–infected and ATL cell lines. (A) Viability of the indicated cells after treatment (2 and 4 days) with 100 μM of the indicated NRTI. MTS values of treated cells relative to untreated cells are shown. Results are expressed as means ± SD of three independent experiments. (B) Viability of the indicated cells after treatment (2 days) with the indicated ABC dose. MTS values of treated cells relative to untreated cells are shown. (C) MT-2 and ED-40515(−) cells were treated with ABC or AZT together with IFN-α (0, 100, or 1000 IU/ml) for 2 days and analyzed with an MTS assay. MTS values of treated cells relative to untreated cells are shown.

**Fig. 2.** ABC induced S/G2-phase arrest and apoptosis. (A) Flow cytometry was used to analyze cell cycle status after propidium iodide (PI) staining. ED-40515(−) and Jurkat cells were treated with and without 100 μM ABC for 2 days and analyzed using flow cytometry. Results are expressed as means ± SD of three independent experiments. (B) Representation of flow cytometric analysis of cell cycle after PI staining. (C) Flow cytometric analysis of apoptosis after annexin V and PI double staining. ED-40515(−), MT-2, Hut-102, ATL-43T, and Jurkat cells were treated with the indicated dose of ABC or Adriamycin (ADR; 1 μg/ml) for 48 hours. Dark and light bars represent early and late apoptosis, respectively. Results are expressed as means ± SD of three independent experiments.
Fig. 3. ABC-induced DSB accumulation in ATL cells as a result of the impairment of DNA repair. (A) Chromosome breaks induced by ABC. ED-40515(−) and Jurkat cells were treated with the indicated doses of ABC and AZT for 2 days. Percentages of chromosome break-positive cells are shown. Results are expressed as means ± SD of three independent experiments. n.s., not significant. (B) Expression kinetics of γH2AX foci after acute exposure to ABC. ED-40515(−) and Jurkat cells were treated with 500 μM ABC for 12 hours. After treatment, the cells were stained with 4',6-diamidino-2-phenylindole (DAPI) and anti-γH2AX at the indicated times. (D) Percentages of γH2AX focus-positive cells at the indicated times. Focus-positive cells are those containing more than five γH2AX foci. (E) Cell proliferation of ED-40515(−) and Jurkat cells based on an MTS assay. MTS values relative to time zero are shown. (F) Expression kinetics of γH2AX foci at the indicated times after 2-Gy γ-irradiation. Irradiated ED-40515(−) and Jurkat cells were stained with DAPI and γH2AX. (G) Percentages of γH2AX focus-positive cells at the indicated times. (H) Cell proliferation of ED-40515(−) and Jurkat cells evaluated by an MTS assay. Data shown are as in (E).
DNA damage checkpoint to induce transient S/G2-phase arrest and apoptosis.

**ABC induces DSBs due to impaired DNA repair in ATL cells**

To characterize DNA damage, we measured chromosome breaks in mitotic ATL cells after exposure to ABC. The exposure significantly increased the number of chromosome breaks in ED-40515(−) cells (Fig. 3A, arrow) but not in Jurkat cells (Fig. 3B). The number of chromosome breaks induced by AZT in ED-40515(−) cells was several times smaller than that induced by ABC (Fig. 3B, left panel), which corresponds with the data in Fig. 1A. We therefore conclude that the therapeutic concentration of ABC as well as that of AZT efficiently induces DSBs in the chromosomal DNA of ATL cells, as does ionizing radiation.

To monitor the kinetics of DSB repair, we transiently exposed cells to a high dose of ABC for 12 hours and then monitored the number of unrepaired DSBs over time by counting the number of subnuclear foci containing the phosphorylated H2A histone variant (γH2AX). ED-40515(−) and Jurkat cells displayed comparable numbers of γH2AX foci after 4 hours, whereas resolution of γH2AX focus formation was indeed delayed in ED-40515(−) cells, but not in Jurkat cells, even after 48 hours (Fig. 3, C and D). Cellular proliferation was also reduced in ED-40515(−) cells (Fig. 3E). Thus, ATL cells are severely deficient in repairing ABC-induced DSBs. We next assessed the performance of the two major DSB repair pathways, homologous recombination (HR) and nonhomologous end-joining, in ATL cells. To this end, we monitored the kinetics of DSB repair after ionizing radiation. ED-40515(−) and Jurkat cells displayed very similar kinetics for DSB repair, indicating the competence of the canonical DSB repair pathways in these cells (Fig. 3, F to H). Because HR is responsible for repairing DSBs generated during DNA replication (21), we examined ongoing HR by measuring the polymerization of RAD51 recombinase, the essential HR factor, at DSB sites in ABC-treated cells. Treatment with ABC induced RAD51 focus formation, colocalizing with γH2AX foci in ED-40515(−) cells (Fig. 4, A and B). This finding suggests that frequent premature termination of DNA replication resulted in both the formation of DSBs and the activation of HR-dependent DSB repair to a significantly higher extent in ATL cells than in non-ATL cells.

