5-Aza-2′-deoxycytidine protects against emphysema in mice via suppressing p16<sup>Ink4a</sup> expression in lung tissue

Zhi-Hui He<sup>1</sup>
Yan Chen<sup>2</sup>
Ping Chen<sup>2</sup>
Sheng-Dong He<sup>2</sup>
Hui-Hui Zeng<sup>2</sup>
Ji-Ru Ye<sup>2</sup>
Da Liu<sup>2</sup>
Jun Cao<sup>3</sup>

<sup>1</sup>Intensive Care Unit, <sup>2</sup>Department of Respiratory Medicine, Second Xiangya Hospital, Central South University, Changsha, <sup>3</sup>Department of Respiratory Medicine, Hunan Provincial People’s Hospital, Changsha, China

**Background:** There is a growing realization that COPD, or at least emphysema, involves several processes presenting in aging and cellular senescence. Endothelial progenitor cells (EPCs) contribute to neovascularization and play an important role in the development of COPD. The gene for p16<sup>Ink4a</sup> is a major dominant senescence one. The aim of the present study was to observe changes in lung function, histomorphology of lung tissue, and expression of p16<sup>Ink4a</sup> in lung tissue and bone marrow-derived EPCs in emphysematous mice induced by cigarette-smoke extract (CSE), and further to search for a potential candidate agent protecting against emphysema induced by CSE.

**Materials and methods:** An animal emphysema model was induced by intraperitoneal injection of CSE. 5-Aza-2′-deoxycytidine (5-Aza-CdR) was administered to the emphysematous mice. Lung function and histomorphology of lung tissue were measured. The p16<sup>Ink4a</sup> protein and mRNA in EPCs and lung tissues were detected using Western blotting and quantitative reverse-transcription polymerase chain reaction, respectively.

**Results:** CSE induced emphysema with increased p16<sup>Ink4a</sup> expression in lung tissue and bone marrow-derived EPCs. 5-Aza-CdR partly protected against emphysema, especially in the lung-morphology profile, and partly protest against the overexpression of p16<sup>Ink4a</sup> in EPCs and lung tissue induced by CSE.

**Conclusion:** 5-Aza-CdR partly protected against emphysema in mice via suppressing p16<sup>Ink4a</sup> expression in EPCs and lung tissue.

**Keywords:** 5-Aza-2′-deoxycytidine, cigarette smoke, emphysema, endothelial progenitor cells, p16<sup>Ink4a</sup>

**Introduction**

Increasing research has indicated that COPD, or at least emphysema, represents premature aging or premature senescence of lung parenchymal cells, which are induced in part by oxidative damage from cigarette-smoke (CS) components, resulting in accelerated lung aging/accelerated lung senescence. Moreover, the key patho-genetic processes involved in COPD are considered to involve those senescent cells, notably progenitor cells, decreasing regenerative properties. Bone marrow-derived endothelial progenitor cells (EPCs), one of the major components of parenchymal cells, provide an alternative source of endothelial cells (ECs) and play a fundamental role in the maintenance of endothelial integrity and function, postnatal vasculogenesis, vascular repair, and tissue regeneration through pivotal bioactivity, differentiating into ECs and secretion of vasoactive substances that promote angiogenesis and maintain vascular homeostasis. The normal function of EPCs is required for tissue repair and

Correspondence: Yan Chen
Department of Respiratory Medicine,
Second Xiangya Hospital, Central South University, Nanyuangong Alley, Yuanjialing Shangguan, Furong Qu, Changsha, Hunan 410011, China
Email chenyan99727@163.com
airway remodeling in lungs.8–10 Our previous study showed decreased and dysfunctional circulating EPCs in patients with COPD.11

p16\(^{ink4a}\) was initially discovered as a tumor-suppressor factor composed of 148 amino-acid residues with molecular weight 16 kD.12 With increased tumor investigations, it was found that relationships between p16\(^{ink4a}\) and tumor cells were not all the same. p16\(^{ink4a}\) is a cyclin-dependent kinase inhibitor that controls cell-cycle progression,13 and could be regarded as a major domineous senescence gene.14 ECs have higher expression rates of p16\(^{ink4a}\), inducing cell senescence in COPD patients.15 p16\(^{ink4a}\) expression in EPCs and emphysematosus lung tissue has been little studied.

