Antitumor activity of dichloroacetate on C6 glioma cell: in vitro and in vivo evaluation

Yu Duan¹
Xin Zhao¹
Wei Ren¹
Xin Wang¹
Ke-Fu Yu¹
Dan Li¹
Xuan Zhang¹
Qiang Zhang¹,²

¹Department of Pharmaceutics, School of Pharmaceutical Sciences, Peking University, Beijing, People’s Republic of China; ²State Key Laboratory of Natural and Biomimetic Drugs, School of Pharmaceutical Sciences, Peking University, Beijing, People’s Republic of China

Abstract: Dichloroacetate (DCA), a small molecule mitochondria-targeting agent, can penetrate the blood–brain barrier, showing potential therapeutic effects on brain tumors. Considering the effects of DCA on tumor cellular metabolism, penetrating across the blood–brain barrier, as well as having potential antitumor activity on brain tumors, the purpose of this study is to investigate the antitumor activity of DCA on C6 glioma cells in vitro and in vivo. DCA inhibited C6 glioma cell proliferation, induced C6 cell apoptosis, and arrested C6 cells in S phase. DCA can inhibit the expression of heat shock proteins 70 (Hsp70) in a dose-dependent and time-dependent manner (P ≤ 0.01). Our in vivo antitumor effect results indicated that DCA markedly inhibited the growth of C6 glioma tumors in both C6 brain tumor-bearing rats and C6 tumor-bearing nude mice (P < 0.01). DCA significantly induced the ROS production and decreased the mitochondrial membrane potential in tumor tissues. Our in vivo antitumor effect results also indicated that DCA has potential antiangiogenic effects. In conclusion, DCA may be a viable therapeutic agent in the treatment of gliomas.

Keywords: dichloroacetate, DCA, C6 glioma, antitumor efficacy, in vitro, in vivo

Introduction
Malignant gliomas are the most common and deadly brain tumors.¹ Approximately 20,000 patients are diagnosed with gliomas each year in the United States.² It was reported that the median survival time of patients with malignant gliomas ranges from 14 to 40–50 weeks despite aggressive multimodality management with surgery, radiation, and chemotherapy.³ Radiotherapy has been of key importance to the treatment of these tumors for decades.⁴ Newer surgical techniques have become important in the management of malignant gliomas.⁵,⁶ Because of the angiogenesis in malignant gliomas,⁷ the antivascular endothelial growth factor therapy has had significant efficacy in gliomas.⁸ In addition, it was reported that heat shock protein 70 (Hsp70) is overexpressed in glioma cells.⁹,¹⁰ Overexpressed Hsp70 could decrease the release of cytochrome c and apoptosis inducing factor (AIF), which can then serve an antiapoptotic function.¹¹ Therefore, down-regulating Hsp70 expression would enhance the release of cytochrome c and AIF, leading to tumor cell apoptosis. It is well known that many chemotherapeutic agents have a low therapeutic index in brain tumors.¹² The failure of chemotherapy is due to the inability of intravenously administered anticancer agents to reach the brain tissue. The blood–brain barrier (BBB) is one of the most important obstacles for preventing the penetration of drugs into the central nervous system.¹³ Therefore, great efforts have been made to develop various strategies for improving the penetration of drugs across the BBB, as well as for improving the targeting effect...
to the brain tumors, including targeting drug delivery systems or conjugates.\textsuperscript{14-16} Compared to the therapy of using chemotherapeutic agents or molecular-targeting therapeutic agents, an alternative antitumor approach may be to consider tumor metabolism.\textsuperscript{17} Now, mitochondria are increasingly recognized as important targets in tumors because of their central roles in apoptotic pathways and in cellular metabolism.\textsuperscript{18} The most common metabolic hallmark of cancer cells is their propensity to metabolize glucose to lactic acid at a high rate even in the presence of oxygen.\textsuperscript{19} Tumor cells produce a clearly increased amount of their energy through glycolysis under aerobic conditions.\textsuperscript{20} This metabolic switch away from mitochondrial respiration towards aerobic glycolysis in tumors could cause tumor cell transformation and suppress apoptosis.\textsuperscript{21} Progress in molecular biology has revealed that energy metabolism is strictly regulated by several protein kinases. Pyruvate dehydrogenase kinase (PDK) is a gatekeeping enzyme that regulates the flux of carbohydrates (pyruvate) into the mitochondria. In the presence of activated PDK, pyruvate dehydrogenase (PDH), a critical enzyme that converts pyruvate to acetyl-CoA instead of lactate in glycolysis, is inhibited, limiting the entry of pyruvate into the mitochondria. Notably, a recent report showed that the entry of pyruvate into the Krebs cycle is inhibited in cancer cells, and aerobic glycolysis is highly increased in cancer cells.\textsuperscript{22} This bioenergetic difference between cancer and normal cells would offer a very selective therapeutic target.\textsuperscript{21}

As many molecules with mitochondrial activity were discovered more than a decade ago, newer targeted screening strategies may uncover many additional compounds with desirable anticancer activities.\textsuperscript{23} Dichloroacetate (DCA) is considered as a small molecule mitochondria-targeting agent.\textsuperscript{24} Clinical studies of DCA have shown reduced lactate levels in patients with congenital lactic acidosis and sepsis.\textsuperscript{24,25} It has been reported that DCA activates the PDH by inhibition of PDK in a dose-dependent manner, and results in increased delivery of pyruvate into the mitochondria.\textsuperscript{26} The antitumor activity of DCA on nonsmall cell lung cancer, breast cancer, glioblastomas, and endometrial and prostate cancer cells has been demonstrated.\textsuperscript{26-28} In addition, the oral bioavailability of DCA is almost 100\%.\textsuperscript{21} DCA can penetrate into the traditional chemotherapy sanctuary sites. Interestingly, it was reported that DCA could penetrate across the BBB,\textsuperscript{20} exhibiting the potential activity for brain therapy.