**A defect in TDP1 is responsible for the cytotoxic effect of ABC on ATL cells**

To identify the factors required for efficient repair of ABC-induced DNA DSBs, we screened a panel of 48 isogenic mutant chicken DT40 cell lines deficient in DNA repair factors (22) for sensitivity to ABC (table S2). The result (fig. S4) was a telling sensitivity profile; the sensitivity of the five mutant cell lines (Tdp1, Poll, Usp1, Gen1, and Rad18) to ABC was more than twofold greater than that of the wild-type DT40 cell line (Fig. 5, A and B, and fig. S5A). Furthermore, the relative sensitivity of Tdp1-deficient DT40 cells to the six individual NRTIs correlated with their cytotoxic effect on ATL cells (Figs. 1A and 5B). These results suggest that a defect in TDP1 might account for the cytotoxic effect of ABC on ATL cells.

We next investigated TDP1 expression in ATL cells [Gene Expression Omnibus (GEO) database] according to the Joint Study on Prognostic Factors of ATL Development (23), and found that the level of TDP1 expression is indeed down-regulated in ATL cells relative to primary CD4+ T cells (Fig. 5, A and C, and fig. S5B). Likewise, the TDP1 protein and mRNA levels in ATL cell lines (Fig. 5D) and in primary ATL cells (Fig. 5, E and F, and table S3) were significantly lower than those in the noninfected T cells and primary CD4+ T cells, respectively. These observations suggest that a defect in TDP1 causes the sensitivity to ABC in ATL cells.

**TDP1 removes ABC from DNA ends in vitro**

To confirm that TDP1 plays a crucial role in ABC-induced DNA damage repair, we first examined whether human TDP1 (hTDP1) removed CBV that was covalently associated with the 3′ end of a DNA
oligonucleotide (Fig. 6A). We incubated total cell lysates from DT40 Tdp1−/− cells or Tdp1+/− cells complemented with hTDP1, with two substrate oligonucleotides containing either 3′-CBV or 3′-Tyr. The latter cellular lysate, but not the former one, efficiently generated the product that migrated one nucleotide shorter than the substrate (Fig. 6B). The processing efficacy of TDP1 for 3′-CBV increased in a dose-dependent manner (Fig. 6C). These biochemical results demonstrate that TDP1 removes 3′-blocking CBV efficiently in vitro.

**TDP1 catalytic activity is required for cellular tolerance to ABC**

To verify the importance of TDP1 in cellular tolerance to ABC, we depleted TDP1 and reconstituted TDP1-deficient cells with a TDP1 transgene. TDP1 small interfering RNA (siRNA)–treated Jurkat cells were more sensitive to ABC than the control siRNA–treated Jurkat cells (Fig. 7A, lanes 4 and 2, respectively). Conversely, reconstitution of MT-2 cells with human wild-type TDP1 markedly increased cellular tolerance to ABC (Fig. 7B and fig. S6, A to C). These data indicate that TDP1 plays an important role in cellular tolerance to ABC. The ectopic expression of TDP1WT did not reverse the sensitivity of MT-2 cells to AZT (Fig. S7), suggesting that hTDP1 eliminates AZT less efficiently than ABC. We conclude that the attenuated function of TDP1 is responsible for a high sensitivity to ABC in ATL cells.

To confirm that TDP1 catalytic activity is required for cellular tolerance to ABC, we prepared two mutant TDP1 transgenes [H263A and H493R (13, 24)] and a wild-type TDP1 transgene (25) and ectopically expressed them in MT-2 cells to generate MT-2/TDP1H263A, MT-2/TDP1H493R, and MT-2/TDP1WT cells, respectively (fig. S6, A to C). Ectopic expression of TDP1WT greatly increased MT-2 cell tolerance to ABC, but not that of the two mutant transgenes (Fig. 7B). Likewise, the phenotypes of the MT-2/TDP1H263A and MT-2/TDP1H493R cells were very similar to that of the parent MT-2 cells in terms of cellular response to ABC, including cell cycle arrest at the S phase (Fig. 7C), apoptosis (Fig. 7D), prolonged γH2AX focus accumulation (Fig. 7, E and F), and inhibition of cellular proliferation (Fig. 7G). We therefore conclude that TDP1 catalytic activity is required for cellular tolerance to ABC. This conclusion is supported by our data shown above and the previous reports showing that TDP1 can remove NRTIs localized at the 3′ ends of single-stranded oligonucleotides in vitro, because TDP1 eliminates trapped TopI from the 3′ ends of DNA strands in vivo (13, 26, 27).