The advent of genome-wide epigenetic studies allowed for more comprehensive study of the epigenome in many diseases. Hypermethylation of genes associated with CS has been reported.16,17 5-Aza-2′-deoxycytidine (5-Aza-CdR), an S-phase-specific inhibitor of DNA methyltransferase, is the most widely used inhibitor of DNA methylation and triggers demethylation, leading to a consecutive reactivation of epigenetically silenced genes in vitro and in vivo.18 In this study, in an attempt to elucidate pathophysiological mechanisms of emphysema with regard to gene hypermethylation, we detected p16\(^{ink4a}\) expression in bone marrow-derived EPCs and lung tissue of mice with emphysema induced by CS extract (CSE) and compared the results with those in mice with emphysema treated with 5-Aza-CdR. Lung function, histomorphology, and apoptosis in lung tissue were the indicators for evaluating the severity of emphysema in mice.

Materials and methods

Animals

A total of 24 C57BL/6J male mice aged 4–6 weeks were randomly enrolled in this study. All animals were purchased from the Shanghai Laboratory Animal Center of the Chinese Academy of Sciences and fed in a cleaning unit at 23°C–25°C and 50%–60% humidity, with a 12-hours light–dark cycle. The study was approved by the institutional review board of Central South University and conformed to the guiding principles for research involving animals and human beings.

Preparation of CSE

CSE was prepared according to a previous publication,19 with some modifications. Briefly, one unfiltered Furong cigarette (tar 13 mg, nicotine 1 mg, carbon monoxide 14 mg/cigarette; China Tobacco Hunan Industrial, Changsha, China) was burned and the smoke passed through 4 mL PBS via a vacuum pump at a constant pressure of −0.1 kPa. This product was further filtered through a filter with 0.22 μM pores (Thermo Fisher Scientific, Waltham, MA, USA) to remove particles and bacteria and used for intraperitoneal injection. The solution was prepared freshly for each injection.

Preparation of 5-Aza-2′-deoxycytidine

5-Aza-CdR powder (5 g; Sigma-Aldrich, St Louis, MO, USA) was dissolved in 2 mL PBS and further diluted to 25 mg/mL, subpackaged, and stored under −80°C until experiments.

Animal modeling

The mouse emphysema model was established as previously described.20 C57BL/6J mice were divided into three groups: controls, CSE, and CSE + 5-Aza-CdR (n=8 per group). The total experimental period was 4 weeks, with intraperitoneal injection of PBS, CSE, or 5-Aza-CdR (Table 1). According to animal weight, intraperitoneal injection doses of PBS, CSE, 5-Aza-CdR were 0.3 mL/20 g, 0.3 mL/20 g, and 2.5 mg/kg (0.3 mL/20 g constant volume), respectively. At day 28, mice were killed for measurement of lung function, detection of histomorphology of lung tissue, and separation of bone marrow-derived EPCs.

Isolation, culture, and identification of EPCs

Ficoll density-gradient centrifugation (Histopaque-1083; Sigma-Aldrich) was used to isolate mononuclear cells (MNCs) from bone marrow of C57BL/6J mice according to a previously published method.21,22 Isolated MNCs were cultured with EGM-2 growth medium in the presence of 5% FBS (SingleQuots; Lonza, Basel, Switzerland) under an atmosphere of 95% humidity, 5% CO\(_2\), and 37°C for EPC culture. Cells were inoculated into culture flasks at a density of 3–5\(\times\)10\(^4\)/mL. Then, culture fluid was replaced totally by fresh culture medium on day 4 of the culture to remove unattached cells. Half replacement with the fresh medium was performed every 3 days. Cell harvesting was performed on day 7 of the culture. To identify EPCs, firstly photos were taken during the culture using phase-contrast

