Lactic Acid Promotes Cell Survival by Blocking Autophagy of B16F10 Mouse Melanoma Cells under Glucose Deprivation and Hypoxic Conditions

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In solid tumors, cancer cells are exposed to microenvironment stress, including hypoxia and insufficient nutrients. An acidic microenvironment in tumors is facilitated by the increase in synthesis of lactic acid; this is known as Warburg effect. We previously showed that B16F10 melanoma cells were induced autophagic cell death by glucose-deprivation stress, and lactic acid suppressed the cell death through the inhibition of autophagy. In this study, effects of lactic acid on cell death of B16F10 cells under hypoxic and glucose-depleted double stress conditions were investigated. The double stress promoted autophagic cell death earlier than glucose-depleted stress alone. Lactic acid repressed the double stress-induced cell death by inhibiting autophagy. These results suggest that lactic acid serves for cell survival under microenvironmental stress conditions in B16F10 melanoma cells.

Key words lactic acid; hypoxia; glucose deprivation; autophagy; cell death

INTRODUCTION

The environment of solid tumors is low-nutrition and hypoxia due to the rapid proliferation of tumor cells and immature vascularization. Glucose is catabolized into lactic acid to enhance aerobic glycolysis, known as Warburg effect, and causes the tumor microenvironment to be acidic. We previously demonstrated that glucose deprivation induced autophagic cell death of B16F10 melanoma cells, but the acid conditions, which are derived from lactic acid or pH-adjustment by HCl, inhibited the cell death mechanism. In vivo, tumor cells are exposed to not only glucose deprivation but also hypoxia. The effects of lactic acid on the double stress of glucose deprivation and hypoxia remain unclear. In this study, we revealed the role of lactic acid in autophagic cell death under double stress conditions in B16F10 melanoma cells.

MATERIALS AND METHODS

Cell Culture, Double Stress Induction, and the Effect on Autophagy Preparation of B16F10 mouse melanoma cells (B16F10 cells), the cell culture, treatment, and analytical methods were previously described. For induction of the double stress of glucose deprivation and hypoxia, B16F10 cells were seeded at 2 × 10⁵ cells/mL medium into 24-well plates. After 24h, the growth medium was changed to a glucose-free medium. The cells were maintained under 20% or 1% O₂ conditions for 16h. The dead cell ratio and viable cell number were calculated by cell counting. Autophagy inhibitor 3-methyladenine (5 mM) and LY294002 (25 µM) was used to analyze the effects of the double stress on autophagy.

Effect of Lactic Acid on B16F10 Cells under the Double Stress Conditions To analyze the effects of lactic acid on the double-stressed B16F10 cells, they were treated with lactic acid (20 mM)-supplemented glucose-free medium for 16h.

Lactic acid-free medium supplemented with sodium lactate (20 mM) was used as a control. In addition, HCl-adjusted Dulbecco’s modified Eagle’s medium (DMEM), which is a same condition as 20 mM lactic acid DMEM (pH 6.9), was used to analyze the effects of acidic environment without lactic acid.

Western Blot Analysis B16F10 cells were cultured in DMEM without glucose under 20 or 1% O₂ conditions at 16h. Western blot analysis using anti-LC3B and anti-β-actin antibodies was performed, as previously described. The ratio of

Fig. 1. Double Stress by Glucose Deprivation and Hypoxia Induces Cell Death and Autophagy in B16F10 Cells

B16F10 cells were subjected to the double stress for 16h. (A) Dead cell rate and (B) relative cell number were evaluated by cell counting. The results are represented by the mean ± standard deviation (S.D.) of triplicate counts. *p<0.01 and **p<0.001 indicate significant differences compared with 20% O₂ conditions (Student’s t-test). (C) LC3B conversion in B16F10 cells was examined by Western blot analysis after 16h of double stress. ‘O₂ conc.’ represents ‘O₂ concentration.’

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**RESULTS**

Effects of the double stress of glucose deprivation and hypoxia on B16F10 cells were compared to the effects of the single stress of glucose deprivation. The double stress of glucose deprivation and hypoxia induced more cell death at 16 h than the glucose deprivation stress alone (Figs. 1A, B). Western blot analysis showed about 2.0-fold higher conversion of LC3B-I to LC3B-II, an autophagy marker, in cells under double stress conditions than in those under glucose deprivation alone (Fig. 1C). Treatment of B16F10 cells under glucose deprivation and hypoxic conditions with autophagy inhibitor LY294002 and 3-MA suppressed cell death (Figs. 2A, B). The ratio of LC3B-II/LC3B-I in B16F10 cells was examined at 16 h after treatment of LY294002 or 3-MA.