**CPT11 and veliparib enhance the lethality of ABC on ATL cells**

CPT11 sensitivity is the classical hallmark of TDP1 loss. We verified a strong cytotoxic effect of CPT11 on ED-40515(−) cells (fig. S8A). We then tested whether a minimal dose of CPT11 synergistically increased the lethality of ABC on ATL cells. CPT11 significantly enhanced the lethality of ABC on ATL cells, with a low combination index (CI < 0.7; fig. S8B).

Furthermore, TDP1 removes trapped TopI from DNA by collaborating with poly(adenosine diphosphate–ribose) polymerase (PARP) (28). We thus theorized that combining ABC with PARP inhibitors might have a synergic cytotoxic effect on ATL cells by completely wiping out the residual activity of TDP1 in ATL cells. We then tested minimum doses of veliparib, which inhibits the catalytic activity of PARP without trapping PARP at DNA damage sites (29). Although veliparib alone had no obvious toxic effect on ATL cells, it significantly enhanced the lethality of ABC on ATL cells, with a low CI (<0.7; fig. S8C). These results suggest that a combination of ABC with CPT11 and/or PARP inhibitors could be a potent new therapeutic strategy against ATL.

**ABC inhibits the growth of ATL cell xenografts in NOD/SCID mice**

Finally, we tested the in vivo effect of ABC on ATL cell growth using an ATL xenograft model. Nonobese diabetic/severe combined...
ABC is converted to the triphosphated CBV in cells and then incorporated into host chromosomal DNA by replicative DNA polymerases, leading to premature termination of DNA replication, collapse of the replication fork, and DSB formation (Fig. 9). Thus, ABC is an effective chain terminator not only in the reverse transcription of retroviral DNA but also in the replication of host chromosomal DNA in ATL cells. This is totally unexpected because long-term treatment with ABC is well tolerated, causing only few side effects in HIV-1 patients. ABC has also been shown to be virtually nontoxic to bone marrow progenitor cells and leukemic cell lines (17). Our study identifies efficient DSB formation as the unique mechanism underlying the specific killing of ATL cells by the antiretroviral agent ABC.

Three mechanisms can inhibit NRTI genotoxicity. First, NRTIs are not potent inhibitors of replicative DNA polymerases (30, 31). Second, the catalytic pocket of replicative DNA polymerases is able to stringently discriminate between triphosphated NRTIs and normal nucleotides, with proofreading activities considered also as a contributing factor. Third, host DNA repair factors may be able to remove NRTIs when they are misincorporated into the 3′ end of primers by replicative DNA polymerases. The third mechanism accounts for the high toxicity of ABC to ATL cells, which, unlike non-HTLV-1–infected cells, are almost entirely unable to repair ABC-induced DSBs (Fig. 3, C and D). Moreover, our study demonstrates that a TDP1 defect is primarily responsible for the potent and selective cytotoxic effect of ABC on ATL cells. TDP1 can eliminate CBV misincorporated by replicative DNA polymerases into the 3′ ends (Fig. 6) because TDP1 eliminates TopI trapped at the 3′ ends of DNA cleavage (13) and oxidative 3′-blocking lesions (32). It is surprising that TDP1 could remove the 3′-blocking nucleotide analog that was not eliminated by the proofreading nuclease of replicative DNA polymerases. TDP1 seems to remove AZT from 3′ ends with considerably less efficiency than ABC because ectopic expression of a TDP1 transgene in MT-2 cells greatly increased cellular tolerance to ABC but not to AZT (Fig. 7B and fig. S7). We thus propose that ABC is a more potent chemotherapeutic agent against ATL than AZT. The current study also suggests that a therapy that combines ABC and CPT11 or a PARP inhibitor such as veliparib would provide a novel therapeutic strategy against ATL by enhancing the cytotoxic effect of ABC with minimum side effects on normal cells (fig. S8).