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**Table 1** Experiment schedule

<table>
<thead>
<tr>
<th>Group</th>
<th>Day 0</th>
<th>Day 11</th>
<th>Day 15</th>
<th>Day 17</th>
<th>Day 19</th>
<th>Day 22</th>
<th>Day 28</th>
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<td>PBS</td>
<td>PBS</td>
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<td>PBS</td>
<td>Disposed</td>
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<tr>
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<td>CSE</td>
<td>CSE</td>
<td>CSE</td>
<td>CSE</td>
<td>CSE</td>
<td>Disposed</td>
</tr>
<tr>
<td>CSE + 5-Aza-CdR</td>
<td>CSE</td>
<td>CSE</td>
<td>Aza</td>
<td>Aza</td>
<td>Aza</td>
<td>Aza</td>
<td>Disposed</td>
</tr>
</tbody>
</table>

**Abbreviations:** CSE, cigarette-smoke extract; 5-Aza-CdR, 5-Aza-2′-deoxycytidine.
microscopic point-count technique at 200 magnification. A total of 36 lines per mouse lung were drawn and measured. The DI was calculated by dividing the defined destructive alveoli by the total number of alveoli counted. Destructive alveolus was defined if at least one of the following alveoli was observed: alveolar wall defects, intraluminal parenchymal rags in alveolar ducts, obviously abnormal morphology, and typically emphysematous changes. Analysis was performed using a microscopic point-count technique at 200× magnification.

Lung-function measurement
Lung-function measurement was performed using small-animal spirometry (PLY3211 system; Buxco Electronics, Wilmington, NC, USA) as previously described with a minor modification.20 Briefly, the mouse was anesthetized by intraperitoneal injection of 10% chloral hydrate (3 mL/kg body weight) and tracheostomized. The trachea was cannulated and the cannula connected to the computer-controlled small-animal spirometer. Airway resistance (Raw), lung dynamic compliance (Cdyn, body weight) and tracheostomized. The trachea was cannulated and the cannula connected to the computer-controlled small-animal spirometer. Airway resistance (Raw), lung dynamic compliance (Cdyn), peak expiratory flow (PEF), and inspiratory time/ expiratory time (Tl/Tc) were measured according to the manufacturer’s instructions.

Histomorphological detection
After lung-function measurement, animals were killed by overdose of anesthetic. The lower-left lobes of lungs were inflated with 4% paraformaldehyde at a pressure of 25 cmH2O, then fixed with 4% paraformaldehyde for 24 hours.19 Fixed lungs were embedded in paraffin (Sigma-Aldrich) and sliced into 4 μm sections. The slices were stained with H&E (Sigma-Aldrich). Pulmonary emphysema was quantified based on the measurement of the mean linear intercept (MLI) and destructive index (DI) in micrometers. The MLI was measured by dividing the length of a line drawn across the lung section by a total number of intercepts counted within this line at 100× magnification. A total of 36 lines per mouse lung were drawn and measured. The DI was calculated by dividing the defined destructive alveoli by the total number of alveoli counted. Destructive alveolus was defined if at least one of the following alveoli was observed: alveolar wall defects, intraluminal parenchymal rags in alveolar ducts, obviously abnormal morphology, and typically emphysematous changes. Analysis was performed using a microscopic point-count technique at 200× magnification.19 Ten randomly selected fields per slice were photographed in a blinded manner. Airways and vascular structures were eliminated from the analysis.

Apoptosis assay
Terminal deoxynucleotidyl transferase dUTP nick-end labeling (TUNEL) was performed to label the DNA-damaged cells in the lungs of experimental mice using an in situ cell-death-detection kit (Hoffman-La Roche, Basel, Switzerland) following the manufacturer’s instructions. The apoptotic index (AI) was calculated as the percentage of TUNEL-positive nuclei in a total of more than 3,000 nuclei randomly counted for each lung at 400× magnification.

