Dual E1A oncolytic adenovirus: targeting tumor heterogeneity with two independent cancer-specific promoter elements, DF3/MUC1 and hTERT

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The therapeutic utility of oncolytic adenoviruses controlled by a single, tumor-specific regulatory element may be limited by the intra- and inter-tumoral heterogeneity that characterizes many cancers. To address this issue, we constructed an oncolytic adenovirus that uses two distinct tumor-specific promoters (DF3/Muc1 and hTERT) to drive separate E1A expression cassettes, in combination with deletion of the viral E1B region, which confers additional tumor selectivity and increased oncolytic activity. The resultant virus, Adeno-DF3-E1A/hTERT-E1A, induced higher levels of E1A oncoprotein, enhanced oncolysis and an earlier and higher apoptotic index in infected tumor cells than following infection with Adeno-hTERT-E1A, which harbors a single hTERT promoter-driven E1A cassette. In isolated U251 human gliosarcoma cell holoclones (putative cancer stem cells), where DF3/Muc1 expression is substantially enriched and hTERT expression is decreased compared with the parental U251 cell population, E1A production and oncolysis were strongly decreased following infection with Adeno-hTERT-E1A, but not Adeno-DF3-E1A/hTERT-E1A. The strong oncolytic activity of Adeno-DF3-E1A/hTERT-E1A translated into superior anti-tumor activity over Adeno-hTERT-E1A in vivo in a U251 solid tumor xenograft model, where hTERT levels were >90% suppressed and the DF3/Muc1 to hTERT expression ratio was substantially increased compared with cultured U251 cells. The enhanced anti-tumor activity of the dual-targeted Adeno-DF3-E1A/hTERT-E1A was achieved despite premature viral host cell death and decreased production of functional viral progeny, which limited tumor cell spread of the viral infection. These findings highlight the therapeutic benefit of targeting oncolytic viruses to heterogeneous tumor cell populations.

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Introduction

Replication-conditional adenoviruses engineered for cancer specificity have shown promising stand-alone anti-tumor activity, as well as potential for therapeutic gene delivery. Various strategies have been used to improve tumor-specific adeno viral activity while reducing toxicity to non-cancerous host tissues. For example, cancer-specific promoter elements can be used to control expression of the E1A oncoprotein, which is critical for the transcriptional activation and replication of adenoviruses. However, even with these improvements, such tumor-targeted adenoviruses have had limited success in the clinic.

The hTERT promoter displays high activity in a majority of human cancers, but not in most host tissues and is considered as an ideal tumor-specific regulator for oncolytic adenoviruses. Previously, we characterized an hTERT promoter fragment, which in the context of the replication-conditional adenovirus Adeno-hTERT-E1A recapitulated high telomerase promoter-based E1A expression and viral activity in cancer cells, but not in primary human hepatocytes. Conceivably, the anti-tumor activity of oncolytic adenoviruses, such as Adeno-hTERT-E1A, might be increased by introducing a second E1A cassette under the control of an independent, cancer-specific promoter element. One such element is derived from the MUC1 gene, which codes for DF3 antigen. DF3/Muc1 is a mucin-like glycoprotein that is overexpressed in ~75% of all human solid tumors, including many late-stage cancers of the pancreas, prostate, breast, ovaries and lungs. DF3/Muc1 functions as an onco gene, is a determinant of resistance to Herceptin, regulates intracellular oxidant levels and apoptosis in response to oxidative stress, and has a role in cellular adhesion, invasion and metastasis. DF3/Muc1 is also associated with tumor-forming stem/progenitor cell-like side populations in MCF-7 breast cancer cells and regulation of human pluripotent stem cell growth.

Presently, we investigated the use of two separate tumor-specific promoter elements to regulate adenoviral
E1A production, and hence viral replication, in a cancer-specific manner. The goal of this strategy was to increase viral host range to encompass a broad array of human tumors, including tumors with cell subpopulations having large differences in DF3/Muc1 and hTERT activity. Tumor cells infected with the dual E1A cassette virus described here, Adeno-DF3-E1A/hTERT-E1A, are shown to express higher levels of E1A oncoprotein and undergo enhanced lysis with an earlier and higher apoptotic index than tumor cells infected with the corresponding single E1A cassette virus, Adeno-hTERT-E1A. However, the increased apoptosis stimulated by Adeno-DF3-E1A/hTERT-E1A is accompanied by decreases in viral release, viral spread and activity as a helper virus compared with Adeno-hTERT-E1A. Nevertheless, the increased oncolytic activity of Adeno-DF3-E1A/hTERT-E1A translated into greater anti-tumor activity in a human gliosarcoma tumor xenograft model where putative cancer stem cells (holoclones—clonally isolated cells that have cancer stem cell-like properties and grow with a holoclone morphology)19 show elevated expression of DF3/Muc1 compared with hTERT. Thus, the dual E1A cassette adenovirus displays therapeutic potential as a stand-alone anticancer agent against heterogeneous populations of tumor cells, albeit with decreased ability to replicate and spread an adenoviral infection.