Considering the effects of DCA on tumor cell metabolism, penetrating across the BBB, as well as potential antitumor activity on brain tumors, in this study, we investigated the antitumor activity of DCA on C6 gliomas in vitro and in vivo. The murine C6 glioma cells were selected as the cell model. The cell proliferation, cell cycle, apoptosis, and in vitro cytotoxicity of the DCA were also investigated. The effect of DCA on Hsp70 expression was analyzed in C6 cells in vitro. The antitumor activity of DCA was evaluated in C6 brain tumor-bearing rats and C6 tumor-bearing nude mice in vivo. Furthermore, the reactive oxygen species (ROS) production and mitochondrial membrane potential in tumor tissues were also investigated. In addition, the antiangiogenic activity of DCA was evaluated in C6 brain tumor-bearing rats in vivo.

**Materials and methods**

**Materials**

The DCA was obtained from Sinopharm Chemical Reagent Beijing Co, Ltd (Beijing, People’s Republic of China) and the solution was adjusted to a pH of 6.8–7.2 by 1 M NaOH solution. Sulforhodamine B (SRB) and Tris-base were purchased from Sigma-Aldrich (St Louis, MO, USA). Cell culture medium, penicillin–streptomycin, fetal bovine serum, horse serum, and L-glutamine were from Gibco\textsuperscript{®}, Life Technologies (Carlsbad, CA, USA). All other chemicals were of analytical grade.

**Cell culture**

Murine C6 glioma cells were purchased from the Institute of Material Medical, Chinese Academy of Medical Sciences and Peking Union Medical College (Beijing, People’s Republic of China). The C6 cells were routinely grown in Ham’s F-10 medium supplemented by 2.5% heat-inactivated fetal bovine serum, 15% horse serum, penicillin (100 U/mL), and streptomycin (100 µg/mL). All cells were maintained at 37°C in humidified environment with 5% CO\textsubscript{2}.

**Animals**

Male Sprague Dawley rats (200 ± 20 g) and male BALB/c nude mice (18–20 g) were obtained from the Experimental Animal Center of Peking University (Beijing, People’s Republic of China), and maintained under natural light/dark conditions. Animals were acclimatized for 7 days prior to the experiment and were allowed free access to a standard diet and water. The temperature and relative humidity were maintained at 25°C and 50\%, respectively. All care and handling of animals were performed with the approval of the Institutional Authority for Laboratory Animal Care of Peking University.
Cell proliferation assay
The effect of DCA on C6 cell proliferation was analyzed with the SRB assay. Briefly, C6 cells were plated in 96-well culture plates (1 x 10^4 cells/well). After 24 hours of incubation, the cells were treated with DCA (0 mM, 5 mM, 10 mM, 20 mM, 40 mM, 80 mM, 160 mM, 320 mM, 640 mM, and 1280 mM) were used. At 5 hours, 12 hours, or 24 hours, the medium was removed, and then cells were collected. Hsp70 were quantified using a commercial Hsp70 enzyme-linked immunosorbent assay (ELISA) kit (Hyperheal Biotech Co, Ltd, Shanghai, People’s Republic of China) as described by the manufacturer’s instruction. The concentration of the DCA (0 mM, 0.05 mM, 0.5 mM, and 5 mM) were used. At 24 hours, the medium was removed, and then cells were fixed with trichloroacetic acid, and washed and stained with SRB. The absorbance was measured at 540 nm using a 96-well plate reader (680; Bio-Rad Laboratories, Hercules, CA, USA). The percentage of surviving cells was calculated using the following formula:

\[ \text{Survival} \% = \frac{(A_{540nm} \text{ for the control cells})}{(A_{540nm} \text{ for the treated cells})} \times 100 \]  

where \( A_{540nm} \) is the absorbance value. Each assay was repeated a minimum of three times, with quadruplicate determinations for each dose level.

In vitro cytotoxicity
In vitro cytotoxicity was measured through the SRB method. Briefly, C6 glioma cells were seeded at a density of 5 x 10^4 cells/well in 96-well transparent plates and incubated for 24 hours. The medium was then changed, and various concentrations of the DCA (0 mM, 1 mM, 2 mM, 4 mM, 8 mM, 16 mM, 32 mM, 64 mM, and 128 mM) were used. At 24 hours, the medium was removed, and then cells were fixed with trichloroacetic acid, and washed and stained with SRB. The absorbance was measured at 540 nm using a 96-well plate reader (680; Bio-Rad Laboratories). The survival percentages were calculated using the following formula:

\[ \text{Survival} \% = \frac{(A_{540nm} \text{ for the treated cells})}{(A_{540nm} \text{ for the control cells})} \times 100 \]  

where \( A_{540nm} \) is the absorbance value. Each assay was carried out in triplicate. Finally, the drug concentration which inhibited the cell growth by 50% (IC_{50}) was determined from semilogarithmic dose–response plots.

