ORIGINAL RESEARCH

The synergistic radiosensitizing effect of tirapazamine-conjugated gold nanoparticles on human hepatoma HepG2 cells under X-ray irradiation

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Abstract: Reductive drug-functionalized gold nanoparticles (AuNPs) have been proposed to enhance the damage of X-rays to cells through improving hydroxyl radical production by secondary electrons. In this work, polyethylene glycol-capped AuNPs were conjugated with tirapazamine (TPZ) moiety, and then thioctyl TPZ (TPZs)-modified AuNPs (TPZs-AuNPs) were synthesized. The TPZs-AuNPs were characterized by transmission electron microscopy, ultraviolet-visible spectra, dynamic light scattering, and inductively coupled plasma mass spectrometry to have a size of 16.6±2.1 nm in diameter and a TPZs/AuNPs ratio of ~700:1. In contrast with PEGylated AuNPs, the as-synthesized TPZs-AuNPs exhibited 20% increment in hydroxyl radical production in water at 2.0 Gy, and 19% increase in sensitizer enhancement ratio at 10% survival fraction for human hepatoma HepG2 cells under X-ray irradiation. The production of reactive oxygen species in HepG2 cells exposed to X-rays in vitro demonstrated a synergistic radiosensitizing effect of AuNPs and TPZ moiety. Thus, the reductive drug-conjugated TPZs-AuNPs as a kind of AuNP radiosensitizer with low gold loading provide a new strategy for enhancing the efficacy of radiation therapy.

Keywords: AuNPs, radiation enhancement, synergistic effect, human hepatoma cells, hydroxyl radical production

Introduction

The estimates of the worldwide incidence and mortality of all cancers have showed that there were 14.1 million new cases and 8.2 million deaths in 2012.[1] Cancer now causes more deaths than all coronary heart diseases or strokes, and has apparently become the leading threat to human health. Radiation therapy is one of the primary modalities of cancer treatment, which provide relieve from symptoms of advanced or metastatic diseases, and is the need of >50% of cancer patients.[2] The major challenge of radiation therapy is maximizing the radiation dose delivered to malignant tissues while minimizing damage to the surrounding normal tissues.[3–5] Improving the efficacy of radiation therapy, like enhancing the radiation effects by increasing the dose specifically to the tumor cells, can benefit a significant number of patients.[6,7]

One way of improving the efficacy of radiation therapy is by expanding photon/Auger electron induction in the physical/chemical stage during irradiation, which will lead to enhancement in reactive oxygen species (ROS) production and subsequently the radiobiological response.[8] The interaction of ionizing radiation with high-Z
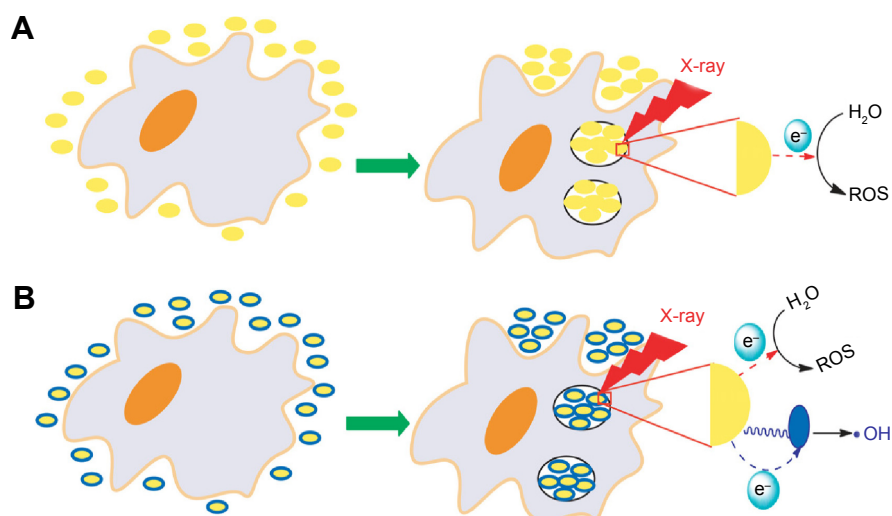
Figure 1 Radiosensitizing principle of drug-modified gold nanoparticles.
Notes: Radiosensitizing principle of nanoparticles based on (**A**) gold- and (**B**) drug-mediated radiosensitization.
Abbreviation: ROS, reactive oxygen species.

(atomic number) materials may result in enhancement of Auger electron emission. When iodine was first used to sensitize cultured cells by Matsudaira et al in 1980, the concept of using high-Z materials to increase the radiation dose given to a tumor during radiation therapy was proposed.[9] Noble metal nanoparticles have had diverse applications in cancer therapy, such as diagnostics[10,11] and treatment reagents,[12] in the last decade. Since the in vivo study on the radiation-enhancing effect of gold nanoparticles (AuNPs) was presented by Hainfeld et al in 2004,[13] gold,[14–16] platinum,[17] and bismuth[18,19] nanoparticles have been proposed as radiation sensitizers under irradiation of X-rays, γ-rays, and electron beams in extensive investigations. Owing to high biocompatibility, facile conjugation with tumor-targeting agents, and high X-ray absorption coefficients, AuNPs have become the primary option for radiation enhancement.[20–22]

In this study, we investigated the radiosensitizing effect of AuNPs under high-linear energy transfer particle radiation as well as low-linear energy transfer conventional radiation.[23,24] AuNPs internalized by cells were found in endosomes/lysosomes instead of nuclei. As a possible mechanism, the dose enhancement in the presence of AuNPs could be attributed to additional photoelectrons and Auger electrons generated from nanoparticles under X-ray irradiation. Photoelectrons are highly energetic, have a long range (up to hundreds of micrometers) in water, and deposit a small fraction of their energy near the nanoparticles. Auger electrons have lower energy and shorter range (<1 μm), allowing most of the energy to be deposited around the nanoparticles in nanoscale.[25] Energy deposition by electrons could result in water hydrolysis

and production of ROS (Figure 1A). We suppose that there is a way of improving the production of free hydroxyl radical via chemical reduction by secondary electrons. The presence of reductive drug-modified AuNPs, which would not only produce ROS via water hydrolysis but also undergo drug reduction by secondary electrons generated from the irradiated AuNPs, might also enhance hydroxyl radical production, leading to an increase in cellular radiosensitivity (Figure 1B).

Tirapazamine (TPZ), a hypoxia-selective radiosensitizer, can be reduced by reductase in cells to produce hydroxyl radical.[26,27] In this study, we designed thioctyl TPZ (TPZs)-modified AuNPs (TPZs-AuNPs) by conjugating TPZ moiety on the surface of AuNPs. With the enhanced permeability and retention effect, TPZs-AuNPs offer a high concentration of TPZ in the microenvironment of a tumor. The TPZ moiety would undergo reduction reaction by secondary electrons generated from AuNPs under X-ray irradiation and amplify the production of hydroxyl radicals, providing an effective way of improving the radiation-enhancing effect of AuNPs. After the successful preparation of TPZs-AuNPs, we investigated their radiosensitizing effect on human hepatoma HepG2 cells exposed to X-rays.

Materials and methods
Reagents and materials
Chloroauric acid trihydrate ($HAuCl_4 \cdot 3H_2O$, 98%) was purchased from Shaanxi Kaida Chemical Engineering Co. (Shaanxi, People's Republic of China). Sodium citrate dihydrate (99%) was obtained from Shanghai Lingfeng Chemical Reagent Co. (Shanghai, People's Republic of China).

