Lidocaine inhibited migration of NSCLCA549 cells via the CXCR4 regulation

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Abstract.
BACKGROUND: Lidocaine is a local anesthetic that wildly used in surgical treatment and postoperative medical care for lung cancers. We hypothesized that lidocaine at clinical plasma concentration can inhibit CXCL12/CXCR4 axis-regulated cytoskeletal remodeling thereby reduce the migration of Non-small-cell lung cancers (NSCLC) cells.

METHODS: We determined the effect of lidocaine at clinical plasma concentration on CXCL12-induced cell viability, apoptosis, cell death, monolayer cell wound healing rate, individual cell migration indicators, expression of CXCR4, CD44, and ICAM-1, intracellular Ca\(^{2+}\) level, and filamentous actin level alteration of NSCLC cells A549 and CXCR4-knocked down A549 cells using CCK-8, Bcl-2 ELISA, Cell death ELISA, wound healing assay, chemotaxis assay, western blotting, QPCR, Fura-2-based intracellular Ca\(^{2+}\) assay, and Fluorescein Phalloidin staining respectively.

RESULTS: Lidocaine did not affect cell viability, apoptosis, and cell death but inhibited CXCL12-induced migration, intracellular Ca\(^{2+}\) releasing, and filamentous actin increase. Lidocaine decreased expression of CXCR4, increased CD44, but had no effect on ICAM-1. CXCL12 induced the increase of CD44 and ICAM-1 but did not affect CD44 in the presence of lidocaine. The knockdown of CXCR4 eliminated all the effects of lidocaine. The overexpression of CXCR4 promoted migration but the migration was inhibited by lidocaine.

CONCLUSION: Lidocaine at clinical plasma concentrations inhibited CXCL12-induced CXCR4 activation, thereby reduced the intracellular Ca\(^{2+}\)-dependent cytoskeleton remodeling, resulting in slower migration of A549 cells.

Keywords: Lidocaine, A549, CXCR, CXCL12, CD44, ICAM-1, cytoskeleton remodeling

List of abbreviations

NSCLC: Non-small cell lung cancers
CXCL12: C-X-C Motif Chemokine Ligand 12
CXCR4: C-X-C chemokine receptor type 4

1. Introduction

Lung cancer is one of the most fatal cancer types with the most global death number among males and the second most global death number among females [1].

As the medical care for lung cancer develops, the mortality of lung cancer decreased in the USA and UK has decreased in recent years. However, in many industrialized nations, the emerging social smoking culture has resulted in higher lung cancer rates in these areas [2]. Among all lung cancer cases, more than four-fifths of clinical lung cancer diagnosed are non-small cell lung cancer (NSCLC) [3], a lung cancer type that has diverse pathological features and with an undesirable prognosis.

Clinical medical advancements have improved cancer therapy extensively in recent years [4]. As clinical diagnosis is critical for disease treatment [5], in-depth analyses of lung cancer subtypes have been developed for potential targeting therapy and customized treatment according to their genetic and cellular heterogeneity.
2. Methods and materials

2.1. Cell line and cells culture

A549 cells were purchased from Biofeng (China). Cells were cultured using Ham’s F12K + 2 mM Glutamine + 10% Foetal Bovine Serum (FBS) in a cell culture incubator at 37°C 5% CO2. Cells were cultured in an FBS-free medium for 24 hours to synchronize before any assay.

2.2. Knockdown and overexpression

CXCR4 knockdown was achieved by transfecting the CXCR4 shRNA (TRCN0000256866) plasmid into A549 cells to silence the expression of CXCR4. The plasmid was purchased from Sigma-Aldrich (USA). The Sequences of shRNA oligonucleotides are as follows: 5’-TCCTGTCCTGCTATTGCATTA-3’. CXCR4 overexpression was achieved by transfecting the CXCR4 expressing (hCXCR4-mTFP1) plasmid into A549 cells to overexpress CXCR4. The plasmid was purchased from Addgene (USA). The transfection method was described previously [28]. The knockdown and overexpression of CXCR4 were validated by western blotting experiments.