Cell cycle analysis
Cell cycle perturbations were analyzed by propidium iodide (PI) DNA staining. Briefly, exponentially growing C6 cells (5 x 10^5) were treated with DCA (0 mM, 5 mM, 20 mM, 40 mM) for 24 hours. At the end of each treatment, cells were collected after a gentle centrifugation at 1000 rpm for 5 minutes and then fixed in 70% ethanol for at least 12 hours at 4°C. Ethanol-suspended cells were diluted with phosphate buffered saline (PBS) and then centrifuged at 1000 rpm for 5 minutes to remove residual ethanol. For cell cycle analysis, the pellets were suspended in 0.4 mL of PBS containing 0.02 mg/mL of PI, 1 mg/mL of DNase-free RNase A and 0.1% of Triton X-100 and incubated at 37°C for 30 minutes. The number of cells used for cell cycle flow cytometry was 10,000. Cell cycle profiles were studied using a flow cytometer (FACSCalibur; Becton Dickinson, Mountain View, CA, USA). The data were analyzed through FCS Express V3 software (De Novo Software, Los Angeles, CA, USA).

Apoptosis assay
The apoptosis was measured through the Annexin V-fluorescein isothiocyanate (FITC) Apoptosis Detection Kit (Biosea Biotechnology Co, Ltd, Beijing, People’s Republic of China) as described by the manufacturer’s instruction. After exposure to DCA (0 mM, 5 mM, 20 mM, 40 mM) for 24 hours, C6 cells were collected, washed twice with PBS, gently resuspended in Annexin V binding buffer, and incubated with Annexin V-FITC/PI in the dark for 30 minutes and analyzed by flow cytometry. In each sample, 10,000 cells were used for analysis.

Hsp70 measurement
Hsp70 were quantified using a commercial Hsp70 enzyme-linked immunosorbent assay (ELISA) kit (Hyperheal Biotech Co, Ltd, Shanghai, People’s Republic of China). Briefly, C6 glioma cells were seeded at a density of 3 x 10^4 cells/well in 175 cm^2 flat-bottom static culture flasks and incubated for 24 hours. The medium was then changed and various concentrations of the DCA (0 mM, 0.05 mM, 0.5 mM, and 5 mM) were used. At 5 hours, 12 hours, or 24 hours, the medium was removed, and then cells were collected. Hsp70 were quantified using commercial Hsp70 ELISA kit as described by the manufacturer’s instruction. The detection limit of these kits for Hsp70 was 0.05 ng/mL. The control group was performed on C6 cells with absent DCA (0 mM DCA). Each sample was run in duplicate and compared with a standard curve. Each assay was carried out in triplicate.

In vivo anti-tumor effect
C6 brain tumor-bearing rats, established according to our previous research, were randomly divided into four groups (each group contained ten animals, n = 10). Animals in group 1 were used as a control group and given distilled water. Animals in groups 2–4 were treated with DCA at the dose of 25 mg/kg, 75 mg/kg, or 125 mg/kg, respectively.
DCA was administrated by oral gavage every day from day 7 after tumor inoculation, and this lasted for 17 consecutive days. On day 23 after tumor inoculation, all rats for each group were sacrificed. The tumors were harvested from the normal brain tissue and weighed. Tumor weight inhibition (TWI) was calculated by the formula:

\[ TWI = \frac{WC - WD}{WC}, \]  

(3)

where WC and WD stand for the tumor weight of animals from the control and treatment groups, respectively.\(^3^3\)

In another separate study, the male BALB/c nude mice were inoculated subcutaneously in the right flank with 0.1 mL of serum-free Ham's F-10 medium C6 cell suspension (4 × 10\(^6\)). After inoculating the C6 cells into the animals, the glioma was developed. Administration began at day 6 after inoculation, when the tumor size reached approximately 100–150 mm\(^3\). On day 6 after tumor inoculation, mice were randomly divided into a control group and three DCA treatment groups. Each group was treated with sterilized water or DCA (25 kg/mg, 75 mg/kg, or 125 mg/kg for 16 consecutive days from day 6 to day 21 after inoculation), respectively. DCA was given by oral gavage. Each group consisted of seven tumor-bearing mice. Throughout the study, mice were weighed and tumors were measured with calipers every two days. Tumor volumes were calculated from the formula:

Tumor volume (mm\(^3\)) = (major axis) \times (minor axis) \times 0.5. \hspace{1cm} (4)

Animals were sacrificed on day 21 after tumor inoculation. The tumors were harvested and weighed. TWI was calculated using the formula:

\[ TWI = \frac{WC - WD}{WC}, \]  

(5)

where WC and WD represent the tumor weight of animals from control and treatment groups, respectively.