Thioctic acid (98%), coumarin-3-carboxylic acid (3-CCA, 97%), *N*-ethyl-*N*′-(3-dimethylaminopropyl) carbodiimide hydrochloride (99%), and *N*-hydroxybenzotrizole (99%) were purchased from J&K Scientific Ltd (Beijing, People's Republic of China). Cyanamide (95%) and *o*-nitroaniline (98%) were purchased from Energy Chemical (Shanghai, People's Republic of China). TPZ was prepared according to the reported procedure.[28] TPZs was prepared by condensation of thioctic acid with TPZ (Supplementary materials). Polyethylene glycol-capped AuNPs (PEG-AuNPs) were prepared by following the reported procedure.[29] TPZs-AuNPs were prepared by adding a TPZs methanol solution into a PEG-AuNPs aqueous solution.

Human hepatoma HepG2 cells were obtained from the Cell Bank of Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences (Shanghai, People's Republic of China). Rosewell Park Memorial Institute (RPMI)-1640 medium, 10% fetal bovine serum (FBS), trypsin, 100 U/mL penicillin, and 100 μg/mL streptomycin were purchased from Hyclone Laboratories, Inc. (Logan, UT, USA). 3-(4,5-Dimethyl-2-thiazolyl)-2,5-diphenyl-2-*H*-tetrazolium bromide (MTT) was purchased from Spectrum Chemical Mfg. Corp. (New Brunswick, NJ, USA). All reagents were used as received. Monoclonal antibodies against γ-H2AX were obtained from Abcam (Cambridge, MA, USA). Rabbit anti-mouse IgG-R antibody was purchased from Santa Cruz Biotechnology Inc. (Dallas, TX, USA). 4′,6-Diamidino-2-phenylindole (DAPI) was purchased from Vector Laboratories (Burlingame, CA, USA). 2′,7′-Dichlorofluorescein diacetate (DCFH-DA, 98%) was bought from Biyuntian Biotechnology Institute (Jiangsu, People's Republic of China). Water (18.2 MΩ) used in the experiments was obtained from an ultrapure water system (Heal Force Co., Hong Kong, People's Republic of China).

The surface plasemonic resonance of AuNPs was recorded with ultraviolet-visible (UV-vis) spectroscopy (Jasco, Japan) dual-beam spectrophotometer (Model V570) from 200 nm to 800 nm with a resolution of 2.0 nm. Transmission electron microscopy (TEM) measurements were performed on a Tecnai TF20 transmission electron microscope (FEI Co., Hillsboro, OR, USA) at 200 kV. The concentrations of AuNPs in solution were measured using IRIS ER/S inductively coupled plasma atomic emission spectrometer (ICP-AES; Thermo Jarrell Ash Corp., Franklin, MA, USA) after AuNPs had been dissolved in aqua regia. The zeta potentials of AuNPs were measured using electrophoretic light scattering, and the hydrodynamic diameter of AuNPs was determined by dynamic light scattering on Zetasizer Nano range (Malvern Instruments, Malvern, UK).

The samples were irradiated with X-rays from an RX-650 X-ray source (Faxitron X-ray Co., Tucson, AZ, USA). The dose rate was measured using a Markus ionization chamber (PTW-Freiburg, Freiburg, Germany) connected to a dosimeter (UNIDOS; PTW-Freiburg). The cell number was counted using a particle counter (Z1; Beckman Coulter Inc., Brea, CA, USA). Fluorescence of the irradiated aqueous solutions at 442 nm was recorded on a microplate reader (Infinite F200/M200; TECAN Co., Männedorf, Switzerland) excited at 395 nm. Fluorescent images of cells were collected on a fluorescence microscope (BX51; Olympus Corporation, Tokyo, Japan).

Cell culture and cell viability assay

Human hepatoma HepG2 cells were cultured in RPMI-1640 medium supplemented with 10% FBS, 100 U/mL penicillin, and 100 μg/mL streptomycin. Cultured HepG2 cells were kept at 37°C in a 5% CO_2 humidified incubator. To quantify cell uptake of AuNPs, the cells were incubated in flasks with 0–20.0 μg/mL of AuNPs for 24 hours. For X-ray irradiation, the cells were cultured in 35 mm Petri dishes with/without AuNPs at a concentration of 10.0 μg/mL for 24 hours. Prior to irradiation and uptake measurement, the cells incubated with AuNPs were washed using phosphate-buffered saline (PBS) to remove the noninternalized AuNPs. Fresh medium with FBS and antibiotics was added to the Petri dishes.

To estimate the cytotoxicity of functionalized AuNPs to cells, cell viability was determined with the MTT assay.[30] Briefly, cells were plated into 96-well tissue culture dishes at a density of 1×10^4 cells/well in 180 μL medium. After plating, HepG2 cells were incubated with AuNPs at a concentration of 1.0–10.0 μg/mL for 24–72 hours, and then 20 μL of MTT (5.0 mg/mL) was added to each well. After incubation at 37°C for 4 hours, the supernatants were removed, and formazan crystals were added in 150 μL dimethyl sulfoxide (DMSO) solution. The well plates were then read on a microplate reader at 570 nm. The viability change of AuNPs-treated cells was expressed as a percentage compared to the cells without AuNPs treatment, and control cells were considered to have a viability value of 100%. Each experiment was conducted in quintuplicate.

Irradiation

For solution and cell exposure, samples were set horizontally to be irradiated. The X-ray irradiation was delivered at 50 kVp and 5 mA, and the dose rate was 0.50 Gy/min. The dose of X-rays delivered to a sample was determined via controlling the irradiation time. For cell exposure, cell

samples in 35 mm Petri dishes were irradiated at doses of 1.0, 2.0, 3.0, 4.0, and 6.0 Gy. All irradiations were performed at room temperature.

Measurement of hydroxyl radical and ROS

Hydroxyl radical production in solution was investigated by using 3-CCA as probe.[31] Following Cheng et al's procedure,[32] 3-CCA was dissolved in PBS (pH 7.4). The solution was filtered by a 0.22 µm membrane filter, and DMSO was added to yield a 3-CCA/DMSO solution, which was then diluted ten times with PBS. PEG-AuNPs and TPZs-AuNPs stock solutions were mixed with the diluted 3-CCA/DMSO solution so as to achieve the same 3-CCA/DMSO molar concentrations and gold concentration (10.0 µg/mL) in samples. An equal volume (365 µL) of the samples was added into different wells of 96-well culture plates and kept away from light. Half an hour after irradiation, the AuNPs in the samples were removed by centrifugation to avoid the fluorescence interference of AuNPs. Fluorescence intensities of the samples were measured to determine the produced 7-hydroxyl-coumarin-3-carboxylic acid, which indicates the yield of hydroxyl radicals.

Intracelluar ROS production was measured by using DCFH-DA, a fluorogenic dye for hydroxyl, peroxyl, and other ROS produced within cells.[33–35] All treatments were carried out in quintuplicate. HepG2 cells were seeded either in 35 mm Petri dishes for qualitative measurements or in 96-well tissue culture plates for quantitative measurements. All samples were cocultured with 10.0 µg/mL AuNPs for 24 hours. After treatment, cells were incubated with 10.0 µM DCFH-DA for 30 minutes followed by irradiation. The cells were then observed under a fluorescence microscope, and all pictures were taken at exactly the same exposure time of 0.2 seconds. For quantitative measurement, the fluorescence intensity at 525 nm was detected using a fluorescence spectrometer excited at 488 nm, and then normalized by the MTT assay. Each experiment was performed in quintuplicate, and the result is represented as the mean ± standard deviation (SD).

Clonogenic survival assay

Cells pretreated with AuNPs for 24 hours were irradiated with X-rays. Immediately after irradiation, the cells were collected by trypsinization, and then were diluted and seeded in 60 mm Petri dishes at an appropriate density for each treatment. The cells were cultured in RPMI-1640 medium for 15 days, and then fixed with alcohol and stained with crystal violet. Colonies containing >50 cells were counted as survivors under a stereomicroscope. Each experiment was performed in quadruplicate, and the result is expressed as the mean ± SD.