2.3. Testing reagents

Lidocaine HCl pre-made parenteral solution was purchased from Hospira Inc. (USA). Human CXCL12 and pertussis toxin were purchased from Sigma-Aldrich (USA). Fura-2 (Fura-2-acetoxymethyl ester) was purchased from Abcam (UK). Cell Counting Kit-8 (CCK-8) was purchased from Dojindo Molecular Technologies, Inc (Kumamoto, Japan).

2.4. CCK-8 assay

The CCK-8 assay is used to determine cell overall viability and cytotoxicity. The CCK-8 assay was conducted as previously described [29]. Briefly, cells were seeded in 96 well plates. After the incubation with testing drugs, 10 µl of the CCK-8 labeling reagent was added to each well. The cells were further incubated for 4 hours. The absorbance at 450 nm was measured using a microplate reader.

2.5. Apoptosis and cell death detection

Apoptosis and cell death detection were determined using Bcl-2 ELISA kit (Abcam, UK) and Cell Death Detection ELISA plus (Roche, USA) respectively. The method of ELISA was described previously [30,31]. The positive control was induced by high-temperature culture (55°C for 30 min) as a previous study [32].

2.6. Western blotting

Membrane protein expression was determined using the western blotting assay. Membrane proteins of A549 were extracted using Mem-PER™ Plus Membrane Protein Extraction Reagent (Thermo Scientific, USA). Membrane protein expression was measured by Western Blotting and quantified using ImageJ software (National Institutes of Health, USA). The expression of CXCR4 was normalized to β-actin expression.
brane Protein Extraction Kit (Thermo Fisher Scientific, USA). The extraction was conducted as previously described [33, 34]. Subsequently, the western blotting assay was conducted as in previous studies [35, 36]. Briefly, SDS-PAGE gel electrophoresis was used to separate proteins and semi-dry protein transfer was conducted as described [37]. The membrane was then incubated with primary and secondary antibodies according to the recommended experimental condition.

Na+/K+ ATPase protein was used as an internal reference gene to normalized the data. Antibodies used including CXCR4 Antibody (PA3-305), Anti-Alpha 1 Na+/K+ ATPase Antibody (#ANP-001), Recombinant Anti-CD44 antibody [EPR1013Y] (ab51037), Anti-ICAM1 antibody [EP1442Y] (ab53013), Goat Anti-Mouse IgG H&L (HRP) (ab6789), and Goat Anti-Rabbit IgG H&L (HRP) (ab6721). ECL reagent (Thermo Scientific, USA) was used to visualize the target protein.

2.7. QPCR

The CXCR4 mRNA level was determined using a QPCR assay. The extraction was conducted as previously described [38, 39]. Briefly, Trizol buffer (Sigma-Aldrich, USA) was used to isolate cellular RNA. An Agilent Bioanalyzer (Agilent Technologies, Inc, USA) was used to assess the concentration and integrity of RNA extracted. QuantiTect Reverse Transcription Kit (QIAGEN, USA) and miRCURY LNA SYBR Green PCR Kit (QIAGEN, USA) were used to conduct RT-PCR. The protocol of PCR reaction was described previously [40]. The GAPDH gene was used as an internal reference gene to normalize the data. CXCR4 Human qPCR Primer Pair (NM_003467) and GAPDH Human qPCR Primer Pair (NM_002046) were purchased from OriGene Technologies (USA).

2.8. Wound healing assay

Cells migration was determined using the wound healing assay. The method was described previously [41, 42]. Briefly, cells were grown in six-well plates at over 90% confluency. A scratch wound was created in the cell monolayer using a 200-µl pipette tip. The cells were cultured with testing reagents and images were recorded every 3 or 6 hours. The wound distance was analyzed using Fiji16 software.

2.9. Chemotaxis assay

Real-time cell migration of individual A549 cells was recorded using the m-Slide chemotaxis system (ibidi, Germany). The method was described previously [43, 44]. Briefly, A549 cells were cultured on the central channel of the chemotaxis slide at 10% confluency for 8 h to allow adherence. Testing drugs were applied when the assay started and the images of cells were recorded for 15 hours with a time-lapse MicroImager. Single-cell tracking was analyzed using the ImageJ software. Spider plots representing the aggregated trajectories of cells. Forward migration indexes and cell velocity were analyzed using the Ibidi software.

