

OBP-301 infection induced death of human cancer cell lines (H1299 human lung cancer and SW620 human colorectal cancer cells) in a dose-dependent manner (Figure 1). Although autophagy, or type II programmed cell death, partially involved in the cell death machinery triggered by OBP-301 infection, oncolytic cells are distinct from apoptotic cells (Supplementary Figure 1).

We next examined whether OBP-301 infection modulated intracellular concentrations of uric acid that might act as a danger signal in tumor cells. Uric acid levels increased in H1299 cells following OBP-301 infection in a time-dependent fashion, although docetaxel slightly upregulated the uric acid concentration 72 h after treatment (Figure 2a). Thus, tumor cells undergoing oncolysis can produce significantly greater amounts of uric acid when compared with apoptotic tumor cells. The uric acid elevation pattern of OBP-301-infected cells almost paralleled that of cells infected with Onyx-015, an E1B 55 kDa-deleted adenovirus engineered to selectively replicate in and lyse p53-deficient cancer cells, and wild-type adenovirus type 5 (Figure 2b), indicating a general effect of adenovirus infection in the regulation of intracellular uric acid levels.

Uric acid is produced during the catabolism of purines and is the end product of this process. Adenoviral replication facilitates the purine catabolism to stimulate the synthesis of progeny DNA, which in turn may increase intracellular uric acid levels by the purine degradation process. In fact, OBP-301 infection significantly increased the amount of uric acid in the cells, whereas replication-deficient dl312 infection had no apparent effect on the levels of uric acid. OBP-301-induced elevation of uric acid levels could be inhibited in the presence of cidofovir (CDV), an acyclic nucleoside phosphonate having potent broad-spectrum anti-DNA virus activity (Figure 2c). CDV has been approved for the treatment of many types of viruses including cytomegalovirus and adenovirus (Lenaerts and Naesens,

2006). We confirmed that CDV at 100 μ M could significantly inhibit replication of OBP-301 in H1299 cells by the real-time quantitative PCR analysis (Supplementary Figure 2). Moreover, as OBP-301 replication was attenuated in telomerase-negative cells, the levels of uric acid could not be altered in normal human lung fibroblasts (NHLF) after OBP-301 infection (Figure 2d). These results suggest that viral replication is required to produce uric acid in infected cells.

Xanthine oxidoreductase (XOR) is a member of the molybdoflavoenzyme family that catalyses the formation of uric acid from xanthine and hypoxanthine (Glantzounis *et al.*, 2005). A strand-specific reverse transcriptase PCR assay demonstrated that XOR mRNA expression gradually decreased in OBP-301-infected cells presumably due to the negative feedback of increased uric acid levels, whereas docetaxel-treated cells yielded consistent bands of the XOR transcripts (Figure 2e). Thus, adenoviral replication could directly stimulate the catalytic DNA turnover, which enables cells to produce more uric acid.

We then examined the ability of OBP-301-infected cells to stimulate immature DCs *in vitro*. DCs generated from HLA-A24⁺ healthy volunteers were co-cultured with HLA-matched H1299 cells (HLA-A32/A24) treated with OBP-301 or docetaxel for 72 h, or freeze thawed. The production of Th1 cytokines such as IFN- γ and IL-12 in the supernatants was then explored by enzyme-linked immunosorbent assay (ELISA) analysis 48 h after the co-culture. DCs incubated with OBP-301-infected cells secreted large amounts of IFN- γ and IL-12, whereas stimulation with docetaxel-treated apoptotic cells induced their secretion at low levels (Figure 3a). The level of cytokine production from DCs incubated with freeze-thawed necrotic cells was similar to that of untreated immature DCs. Moreover, we confirmed that addition of OBP-301 alone without target tumor cells did not affect the cytokine secretion of DCs into the supernatant, indicating that infection of OBP-301 itself had no apparent effect on DCs. Thus, DCs stimulated with oncolytic tumor cells preferentially secrete high-level Th1 cytokines. Flow cytometry demonstrated that the increase in the expression of CD83, which is expressed on mature DCs, was slightly higher on DCs incubated with oncolytic cells than those with apoptotic or necrotic cells, indicating that oncolytic tumor cells seems to have a positive influence on DC maturation (Supplementary Figure 3).

In the next step, we investigated the effects of oncolytic tumor cells on T-cell activation in the presence of DCs. H1299 cells were infected with OBP-301 over 72 h, and then co-incubated with HLA-matched HLA-A24⁺ peripheral blood mononuclear cells (PBMCs) for another 48 h in mixed lymphocyte tumor culture (MLTC). In other tests, H1299 cells were exposed to docetaxel for 72 h or freeze thawed, and then co-cultured with PBMCs. We examined the secretion of IFN- γ and IL-12 into the supernatants after MLTC for 7 days. Stimulation with OBP-301-infected cells induced the secretion of high levels of IFN- γ and IL-12 into MLTC supernatants, which was significantly higher

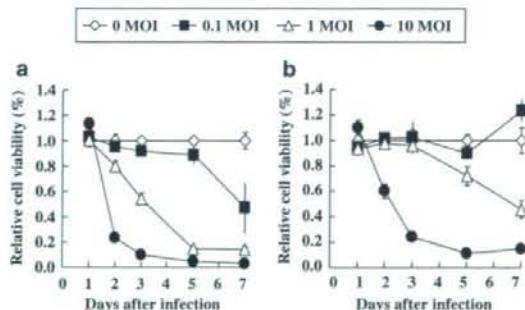


Figure 1 *In vitro* cytopathic effects of OBP-301 on human cancer cells. H1299 human non-small cell lung cancer (a) and SW620 human colorectal cancer cells (b) were infected with OBP-301 at indicated multiplicity of infection (MOI) values, and surviving cells were quantitated over 7 days by XTT assay. The cell viability of mock-treated cells on day 1 was considered 1.0, and the relative cell viability was calculated. Each data represent the mean \pm standard deviation (s.d.) of triplicate experiments.

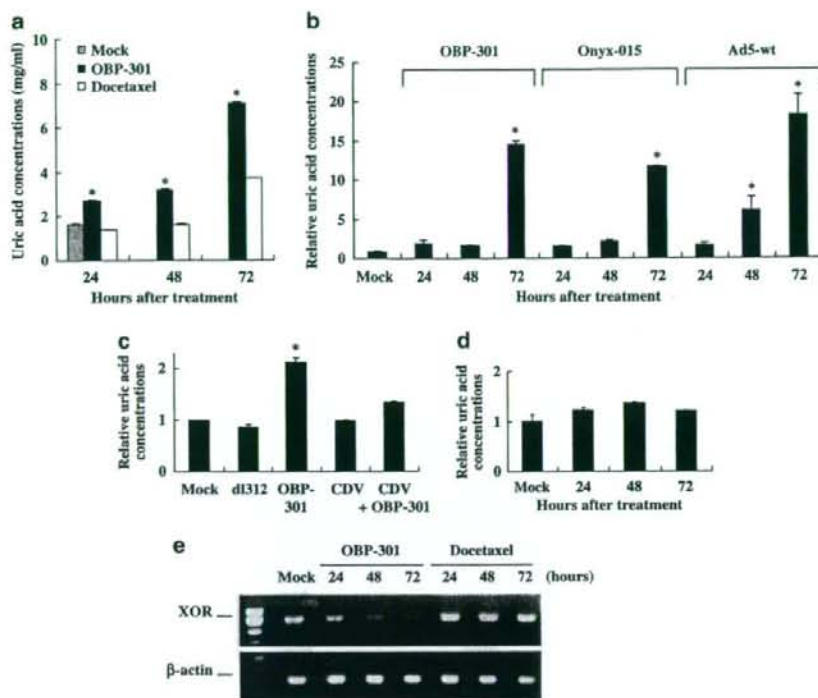


Figure 2 (a) Uric acid concentrations in H1299 cells treated with OBP-301 or docetaxel. H1299 cells were infected with 1.0 MOI of OBP-301 or treated with 10 nM of docetaxel for indicated time periods, and uric acid concentrations were determined enzymatically in the cell homogenates. Single asterisk indicates $P < 0.01$, significantly different from docetaxel-treated cells. (b) Uric acid levels in H1299 cells treated with OBP-301, Onyx-015 or wild-type adenovirus. H1299 cells were harvested at indicated time points over 72 h after infection with 10 MOI of viruses, and subjected to the measurement of uric acid concentrations. The levels of uric acid concentration are defined as the fold-increase for each sample relative to that of mock-treated cells (mock equals 1). Single asterisk indicates $P < 0.01$, significantly different from mock-treated cells. (c) Uric acid concentrations in H1299 cells infected with 1.0 MOI of OBP-301 or replication-deficient dl312 adenovirus were measured 24 h after infection. Uric acid production was also assessed in H1299 cells infected with 1.0 MOI of OBP-301 in the presence of 100 μM of anti-virus agent cidofovir (CDV). H1299 cells treated with 100 μM of CDV were subjected to the assay as a control. All uric acid levels are normalized to that of mock-treated cells (mock equals 1). Single asterisk indicates $P < 0.01$, significantly different from mock-treated cells. (d) Uric acid levels in NHLF infected with OBP-301. NHLF cells were infected with 1.0 MOI of OBP-301 for indicated time periods, and uric acid concentrations were measured. The uric acid levels are normalized to that of mock-treated cells. (e) Detection of xanthine oxidoreductase (XOR) mRNA expression in OBP-301-infected H1299 cells by RT-PCR analysis. Cells were infected with 1.0 MOI of OBP-301 or treated with 10 nM of docetaxel, and then collected at the indicated time points. First-strand DNA generated from RNA was amplified using either the primers specific for XOR sequence or the primers that recognize β -actin sequences as an internal control.

than that with docetaxel-treated or freeze-thawed H1299 cells (Figure 3b). Thus, oncolytic tumor cells can accelerate the cleavage of tumor antigen peptides that can be associated with MHC class I molecules via IFN- γ secretion by immune cells.

Stimulation of cells with IFN- γ is known to induce the expression of PA28, a proteasome activator that accelerates the *in vitro* processing of MHC class I ligands from their polypeptide precursors (Sun *et al.*, 2002). We investigated whether PA28 expression was upregulated in H1299 cells by adding the supernatants of co-cultures of PBMCs and OBP-301-infected H1299 cells. Western blot analysis for PA28 demonstrated that, following heat inactivation of residual OBP-301, MLTC supernatants with oncolytic tumor cells induced a strong endogenous PA28 expression in H1299 cells. In contrast, exposure to the supernatants of PBMCs alone, PBMCs with untreated H1299 cells, and PBMCs with oncolytic

tumor cells without heat inactivation resulted in no apparent changes in the expression levels of PA28 (Figure 4).