DNA damage measurement

Cells (1×10^5) were plated on coverslips and allowed 24 hours to attach, and then were pretreated with 10.0 µg/mL AuNPs for 24 hours followed by X-ray irradiation. At different times post-irradiation, the cells were fixed with 4% paraformaldehyde for 10 minutes. The fixed cells were permeabilized with 0.1% Triton X-100 and subsequently blocked with 5% bovine serum albumin at room temperature for 1 hour. The cells were then incubated overnight at 4°C with primary monoclonal antibodies against γ-H2AX (200-fold dilution). After primary antibody incubation, the cells were washed with PBS and incubated with rabbit anti-mouse IgG-R secondary antibody (300-fold dilution) for 1 hour at room temperature. Nuclei were counterstained with mounting medium DAPI at a concentration of 1.5 µg/mL. Finally, γ-H2AX foci were detected with a fluorescence microscope.

Cell uptake of nanoparticles

After cell incubation with AuNPs for 24 hours, the medium was removed. HepG2 cells were washed with PBS and harvested using trypsin. Then, the cell number was counted. Finally, the cell suspension was centrifuged for 20 minutes at 12,000 rpm. The pellets were dissolved in aqua regia and diluted with ultrapure water. The concentration of gold was determined by means of ICP-AES. The results presented were normalized by the cell numbers. Data are represented as the mean ± SD of the triplicate samples.

Results and discussion

PEG-AuNPs were reported with excellent colloidal stability in different in vitro environments.[21,36,37] We fabricated PEG-AuNPs by place-exchange reaction to synthesize aqueous TPZs-AuNPs (Figure 2). Methoxylpolyethyleneglycolthiol was adhered to AuNP surface, and TPZs was anchored on the surface via Au–S bonds. As-synthesized TPZs-AuNPs were spherical, relatively monodispersed, and exhibited aqueous colloidal stability for several months. TEM micrographs of the synthesized PEG-AuNPs and TPZs-AuNPs are shown in Figure 3A and B. No change in average size was observed after functionalization. The average particle diameter of TPZs-AuNPs was 16.6±2.1 nm, as derived from the TEM micrographs (Figure 3C). In the UV-vis spectra of TPZs-AuNPs, the characteristic absorption at 526 nm and a
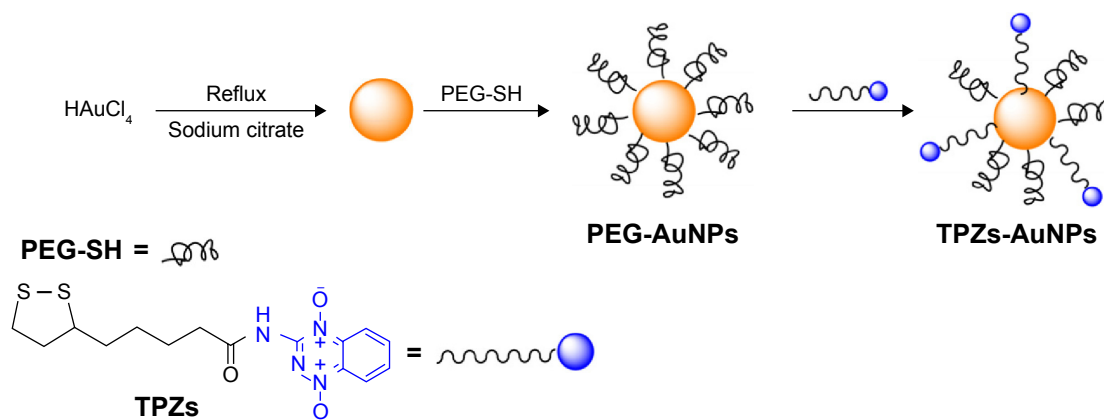
Figure 2 Preparation of TPZs-AuNPs.
Abbreviations: TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles; PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs, thioctyl tirapazamine.

shoulder band at 264 nm were in accordance with the surface plasmonic resonance of PEG-AuNPs at 522 nm and the characteristic band of TPZs at 270 nm, respectively (Figure 3D). ICP-AES was used to measure the Au and S concentrations in PEG-AuNPs and TPZs-AuNPs. In TPZs-AuNPs, the ratio of TPZs to AuNPs was predicted to be 700:1 by ICP-AES, which was close to the ratio of 732:1 obtained by UV-vis measurement (Figure S1). Therefore, the concentration of TPZs was ~3.5×10^{-7} M in a 10.0 μg/mL TPZs-AuNPs solution. The modification of nanoparticles always leads

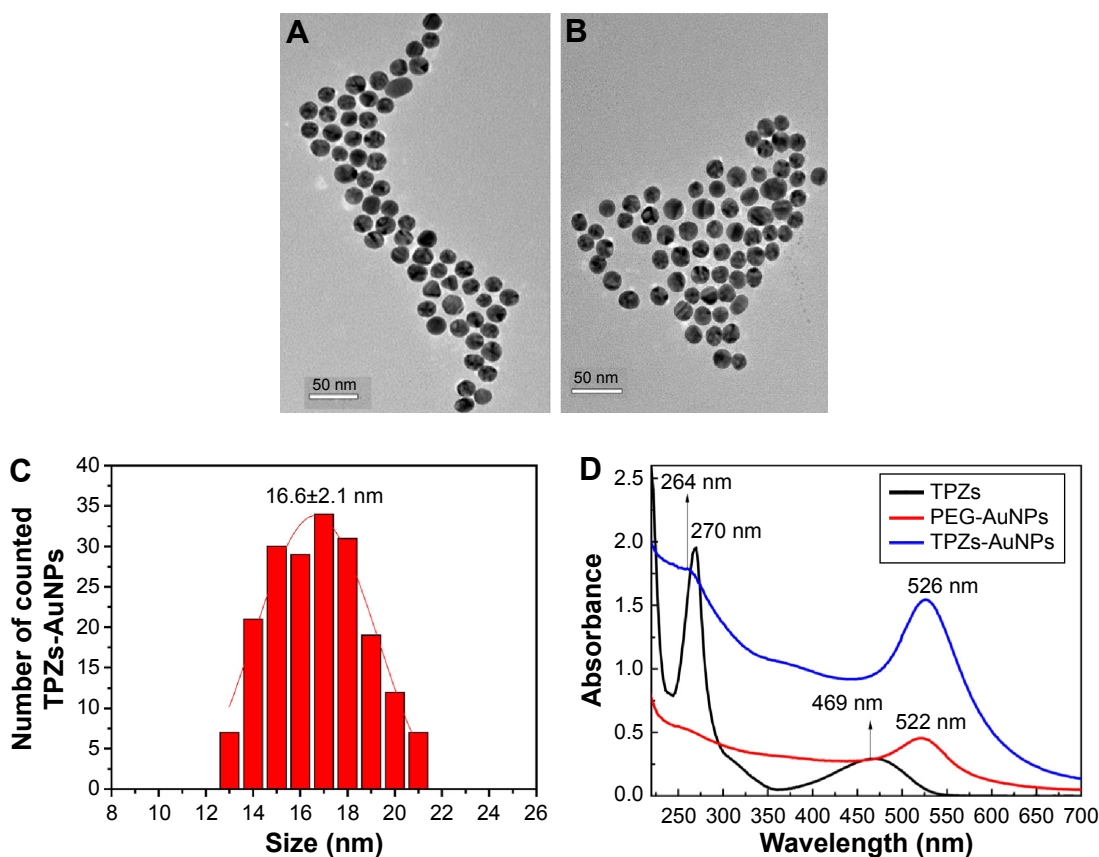**Figure 3** Characterization of gold nanoparticles.
Notes: The TEM image of (**A**) PEG-AuNPs and (**B**) TPZs-AuNPs. (**C**) The size distribution of TPZs-AuNPs. (**D**) The UV-vis spectra of PEG-AuNPs and TPZs-AuNPs in this study.
Abbreviations: TEM, transmission electron microscopy; PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles; TPZs, thioctyl tirapazamine; UV-vis, ultraviolet-visible.

to changes in surface chemical activity and hydrodynamic diameter. The zeta potential of 16.6 nm naked AuNPs was −45.4 mV, which was due to citric acid coating and negative charge on the surface. PEG-SH coating and TPZs conjugation induced the increase of zeta potential to −30.3 and −13.9 mV, respectively. The hydrodynamic diameter of TPZs-AuNPs was 37.8 nm, which was 2.5 nm less than the size of 40.3 nm for PEG-AuNPs, as determined by the dynamic light scattering measurements after the conjugation of TPZs on the PEG-AuNPs surface.

