Nasopharyngeal Carcinoma Column

Regulation of hypoxia-induced mRNA expressions of HIF-1α and osteopontin and in vitro radiosensitization by tirapazamine in human nasopharyngeal carcinoma HNE-1 and CNE-1 cells

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[Abstract] Background and Objective: Combined hypoxic cytotoxic drugs and chemoradiotherapy is an important mean of oncotherapy, and Tirapazamine (TPZ) is one of the most remarkable drugs. It has been shown that TPZ has a synergistic effect with radiotherapy on tumor cells, but whether TPZ would down-regulate the expression of the hypoxia-induced genes has not been reported. This study was to investigate the hypoxia-induced mRNA expressions of hypoxia inducible factor-1α (HIF-1α) and osteopontin (OPN) in human nasopharyngeal carcinoma HNE-1 and CNE-1 cells and the radiosensitization of TPZ, a hypoxia-specific drug, on HNE-1 and CNE-1 cells in vitro. Methods: The IC₅₀ values of TPZ for HNE-1 and CNE-1 cells were measured using MTT assay, and the mRNA expressions of HIF-1α and OPN in HNE-1 and CNE-1 cells was determined using RT-PCR under aerobic and hypoxic conditions, respectively. The survival rates of HNE-1 and CNE-1 cells treated with or without TPZ at IC₅₀ in the presence or absence of oxygen for 6 h were determined using colony formation assay following exposure to 1–6 Gy of ⁶⁰Co radiation. The dose-survival curves were plotted and the values of D₀, Dα and SER were calculated as a single-hit multitarget model. Results: The IC₅₀ values of TPZ were 34.81 μmol/L and 35.02 μmol/L in HNE-1 and CNE-1 cells under aerobic condition, and 30.20 μmol/L and 28.48 μmol/L under hypoxic condition, respectively. The expressions of HIF-1α and OPN mRNA were reduced by TPZ in HNE-1 cells, but not in CNE-1 cells under hypoxic condition. For the HNE-1 cells, the respective values of D₀ and Dα were 0.89 Gy and 0.28 Gy following normoxic Irradiation versus 1.47 Gy and 0.44 Gy following hypoxic irradiation. For the CNE-1 cells, the respective values of D₀ and Dα were 0.72 Gy and 0.68 Gy following normoxic irradiation versus 0.95 Gy and 0.56 Gy following hypoxic irradiation. The values of D₀ and Dα for HNE-1 and CNE-1 cells treated with TPZ under hypoxic condition following Irradiation were 0.66 Gy, 0.21 Gy and 0.85 Gy, 0.79 Gy, respectively. Conclusion: TPZ can down-regulate hypoxia-induced expression of HIF-1α and OPN mRNA of HNE-1 cells and radiosensitize the HNE-1 cells but not CNE-1 cells, and act as a hypoxia modifier.

Key words: Tirapazamine, nasopharyngeal carcinoma, irradiation, HIF-1α, osteopontin

Tumor hypoxia is one of the main reasons for radiotherapy and chemotherapy failure in solid tumors (chemotherapy and radiotherapy resistance). The hypoxic cells account for 10%–50% in solid tumors and their tolerance to radiation and chemotherapy is 2.5–3 times stronger than that of aerobic cells, which becomes one of the important factors making cancer difficult to cure, and easy to recur and metastasize. Therefore, the toxic drugs to hypoxic cells in combination with chemoradiotherapy is an important regimen for cancer treatment. Tirapazamine (TPZ) is one of the most remarkable toxic drugs to hypoxic cells. TPZ enters into the cell, forms a high activity of free radicals through the role of intracellular reductase, and reduces the intracellular oxygen ion into oxygen. Under hypoxic condition, the highly active TPZ free radical can obtain a hydrogen atom from the DNA through the mediation of topoisomerase II, cause single/double-strand breaks and chromosome damage, thus resulting in cell death. Studies have shown that TPZ combined with radiotherapy has synergistic effects on tumor cells, but whether it can down-regulate hypoxia-induced genes under hypoxic condition has not been reported. In this experiment, HNE-1 and CNE-1 human nasopharyngeal carcinoma cells were used as an in vitro model to explore the expression differences of hypoxia inducible factor-1α (HIF-1α) and OPN in these two cell lines under normoxic and hypoxic conditions and
the impact of TPZ on the expression, and to detect the radiation sensitizing effects of TPZ on the two nasopharyngeal carcinoma cell lines under hypoxic condition by colony-forming method so as to provide a new sensitizing means for clinical radiotherapy of nasopharyngeal carcinoma.