Finally, the cytotoxic T-lymphocyte (CTL) response against human cancer cells was assessed by a standard 6-h ^{51}Cr release assay after a 7-day MLTC using various forms of H1299 cells. The lytic activity of CTLs induced by apoptotic or necrotic H1299 cells was comparable with that of human lymphokine-activated killer (LAK) cells; CTLs stimulated with oncolytic H1299 cells, however, more efficiently killed target H1299 cells (Figure 5). In contrast, LAK cells effectively lysed SW620 cells, whereas these cells were minimally killed by CTLs stimulated with apoptotic, necrotic or oncolytic H1299 cells. Furthermore, HLA-unmatched, HLA-A26/A30+ A549 human lung cancer cells were not sensitive to oncolytic tumor cell-induced cytotoxicity (data not shown), suggesting that effector cells stimulated with

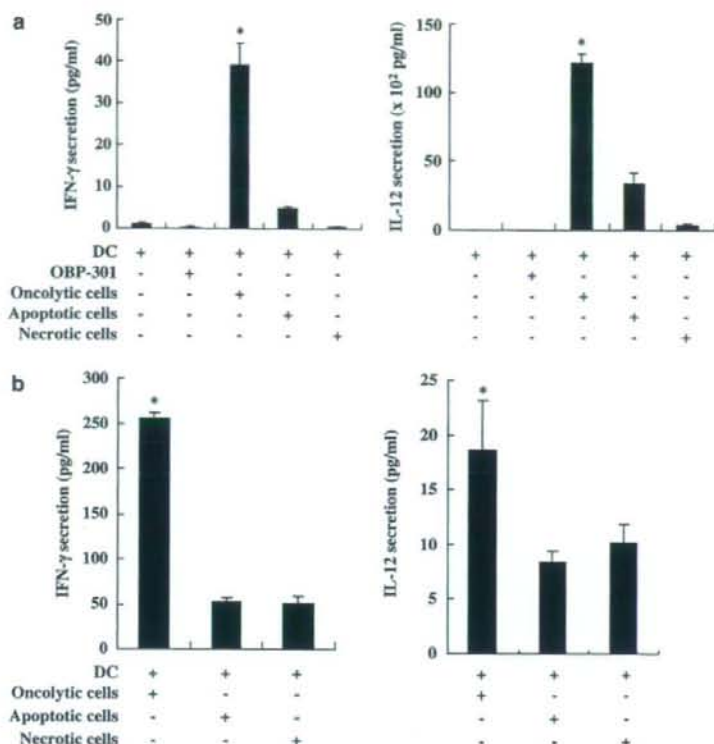


Figure 3 (a) Secretion of Th1-type cytokines by oncolytic, apoptotic or necrotic tumor cells. H1299 cells were treated with 1.0 MOI of OBP-301 or 50 nM of docetaxel for 72 h, or freeze thawed, and then co-cultured with immature dendritic cells (DCs) obtained from monocytes for additional 48 h. The culture supernatants were harvested and tested by ELISA for interferon (IFN)- γ (left) and interleukin (IL)-12 (right) concentrations. As a control, the supernatants of immature DCs alone or with OBP-301 at an MOI of 1.0 were also examined. Data are mean \pm s.d. of triplicate experiments. Single asterisk indicates $P < 0.01$, significantly different from other groups. (b) Tumor-specific CTL induction in MLTC with oncolytic, apoptotic or necrotic tumor cells. IFN- γ (left) and IL-12 (right) concentrations in the supernatants of MLTC analysed by ELISA. H1299 cells were treated with 1.0 MOI of OBP-301 or 50 nM of docetaxel for 72 h, or freeze thawed, and then co-cultured with PBMCs obtained from HLA-A24* healthy volunteers for 48 h in MLTC. Data are mean \pm s.d. of triplicate experiments. Single asterisk indicates $P < 0.01$, significantly different from other groups.

OBP-301-infected tumor cells exhibit MHC class I-restricted reactivity.

Discussion

In the present study, our goal was to determine whether oncolytic virus is effective not only as a direct cytotoxic drug but also as an immunostimulatory agent that could induce specific CTL for the remaining antigen-bearing tumor cells. Several groups have debated whether necrotic or apoptotic cells can stimulate DCs to cross-present cell-derived peptides, with subsequent enhancement of tumor immunogenicity. Furthermore, it has been reported recently that the immunogenicity of tumors is not regulated by signals associated with apoptotic or necrotic cell death, but is an intrinsic feature of the tumor itself (Bartholomae et al., 2004). Our data indicate that viral oncolysis could efficiently load tumor antigen on DCs, and then generate CTL response as judged from

the production of cytokines. Moreover, the CTL activity against untreated tumor cells suggests that CTLs are specific to tumor antigens, but not to adenovirus proteins.

DCs are known to ingest dying tumor cells and initiate tumor-specific responses when associated with appropriate danger signals, which are endogenous activation signals liberated by dying cells. Recent studies have shown that some intrinsic biochemical factors, such as uric acid, bradykinin and heat shock protein (HSP110) act as danger signals through their interaction with DCs, and influence the subsequent immune response (Aliberti et al., 2003; Shi et al., 2003; Manjili et al., 2005). Large amounts of uric acid can be produced following tissue injury *in vivo*, and activate the immune response against injured cells and dying tissues. We found that OBP-301 infection increased intracellular uric acid levels in human tumor cells compared with apoptosis- or necrosis-inducing stimuli, suggesting that viral replication itself can enhance tumorigenicity.

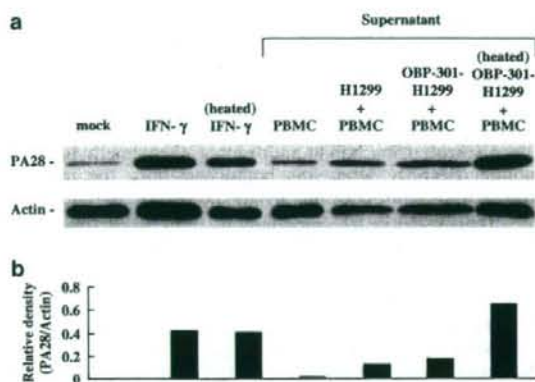


Figure 4 (a) Western blot analysis of PA28 in H1299 cells exposed to the supernatants of MLTC. peripheral blood mononuclear cells (PBMCs) were incubated with mock, untreated H1299 cells or H1299 cells treated with 10 MOI of OBP-301 for 72 h in MLTC, and the supernatants were harvested 48 h after the co-culture. H1299 cells were further incubated with the supernatants for 72 h with or without heat inactivation of residual virus (56 °C, 10 min). H1299 cells were also incubated with 5 ng ml⁻¹ of interferon (IFN)- γ with or without heating for 72 h. Equivalent amounts of protein obtained from whole cell lysates were loaded in each lane, probed with anti-PA28 antibody and then visualized by using an ECL detection system. Equal loading of samples was confirmed by stripping each blot and reprobing with anti-actin antiserum. (b) PA28 protein expression was quantified by densitometric scanning using NIH Image software and normalization by dividing the actin signal.

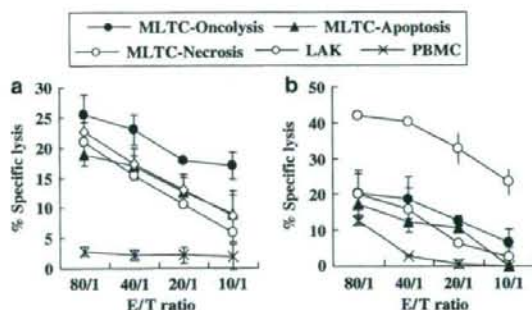


Figure 5 Cytolytic reactivity against H1299 (a) and SW620 (b) human cancer cells was assessed after 7-day mixed lymphocyte tumor culture (MLTC) with oncolytic, apoptotic or necrotic H1299 cells treated the same as above by 6-h standard ⁵¹Cr-release assay. Lymphokine-activated killer (LAK) cells were generated from peripheral blood mononuclear cells (PBMCs) in the presence of interleukin (IL)-2 (100 U ml⁻¹) for 3 days. The CTLs were compared with LAK cells and untreated PBMCs, which served as positive and negative controls, respectively. Data represent the mean \pm s.d. of three wells at four different effector-to-target (E/T) ratios.

Viral oncolysis increases the immunogenicity of tumor cells presumably by the release of proinflammatory cytokines (Lindenmann and Klein, 1967). We showed that OBP-301-infected oncolytic tumor cells

efficiently stimulated immature DCs to produce greater amounts of IFN- γ and IL-12 than apoptotic and necrotic cells, and that such stimulation led to DC maturation. Viral infection itself has been reported to activate DCs to secrete pro- or anti-inflammatory cytokines, which can drive DCs to undergo the maturation process (Ho *et al.*, 2001); the observation that OBP-301 alone had no effect on cytokine production by DCs, however, indicates that OBP-301 itself may be less infective or stimulatory to DCs. The result is consistent with our finding that OBP-301 attenuated replication as well as cytotoxicity in human normal cells.

It will be of interest to more mechanistically define why viral oncolysis efficiently induces CTL activity against tumor cells. We hypothesized that viral replication itself or the released cytokines by immune cells positively influences tumor cell immunogenicity. The IFN- γ -inducible proteasome modulator complex PA28 participates in the generation of antigenic peptides required for MHC class I antigen presentation (Sijts *et al.*, 2002). As expected, the supernatants of MLTC with OBP-301-infected tumor cells, in which IFN- γ secretion was detected, induced a strong expression of endogenous PA28. Thus, oncolytic tumor cells can accelerate the cleavage of tumor antigen peptides that can be associated with MHC class I molecules via IFN- γ secretion by immune cells. In fact, it has been reported that restoration of PA28 expression in PA28-deficient melanoma cells rescues the melanoma antigen epitope presentation (Sun *et al.*, 2002); our preliminary experiments however demonstrated that human tumor cells transfected with PA28 α expression vector were less sensitive to tumor-specific CTLs (data not shown). These observations suggest that antigen peptide production alone does not seem sufficient to enhance tumor immunogenicity.

In conclusion, we provide for the first time evidence that oncolytic virus replication induces tumor-specific immune responses by stimulating uric acid production as a danger signal as well as accelerating tumor antigen cleavage by IFN- γ -inducible PA28 expression. Since the induction of systemic immunity has rarely been observed in clinical trials with other conditionally replication-competent viruses, more *in vivo* experiments are clearly required to support the induction of antitumor immunity by OBP-301 treatment. Our data, however, suggest that the antitumor effect of OBP-301 might be potentially both direct and indirect as well as systemic rather than local.

Materials and methods

Cell lines and reagents

The human non-small lung cancer cell lines H1299 (HLA-A32/A24) and the human colorectal carcinoma cell lines SW620 (HLA-A02/A24) were maintained *in vitro* in RPMI 1640 supplemented with 10% fetal calf serum, 100 U ml⁻¹ penicillin and 100 mg ml⁻¹ streptomycin. Recombinant human cytokines granulocyte/macrophage colony-stimulating factor (GM-CSF), IL-4, TNF- α and IL-7 were purchased from Genzyme

Techne (Minneapolis, MN, USA), IFN- γ from Peptrotech (Rocky Hill, NJ, USA) and IL-2 from Roche (Mannheim, Germany). [^{51}Cr] sodium chromate was obtained from NEN Life Science Products (Boston, MA, USA). Docetaxel (taxotere) was kindly provided by Aventis Pharma (Tokyo, Japan).

Adenovirus

The recombinant replication-selective, tumor-specific adenovirus vector OBP-301 (Telomelysin), in which the hTERT promoter element drives the expression of *E1A* and *E1B* genes linked with an IRES, was constructed and characterized previously (Kawashima et al., 2004; Umeoka et al., 2004; Taki et al., 2005; Watanabe et al., 2006). Onyx-015 (dl1520) is an E1B 55 kDa-deleted adenovirus engineered to selectively replicate in and lyse p53-deficient cancer cells, and kindly provided by Dr Frank McCormick (UCSF Comprehensive Cancer Center and Cancer Research Institute). The E1A-deleted adenovirus vector lacking a cDNA insert (dl312) was also used as a control vector. The viruses were purified by CsCl₂ step gradient ultracentrifugation followed by CsCl₂ linear gradient ultracentrifugation.

Cell viability assay

XTT assay was performed to measure cell viability. Briefly, cells were plated on 96-well plates at 5×10^3 per well 24 h before treatment and then infected with OBP-301 or exposed to docetaxel. Cell viability was determined at the times indicated by using a Cell Proliferation Kit II (Roche Molecular Biochemicals) according to the protocol provided by the manufacturer.

Reverse transcription (RT)-PCR

Total RNA was isolated from mock-, OBP-301- and docetaxel-treated cells using RNeasy (Qiagen/BioTeck, Friendswood, TX, USA) in a single-step phenol-extraction method and used as templates. Reverse transcription was performed at 22°C for 10 min and then 42°C for 20 min using 1.0 μg of RNA per reaction to ensure that the amount of amplified DNA was proportional to that of specific mRNA in the original sample. PCR was performed with specific primers in volumes of 50- μl according to the protocol provided by the manufacturer (PCR kit; Perkin-Elmer/Cetus, Norwalk, CT, USA). The specific primers used for XOR were 5'-GCG AAG GAT AAG GTT ACT TGT-3' (forward) and 5'-CTC CAG GTA GAA GTG CTC TTG-3' (reverse); and for β -actin were 5'-ATG GTG GGA ATG GGT CAG AAG-3' (forward) and 5'-GCA GCT CAT TGT AGA AGG-3' (reverse). The reaction conditions were denaturing at 94°C for 2 min followed by 30 cycles consisting of denaturing at 94°C (30 s), annealing at 65°C (15 s) and extension at 72°C (10 s) using a thermal cycler (Perkin-Elmer, Foster City, CA, USA). The reactions were completed by a final 2-min extension at 72°C. The PCR products were resolved on 1% agarose gels and visualized by SYBR Gold Nucleic Acid Gel Stain (Molecular Probes Inc., Eugene, OR, USA).