2.10. Fura-2-based intracellular Ca$^{2+}$ assay

Fura-2-based fluorescence was used to determine the intracellular Ca$^{2+}$ level. Fura-2 has been wildly used in the indication of intracellular Ca$^{2+}$ [45, 46]. A549 cells were cultured on poly-D-lysine coated TC-treated Cell Culture Dishes (Falcon) with lidocaine or vehicle for 24 hours. Cells of the Gai blocker group were pretreated with 1 mg/ml pertussis toxin for 2 hours before the Fura-2-loading. Cells were loaded with Fura-2-loading buffer (1 mM Fura-2, 140 mM NaCl, 5 mM KCl, 20 mM HEPES, and 1 mM CaCl$_2$, pH 7.4.) for 20 min at 37°C. Then the loaded cells were gently washed with a washing buffer (140 mM NaCl, 5 mM KCl, 20 mM HEPES, and 1 mM CaCl$_2$, pH 7.4.). Images were recorded real-timely on an Axiovert 200 inverted microscope with an excitation at 340 and 380 nm, CXCL12 was perfused 50 s after the recording start and the monitoring continued until 150 s. The F340/380 ratio was calculated and plotted against time to indicate the changes in intracellular Ca$^{2+}$ level.

2.11. Actin polymerization detection

The cytoskeleton remodeling was indicated by the determination of actin polymerization using Fluorescein Phalloidin staining. The method was described previously [47]. Briefly, cells were fixed in 4% paraformaldehyde for 10 min at 4°C, then were permeabilized with 0.01% Triton-X-100 for 5 min. Filamentous actin was stained using Fluorescein Phalloidin (F432, Thermo Fisher Scientific, USA) for 40 min. Images of fluorescence were recorded and the fluorescence intensity was analyzed using the Fiji software.

2.12. Plotting and statistical analysis

Means and standard deviations are displayed in the figures. A T-test or ANOVA was used to analyze the significance of the difference ($p < 0.01$). Dunnett’s post hoc tests were used to test the difference between groups. GraphPad Prism (version 8) was used to plot the data and calculate statistics. Figure 7 was plotted using the BioRender online tools.
3. Results

3.1. Lidocaine did not affect A549 cell viability at the clinical plasma concentrations

The anti-arrhythmia plasma concentration and approximately equipotent nerve block concentrations of lidocaine are around 10 µM [48]. We first evaluated the toxicity of lidocaine at the clinical plasma concentration range (1–100 µM) on NSCLCA549 cells. Results showed that at these concentrations, lidocaine did not affect cell viability, apoptosis, and cell death (Fig. 1A–C). Therefore, we suggested that Lidocaine at clinical plasma concentrations did not inhibit proliferation or induce apoptosis or necrosis of A549. In addition, CXCL12 at 100 nM also failed to affect A549 viability (Fig. 1D). In this study, we performed a knockdown experiment of CXCR4 in A549. Results showed that CXCR4 knockdown did not affect the surface expression of CXCR4 in A549 cells. A549 were exposed to tested drugs for 24 hours. A. Effect of clinical-plasma-dose lidocaine on A549 viability. B. Effect of lidocaine on A549 apoptosis. C. Effect of lidocaine on A549 cell death. PC: positive control induced by high-temperature culture (55°C for 30 min). D. Effect of CXCL12 and lidocaine on the viability of A549. E. Effect of CXCL12 and lidocaine on the viability of CXCR4 knocked down A549. (Significant differences are indicated by “p < 0.01, NS = not significant).
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3.3. Lidocaine inhibited CXCL12 induced migration

CXCL12 is the agonist of the CXCR4 and has been wildly used for studying chemokinesis or chemotaxis of cells [53]. To tested if lidocaine affected CXCL12 induced migration of A549 cells, we performed both the wound healing assay to observed the migration of monolayer A549 cells as a group and in vitro chemotaxis assay to observe the migration of individual A549 cells. Results showed that CXCL12 significantly increased the A549 wound healing rate. In addition, CXCL12 also increased the migration index and velocity of individual cells. As shown in the aggregated trajectories of individual A549 cells, A549 migrated a longer distance at the presence of CXCL12. At the presence of lidocaine, the A549 wound healing rate was the same as control, but the wound healing rate decreased compared with CXCL12-induced cells (Fig. 3). These results revealed that lidocaine did not affect A549 cell migration but blocked the stimulation of CXCL12.