Materials and methods

Drugs and reagents

TPZ was purchased from the U.S. Biotechnology Co., Ltd.; methilthiazolyl tetrazolium (MTT) and dimethyl sulfoxide (DMSO) from Sigma; RPMI-1640 medium from Gibco; calf serum from China Lanzhou Ming-Hai Bio-Engineering Company; Trizol kit from Shanghai Bio-Engineering Co., Ltd.; and RT-PCR kit from Chengdu Boracker Biotechnology Co., Ltd.

Cells and culture conditions

Human nasopharyngeal carcinoma cell lines HNE-1 (EBV-positive) and CNE-1 (EBV-negative) were obtained from the Sichuan Provincial Tumor Institute and were cultured in RPMI-1640 culture medium containing fetal calf serum (10%), penicillin (100 u/mL) and gentamicin (40 u/mL) under conventional conditions (37°C, 5% CO₂); for hypoxic culture, conventionally cultured cells were placed in a hypoxic culture device containing 5% CO₂ and 0.01% O₂. All experiments were carried out in the cell exponential growth phase.

Detection of cell growth inhibition by MTT

The HNE-1 and CNE-1 cells in exponential growth phase were collected, adjusted to a cell concentration of 5 × 10⁵/mL with RPMI-1640 medium containing 10% fetal calf serum, and were inoculated onto a 96-well plate, 0.1 mL/well. After cell adherence, different concentrations of TPZ (a final concentration of 5–50 μmol/L) were added, the drug of each concentration was inoculated into four holes. After 72 h under normoxic and hypoxic conditions, 20 μL MTT (5 mg/mL) was added, supernatant was discarded after 4h incubation at 37°C, and added with 150 μL dimethyl sulfoxide (DMSO). After full blending, DF-M3000 enzyme-labeled instrument was used to measure the absorbance (A value) at 570 nm wavelength and the cell survival rate was calculated by the following formula, as well as the concentration-survival regression equation and half-effective inhibitory concentration IC₅₀ and IC₅₀. Each experiment was repeated 3 times.

Cell survival rate = \((A_{test \ group} - A_{blank \ group}) / (A_{control \ group} - A_{blank \ group})\)×100%

Detection of HIF-1α and OPN expression by RT-PCR

The HNE-1 and CNE-1 cells treated or untreated by TPZ and cultured under hypoxic or normoxic conditions were collected, the medium was discarded, washed by PBS for twice and added with 1 mL Trizol reagent. The total RNA was extracted according to the instructions from Trizol kit, and quantitatively examined by 1.5% agarose gel electrophoresis, which showed two brightness diminishing clear bands of 28S and 18S without obvious sign of dispersion, and the brightness ratio of 28S and 18S band was about 2:1. The total RNA (2 μg) with a final reaction volume of 20 μL was reversibly transcribed into cDNA. Two μL cDNA was used for PCR with β-actin as internal reference, and the PCR primers were designed using Primer premier 4.0 software. β-actin primers: 5’-AACAGATGGCCACGGCTGCT-3’ for upstream primer, and 5’-GACTCGTCATACTCCTGCTTGG-3’ for downstream primer (421 bp); HIF-1α primers: 5’-TTGATTTGACATCCATCTCTC-3’ for upstream primer, and 5’-TCCGTTTTCTCTGAGCATTTC-3’ for downstream primer (249 bp); OPN primers: 5’-GGAGGAG GCAGAGCACAG-3’ for upstream primer, 5’-CGTTGGACT TACTGGAAG-3’ for downstream primer (331 bp). HIF-1α reaction conditions: 94°C for 2 min, followed by 35 cycles of 94°C for 0.5 min, 56°C for 0.5 min and 72°C for 1 min, and then thermal insulation at 22°C. OPN reaction conditions: 94°C for 2 min, followed by 35 cycles of 94°C for 0.5 min, 55°C for 0.5 min and 72°C for 1 min, and then thermal insulation at 22°C. β-actin reaction conditions were the same with HIF-1α and OPN. Six μL PCR product was used for 1.5% agarose gel electrophoresis, the gray stripe was measured quantitatively by ultraviolet imaging system, and the ratio of HIF-1α and OPN to β-actin was analyzed respectively. Each experiment was repeated four times.