Preparation of tumor cells

For induction of oncolysis, tumor cells were infected with OBP-301 at a multiplicity of infection (MOI) of 1–10, and then collected 24–72 h after infection. Apoptotic tumor cells were obtained after 24–72-h exposure to 50–100 nM of docetaxel. For induction of necrosis, tumor cells suspended in phosphate-buffered saline (PBS) were subjected to rapid four freeze/thaw cycles using a 60°C water bath and liquid nitrogen.

Measurement of uric acid concentration

Cultured cells were harvested after treatment and rinsed three times with PBS. These cells were resuspended in lysis buffer at a density of 200×10^6 cells per 100 μl . The buffer contained 10 mM Tris-HCl (pH 7.5), 150 mM NaCl, 50 mM NaF, 1 mM ethylenediaminetetraacetic acid (EDTA), 0.5 mM Na₂VO₄, 10% glycerol, 0.5% NP-40 and 0.1 mM phenylmethylsulfonyl fluoride (PMSF). After 10-s homogenization, the resulting extracts were kept on ice for 30 min and then were centrifuged for 15 min at 2000 g. The supernatants from treated tumor cells were assayed for uric acid using Uric Acid C test (Wako, Osaka, Japan).

Preparation of DCs

Peripheral blood samples were obtained from normal HLA-A24 positive healthy volunteers and PBMC were isolated by sedimentation over Ficoll-Hypaque. They were subsequently allowed to adhere in culture flasks for 1 h at 37°C at a density of 4.0×10^7 cells per plate. Non-adherent cells in the plate were removed and the remaining (adherent) cells were cultured for 7 days in AIM-V (Gibco, Rockville, MD, USA) containing 2% heated-inactivated autologous serum supplemented with GM-CSF (50 ng ml⁻¹) and IL-4 (50 ng ml⁻¹).

Cytokine production assay

DCs were co-cultured with treated tumor cells at a ratio of 3:1 (DC/tumor cell) in a culture medium containing GM-CSF (50 ng ml⁻¹) and IL-4 (50 ng ml⁻¹). After 24-h incubation, the supernatant was collected and stored at -80°C until the assay. The concentrations of IFN- γ and IL-12 (p40 and p70) were measured with appropriate ELISA kits (BioSource, Camarillo, CA, USA).

MLTC and CTL assay

PBMCs were co-cultured with treated tumor cells at a ratio of 20:1 in the presence of IL-2 (Roche) (10 U ml⁻¹) and IL-7 (Genzyme Techne) (5 ng ml⁻¹) for 7 days. Cultured cells were then used as effector cells in a standard 4 h-⁵¹Cr release assay and the percentage of lysed cells was calculated. Percent specific lysis = ((experimental cpm - spontaneous cpm) / (maximal cpm - spontaneous cpm)) \times 100. Supernatants from MLTC performed as above were also assayed for IFN- γ and IL-12 by ELISA assays (BioSource).

Western blot analysis

The primary antibodies against proteasome activator PA28 (ZMD353; Invitrogen, Carlsbad, CA, USA), actin (AC-40; Sigma Chemical Co., St. Louis, MO, USA) and peroxidase-linked secondary antibody (Amersham, Arlington Heights, IL, USA) were used. Cells were washed twice in cold PBS and collected, then lysed in lysis buffer (10 mM Tris (pH 7.5), 150 mM NaCl, 50 mM NaF, 1 mM EDTA, 10% glycerol and 0.5% NP40) containing proteinase inhibitors (0.1 mM PMSF and 0.5 mM Na₂VO₄). After 20 min on ice, the lysates were spun at 14000 rpm in a microcentrifuge at 4°C for 10 min. The supernatants were used as whole cell extracts. Protein concentration was determined using the Bio-Rad protein determination method (Bio-Rad, Richmond, CA, USA). Equal amounts (50 μg) of proteins were boiled for 5 min and electrophoresed under reducing conditions on 6–12.5% (w/v) polyacrylamide gels. Proteins were electrophoretically transferred to a Hybond-polyvinylidene difluoride transfer membranes (Amersham Life Science, Buckinghamshire, UK), and incubated with the primary antibody, followed by peroxidase-linked secondary antibody. An Amersham ECL

chemiluminescent western system (Amersham) was used to detect secondary probes.

Statistical analysis

Data are expressed as mean \pm s.d. The Student's *t*-test was used to compare differences. Statistical significance was defined when *P* was <0.05.

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Supplementary Information accompanies the paper on the Oncogene website (<http://www.nature.com/onc>).

Establishment of biological and pharmacokinetic assays of telomerase-specific replication-selective adenovirus

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The use of replication-selective tumor-specific viruses represents a novel approach for the treatment of neoplastic disease. We constructed an attenuated adenovirus, telomerase-specific replication-selective adenovirus (TRAD), in which the human telomerase reverse transcriptase promoter element drives the expression of the *E1A* and *E1B* genes linked with an internal ribosome entry site (IRES). Forty-eight hours after TRAD infection at a multiplicity of infection of 1.0, the cell viability of H1299 human lung cancer cells was consistently less than 50% and therefore this procedure could be used as a potency assay to assess the biological activity of TRAD. We also established a quantitative real-time polymerase chain reaction (PCR) analysis with consensus primers for either the adenovirus *E1A* or IRES sequence. The linear ranges of quantitation with *E1A* and IRES primers were 10^3 – 10^8 and 10^2 – 10^8 plaque-forming units/mL in the plasma, respectively. The PCR analysis demonstrated that the levels of *E1A* in normal tissues were more than 10^3 lower than in the tumors of A549 human lung tumor xenografts in *nu/nu* mice at 28 days after intratumoral injection. Our results suggest that the cell-killing assay against H1299 cells and real-time PCR can be used to assess the biological activity and biodistribution of TRAD in clinical trials. (*Cancer Sci* 2008; 99: 385–390)

The emerging fields of functional genomics and functional proteomics provide an expanding repertoire of clinically applicable targeted therapeutics.⁽¹⁾ Replication-selective oncolytic viruses provide a new platform for treatment of a variety of human cancers.^(2,3) Promising clinical trials have shown the antitumor potency and safety of mutant or genetically modified adenoviruses.^(4,5) We constructed previously an adenovirus vector, TRAD, in which the hTERT promoter element drives the expression of the *E1A* and *E1B* genes linked with an IRES. We showed that TRAD caused efficient selective killing of human cancer cells, but not normal cells.⁽⁶⁾ Many studies have demonstrated that the majority of malignant tumors express telomerase activity,⁽⁷⁾ suggesting that TRAD can potentially kill most human cancer cells.

TRAD can replicate and then lyse cancer cells, infect neighboring cancer cells, and subsequently induce oncolysis throughout the whole tumor mass *in vivo*. As preclinical models showed that TRAD could spread into the bloodstream, it is important to monitor carefully the amount of TRAD in the circulation after intratumoral injection of TRAD to avoid serious adverse events due to viremia. Although we used vector-specific primers that detected the p53 open reading frame–adenoviral DNA junction in a phase I clinical trial of a replication-deficient adenoviral vector expressing the wild-type p53 gene (Advexin),⁽⁸⁾ no appropriate method has been established to detect TRAD quantitatively. In addition, there is also a need for a procedure that can evaluate the biological activity of TRAD for clinical application.

In the present study, we characterized a potent antitumor viral agent, TRAD, to establish a biological assay and developed a

single quantitative PCR method that can be used to assess the number of viral genomes present in the plasma as well as tissues.

Materials and Methods

Cells and culture conditions. H1299 (a human non-small-cell lung cancer cell line), H460 (a human large-cell lung cancer cell line), A549 (a human lung adenocarcinoma cell line), LNCap (a human metastatic prostatic carcinoma cell line), MKN28 and MKN45 (human gastric adenocarcinoma cell lines), PC-3 (a human prostate adenocarcinoma cell line), SW620 (a human colorectal carcinoma cell line), and TE8 and T.Tn (human esophagus squamous carcinoma cell lines) were propagated to monolayer cultures in RPMI-1640 supplemented with 10% FBS, and 100 units/mL PG and 100 µg/mL SM. HeLa (a human cervical adenocarcinoma cell line), HepG2 (a human hepatocellular carcinoma cell line), Panc-1 (a human pancreatic epithelioid carcinoma cell line), and 293 (a transformed embryonic kidney cell line) were grown in DMEM containing high glucose (4.5 g/L) (high) with 10% FBS and PG/SM. HT-29 (a human colorectal adenocarcinoma cell line) was grown in McCoy's 5a with 10% FBS and PG/SM. MCF-7 (a human mammary gland adenocarcinoma cell line) was grown in Earle's Minimum Essential Medium with 10% FBS, PG/SM, and 2 mM L-glutamine. OST, SaOS2, and HOS (human osteosarcoma cell lines) were grown in DMEM (high) with 10% FBS and PG/SM. HSC-3 and HSC-4 (human tongue squamous carcinoma cell lines) were obtained from the Health Science Resources Bank (Osaka, Japan) and grown in DMEM (high) with 10% FBS and PG/SM. SCC-4 and SCC-9 (human tongue squamous carcinoma cell lines) were obtained from American Type Culture Collection (ATCC, Rockville, MD, USA) and grown in DMEM containing Nutrient Mixture (Ham's F-12) with 10% FBS, PG/SM, and 400 ng/mL hydrocortisone. U-2OS (a human osteosarcoma cell line) was obtained from ATCC and grown in McCoy's 5a with 10% FBS and PG/SM. NHLF was purchased from Takara Biomedicals (Kyoto, Japan) and cultured in the medium recommended by the manufacturer.

Recombinant adenoviruses. The recombinant replication-selective tumor-specific adenovirus vector TRAD was constructed and

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Abbreviations: ATCC, American Type Culture Collection; DMEM, Dulbecco's modified Eagle's medium; FBS, fetal bovine serum; hTERT, human telomerase reverse transcriptase; ID₅₀, the multiplicity of infection that causes 50% growth inhibition; IRES, internal ribosome entry site; NHLF, normal human lung fibroblasts; MOI, multiplicity of infection; PCR, polymerase chain reaction; PFU, plaque-forming units; PG, penicillin; SM, streptomycin; TRAD, telomerase-specific replication-selective adenovirus; XTT, sodium 3'-[1-(phenylamino)carbonyl]-3,4-tetrazolium]-bis(4-methoxy-6-nitro)benzene sulfonic acid hydrate.

characterized as described previously.^(6,9-11) The virus was purified by CsCl₂ step-gradient ultracentrifugation followed by CsCl₂ linear-gradient ultracentrifugation. The virus particle titer and infectious titer were determined spectrophotometrically and by plaque assay, respectively, in 293 cells.

Cell-viability assay. The XTT assay was carried out to measure cell viability. Cells were plated on 96-well plates at 1×10^5 cells/well 20 h before viral infection. HSC-4, SCC-4, and SCC-9 cells were then infected with TRAD at MOI of 0, 1, 10, and 50 PFU/cell. Other cell lines were infected with TRAD at MOI of 0, 0.1, 1, and 10 PFU/cell. Cell viability was determined at 1, 2, 3, and 5 days after virus infection using Cell Proliferation Kit II (Roche Molecular Biochemicals, Indianapolis, IN, USA) according to the protocol provided by the manufacturer. Using the cell viability data at 3 days after virus infection, we determined the TRAD ID₅₀ of each cell line.

Cell-killing assay. H1299 cells were plated at 5×10^4 cells/well on 24-well plates and infected with TRAD at MOI of 0, 0.01, 0.1, 1, and 10 PFU/cell. Forty-eight hours later, the number of cells in each well was counted. Experiments were carried out in triplicate for each MOI, and cell viability was assessed by the trypan blue dye exclusion assay.

