Amide-linked local anesthetics induce apoptosis in human non-small cell lung cancer

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Background: A retrospective analysis of patients undergoing cancer surgery suggested that using local anesthetics could reduce cancer recurrence and improve survival rate. Previous studies have indicated that local anesthetics may induce apoptosis in several kinds of cells in vitro, but the mechanism is unclear.

Methods: Cell viability was analyzed by MTS; reactive oxygen species (ROS), mitochondrial membrane potential (MMP, ∆Ψm), cell cycle distribution, and cell apoptosis assay were detected by flow cytometry; DNA damage was measured by comet assay; cell invasion and migration were observed by microscopy; The expression level of related proteins was detected by western blot assay.

Results: The results indicated that lidocaine and ropivacaine could decrease viability, induce G0/G1 phase arrest and apoptosis in human non-small cell lung cancer (NSCLC) cells A549 and H520. Invasion and migration were suppressed. Western blot indicated the related apoptotic pathways proteins changed accordingly. Additionally, lidocaine and ropivacaine downregulated ∆Ψm, provoked DNA damage, upregulated ROS production and activated mitogen-activated protein kinase (MAPK) pathways in A549 and H520 cells.

Conclusions: The cytotoxic effect of amide-linked local anesthetics on NSCLC cells were mainly due to apoptosis. The antitumor mechanism of lidocaine and ropivacaine may involve apoptotic pathways and MAPK pathways.

Keywords: Local anesthetics; apoptosis; apoptotic pathways; mitogen-activated protein kinase pathways (MAPK pathways); non-small cell lung cancer (NSCLC)

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Introduction

Local anesthetics have a broad pharmacological roles that go beyond the known analgesia and antiarrhythmic (1). They are used in a wide range of clinical situations to prevent or reduce acute pain, chronic pain and cancer pain (2). A retrospective analysis of patients undergoing cancer surgery suggests that using regional anesthesia may reduce cancer recurrence and improve survival rate (3,4). Recently, some studies have demonstrated that local anesthetics inhibit proliferation, suppress invasion and migration, and induce apoptosis at a range of certain concentrations (5-7). The mechanisms are still unclear. It seems to be unrelated to the sodium-channel blockade (8-10), while in other reports local anesthetics work in the manner of inhibiting the activity of sodium channels (11,12). Previous study has suggested that local anesthetics could induce apoptosis in human thyroid cancer cells, which is associated with mitogen-activated protein kinase (MAPK) pathways (13). Another report has suggested that inhibition of MAPK pathways protects against local anesthetics-induced neurotoxicity (14). However, little
is known about local anesthetics-induced cytotoxicity in human non-small-cell lung cancer (NSCLC) cells.

Methods

Cells lines and culture conditions

Human NSCLC cell lines A549 and H520 were purchased from the Institute of Biochemistry and Cell Biology, Chinese Academy of Science, China. Cells were cultured in a RPMI-1640 medium supplemented with 10% fetal bovine serum, 1% penicillin (100 U/mL) and streptomycin (100 μg/mL) (Gibco, NY, USA). Both cell lines were incubated at 37 °C in a humid incubator with 5% CO₂.

MTS assay

Approximately 5×10³ cells/well were placed in a 96-well plate and then treated with lidocaine or ropivacaine individually at concentrations ranging from 0 to 32 mM for 24, 48, or 72 h. In each well, 100 μL MTS reagent (Promega, WI, USA) was added, followed by incubated in the dark at 37 °C for 1.5 h. The absorbance values were measured at 490 nm using Varioskan Flash (Thermo Fisher Scientific Inc., MA, USA). The median 50% effective dose (ED₅₀) values were calculated using the probit method of Miller and Tainter.

