Adenovirus-mediated delivery of interferon-γ gene inhibits the growth of nasopharyngeal carcinoma

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Abstract

Background: Interferon-γ (IFN-γ) is regarded as a potent antitumor agent, but its clinical application is limited by its short half-life and significant side effects. In this paper, we tried to develop IFN-γ gene therapy by a replication defective adenovirus encoding the human IFN-γ (Ad-IFNγ), and evaluate the antitumoral effects of Ad-IFNγ on nasopharyngeal carcinoma (NPC) cell lines in vitro and in xenografts model.

Methods: The mRNA levels of human IFN-γ in Ad-IFNγ-infected NPC cells were detected by reverse transcription-polymerase chain reaction (RT-PCR), and IFN-γ protein concentrations were measured by enzyme-linked immunosorbent assay (ELISA) in the culture supernatants of NPC cells and tumor tissues and bloods of nude mice treated with Ad-IFNγ. The effects of Ad-IFNγ on NPC cell proliferation was determined using MTT assay, cell cycle distribution was determined by flow cytometry analysis for DNA content, and cells apoptosis were analyzed by Annexin V-FITC/7-AAD binding assay and hoechst 33342/PI double staining. The anti-tumor effects and toxicity of Ad-IFNγ were evaluated in BALB/c nude mice carrying NPC xenografts.

Results: The results demonstrated that Ad-IFNγ efficiently expressed human IFN-γ protein in NPC cell lines in vitro and in vivo. Ad-IFNγ infection resulted in antiproliferative effects on NPC cells by inducing G1 phase arrest and cell apoptosis. Intratumoral administration of Ad-IFNγ significantly inhibited the growth of CNE-2 and C666-1 cell xenografts in nude mice, while no significant toxicity was observed.

Conclusions: These findings indicate IFN-γ gene therapy mediated by replication defective adenoviral vector is likely a promising approach in the treatment of nasopharyngeal carcinoma.

Keywords: Gene therapy, Interferon-γ, Nasopharyngeal carcinoma, Adenoviral vector

Background

Nasopharyngeal carcinoma (NPC) is a rare tumor arising from the epithelium of the nasopharynx. However, it has a high incidence rate in South China and Southeast Asia [1]. Radiation therapy is the main strategy for local control of NPC [2], but the 5-year survival for stage IV NPC is only 30%. The poor survival is often associated with high incidences of local, regional and systemic recurrences.

Although concurrent chemoradiotherapy is developed as a standard treatment approach for advanced NPC, the outcome isn’t still satisfactory [3]. Therefore, the development of multidisciplinary therapeutic approaches is crucial for improvement of survival in NPC patients.

Interferon-γ (IFN-γ), a multifunctional cytokine produced mainly by T helper cells, cytotoxic T cells and natural killer cells [4], exerts antiviral, antiproliferative, immunomodulatory and antiangiogenesis effects [5]. IFN-γ inhibits the growth of numerous tumors [6-10] and has been tried to use in the clinical management of tumors [11-14]. However, the clinical application of recombinant IFN-γ protein is hampered by its short half-life and significant side effects. IFN-γ Gene therapy can continuously...
produce and release therapeutic protein in local focus to overcome the obstacle of recombinant IFN-γ protein [9,10,15,16]. In this study, we tried to develop IFN-γ gene therapy by a replication defective adenovirus encoding the human IFN-γ (Ad-IFNγ) on nasopharyngeal carcinoma (NPC), our results showed Ad-IFNγ effectively expressed in NPC cells, significantly inhibited tumor cell proliferation and induced cell apoptosis in vitro, inhibited the growth of xenografts in nude mice.