**ROS in tumor tissues**

The ROS production in tumor tissues was assayed using the biopsy ROS kit (Genmed Scientifics, Inc, Wilmington, DE, USA). The tumor tissues of nude mice obtained from in vivo antitumor activity experiments were immediately incubated with JC-1 at 37°C for 20 minutes. Then, these tissues were homogenized with a glass homogenizer, and the homogenate was centrifuged at 1500 g for 10 minutes to attain the supernatant. The fluorescence was monitored by a fluorospectrophotometer. The excitation wavelength was set at 490 nm, while the emission wavelength was set at 590 nm and 530 nm for red and green fluorescence, respectively. The MMP was represented by the ratio of red and green fluorescence intensity.

**MMP in tumor tissues**

The determination of mitochondrial membrane potential (MMP) in tumor tissues was assayed using a biopsy MMP kit (Genmed Scientifics, Inc). The tumor tissue slices of nude mice obtained from in vivo antitumor activity experiments were immediately incubated with JC-1 at 37°C for 20 minutes. Then, these tissues were homogenized with a glass homogenizer, and the homogenate was centrifuged at 1500 g for 10 minutes to attain the supernatant. The fluorescence was monitored by a fluorospectrophotometer. The excitation wavelength was set at 490 nm, while the emission wavelength was set at 590 nm and 530 nm for red and green fluorescence, respectively. The MMP was represented by the ratio of red and green fluorescence intensity.

**Statistical analysis**

Data are presented as the mean ± standard deviation. One-way analysis of variance was used to determine
significance among groups, after which post-hoc tests with the Bonferroni correction were used for comparisons between individual groups. Statistical significance was established at $P < 0.05$.

**Results**

**DCA induces cell cycle arrest and apoptosis in C6 glioma cell lines**

The inhibitory effect of DCA on C6 cell growth was determined by SRB assay. As shown in Figure 1, DCA inhibited cell growth in the C6 cell line in a concentration-dependent manner. The value of the antiproliferative ratio (100% – variability) at 10 mM and 25 mM was 80.6% and 50.5%, respectively. According to the results of in vitro cytotoxicity, the calculated IC$_{50}$ values of DCA for C6 cells were 27.0 ± 3.0 mM.

The C6 cells were incubated with DCA for 24 hours. The cells were fixed, stained with PI, and then analyzed by flow cytometry. Figure 2 showed the effect of DCA on the C6 cell cycle progression. It can be seen that treatment of C6 cells with DCA resulted in an enhancement of the S-G$_2$/M cell cycle arrest. The amount of cells in the S-G$_2$/M phase increased from ∼17.7% (control cells, 0 mM DCA) to ∼34.8%, 34.6%, and 36.7% for C6 cells treated with 5 mM, 20 mM, and 40 mM DCA, respectively.

Apoptosis of C6 cells were analyzed by flow cytometry using Annexin V-FITC/PI. The data indicate that DCA increased the percentage of total apoptotic cells in a dose-dependent manner after treatment with DCA ($P < 0.01$), as shown in Figure 3. The percentage of total apoptotic cells increased from ∼5.5% (control cells, 0 mM DCA) to ∼7.0%, 18.6%, and 24.2% for C6 cells treated with 5 mM, 20 mM, and 40 mM DCA, respectively.

**DCA inhibits Hsp70 expression in C6 cell line**

Hsp70 were quantified using a commercial Hsp70 ELISA kit. As shown in Figure 4, a dose-dependent and time-dependent inhibition of DCA on the level of Hsp70 has been found. Compared with the control group (0 mM DCA), the level of Hsp70 was significantly decreased, except for 0.05 mM at the 5-hour incubation time point ($P < 0.01$). The level of Hsp70 after 24 hours of incubation was also significantly decreased.
Figure 4 Effects of DCA on the level of Hsp70 in C6 cells.

Notes: C6 glioma cells were incubated with DCA (0 mM, 0.05 mM, 0.5 mM, and 5 mM) for 5 hours, 12 hours, or 24 hours, respectively. The control group was performed on C6 cells with absent DCA (0 mM DCA). Hsp70 were quantified using a commercial Hsp70 ELISA kit. Each sample was run in duplicate and compared with a standard curve. Each assay was carried out in triplicate. *P < 0.01 versus the control group (0 mM); †P < 0.05 or ††P < 0.01 versus the 5-hour incubation group; ‡P < 0.01 versus the 12-hour incubation group.

Abbreviations: DCA, dichloroacetate; ELISA, enzyme linked immunosorbent assay.

decreased compared with that after 5 hours or 12 hours of incubation (P < 0.01).

DCA inhibits C6 glioma tumor growth in vivo

In the in vivo antitumor activity experiments, the antitumor activity of DCA was evaluated by measuring the tumor weight in C6 glioma tumor-bearing rats after C6 cell implantation. In the present study, at days 19 and 21 posttumor inoculation, two tumor-bearing rats in the control group died with a tumor weight of 680 mg and 446 mg, respectively.

As shown in Figure 5A, DCA markedly inhibited the growth of C6 glioma tumors at doses of 25 mg/kg, 75 mg/kg, and 125 mg/kg (P < 0.01). There were no significant differences among the DCA treatment groups (25 mg/kg, 75 mg/kg, and 125 mg/kg). The average tumor weight in the distilled water, 25 mg/kg, 75 mg/kg, and 125 mg/kg DCA treatment groups at day 23 after C6 glioma cell implantation was 436 mg, 124 mg, 129 mg, and 125 mg, respectively. The values of TWI (%) in the 25 mg/kg, 75 mg/kg, and 125 mg/kg DCA treatment groups compared with that in the control group were ~71.5%, ~70.4%, and ~71.3%, respectively.