Cell cycle assay

Cells were exposed to lidocaine or ropivacaine for 24 h at the concentration of 24 h ED₅₀, then washed by phosphate-buffered saline (PBS) and fixed with 75% ethanol overnight at 4 °C. After that, cells were incubated with Rnase (0.1 mg/mL) for 30 min, followed by stained with 40 μg/mL propidium iodide (PI) for another 30 min. Analysis was performed using the Cell Quest software of a Becton-Dickinson fluorescence-activated cell sorting (FACS) Calibur flow cytometer (Becton-Dickinson, CA, USA), the excitation wavelength was set at 488 nm.

Apoptosis assay

Cells were collected after treated with lidocaine or ropivacaine for 24 h, and a minimum of 15,000 cells were analyzed in each measurement. Cells were stained with FITC-conjugated anti-annexin V antibody and PI, then quantified by FACS Calibur flow cytometer with a 488 nm argon laser. The cells in the early state of apoptosis were stained with annexin V, while the late state of apoptosis was stained with PI.

Invasion and migration assays

The upper chamber of a 6.5 mm Transwell polycarbonate membrane were coated with diluted matrigel (3.9 mg/mL, 60–80 μL) inserting with 8 μm pores (Coster, MA, USA). Cells were resuspended in 300 μL serum-free RPMI-1640 medium with no other supplements or with local anesthetic were incubated for 15 min at room temperature. The inserts were then placed into 500 μL complete medium (RPMI-1640, 10% fetal bovine serum, 1% penicillin and streptomycin plus the same concentration of local anesthetic as present in the upper chamber) in a 24-well plate. After incubation with lidocaine or ropivacaine for 24 h, the upper surface cells were scraped, however, the lower side cells were fixed with 75% ethanol, followed by stained with crystal violet. The migration assay was conducted similarly, with no matrigel on the upper chamber. The inverted microscope was used to count the number of cells at three randomly selected visual fields with 400× magnification.

Detection of intracellular reactive oxygen species (ROS)

The intracellular ROS level was detected using an oxidation-sensitive fluorescent probe. The cells were plated at a density of 3×10⁵ cells/well in 6-well plates and treated with lidocaine or ropivacaine for another 24 h. Subsequently, cells were centrifuged and resuspended in 500 μL of 2,7-dichlorofluorescin diacetate (10 μM, DCFH-DA, Molecular Probes, OR, USA) for ROS detection. After incubated at 37 °C for 30 min, cells from each treatment were analyzed by flow cytometry.

Detection of mitochondrial membrane potential (MMP, ΔΨm)

MMP was determined by flow cytometry using the ΔΨm-dependent fluorescent dye JC-1 (Sigma, MO, USA). The cationic dye JC-1 was a highly specific probe for detecting changes in ΔΨm to evaluate mitochondrial membrane integrity for which could selectively enter into mitochondria and undergo a reversible change in fluorescence emission according to the ΔΨm. Approximately 3×10⁵ cells/well were cultured in a 6-well plate. After treated with local anesthetic for 24 h, cells were harvested and incubated with JC-1 for 20 min at 37 °C according to the manufacturer's
instructions (Sigma, MO, USA). Then the samples were subjected to ΔΨm determination by flow cytometry as JC-1 formed red fluorescence in intact mitochondria, while green fluorescence was formed in JC-1 monomers at low ΔΨm.

**Comet assay**

Comet assay was used to evaluate the effect of local anesthetics on DNA damage of NSCLC cells. Cells were harvested and resuspended at a density of 1×10⁵/mL to spread on microscopic slides, precoated with a thin layer of 0.5% melting agarose. After gelling for 15 min at 4 °C, slides were incubated in lysis solution for 1.5 h. Afterward, each slide was placed in a tank containing balanced solution for another 20 min. Subsequently, each slide was subjected to electrophoresis at 30 V for 40 min. After immersed in neutralization buffer for 5 min and stained with SYBR Green (Invitrogen, CA, USA), the slides were observed with an FV-1000 laser scanning fluorescence microscope (Carl Zeiss, Oberkochen, Germany). The tails of the comet reflected the DNA damage.