The molecular mechanism underlying reduced TDP1 gene expression is unclear. However, the current study reveals that the transcriptional status of the TDP1 gene may be a reliable biomarker for predicting the efficacy of ABC in anti-malignancy therapy. In addition, a recent report showing the defect of TDP1 in several lung cancer cell lines suggests the much broader indication of ABC to other cancers (33).

Finally, we demonstrate that ABC efficiently inhibits the growth of ATL cell xenografts in NOD/SCID mice. We administered ABC at 75 mg/kg per day orally. The plasma concentration of ABC in the mice was comparable to that in HIV-1 patients taking orally 410 mg of ABC once daily (34). Thus, the daily ABC dose given to HIV-1 patients might be sufficient for treating ATL, and our findings establish a rationale to expand its use for ATL patients. Moreover, ABC might prove useful for treating HTLV-1 asymptomatic carriers. A high HTLV-1 proviral load in the asymptomatic carrier promotes the progression to ATL and other HTLV-1–associated diseases (35). We here show that ABC is useful for suppressing HTLV-1–infected cells other than ATL cells (Fig. 1). In addition, ABC has very few side effects, even during long-term treatment. Thus, ABC would be an effective prophylactic agent in both HTLV-1 carriers with a high proviral immunodeficient (NOD/SCID) mice were inoculated with ED-40515(−) cells and treated daily with ABC via oral gavage at a dose of 75 mg/kg per day, which is comparable to 410 mg once daily of orally administered ABC in humans. Orally administered ABC significantly inhibited the growth of ATL tumor grafts (Fig. 8A) and improved survival (Fig. 8B).

**DISCUSSION**

In this report, we demonstrate that ABC specifically and potently kills ATL cells. ABC triggers apoptosis by inducing DSBs in host chromosomal DNA. The most likely mechanism for DSB induction is that ABC is a novel therapeutic strategy against ATL by enhancing the cytotoxic effect of ABC with minimum side effects on normal cells (fig. S8).
Depletion of TDP1 expression was confirmed by Western blot analysis 48 hours after transfection (right panel). Jurkat cells transfected with control siRNA or siTDP1 were treated with or without 300 μM ABC for 48 hours. MTS values of treated relative to untreated cells are shown (left panel). Results are expressed as means ± SD of three independent experiments. (B) MT-2 cells stably transfected with either wild-type (WT) TDP1, H263A TDP1, or H493R TDP1 transgenes. Stably transfected clones and control cells (mock) were treated with the indicated dose of ABC for 48 hours. Western blot analysis was conducted for expression of transgenes (right panel). Data shown as in (A, left panel). (C) Cell cycle analysis is shown as in Fig. 2A. MT-2/TDP1 WT, MT-2/TDP1 H263A, and MT-2/TDP1 H493R cells were treated with or without 100 μM ABC for 24 hours and subjected to flow cytometry. (D) The extent of apoptosis is shown as the percentage of annexin V–positive cells (y axis). Cells were treated as in (C). (E) The indicated cells were treated as in Fig. 3C. Data shown as in Fig. 3C. (F) Statistical analysis of data in (E). MT-2/TDP1 WT cells, but not MT-2/TDP1 H263A or MT-2/TDP1 H493R cells, are resistant to ABC. (G) Proliferation of the indicated cells is shown as in Fig. 3E.
Fig. 8. ABC inhibits the growth of ATL cell xenografts in NOD/SCID mice. (A) Tumor sizes of xenografted NOD/SCID mice treated with or without ABC. The mean sizes of individual tumors were plotted (control group, n = 7; ABC group, n = 6). P values were determined using Student’s t test (*P < 0.05). Error bars represent the SEM. (B) Kaplan-Meier survival curves for the ABC-treated and control mice. P values were determined using the Wilcoxon test.

Fig. 9. Specific lethality of ABC on ATL cells due to a defect in TDP1. ABC is phosphorylated in a unique stepwise anabolism and is converted to the triphosphate of CBV. During DNA synthesis, triphosphorylated ABC was incorporated into host chromosomal DNA by replicative DNA polymerases, leading to premature termination of DNA replication. In normal cells, TDP1 removes ABC quickly and DNA synthesis continues. However, in HTLV-1(+) cells, the collapse of the replication fork is induced because of the deficiency of TDP1, leading to DSB formation and apoptosis.
load and patients with HTLV-1–associated myelopathy/tropical spastic paraparesis (HAM/TSP), preventing the progression of these diseases.