Methods
Materials
Dulbecco’s Modified Eagle Medium (DMEM), fetal bovine serum, propidium iodide (PI), and TRIzol reagent were from Invitrogen (Carlsbad, CA, USA). Reverse Transcription System and GoTaq® DNA Polymerase were from Promega (Beijing) Biotech Co., Ltd (Beijing, China). Annexin V-FITC/7-AAD apoptosis detection kit was from Beckman Coulter, Inc. (Marseille, France), while Double Stain Apoptosis Detection Kit (Hoechst 33342/PI) from GenScript USA Inc. (Piscataway, NJ, USA). Rabbit anti-human Ki-67 polyclonal antibody was from NeoMarkers For Lab Vision Corporation (Fremont, CA, USA), and Streptavidin-Horseradish Peroxidase kit from Beijing Zhongsan Golden Bridge Biotechnology Co. (Beijing, China). In Situ Cell Death Detection kit was purchased from Roche Applied Science (Mannheim, Germany). Human interferon-γ ELISA kit was purchased from Boster (Wuhan, China), recombinant human interferon-γ (rhIFN-γ) protein was from Shanghai Clonbiotech. Co., Ltd (Shanghai China). (3,4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) and all other reagents were of molecular biology grade and obtained from Sigma-Aldrich (Shanghai, China).

Cell lines, recombinant adenoviruses and infection
Human nasopharyngeal carcinoma (NPC) cell lines CNE-1, CNE-2 and C666-1 were maintained in DMEM containing 10% fetal bovine serum (FBS) at a humidified atmosphere with 5% CO2 at 37°C. Replication defective adenoviruses encoding human interferon-γ (Ad-IFNγ) and β-galactosidase (Ad-LacZ), kindly provided by Guangzhou Double Bioproducts Co., Ltd., were stored at -80°C for use. For adenovirus infection, NPC cells were seeded and cultured for 24 h, and then removed culture medium, washed with phosphate buffered saline (PBS) (pH 7.4), followed by infection with adenovirus in serum-free DMEM for 3 h. After removal of residual virus by PBS washing, cells were cultured in normal medium for indicated time before the analyses of IFN-γ expression, cell proliferation and apoptosis.

Reverse transcription-polymerase chain reaction (PCR)
Cells were harvested and total RNAs were extracted using TRIzol® reagent according to the manufacture’s instruction. mRNAs were transcribed into cDNAs and PCR reactions were carried out using their specific primer pairs: IFN-γ’s sense, 5’- TTACTGGGATGCTCTTCGAC-3’ (amplicon 473 bp); β-actin’s sense, 5’-CGTCTTCCCCCTCATCGTG-3’, antisense, 5’-TAGACAGCCTGGATAGCAAC-3’ (amplicon 334 bp). Amplification was done with an initial cycle of 95°C for 4 min, followed by 30 cycles of 95°C for 30 s, 55°C for 40 s, 72°C for 1 s, with a final extension at 72°C for 10 min. PCR products were analyzed by ethidium bromide staining on 1.5% agarose gels.

Enzyme-linked Immunosorbent Assay (ELISA)
NPC cells were seeded into 6-well plate at a density of 1×10⁵ cells/well for 24 hours and then infected with adenoviruses as described above. The culture supernatants were collected at different time points and IFN-γ concentration was determined by ELISA according to the manufacture’s instruction (Boster, Wuhan, China) (the sensitivity is 15.6 pg/mL).

Cell proliferation analysis
The effect of Ad-IFNγ on NPC cell proliferation was determined using MTT assay as previously described [17]. Briefly, cells were seeded in 96-well plates at a density of 2000 cells/well for 24 hours and then infected with adenoviruses as described above, followed by incubation for 72 h. Viable cells were stained with MTT for 4 hours and followed by determination of OD₅₇₀ nm with a reference wavelength at 630 nm.

Cell cycle distribution and apoptosis analysis
Cell cycle distribution was determined by flow cytometry analysis for DNA content. Briefly, cells were harvested by trypsinization and washed by PBS, and followed by fixed in cold 70% ethanol for 1 hour at 4°C. Cell suspensions were washed twice in PBS, treated by ribonuclease and followed by PI staining. And then cells were performed flow cytometry analysis for DNA content and cell cycle distribution. Cells apoptosis were analyzed by Annexin V-FITC/7-AAD binding assay and hoechst 33342/PI double staining. Both floating and adherent cells were harvested and washed by PBS, then followed by staining according to the manufacture’s instruction. Cells stained with Annexin V-FITC/7-AAD were analyzed by flow cytometry, while those stained with hoechst 33342/PI were analyzed by fluorescence microscopy.