Materials and methods

Cell lines and reagents

Cyclophosphamide (CPA) was purchased from Sigma Chemical (St Louis, MO). Fetal bovine serum (FBS) and Roswell Park Memorial Institute (RPMI) culture medium was purchased from Invitrogen (Frederick, MD). Human tumor cell lines and primary hepatocytes were those used previously.7 Human tumor cells were grown at 37 °C in a humidified, 5% CO₂ atmosphere in RPMI 1640 culture medium containing 5% FBS, 100 units ml⁻¹ penicillin and 100 µg ml⁻¹ streptomycin. U251 gliosarcoma cells implanted in scid mice (see below) were grown in RPMI/10% FBS and antibiotics. Adeno-2B11, a replication-deficient adenovirus encoding the cytochrome P450 gene CYP2B11, was previously described.20

DF3/Muc1 promoter activity assays

A DF3/Muc1 core promoter fragment (nucleotides −699 to +27) was PCR-amplified from MCF-7 genomic DNA cells using the primes: 5’-TATCTCGGACACCTAGGGTTCACTCGGAG-3’ and 5’-TATCTCGGATTCAAGGGCAGGCCGTGGC-3’, where lower case letters designate engineered restriction enzyme sites. The TATA box-containing DF3/Muc1 core promoter was verified by sequencing and sub-cloned into the reporter plasmid pGL3-Basic (Promega, Madison, WI) upstream of the luciferase gene using the restriction enzymes NheI and Xhol (New England Biolabs, Beverly, MA). The resulting luciferase reporter vector, pGL3-DF3-Luc, was transfected with TransIT-LT1 (Cat. #MIR 2300; Mirus, Madison, WI) into individual tumor cell lines followed by dual luciferase analysis.7

qPCR analysis

DF3/Muc1 and hTERT RNAs were quantified in tumor cell lines and in solid tumors grown subcutaneously in scid mice by qPCR as described7 using the following human-specific primers: 5’-ACGGCGAATGGGAAAGAAAC-3’ (hTERT sense), 5’-CAGCTGCTTCCGAAGTTCAC-3’ (hTERT antisense), 5’-TTTCTCTTGACTCTGCTGCTCC-3’ (DF3/Muc1 sense) and 5’-AGCCGAAGTCTCCTTCTTCTCCA-3’ (DF3/Muc1 antisense). qPCR analysis of interleukin-8 and ALDH1A1 RNAs used the following primers: 5’-TGTTGTATGCTGATGCGACTT-3’ (AL DH1A1 sense), 5’-CCATGGTGCTGAAATTTCAACAG-3’ (ALDH1A1 antisense), 5’-CTGGGTGCAGAGGTTTG TGGAGA-3’ (interleukin-8 sense) and 5’-TGCGAACC TACACAGACCCCA-3’ (interleukin-8 antisense). Results were analyzed using the comparative C_T method as described by the manufacturer. Data were normalized to the 18S RNA content of each sample, determined by qPCR as described.7

Construction of Adeno-DF3-E1A/hTERT-E1A

The single E1A cassette Adeno-hTERT-E1A was prepared as described.7 To construct a DF3-E1A expression cassette, luciferase complementary DNA was excised from pGL3-DF3-Luc using HindIII and XhoI and the vector was religated to generate pGL3-DF3. Next, the viral E1A gene was PCR-amplified from the plasmid pXC1 (Microbix, Toronto, ON, Canada) using the primers 5’-TAGAGATCTGGCACTCGCC-3’ (DF3/Muc1 antisense) and 5’-TAGAGATCTCCTCACCTG-3’ (reverse). The PCR product was BglII digested and ligated into BglII-digested pGL3-DF3 to yield pGL3-DF3-E1A. The DF3- and hTERT-E1A cassettes used to construct the dual E1A cassette adenovirus both contain the same E1A gene sequence and were verified for sequence integrity and functionality as described.7 To generate the final adenoviral construct, Adeno-DF3-E1A/hTERT-E1A, the entire DF3-E1A cassette, including an SV40 late polyA signal, was excised using BamHI, blunt-ended, digested with MluI for complete excision and inserted into a SpeI-digested, blunt-ended and then MluI-digested pHShuttle-hTERT-E1A.7 This cloning step ensured directionality of the incorporated DF3-E1A-polyA cassette and insertion ~2 kb upstream of the previously positioned hTERT-E1A-polyA cassette. The entire ICeul-PISceI pShuttle fragment containing the DF3-E1A and hTERT-E1A expression cassettes was then ligated by Dr Y Jouanaidi of this laboratory into the adenoviral genome using the Adeno-X expression system, as per the manufacturer’s instructions (Clontech Labs, Mountain View, CA). The Adeno-X genome is E1- and E3-region deleted, leaving room for up to ~8.3 kb of sequence insertion; the cassettes for DF3-E1A-polyA (~2.1 kb), hTERT-E1A-polyA (~1.75 kb) and pHShuttle (~1.1 kb, including an unused cytomegalovirus promoter insertion site) constitute ~4.95 kb of that space, allowing for up to 3 kb additional sequence downstream of the cytomegalovirus promoter (Figure 1c). Adenoviral stocks were propagated, amplified and purified.21 Viral titers were quantified and
verified in triplicate in HEK 293 cells using the Adeno-X Rapid Titer kit (Clontech Labs); as such, they represent functional titers on the basis of the final hexon production. Virus aliquots were stored at −80°C. Genomic integrity of the final virus preparation was verified by PCR and DNA sequencing.