**MATERIALS AND METHODS**

**Cell culture**

ED-40515(+) is an interleukin-2 (IL-2)–dependent T cell line of leukemic cell origin established from an ATL patient (36). ED-40515(−) is an IL-2–independent subclone of ED-40515(+) (37). SY is an IL-2–dependent HTLV-1–infected T cell line derived from a nonleukemic cell clone of ED-40515(+) from an ATL patient. ATL-43T is an IL-2–dependent T cell line established from double-negative (CD4−/CD8−) leukemic cells from a patient with ATL (38). SYK-11L(+) is a leukemic cell line from ATL tumor cells in an in vivo cell proliferation model using SCID mice (39). Hut-102 is an IL-2–independent T cell line from a patient with mycosis fungoides (40). MT-2 is a stably transformed HTLV-1–infected cell line, as previously reported (36). A summary of HTLV-1–infected and ATL cell lines is presented in table S4. Jurkat and H9 are IL-2–independent non–HTLV-1–infected human T cell lines. SU-DHL-6 is a non–HTLV-1–infected human B cell line. These cells were maintained in RPMI 1640 medium (Nacalai Tesque) containing 10% fetal bovine serum (FBS) and penicillin (100 U/ml), streptomycin (100 μg/ml), and L-glutamine (0.292 mg/ml) (PSG) (Invitrogen). The IL-2–dependent cell lines were maintained in the same medium with 0.5 mM recombinant human IL-2 (a gift from Shionogi Pharma). Human embryonic kidney (HEK) 293T cells were maintained in Dulbecco’s modified Eagle’s medium (Nacalai Tesque) containing 10% FBS and 1% PSG. The chicken DT40 cell lines used in this study were obtained from the Laboratory of Radiation Genetics, Graduate School of Medicine, Kyoto University (Kyoto, Japan). All mutant cell lines were previously authenticated (references provided in table S2). DT40 Tdp1+/− and Tdp1−/− hTDP1 cells (32) were gifts from Y. Pommier [Laboratory of Molecular Pharmacology, Center for Cancer Research, National Institutes of Health (NIH), Bethesda, MD]. The DT40 cells were cultured at 39.5°C with 5% CO2 in RPMI 1640 medium supplemented with 1% chicken serum (Gibco BRL), 10% FBS (Sigma-Aldrich), 10−5 M β-mercaptoethanol (Nacalai Tesque), penicillin (100 U/ml), streptomycin (100 μg/ml; Nacalai Tesque), and L-glutamine (0.292 mg/ml; Nacalai Tesque).

**Reagents and antibodies**

ABC, AZT, ddI, d4T, 3TC, and TDF were donated by the NIH through the AIDS reagent program. ABC was purchased from Sigma-Aldrich (SML0089) and Carbosynth (NA10019). Camptothecin (TG4110) and veliparib (ABT-888) were purchased from TopoGEN and Selleck Chemicals, respectively. These chemicals were dissolved in 100% dimethyl sulfoxide (DMSO, Nacalai Tesque) to a stock concentration of 10 to 500 mM and were stored at −20°C. Natural-type human IFN-α (Sumiferon) was purchased from Dainippon Sumitomo Pharma. Mouse anti-γH2AX (JBW301) was purchased from Merck Millipore, rabbit anti-RAD51 (H-92) was purchased from Santa Cruz Biotechnology, and rabbit anti-RAD51 (70–001) and mouse anti-RAD51 (B01P) were purchased from BioAcademia and Abnoba, respectively. Phycocerythrin– annexin V was purchased from BD Biosciences, and fluorescein isothiocyanate (FITC)–annexin V from BioLegend. PI (Nacalai Tesque) was dissolved in phosphate-buffered saline (PBS) to a stock concentration of 2 mg/ml. Rabbit anti-TDP1 (ab4166) was purchased from Abcam. Anti-β-actin (AC-15, A5441) was purchased from Sigma-Aldrich.

**Human cell killing assay**

To assess the sensitivity of human cell lines to the NRTIs with or without IFN-α or veliparib, 1×10⁴ to 2×10⁴ cells were seeded in 96-well flat-bottomed culture plates (final volume of 100 μl per well) and cultured for 0 to 96 hours at 37°C. Twenty microliters of CellTiter Glo 96 AQueous One solution from a cell proliferation assay kit (Promega) was added to each well, after which cell survival was determined by measuring the absorbance intensity at 490 nm using an ALVO X-3 (PerkinElmer). The absorbance values were normalized to those of controls.