Animal model and experimental design
Female BALB/c nude mice (5-6 weeks old, 18-20 g) were obtained from Shanghai Slike Experimental Animals Co.
PBS, 2×10^9 pfu (plaque-forming unit) of Ad-LacZ, 2×10^9 were treated by weekly intratumoral injection of 100 μL of PBS when CNE-2 xenografts reached an approximate diameter of 7 mm. Tumor tissues and bloods were collected at different time. IFN-γ concentration in the samples was measured by ELISA.

To assess antitumor effects of Ad-IFNγ in vivo, mice were intratumorally injected with 2×10^9 pfu in 100 μL of PBS when CNE-2 xenografts reached an approximate diameter of 7 mm. Tumor sizes and bloods were collected at different time. IFNγ concentration in the samples was measured by ELISA.

To assess antitumor effects of Ad-IFNγ in vivo, mice were randomly assigned into eight groups (7-8 mice per group) when xenografts reached 4-5 mm diameter. Mice were maintained by serial subcutaneous transplantation in BALB/c nude mice, were subcutaneously transplanted into the flanks of mice to construct xenograft model.

To assess antitumor effects of Ad-IFNγ in vivo, mice were randomly assigned into eight groups (7-8 mice per group) when xenografts reached 4-5 mm diameter. Mice were treated by weekly intratumoral injection of 100 μL of PBS, 2×10^9 pfu (plaque-forming unit) of Ad-LacZ, 2×10^9 pfu, 1×10^8 pfu, 5×10^7 pfu, or 1×10^6 pfu of Ad-IFNγ (in 100 μL of PBS) per dose respectively; while mice in other two groups were treated by daily intratumoral injection of 100 μL of normal saline or 1×10^6 IU/(kg body weight) of rhIFN-γ protein per dose respectively. The treatment was performed for three consecutive weeks, every injection was distributed equally into each half (8-12 mm of diameter) or quadrants (>12 mm of diameter) of tumors. Body weight and tumor size were measured every 4-5 days, and tumor xenografts were weighed at the end point of experiments.

Histological analysis
Tumor tissue was fixed in buffered formalin and embedded in paraffin. Sections (5 μM thick) were mounted on Poly-L-Lysine treated slides, standard H&E staining was utilized for histopathological assay. Human Ki67 immunohistochemistry was performed for cell proliferation assay, while TUNEL labeling was done using a fluorescent in situ cell death detection kit for apoptosis analysis in tumor tissues.

Statistical analysis
All experiments were repeated at least three times. The data were analyzed with One-way ANOVA or t test by using SPSS 10.0 for Windows software (SSPS Inc., Chicago, IL, USA). P < 0.05 was considered statistically significant.

Results
Ad-IFNγ efficiently expressed hIFNγ in NPC cells
To evaluate the capability of Ad-IFNγ expressed transgenic product in nasopharyngeal carcinoma (NPC) cells, we firstly investigated IFNγ mRNA levels in CNE-2 cells after Ad-IFNγ infection at different multiplicities of infection (MOIs) at different time. The results showed that Ad-IFNγ efficiently transcribed human IFNγ gene in CNE-2 cells at a dose-dependent manner (Figure 1A). hIFNγ mRNA was detected as early as 8 hours after Ad-IFNγ infection, the levels reached the top at 48 hours after infection, and then gradually decreased, while No IFNγ mRNA was detected in Ad-LacZ-infected cells (Figure 1B).

We investigated IFNγ expression at protein levels in CNE-1, CNE-2 or C666-1 infected by Ad-IFNγ, and found that IFNγ protein concentration in the supernatants of NPC cells continuously increased within 72 hours post-infection (Figure 1C). The differences from IFNγ mRNA which peaked at 48 hours may be due to the instability of the mRNA. No IFNγ proteins were found in the supernatants of Ad-LacZ-infected NPC cells (data not shown).