In another separate study, the male BALB/c nude mice were inoculated subcutaneously with C6 glioma cell suspension. Mice were randomly divided into a sterilized water group and three DCA treatment groups (25 mg/kg, 75 mg/kg, and 125 mg/kg). Each group consisted of seven tumor-bearing mice. As shown in Figure 5B, DCA markedly inhibited the growth of C6 glioma tumors at doses of 25 mg/kg, 75 mg/kg, and 125 mg/kg compared with that in sterilized water group (P < 0.01). There were also no significant differences among the DCA treatment groups (25 mg/kg, 75 mg/kg, and 125 mg/kg). At day 21 after C6 glioma cell implantation, the average tumor volume in sterilized water and DCA treatment groups (25 mg/kg, 75 mg/kg, and 125 mg/kg) were measured with calipers every 2 days. Tumor volumes were calculated from the formula:

\[
\text{Tumor volume (mm}^3) = (\text{major axis}) \times (\text{minor axis})^2 \times 0.5.
\]

Animals were sacrificed on day 21 posttumor inoculation. The tumors were harvested and weighed. Control group, sterilized water (n = 7); DCA treatment groups, at doses of 25 mg/kg, 75 mg/kg, or 125 mg/kg, respectively (n = 7). Columns or point, mean; bars, SD. *P < 0.01, versus control group.

Abbreviations: DCA, dichloroacetate; n, number; SD, standard deviation.
and 125 mg/kg) was 1940 ± 441 mm³, 801 ± 182 mm³, 845 ± 156 mm³, and 596 ± 107 mm³, respectively. The average tumor weight in the sterilized water, 25 mg/kg, 75 mg/kg, and 125 mg/kg DCA treatment groups at day 21 after C6 glioma cell implantation was 2539 mg, 924 mg, 1059 mg, and 864 mg, respectively (Figure 5C). The values of T2WI (%) in the 25 mg/kg, 75 mg/kg, and 125 mg/kg DCA treatment groups compared with that in sterilized water group were ~60.2%, ~54.5%, and ~62.9%, respectively.

**DCA increases ROS production, decreases MMP, and inhibits angiogenesis in tumor tissues in vivo**

The ROS production in tumor tissues after DCA administration to C6 tumor-bearing nude mice was evaluated. When C6 tumor-bearing nude mice were treated with DCA for 16 consecutive days (from day 6 to day 21), on day 21 after tumor inoculation, the ROS production in the tumor tissues was significantly increased compared with that in untreated animals (P < 0.01), as shown in Figure 6A. The ROS production in 25 mg/kg, 75 mg/kg, or 125 mg/kg DCA treatment groups was 2.5-, 3.1-, and 2.4-fold higher than that in the untreated group, respectively.

The MMP in tumor tissues after DCA administration to C6 tumor-bearing nude mice was also investigated. When tumor-bearing nude mice were treated with DCA for 16 consecutive days (from day 6 to day 21), on day 21 after tumor inoculation, the MMP in the tumor tissues was significantly decreased compared with that in untreated animals (P < 0.01), as shown in Figure 6B. The red/green fluorescence ratio in 25 mg/kg, 75 mg/kg, or 125 mg/kg DCA treatment groups was 77%, 77%, and 83% lower than that in the untreated group, respectively.

To evaluate the antiangiogenic activity of DCA in vivo, microvessel vessel density in tumor tissues after DCA administration to tumor-bearing rats was assessed by immunohistochemistry. As shown in Figure 7, microvessels were easily observed by CD31 staining. The numbers of the microvessels in the DCA treatment groups (25 mg/kg, 75 mg/kg, and 125 mg/kg) were 11 ± 1.4, 2.3 ± 0.6 and 1.3 ± 0.6 respectively, which were significantly lower than the number in the untreated group (14 ± 2.8).

**Discussion**

As a small molecule mitochondria-targeting agent, DCA can penetrate the BBB, showing potential therapeutic effects on brain tumors. In addition, the oral bioavailability of DCA is nearly 100%; therefore, the objective of the present study was to investigate the potential antitumor activity of DCA on brain tumors in vitro and in vivo.

The unique metabolism of most solid tumors integrates many proximal pathways and results in a consideration of mitochondria. The characteristic of tumor metabolism indicated that tumor cells rapidly use glucose and convert the majority of it to lactate, even in the presence of oxygen, which is the so-called aerobic glycolysis or Warburg effect. PDH is a metabolic switch that determines whether or not mitochondrial respiration or aerobic glycolysis should occur. It converts pyruvate...
DCA cells. 

explain DCA-inducing apoptosis and antiproliferation in suggested that the results of this cell cycle arrest may partly arrest with an accompanying decrease in the G1 phase. We revealed that DCA-induced C6 cells in the S phase cell cycle the induction of apoptosis.

in C6 cells, 

activity of PDH, induce ROS production, and decrease MMP 

 ROS are associated with opening of the mitochondrial transition pore. Opening of the MMP-sensitive mitochondrial transition pore allows for the efflux of cytochrome c and AIF from the mitochondria into the cytoplasm and induces apoptosis. 