Irradiation method and dose-survival curve

¹³C̃O γ-ray was used with an open-field size of 5 cm × 5 cm, a depth of 80 cm (SSD) and a liquid depth of 0.5cm. Radiation doses were 0, 0.2, 0.4, 1, 2, 3, 4, 5 and 6 Gy. The HNE-1 and CNE-1 cells in exponential growth phase were cultured and inoculated into 12-well plates, 2.5 × 10⁴ cells/well, added with TPZ (IC₅₀), and irradiated after 6 h under normoxic and hypoxic conditions. After irradiation, the cells were collected immediately with 0.25% trypsin-EDTA, counted and inoculated at a 60 mm dish in diameter by adding 4 mL RPMI-1640 medium containing 10% fetal calf serum, cultured for 14 days at 37°C, 5% CO₂, the medium was discarded and stained by solution with crystal violet (0.5%) and methanol (30%) for 1 min, and then the clone numbers in each dose group were counted (for colonies with more than 50 cells) and the cell survival fraction (SF) was calculated. The experiment was repeated three times. The single-hit multi-target mathematical model was used to draw a dose-survival curve, D₀, D₅₀, SF₂ and radiosensitization ratios (SER) were obtained, and the radiosensitivity of each group was analyzed and compared.

Statistical analysis

The experimental data were expressed as mean ± SD and tested by t test. Analysis of variance was used to compare two groups. SPSS13.0 statistical software was used and the test level was 0.05.

Results

Growth inhibition of nasopharyngeal carcinoma cell lines by TPZ

The IC₅₀ of TPZ in HNE-1 and CNE-1 cells under normoxic condition was 34.81 ± 4.18 μmol/L and 35.02 ± 4.20 μmol/L; the IC₅₀ in HNE-1 and CNE-1 cells under hypoxia condition was 30.20 ± 3.32 μmol/L and 28.48 ± 3.13 μmol/L, respectively. Compared with normoxic group serving as control, the sensitivity to TPZ was increased in hypoxic groups and the IC₅₀ ratio was
Expressions of HIF-1α and OPN in HNE-1 and CNE-1 nasopharyngeal carcinoma cell lines

Semi-quantitative RT-PCR test showed that under normoxic condition, HIF-1α mRNA was expressed in two kinds of nasopharyngeal carcinoma cells, but at low level and the expression was cell-specific, which was higher in CNE-1 cells. After culture under hypoxic condition for 24 h, the expression of HIF-1α mRNA significantly increased in HNE-1 cells, but slightly decreased in CNE-1 cells. After treatment by different concentrations of TPZ, the expression level in HNE-1 cells became lower than in the hypoxic group, but without significant dependency on concentration, while that in CNE-1 cells increased, as shown in Figures 1 and 2.

Under normoxic condition, OPN mRNA was expressed in HNE-1 cells at a very low level and was not expressed in CNE-1 cells. Under hypoxic condition, it was expressed in two kinds of nasopharyngeal carcinoma cells at a significantly increased level, especially in HNE-1 cells. After treatment with different concentrations of TPZ, the expression levels in the two kinds of cells gradually decreased with increased concentration as compared with those in hypoxia group (Figures 3 and 4).