Cell line infectivity and oncolysis
The intrinsic adenoviral infectivity of each cell line was determined by infection with Adeno-β-Gal. To compare the cytolytic activity of Adeno-hTERT-E1A to that of Adeno-DF3-E1A/hTERT-E1A, cells were plated overnight in triplicate at 14,000 cells per well of a 24-well plate. After 24 h the cells were infected for a 4-h period with virus in 200 μl culture medium containing the indicated multiplicity of infection (MOI) of Adeno-hTERT-E1A or Adeno-DF3-E1A/hTERT-E1A, after which 1 ml of fresh medium was added to each well. Cells remaining 3 or 6 days later were stained and quantified using crystal violet. Data are expressed as cell number (A595) relative to uninfected cell controls, mean ± s.d. for triplicate samples.

Western blotting and caspase assays
Western blotting for E1A and PARP-p85 protein used extracts prepared from adenovirus-infected cells seeded at 300,000 cells per well in a 6-well plate 24 h before infection. To assay virus-induced apoptosis, tumor cells were seeded in triplicate at 5000 cells per well in a 96-well plate. The next day, cells in each well were infected with Adeno-hTERT-E1A or Adeno-DF3-E1A/hTERT-E1A at MOIs of 0, 1, 10 and 100 in a total vol of 50 μl per well. After a 30 h infection, 50 μl of reconstituted Promega Caspase-Glo luminescence assay reagent (Cat#G8091, Promega) was added to each well, incubated for 1 h, and then luminescence readings were taken on a Wallac VICTOR® 1420 Multilabel Counter plate-reader (Cat# 1420–032, Perkin Elmer, Waltham, MA). Cell line-specific
background activity, determined from 0 MOI control samples, was subtracted from each sample.

**Human tumor xenografts**
A 5-week-old (24–26 g) male ICR/Fox Chase scid mice (Taconic Farms, Germantown, NY) were housed in the Boston University Laboratory of Animal Care Facility and treated in accordance with approved protocols and federal guidelines. Mice were injected subcutaneously on each posterior flank with 6 × 10⁶ U251 cells in 0.2 ml serum-free RPMI using a 0.5-inch 29-gauge needle and 1 ml insulin syringe. Tumor areas (length × width) were measured twice weekly using Vernier calipers (VWR International, Cat# 62379-531) and tumor volume was calculated on the basis of \( V_{ol} = (\pi/6)(L \times W)^{3/2} \). When the tumors reached ~150 mm³ in vol, the mice were divided into three groups (5 mice, 10 tumors per group): (a) control group, phosphate-buffered saline (vehicle) injected intratumorally, 90 µl per tumor per day on two consecutive days; (b) Adeno-hTERT-E1A, intratumorally 5 × 10⁸ plaque forming units injected at a single site in a volume of 90 µl per tumor per day on two consecutive days (total 1 × 10⁸ plaque forming units); and (c) Adeno-DF3-E1A/hTERT-E1A, intratumorally 5 × 10⁸ plaque forming units injected in 90 µl per tumor per day on two consecutive days (total 1 × 10⁹ plaque forming units). A second series of two daily virus or vehicle injections was administered to each tumor 15 days after the first treatment cycle. Tumor growth rates before virus administration were similar in all three groups. One-way analysis of variance analysis using the Bonferroni multiple comparison was carried out using GraphPad Prism 4 software (San Diego, CA).

**U251 holoclone studies**
Holoclones were identified on the basis of the colony morphology after single cell clonal seeding in RPMI + 7% FBS (1 cell per well in 96-well plates).19 Holoclones represented ~10% of the original U251 cell population used to select the holoclones; the other U251 cells displayed a meroclone (~30%) or paraclone morphology (~60%). RNA analysis of individual holoclones (clones designated A1, B3 and B6) was carried out in comparison to U251 parental cells and mixed meroclone/paraclone subpopulations by qPCR using RNA extracted from 300,000 cells per well of a 6-well plate for each clone. The intrinsic adenoviral infectivity of the U251 clones, and the parental U251 cells, was determined by X-gal staining after a 48-h infection with Adeno-β-gal at MOIs of 0, 0.25, 1 and 5. X-gal-stained cells were eluted with dimethyl sulfoxide (DMSO) and \( A_{560} \) was determined.

**Titration of functional virus particles released into culture supernatant**
Adenovirus titers in culture supernatants were determined by hexon staining of 293 cells incubated for 30 h with the culture supernatant from adenovirus-infected U251 and A549 cells2 with the following modifications. Cells were infected for 4 h with either Adeno-hTERT-E1A or Adeno-DF3-E1A/hTERT-E1A at an MOI of 5, washed with 1 ml of fresh RPMI/5% FBS growth medium, refreshed with 1 ml of RPMI/5% FBS and incubated for the indicated times (1, 3 or 5 days for U251 cells, and 3, 5 or 7 for A549 cells). At each time point, supernatant from each well (~1 ml) was used to prepare 10-fold serial dilutions (10⁰, 10⁻¹ and 10⁻² for U251 supernatants; 10⁻², 10⁻³ and 10⁻⁴ for A549 supernatants). Each dilution (0.5 ml) was used to infect 293 cells (seeded in 24-well plates 24 h earlier at 100,000 cells per well) for a 4 h period, followed by the addition of 1 ml of fresh RPMI/5% FBS medium per well. The infected 293 cells were incubated for a further 26 h (30 h total infection), methanol-fixed, stored at −20°C and processed for hexon immunostaining.7