With TPZs-AuNPs in hand, we investigated their radiation-enhancing effect through hydroxyl radical production in aqueous solution exposed to X-rays. The radiation enhancements of PEG-AuNPs and TPZs-AuNPs in solutions were evaluated by hydroxyl radical production using 3-CCA as probe following the procedure presented by Cheng et al.[32] The fluorescence intensity of irradiated solution increased along with the radiation dose (Figure S2). The radiation enhancement ratio could be calculated using the equation $E_r = F_i/F_0$, where E_r, F_i, and F_0 stand for enhancement ratio, fluorescence intensity of the irradiated solution with or without AuNPs, and fluorescence intensity of the nonirradiated 3-CCA solution without AuNPs (control), respectively. The dependence of the radiation enhancement ratios of PEG-AuNPs and TPZs-AuNPs on the radiation dose is shown in Figure 4A. Clearly, the fluorescence intensities did not change when PEG-AuNPs and TPZs-AuNPs were added to 3-CCA solutions. When the solutions were irradiated with X-rays of 2.0 Gy, the enhancement ratio increased to 1.29 for PEG-AuNPs, which was slightly higher than 1.27 for

X-rays alone but was 21% lower than 1.56 for TPZs-AuNPs. The same phenomenon was observed when the exposure dose increased to 4.0 Gy. TPZs-AuNPs showed an enhancement ratio of 2.56 in terms of hydroxyl radical production, which was 23.7% higher than 2.07 for PEG-AuNPs. The presence of 50 mM DMSO in solution resulted in hydroxyl radical quenching, and thus the sharp decrease in enhancement ratio to the control level. These results demonstrated that TPZs-AuNPs presented 20% more radiation enhancement in terms of hydroxyl radical production in aqueous solution induced by X-rays than PEG-AuNPs at the same gold concentration.

Since TPZ moiety might also induce hydroxyl radical production under X-ray irradiation, we investigated the hydroxyl radical production of TPZ solution exposed to X-rays. No enhancement was observed even up to the concentration of 2.5 μM (Figure S3), which was approximately sixfold higher than the TPZ moiety concentration in 10.0 μg/mL TPZs-AuNPs solution. Furthermore, we studied the cytotoxicity of TPZ and TPZs (Figure S4A and B) to HepG2 cells. After coculturing with 0–1.0 μM TPZ and TPZs for 24 hours, no apparent decrease in cell viability was observed. When the coculture time was prolonged to 48 and 72 hours, obvious cell viability inhibition of TPZ and TPZs was observed. This might be due to the accumulation of TPZ moiety in cells with increasing the coculture time, leading to endogenous hydroxyl radical improvement and subsequently cell apoptosis. Besides free TPZs moiety, we further studied the cytotoxicity of TPZs-AuNPs as well as PEG-AuNPs to HepG2 cells. As shown in Figure 4B, the viability of HepG2 cells
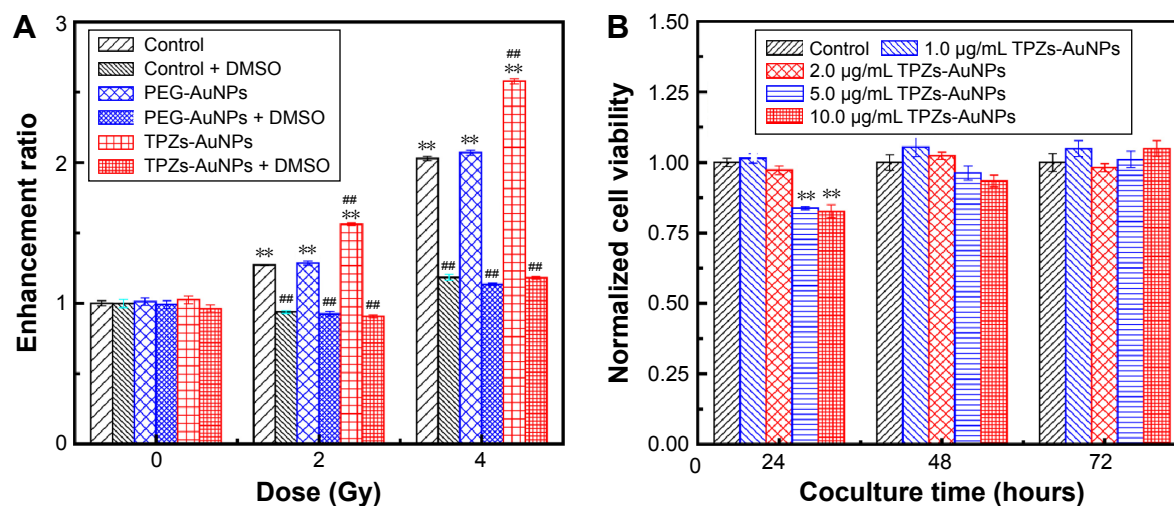

Figure 4 Radiation enhancement in solution and cell viability of gold nanoparticles.
Notes: (**A**) The radiation enhancement ratio of PEG-AuNPs and TPZs-AuNPs concerning hydroxyl radical production in solutions exposed to X-rays (50 kVp) at the gold concentration of 10.0 μg/mL. (**B**) The normalized viability of HepG2 cells cocultured with TPZs-AuNPs at various concentrations for 24, 48, and 72 hours. ******$P<0.01$ versus control group without irradiation and **##**$P<0.01$ versus control group with X-ray alone, determined by the two-tailed Student's *t*-test.
Abbreviations: PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles; DMSO, dimethyl sulfoxide.

did not decrease when cocultured with 0–2.0 µg/mL TPZs-AuNPs for 24 hours. The cell viability reduced to ~83% after coculturing with 10.0 µg/mL TPZs-AuNPs for 24 hours but recovered to 93% and 100% when the coculture time was prolonged to 48 and 72 hours, respectively. In addition, the viability of HepG2 cells decreased by 2%–8% after coculturing with 0–10.0 µg/mL PEG-AuNPs for 24 hours. Further slight decrement in cell viability was observed for 48- and 72-hour coculture times (Figure S4C). Clearly, PEG-AuNPs and TPZs-AuNPs did not affect the viability of HepG2 cells significantly. When TPZ moiety was conjugated to the surface of AuNPs, the cytotoxicity of TPZ moiety was reduced.