Our previous in vitro results indicated that DCA can increase the activity of PDH, induce ROS production, and decrease MMP in C6 cells, 

 showing that DCA can induces the apoptosis of C6 cells through the activation of the mitochondrial pathway.

Cell cycle control represents a major regulatory mechanism of cell growth. 

 Blockade of the cell cycle is regarded as an effective strategy for the development of novel cancer therapies. 

It has been reported that DCA treatment resulted in an increase in the proportion of tumor cells in the S phase, showing a decrease in proliferation as well as the induction of apoptosis. 

Our cell cycle analysis results revealed that DCA-induced C6 cells in the S phase cell cycle arrest with an accompanying decrease in the G1 phase. We suggested that the results of this cell cycle arrest may partly explain DCA-inducing apoptosis and antiproliferation in C6 cells.

Heat shock proteins (HSPs) are involved in protein folding, aggregation, transport, and/or stabilization by acting as a molecular chaperone, leading to the inhibition of apoptosis by both caspase-dependent and/or independent pathways. 

HSPs are overexpressed in a wide range of human cancers and are implicated in tumor cell proliferation, differentiation, invasion, and metastasis. It has been reported that Hsp70 expression was much more abundant in glioma tumor cells, such as C6 cells, than in normal brain tissues. Considering the fact that high expression of HSPs is essential for cancer survival, the inhibition of HSPs is an important strategy of anticancer therapy. Our current results indicated that DCA could down-regulate Hsp70 expression in C6 cells in a dose-dependent and time-dependent manner, showing a new pathway of DCA-induced C6 cell apoptosis.

Unlike cytotoxic chemotherapeutic agents, the half maximal inhibitory concentration (IC50) values of DCA were about 27 mM for C6 cells. Similar IC50 values of DCA (15–30 mM) for U251, SKOV-3, A549, and MDA-MB-231 cell lines were also observed (data not shown). These results indicated that DCA has unique antitumor mechanisms on tumor cell lines.

In the present research, the antitumor activity of DCA after oral administration was investigated in C6 brain tumor-bearing rats and C6 tumor-bearing nude mice in vivo. Our in vivo antitumor activity results indicated that DCA markedly inhibited the growth of C6 glioma tumor in both C6 brain tumor-bearing rats and C6 tumor-bearing nude mice (P < 0.01). We also found no significant difference in the antitumor activity among the DCA treatment groups (25, 75 and 125 mg/kg). It was reported that DCA administered at 35–50 mg/kg decreases lactate levels by more than 60% and directly activates PDH by threelfold to sixfold. 

Our in vivo antitumor activity results suggested that DCA might possibly produce the antitumor activity at a dose lower than 25 mg/kg. In a clinical experiment, each of five glioblastoma patients was treated with oral DCA for up to 15 months. The starting oral dose of DCA was 12.5 mg/kg twice a day for 1 month, at which point the dose was increased to 25 mg/kg orally twice a day, if dose-limiting toxicity occurred, decreasing the dose by 50%. 

The efficacy and safety of DCA on the treatment of glioblastoma was confirmed, even at a dose of 6.25 mg/kg orally twice a day. 

In addition, after 5 years of continued treatment with oral DCA at a dose of 25 mg/kg, the serum DCA levels are only slightly increased compared with the levels after the first several doses, also showing its safety for oral administration at this dose.
DCA can enter the circulation rapidly after oral administration and then generate the stimulation of PDH activity generally within minutes. Our in vivo results in tumor tissues indicated that DCA significantly induced ROS production and decreased MMP in tumor tissues, showing similar mechanisms with those observed in vitro.

Histologically, glioblastomas are highly angiogenic and characterized by microvascular proliferations. In the present research, the angiogenic activity of DCA was investigated in vivo by immunohistochemistry. The numbers of microvessels in the DCA treatment groups were significantly decreased, suggesting the potential antiangiogenic effect of DCA. It has been reported that early carcinogenesis often occurs in a hypoxic environment. Under hypoxic conditions, hypoxia-inducible factor (HIF-1α) is activated and induces angiogenesis. In addition, HIF-1α can also induce the expression of PDK, which can inhibit the activity of PDH. The inhibition effect of DCA on HIF-1α would decrease vascular endothelial growth factor and inhibit angiogenesis. In addition, because the cancer cells were “normalized” following the increase in mitochondrial respiration and mitochondrial depolarization, we suggested that the hypoxic microenvironment in or around tumor tissue would be improved. We also found that, unlike the results obtained from in vivo antitumor experiments, the antiangiogenic effect in the 25 mg/kg treatment group was lower than that in 75 mg/kg or 125 mg/kg treatment groups. Therefore, we suggested that the inhibition of DCA on HIF-1α or vascular endothelial growth factor would be further investigated.

In conclusion, DCA induces the apoptosis of C6 cells through the activation of the mitochondrial pathway, arresting the cell cycle of C6 cells in S phase and down-regulating Hsp70 expression. Our in vivo antitumor experiment results indicated that DCA markedly inhibited the growth of C6 glioma tumors in both C6 brain tumor-bearing rats and C6 tumor-bearing nude mice ($P < 0.01$). DCA significantly induced the ROS production and decreased the MMP in tumor tissues. Our in vivo antitumor activity results also indicated that DCA has an antiangiogenic effect.