Ad-IFNγ inhibited the proliferation of NPC cells in vitro
We evaluated the effects of Ad-IFNγ on the proliferation of nasopharyngeal carcinoma (NPC) cells by MTT assay after 72 h after Ad-IFNγ infection. The anti-proliferative effects of Ad-IFNγ on NPC cells were shown in Figure 2A. Ad-IFNγ inhibited the proliferation of CNE-1, CNE-2 and C666-1 NPC cell lines from 1 to 100 MOIs at a dose-dependent manner. No significant inhibition was observed on NPC cells after infection with 50 MOIs of Ad-LacZ.

To explore the mechanism involved in proliferation inhibition of Ad-IFNγ on NPC cells, the cell cycle profiles were analyzed by flow cytometry for DNA contents on NPC cells after infection with 100 MOIs of adenoviruses for 72 h. The results showed a significantly higher percentage at the G1 phase in the cells infected with Ad-IFNγ than those in Ad-LacZ infected cells (CNE-2: 70.8% ± 6.3% vs 56.1% ± 5.9%, p<0.05; C666-1: 75.8% ± 6.9% vs 62.0% ± 4.8%, p<0.05) (Figure 2B). These indicated that Ad-IFNγ possesses an effect of G1 phase arrest on NPC cells.

Ad-IFNγ induced the apoptosis of NPC cells in vitro
To examine the fate of Ad-IFNγ-infected NPC cell lines, an Annexin V/7-AAD binding assay and Hoechst 33342/PI double staining were performed after treatment with 100 MOIs of adenoviruses for 72 h. The results showed that there was an increased fraction of Annexin V+/7-AAD− (early apoptosis) and Annexin V+/7-AAD+ (late apoptosis) in Ad-IFNγ-infected NPC cells than those of Ad-LacZ-infected cells (CNE-2, 21.2% vs 5.3%, p<0.01; C666-1, 16.3% vs 6.4%, p<0.01) (Figure 3A). In Hoechst 33342/PI double staining assay, there were higher percentages of bright blue cells (apoptosis) in NPC cells infected with Ad-IFNγ than those with Ad-LacZ (CNE-2, 25.2% vs 4.8%, p<0.01; C666-1, 19.5% vs 3.0%, p<0.01) (Figure 3B). These data suggested that Ad-IFNγ induced the apoptosis in NPC cells.
Ad-IFNγ efficiently expressed human IFNγ in NPC xenografts in nude mice

To evaluate the dynamic expression of AdIFNγ in vivo, we measured the levels of hIFNγ in tumor and blood samples collected from nude mice carrying CNE-2 NPC xenografts on days 1, 3, 5 and 7 after intratumoral injection with Ad-IFNγ. The results showed that Ad-IFNγ efficiently expressed hIFNγ in NPC xenografts, the concentration of hIFNγ in tumor tissue was (136.5±25.5) pg/(mg tissue) at 24 hours post-injection of AdIFNγ, reached the peak of (265.8±26.2) pg/(mg tissue) at 72 hours, and then decreased gradually to (61.1±8.3) pg/(mg tissue) at day 7 (Figure 4A). hIFNγ were also detected in blood. As same as in tumor tissue, the concentration of hIFNγ reached the peak of (64.9±15.9) pg/mL after 3 days post-injection, and then decreased gradually to (18.9±7.3) pg/mL at day 7.