3.4. CXCR4 knockdown block lidocaine effect in migration

To investigate whether lidocaine affects migration through CXCR4, we also performed migration assays in CXCR4 knockdown A549 cells. Results showed that knockdown of CXCR4 eliminated the effects of CXCL12 or lidocaine on A549 migration. The exposure of CXCL12 or lidocaine had no significant effect on the wound healing speed and the migrate index and velocity of individual cells (Fig. 4). This suggested that CXCR4 was essential for the effects of CXCL12 and lidocaine.

3.5. Lidocaine inhibited migration caused by CXCR4 overexpression

To further investigate the effect of lidocaine on CXCR4 mediated migration, we also performed migration assays in CXCR4 overexpressing A549 cells. Results showed that CXCL12 significantly increased CXCR4 overexpressing the A549 wound healing rate. In addition, CXCL12 also increased the migration index and velocity of individual CXCR4 overexpressing A549 cells. As shown in the aggregated trajectories of individual CXCR4 overexpressing A549 cells, these cells migrated a longer distance at the presence of CXCL12. At the presence of lidocaine, CXCR4 overexpressing the A549 wound healing rate was the same as control.
Fig. 3. Effect of lidocaine on CXCL12-induced A549 migration. A. Effect of lidocaine on CXCL12-induced A549 monolayer wound healing. B. Representative images of wound healing assay. C. Effect of lidocaine on CXCL12-induced A549 forward migration index. D. Effect of lidocaine on CXCL12-induced A549 migration velocity. E. Aggregated trajectories of individual A549 cells at different experimental conditions (10 hours). (Significant differences are indicated by *p < 0.01, NS = not significant).

but the wound healing rate decreased compared with CXCL12-induced cells (Fig. 5). These results revealed that lidocaine did not affect CXCR4 overexpressing A549 cell migration but blocked the stimulation of CXCL12 (Fig. 6). These results demonstrated that the effect of lidocaine was mediated by CXCR4/CXCL12 signal.

3.6. Regulations of lidocaine on CD44 and ICAM-1

To further explore the potential mechanisms for lidocaine action on A549 migration, we investigated two critical adhesion molecules on the epithelial cell membrane, CD44 and ICAM-1. CD44 has been widely used as a key migration-related biomarker for lung cancers [54,55]. Another migration-related extracellular molecule, ICAM-1, is also thought to be critical in lung cancer cell migration [56–58]. Both CD44 and ICAM-1 levels were significantly increased by CXCL12. Lidocaine increased CD44 in both wild-type A549 and CXCR4 knockdown A549 cells in the absence or presence of CXCL12, but it did not affect ICAM-1 (Fig. 6). These results suggested that lidocaine can up-regulate
Fig. 4. Effect of lidocaine on CXCL12-induced CXCR4 knocked down A549 cells migration. A. Effect of lidocaine on CXCL12-induced CXCR4-knocked down A549 monolayer wound healing. B. Representative images of wound healing assay. C. Effect of lidocaine on CXCL12-induced CXCR4-knocked down A549 forward migration index. D. Effect of lidocaine on CXCL12-induced CXCR4-knocked down A549 migration velocity. E. Aggregated trajectories of individual CXCR4-knocked down A549 cells at different experimental conditions (10 hours). (Significant differences are indicated by \( p < 0.01 \), NS = not significant).

CD44 but not ICAM-1. However, the regulation of lidocaine on CD44 was not essential for CXCL12 induced migration.