**TPZ radiosensitization**

The HNE-1 cell line had a D0 value of 0.89 Gy and a Dq value of 0.28 Gy in normoxic irradiation group, a D0 value of 1.47 Gy and a Dq value of 0.44 Gy in hypoxic irradiation group, and a D0 value of 0.66 Gy and a Dq value of 0.21 Gy in hypoxic drug-added irradiation group. The CNE-1 cell line had a D0 value of 0.72 Gy and a Dq value of 0.68 Gy in normoxic irradiation group, a D0 value of 0.95 Gy and a Dq value of 0.56 Gy in hypoxic irradiation group, and a D0 value of 0.85 Gy and a Dq value of 0.79 Gy in hypoxic drug-added irradiation group. Figures 5 and 6 showed the radiation survival curves of HNE-1 and CNE-1 in normoxic group, hypoxic group and hypoxia drug-added group, respectively. It could be seen that HNE-1 cells had a significantly lower radiosensitivity after treatment by hypoxia (P < 0.05), while the radiosensitivity significantly increased after TPZ treatment (P < 0.05); CNE-1 cells had a slightly lower radiosensitivity after hypoxic treatment (P > 0.05), while the radiosensitivity slightly increased after TPZ administration (P > 0.05).

**Discussion**

In this study, the results showed that under hypoxic condition, TPZ could enhance the cytotoxicity in HNE-1 and CNE-1 cells, especially in HNE-1 cells. Studies have shown that TPZ enters into the cell, forms a highly active free radical through the action of intracellular reductase, and the intermediate of free radical is rapidly oxidized into the parent molecule under aerobic condition, while producing a super-oxide free base, leading to occurrence of side effects such as muscle spasm when patients receive TPZ treatment. Under hypoxic condition, the highly active TPZ free radical can obtain a hydrogen atom from adjacent large molecule, resulting in DNA single/double-strand breaks and chromosome damage mediated by topoisomerase II, and causing cell death. In the hypoxic microenvironment in solid tumors, tumor cells maintain their metabolic stability and promote their growth and
Increasing the oxygen concentration of cell microenvironment. Producing oxygen, and down-regulating the HIF-1 because under hypoxic condition, through the role of intracellular (LMP1) promote HIF-1 cell lines significantly increased because the EB virus could also specifically over-expressed, increases the expression of vascular endothelial growth factor (VEGF), and promotes tumor proliferation, resulting in decreased radiosensitivity. Many previous studies using TPZ for hypoxic cells have shown that TPZ can enhance the radiosensitivity of hypoxic cells because under normoxic and hypoxic conditions confirmed that, under hypoxic condition, the radiosensitivity of both nasopharyngeal carcinoma cells decreased, especially the HNE-1 cells, which was similar to the results from most of the other clinical reports. The increased radiosensitivity after the use of TPZ under hypoxic condition may be due to the increased oxygen concentration of cell micro-environment and the inhibition of HIF-1α expression.

Tumor hypoxia has been one of the main factors compromising the efficacy of radiotherapy and chemotherapy. Under hypoxic condition, on one hand, the radiation-generated oxygen free radicals in tumor cells are reduced, lessening the radiotherapy-induced DNA breakage; on the other hand, HIF-1 is specifically over-expressed, increases the expression of vascular endothelial growth factor (VEGF), and promotes tumor proliferation, resulting in decreased radiosensitivity. Many previous studies using TPZ for hypoxic cells have shown that TPZ can enhance the radiosensitivity of hypoxic cells. Our study on radiosensitivity of HNE-1 and CNE-1 cells under normoxic and hypoxic conditions confirmed that, under hypoxic condition, the radiosensitivity of both nasopharyngeal carcinoma cells decreased, especially the HNE-1 cells, which was similar to the results from most of the other clinical reports. The increased radiosensitivity after the use of TPZ under hypoxic condition may be due to the increased oxygen concentration of cell micro-environment and the inhibition of HIF-1α expression.

In this study, for the first time, TPZ is found to up-regulate the mRNA expressions of hypoxia-related gene HIF-1α and OPN, while enhancing the radiosensitivity through increasing the oxygen concentration in cell micro-environment, so it can be used as a kind of oxygen additive to increase the radiosensitivity of human nasopharyngeal carcinoma hypoxic cell lines. However, the role of TPZ in the protein expression of hypoxia-related gene HIF-1α and OPN, as well as downstream proteins needs to be further studied. At present, this drug is in phase III clinical trials abroad, but part of the clinical trials show that TPZ has obvious side effects such as muscle spasm, therefore, how to further overcome its toxicity and target hypoxic cells is still a focus of therapy targeting tumor hypoxia.

References