The low cytotoxicity of TPZs-AuNPs and good stability encouraged us to investigate their intracellular ROS enhancement. ROS produced in normal cellular environments are essential for life and become harmful when overproduced. The intracellular ROS were visualized via DCFH-DA probes for HepG2 cells with or without 10.0 µg/mL AuNPs as shown in Figure 5A. Without X-ray irradiation, little visible fluorescence was observed after cells were treated with AuNPs for

20 minutes. After irradiation, a strong fluorescence was emitted by the AuNPs-treated cells. The enhancement ratio of ROS upon irradiation is shown in Figure 5B. Without irradiation, the ROS production in the presence of TPZs-AuNPs was ~12% higher than the control, which was consistent with the cytotoxicity result. No significant enhancement of ROS production was observed in HepG2 cells with PEG-AuNPs under X-ray irradiation compared to the control. The presence of TPZs-AuNPs led to ~23% higher fluorescence intensity than PEG-AuNPs at the dose of 2.0 Gy. When the radiation dose reached 4.0 Gy, the intracellular ROS yield increased to 2.6-fold in the presence of PEG-AuNPs, which was only 4% higher than the control without AuNPs. In contrast, the yield increased to 3.1-fold in the presence of TPZs-AuNPs, a 19.2% increment in ROS production than PEG-AuNPs. These results demonstrated that the presence of TPZs-AuNPs contributed to ~20% increment in ROS production in cells compared with PEG-AuNPs. This is in accordance with the result of hydroxyl radical production, indicating that the intracellular ROS increment mainly came from the increase of hydroxyl radical yield.
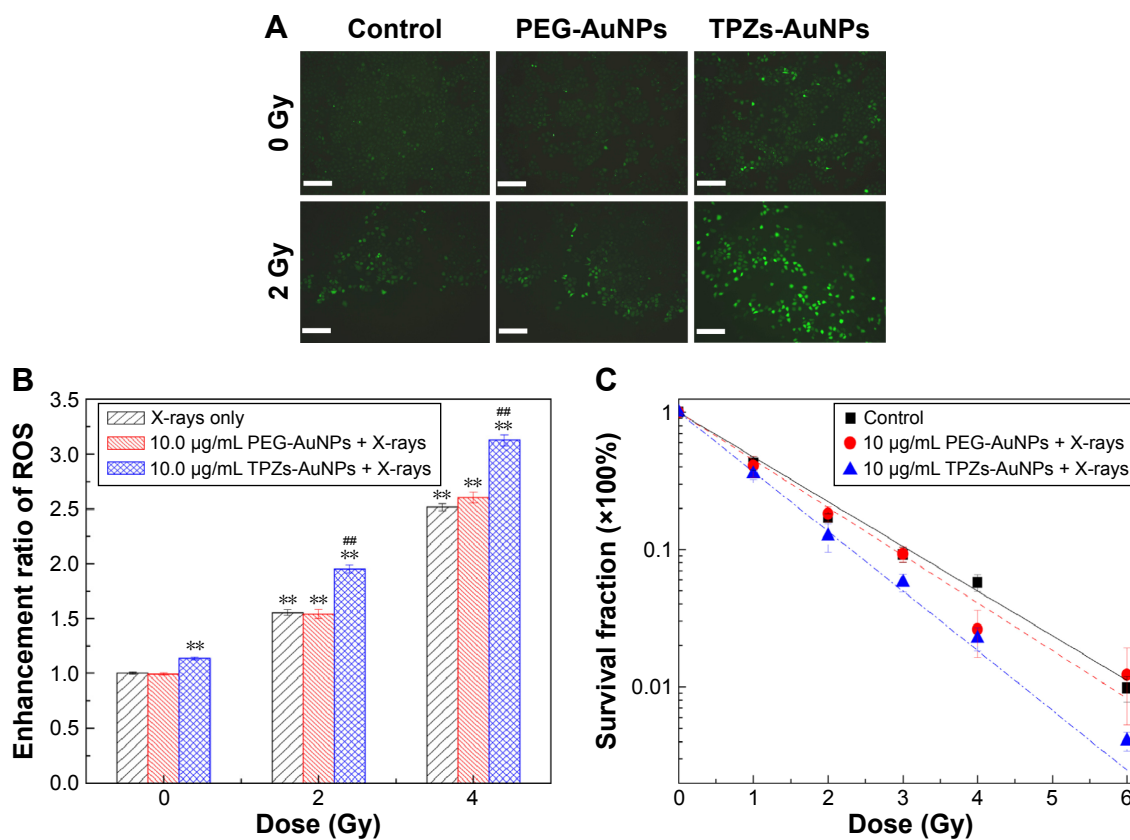

Figure 5 Radiation enhancement in cells.
Notes: (**A**) The fluorescence images of ROS with DCFH-DA in HepG2 cells after X-ray irradiation in the presence of PEG-AuNPs and TPZs-AuNPs. All scale bars represent 200 µm. (**B**) The enhancement ratio of ROS in HepG2 cells after X-ray irradiation in the presence of PEG-AuNPs and TPZs-AuNPs compared to control. (**C**) The survival curves of HepG2 cells in the presence of PEG-AuNPs and TPZs-AuNPs after exposure to 50 kVp X-rays. *******P*<0.01 versus control group and *##P*<0.01 versus X-rays alone, determined by the two-tailed Student's *t*-test.
Abbreviations: ROS, reactive oxygen species; DCFH-DA, 2′,7′-dichlorofluorescein diacetate; PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles.

The satisfactory result in ROS enhancement in HepG2 cells induced by X-rays in combination with TPZs-AuNPs prompted us to investigate the subsequent cell survival effect under X-ray irradiation combined with AuNPs. To test the role of TPZ-AuNPs in radiation-induced cytotoxicity enhancement, cell clonogenic survival experiments were carried out. The survival curves of HepG2 cells in the presence of AuNPs irradiated with X-rays are displayed in Figure 5C. Clearly, the cell survival fraction decreased exponentially with increase in the radiation dose. The addition of 10.0 µg/mL TPZs-AuNPs led to a sharper decrease in survival fraction than PEG-AuNPs along with a decrease in the radiation dose. The doses of X-rays required to obtain 10% cell survival fraction for HepG2 cells were 3.10, 2.92, and 2.32 Gy for the control without AuNPs, PEG-AuNPs, and TPZs-AuNPs, respectively. The sensitizer enhancement ratios of PEG-AuNPs and TPZ-AuNPs were 1.05 and 1.25 at the concentration of 10.0 µg/mL, respectively (Table 1). Compared to PEG-AuNPs, TPZs-AuNPs showed an increase of 19% in sensitizer enhancement ratio at 10% survival level for HepG2 cells under X-ray irradiation. This result is in accordance with the enhancement in hydroxyl radical production and intracellular ROS. As mentioned earlier, the addition of DMSO led to the hydroxyl radical quenching. Consequently, we deduced that the survival fraction decrease of HepG2 cells with TPZs-AuNPs under X-ray irradiation was mainly due to the increase in hydroxyl radical production. Therefore, the TPZ moiety on AuNPs surface contributed to the increase in ROS production and ultimately decrease in cell survival.

Cell death or apoptosis induced by ionizing radiations might be mainly attributed to the nucleus damages such as base damages, DNA–protein cross-links, and DNA double-strand breaks (DSBs).[38] DNA DSBs in HepG2 cells exposed to X-rays in the presence of TPZs-AuNPs and PEG-AuNPs were then investigated to clarify whether the increased cell lethality was derived from nucleus or cytoplasm damages.

Table 1 Required dose for 10% survival level and sensitizer enhancement ratio at 10% survival level for HepG2 cells cocultured with PEG-AuNPs and TPZ-AuNPs at a concentration of 10.0 µg/mL

Items		Control	PEG-AuNPs	TPZs-AuNPs
SF =10%	Dose required for X-rays (Gy)	3.10	2.92	2.32
	Sensitizer enhancement ratio	1.00	1.05	1.25

Abbreviations: PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles; SF, survival fraction.