3.7. Lidocaine inhibited CXCL12-induced intracellular \( \text{Ca}^{2+} \) releasing and cytoskeleton remodeling

Another potential mechanism we hypothesized underlying lidocaine’s effect on migration is the regulation of the cytoskeleton remodeling. To test this hypothesis, we monitored the effect of lidocaine on the intracellular \( \text{Ca}^{2+} \) level alteration following the addition of CXCL12. The pertussis toxin was used as a control to block the Gai subunit of CXCR4 [47]. The CXCR4 knockdown A549 cells were used as another control. Results showed that lidocaine decreased the peak of the \( \text{Ca}^{2+} \) curve by about 50%. CXCR4 knockdown A549 cells reacted slightly (less than 5% of the peak of \( \text{Ca}^{2+} \) curve in the control group) to CXCL12 and pertussis toxin blocked almost all the effects of CXCL12 (Fig. 7). To further study the CXCL12-induced cytoskeleton remodeling of A549 at the presence of lidocaine, we stained filamentous actin with phalloidin coupled to fluorescein isothiocyanate to observe the rearrangement of cortical actin fibers during CXCL12 stimulation. Results showed that lidocaine reduced cytoskeleton remodeling by about 50% and both the pertussis toxin group and CXCR4 knockdown group failed to show any...
Fig. 5. Effect of lidocaine on CXCL12-induced CXCR4 overexpressing A549 cells migration. A. Effect of lidocaine on CXCL12-induced CXCR4 overexpressing A549 monolayer wound healing. B. Representative images of wound healing assay. C. Effect of lidocaine on CXCL12-induced CXCR4 overexpressing A549 forward migration index D. Effect of lidocaine on CXCL12-induced CXCR4 overexpressing A549 migration velocity. E. Aggregated trajectories of individual CXCR4 overexpressing A549 cells at different experimental conditions (10 hours). (Significant differences are indicated by *p < 0.01, NS = not significant).

reaction following CXCL12 stimulation. These results suggested that lidocaine inhibited CXCL12-induced intracellular Ca\(^{2+}\) releasing thereby reduced cytoskeleton remodeling.

4. Discussion

The therapeutic effect of surgery in metastatic NSCLC has been controversial [59,60]. During the surgery, many clinical factors might contribute to the metastasis and recurrence of lung cancer surgery The application of lidocaine during the surgery or preoperative treatment results in a plasma lidocaine microenvironment for lung cancer cell migration and survival. Hence, the potential impact of lidocaine on metastasis and recurrence of lung cancer surgery should be further studied. However, so far, the effect of lidocaine at plasma concentration on cancers was less studied. The doses of lidocaine used in most previous studies are much higher than the clinical plasma concentration. Therefore, although many previous studies demonstrated the effect of lidocaine on lung cancer cells, they fail to convince clinical surgeons that lidocaine exerts a considerable impact on the surgery outcome. Lidocaine is known as a sodium channel blocker. Although the major target of lidocaine, the voltage-gated sodium channels, has been found to play a role in cancer developments [61], lidocaine might also affect cancer independent of sodium channel blockade.
Fig. 6. Effect of lidocaine on membrane expression of CD44 and ICAM-1 in A549. A. Representative images of western blotting. B. membrane expression of CD44 protein in A549. C. membrane expression of ICAM-1 protein in A549. (Significant differences are indicated by *p < 0.01, NS = not significant)

Although a previous study suggested that lidocaine at the “mM” concentration range inhibited proliferation [15] and induced apoptosis of A549 [62], our result showed that, at plasma concentration, lidocaine had almost no effect on cell proliferation, apoptosis, and cell death. The previous study also suggested that the migration of A549 was inhibited by lidocaine at 8 mM, but as shown by our data, the effect of lidocaine on migration was not significant when the doses of lidocaine decreased to 100 µM. Lidocaine at clinical plasma concentrations only reduced CXCL12-induced migration in A549. The western blotting assay showed that lidocaine decreased the expression of CXCR4 especially the membrane expression of CXCR4. We suggested that the reduction of CXCR4 on the membrane surface impaired the sensitivity of cells to CXCL12 stimulations. A previous study has suggested that lidocaine can regulate CXCR4 sensitivity to CXCL12 in breast cancer cells MDA-MB-231 [22], which was similar to A549 according to our results.