**Helper virus studies**
U251 cells were seeded at 40,000 cells per well in 12-well plates. 24 h later the cells were infected with Adeno-2B11 (at MOI 8) in combination with Adeno-hTERT-E1A, Adeno-DF3-E1A/hTERT-E1A, or ONYX-015 (at MOIs of 0, 1.5 and 5), and in a total volume of 0.4 ml for an initial period of 4 h, after which 1 ml of fresh medium was added per well. After 72 h, U251 supernatants (~1.5 ml per well) were extracted, mixed thoroughly and divided into seven aliquots of 0.2 ml each. Aliquots were applied to fresh U251 cells, seeded 24 h earlier in 24-well plates at 4000 cells per well, to assay replicating virus-dependent CPA cytotoxicity. A sample of each supernatant (0.2 ml) was also applied to fresh 293 cells to titrate adenovirus release, as described above. Each aliquot of U251 culture supernatant was applied to the fresh U251 cells for a 4-h period, after which 0.5 ml fresh medium was added per well. The next day (day 4), the culture medium was aspirated and replaced by 1 ml of fresh medium containing CPA at 0–1000 µM, final concentrations. Cells were incubated for 48 h, the time at which culture medium was changed to fresh-CPA-containing medium and the cells were incubated for another 4 days (10 days total). Cells were stained with crystal violet and relative cell numbers were determined by \( A_{595} \).

**Adeno-2B11 spread and high-performance liquid chromatography analysis of 4-OH-CPA production**
Cells were seeded in 6-well plates at 75,000 cells per well. 24 h later, the cells were infected for 4 h with Adeno-2B11 (MOI 0 (control), MOI 8 (U251 cells) or MOI 32 (MCF-7/4HC and MDA-MB-231 cells), on the basis of each cell line’s intrinsic adenoviral infectivity) in combination with either Adeno-hTERT-E1A or ONYX-015 at a range of MOIs in 1 ml RPMI 1640 containing 5% FBS. Fresh culture medium (2 ml) was then added directly to each well and the incubation with virus was continued for 48 h total. The cells were then treated for 4 h with 250 µM CPA in 3 ml fresh RPMI 1640 containing 5% FBS and 5 mM semicarbazide. Culture medium (0.5 ml) was then removed and assayed for 4-OH-CPA by high-performance liquid chromatography.20 The remaining cells were quantified by crystal violet staining. Controls, either incubated without CPA or uninfected cells, were set to 100% cell survival.

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Results

Tumor cell transcriptional activity of core DF3-promoter

The DF3/Muc1 core promoter (nucleotides −699 to +27) contains an E-box, several GC-rich regions and Sp1 sites and a TATA box. To ascertain whether this promoter fragment could confer strong expression in tumor cells, a luciferase reporter plasmid driven by this promoter was transfected into a panel of human tumor cell lines (Figure 1a). DF3/Muc1 promoter-regulated luciferase activity had a cell line expression profile similar to that of endogenous DF3/Muc1 RNA (Figure 1b) in most of the cell lines tested, suggesting that the core promoter activity is an important determinant of cellular DF3/Muc1 RNA levels. However, in the case of BT-549 and A549 cells, high luciferase promoter activity but low endogenous RNA was observed, suggesting negative regulatory sequences outside of the core DF3 promoter construct may be important for DF3 expression in these two cell lines. DF3/Muc1 RNA varied between cell lines but was most highly expressed in the breast cancer cell lines (Figure 1b; first six cell lines). DF3/Muc1 RNA was barely detectable in human hepatocytes, where adenoviruses have a strong natural tropism, supporting the use of DF3/Muc1 core promoter to facilitate cancer-specific adenoviral expression.

Oncolytic profile of Adeno-DF3-E1A/hTERT-E1A

We constructed Adeno-DF3-E1A/hTERT-E1A (Figure 1c), which includes a DF3/Muc1 promoter-driven E1A cassette in addition to the hTERT promoter driven E1A cassette found in Adeno-hTERT-E1A. The oncolytic profile of this dual E1A cassette virus was compared with that of the single E1A cassette Adeno-hTERT-E1A (Figure 2). In U251 human glioblastoma cells, somewhat higher MOIs of Adeno-hTERT-E1A were required to achieve a comparable level of oncolysis, as seen after 3 and 6 days (Figure 2a). Although U251 cells have high endogenous hTERT activity and are deficient in DF3/Muc1 RNA (Figure 1b), the activity of the core DF3/Muc1 promoter incorporated into our adenovirus (Figure 1a) although low, is apparently sufficient to increase Adeno-DF3-E1A/hTERT-E1A-dependent oncolysis compared with Adeno-hTERT-E1A-infected cells. A greater increase in oncolytic activity with the dual E1A cassette virus was seen in A549 lung cancer cells and in MCF-7/4HC breast cancer cells (Figures 2b and c), where DF3/Muc1 promoter activity is high (Figure 1a). Oncolysis was weaker in MDA-MB-231 cells (Figure 2d), which show low DF3/Muc1 promoter activity (Figure 1a) and low intrinsic adenoviral infectivity.