γ-H2AX on Ser139 is one of the key events of DNA damage response and considered to be a marker of DSBs.[39] Figure 6A shows the results of γ-H2AX foci detection acquired in this study for HepG2 cells in the presence of PEG-AuNPs and TPZs-AuNPs. The number of γ-H2AX foci in HepG2 cells was 3.2±1.1, while it slightly increased to 4.1±1.8 and 5.1±1.5 when cells were pretreated with PEG-AuNPs and TPZs-AuNPs for 24 hours, respectively. When HepG2 cells were irradiated with X-rays of 2.0 Gy, the number of γ-H2AX foci significantly increased to 22.4±5.3. No obvious difference in the number of γ-H2AX foci in the cells with or without AuNPs was observed. The number of γ-H2AX foci slightly increased by 1.7 and 2.4 for cells with PEG-AuNPs and TPZs-AuNPs at 12 hours post-irradiation, respectively, suggesting that DNA damage was mainly exacerbated by X-rays and slightly improved by addition of PEG-AuNPs and TPZs-AuNPs. No obvious difference in DNA DSBs reflected by the γ-H2AX foci was found for HepG2 cells with TPZs-AuNPs or PEG-AuNPs under X-ray irradiation. Probably, the difference in the damages between HepG2 cells with TPZs-AuNPs and PEG-AuNPs induced by X-rays mainly came from the damages in cytoplasm instead of nucleus. This is also coincident with the accumulation of AuNPs in cytoplasm observed in our previous study.[23]

The conjugation of TPZ moiety to PEG-AuNPs surface resulted in changes in particle size and zeta potential, thus possibly influencing the cell uptake. The concentration of AuNPs, especially cell uptake of AuNPs, is the key factor for the enhancement of radiation effect. As shown in Figure 6B, the cell uptake increased with increasing gold concentration in the range of 0–20.0 µg/mL. The cell uptake for TPZs-AuNPs at 10.0 µg/mL concentration was 17.32 fg/cell, while it was only 8.08 fg/cell for PEG-AuNPs. The HepG2 cell uptake of TPZs-AuNPs was approximately tenfold lower than HeLa cell uptake of citrate-capped AuNPs shown in our previous study.[23] The experiments were conducted at the same coculture concentration of AuNPs, but the cell uptake of TPZs-AuNPs was twice more than that of PEG-AuNPs. Whether the radiation enhancement of TPZs-AuNPs is derived from the higher cell uptake of AuNPs is not known.

We then investigated the enhancement in intracellular ROS production for HepG2 cells cocultured with 5.0 µg/mL TPZs-AuNPs and 20.0 µg/mL PEG-AuNPs under X-ray irradiation (Figure S5). The coculture concentrations of these two AuNPs corresponded to 14.75 and 13.43 fg/cell in terms of gold concentration in HepG2 cells, respectively, that is, almost the same gold concentration in cells. The enhancement ratio of ROS production was ~1.24 for TPZs-AuNPs without
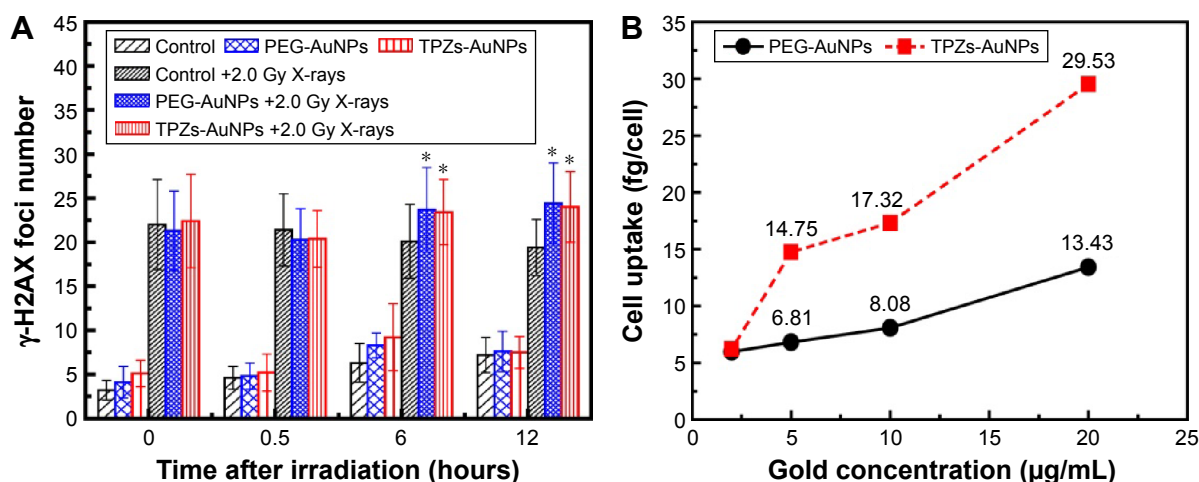
Figure 6 The dependence of γ-H2AX foci and cell uptake of gold nanoparticles.
Notes: (**A**) The numbers of γ-H2AX foci in AuNPs-pretreated HepG2 cells under X-ray irradiation at different times post-irradiation. (**B**) The dependence of uptake of PEG-AuNPs and TPZs-AuNPs by HepG2 cells on coculture concentration. *$P<0.05$ versus control group, determined by the two-tailed Student's *t*-test.
Abbreviations: AuNPs, gold nanoparticles; PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles.

irradiation, which was 24% higher than that in cells without TPZs-AuNPs. This might be due to the endogenous hydroxyl radical production elicited by the accumulated TPZs-AuNPs. When irradiated with X-rays at 2.0 Gy, the radiation enhancement ratios were 1.77, 1.80, and 2.43 for the control, PEG-AuNPs-pretreated cells, and TPZs-AuNPs-pretreated cells, respectively. Therefore, the addition of PEG-AuNPs and TPZ-AuNPs led to 1.7% and 37.3% increments in radiation enhancement ratio, respectively. When the radiation dose increased to 4.0 Gy, TPZ-AuNPs also exhibited 38% increment in radiation enhancement ratio. These results further proved that the conjugation of TPZ moiety on the surface of PEG-AuNPs elicited a greater radiation enhancement than PEG-AuNPs alone under X-ray irradiation.

The radiosensitizing effect of AuNPs has been shown to depend very strongly on the concentration, size, shape, and coating of AuNPs.[40] In our cell radiobiological experiment, 10.0 µg/mL (~50 µM gold concentration) TPZs-AuNPs were used, which was one-tenth of the concentration of PEG-AuNPs used by Liu et al, where 10% radiation enhancement was obtained for 150 kVp X-rays.[36] As far as the gold concentration in a cell is concerned, it was 17.32 fg/cell for the coculture concentration of 10.0 µg/mL, that is, 17.32 ppm (the weight of 10^9 cells is ~1.0 g) in HepG2 cells. As proposed, the accumulated TPZs-AuNPs in endosome/lysosome provided a relatively high concentration of AuNPs and TPZs. On one hand, accumulated AuNPs and TPZs might absorb X-rays to directly produce ROS. On the other hand, plenty of secondary electrons, generated from the irradiated AuNPs, could be captured by TPZ moiety on the AuNPs surface, undergoing a reductive reaction to produce hydroxyl radicals (Figure 7).[27] It could be treated as an intramolecular electron transfer system, which was much more
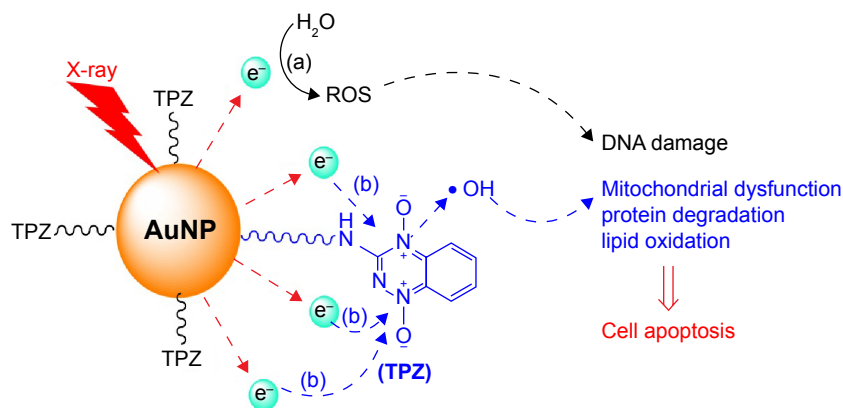

Figure 7 The radiation enhancement mechanism of TPZs-AuNPs proposed in this study.
Abbreviations: TPZ, tirapazamine; AuNP, gold nanoparticle; ROS, reative oxygen species; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles.

efficient than intermolecular electron transfer. Therefore, the synergetic effect between TPZ moiety and AuNPs contributed to the significant radiation enhancement, although a very low loading of TPZs-AuNPs was used. Further study on the protein expression and signal transduction induced by TPZs-AuNPs is needed to disclose the molecular mechanisms underlying the enhanced cell-killing effect. Additionally, because TPZ is a specific drug for hypoxia cells, the radiosentizing effect of TPZs-AuNPs on cells under hypoxia condition as well as in vivo will be investigated in our future study.