Different membrane surface proteins expressed on lung cancer cells as adhesion molecules can be critical in the migration of cells [63] CD44 has been reported to play roles in the metastasis of NSCLC cells [54]. In this study, the expression of CD44 was promoted by CXCL12 stimulation. Our results also showed that the
Fig. 7. Effect of lidocaine on CXCL12-induced intracellular Ca\(^{2+}\) releasing and cytoskeleton remodeling. A. Real-time monitoring of the intracellular Ca\(^{2+}\) concentration in response to CXCL12 addition in A549 or CXCR4-knocked down A549 cells pre-incubated with testing reagents for 2 hours. B. Fluorescence intensity of filamentous actin before and after the exposure of CXCL12. The data were displayed in the Relative Fluorescence Unit. C. Representative images of fluorescence of filamentous actin. (Significant differences are indicated by \(^*\) p < 0.01, NS = not significant).

Surface expression of CD44 on A549 was up-regulated by lidocaine. However, the lidocaine did not further increase the CD44 expression at the presence of CXCL12 and the increase of CD44 in the lidocaine alone group did not affect cell migration. Thus, we suggested that CD44 was directly up-regulated by lidocaine bypassing the CXCL12/CXCR4 axis but the increase of CD44 was not essential for the migration of A549. In addition, we also determined another critical adhesion molecule for migration, the ICAM-1. The expression of ICAM-1 has been associated with lung cancer progression and prognosis [64]. A previous study reported that lidocaine affects the migration of a lung cancer cell line H838 by reducing ICAM-1 [65]. However, in this study, the lidocaine at plasma concentration did not affect ICAM-1 expression. The ICAM-1 expression was up-regulated by CXCR4 activation by CXCL12, hence, we suggested ICAM-1 might be a potential downstream target of lidocaine.
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Fig. 8. Effect of lidocaine on A549 migration.

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Lidocaine/CXCR regulation (Fig. 8). Further validation is required in the future.

Another activity that might impact the migration of A549 was the remodeling of the cytoskeleton. The activation of CXCR4 by CXCL12 can trigger intracellular Ca\(^{2+}\) releasing [66]. Subsequently, the released intracellular Ca\(^{2+}\) facilitated many Ca\(^{2+}\)-dependent activities that are involved in cytoskeleton remodeling [67,68]. Results showed that lidocaine significantly reduced the intracellular Ca\(^{2+}\) releasing triggered by CXCL12 stimulated CXCR4 activation. We further determined the level of filamentous actin to observe the rearrangement of cortical actin fibers during CXCL12 stimulation. Actin is an important part of the cytoskeleton in most eukaryotic cells [69]. There are two types of actin: globular G-actin and filamentous F-actin [69]. The G-actin can polymerize into the actin filaments, the F-actin polymer filaments that form the cytoskeleton [69]. Thus, the level of F-actin can indicate the rate of cytoskeleton rearrangement. Our results showed that lidocaine reduced cytoskeleton remodeling by about 50%. Therefore, we suggested that lidocaine inhibited cytoskeleton remodeling via suppressing CXCL12-induced intracellular Ca\(^{2+}\) releasing (Fig. 8). However, lidocaine, as a multiple channel blocker, might have other effects on intracellular Ca\(^{2+}\) levels. Many cancer-related ion channels, such as TRP [70] and TPCs [71], might contribute to lidocaine effects on cancer cells and need further investigation.

5. Conclusion

This study demonstrated that lidocaine at clinical plasma concentrations showed a significant inhibition effect on CXCL12-induced CXCR4 activation, thereby reduced the intracellular Ca\(^{2+}\) dependent cytoskeleton remodeling, resulting in slower migration of A549 cells. This investigation can provide a better understanding of the pharmacological effects of lidocaine at clinical plasma concentrations on the migration of NSCLC cells and can optimize the clinical application of lidocaine in lung cancer surgery.

Ethics approval and consent to participate

This work was approved and consented by the Ethical Committee of Changzhi Medical College Affiliated Heping hospital.

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Availability of data and materials

The raw data of this study are provided from the corresponding author with a reasonable request.

Authors’ contributions

INTERPRETATION OR ANALYSIS OF DATA: Baichun Xing
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Conflict of interest

The authors claimed that there is no conflict of interest.

Consent for publication

All the author consent for this publication.
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regulating CXCR4 to suppress RhoA/ROCK signaling path-