Ad-IFNγ inhibited the growth of human NPC xenografts

Administration of Ad-IFNγ resulted in significant growth suppression of NPC xenografts compared to the control groups. As shown in Figure 5A and 5B, tumor growth in the Ad-IFNγ-treated groups was significantly slower than that in Ad-LacZ (vector control) group or PBS (medium control) group. The inhibition effects showed a clear dose-dependent manner. At the end of experiment, the tumor weights in Ad-IFNγ groups were significantly lighter than those in Ad-LacZ or PBS group (Figure 5C and 5D), 1×10^8, 5×10^8, 1×10^9 and 2×10^9 pfu/dose of Ad-IFNγ treatment respectively produced 24.1%, 43.2%, 64.8%, and 73.5% of growth inhibition for CNE-2 xenografts (Figure 5C), 42.6%, 54.3%, 71.3% and 81.7% of growth inhibition for C666-1 xenografts (Figure 5D). However, 2×10^9 pfu/dose of Ad-LacZ barely inhibited tumor growth. The efficacy of daily intratumoral injection of 1×10^6 IU/(kg body weight)/day of rhIFNγ was likely similar with that of the administration of 5×10^9 pfu/dose/week of Ad-IFNγ. There were no significant differences on body weights between Ad-IFNγ-treated and Ad-LacZ or PBS-treated mice (P>0.05, data not shown), which indicated low general toxicity on Ad-IFNγ treatment.

Representative tumors harvested from each group were processed for histological analyses. Pathologic analysis (via H&E staining) found that there were large areas of necrosis in the tumor tissue treated with Ad-IFNγ, while few necrotic areas were observed in the Ad-LacZ-treated and PBS-treated tumors (data not shown). Cell proliferation was estimated by the immunohistochemical assessment of the nuclear antigen Ki-67. The results showed that Ki-67 was significantly lower in tumor tissue treated with Ad-IFNγ than those in Ad-LacZ group and PBS group. For apoptosis analysis, TUNEL positive ratios were significantly higher in Ad-IFNγ groups than those in Ad-LacZ group and PBS group. The representative pictures for Ki-67 and TUNEL staining in PBS group, Ad-LacZ group, and 1×10^9 pfu Ad-IFNγ group were shown in Figure 6.
Discussion
Interferon-γ (IFN-γ), as a multifunctional cytokine, exerts diverse biological functions related to host defense and immune regulation, such as inflammation, innate and acquired immunity, cell cycle and apoptosis [5]. IFN-γ plays a critical role in promoting protective host responses to tumors, which proposed mechanisms include, (a) anti-proliferative and pro-apoptotic actions, (b) anti-angiogenesis in tumors, and (c) promoting both the innate and adoptive immune responses against tumors [20,21].

Although IFN-γ has been investigated as a potential therapeutics for various types of tumors [6-13], the attempts to improve antitumor efficacy by increasing the dose or by repetitive continuous administration resulted in higher toxicity and low efficacy [22]. Considerable experimental data from our group and other investigators demonstrated that intratumoral IFN-γ gene transfer to achieve long-term, continuous locoregional exposure of IFN-γ is likely an appropriate approach for improving efficacy and reducing toxicity of IFN-γ [9,10,15,16,23,24].

In this study, we reported that adenovirus-mediated IFN-γ gene transfer (Ad-IFNγ) inhibited tumor growth of human nasopharyngeal carcinoma (NPC) cells (Figure 2 and Figure 5). Here the anti-NPC activity of IFN-γ included not only direct anti-proliferative and pro-apoptotic actions, also indirect mechanisms, such as immunomodulation and antiangiogenesis.

\[ \text{Ad-IFN}\gamma \text{efficiently expressed human IFN-γ in nasopharyngeal carcinoma cells in vitro and in vivo (Figure 1 and Figure 3), and exhibited strong antiproliferative effects (Figure 2A and Figure 6).} \]

According to previous reports, the antiproliferative mechanisms of IFN-γ seem to be cell type specific [9,25-29], either induction of cell

**Figure 2** Ad-IFNγ inhibited the proliferation of NPC cells by inducing G1 phase arrest in vitro. (A) Cell viability assay. CNE-2 or C666-1 cells were infected with Ad-IFNγ at different MOIs for 72 h, followed by MTT assay for cell viability. (B) Cell cycle distribution. CNE-2 or C666-1 cells were infected with Ad-IFNγ at 50 MOIs for 48 h, followed by flow cytometry analysis for DNA content and cell cycle distribution as described in “Materials and methods”. * p < 0.05, ** p < 0.01 while compared with corresponding cells infected with Ad-LacZ, N=3.
cycle arrest or apoptosis. Here we found that both G1 phase arrest and apoptosis contributed to Ad-IFNγ-mediated growth suppression in NPC cell lines (Figure 2B, Figure 3 and Figure 6), consistent with the report that the antiproliferative effects of minicircle-IFNγ on NPC cell lines could be attributed to G1 arrest and apoptosis [9]. The JAK/STAT pathway may be responsible for most biological effects mediated by IFN-γ [20,30], which regulates different cell cycle-associated proteins that control the G1-S checkpoint [25], or induce cell apoptosis through up-regulating the expression of various apoptosis-related proteins in different cell types [31-33].