Quantitative real-time PCR assay. Viral DNA from serially diluted viral stocks and tumor cells infected with TRAD were extracted using QIAamp DNA Mini Kit (Qiagen, Valencia, CA, USA), and quantitative real-time PCR assay for either the *E1A* gene or the IRES sequence was carried out using a LightCycler instrument and a LightCycler DNA Master SYBR Green I kit (Roche Molecular Biochemicals). Typical amplification mixes (20 μ L) contained 3 mM MgCl₂, 0.3 μ M of each primer for IRES or 0.5 μ M for *E1A*, and 2 μ L of 10 \times LightCycler FastStart DNA Master SYBR Green I. The sequences of the specific primers used in this experiment were: IRES, 5'-GAT TTT CCA CCA TAT TGC CG-3' and 5'-TTC ACG ACA TTC AAC AGA CC-3'; *E1A*, 5'-CCT GTG TCT AGA GAA TGC AA-3' and 5'-ACA GCT CAA GTC CAA AGG TT-3'. PCR amplifications were carried out in glass capillary tubes. PCR amplification for IRES began with a 10-min denaturation step at 95°C and then 40 cycles of denaturation at 95°C for 10 s, annealing at 60°C for 10 s, and extension at 72°C for 6 s. PCR amplification for *E1A* began with a 10-min denaturation step at 95°C and then 40 cycles of denaturation at 95°C for 10 s, annealing at 58°C for 15 s, and extension at 72°C for 8 s. Data analysis was carried out using LightCycler Software (Roche Molecular Biochemicals).

In vivo human tumor model. A549 human lung cancer cells (5×10^6 cells/mouse) were injected subcutaneously into the flank of 7- to 9-week-old female BALB/c *nu/nu* mice and permitted to grow to approximately 5–6 mm in diameter. At that stage, a 100- μ L solution containing 1×10^8 PFU of TRAD was injected into the tumor. The tumors and organs were harvested 28 and 70 days later and DNA was extracted from each tissue. To compare viral replication in the tumor and other normal organs, quantitative real-time PCR for the *E1A* gene was carried out using a LightCycler instrument. The experimental protocol was approved by the Ethics Review Committee for Animal Experimentation of Okayama University School of Medicine.

Statistical analysis. All data were expressed as mean \pm SD. Differences between groups were examined for statistical significance using Student's *t*-test. A *P*-value less than 0.05 denoted the presence of a statistically significant difference.

Results

In vitro cytopathic efficacy of TRAD in human cancer cell lines derived from different organs. To determine whether TRAD infection induces broad-spectrum selective cell lysis, 23 tumor cell lines derived from 11 different organs (head and neck, lung, esophagus, stomach, colon, liver, pancreas, breast, prostate,

uterus, and bone) were infected with TRAD at various MOI. Previous studies using a real-time reverse transcription-PCR method have demonstrated that these cell lines express detectable levels of hTERT mRNA.^(6,9) Cytotoxicity was then assessed using the XTT cell-viability assay over 5 days after infection. As shown in Figure 1a, TRAD infection induced cell death in all cell lines except T.Tn esophageal cancer cells in a dose-dependent manner. Calculated ID₅₀ values confirmed that all cell lines except T.Tn could be killed efficiently by TRAD at an MOI of less than 25 (Fig. 1b). These results suggest the broad-spectrum antitumor potency of TRAD.

Establishment of a standard assay to assess the biological activity of TRAD. H1299 human lung cancer cells and LNCap human prostate cancer cells were the most sensitive cell lines to TRAD-induced cell death (Fig. 1b). Accordingly, we used H1299 cells to evaluate the biological activity of TRAD. To test whether the selective replication of TRAD translates into selective oncolysis, we compared the cytopathic effects of TRAD in H1299 cells and NHLF at 5 days after infection. The dose-response curve of the relative cell viability in H1299 cells was shifted to the left compared to that in NHLF, suggesting that TRAD killed H1299 cells 10²–10³ more efficiently than NHLF (Fig. 2a).

We next determined the minimal dose of TRAD that could induce more than 50% of cell death in H1299 cells. As shown in Figure 2b, the cell viability of H1299 cells was less than 40% at 48 h after their infection with TRAD at a MOI of 1.0, but was 60% after infection with a MOI of 0.1. We also confirmed that H1299 cells at various passages (5th to 20th after purchase from ATCC) could be killed by TRAD in a similar fashion (data not shown). Therefore, TRAD could be considered biologically active, if TRAD at a MOI of 1 reduces the cell viability of H1299 cells by more than 50% at 48 h after infection. To estimate the utility of this assay, we examined the biological activity of heat-inactivated TRAD. Infection with intact TRAD at a MOI of 10 induced approximately 90% reduction in H1299 cell viability at 48 h after infection, whereas the antitumor activity was completely inhibited when it was preheated at 56°C for 5 or 10 min (Fig. 2c).

Development of quantitative PCR assay to detect copy numbers of TRAD. We used real-time PCR for quantitative detection of TRAD. Oligonucleotide primers were designed to achieve DNA amplification of the adenoviral *E1A* or IRES sequences in the TRAD genome (Fig. 3a). To generate accurate standard curves, TRAD at a known concentration was serially diluted and used as a template for real-time PCR analysis. Detection of IRES and *E1A* genome copies was achieved consistently and reproducibly by the PCR cycle values used. A linear relationship could be obtained between the number of cycles and the log₁₀ dilution when 10²–10⁸ IRES copies and 10³–10⁸ *E1A* copies were assayed. Regression analysis of IRES and *E1A* curves resulted in very high correlation coefficients (0.99 and 1.00, respectively) for these concentration ranges (Fig. 3b). In addition, the dilution of TRAD virus in the plasma did not affect the sensitivity and dynamic ranges of quantification (Fig. 3b), suggesting that this method can be used to detect TRAD in the blood circulation.

In vitro quantification and replication monitoring of TRAD in infected human tumor and normal cells. We next examined the replication ability of TRAD in different cell lines by measuring the relative amounts of IRES and *E1A* copy numbers. LNCap and NHLF cells were harvested at the indicated time points over 5 and 7 days, respectively, after infection with TRAD, and subjected to quantitative real-time PCR analysis using IRES and *E1A* primers. The ratios were normalized by dividing the value of cells obtained at 2 h after viral infection. As shown in Figure 4a, TRAD replicated 10³–10⁴ by 5 days after infection; its replication, however, was attenuated to less than 10³ in normal NHLF cells. We previously reported that TRAD could replicate 10⁵–10⁶ by 3 days after infection in H1299 cells;^(6,10) however, as

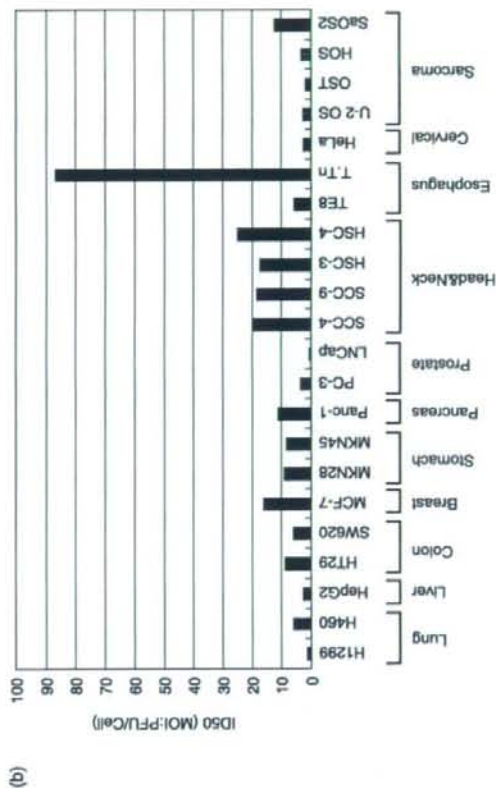
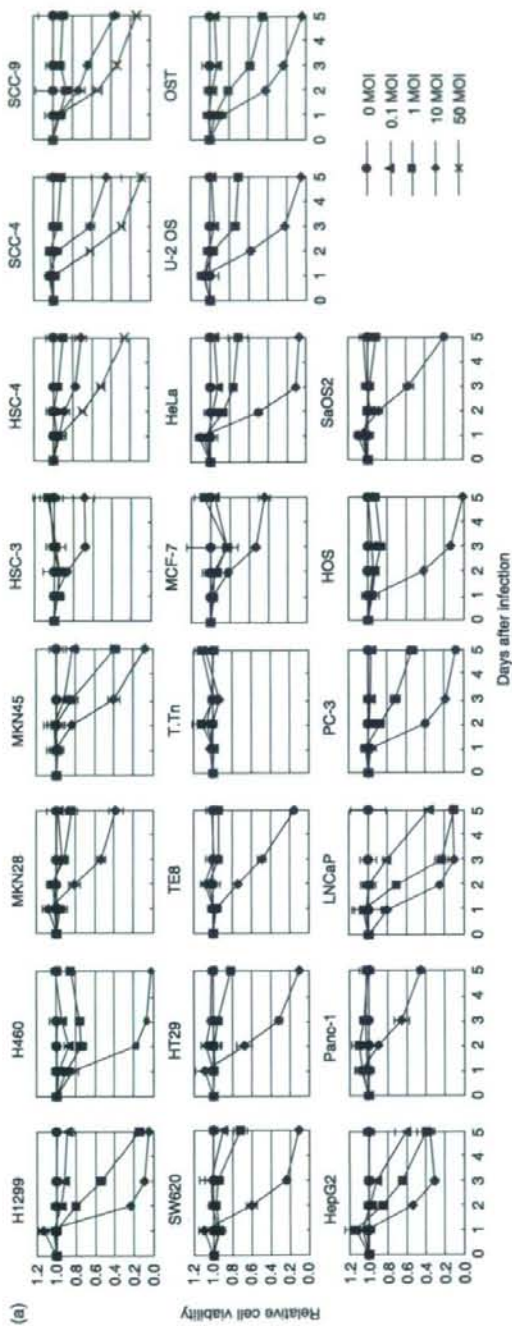


Fig. 1. Oncolytic effects of telomerase-specific replication-selective adenovirus (TRAD) *in vitro* on a variety of human cancer cell lines. (a) Cells were infected with TRAD at indicated multiplicity of infection (MOI) values, and surviving cells were quantitated over 5 days by XTT assay. Data are mean \pm SD. (b) The 50% inhibiting doses of TRAD on cell viability at 3 days after infection were calculated and expressed as ID₅₀ values. PFU, plaque-forming units; XXT, sodium 3-[1-(phenylamino)carbonyl]-3,4-tetrahydro-2H-pyridin-2-one)benzene sulfonic acid hydrate.

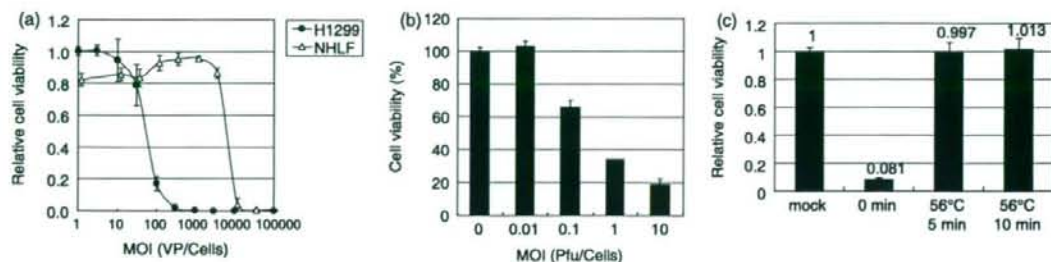


Fig. 2. Antitumor effects of telomerase-specific replication-selective adenovirus (TRAD) on H1299 non-small-cell lung cancer cells *in vitro*. (a) Effects of various concentrations of TRAD on H1299 cancer cells and normal human lung fibroblasts (NHLF) assessed at 5 days after treatment with XTT assay. Results are expressed as the percentage of untreated control. (b) H1299 cells were cultured as monolayers in triplicate in 24-well culture plates, infected with TRAD at the indicated multiplicities of infection (MOI), and assessed for cell viability 48 h after infection. Mock-infected cells were used as a control. (c) H1299 cells were plated on 96-well plates and infected with 10 MOI of TRAD heated at 56°C for 5 or 10 min, or non-treated TRAD. An XTT assay was carried out at 3 days after virus infection. Mock-infected cells were used as a control. Data represent the mean \pm SD of triplicate experiments. PFU, plaque-forming units; XTT, sodium 3'-[1-(phenylaminocarbonyl)-3,4-tetrazolium]-bis(4-methoxy-6-nitro) benzene sulfonic acid hydrate; VP, virus particles.