Acknowledgments

The authors gratefully acknowledge financial support from the National Natural Science Foundation of China (No 81172992) and the National Basic Research Program of China (973 Program 2009CB930300 and 2013CB932501) and Innovation Team of the Ministry of Education (No BMU20110263).

Disclosure

The authors report no conflicts of interest in this work.

References

1. Van Meir EG, Hadjipanayis CG, Norden AD, Shu HK, Wen PY, Olson JJ. Exciting new advances in neuro-oncology: the avenue to a cure for malignant glioma. CA Cancer J Clin. 2010;60(3):166–193.
2. Khasraw M, Lassman AB. Advances in the treatment of malignant gliomas. Curr Oncol Rep. 2010;12(1):26–33.
3. Avgeropoulos NG, Batchelor TT. New treatment strategies for malignant gliomas. Oncologist. 1999;4(3):209–224.
4. Stieber VW, Mehta MP. Advances in radiation therapy for brain tumors. Neurol Clin. 2007;25(4):1005–1033, ix.
5. Souweidane MM, Luther N. Endoscopic resection of solid intraventricular brain tumors. J Neurosurg. 2006;105(2):271–278.
6. Stummer W, Pichlmayer U, Meinel T, Wiestler OD, Zanella F, Reulen HJ; for ALA-Glioma Study Group. Fluorescence-guided surgery with 5-aminolevulinic acid for resection of malignant glioma: a randomised controlled multicentre phase III trial. Lancet Oncol. 2006;7(5):392–401.
7. El Hallani S, Boisselier B, Peglion F, et al. A new alternative mechanism in glioblastoma vascularization: tubular vasogenic mimicry. Brain. 2010;133(Pt 4):973–982.
8. Buie LW, Valgus J. Bevacizumab: a treatment option for recurrent glioblastoma multiforme. Ann Pharmacother. 2008;42(10):1486–1490.
9. Liu Y, Zheng T, Zhao S, et al. Inhibition of heat shock protein response enhances PS-341-mediated glioma cell death. Ann Surg Oncol. 2012;19 Suppl 3:S421–S429.
10. Fedrigo CA, Grivicich I, Schumemann DP, et al. Radiosensitivity of human glioma spheroids and expression of HSP70, p53 and EGFr. Radiat Oncol. 2011;6:156.
11. Garrido C, Gurbuxani S, Ravagnan L, Kroemer G. Heat shock proteins: endogenous modulators of apoptotic cell death. Biochem Biophys Res Commun. 2001;286(3):433–442.
12. Koo YE, Reddy GR, Bhojani M, et al. Brain cancer diagnosis and therapy with nanoplatforms. Adv Drug Deliv Rev. 2006;58(14):1556–1577.
13. Partridge WM. BBB-Genomics: creating new openings for brain-drug targeting. Drug Discov Today. 2001;6(8):381–383.
14. Gan CW, Feng SS. Transferrin-conjugated nanoparticles of poly(lactide)-D-alpha-tocopheryl polyethylene glycol succinate diblock copolymer for targeted drug delivery across the blood-brain barrier. Biomaterials. 2010;31(30):7748–7757.
15. Chertok B, David AE, Yang VC. Polyethyleneimine-modified iron oxide nanoparticles for brain tumor drug delivery using magnetic targeting and intra-carotid administration. Biomaterials. 2010;31(24):6317–6324.
16. Ke XY, Zhao BJ, Zhao X, et al. The therapeutic efficacy of conjugated linoleic acid—paclitaxel on glioma in the rat. Biomaterials. 2010;31(22):5855–5864.
17. Pan JG, Mak TW. Metabolic targeting as an anticancer strategy: dawn of a new era? Sci STKE. 2007;2007(381):pe14.
18. Samudio I, Fiegl M, Andreeff M. Mitochondrial uncoupling and the Warburg effect: molecular basis for the reprogramming of cancer cell metabolism. Cancer Res. 2009;69(6):2163–2166.
19. Puthenveedu P, Millard M, Neamati N. Opportunities in discovery and delivery of anticancer drugs targeting mitochondria and cancer cell metabolism. Adv Drug Deliv Rev. 2009;61(14):1250–1275.
20. Warburg O. On the origin of cancer cells. Science. February 24, 1956; 123(3191):309–314.
21. Michelakis ED, Webster L, Mackey JR. Dichloroacetate (DCA) as a potential metabolic-targeting therapy for cancer. Br J Cancer. 2008;99(7):898–999.
22. Yeung SJ, Pan J, Lee MH. Roles of p53, MYC and HIF-1 in regulating glycolysis—the seventh hallmark of cancer. Cell Mol Life Sci. 2008;65(24):3981–3999.
23. Fulda S, Galluzzi L, Kroemer G. Targeting mitochondria for cancer therapy. *Nat Rev Drug Discov*. 2010;9(6):447–464.
24. Coude FX, Saudubray JM, DeMaugre F, Marsac C, Leroux JP, Charpentier C. Dichloroacetate as treatment for congenital lactic acidosis. *N Engl J Med*. 1978;299(24):1365–1366.