**Statistical Analysis**  
Student’s t-test or one-way ANOVA (post-hoc test: Tukey–Kramer test) were used to evaluate the statistical significance of the rate of cell death and cell number. A $p < 0.05$ was considered significant.

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Fig. 2. Inhibition of Autophagy Suppresses Cell Death in B16F10 Cells under the Stress of Glucose Deprivation and Hypoxic Conditions

(A) Dead cell rate and (B) relative cell number of B16F10 cells were calculated 16 h after treatment with 25 µM LY294002 or 5 mM 3-MA. The bars represent the mean ± S.D. of triplicate counts. $p < 0.001$ (comparison to 20% O$_2$ conditions), $p < 0.001$ (comparison to 1% O$_2$ conditions without treatment of autophagy inhibitor) indicate significant differences (Tukey–Kramer test). 1% O$_2$ (−) and LY indicate 20% O$_2$ conditions and LY294002 treatment, respectively. (C) LC3B conversion in B16F10 cells was examined at 16 h after treatment of LY294002 or 3-MA.

Fig. 3. Lactic Acid Represses Double Stress-Induced Cell Death and Autophagy in B16F10 Cells

B16F10 cells were cultured under glucose deprivation and 1% O$_2$ conditions with (+) or without (−) 20 mM lactic acid for 16 h. (A) Dead cell rate and (B) relative cell number were analyzed. The results represent the mean ± S.D. in triplicate. $p < 0.001$ (compared to 20% O$_2$ without lactic acid conditions), $p < 0.001$ and $p < 0.05$ (compared to 1% O$_2$ without lactic acid conditions) indicate significant differences (Tukey–Kramer test). (C) Western blot analysis with anti-LC3B and anti-β-actin antibodies was performed using B16F10 cells under double stress conditions.

Fig. 4. HCl-Adjusted Acidic Condition Suppresses Cell Death and Autophagy under Glucose Deprivation and Hypoxic Conditions

B16F10 cells were cultured in DMEM (pH 6.9) adjusted by HCl under glucose deprivation and 1% O$_2$ conditions for 16 h. (A) Dead cell rate and (B) relative cell number were analyzed. The results represent the mean ± S.D. in triplicate. $p < 0.001$ (compared to 20% O$_2$ without HCl conditions), $p < 0.001$ and $p < 0.05$ (compared to 1% O$_2$ without HCl conditions) indicate significant differences (Tukey–Kramer test). (C) LC3B conversion in B16F10 cells was examined by Western blot analysis with anti-LC3B antibody after 16 h of double stress.
was about 40 or 10% lower than untreated-cells under glucose deprivation and 1% O₂ conditions (Fig. 2C).

The effects of lactic acid on the double stress-inducing autophagic cell death in B16F10 cells were examined. As expected, lactic acid inhibited cell death and increased survival cell number (Figs. 3A, B). The LC3B conversion in B16F10 cells treated with 20 mM lactic acid conditions was about 60% lower than untreated-cells under glucose deprivation and 1% O₂ conditions (Fig. 3C). Furthermore, acidic DMEM adjusted by HCl without lactic acid also inhibited cell death under the double stress conditions (Figs. 4A, B). Western blot analysis showed that LC3B conversion of B16F10 cells cultured in DMEM adjusted by HCl was over 90% decrease compared to 1% O₂ without pH-adjustment by HCl (Fig. 4C).

DISCUSSION

In this study, we analyzed the effects of lactic acid on B16F10 cells that were subjected to the microenvironment stresses of glucose deprivation and hypoxia. Autophagy is induced when cells are subjected to stresses such as nutrient deprivation. Intracellular proteins and organelles are degraded and recycled as an alternative energy source. Therefore, autophagy is important for cell survival.1,9 However, prolonged autophagy induces self-digestion and triggers cell death.10,11 In tumor cells, the role of autophagy in response to microenvironmental stress including glucose deprivation and hypoxia has been reported both cell survival and cell death.12-16 We previously showed that glucose deprivation stress induced cell death by autophagy in B16F10 cells.7 In this study, the induction of hypoxic stress to glucose depletion induced autophagic cell death more than glucose depletion stress alone (Fig. 1). Although inhibition of autophagy by LY294002 and 3-MA suppressed glucose deprivation-induced cell death and autophagy in B16 melanoma cells. Biochem. Biophys. Res. Commun., 496, 1357–1361 (2018).

The authors declare no conflict of interest.

REFERENCES