Although antiproliferation and apoptosis induction were main effects of Ad-IFNγ-mediated anti-NPC in this study, antiangiogenesis may contribute also to growth inhibition of NPC xenografts in nude mice. Because the microvessel densities (MVDs) were found to be decreased in the xenografts treated with Ad-IFNγ compared with those treated with Ad-LacZ or PBS in our study (data not shown). Previous studies reveal that IP-10, an inhibitor of angiogenesis, could be induced by IFN-γ in endothelial cells and exert potent antiangiogenesis activity by inhibiting endothelial cell differentiation, motility and tube formation [34,35]. However, due to highly species specificity, human unlikely exerts direct effects on murine vascular
system, but likely acts on tumor cells to indirectly regulate angiogenesis. It was reported that IFN-\(\gamma\) acted as antiangiogenic cytokine by inhibiting the expression of angiogenic factors, such as VEGF or perlecan, in renal cell carcinoma [36], stromal fibroblasts [37], human cornea [38], and WiDr/HT29 colon carcinoma cells[39], or by inducing the expression of anti-angiogenic factor monokine induced by interferon-gamma (MIG, or CXCL9) in nonsmall cell lung carcinom [40]. Here we would like to make a hypothesis that IFN-\(\gamma\) may exert antiangiogenic effect in nude mice carrying NPC xenografts by regulating the expression of some angiogenic or antiangiogenic factors in NPC cells. This hypothesis, of course, needs to be tested and verified.

Human and murine IFN-\(\gamma\) display low level of sequence homology (only 40%) at protein level, which explains why the human and murine proteins display a strict species specificity in their ability to bind to and activate human and murine cells [41]. So human IFN-\(\gamma\) has less activity in the nude mouse host, though nude mice display potent macrophage and NK cell activity [42-44] and remain some basal T-cell function [45]. Direct immunomodulation on mouse immune system by human IFN-\(\gamma\) unlikely contributes to its antitumor effects. However, we cannot exclude

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**Figure 4** Ad-IFN\(\gamma\) effectively expressed human IFN-\(\gamma\) in NPC xenografts in nude mice. Mice were intratumorally injected with 2x10^9 pfu of Ad-IFN\(\gamma\), and then collected tumor tissues (A) and bloods (B) 1 day, 3, 5 and 7 days after Ad-IFN\(\gamma\) injection (mice before injection marked "day 0" as negative control, 3-4 mice per time point). Tumor tissues were made into homogenates in cold PBS, and then human IFN-\(\gamma\) concentrations in the tissue homogenates and blood plasmas were measured by using human interferon-\(\gamma\) ELISA kit (Boster, Wuhan, China).

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**Figure 5** Ad-IFN\(\gamma\) inhibited the growth of human NPC xenografts. When CNE-2 or C666-1 xenografts reached 4-5 mm diameter, mice were randomly assigned (7~8 mice per group) and treated by weekly intratumoral injection of 100 \(\mu\)L of PBS, Ad-LacZ or Ad-IFN\(\gamma\) (in 100 \(\mu\)L of PBS) or by daily intratumoral injection of 100 \(\mu\)L of normal saline (NS) or rhIFN-\(\gamma\) protein (in 100 \(\mu\)L of NS), respectively. (A, B) Tumor growth curves. Tumor size was measured every 4 days for CNE-2 xenografts (A) or 5 days for C666-1 (B), and the volume (V) was calculated according to the following formula: \(V = 0.52 \times \text{length} \times \text{width}^2\) [18,19]. (C, D) Tumor weights. After 3 weeks of treatment, mice with CNE-2 (C) or C666-1 (D) xenografts were sacrificed and tumor tissues were resected followed by weighing. (* \(p < 0.05\), ** \(p < 0.01\) compared with corresponding media).
the possibility that other indirect effects are involved in Ad-IFNγ-mediated antitumor immune response. Firstly, human IFN-γ could modify the expression of MHC and costimulatory molecules or cytokines, chemokines on human NPC cells [21,46], which may activate the residual immune system of nude mice. Secondly, adenoviral vector and heterogenous IFN-γ expression likely induce a nonspecific immune response directed by NK and/or microphage cells in nude mice [47].