Increased expression of E1A and enhanced tumor cell apoptosis by dual E1A cassette virus

E1A protein production, a prerequisite for adenovirus replication and an indicator of the potential of an adenovirus to lyse infected cells, was monitored in tumor cells infected with Adeno-DF3-E1A/hTERT-E1A and Adeno-hTERT-E1A. In accord with the oncolytic activity profiles shown in Figure 2, E1A protein levels were higher in cells infected with the dual cassette virus (Figure 3). Thus, addition of the second, DF3 promoter-regulated E1A cassette resulted in increased E1A expression in each cell line.

Although adenoviral infection typically leads to cell death by a non-apoptotic mechanism, the oncolytic viruses described here both lack E1B region genes that code for anti-apoptotic factors, E1B-19 kDa and E1B-55 kDa. Deletion of these genes may increase selectivity for cancer cells, where apoptotic pathways are mutated or suppressed. The increased expression of E1A in tumor cells infected with Adeno-hTERT-E1A, combined with the E1B gene deletion, leads to tumor cell killing by an E1A-induced apoptotic mechanism. Tumor cells infected with Adeno-DF3-E1A/hTERT-E1A were assayed and compared with cells infected with Adeno-hTERT-E1A for PARP cleavage, an early apoptotic event. PARP cleavage to yield PARP-p85 occurred at lower MOIs of Adeno-DF3-E1A/hTERT-E1A as compared with Adeno-hTERT-E1A in MCF-7/4HC and A549 cells (Figures 3a and b). In U251 cells, PARP-p85 levels (data not shown) and caspase 3/7 activity (Figure 3c), which reflects downstream apoptotic events, were very similar after infection with either virus, consistent with the similar, and near complete, oncolytic activity of each virus seen in Figure 2a. Addition of the second E1A cassette significantly increased caspase-3/7 activity in MDA-MB-231, MCF-7/4HC and A549 cells (Figure 3c), consistent with the more substantial increases in PARP cleavage and oncolysis seen in these cells.

Activity in a U251 tumor xenograft model

The anti-tumor activity of Adeno-DF3-E1A/hTERT-E1A was compared with that of Adeno-hTERT-E1A in scid mice bearing subcutaneous human U251 tumor xenografts (Figure 4a). Significant tumor growth inhibition was achieved with Adeno-DF3-E1A/hTERT-E1A (23.4 ± 7.6% inhibition after treatment period 1 (days 0–14) (P < 0.05 vs control; one-way analysis of variance analysis with Bonferroni multiple comparison) and 36.9 ± 11.2% inhibition (P < 0.01 vs control) by the end of treatment period 2 (days 14–37), but not with Adeno-hTERT-E1A. The aggressive growth of U251 tumor cells resulted in only a short period of growth stasis, which was apparent after the second series of Adeno-DF3-E1A/hTERT-E1A injections. The tumor growth trends were significantly different for each virus (P < 0.05) but only the Adeno-DF3-E1A/hTERT-E1A treatment group was significantly different from the virus-free control (P < 0.01). No significant host toxicity was observed with either virus, as determined by monitoring body weights (data not shown).

Next, we investigated why the dual E1A cassette virus was more active than the single, hTERT-E1A cassette virus in U251 tumors, given the high hTERT promoter activity and low DF3/Muc1 promoter activity seen in U251 cells (Figure 1a). Whereas cultured U251 cells express DF3/Muc1 RNA at a low level, we observed ~43-fold higher levels of DF3/Muc1 RNA in U251 tumor xenografts (Figure 4b). Conversely, hTERT RNA
was ~10-fold lower in U251 tumors than in cultured U251 cells (Figure 4b). These tumor microenvironment-dependent increases in DF3/Muc1 and decreases in hTERT expression explain the enhanced in vivo antitumor activity of Adeno-DF3-E1A/hTERT-E1A as compared with Adeno-hTERT-E1A. A very similar pattern of increased DF3/Muc1 expression (~21-fold) and decreased hTERT expression (~15-fold) was seen in A549 tumors compared with cells (Figure 4c). In MDA-MB-231 tumors, DF3/Muc1 expression was also increased, by ~10-fold, however, hTERT expression was unchanged compared with cultured tumor cells (Figure 4d).

Activity of Adeno-DF3-E1A/hTERT-E1A against U251 holoclones
Cultured tumor cells that grow with a holoclone morphology are enriched in cancer stem-like cells. U251 holoclones were isolated and characterized with respect to DF3/Muc1 and hTERT expression. Figure 5a shows a 10-fold increase in DF3/Muc1 RNA and a 5-fold increase in hTERT RNA in U251 holoclones compared with parental U251 cells.

Figure 2. Oncolytic activity of Adeno-DF3-E1A/hTERT-E1A and Adeno-hTERT-E1A—four human tumor cell lines: U251 (a), A549 (b), MCF-7/4HC (c), and MDA-MB-231 (d), were infected with Adeno-DF3-E1A/hTERT-E1A or Adeno-hTERT-E11A at the indicated MOIs. The cells were stained with crystal violet 3 (left) or 6 days (right) later to quantify remaining cells. Data point represents mean ± s.d. values on the basis of n = 3 replicates, with the uninfected cell controls set = 100%.

