Hydroxyapatite nanoparticles inhibit the growth of human glioma cells in vitro and in vivo

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Abstract: Hydroxyapatite nanoparticles (nano-HAPs) have been reported to exhibit antitumor effects on various human cancers, but the effects of nano-HAPs on human glioma cells remain unclear. The aim of this study was to explore the inhibitory effect of nano-HAPs on the growth of human glioma U251 and SHG44 cells in vitro and in vivo. Nano-HAPs could inhibit the growth of U251 and SHG44 cells in a dose- and time-dependent manner, according to methyl thiazolotetrazolium assay and flow cytometry. Treated with 120 mg/L and 240 mg/L nano-HAPs for 48 hours, typical apoptotic morphological changes were noted under Hoechst staining and transmission electron microscopy. The tumor growth of cells was inhibited after the injection in vivo, and the related side effects significantly decreased in the nano-HAP-and-drug combination group. Because of the function of nano-HAPs, the expression of c-Met, SATB1, Ki-67, and bcl-2 protein decreased, and the expression of SLC22A18 and caspase-3 protein decreased noticeably. The findings indicate that nano-HAPs have an evident inhibitory action and induce apoptosis of human glioma cells in vitro and in vivo. In a drug combination, they can significantly reduce the adverse reaction related to the chemotherapeutic drug 1,3-bis(2-chloroethyl)-1-nitrosourea (BCNU).

Keywords: glioma, hydroxyapatite nanoparticles, growth mechanism

Introduction

Glioma is the most common, and the primary, malignant brain tumor of the central nervous system. In recent years, great progress has been made in diagnosing and treating glioma, but its recurrence after resection still makes long-term prognosis unsatisfactory.¹⁻³ Looking for new treatments and further studying its mechanism remain very important. With the development of nanotechnology, a new inorganic material, 1 nanoparticles (nano-HAPs), was found capable of inhibiting the proliferation of tumor cells.^{4,5} However, there is yet no report about the inhibitory effect of nano-HAPs on human glioma growth. In the current study, we sought to investigate the effect of nano-HAPs on the growth of human glioma U251 and SHG44 cells in vitro and in vivo and the mechanism involved, to provide a theoretical basis for clinical use.

Materials and methods

Cell culture

Human glioma U251 (human glioma cells, from Wuhan University of China, Wuhan, China) and SHG44 cells (human glioma cells, from Fourth Military Medical University of China, Xi'an, China) were cultured in RPMI-1640 (Gibco Life

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Technologies, Paisley, Scotland, UK) supplemented with 10% fetal bovine serum, 100 µg/mL penicillin, and 100 µg/mL streptomycin.

Measurement of cell growth

Cell proliferation was measured using the methyl thiazolotetrazolium (MTT) assay.^{6,7} Cells were seeded in plates at a density of 1×10^4 cells/well. When U251 and SHG44 cells were anchored to 96-well plates, the culture medium containing various nano-HAPs (East China University of Science and Technology Institute of Biomaterials, Shanghai, China) concentrations (0, 15, 30, 60, 120, and 240 mg/L) replaced the RPMI-1640 culture medium, the cells were incubated for another 24, 48, or 72 hours, and then 200 µL 5 mg/µL MTT (Sigma, St Louis, MO) in phosphate-buffered saline (PBS) was added to each well and incubated for 4 hours; the precipitate was solubilized in 100 µL 100% dimethylsulfoxide (Sigma) and shaken for 15 minutes. Absorbance values were determined using an enzyme-linked immunosorbent assay reader (Model 318; Shanghai Sunke Instruments Ltd., Shanghai, China) at 540 nm. Each assay was performed nine times.

Hoechst staining

Typical apoptotic morphological changes were observed under Hoechst staining.⁸ Cells treated with 0 mg/L, 120 mg/L, and 240 mg/L of nano-HAPs were harvested for 48 hours, fixed in 4% (v/v) paraformaldehyde for 10 minutes, and then incubated for 10 minutes with Hoechst 33342 dye (Sigma) (10 µg/mL). U251 and SHG44 cells were observed under an inverted fluorescence microscope (IX70; Olympus, Tokyo, Japan) after being washed with PBS.

Transmission electron-microscopic assay

Typical apoptotic morphological changes were also observed by transmission electron-microscopic assay.⁹ Cells were seeded in flasks. When U251 and SHG44 cells were anchored to plates, the different concentrations (0, 120, and 240 mg/L) of nano-HAPs were added, and the U251 and SHG44 cells were incubated for 48 hours; the specimens were postfixed in 1% osmium tetroxide/0.1 M sodium cacodylate buffer, dehydrated in a graded series of ethanol, and embedded after rinsing three times.¹⁰ The ultrathin sections of the specimens were stained with lead citrate and uranyl acetate, and observed with a transmission electron microscope. Each assay was performed in triplicate.