**Cell cycle analysis**

The pretreated cells were fixed with chilled 70% ethanol for 30 min at 4°C and then stained with PI (5 μg/ml; Nacalai Tesque) in the presence of 1% bovine serum albumin (BSA; Nacalai Tesque) and ribonuclease (100 μg/ml; Sigma–Aldrich) for 30 min at room temperature. Immediately after staining, evaluation was performed using a FACS-Calibur analyzer, and the data were analyzed using ModFit software. DMSO treatment was used as a negative control, and Adriaemycin (1 μg/ml) treatment was used as a positive control.

**Apoptosis assay**

The pretreated cells were stained with FITC–annexin V (BioLegend) and PI for 10 min at 4°C. After staining, analysis was performed immediately using a FACSCalibur analyzer. DMSO treatment was used as a negative control, and Adriaemycin (1 μg/ml) treatment was used as a positive control.

**Chromosomal analysis**

The cells were cultured with colcemid (50 ng/ml) for 2 hours at 37°C, fixed with Carnoy’s solution, and placed on a glass slide. The slide was stained with 3.5% Giemsa (Nacalai Tesque) for 10 min, and the cells were examined by microscopy. Cells containing one chromosome break were defined as chromosome break–positive.

**Cell irradiation**

Cells were irradiated with 2 Gy of 135Cs γ–radiation using a Gamma-cell 40 Exactor (Best Theratronics).

**Immunofluorescence staining**

Prepared cells were placed on a glass slide via Cytospin. The cells were fixed with 4% formaldehyde (Nacalai Tesque) and washed with PBS. Next, the fixed cells were permeabilized using 0.1% NP-40/PBS for 10 min. After blocking with 3% BSA/PBS–Tween 20 for 15 min, the cells were incubated with the primary antibody at 37°C. Cells were then stained with Alexa Fluor 594 goat anti-mouse immunoglobulin G (IgG) (H+L) antibody, Alexa Fluor 488 goat anti-rabbit IgG (H+L) antibody, or Alexa Fluor 488 goat anti-mouse IgG (H+L) antibody (Molecular Probes) and DAPI (H-1200, Vector Laboratories) and observed using a fluorescence microscope (BZ-8100, Keyence).

**Evaluation of relative cellular sensitivity to NRTIs**

To assess the sensitivity of DT40 cells to various NRTIs, 1×10⁵ DT40 cells were seeded in 96-well black dishes (final volume of 100 μl per well) and continuously exposed to 25 μM of the indicated drug for 48 hours. CellTiter-Glo Luminescent Cell Viability Assay solution
These two patients mononuclear cells isolated from two acute-type ATL patients through Ficoll gradient centrifugation. Peripheral blood mononuclear cells were isolated from healthy donors and 10 University Hospital, and all participants provided written informed consent. Isolation of primary CD4 T cell and ATL cell samples

Total RNA was isolated from human samples using a High Pure RNA Isolation Kit (Roche). First-strand complementary DNA (cDNA) was generated using the PrimeScript II 1st strand cDNA Synthesis Kit (Takara). cDNA was amplified in a Thermal Cycler Dice Real Time System II (Takara) using SYBR Premix Ex Taq II (Takara). The data were normalized to the corresponding values for glyceraldehyde-3-phosphate dehydrogenase (GAPDH). The expression levels of the cells relative to normal cells (=1) are shown in Fig. 5 and fig. S5. Error bars represent SDs. The primers used were TDP1 forward, 5'-AGG-CAGCCCTTGGACAGATT-3'; TDP1 reverse, 5'-GGTACGCTGAGATTCGTGCC-3'; GAPDH forward, 5'-GAAGGTGAAGGGTCGAGTC-3'; and GAPDH reverse, 5'-GAAGATGGTGATGGATTTC-3'.

Western blot

A total of 5 × 10⁶ cells were lysed using ice-cold lysis buffer (MPER, Thermo Scientific) containing 1 mM phenylmethylsulfonyl fluoride, phosphatase inhibitor cocktail (Roche), and protease inhibitor cocktail (Nacalai Tesque), followed by centrifugation at 13,000 rpm for 15 min at 4°C. The cell lysates were mixed with an equal volume of twofold concentrated sample buffer (Bio-Rad Laboratories) containing β-mercaptoethanol (Nacalai Tesque) and were treated for 5 min at 100°C. Western blot was performed as described previously (41).

Isolation of primary CD4 T cell and ATL cell samples

This project was approved by the Institutional Review Board of Kyoto University Hospital, and all participants provided written informed consent in accordance with the Declaration of Helsinki. Peripheral blood mononuclear cells were isolated from healthy donors and 10 ATL patients through Ficoll gradient centrifugation. Peripheral blood mononuclear cells isolated from two acute-type ATL patients through Ficoll gradient centrifugation were used for Western blot (Fig. 5F). These two patients' characteristics are provided in table S3. The CD4⁺ T cell population was isolated using an EasySep Human CD4⁺ T Cell Enrichment Kit (STEMCELL Technologies).