Conclusion

We successfully fabricated AuNPs with TPZ moiety by conjugating TPZs on their surface using place-exchange reaction. Our study showed that HepG2 cell lethality induced by X-rays could be enhanced >20% in vitro in the presence of TPZs-AuNPs compared to PEG-AuNPs at almost the same gold concentration. The ROS measurement proved that the reduction of TPZ could be chemically improved by secondary electrons from AuNPs under X-ray irradiation. The synergetic effect between TPZ moiety and AuNPs led to the significant radiation enhancement of TPZs-AuNPs. The improvement which we achieved in this study using drug-modified AuNPs with low gold loading suggests that this methodology is potentially useful for designing more efficient radiation sensitizers for clinical use in the future.

Acknowledgments

Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials
The preparation process of thioctyl tirapazamine
The synthesis of 3-amine-1,2,4-benzotriazole-1-oxide
2-Nitroaniline (4.3 mmol) and cyanamide (22 mmol) were melted together at 100°C, and then cooled to 50°C. Concentrated HCl (5 mL) was added carefully. The mixture was stirred until the exotherm subsided and then stirred at 100°C for 2 hours. Cyanamide (22 mmol) was added, and the mixture was stirred at 100°C for 4 hours. The mixture was cooled to 20°C and made strongly basic with 7.5 M NaOH solution (50 mL), and heated at 100°C for 1 hour. Then, the mixture was cooled to 20°C and diluted with water (100 mL). The precipitate was filtered, washed with water (2×10 mL) and ether (2×10 mL), and dried to obtain the corresponding 3-amine-1,2,4-benzotriazole-1-oxide.

The synthesis of 3-amine-1,2,4-benzotriazole-1,4-dioxide (tirapazamine)
Hydrogen peroxide (30%, 12 mL) was added dropwise to a suspension of 3-amine-1,2,4-benzotriazole-1-oxide (2.60 mmol) in HOAc (25 mL), and the suspension was stirred at 50°C for 4 hours. The solution was diluted with water (100 mL) and carefully neutralized with solid $NaHCO_3$. The solution was extracted with chloroform ($CHCl_3$) (5×50 mL), the combined organic extracts were dried, and the solvent was evaporated. The residue was chromatographed, by eluting with a gradient (0%–5%) of MeOH/$CHCl_3$, to obtain the corresponding 1,2,4-benzotriazin-3-amine-1,4-dioxide (tirapazamine, TPZ) (melting point 228°C–230°C). ^{1}H NMR (400 MHz, CDCl_3) δ 8.57 (dd, J_1=12.8 Hz, J_2=8.6 Hz, 2H), 8.14–8.03 (m, 1H), 7.97 (m, 1H), 4.08 (s, 2H). MS (m/z) 178.05 (M$^+$, 100%).

The synthesis of thioctyl tirapazamine
Thioctic acid (0.412 g, 2 mmol) was dissolved in dichloromethane (20 mL), followed by addition of triethylamine (0.6 mL), and then N-ethyl-N'-(3-dimethylaminopropyl) carbodiimide hydrochloride (0.460 g, 2.4 mmol) and N-hydroxybenzotrizole (0.324 g, 1.2 mmol) were added. The mixture was stirred at room temperature for 1 hour. TPZ (0.356 g, 2 mmol) was added. The resulting mixture was stood overnight. After treatment, the residue was purified by silicon column chromatography to give thioctyl tirapazamine (TPZs; 0.439 g, 60% yield). ^{1}H NMR (400 MHz, CDCl_3) δ 9.08 (s, 1H), 8.57 (d, J=8.4 Hz, 1H), 8.54 (d, J=8.4 Hz, 1H), 8.10 (dd, J_1=8.4 Hz, J_2=7.2 Hz, 1H), 7.97 (dd, J_1=8.4 Hz, J_2=7.2 Hz, 1H), 3.61 (m, 1H), 3.18 (m, 2H), 2.49 (m, 1H), 2.41 (t, J=6.8 Hz, 2H), 1.95 (m, 1H), 1.68–1.85 (m, 4H), 1.658 (m, 2H). HR-MS (M+H)$^+$ m/z: 366.0820, found: 366.0807.

The measurement of TPZs number in TPZs-modified AuNPs using UV-vis spectroscopy
A standard curve of TPZs
TPZs (5.0 mg) was dissolved in methanol (10 mL) as mother solution. Dilution of mother solution with methanol gave various concentrations of TPZs (1×10^{-3}, 5×10^{-4}, 4×10^{-4}, 2×10^{-4}, 1×10^{-4}, 8×10^{-5}, 5×10^{-5}, 4×10^{-5}, 2×10^{-5}, 1×10^{-5} M). The absorption spectra of the resulting TPZs solutions (100 μL) were measured. The dependence of absorbance at 460 nm on TPZs concentration is shown in Figure S1 as a standard curve.

The measurement of TPZs concentration in TPZs-modified AuNPs mother solution
The concentration of TPZs-modified AuNPs (TPZs-AuNPs) stock was measured by ICP-AES (TJA Co., Franklin, MA, USA). A TPZs-AuNPs solution (0.5 mL) at gold concentration of 3,652 mg/mL (18.54 mM) was dissolved in aqua regia (0.5 mL) and diluted with methanol (4.0 mL). The absorbance at 460 nm of the resulting TPZs solution (100 μL) was 0.07958. The concentration of TPZs in the solution under measurement was 1.349×10^{-5} M, calculated using an external standard method. In the stock TPZs-AuNPs solution, the TPZs concentration was 1.349×10^{-4} M.

The calculation of TPZs number in TPZs-AuNPs
According to the reference, the number of gold atoms in a 15 nm gold nanoparticle was ~100,594. In the mother solution, the gold concentration was 18.54 mM, and the concentration of TPZs was 1.349×10^{-4} M. Thus, the calculated number ratio of TPZs to AuNPs was 732:1.
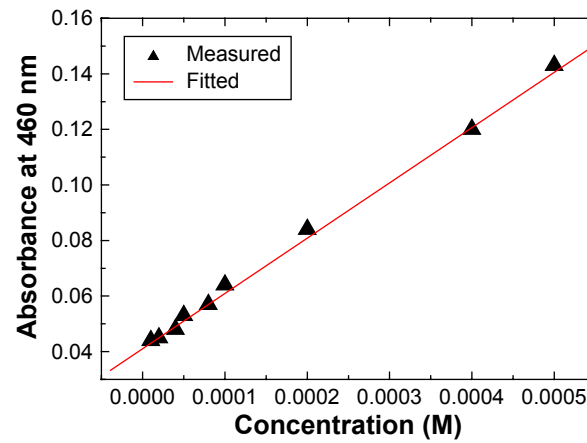
Figure S1 The absorbance at 460 nm of TPZs methanol aqueous solution at various concentrations.
Abbreviation: TPZs, thioctyl tirapazamine.
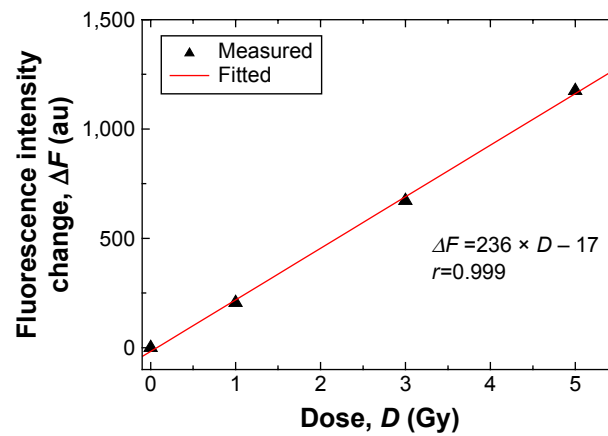