Taken together, Ad-IFNγ displayed efficient anti-NPC activities by inhibiting tumor cell proliferation and induced cell apoptosis in this study. Additional indirect effects on antiangiogenesis and immunomodulatory may also be involved in this antitumor activity. So IFN-γ gene therapy by a replication defective adenovirus encoding the human IFN-γ (Ad-IFNγ) is likely a potential novel therapeutics on comprehensive therapy of nasopharyngeal carcinoma.

Nevertheless, recent reports have shown that the immune response activated by IFN-γ plays a dual role in cancer: It can not only suppress tumor growth by destroying cancer cells or inhibiting their outgrowth but also promote tumor progression either by selecting for tumor cells that are more fit to survive in an immunocompetent host or by establishing conditions within the tumor microenvironment that facilitate tumor outgrowth [48]. IFN-γ treatment is a double-edged sword whose anti- and protumorigenic activities are dependent on the cellular, microenvironmental, and/or molecular context [30]. Thereby, more investigations should be carried out to clarify the influences of cellular, microenvironmental, immunological and molecular events on the anti-NPC effects of Ad-IFNγ before its clinical application.

Conclusions
The results of the present study indicate that Ad-IFNγ likely performs potent anti-nasopharyngeal carcinoma effects by inhibiting cell proliferation, inducing G1 phase arrest and cell apoptosis. These findings have implications for the use of IFN-γ gene therapy mediated by replication defective adenoviral vector as a promising approach in the treatment of nasopharyngeal carcinoma.

Abbreviations
IFN-γ: Interferon-gamma; rhIFNγ: Recombinant human interferon-gamma; Ad-IFNγ: A replication defective adenovirus encoding the human IFN-γ; Ad-LacZ: A replication defective adenovirus encoding β-galactosidase; NPC: Nasopharyngeal carcinoma; RT-PCR: Reverse transcription-polymerase chain reaction; ELISA: Enzyme-linked immunosorbent assay; MT: 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; PI: Propidium iodide; DMEM: Dulbecco’s Modified Eagle Medium; PBS: Phosphate buffered saline; FBS: Fetal bovine serum; H&E: Hematoxylin and eosin; MOC: Multiplicities of infection; TUNEL: TdT-mediated dUTP-biotin nick end labeling; MVD: Microvessel density; MIG: Monokine induced by interferon-gamma; NK cell: Natural killer cell.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
RL carried out the cell proliferation and apoptosis analysis, collected and analyzed all of results, and designed and drafted the article. ZW, LCZ, and HL performed the animal experiments. LZ evaluated the expression of IFN-γ by RT-PCR and ELISA. PZ carried out immunohistochemical assay. HL and LX made animal model. HH performed and validated the statistical data. JW and WH conceived and coordinated the work, and helped in drafting the manuscript. All authors have read and approved the final manuscript.

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Figure 6 Histological analysis in C666-1 xenografts. C666-1 xenografts were resected at the end point of experiments, fixed and embedded in paraffin. Sections (5 μm thick) were mounted on Poly-L-Lysine treated slides. Upper, K667 expression assay. Sections were microwaved for 10 min for “antigen retrieval” and incubated with K667 antibody (1:200), followed by visualization with DAB using a Streptavidin-Horseradish Peroxidase kit. Lower, TUNEL labeling for apoptosis analysis. TUNEL labeling was performed with a fluorescent in situ cell death detection kit, according to the manufacturer’s instruction (Roche Applied Science, Germany).