Measurement of apoptosis by flow cytometry

The suspended single cells of U251 and SHG44, treated 48 hours with nano-HAPs at various concentrations (0, 120, and 240 mg/L), were harvested. After washing with PBS fixed in 70% cold ethanol treated with 10 g/L RNase suspended and stained with 10 g/L propidium iodide. U251 and SHG44 cells were stained directly with PI at a concentration of 10 µg/mL and 2% Annexin-V-Fluos (Roche, Basel, Switzerland) in incubation buffer for 10 minutes. U251 and SHG44 cells were acquired with the FACS calibrator (BD) after setting the instrument with the controls (nontreated, stained cells), after two washes in PBS. In this experiment, U251 and SHG44 cells with early apoptotic signals, stained with annexin-V, and cells with late death signals, stained with PI, were considered and quantified, and the apoptotic cells were analyzed using CellQuest software.^{11,12} Each assay was performed in triplicate.

Murine xenograft model

Male, 4-to-6-week-old BALB/c athymic nude mice were subcutaneously injected with 2×10^6 human glioma cells. The 60 human glioma-bearing nude mice were randomly divided into five groups. Drug injection was through the tail vein every other day for 3 weeks. Group A (control group) was injected with 2 mL normal saline each time; Group B was injected with a nano-HAP solution of 25 mg/kg; Group C was injected with a nano-HAP solution of 50 mg/kg; Group D was injected with the chemotherapy drug 1,3-bis(2-chloroethyl)-1-nitrosourea (BCNU) at 25 mg/kg; and Group E was injected with a mixture of nano-HAP solution (25 mg/mL) and BCNU (25 mg/mL). The doses were both 25 mg/kg. Tumor diameters were measured at regular intervals with digital calipers, and the tumor volume in cubic millimeters was calculated by this formula: volume = (width)² × length/2. The animal experiments in this study were performed in compliance with the guidelines of the Institute for Medical School Institutes at Shanghai Jiao Tong University and Wuhan University.

Monitoring the growth state of human glioma-bearing nude mice

The human glioma-bearing nude mice were housed in a standard animal laboratory; the general growth-state changes of the animals, such as water-drinking, food intake, color pattern, nasal and oral discharge, and activity, were followed up every day. The human glioma-bearing nude mice were weighed, before feeding, every 3 days in the early morning.¹³

Transferase-mediated dUTP nick end labeling staining

Sections of glioma tissues of nude mice were deparaffinized in xylene, rehydrated, washed with PBS, and treated with 20 µg/mL of Proteinase K (Roche) for 10 minutes at room temperature.¹⁴ Terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling reactions were conducted using Apotag (Chemicon) according to the manufacturer's instructions, and detected using Cy3-anti-digoxigenin (Jackson). Microscopy was done on a Nikon E-600 microscope equipped with a QICam Fast 1394 camera (Q Imaging) and IVision imaging software (BioVision Technologies).

Western blotting analysis

The suspended single U251 and SHG44 cell treatment with nano-HAPs at various concentrations (0, 120, and 240 mg/L) for 48 hours were harvested. U251 and SHG44 cells were washed in PBS and lysed in buffer using the standard methods.^{15,16} The samples of glioma tissues of nude mice in the five groups were homogenized in an RIPA lysis buffer. Lysates were cleared by centrifugation (14,000 rpm) at 4°C for 30 minutes. Protein samples (approximately 40 µg) were separated by SDS-PAGE (15% gel), transferred to a PVDF membrane, and nonspecific binding sites blocked by incubation in 5% nonfat milk for 60 minutes. Membranes were incubated at 4°C overnight with anti-c-Met antibody (1:400 dilution), anti-SLC22A18 antibody (1:1000 dilution), anti-SATB1 antibody (1:200 dilution), anti-caspase-3 antibody (1:500 dilution), anti-bcl-2 antibody (1:300 dilution), or anti-Ki-67 antibody (1:100 dilution). Then the membranes were washed three times with TBST for 10 minutes and probed with the HRP-conjugated secondary antibody (at 1:2000 dilution) at room temperature for 30 minutes. The membrane was developed by an enhanced chemiluminescence system (ECL, Pierce) after being washed three times.

Statistical analysis

Statistical analyses were done and graphs created using the Statistical Package for the Social Sciences (version 12.0 for Windows; SPSS, Chicago, IL). Quantitative values were expressed as means ± standard deviation. Statistical differences between groups were examined using the Fisher's exact test. *P*-values less than 0.05 were considered statistically significant.