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decrease in hTERT RNA for three independent U251 holoclones as compared with either the parental U251 cell line or a mixed U251 merocline/paraclone subpopulation, which comprises ~90% of the parental cell population. DF3/Muc1 is associated with tumor stem-like cells in breast cancer. Expression of ALDH1A1, an established tumor stem-like cell marker, was also enriched ~10-fold in all three holoclones, whereas the very high parental cell expression of interleukin-8, which is associated with tumor progression and metastasis, was abolished (data not shown), consistent with the cancer stem cell-like nature of these holoclones. In agreement with the observed changes in hTERT and DF3/Muc1 expression, U251 holoclones showed decreased response to Adeno-hTERT-E1A, as indicated by the substantially lower E1A protein level in U251 holoclones infected with Adeno-hTERT-E1A compared with Adeno-DF3-E1A/hTERT-E1A (Figure 5b). Adeno-hTERT-E1A also showed substantially reduced oncolysis of the holoclones compared with parent U251 cells (Figure 5c), whereas Adeno-DF3-E1A/hTERT-E1A oncolytic activity showed a small decrease in activity (Figure 5d), which can be explained by the reduced adenoviral infectivity of the holoclones, as determined by
Adeno-hTERT-E1A disrupts its ability to produce active, functional viral particles and thereby spread the viral infection within a tumor cell population. Supernatants from tumor cells infected with Adeno-DF3-E1A/hTERT-E1A or Adeno-hTERT-E1A were collected and titred for functional virus. Adeno-hTERT-E1A was more efficient than Adeno-DF3-E1A/hTERT-E1A in terms of the release of functional virus, as seen at various time points after infection, in both U251 cells and A549 cells (Figure 6a). This finding suggests that the second E1A cassette of Adeno-DF3-E1A/hTERT-E1A induces premature host cell death, reducing the overall yield of functional viral progeny.

To determine how the decrease in adenovirus release affects the expression of a co-administered replication-deficient adenovirus, studies were carried out with Adeno-2B11, which codes for the prodrug-activating cytochrome P450 enzyme 2B11 (CYP2B11), an efficient catalyst of CPA activation.29 U251 cells were infected with Adeno-2B11 in combination with Adeno-hTERT-E1A, Adeno-DF3-E1A/hTERT-E1A or another oncolytic virus, ONYX-015, at various MOIs. Around 3 days later, the culture supernatants were applied to fresh U251 cells in the presence of CPA, a CYP2B11-activated prodrug. Adeno-hTERT-E1A was the most effective in facilitating Adeno-2B11 replication, resulting in CPA-induced cell death, whereas ONYX-015 showed moderate activity (Figure 6b). Adeno-DF3-E1A/hTERT-E1A was almost inactive (Figure 6b), consistent with the early oncolysis and reduced viral release (Figure 6a) comprising the Adeno-2B11 helper virus activity. Analysis of culture supernatants for the release of 4-OH-CPA, the active metabolite of CPA, confirmed the superior ability of Adeno-hTERT-E1A to serve as a helper virus in U251 cells, as evidenced by the lower MOIs required to produce high levels of 4-OH-CPA (Figure 6c). At higher MOIs, Adeno-hTERT-E1A induced substantial cell death (data not shown), thereby decreasing the number of cells capable of converting CPA to 4-OH-CPA. Adeno-hTERT-E1A and ONYX-015 exhibited similar helper virus activity in MCF-7/4HC cells (Figure 6d), whereas in MDA-MB-231 cells, which are intrinsically difficult to infect, Adeno-hTERT-E1A showed low helper virus activity, whereas ONYX-015 was inactive (data not shown).

Enhanced tumor cell apoptosis suppresses viral release and helper virus activity of dual cassette virus

Next, we investigated whether the earlier and more extensive apoptosis induced by Adeno-DF3-E1A/hTERT-E1A on each of two consecutive days (arrows), with the virus treatment repeated 14 days later. (a) Relative tumor volume, normalized to the tumor volume on the first day of virus infection (day 0) (mean ± s.e. for n = 5 mice and n = 10 tumors per group). Results of one-way analysis of variance analysis of tumor volume data with Bonferroni multiple test correction are as shown (*P < 0.05; **P < 0.001). (b-d) DF3/Muc1 (left) and hTERT (right) RNA were assayed by qPCR in solid tumor xenografts grown in aicd mice from U251, A549 and MDA-MB-231 cells (four different tumor RNA samples) and in four separate RNA preparations from the corresponding cell lines grown in culture. RNA levels were set relative to the mean RNA level of either the four cell-culture (for DF3) or the four tumor (for hTERT) samples. Data shown are mean ± s.d. for n = 3 replicates. Statistical analysis by two-tailed unpaired Student’s t-test with 95% confidence intervals comparing grouped tumor RNA levels to grouped cell culture RNA levels: P < 0.001 (**); P < 0.0001 (***); and P < 0.05 (*).

Discussion

Cancer-specific promoters can be used to control viral regulatory genes, such as adenoviral E1A, to restrict the replication of oncolytic adenoviruses to malignant cells and tissues. Dual-specificity adenoviral promoters that regulate E1A expression in response to multiple stimuli, for example, estrogens and hypoxia, have also been described.31 The cancer-specific promoters DF3/Muc1 and hTERT can both confer tumor-specific replication, and several tumor cell-replicating, DF3/Muc1-driven and hTERT-driven adenoviruses have been described.8,32 However, none of these viruses combine both promoter elements into a single virus to regulate E1A expression and viral replication. As hTERT is expressed in > 90% of cancers,5,6 and DF3/Muc1 displays increased activity in 75% of solid tumors and in cancer stem cell-associated subpopulations,17,18 an oncolytic virus that combines both of these features has the potential to induce oncolytic activity across a broad range of human tumors and tumor cell populations. The heterogeneity of DF3 and hTERT expression within a single tumor, discussed below, could further increase the utility of such a virus. Presently, we describe a novel adenovirus, Adeno-DF3-E1A/hTERT-E1A, where DF3/Muc1 and hTERT gene regulatory sequences are combined into a single virus to develop a therapeutic approach to targeting heterogeneous populations of tumor cells, including cancer stem-like cells, within a given patient’s tumor.