**Western blot assay**

Equivalent amount of proteins (30 μg) were individually subjected to gel electrophoresis. The proteins were incubated with the indicated primary antibodies: Fas and FasL, Bax and Bcl-2, endonuclease G (Endo G), apoptosis-inducing factor (AIF), cytochrome c, caspase and cleaved caspase-3, -8, -9, poly ADP-ribose polymerase (PARP) and cleaved PARP, cyclin D1, total extracellular signal-regulated protein kinases (ERKs), total c-Jun NH2-terminal kinases (JNKs), total p38 MAPK, p-ERKs, p-JNKs, and p-p38. Anti-β-actin was used as a loading control. Corresponding horseradish peroxidase-conjugated secondary antibodies were used against each primary antibody. Proteins were detected using the chemiluminescent detection reagents.

**Statistics analysis**

Data were showed as mean ± standard deviation (SD). Statistical analysis of the differences between two groups was evaluated using the one-way analysis of variance, followed by Student's t-test using the SPSS 16.0 software (SPSS Inc., IL, USA). Values of P<0.05 were considered significant differences (*P<0.05; **P<0.01; ***P<0.001).

**Results**

**Local anesthetics suppressed NSCLC cell viability**

The cell viability assay showed that local anesthetics
suppressed the growth of NSCLC cells in a dose- and time-dependent manner (Figure 1). The ED$_{50}$ values of lidocaine and ropivacaine in A549 cells were higher than that in H520 cells. The ED$_{50}$ of lidocaine was 9.51 and 6.14 mM for A549 and H520 cells at 24 h, respectively. In addition, the ED$_{50}$ of ropivacaine at 24 h was 4.06 and 2.62 mM for A549 and H520 cells, respectively.

**Local anesthetics arrested NSCLC cell cycle at the G0/G1 phase**

As shown in Figure 2A, the percentage of cells in the G0/G1 phase in the treated groups were significantly increased. The cell cycle distribution analysis indicated 91.51%±1.53% A549 and 87.72%±1.60% H520 for lidocaine-treated groups compared with 69.53%±1.62% A549 and 64.94%±1.44% H520 for negative control (NC) groups at the G0/G1 phase (**P<0.001). Simultaneously, ropivacaine-treated groups were 92.57%±1.57% A549 and 86.65%±1.27% H520 at the G0/G1 phase (**P<0.001) (Figure 2A).

**Local anesthetics induced NSCLC cells apoptosis**

The total percentage of apoptosis (including early and late apoptosis) was significantly increased in the treated groups (**P<0.001) (Figure 2B).

**Local anesthetics inhibited invasion and migration of NSCLC cells**

The invasion capability of local anesthetics-treated groups decreased in comparison with NC groups, as the number of cells invading through the membrane matrix was obviously decreased (**P<0.001) (Figure 3). Similar to invasion, the migration was also drastically suppressed in local anesthetics-treated groups than in NC groups (**P<0.001) (Figure 3).

**Local anesthetics induced mitochondrion and DNA damage**

The level of ROS increased and ΔΨm decreased
After local anesthetics treated for 24 h at the concentrations of ED$_{50}$, the number of cells was counted at 3 randomly selected visual fields by an inverted microscope at 400× magnification. (A) Local anesthetics inhibited migration and invasion of A549 cells; (B) local anesthetics inhibited migration and invasion of H520 cells (**P<0.001).