Results

Effect of nano-HAPs on the growth of cells

MTT assays demonstrated that cell viability was reduced after 24-hour incubation, and the viability of U251 and SHG44 cells decreased when the concentration and time period increased. There were significant differences in the viability of the U251 and SHG44 cells incubated with the various concentrations (0, 15, 30, 60, 120, and 240 mg/L) of nano-HAPs, and the viability of cells was also significantly inhibited by nano-HAPs as time increased. When cells were exposed to 240 mg/L HAP for 72 hours, the U251 and SHG44 cells' viability decreased greatly (to $31.73\% \pm 2.03\%$ and $32.28\% \pm 2.12\%$, respectively, compared with the control group's viability). ANOVA analysis and Dunnett's test revealed that nano-HAPs inhibited the U251 and SHG44 cells' growth in a dose- and time-dependent manner (Figure 1).

Effect of nano-HAPs on the morphology of cells

The nucleus in the control group was round and big, with a smooth nuclear membrane, free of condensation and fragmentation (Figure 2). But typical apoptotic morphological changes were found in the U251 and SHG44 cells 48 hours after treatment with 120 and 240 mg/L nano-HAPs. The higher the concentration of nano-HAPs, the more apoptotic cells there were. The changes included detachment, deformation, and shrinkage from the culture dishes. Chromatin margination and nuclear condensation were obvious from Hoechst staining (Figure 2A), and segmentation, nuclear

Figure 1 Effects of HAP nanoparticles on the viability of U251 and SHG44 cells determined by MTT assay.

Abbreviations: HAP, hydroxyapatite; MTT, methyl thiazolotetrazolium; SHG44, human glioma SHG44 cells; U251, human glioma U251 cells.

Figure 2 Effect of HAP nanoparticles on the morphologic changes of cells. (A) Hoechst 33342 staining. The apoptotic cells were marked by green fluorescence; scale bar, 20 μ m. (B) Transmission electron microscopy. **Notes:** Original magnification $\times 6000$; scale bar, 1 μ m. **Abbreviations:** HAP, hydroxyapatite; SHG44, human glioma SHG44 cells; U251, human glioma U251 cell.

condensation, and chromatin margination were observed by transmission electron microscopy (Figure 2B).

Effect of nano-HAPs on the apoptosis of cells in vitro

To quantitate nano-HAP-induced apoptotic cell death in U251 and SHG44 cells, approximately 1×10^6 U251 and SHG44 cells were double stained with Annexin-V-FITC and propidium iodide (PI) at different times, post-transfection. Apoptotic cell death was detected 48 hours after treatment with 120 and 240 mg/L nano-HAPs (Figure 3A). FACS analysis identified significantly higher numbers of apoptotic cells in nano-HAP-treated groups than in the control group (Figure 3B).

Effect of nano-HAPs on the expression of c-Met, SLC22A18, and SATB1 protein in U251 and SHG44 cells

Total protein was extracted from the U251 and SHG44 cells and analyzed by western blotting. The c-Met protein inhibition rates of 240 mg/L nano-HAPs in glioma U251 and SHG44 cells were, respectively, 82.65% and 76.04%, compared with the control group (Figure 4). Each treatment

Figure 3 FACS analysis of Annexin-V staining of cells after treatment with 120 and 240 mg/L HAP nanoparticles. (A) Representative FACS scatter plots of cells. (B) Percentages of apoptotic cells in cells after treatment. **Abbreviations:** HAP, hydroxyapatite; PI, propidium iodide; SHG44, human glioma SHG44 cells; U251, human glioma U251 cells.

group revealed that the expression of SLC22A18 increased, whereas the expression of SATB1 and Ki-67 decreased in glioma U251 and SHG44 cells (Figure 4).

Change of the growth state of human glioma-bearing nude mice

Before drug injection in each group, the weights between all the groups had no significant differences, the general state of health of the nude mice was very good, and activity and food intake were normal. The nude mice in the control group started to have symptoms of gradual failure two weeks after injection, the mice's food intake and activity worsened, their nasal and oral discharge increased, and their stools became loose. Their weight in the two nano-HAP-treated groups was higher than in the control group (Figure 5), and the time it took the two nano-HAP-treated groups to have symptoms of gradual failure was also longer than in the control group. At 1 week after injection, the weight of the nude mice was lower than that in the control group, and at 3 weeks after injection, the weight of the nude mice was lower than other groups. The activity and food intake of the nude mice decreased significantly after three drug injections. After these injec-

Figure 4 Effect of HAP nanoparticles on the expression of c-Met, SLC22A18 and SATB1 protein in U251 and SHG44 cells in vitro. (A and B) Representative images of western blotting analysis of c-Met, SLC22A18 and SATB1 expression in U251 and SHG44 cells. (C and D) Level of the c-Met, SLC22A18 and SATB1 protein expression in U251 and SHG44 cells.