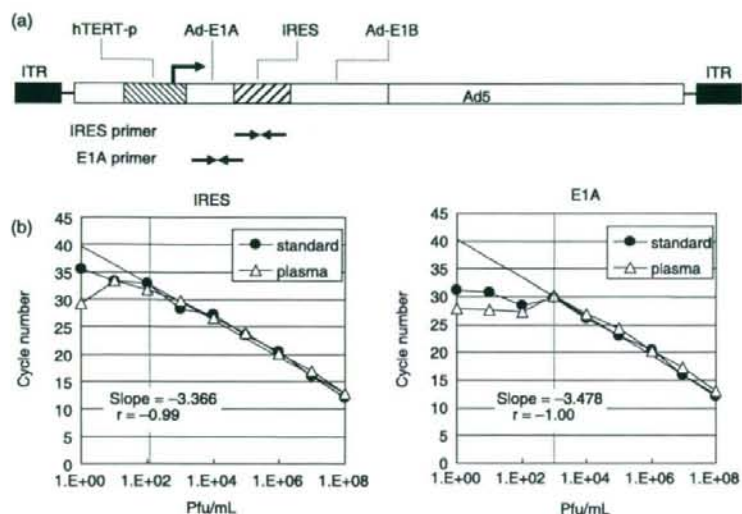


Fig. 3. Detection of normal human lung fibroblasts (TRAD) using quantitative polymerase chain reaction (PCR) assay. (a) Schematic diagram of the DNA structure of TRAD. TRAD contains the human telomerase reverse transcriptase (hTERT) promoter sequence inserted into the adenovirus genome to drive transcription of the *E1A* and *E1B* bicistronic cassette linked by the internal ribosome entry site (IRES) structure. Sites to which PCR primers (IRES and *E1A*) were targeted are indicated. Two primer pairs of IRES and *E1A* were designed to detect the TRAD genome. (b) Standard calibration curves of threshold cycle values and copy numbers are shown using serial dilution of TRAD virus stock. The coefficient of correlation (r^2) and slope are indicated for assays with IRES and *E1A* primers. ITR, inverted terminal repeats; PFU, plaque-forming units.

LNCap cells were more sensitive to TRAD-mediated cytotoxicity than H1299 cells (Fig. 1a), viral replication reached a plateau phase around 10^4 when LNCap cells started to die. Moreover, PCR targeting IRES and *E1A* showed similar replication profiles for TRAD in MCF-7 human breast cancer cells (Fig. 4b). To monitor the long-term viral replication, MCF-7 cells that were less sensitive to the cytopathic effect of the virus were used.

In vivo determination of TRAD genomes in tissue samples after intratumoral injection. To evaluate selective replication of TRAD *in vivo*, we examined mouse tissues, including implanted tumors, for the presence of viral DNA by quantitative real-time PCR, following intratumoral viral injection. Mice with established subcutaneous A549 human lung tumor xenografts received a single intratumoral injection of 1×10^8 PFU of TRAD, and were killed 28 or 70 days after injection. To obtain the sufficient amounts of tumor tissues for analysis, we chose to use A549 cells. Our preliminary experiments demonstrated that intratumoral administration of TRAD suppressed tumor growth significantly compared with mock-treated tumors at 42 days after initiation of treatment ($P < 0.05$); however, the *in vivo* antitumor effect against A549 tumors was less than that against H1299 or LNCap

tumors (data not shown). Although *E1A* DNA was detected in serum and some normal tissues examined (brain, heart, lung, ovary, liver, uterus, kidneys, bladder, colon, and axillary and mesenteric lymph nodes), tumors injected with TRAD contained at least 1000-fold more *E1A* copies (Fig. 5). These results suggest that quantitative real-time PCR allows detection and quantification of the number of TRAD genomes present in tissue samples after intratumoral injection of TRAD *in vivo*.

Discussion

Oncolytic viruses have been developed as anticancer agents based on the advantage of selective killing of tumor cells by controlled replication of the virus in the tumors, resulting in minimal undesired effects on normal cells.⁽⁷⁾ Furthermore, amplified viruses can infect adjacent tumor cells as well as reach distant metastatic tumors through the blood circulation. Although this might be a potential advantage of oncolytic viruses, systemic dissemination of large amounts of virus may induce virus-related symptoms including fever, diarrhea, pneumonia, and hepatitis, eventually leading to death. Therefore, virus shedding and distribution have to be evaluated by appropriate and suitable methods. In addition,

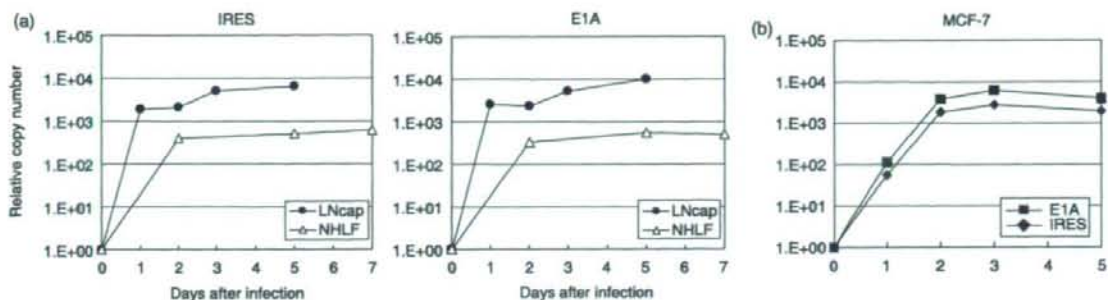


Fig. 4. Quantitative measurement of viral DNA replication in human cancer and normal cells *in vitro* by quantitative polymerase chain reaction (PCR) assay. (a) LNCap human prostate cancer cells and normal human lung fibroblast (NHLF) cells were infected with telomerase-specific replication-selective adenovirus (TRAD) at a multiplicity of infection (MOI) of 1 for 2 h. Following the removal of virus inoculum, cells were further incubated for the indicated periods of time, and then subjected to the real-time quantitative PCR assay. The amounts of viral internal ribosome entry site (IRES) and E1A copy number was defined as the fold increase for each sample relative to that at 2 h (2 h equals 1). (b) MCF-7 human breast cancer cells were infected with TRAD at a MOI of 1 and subjected to the PCR assay at the indicated time points. The relative TRAD DNA levels detected by IRES and E1A primers were plotted.

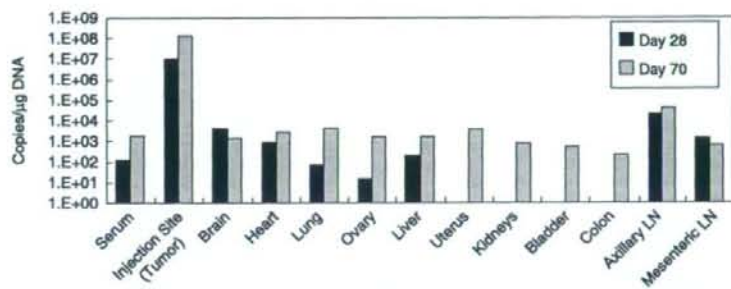


Fig. 5. Spread and replication of telomerase-specific replication-selective adenovirus (TRAD) following intratumoral administration in *nu/nu* mice transplanted with A549 tumor cells. A549 tumor cells were injected subcutaneously into the right flank of mice at 5×10^6 cells/mouse. Mice received intratumoral injection of 1×10^8 plaque-forming units of TRAD when the tumor reached a size of approximately 5–6 mm in diameter. DNA was extracted from the subcutaneous tumor and various tissues of *nu/nu* mice at 28 or 70 days after infection. Viral DNA was detected by quantitative polymerase chain reaction amplification of the adenoviral E1A sequence. The amounts of TRAD genome were defined as viral E1A copy number per μ g DNA. LN, lymph nodes.

to avoid unexpected infectious disease due to viral overdose, we need assays that accurately detect the biological activity of viruses. In the present study, for clinical trials of TRAD, we developed an assay designed to estimate the biological activity of TRAD and to detect the copy number of TRAD in the plasma as well as tissues.

Although telomerase-specific TRAD exhibited a broad cytopathic effect against human cancer cell lines of different tissue origins, a human non-small-cell lung cancer cell line, H1299, was chosen for the biological assay of TRAD. H1299 was one of the most sensitive cell lines to TRAD-mediated cell death ($ID_{50} = 0.94$ MOI) and could be killed efficiently by TRAD infection in a dose-dependent fashion (Fig. 1). Because H1299 cells can be obtained from ATCC, they can be used in clinical laboratories to assess the biological activity of TRAD with a qualified standard protocol. In addition, although adenoviral E1B-55 kDa protein is known to bind to the tumor suppressor p53 protein,⁽¹²⁾ H1299 cells are p53-null and therefore the interaction of E1B-55 kDa with p53, which in turn results in transcriptional modulation, can be ignored in this cell line. Thus, H1299 is considered an appropriate cell line for assessment of TRAD activity in certain preparations. In the present study, we considered TRAD to be active when the viability of H1299 cells was reduced by more than 50% at 48 h after TRAD infection at an MOI of 1. Using this biological assay, we confirmed that heat

treatment of aliquots of TRAD at 56°C for 5 min is sufficient to inactivate its antitumor potential (Fig. 2c). These results advocate the use of the H1299 cell-based cytotoxicity assay as a standard method for quantitative assessment of the biological activity of TRAD in virus stocks for clinical trials.

Various biological methods, such as determination of infectious units in plaque assays, have been used routinely in clinical trials to monitor viral loads in the peripheral circulation.⁽⁸⁾ These methods are useful for evaluating safety because the viral titers directly reflect the infectivity of viruses. However, because the plaque assay consists of labor-intensive and time-consuming steps, real-time monitoring of the biodistribution of the virus might be difficult. Here we described the development of a quantitative real-time PCR assay that can accurately quantify genome copy numbers of TRAD over a large linear range. Using primers targeting TRAD-specific sequences, such as adenoviral E1A and IRES, real-time PCR could accurately detect the number of TRAD genomes in the plasma as well as in the cells (Figs 3,4). The assay showed that TRAD replicated even in NHLF, although the level was much lower than that in tumor cells. It is usually difficult to maintain the normal cells primarily isolated from human tissues such as human hepatocytes in the culture; however, commercially available NHLF could be cultured for several passages, suggesting that NHLF may have some characteristics different from primary isolated normal cells, including

telomerase activity. We also found that the number of viral genomes could be measured in genomic DNA purified from tissues of mice *in vivo* after injection of TRAD into the xenografts (Fig. 5). Although viral DNA could be detected even in normal tissues 70 days after intratumoral injection of TRAD, the absence of infectious virus as assessed by the plaque assay suggests that there are only DNA fragments in tissues. Our preliminary experiments have demonstrated that DNA could be isolated from tumors as small as 5 mm in diameter (data not shown). Therefore, the real-time PCR method with E1A and IRES primers permits rapid and quantitative detection of TRAD DNA in clinical samples.

We have shown recently the antiviral activity of cidofovir against TRAD *in vitro*. Cidofovir is an acyclic nucleoside phosphonate with potent broad-spectrum anti-DNA viral activity and has been approved for the treatment of many types of viruses, including cytomegalovirus and adenovirus.⁽¹³⁾ Although viremia after TRAD administration is extremely rare because of the anti-adenovirus antibodies expected to be present in most patients, a

real-time PCR-based pharmacokinetic assay can allow the early detection of disseminated virus, and thus its use could provide an indication for commencement of cidofovir treatment in clinical trials.

In summary, we have established a fast, reliable, and sensitive assay to assess the biological activity of TRAD *in vitro* and to detect the viral genome in the plasma as well as tissues *in vivo*. A phase I clinical trial of TRAD targeting advanced solid tumors is currently underway in the USA following the approval of the Food and Drug Administration. Such an assay has been used in this ongoing trial and the data will be analyzed in the near future for the assessment of the safety, efficacy, and bio-distribution of TRAD.