25. Stacpoole PW, Wright EC, Baumgartner TG, et al. A controlled clinical trial of dichloroacetate for treatment of lactic acidosis in adults. *The Dichloroacetate-Lactic Acidosis Study Group. N Engl J Med*. 1992;327(22):1564–1569.
26. Bonnet S, Archer SL, Allalunis-Turner J, et al. A mitochondria-K+ channel axis is suppressed in cancer and its normalization promotes apoptosis and inhibits cancer growth. *Cancer Cell*. 2007;11(1):37–51.
27. Wong JY, Huggins GS, Debidda M, Munshi NC, De Vivo I. Dichloroacetate induces apoptosis in endometrial cancer cells. *Gynecol Oncol*. 2008;109(3):394–402.
28. Cao W, Yacoub S, Shiverick KT, et al. Dichloroacetate (DCA) sensitizes both wild-type and over expressing Bel-2 prostate cancer cells in vitro to radiation. *Prostate*. 2008;68(11):1223–1231.
29. Kuroda Y, Toshima K, Watanabe T, et al. Effects of dichloroacetate on pyruvate metabolism in rat brain in vivo. *Pediatr Res*. 1984;18(10):936–938.
30. Abemayer E, Kovachich GB, Haugaard N. Effects of dichloroacetate on brain pyruvate dehydrogenase. *J Neurochem*. 1984;42(1):38–42.
31. Viechtl V, Kirtikara K. Sulfuric acid B colorimetric assay for cytotoxicity screening. *Nat Protoc*. 2006;1(3):1112–1116.
32. Huang Y, Chen XM, Zhao BX, et al. Antiangiogenic activity of sterically stabilized liposomes containing paclitaxel (SSL-PTX): in vitro and in vivo. *AAPS Pharm Sci Tech*. 2010;11(2):752–759.
33. Petrangolini G, Supino R, Pratesi G, et al. Effect of a novel vascular-H+-ATPase inhibitor on cell and tumor response to camptothecins. *J Pharmacol Exp Ther*. 2006;318(3):939–946.
34. Zhao BJ, Ke XY, Huang Y, et al. The antiangiogenic efficacy of NGR-modified PEG-DSPE micelles containing paclitaxel (NGR-M-PTX) for the treatment of glioma in rats. *J Drug Target*. 2011;19(5):382–390.
35. Shannugam M, McBrayer SK, Rosen ST. Targeting the Warburg effect in hematological malignancies: from PET to therapy. *Curr Opin Oncol*. 2009;21(6):531–536.
36. Stacpoole PW, Henderson GN, Yan Z, James MO. Clinical pharmacology and toxicity of dichloroacetate. *Environ Health Perspect*. 1998;106 Suppl 4:989–994.
37. Zhao X, Wang X, Yu KF, et al. Effects of dichloroacetate on the activation of the mitochondrial pathway in C6 cells in vitro. *Journal of Chinese Pharmaceutical Sciences*. 2011;20(5):460–465.
38. Jeong HW, Han DC, Son KH, et al. Antitumor effect of the cinnamaldehyde derivative CB403 through the arrest of cell cycle progression in the G2/M phase. *Biochem Pharmacol*. 2003;65(8):1343–1350.
39. Buolamwini JK. Cell cycle molecular targets in novel anticancer drug discovery. *Curr Pharm Des*. 2000;6(4):379–392.
40. McDonald ER 3rd, El-Deiry WS. Cell cycle control as a basis for cancer drug development (Review). *Int J Oncol*. 2000;16(5):871–886.
41. Madhok BM, Yeluri S, Perry SL, Hughes TA, Jayne DG. Dichloroacetate induces apoptosis and cell-cycle arrest in colorectal cancer cells. *Br J Cancer*. 2010;102(12):1746–1752.
42. Sarkar R, Mukherjee S, Biswas J, Roy M. Sulphoraphane, a naturally occurring isothiocyanate induces apoptosis in breast cancer cells by targeting heat shock proteins. *Biochem Biophys Res Commun*. 2012;427(1):80–85.
43. Howlett RA, Heigenhauser GJ, Hultman E, Hollidge-Horvat MG, Spriet LL. Effects of dichloroacetate infusion on human skeletal muscle metabolism at the onset of exercise. *Am J Physiol*. 1999;277(1 Pt 1):E18–E25.
44. Parolin ML, Spriet LL, Hultman E, et al. Effects of PDH activation by dichloroacetate in human skeletal muscle during exercise in hypoxia. *Am J Physiol Endocrinol Metab*. 2000;279(4):E752–E761.
45. Michelakis ED, Sutendra G, Dromparis P, et al. Metabolic modulation of glioblastoma with dichloroacetate. *Sci Transl Med*. 2010;2(31):31ra34.
46. Mori M, Yamagata T, Goto T, Saito S, Momoi MY. Dichloroacetate treatment for mitochondrial cytopathy: long-term effects in MELAS. *Brain Dev*. 2004;26(7):453–458.
47. Louis DN, Ohgaki H, Wiestler OD, et al. The 2007 WHO classification of tumours of the central nervous system. *Acta Neuropathol*. 2007;114(2):97–109.
48. Cairns RA, Papandreu I, Supthin PD, Denko NC. Metabolic targeting of hypoxia and HIF1 in solid tumors can enhance cytotoxic chemotherapy. *Proc Natl Acad Sci U S A*. 2007;104(22):9445–9450.