Abbreviations: HAP, hydroxyapatite; U251, human glioma U251 cells; SHG44, human glioma SHG44 cells.

tions, the weight of the drug-combination group decreased, but there was no significant difference compared with the group receiving the chemotherapy drug BCNU.

Effects of the nano-HAPs on tumor growth in vivo

As shown in Figure 6, the control group's U251 and SHG44 xenograft tumors formed and grew very rapidly. However, xenograft tumor formation treated with both nano-HAPs and BCNU was significantly delayed. At the end of

Figure 5 Effects of HAP nanoparticles on weight of tumor-bearing nude mice.

Notes: Lane 1, the control group; lane 2, nano-HAP solution (25 mg/kg) group; lane 3, nano-HAP solution (50 mg/kg) group; lane 4, BCNU (25 mg/kg) group; lane 5, nano-HAP solution (25 mg/mL) and BCNU (25 mg/mL) group.

Abbreviations: Nano-HAPs, hydroxyapatite nanoparticles; BCNU, 1,3-bis(2-chloroethyl)-1-nitrosourea; SHG44, human glioma SHG44 cells; U251, human glioma U251 cells.

the experiment, the U251- and SHG44-cell tumors treated with both nano-HAPs and BCNU were significantly smaller than the tumors from the other groups.

Transferase-mediated dUTP nick end labeling assay

As shown in Figure 7, visualized using the DAB substrate, the brown color indicates apoptotic nuclei. Apoptosis was calculated as percentage of at least 1000 cells. After 28 days, the mixed nano-HAP and BCNU treatment of glioma U251 and SHG44 cells resulted in significantly higher apoptotic indexes (68.5% and 67.7%, respectively), compared to those of the other groups.

Caspase-3 and bcl-2 protein expression in tumor tissues

Total protein was exacted from tumor tissues from the five groups and analyzed by western blotting. Each treatment group revealed that the expression of caspase-3 increased in

Figure 6 Effect of HAP nanoparticles on tumor growth in vivo. (A and B) Subcutaneous tumor model. (C and D) Tumor growth curves of each group over 28 days. **Abbreviation:** Nano-HAPs, hydroxyapatite nanoparticles; BCNU, 1,3-bis(2-chloroethyl)-1-nitrosourea; U251, human glioma U251 cells; SHG44, human glioma SHG44 cells.

the glioma U251 and SHG44 cells of nude mice (Figure 8). The bcl-2 protein inhibition rates for nano-HAP plus BCNU, as compared with the control group, were 81.63% in the glioma U251 cells and 85.11% in the SHG44 cells (Figure 8).

Discussion

Hydroxyapatite has been widely used as a new biomaterial for oral cavity medicine and bone damage, is used as a medicine carrier in the body, and has good tissue compatibility both outside and inside the body.^{17–19} Nano-HAP has little excitatory effect on blood vessels: nano-HAP medicine could be delivered by intravenous injection.^{20,21} In this study,

Figure 7 In vivo assessment of apoptosis by TUNEL assay in the glioma tissues of nude mice at one month after treatment. (A) Representative images of TUNEL assay. Original magnification: $\times 400$. (B) Apoptotic index in the glioma U251 and SHG44 cell tissues of nude mice.

Abbreviations: TUNEL, transferase-mediated dUTP nick end labeling; U251, human glioma U251 cells.

nano-HAPs were synthesized by the sol-coagel method, which had a good dispersive effect, and the very uniform size of nano-HAPs was about 50 nm, which had the advantages of providing high surface energy and so on. Hydroxyapatite nanoparticles were found capable of inhibiting the proliferation of tumor cells.^{4,5}

In this study, it was found that different concentrations of nano-HAPs significantly inhibited the proliferation of human glioma U251 and SHG44 cells in a dose- and time-dependent manner, as observed using MTT assays. Typical apoptotic morphological changes such as nuclear chromatin fragmentation and condensation, cytoplasmic budding, cell shrinkage, and apoptotic body formation were observed after U251 and SHG44 cells were treated for 48 hours with HAP nanoparticles, as observed using Hoechst staining and transmission electron microscopy. Quantified by flow cytometry, the higher the concentration of HAP nanoparticles, the more apoptotic cells there were. Moreover, this study has shown that nano-HAPs had a significantly inhibitory effect on the growth of the human glioma U251 and SHG44 cells in nude mice, and induced apoptosis after its intravenous injection into same. The chemotherapy drug BCNU's inhibition of human glioma growth was greater than that of nano-HAP alone; however, adverse reaction to BCNU was also evident: the activity and food-intake of nude mice were evidently reduced after being continuously injected with BCNU. And their body weights were significantly lower than those