Acknowledgments

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Combination of oncolytic adenovirotherapy and Bax gene therapy in human cancer xenografted models. Potential merits and hurdles for combination therapy

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Cancer gene therapy and oncolytic virotherapy have been studied extensively. However, their clinical application is hampered by their weak anticancer activity. We previously constructed a replicating adenovirus (OBP-301, Telomelysin), in which the human telomerase reverse transcriptase (hTERT) promoter drives expression of the adenoviral E1 genes, and causes selective lysis of human cancer cells. We hypothesized that combination adenoviral therapy containing OBP-301 and a nonreplicating adenovirus expressing the proapoptotic Bax gene could overcome the weakness and augment the anticancer efficacy of each modality. Combination treatment resulted in marked Bax protein expression and enhanced efficacy *in vitro* cell viability assay, when compared with either single treatment. However, combination treatment was not as effective in suppressing both subcutaneous and pleural disseminated tumors compared with OBP-301 treatment alone. Further investigation revealed that combination treatment resulted in suppressed E1A protein expression associated with reduced viral replication. Our results suggest that Bax gene therapy in combination with oncolytic adenovirotherapy potentially augments their antitumor activity, but further improvements may be required to maximize the combinatorial effect *in vivo*, for the Bax gene expression to avoid interference with production of the oncolytic virus.

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Key words: gene therapy; oncolytic virus; bax; adenovirus

Oncolytic adenoviruses that can selectively replicate in tumor cells and cause lysis of infected cells have been extensively investigated as novel anticancer agents.^{1,2} Recently, such vectors have been approved for clinical trials.^{3–8} However, preclinical and clinical studies have revealed that the clinical application of these agents is hampered by their weak anticancer activity. Therefore, the development of strategies that maximize their anticancer activity is essential to the success of these agents in treating cancer.

One of the approaches to overcome this weakness is combination therapy of the oncolytic virus with a virus expressing therapeutic genes such as proapoptotic genes to augment the killing effect on cancer cells, which may lead to the future development of an arming oncolytic virus as a cancer therapeutic.

Bax is a strong proapoptotic gene that causes cytochrome C release from mitochondria, activates the caspase pathway and leads to apoptosis.^{9,10} We previously constructed a binary adenoviral vector system (Ad/PGK-GV16 + Ad/GT-Bax: Ad/Bax) for Bax overexpression¹¹ that can transfer the Bax gene to cancer cells *in vitro* and *in vivo* and induce effective apoptosis.^{12,13} However, because this system consists of E1-deleted, replication-defective adenovirus, its cell killing activity is theoretically limited to the transduced cells.

We also constructed a novel oncolytic adenovirus (OBP-301, Telomelysin), in which the human telomerase reverse transcriptase (hTERT) promoter drives expression of the adenoviral E1 genes. OBP-301 thus replicates preferentially in human cancer cells and causes their selective lysis.¹⁴ However, as this virus also does not contain any therapeutic genes, clinical weakness is anticipated.

We hypothesize that Bax gene therapy in combination with an oncolytic adenovirus could augment anticancer efficacy by overcoming the weakness of each modality. The replication-defective

adenovirus with E1 deletion can become replication-competent when cotransduced with an oncolytic adenovirus supplying E1 *in trans*. Thus, the expression of the Bax gene and the viral copies would increase, which facilitates cancer cells to undergo apoptosis, release virus progenies and further spread them within the tumor.

In this study, we evaluated the transgene expression, viral replication and antitumor effect of Bax gene therapy combined with OBP-301 oncolytic adenovirotherapy in human cancer cell lines.

Material and methods

Cell lines and cell culture

Human bronchioloalveolar carcinoma A549 cells were propagated in a monolayer culture in Dulbecco's modified Eagle's medium containing Ham's F-12 nutrient mixture supplemented with 10% fetal calf serum (FCS) and antibiotics. Human gastric cancer MKN45 cells were cultured in RPMI 1640 medium containing 10% FCS and antibiotics. The cells were cultured at 37°C in a humidified incubator containing 5% CO₂.

Recombinant adenoviruses

The recombinant replication-selective, tumor-specific adenovirus vector OBP-301 (Telomelysin), and nonreplicating E1-deleted binary adenovirus vector system expressing the proapoptotic Bax gene (Ad/PGK-GV16 + Ad/GT-Bax: Ad/Bax) were previously constructed and characterized.^{11,14} In a binary vector system, expression of the Bax gene can be induced by transferring Ad/GT-Bax into target cells along with Ad/PGK-GV16 at a ratio of 1:1. All of the viruses were propagated in a package containing 293 cells and purified by CsCl₂ step gradient ultracentrifugation followed by CsCl₂ linear gradient ultracentrifugation. Determination of virus particle titer and infectious titer (plaque-forming unit: PFU) was accomplished spectrophotometrically by the method of Maizel *et al.*¹⁵ and by the method of Kanegae *et al.*,¹⁶ respectively. The particles: plaque ratios were between 23:1 and 35:1.

Cell viability assay

Cells were plated in 96-well plates at a density of 1,000 cells/well and infected with OBP-301 alone, Ad/Bax alone, or their combination 15 hr later. Cell viability was assessed using a colorimetric XTT assay with the Cell Proliferation Kit II (Roche Molecular Biochemicals, Indianapolis, IN) according to the manufacturer's protocol. We determined the combination index (CI) at

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different multiplicities of infection (MOIs) in A549 cells with CalcuSyn software (BioSoft, Cambridge, UK).

Cell cycle analysis

A549 and MKN45 cells were seeded on 6-well plates at 1×10^5 cells/well and were infected with OBP-301 at an MOI of 1 PFU/cell, Ad/Bax at an MOI of 5 or 50 PFU/cell, or OBP-301 at an MOI of 1 PFU/cell in combination with Ad/Bax at an MOI of 5 PFU/cell 15 hr later. At 24 and 96 hr after infection, adherent cells were harvested with trypsin-EDTA and nonadherent cells were also harvested. All cells were washed with PBS, fixed and permeated with 70% ice-cold ethanol at 4°C for 12 h. Cells were centrifuged at 1,200 rpm, resuspended in freshly prepared propidium iodide (PI) staining buffer (0.1% Triton X-100, 200 µg/ml RNase, and 50 µg/ml PI in PBS), and incubated for 30 min at 4°C in the dark. Single color fluorescent flow cytometry was performed with a FACS caliber flow cytometer (Becton Dickinson, San Jose, CA). The histograms were analyzed with FlowJO software (Tree Star, Ashland, OR).

Western blot analysis

A549 cells were seeded and infected with OBP-301 alone, Ad/Bax alone, or their combination at an MOI of 1 PFU/cell 15 hr later. The cells were incubated at 37°C and harvested for Western blot analysis at 72 and 120 hr. The primary antibodies against Bax (Santa Cruz Biotechnology, Santa Cruz, CA), caspase 3, E1A (BD Biosciences, San Jose, CA) and β -actin (Sigma, St. Louis, MO) and peroxidase-linked secondary antibody (Amersham, Arlington Heights IL) were used. Cells were washed twice in cold PBS, collected and lysed in lysis buffer [10 mM Tris (pH 7.5), 150 mM NaCl, 50 mM NaF, 1 mM EDTA, 10% glycerol, 0.5% NP40] containing proteinase inhibitors (0.1 mM phenylmethylsulfonyl fluoride, 0.5 mM Na3VO4). After 20 min on ice, the lysates were spun at 14,000 rpm at 4°C for 10 min. The supernatants were used as whole cell extracts. Protein concentration was determined using the Bio-Rad protein determination method (Bio-Rad, Richmond, CA). Equal amounts of proteins were boiled for 5 min and electrophoresed under reducing conditions on 4–12% (w/v) polyacrylamide gels. Proteins were electrophoretically transferred to Hybond polyvinylidene difluoride transfer membranes (Amersham Life Science, Buckinghamshire, UK) and incubated with the primary followed by peroxidase-linked secondary antibody. An ECL chemiluminescent Western blot system (Amersham) was used to detect secondary probes.

Animal experiments

A549 xenografts were established in 6-week-old female nude mice (Balb/c nu/nu) through subcutaneous inoculation of 2×10^6 A549 cells into the dorsal flank. Once each tumor reached a diameter of ~3–9 mm, the mice were randomly assigned into 4 groups and a 50-µl solution containing OBP-301, Ad/Bax, or their combination, at a dose of 1×10^7 PFU/body or PBS was injected into the tumor on 3 consecutive days. Tumors were measured 2–3 times a week and tumor volume was calculated using the equation $a \times b^2 \times 0.5$, in which a and b are the largest and smallest diameters, respectively. Animals were killed when their tumor reached a diameter of ~15 mm. To develop pleural disseminated xenografts, mice were inoculated with 2×10^6 A549 cells into the pleural cavity through a 27-gauge needle. Also, to assess the efficiency of adenoviral gene transfer into the pleural disseminated tumors, 24 hr after tumor injection, 100 µl of 1.0×10^7 PFU of OBP-301, Ad/Bax, both of them, or PBS was injected into the thoracic space by the same technique. The procedure was repeated over 3 consecutive days. Three weeks after cell inoculation, the mice were killed and their thoracic spaces were examined macroscopically for any growths. Tumors in the thoracic spaces were removed and weighed. The experimental protocol was approved by the Ethics Review Committee for Animal Experimentation of Okayama University School of Medicine, Okayama, Japan. The tumor growth

of each group was statistically analyzed by Student *t*-test and a *p*-value of ≤ 0.05 was considered significant.

Viral replication assay in vivo

A549 cells were injected into the dorsal flank of nude mice. When the tumor reached a size of ~5 mm, 1×10^7 PFU of OBP-301, OBP-301 + Ad/Bax, OBP-301 + Ad/LacZ, or Ad/Bax was injected to the tumors ($n = 5$, each group). Tumors were harvested 7 days after viral injection, immediately frozen and milled in PBS by using Micro Smash MS-100 (Tomy Digital Biology, Tokyo, Japan). The virus was then extracted by freezing and thawing 3 times and cell debris was spun down. The supernatant was collected and subjected to titer assay for viral PFU. The final result was expressed as a total PFU from 1 whole tumor.

Quantitative real-time polymerase chain reaction

A549 cells were seeded on 25-cm² flasks at 5×10^5 cells 15 hr before infection. Cells were infected with OBP-301 alone or in combination with Ad/Bax or Ad/LacZ 15 hr later. The cells were incubated at 37°C, trypsinized and harvested for replication analysis at 48 hr. DNA purification was performed using a QIAmp DNA mini kit (Qiagen, Valencia, CA). The E1A DNA copy number was determined by quantitative real time polymerase chain reaction (PCR), using a LightCycler instrument and LightCycler-DNA Master SYBR Green I (Roche Diagnostics).

Results

Combination of oncolytic virotherapy and Bax gene therapy enhanced cell killing in vitro

To compare the efficacy of the combined use of oncolytic virotherapy with Bax gene therapy *in vitro*, A549 and MKN45 human cancer cells were treated with OBP-301 alone, Ad/Bax alone, or the combination. Based on each optimal condition, A549 was infected with OBP-301 at 1 MOI and Ad/Bax at 10 MOI, and MKN45 was infected at 10 MOI in both viruses. In both cell lines, Ad/Bax treatment showed only minimal suppression of cell viability and OBP-301 treatment showed a modest and delayed oncolytic effect, whereas combination treatment with OBP-301 and Ad/Bax induced rapid and massive cell death, and almost complete cell killing (Fig. 1).

To evaluate the synergistic effect of the combined oncolytic virotherapy and Bax gene therapy, the CI was determined at different MOIs (Table I) with CalcuSyn software (BioSoft, Cambridge, UK). We found that in A549 cells, the combination of OBP-301 with Ad/Bax led to a strong synergism (CI < 0.3).