$$\Delta F = 236 \times D - 17$$
$$r = 0.999$$

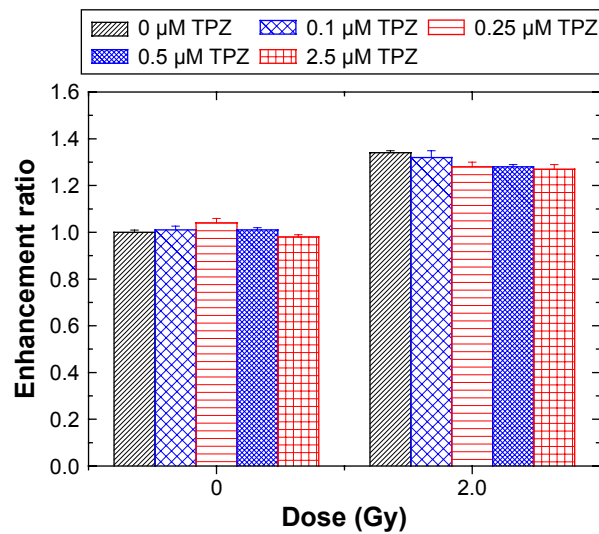**Figure S2** The dependence of fluorescence intensity change on the dose of X-rays (50 kVp).

Figure S3 The enhancement ratio of hydroxyl radical production induced by tirapazamine under X-ray irradiation.
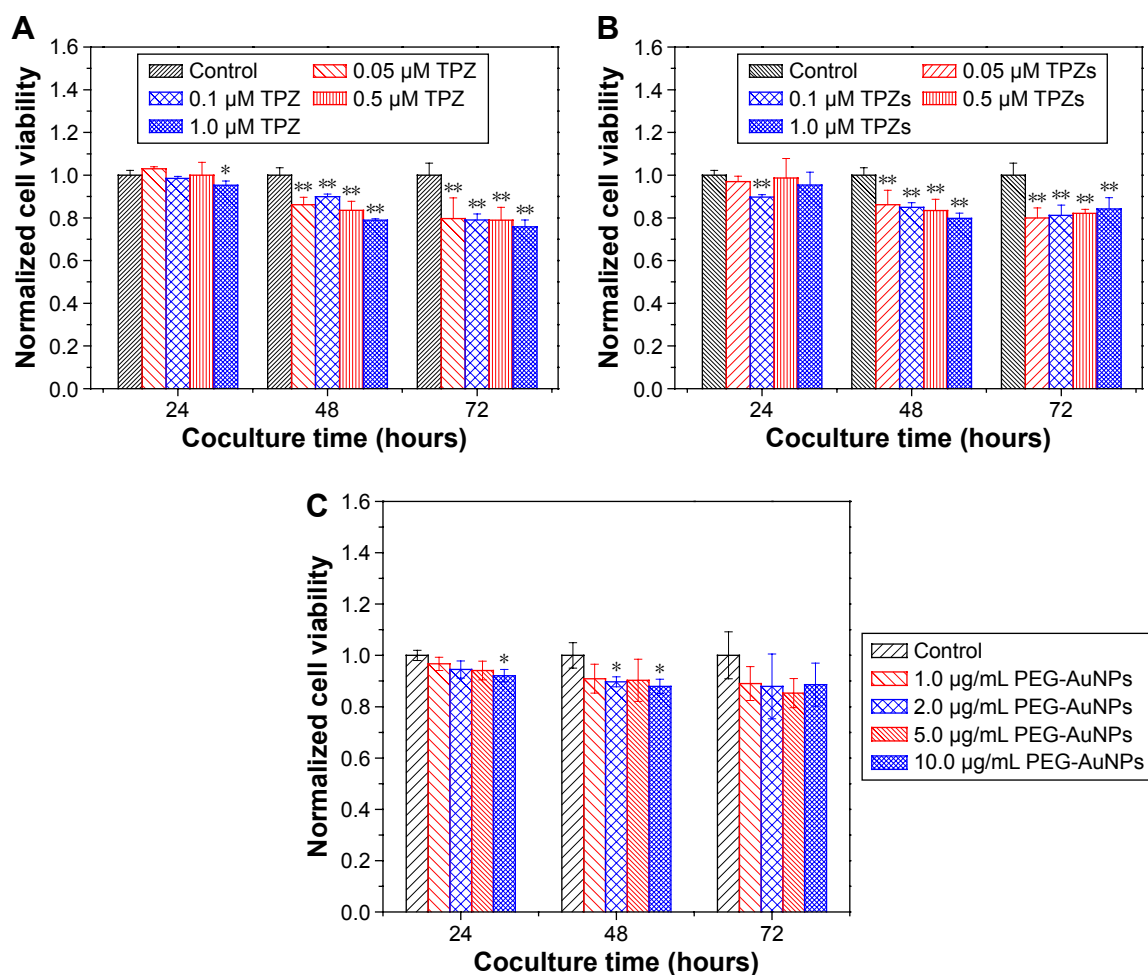**Abbreviation:** TPZ, tirapazamine.

Figure S4 The Cytotoxicity of TPZ, TPZs and PEG-AuNPs.

Notes: The normalized cell viability of HepG2 cells cocultured with (**A**) TPZ, (**B**) TPZs, and (**C**) PEG-AuNPs at various concentrations relative to that of untreated HepG2 cells at 24, 48, and 72 hours. *$P<0.05$ and **$P<0.01$, determined by the two-tailed Student's t-test.

Abbreviations: TPZ, tirapazamine; TPZs, thioctyl tirapazamine; PEG-AuNPs, polyethylene glycol-capped gold nanoparticles.
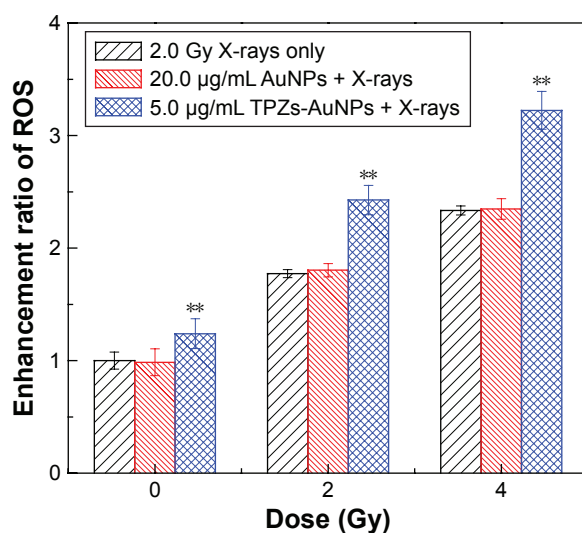
Figure S5 The enhancement ratio of ROS in HepG2 cells after X-ray irradiation in the presence of PEG-AuNPs (20.0 μg/mL) and TPZs-AuNPs (5.0 μg/mL).

Note: **$P<0.01$ versus control group, determined by the two-tailed Student's t-test.

Abbreviations: ROS, reactive oxygen species; PEG-AuNPs, polyethylene glycol-capped gold nanoparticles; TPZs-AuNPs, thioctyl tirapazamine-modified gold nanoparticles; AuNPs, gold nanoparticles.