Adeno-hTERT-E1A is a single hTERT-E1A cassette adenovirus that uses an hTERT core promoter to drive adenoviral E1A expression in a manner similar to that of the endogenous full-length hTERT promoter.7 Furthermore, Adeno-hTERT-E1A combines hTERT regulation of E1A with deletion of the E1B region genes E1B-19 kDa and C2B11,20 which codes for the prodrug-activating cytochrome P450 enzyme 2B11 (CYP2B11), an efficient catalyst of CPA activation.29 U251 cells were infected with Adeno-2B11 in combination with Adeno-hTERT-E1A, Adeno-DF3-E1A/hTERT-E1A or another oncolytic virus, ONYX-015, at various MOIs. Around 3 days later, the culture supernatants were applied to fresh U251 cells in the presence of CPA, a CYP2B11-activated prodrug. Adeno-hTERT-E1A was the most effective in facilitating Adeno-2B11 replication, resulting in CPA-induced cell death, whereas ONYX-015 showed moderate activity (Figure 6b). Adeno-DF3-E1A/hTERT-E1A was almost inactive (Figure 6b), consistent with the early oncolysis and reduced viral release (Figure 6a) comprising the Adeno-2B11 helper virus activity. Analysis of culture supernatants for the release of 4-OH-CPA, the active metabolite of CPA, confirmed the superior ability of Adeno-hTERT-E1A to serve as a helper virus in U251 cells, as evidenced by the lower MOIs required to produce high levels of 4-OH-CPA (Figure 6c). At higher MOIs, Adeno-hTERT-E1A induced substantial cell death (data not shown), thereby decreasing the number of cells capable of converting CPA to 4-OH-CPA. Adeno-hTERT-E1A and ONYX-015 exhibited similar helper virus activity in MCF-7/4HC cells (Figure 6d), whereas in MDA-MB-231 cells, which are intrinsically difficult to infect, Adeno-hTERT-E1A showed low helper virus activity, whereas ONYX-015 was inactive (data not shown).
E1B-55 kDa to further increase tumor cell-specificity. \(^7\) Adeno-DF3-E1A/hTERT-E1A contains the same hTERT-E1A expression cassette and E1B gene deletions as Adeno-hTERT-E1A, but in addition, places a second E1A gene cassette under control of a core DF3/Muc1 promoter element. This promoter element was chosen on the basis of the earlier promoter deletion studies\(^{23-25}\) and is shown here to regulate expression in a cell line-dependent manner similar to that of endogenous DF3/Muc1 RNA. Incorporation of a second E1A gene cassette did not lead to recombination or instability of the dual DF3/hTERT adenovirus genome following viral amplification. Functionally, the second E1A cassette conferred on Adeno-DF3-E1A/hTERT-E1A the ability to induce higher E1A expression, more extensive apoptosis and greater oncolysis as compared with Adeno-DF3-E1A/hTERT-E1A.

Figure 5  Gene expression and adenoviral activity in U251 holoclones—(a) qPCR analysis of DF3/Muc1 (left) and hTERT (right) RNA in U251-derived holoclones (designated A1, B3 and B6), the parental U251 cell population and a meroclone/paraclone (M/P) mixed cell population. RNA levels were set relative to the U251 parental cells and are mean ± s.d. values for \(n = 3\) replicates. Statistical analysis of RNA levels in holoclones vs U251 parental and meroclone/paraclone population: **\(P < 0.01\); ***\(P < 0.001\). (b) Western blot analysis of E1A (top) and PARP p85 protein (bottom) from total cell lysates (50 \(\mu\)g per lane) prepared from U251 holoclones and parental cells infected for 24 h with Adeno-hTERT-E1A (‘hT’) or Adeno-DF3-E1A/hTERT-E1A (‘D/hT’), as indicated. (c) and (d) U251 holoclones and parental cells were seeded 24 h before infection with Adeno-hTERT-E1A (c) or Adeno-DF3-E1A/hTERT-E1A (d) at the indicated MOIs. The cells were stained with crystal violet 3 days later to quantify remaining cells. Data shown are mean ± s.d. values based on \(n = 3\) replicates, with the uninfected cell controls set = 100%. One-way analysis of variance with Bonferroni multiple test correction indicated no significant difference in Adeno-DF3-E1A/hTERT-E1A-induced cytotoxicity profiles for the holoclones vs parental U251 cells (d), whereas cytotoxicity profiles were significantly different for Adeno-hTERT-E1A-infected holoclones B3 and B6 (but not holoclone A1) compared with parental U251 cells (\(P < 0.05\)) (c). Adeno-hTERT-E1A-induced cytotoxicity profiles were significantly different for holoclone A1 compared with U251 parental cells when analyzed at individual MOIs, ranging from 5 to 20 MOI (\(P < 0.05\)).
hTERT-E1A. The studies reported here were all carried out in parallel using carefully titered batches of the single hTERT and the dual DF3/hTERT adenoviruses, making it highly likely that the additional activity of Adeno-DF3-E1A/hTERT-E1A is due to the second, DF3 promoter-regulated E1A cassette. Efforts to prepare a single DF3 virus to verify this were unsuccessful.