Altered expression of related proteins

Western blot assay showed that the expression level of cyclin D1 decreased in local anesthetics-treated groups in comparison with NC groups, which was inconsistent with the result of G0/G1 phase arrest (Figure 6A). In addition, the expression level of the intrinsic mitochondrial pathway proteins Bax, Bcl-2, cleaved caspase-9, cytochrome c, AIF and Endo G changed accordingly (Figure 6B,C). Caspase-3 and PARP, the common proteins of two apoptotic pathways, were cleaved after local anesthetics treatment. Furthermore, the expression levels of the extrinsic death receptor pathway proteins Fas and its receptor FasL, cleaved caspase-8 were upregulated (Figures 6C). Moreover, local anesthetics did not alter the expression of p38 MAPK and its phosphorylation (Figure 6D). Unlike p38 MAPK, lidocaine and ropivacaine had no obvious effect on total ERK1/2 and JNK levels; however, the increased phosphorylation of ERK1/2 and JNK were observed (Figure 6D). In total, lidocaine and ropivacaine triggered apoptosis in human NSCLC cells via apoptotic pathways and MAPK pathways (Figure 7).

Discussion

As the leading cause of cancer-related mortality globally, the annual burden of lung cancer is larger than that of any other cancers, for which more than 85% of those cases are currently classified as NSCLC (15-17). Despite recent advances in diagnosis and treatment, the predicted 5-year survival rate is only 15.9% which has only marginally improved during the past decades (18). Thus, the underlying molecular mechanisms and new therapeutic strategies are urgently required in lung cancer.

Retrospective studies of patients undergoing cancer surgeries suggest that using regional anesthesia reduces the risk of tumor metastasis and recurrence, but the mechanism remains unclear (19-21). The benefits may be due to the attenuation of immunosuppression by regional anesthesia (12,22). Some in vitro animal data demonstrate that opioids promote tumor growth and metastasis, largely by inducing mitogenesis and angiogenesis (23,24). Regional anesthesia, in part, reduces the use of opioids, and thus may reduce tumor recurrence and improve survival. However, Doornbal et al. study shows that morphine does not facilitate breast cancer progression (25). Thus, further studies need to be conducted for the specific mechanisms of opioids on cancer. Apart from the preservation of immune system and the reduction in opioids requirement, systemic administration of local anesthetics during surgery plays a
Figure 4 Cells were treated with local anesthetics for 24 h at the concentrations of ED\textsubscript{50}. (A) Local anesthetics increased ROS production in A549 and H520 cells; (B) local anesthetics downregulated ∆Ψ\textsubscript{m} in both A549 and H520 cells (**P<0.001).

Figure 5 Cells were treated with local anesthetics for 24 h at the concentrations of ED\textsubscript{50}. The tails of the comet reflected the DNA damage.
Figure 6 Cells were treated with local anesthetics at the concentrations of ED50 for 24 h and then total proteins were detected by western blot using beta-actin as an internal control. (A) Local anesthetics downregulated the expression level of cyclin D1 compared with NC groups; (B) local anesthetics affected the protein levels of the intrinsic mitochondrial pathway; (C) local anesthetics affected the protein levels of the extrinsic death receptor pathway; (D) local anesthetics activated MAPK pathways.

Figure 7 Amide-linked local anesthetics induce apoptosis through apoptotic pathways (the extrinsic death receptor pathway and the intrinsic mitochondrial pathway) and MAPK pathways in human non-small-cell lung cancer cells.
role of anti-hyperalgesic and anti-inflammatory (26,27). One paramount benefit of local anesthetics is that they may induce apoptosis in tumor cells but not in normal tissues (23).

The effects of lidocaine and ropivacaine on NSCLC cells in vitro were examined in the present study, as they are the two most commonly used amide-linked local anesthetics in China. Our study demonstrated that lidocaine and ropivacaine inhibited cell growth and arrested cell cycle at G0/G1 phase. Once the cells from the G1 phase moved into the S phase, they could no longer rely on external stimuli, and complete the cell division automatically (28). In all known cell cycle proteins, cyclin D1 was the most critical checkpoint protein in regulating G1 phase to S phase (28). Our study demonstrated that the expression of cyclin D1 was downregulated which could prevent cells move from G1 to S phase thus inhibiting cell growth. The overexpression of cyclin D1 was associated with poor prognosis, and could significantly reduce postoperative long-term survival rate (28). Thus, downregulation the expression and function of cyclin D1 have become one of the important hot areas targeting the drug antitumor research.