Bax gene therapy in combination with oncolytic virotherapy increased the production of Bax protein in vitro and induction of apoptosis

To examine Bax protein expression in cells infected with the replication-deficient Ad/Bax in combination with the replicating OBP-301, A549 cells were infected with OBP-301 alone, Ad/Bax alone, or OBP-301 in combination with Ad/Bax or Ad/LacZ and then harvested on day 3 or 5. Cells treated with OBP-301 or the combination of OBP-301 with Ad/LacZ resulted in minimal Bax protein expression, whereas treatment with Ad/Bax and the combination treatment with OBP-301 plus Ad/Bax showed higher expression of Bax protein. Densitometric measurement documented increments of Bax expression by combination of Ad/Bax with OBP-301 (Fig. 2). In addition, treatment with OBP-301 plus Ad/Bax resulted in caspase-3 activation, while treatment with OBP-301 only or OBP-301 plus control virus Ad/LacZ did not. This means that the combination with Ad/Bax activates the apoptotic cascade in cancer cells cotreated with oncolytic virus.

Bax gene therapy in combination with oncolytic virotherapy increased apoptotic cells in vitro

To quantify the apoptotic effects of Ad/Bax treatment in combination with OBP-301 treatment, sub-G1/G0 fractions were

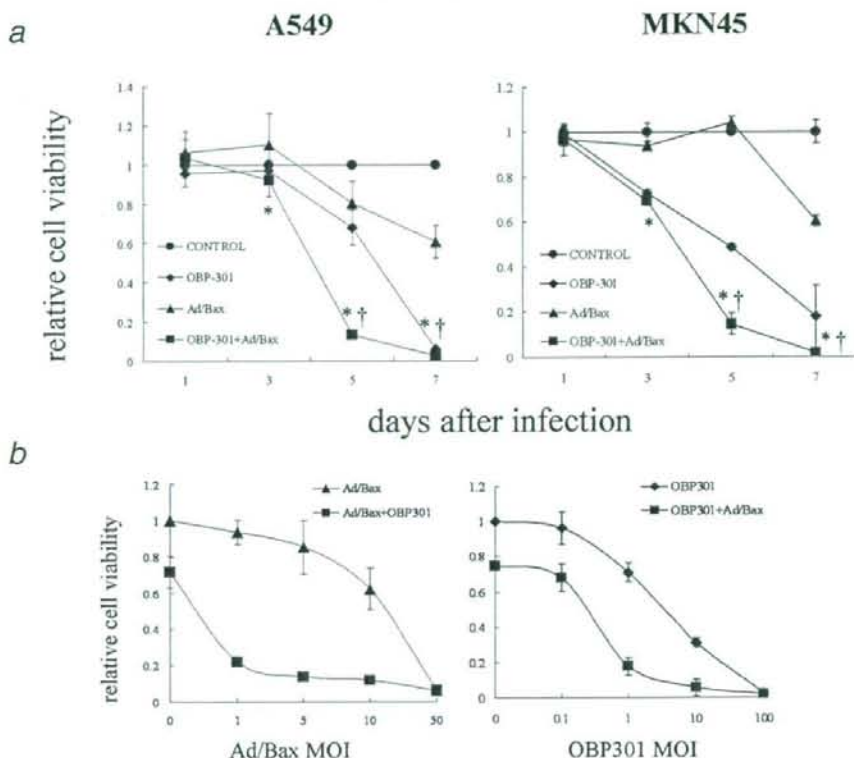


FIGURE 1 – Cytotoxicity effects of OBP-301 combined with Ad/Bax in human cancer cell lines. (a) The indicated cancer cells were infected with OBP-301 alone, Ad/Bax alone, or in combination at the indicated MOI. Cell viability was assessed by XTT assay over 7 days. Bars, standard deviation (SD). *, $p < 0.05$, OBP-301 + Ad/Bax versus Ad/Bax alone; †, $p < 0.05$, OBP-301 + Ad/Bax versus OBP-301 alone. (b) A549 cells were infected with 1 MOI of OBP-301 and Ad/Bax at the indicated MOI (left), or 10 MOI of Ad/Bax and OBP-301 at the indicated MOI (right). Cell viability was assessed by XTT assay at 5 days after infection. Bars, SD.

TABLE 1 – COMBINATION INDEX (CI) OF OBP-301 AND Ad/Bax

Ad/Bax (MOI)	CI
1	0.122
5	0.111
10	0.132
50	0.217

OBP301 (MOI)	CI
0.1	1.054
1	0.144
10	0.197
100	0.592

determined by flow cytometry. A549 cells were infected with OBP-301, Ad/Bax, or OBP-301 plus Ad/Bax, and then subjected to the analysis at day 4. Although treatment with OBP-301 or Ad/Bax resulted in only background levels of apoptotic cells similar to that of mock infection, treatment with Ad/Bax plus OBP-301 markedly increased the apoptotic cells (Fig. 3).

Combination with Bax gene therapy did not augment antitumor activity of OBP-301-mediated oncolytic virotherapy in vivo

Based on the *in vitro* favorable combination effect of Ad/Bax and OBP-301, the antitumor effect on established tumors was further assessed. The A549 tumors were established in the dorsal

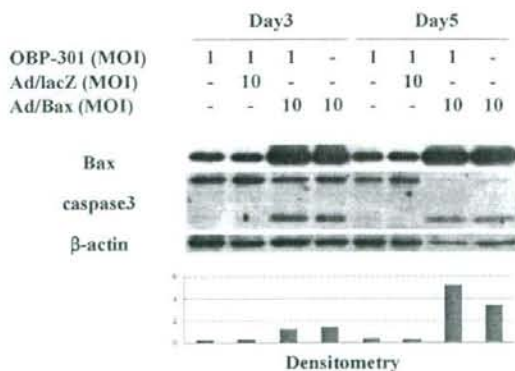


FIGURE 2 – Expression of Bax and caspase 3 in cells infected with Ad/Bax in combination with OBP-301. A549 cells were infected at 1 MOI of OBP-301 alone, 10 MOI of Ad/Bax alone, or 1 MOI of OBP-301 in combination with 10 MOI of Ad/Bax or Ad/lacZ, and then harvested at day 3 or 5. Lysates were subjected to immunoblot analysis with antibodies recognizing Bax, caspase 3, or β -actin. The lower panel shows the intensity of each band of Bax determined by densitometric scanning using NIH image software and normalized by dividing the actin signal.

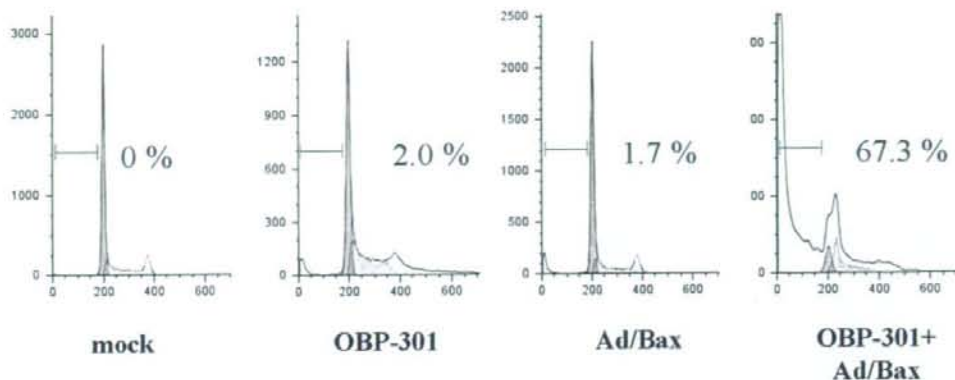


FIGURE 3 – Cell cycle analysis in A549 cells infected with OBP-301 in combination with Ad/Bax. Indicated cancer cells were infected with mock solution, 1 MOI of OBP-301, 10 MOI of Ad/Bax, or 1 MOI of OBP-301 plus 10 MOI of Ad/Bax, and harvested at day 4; cell cycle analysis using flow cytometry was then performed. The histograms were analyzed with FlowJO software.

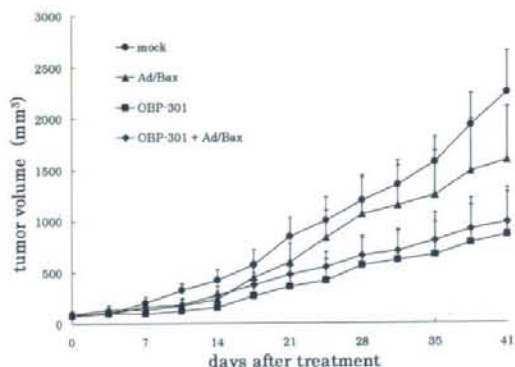


FIGURE 4 – Tumor xenograft size after infection with OBP-301, Ad/Bax, or their combination. A549 xenografts were established in nude mice (Balb/c nu/nu) through subcutaneous inoculation of 2×10^6 A549 cells into the dorsal flank. When the tumor reached a diameter of ~ 5 mm, the virus solution was injected into the tumor at a dose of 1×10^7 PFU/body on 3 consecutive days. Tumor volume was calculated using the equation $a \times b^2 \times 0.5$, in which a and b are the largest and smallest diameters, respectively. Bars, standard error (SE). *, $p < 0.05$, compared with mock-treated tumors.

flank of nude mice and the virus was injected into the tumor at a dose of 1×10^7 PFU/body on 3 consecutive days. Tumor size was measured 2–3 times a week and followed for up to 41 days. Treatment with Ad/Bax alone did not suppress tumor growth significantly when compared with mock treatment because of a suboptimal dose, whereas treatments with OBP-301 and OBP-301 plus Ad/Bax significantly suppressed tumor growth when compared with other controls ($p < 0.05$; Fig. 4). However, the effect of adding Ad/Bax to OBP-301 treatment was less than expected, and the difference was not significant.

We also examined the antitumor effects of this combination treatment in mice with pleurally disseminated A549 tumors. The mice were injected with 100 μ l of 1.0×10^7 PFU of OBP-301, Ad/Bax, or both into the thoracic space on 3 consecutive days. Three weeks after cell inoculation, the mice were killed and their thoracic spaces were examined macroscopically for any growths. To quantitate the reduction of tumor burden in the thoracic spaces, the tumors were removed and weighed. Although treatment with OBP-301 alone or Ad/Bax in combination with

OBP-301 significantly suppressed tumor weight, there was no significant difference between them (Figs. 5a and 5b). These results suggest that Bax gene therapy does not augment oncolytic virotherapy *in vivo*.

Combination treatment resulted in suppressed E1A protein expression due to reduced viral replication *in vitro*

To explore the underlying mechanism by which oncolytic virotherapy did not work cooperatively with proapoptotic Bax gene therapy, E1A protein expression was examined by Western blot analysis. A549 cells were treated with each virus singly or in combination and then subjected to the analysis at days 3 and 5. Treatment with OBP-301 alone and in combination treatment with Ad/lacZ demonstrated increased E1A protein expression, meaning efficient viral replication. Treatment with Ad/Bax alone showed no E1A protein expression because of the E1-deficient replication incompetent virus. Of note, treatment with OBP-301 plus Ad/Bax showed suppressed expression of E1A protein when compared with treatment with OBP-301 alone, meaning that the combination with Ad/Bax may interfere with viral replication (Fig. 6).

To further document the effect of Bax expression on the process of viral propagation, the replication and release of viral progenies were measured by quantitative real time PCR. A549 cells were infected with viruses and the supernatants of the cell groups were collected. DNA was extracted from them and subjected to the assay. We found suppressed viral copy numbers in cells treated with the combinations with Ad/Bax and Ad/lacZ (Fig. 7). The inhibition of E1A copy numbers in a group of Ad/Bax combination was about 10 times of inhibition in a group of Ad/lacZ combination. Our results suggest that Bax gene therapy may interfere with viral production and thus be not conducive to oncolytic virotherapy. To further examine viral replication *in vivo*, the amounts of infectious viruses produced in the tumors were analyzed. Titers were very low for tumors treated with Ad/Bax because of a lack of replication in the absence of E1 gene, while tumors treated with OBP-301 produced higher amount of infectious virus than those treated with Ad/Bax alone, suggesting viral replication in tumors. However, because of the large variation in data from each group, there was no significant difference between treatment groups (data not shown).