The oncolytic activity of Adeno-DF3-E1A/hTERT-E1A was increased over that of Adeno-hTERT-E1A in several cultured tumor cells, with the increased activity also apparent in vivo in a U251 gliosarcoma solid tumor xenograft model (Figure 4a). Given the high adenoviral infectivity and high hTERT promoter activity of U251 cells, Adeno-hTERT-E1A and Adeno-DF3-E1A/hTERT-E1A were both expected to induce strong anti-tumor responses. However, Adeno-hTERT-E1A, which induced extensive U251 cell death in culture, showed no significant anti-tumor activity in U251 tumor xenografts, in contrast to Adeno-DF3-E1A/hTERT-E1A. The ineffectiveness of Adeno-hTERT-E1A, but not Adeno-DF3-E1A/hTERT-E1A, against U251 tumors in vivo may be explained by the 10-fold decrease in hTERT expression, coupled with the 43-fold increase in DF3/Muc1 expression that we observed in U251 solid tumors compared with cultured U251 cells (Figure 4b). Thus, incorporation of a DF3/Muc1 promoter, which was previously shown to have augmen-

Figure 6 Adeno-DF3-E1A/hTERT-E1A exhibits decreased viral spread and diminished helper virus activity towards Adeno-2B11-mediated CPA prodrug-activation—(a) U251 cells (left) and A549 cells (right) were infected with Adeno-hTERT-E1A (hTERT) or Adeno-DF3-E1A/hTERT-E1A (DF3/hTERT) at MOI 5. The cell culture supernatant was removed 1–7 days later, as indicated, and applied to 293 cells to titrate adenoviral particles released into the supernatant. Data shown are mean ± s.d. values based on duplicate titerings from three 10-fold serial dilutions (10^0, 10^-1 and 10^-2) of the virus supernatants. Numbers over each bar indicate the ratio of adenoviral titer between Adeno-hTERT-E1A and Adeno-DF3-E1A/hTERT-E1A. Statistical analysis by two-tailed unpaired Student’s t-test with 95% confidence intervals for Adeno-hTERT-E1A vs Adeno-DF3-E1A/hTERT-E1A: *P < 0.05; **P < 0.001. (b) CPA dose-dependent killing of U251 cells infected with cell culture supernatants from U251 cells previously infected with Adeno-2B11 (MOI 6) in combination with Adeno-hTERT-E1A (left), ONYX-015 (middle) or Adeno-DF3-E1A/hTERT-E1A (right) at MOIs of 0, 1.5 and 5. Cells treated with virus in the absence of CPA were set as 100%. (c) and (d) The active metabolite of CPA, 4-OH-CPA, was assayed by high-performance liquid chromatography in U251 and MCF-7/4HC cells infected with Adeno-2B11 + Adeno-hTERT-E1A (white bars) or with Adeno-2B11 + ONYX-015 (black bars) for 2 days followed by a 4-h incubation with 250 μM CPA + semicarbazide (5 mM). Data shown in (b) and (c) are mean ± s.d. values based on n = 2 replicates. All experiments were performed at least in duplicate.
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zation of U251 cells to CPA and by the increased production of active prodrug metabolites (4-OH-CPA) seen in U251 cells. The loss of helper virus activity in the case of Adeno-DF3-E1A/hTERT-E1A demonstrates the importance of balancing E1A production with the extent and the timing of virus-induced apoptosis. Indeed, to be most effective, P450 and other prodrug enzyme-based gene therapy strategies require delayed, rather than accelerated death of tumor cells that express the prodrug-activating enzyme.\(^5\)

In conclusion, we have engineered a novel dual E1A cassette oncolytic adenovirus that displays improved oncolysis in many cancer cell lines compared with the corresponding single E1A-cassette adenovirus and has the potential to target a heterogeneous tumor cell population, including cancer stem-like cells, which display strong DF3/Muc1 promoter activity. Despite the increased activity of the dual E1A cassette virus as a stand-alone anti-cancer agent, the premature apoptosis of infected tumor cells decreased its potential to replicate and to facilitate helper virus activity. Future studies may examine adenoviral genomes that combine the dual promoter E1A gene control described here with the retention of adenoviral E1B and/or E3 region genetic elements to prevent premature host cell apoptosis and thereby facilitate more efficient production of viral progeny while maintaining activity in a wide range of cancer types and tumors with heterogeneous tumor cell populations.

Conflict of interest

The authors declare no conflict of interest.

Abbreviations

4-OH-CPA, 4-hydroxycyclophosphamide; CMV, cytomegalovirus; CPA, cyclophosphamide; E1A, early adenoviral region 1A; FBS, fetal bovine serum; hTERT, human telomerase; MOI, multiplicity of infection; PARP, poly (ADP-ribose) polymerase; pfu, plaque forming units; qPCR, quantitative PCR; RPMI, Roswell Park Memorial Institute.

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References