In vitro biochemical assays for TDP1 activity

All single-standard DNA substrates share the commons sequence of 5'-TCCGTGGAAGCCTGTTT-3'. Oligonucleotides were synthesized in the reversed direction, as described previously (42). CBV was prepared according to the literature (43). Total cell lysates from DT40 Tdp1⁻/⁻ and Tdp1⁻/⁻ hTDP1 cells were prepared in the same manner as previously described (44). The protein concentration of lysates was determined by Bradford protein assay (Quick Start protein assay, Bio-Rad Laboratories). To prepare DNA substrate, oligonucleotides with 3-phosphotyrosine and CBV linkages synthesized were incubated with 1 M l-lysine, 1 M dithiothreitol, 1 M mercaptoethanol (Nacalai Tesque) and were treated for 5 min at 4°C. The cell lysates were mixed with an equal volume of twofold concentrated sample buffer (Bio-Rad Laboratories) containing 1 mM phenylmethylsulfonyl fluoride, phosphatase inhibitor cocktail (Roche), and protease inhibitor cocktail (Nacalai Tesque), followed by centrifugation at 13,000 rpm for 15 min at 4°C. The cell lysates were mixed with an equal volume of twofold concentrated sample buffer (Bio-Rad Laboratories) containing β-mercaptoethanol (Nacalai Tesque) and were treated for 5 min at 100°C. Western blot was performed as described previously (41).

Western blot

A total of 5 × 10⁶ cells were lysed using ice-cold lysis buffer (MPER, Thermo Scientific) containing 1 mM phenylmethylsulfonyl fluoride, phosphatase inhibitor cocktail (Roche), and protease inhibitor cocktail (Nacalai Tesque), followed by centrifugation at 13,000 rpm for 15 min at 4°C. The cell lysates were mixed with an equal volume of twofold concentrated sample buffer (Bio-Rad Laboratories) containing β-mercaptoethanol (Nacalai Tesque) and were treated for 5 min at 100°C. Western blot was performed as described previously (41).

Isolation of primary CD4 T cell and ATL cell samples

This project was approved by the Institutional Review Board of Kyoto University Hospital, and all participants provided written informed consent in accordance with the Declaration of Helsinki. Peripheral blood mononuclear cells were isolated from healthy donors and 10 ATL patients through Ficoll gradient centrifugation. Peripheral blood mononuclear cells isolated from two acute-type ATL patients through Ficoll gradient centrifugation were used for Western blot (Fig. 5F). These two patients' characteristics are provided in table S3. The CD4⁺ T cell population was isolated using an EasySep Human CD4⁺ T Cell Enrichment Kit (STEMCELL Technologies).

In vitro biochemical assays for TDP1 activity

All single-standard DNA substrates share the commons sequence of 5'-TCCGTGGAAGCCTGTTT-3'. Oligonucleotides were synthesized in the reversed direction, as described previously (42). CBV was prepared according to the literature (43). Total cell lysates from DT40 Tdp1⁻/⁻ and Tdp1⁻/⁻ hTDP1 cells were prepared in the same manner as previously described (44). The protein concentration of lysates was determined by Bradford protein assay (Quick Start protein assay, Bio-Rad Laboratories). To prepare DNA substrate, oligonucleotides with 3-phosphotyrosine and CBV linkages synthesized were incubated with [γ-³²P]adenosine triphosphate (NEG502A, PerkinElmer Life Sciences) and T4 polynucleotide kinase (Takara) for 5'-end labeling. After being purified using a nucleotide removal kit (Qiagen), 1 nM labeled oligonucleotide was incubated with cell lysates at 25°C for 15 min. Samples were separated by 16% denaturing polyacrylamide gel electrophoresis (7 M, urea). Dried gel was exposed on the imaging plate (FujiFilm) and then scanned in Fuji Bas 2500 system. Quantification was performed by Image Gauge software (FujiFilm).

RNA interference

To evaluate the role of TDP1 in the cell sensitivity to ABC, we knocked down TDP1 in Jurkat cells using specific siRNA. Transient TDP1 knockdown was achieved using the siGENOME SMARTpool (Thermo Scientific). A total of 5 × 10⁶ Jurkat cells were electroporated with either nontargeted siRNA (Stealth RNAi NEGATIVE CONTROL, Invitrogen) or hTDP1-specific siRNA (20 pmol/µl) using the Amazax Cell Nucleofector System (Lonza). Knockdown efficiency was determined by quantitative PCR 24 hours after transfection and Western blot 48 hours after transfection.