Additionally, invasion and migration were suppressed by lidocaine and ropivacaine treatment at a certain range of concentrations which meant the reduction of tumor malignancy.

Furthermore, lidocaine and ropivacaine treatment induced apoptosis. Apoptotic pathways include two major signaling routes: the extrinsic death receptor pathway and the intrinsic mitochondrial pathway (29,30). Apoptosis was mainly controlled by caspases, a family of intracellular cysteine proteases, which were grouped into initiators (caspase-2, -8, -9, and -10) and effectors (caspase-3, -6, and -7) (31,32).

Caspases could activate through being cleaved. Firstly, lidocaine and ropivacaine could activate the extrinsic death receptor pathway. Protein ligand Fas bound to its receptors FasL activating the initiator caspase-8 (31). Moreover, Bcl-2 family participated in the apoptotic process, functioning as promoters (Bax) or inhibitors (Bcl-2). Activated Bax could form an oligomeric pore, resulting in the permeabilization of the mitochondrial outer membrane along with a concomitant decrease in the Bcl-2 level (30,33). An increase of Bax/Bcl-2 ratio could contribute to increased sensitivity of cells to apoptosis. A decrease in ∆Ψm was an early event indicating apoptosis, simultaneously with the increase of Bax/Bcl-2 ratio (30). Lidocaine and ropivacaine downregulated ∆Ψm resulting in mitochondrial dysfunction. The dysfunction of mitochondrion released apoptogenic proteins cytochrome c from mitochondria to the cytosol, resulting in the activation of downstream caspases which was ultimately required to induce apoptosis. Endo G and AIF were also released from mitochondria, and then translocated to the nuclei to induce apoptosis via caspase-independent mitochondrial apoptotic pathway. All in all, these results suggested that local anesthetics could activate the mitochondrial apoptotic pathway (34).

Cleaved caspase-3, the active form of caspase-3, was the capital cleavage enzyme in apoptosis (13). Apoptosis was characterized by the nuclear DNA degradation in response to a variety of apoptotic stimuli (35,36). PARP could be cleaved by caspase-3 and -7 during apoptosis which was involved in DNA damage and repair. This cleavage inactivated PARP contributed to cells’ apoptosis (8). Increased PARP cleavage was observed in NSCLC cells after treated with lidocaine or ropivacaine.

In addition to the two classical apoptotic pathways, ROS production was upregulated, which was an explicit indicator of apoptosis (34). The increased ROS production was a clear indication of apoptosis via activating endoplasmic reticulum (ER) stress pathway, which included MAPK pathways (34). The members of MAPK family, including ERKs, JNKs, and p38 MAPK, were activated by phosphorylation on threonine and tyrosine residues by upstream dual-specificity kinases (37). The results showed the phosphorylation of ERK1/2 and JNK increased, suggesting that ERK1/2, JNK, and p38 MAPK may have different effects on local anesthetics induced NSCLC cells apoptosis.

In summary, local anesthetics affect the outcomes of NSCLC in a variety of aspects, including arrest cell cycle, induce apoptosis, and inhibit invasion and migration. In addition, local anesthetics may attenuate the neuroendocrine response due to surgery, thus improve the preservation of immunocompetence. Furthermore, local anesthetics may make tumor cells more sensitive to the effects of chemotherapy. Taking into account that local anesthetics used for postoperative pain relief specially via intrapleural analgesia after minimally invasive thoracoscopic surgery (38,39), our study indicate the additional benefits of local anesthetics in lung cancer surgery which may have substantial clinical implications.

Conclusions

Our study indicates that amide-linked lidocaine and ropivacaine trigger apoptosis in human NSCLC cells via apoptotic pathways and MAPK pathways. The results reveal the beneficial actions of amide-linked local anesthetics.
and call for further studies to their use during lung cancer surgery.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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