Discussion

Replication-competent oncolytic adenovirus has shown a remarkable safety profile but efficacy for cancer therapy continues to be a major obstacle. To eliminate cancer cells, the virus also must spread throughout the bulk of the tumor and induce cell

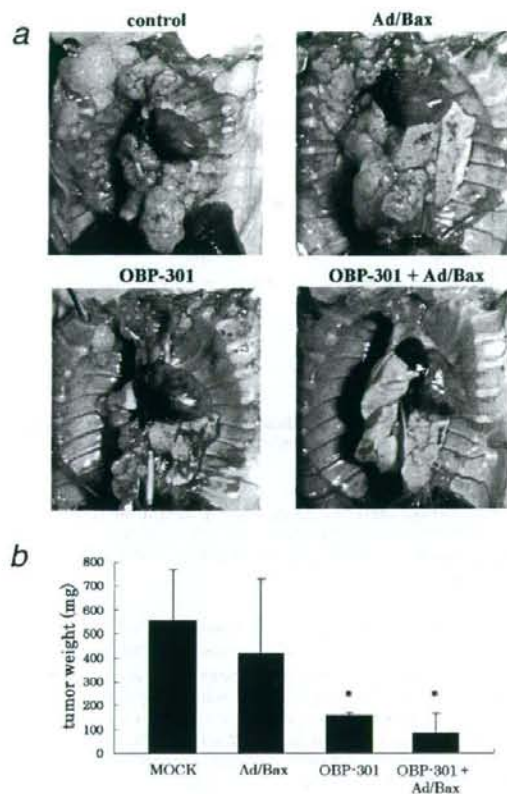


FIGURE 5 – Antitumor effects of OBP-301 combined with Ad/Bax in mice with A549 pleural dissemination. Mice were inoculated with 2×10^6 A549 cells into the pleural cavity, and 24 hr after tumor injection, $100 \mu\text{l}$ of 1.0×10^7 PFU of OBP-301, Ad/Bax, both of them, or PBS was injected into the thoracic space. The procedure was repeated over 3 consecutive days. (a) Three weeks after cell inoculation, mice were killed and their thoracic spaces were examined macroscopically for any growths. (b) Tumors in the thoracic spaces were removed and weighed. Bars, SD. *, $p < 0.05$, compared with mock-treated tumors.

death. Even with replicating adenoviral systems, the necessity to infect all cancer cells remains a major challenge.

In this study, we hypothesized that the oncolytic adenovirotherapy combined with Bax gene therapy could enhance the oncolytic potency. As expected, the combination treatment resulted in Bax overexpression, induced early apoptosis and enhanced efficacy in the cell viability assay *in vitro*. However, disappointingly, combination treatment did not result in further reductions in flank tumor size and pleural disseminated tumor weight, which was associated with suppressed E1A protein expression and reduced viral replication of OBP-301.

The antitumor effect of the oncolytic virus depends on the cytopathic effect intrinsic to adenovirus replication and dissemination throughout the tumor mass. Because viral infection of the tumor bulk depends on cell-to-cell viral spread, there may be a race between rapid tumor growth and viral spread. Therefore, to improve the efficacy of oncolytic virotherapy, the accelerated viral release and induction of cell death must be necessary. A previous study has shown that loss of E1b-19kD function in a replicating adenovirus enhances early viral release, leading to accelerated cell-to-cell viral spread.¹⁷ In turn, Chiou and White reported that inhibition of apoptosis could severely attenuate viral release.¹⁸ Taken together, this suggests that a combination of proapoptotic

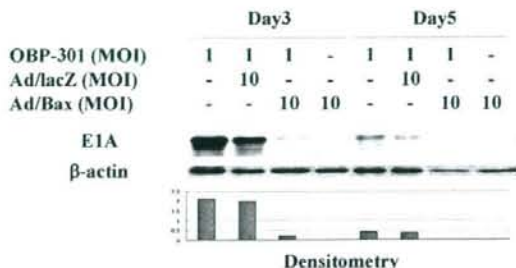


FIGURE 6 – Expression of E1A in cells infected with Ad/Bax in combination with OBP-301. A549 cells were infected at 1 MOI of OBP-301 alone, 10 MOI of Ad/Bax alone, or 1 MOI of OBP-301 in combination with 10 MOI of Ad/Bax or Ad/lacZ, and then harvested at day 3 or 5. Lysates were subjected to immunoblot analysis with antibodies recognizing E1A or β -actin. The lower panel shows the intensity of each band of Bax determined by densitometric scanning using NIH image software and normalized by dividing the actin signal.

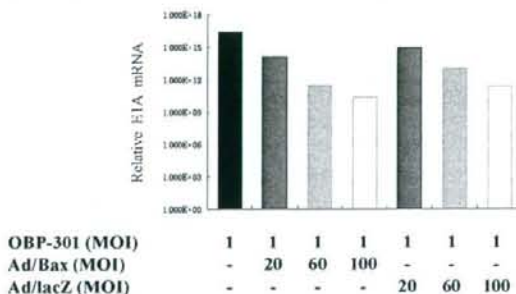


FIGURE 7 – Assessment of viral DNA replication in cells infected with Ad/Bax in combination with OBP-301. A549 cells were seeded on 25-cm^2 flasks at 5×10^5 cells 15 hr before infection. Cells were infected with OBP-301 alone, Ad/Bax alone, or their combination at an MOI of 1 PFU/cell 15 hr later. The cells were harvested at 48 hr and then subjected to real-time quantitative PCR assay. The amounts of viral E1A copy number are defined as the fold increase for each sample relative to that at 2 hr (2 hr equals 1).

gene therapy and oncolytic virotherapy may improve the antitumor efficacy by enhancing cell-to-cell spread of the virus. The other rationale of the combination of proapoptotic gene therapy is to cope with the cancer resistance to the oncolytic virotherapy. Because heterogeneity in cancer cell populations drives the development of resistance, the approach of killing cells by multiple nonoverlapping mechanisms could be a solution. Although an oncolytic virus can kill cells by apoptosis-independent mechanisms such as direct cell lysis and necrosis, cells treated with combination therapy showed a pattern of apoptosis evidenced by analysis of the cell cycle. However, because induction of apoptosis and premature cell lysis may potentially compromise viral yield, reduced viral production could be a concern in using Bax-expressing virus in combination.

In the *in vitro* cell viability assay, the combination of Bax-expressing adenovirus with a replicating adenovirus leads to increased apoptosis at an earlier phase as theoretically anticipated. In turn, we have demonstrated a reduced viral yield in A549 cells. The early apoptotic cell death induced by Bax overexpression may limit an increase of virus and disturb viral replication. Our results are consistent with those of Lambright *et al.*,¹⁹ who found that the addition of ganciclovir therapy to a replicating vector containing the HSVtk suicide gene did not augment efficacy, despite the enhanced production of the transgene. Viral replication and timely cell-death induction thus need to be well coordinated, and forced

induction of early apoptosis could be a cause of losing the combinative effect. Our data suggest that impaired viral replication may offset the benefits due to enhanced transgene spread *in vivo*.

In this study, we have shown the discrepancy between *in vitro* and *in vivo* findings, but this would not result solely from the reduced viral replication. Although viruses can rapidly and evenly spread in cell culture monolayer, this would not be expected in a solid tumor. In a tumor mass, it is very difficult to distribute viral progenies to even the majority of tumor cells. Because a cancer-targeting oncolytic virus is designed to selectively replicate in cancer cells, the normal interstitial cells would be obstacles. The virus would also face other obstacles for viral distribution, including tight intercellular space and continuous drainage by the circulatory and lymphatic systems. In this circumstance, the intratumoral dispersion is confined to one part of the tumor, and thus the requirement of double infection in a complementary strategy curtails the efficacy at low multiplicities of infection.

Although our data suggest that Bax gene therapy in combination with oncolytic adenovirotherapy is not likely to be therapeutically beneficial, the concept of "armed" replicating adenovirotherapy still has potential merit. Overexpression of the adenoviral death pro-

tein, expression of TRAIL and deletion of the E1B-19kD gene have all been shown to improve viral spread and efficacy of replicating adenoviruses *in vitro* and *in vivo*.^{5,17,20-25} Various suicide genes or cytokines have also been expressed with replication-competent vectors to improve cell killing.²⁶⁻³⁰ One important factor to keep in mind is to enhance antitumor effect without inhibiting the proliferative capacity of viruses.

In summary, oncolytic adenovirotherapy in combination with Bax gene therapy resulted in augmented cell killing *in vitro*. However, in a xenografted tumor model, oncolytic efficacy was not improved; this was associated with suppressed E1A protein expression and reduced viral replication. Bax gene therapy may possibly interfere with viral production and thus be not conducive to oncolytic virotherapy.

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SHORT COMMUNICATION

Autophagy-inducing agents augment the antitumor effect of telomerase-selective oncolytic adenovirus OBP-405 on glioblastoma cells

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Oncolytic adenoviruses are a promising tool in cancer therapy. In this study, we characterized the role of autophagy in oncolytic adenovirus-induced therapeutic effects. OBP-405, an oncolytic adenovirus regulated by the human telomerase reverse transcriptase promoter (hTERT-Ad, OBP-301) with a tropism modification (RGD) exhibited a strong antitumor effect on glioblastoma cells. When autophagy was inhibited pharmacologically, the cytotoxicity of OBP-405 was attenuated. In addition, autophagy-deficient Atg5^{-/-} mouse embryonic fibroblasts (MEFs) were less sensitive than wild-type MEFs to OBP-405. These findings indicate that OBP-405-induced autophagy is a cell killing effect. Moreover, autophagy-inducing therapies

(temozolomide and rapamycin) synergistically sensitized tumor cells to OBP-405 by stimulating the autophagic pathway without altering OBP-405 replication. Mice harboring intracranial tumors treated with OBP-405 and temozolomide survived significantly longer than those treated with temozolomide alone, and mice treated with OBP-405 and the rapamycin analog RAD001 survived significantly longer than those treated with RAD001 alone. The observation that autophagy inducers increase OBP-405 antitumor activity suggests a novel strategy for treating patients with glioblastoma.

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Oncolytic adenoviruses have recently been developed as a novel cancer therapy.^{1,2} We have shown that the oncolytic adenovirus regulated by the human telomerase reverse transcriptase promoter (hTERT-Ad) induces nonapoptotic autophagy in glioblastoma cells.³ However, the molecular machinery underlying autophagy-induced cell death remains unclear.^{4–6} Furthermore, the processes determining whether autophagy in cancer cells causes death or acts as a protective mechanism activated during cellular distress are unknown. In this study, we elucidated the therapeutic role of autophagy in hTERT-Ad infection.

The infection efficiency of adenoviral constructs, which are derived from human adenovirus serotype 5, varies widely depending on the cellular expression of the coxsackievirus and adenovirus receptor (CAR).⁷ Modification of the adenoviral fiber protein is one approach to overcoming the limitation imposed by this depen-

dence on CAR.⁸ A recent study demonstrated the activity of OBP-405, which is an hTERT-Ad with a tropism modification (RGD): OBP-405 had a profound oncolytic effect *in vitro* and *in vivo* on lung and colon cancer cells regardless of the CAR expression level.⁹ In addition, for cells with low CAR expression levels, OBP-405 had a higher rate of infection than did OBP-301, an hTERT-Ad that expresses the *E1A* gene under the control of a 455-bp hTERT promoter.⁹ The hTERT-Ad that we used previously has a 255-bp hTERT promoter.³

Good manufacturing practice OBP-301 (Telomelysin) and OBP-405 (Telomelysin-RGD) were used in the current study. We first determined the expression levels of CAR and integrins ($\alpha v \beta 3$ and $\alpha v \beta 5$) on the cell surface of glioblastoma cells. To predict the response of human glioblastoma to OBP-301 or OBP-405, the established cell lines U87-MG, U251-MG, D54, and U373-MG and primary MDC-01 cells isolated from glioblastoma tissue were used. The U87-MG, D54 and MDC-01 cells expressed the lowest levels of CAR, whereas the U251-MG cells expressed the highest level of CAR (Figure 1a). In contrast, $\alpha v \beta 3$ and $\alpha v \beta 5$ integrins were highly expressed in each of these glioblastoma cells. Cells that expressed low levels of CAR were resistant to OBP-301 infection, whereas OBP-405 effectively decreased viabi-

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