Plasmid constructs

Plasmids encoding FLAG-TDP1 wild type, FLAG-TDP1 H263A, and FLAG-TDP1 H493R were a gift from Y. Pommier (NIH). H263A and H493R are inactive TDP1 mutants; the latter causes spinocerebellar ataxia with axonal neuropathy (SCAN1). Wild-type, H263A, and H493R FLAG-TDP1 were cloned into the lentiviral vector CSII-CMV-MCS-IRE-S-hrGFP.

Generation of MT-2 cells reconstituted with hTDP1

Wild-type, H263A, and H493R FLAG-TDP1-expressing lentiviral vectors were prepared as described previously (41). MT-2 cells were infected with each viral vector. Confirmation of infectivity was based on human recombinant green fluorescent protein (hrGFP) expression, and the expression level of TDP1 in each group of infected MT-2 cells was verified by Western blot with rabbit anti-TDP1.

Preparation of ATL model mice

The rapid tumor formation by the ATL cell line in NOD/SCID mice has been previously established (38, 45). Four-week-old NOD/SCID mice were purchased from CLEA Japan. Fourteen mice were anesthetized with ether, and 4 × 10⁶ ED-40515(−) cells were subcutaneously inoculated into the posterior cervical lesion. Beginning 2 weeks after inoculation, the long and short axes were measured weekly. The tumor volume was approximated as (long axis) × (short axis)². All experiments were performed under the approved protocols of the Institute of Laboratory Animals, Graduate School of Medicine, Kyoto University.

Administration of ABC

ABC dissolved in water was administered orally to mice at a dose of 75 mg/kg per day on weekdays from day 0 to day 20. The control mice received an equivalent amount of water.

Analysis of combined treatment effects

The synergistic effects of the combined treatments were analyzed using the Chou-Talalay method (46). The CI of each combined treatment was calculated using CompuSyn software, and CIs of 0.3 to 0.7, 0.1 to 0.3, and <0.1 were defined as synergism, strong synergism, and very strong synergism, respectively (47).

Statistical analysis

The statistical significance of the differences between the drug-treated and control cultures was determined using Student’s t test or the χ² test. The statistical significance of the survival rates was determined using the Wilcoxon test.


Acknowledgments: We thank M. Maeda for providing the ED-40515(−), ED-40515(+), and ATL-43T cells, and M. Takada (Kyoto University) for providing the Fancc$^{-/-}$, Fancg$^{-/-}$, Smn$^{-/-}$, and Artemis$^{-/-}$ DT40 cells. We also thank K. Imada for his suggestion regarding the use of the mouse xenograft model. We are grateful to H. Mitsuya, T. Watanabe, and J. Murai for helpful discussions. We also thank J. Hejna for English language editing. Funding: This work was partially supported by the Japan Society for the Promotion of Science (JSPS) KAKENHI program; grants-in-aid from the Ministry of Education, Culture, Sports, Science, and Technology of Japan; and Health and Labour Sciences research grants from the Ministry of Health, Labour, and Welfare of Japan. Author contributions: K.T. and M.K. designed and performed most of the experiments and wrote the manuscript. Y.T., F.I., T.S., K.N., M.S., K.H., J.Y., S.I., and K.I. performed some of the experiments, and K.S., M.H., K.S., and N.K. analyzed the data and interpreted the results. H.S. and S.T. participated in the experimental design and provided materials. A.T.-K. participated in the experimental design, supervised and funded the project, and wrote the manuscript. Competing interests: The authors declare that they have no competing interests.
Abacavir, an anti–HIV-1 drug, targets TDP1-deficient adult T cell leukemia

Kohei Tada, Masayuki Kobayashi, Yoko Takiuchi, Fumie Iwai, Takashi Sakamoto, Kayoko Nagata, Masanobu Shinhara, Katsuhiro Ito, Kotaro Shirakawa, Masakatsu Hishizawa, Keisuke Shindo, Norimitsu Kadowaki, Kouji Hirota, Junpei Yamamoto, Shigenori Iwai, Hiroyuki Sasumuma, Shunichi Takeda and Akifumi Takaori-Kondo

Sci Adv 1 (3), e1400203.
DOI: 10.1126/sciadv.1400203