Figure 8 Effect of HAP nanoparticles on the expression of caspase-3 and bcl-2 protein in U251 and SHG44 cells in vivo. (A) Representative images of western blotting analysis of caspase-3 and bcl-2 expression. (B and C) Level of the caspase-3 and bcl-2 protein expression in tumors.

Notes: Lane 1, the control group; lane 2, nano-HAP solution (25 mg/kg) group; lane 3, nano-HAP solution (50 mg/kg) group; lane 4, BCNU (25 mg/kg) group; lane 5, nano-HAP solution (25 mg/mL) and BCNU (25 mg/mL) group.

Abbreviations: Nano-HAP, hydroxyapatite nanoparticles; U251, human glioma U251 cells; SHG44, human glioma SHG44 cells; BCNU, 1,3-bis(2-chloroethyl)-1-nitrosourea.

in the other groups. These findings showed no evidence of adverse bodily reactions caused by nano-HAPs. Therefore, the present research indicates that nano-HAPs can inhibit the growth of human glioma cells and induce apoptosis, in vitro and in vivo.

In this study, it was also shown that not only do nano-HAPs have an antitumor effect, but they also cooperate strongly with other antitumor drugs. In the drug combination group, the inhibition rates of human glioma U251 and SHG44 cells were the highest in the in vivo experiments. And the general state of

the mice in the two groups was quite different after their being injected with the same dosage of BCNU. Dynamic monitoring of body weight showed that the effect of BCNU on the physiological function of the mice in the drug combination group was evidently less than in BCNU-only group. The mice's general state in the former group was also obviously better than it was in the latter group. Combined with nano-HAPs (whose high surface energy and large surface area make them ideal drug carriers), BCNU could play an important role in tumor treatment.

We have previously found that HGF and its receptor, c-Met, played an important role in the formation, progression, and angiogenesis of glioma, that they could promote tumor proliferation and intratumoral microvascular formation, and that they were closely related to the prognosis of the patients.³ Antisense oligodeoxynucleotide targeting of the c-Met gene or c-Met-targeted RNA interference downregulated or inhibited c-Met expression, which inhibited U251 glioma cells growth and metastasis, induced cell apoptosis, and increased the sensitivity of human glioma cells to paclitaxel.^{22–25} Recently, we have found that SLC22A18 downregulation via promoter methylation was associated with the development and progression of glioma, that it represented a candidate biomarker, and that the elevated expression of SLC22A18 increased the sensitivity of U251 glioma cells to BCNU.^{1,2,26}

In recent years, SATB1 has attracted considerable attention because of its high expression in tumor tissues as a variety of malignancies, which suggests its important role in promoting tumor growth, invasion, and metastasis; it may also have potential value as a candidate for cancer therapy.^{27–29} Our results showed that the expression of c-Met, SATB1, and Ki-67 protein decreased and that SLC22A18 protein in glioma U251 and SHG44 cells increased after the cells were treated with various concentrations of hydroxyapatite nanoparticles in vitro. Owing to the function of nano-HAPs, the caspase-3 protein in tumor tissue increased, and the expression of bcl-2 protein clearly decreased. This might be the first report about the antitumor mechanism of nano-HAPs decreasing the expression of c-Met, SATB1, Ki-67, and bcl-2 protein while increasing the expression of SLC22A18 and caspase-3 protein. This could provide a basis for its clinical application.

Conclusion

In this study we have shown that not only do hydroxyapatite nanoparticles have an obvious antineoplastic function in vitro and in vivo, but also that they reduce the poisonous, adverse reactions to BCNU, strongly cooperate with certain other chemotherapy drugs, decrease the toxicity of certain

other chemotherapy drugs, and might become a new clinical antineoplastic drug. They might work by reducing the expression of c-Met, SATB1, Ki-67, and bcl-2 and by increasing the expression of SLC22A18 and caspase-3.

Acknowledgments

The authors would like to thank East China University of Science and Technology Institute of Biomaterials for providing the hydroxyapatite nanoparticles. This work was supported by grants from the Innovation Program of Shanghai Municipal Education Commission (12YZ046) and the Natural Science Foundation of China (30901535).

Disclosure

The authors report no conflicts of interest in this work.

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