Western blotting
Briefly, EPCs were washed three times with ice-cold PBS, then lysed in radioimmunoprecipitation-assay lystate (Applygen Technologies Beijing, China) for 30 minutes on ice. Lung tissues were homogenized manually in a glass homogenizer and lysed in radioimmunoprecipitation-assay lystate for 30 minutes on ice. Solutions of EPCs or lung tissue were centrifuged at 4°C, 12,000 g for 5 minutes. A BCA protein-quantification kit (Wellbio, Changsha, China) was used for protein measurement. Protein (30–60 μg) was mixed 1:1 with 2× sodium dodecyl sulfate (SDS) loading buffer (20% glycerol, 4% SDS, 3.12% dithiothreitol DDT, 0.2% bromophenol blue, and 0.1 mol/L Tris HCl, pH 6.8; all Sigma-Aldrich) and incubated at 100°C for 4 minutes. Equal amounts of protein for each sample were separated by 10%–12% SDS–polyacrylamide gel run at 120 V for 90 minutes and blotted onto a polyvinylidene difluoride microporous membrane (EMD Millipore, Billerica, MA, USA). Membranes were incubated with a 1:200 dilution of primary antibody (mouse monoclonal antibody; Santa Cruz Biotechnology, Dallas, TX, USA) overnight, then washed for three times with Tris-buffered-saline with Tween (TBS-T) and revealed using secondary antimouse antibody with horseradish peroxidase conjugate (1:3,000, 1 hour), followed by washing with TBS-T again. Immunoreactive bands were developed using enhanced chemiluminescence substrate (Thermo Fisher Scientific).

RNA extraction and quantitative RT-PCR
p16INK4a mRNA expression in bone marrow-derived EPCs and lung tissue was detected by quantitative reverse-transcription polymerase chain reaction (RT-PCR). Total RNA was extracted from cells or tissues using Trizol reagent (Thermo Fisher Scientific). First-strand cDNA was synthesized
using a RevertAid first-strand cDNA-synthesis kit (Thermo Fisher Scientific) according to the manufacturer’s instructions, and used as the template for quantitative RT-PCR analysis. DNase-treated samples were subjected to RT-PCR analysis using SYBR Green quantitative PCR master mix (Thermo Fisher Scientific) on a CFX96 real-time system (Bio-Rad Laboratories, Hercules, CA, USA), with β-actin used as an internal control. The PCR-amplification conditions were 10 minutes at 95°C followed by 40 cycles of denaturation at 95°C for 15 seconds and annealing and extension for 1 minute at 60°C. Data were analyzed using comparative C	extunderscore iT. The relative expression level of p16

\[ \text{p16}_{\text{ink4a}} \]

was calculated by determining the ratio of p16

\[ \text{p16}_{\text{ink4a}} \]

to that of the internal control. Melting-curve analysis (65°C–95°C) was used to determine melting temperatures of specific amplification products and primer dimmers. Each experiment was repeated twice in triplicate. Primer sequences were: β-actin, 5′-CATCCTGCCTGAGCCTGGG-3′ (forward), 5′-TAATGTCACGCACGATTTC-3′ (reverse); p16

\[ \text{p16}_{\text{ink4a}} \]

, 5′-CCGCTCAGCGCCGCTTCTTT-3′ (forward), 5′-CCGCCCCCTTGCATTGT-3′ (reverse).

### Statistical analysis

Analyses were performed using SPSS for Windows 16.0 (SPSS, Chicago, IL, USA). All data are expressed as means ± SD. Analyses of differences among groups were performed using one-way analysis of variance, followed by post hoc analysis as appropriate. Values of \( P < 0.05 \) were considered statistically significant.

### Results

#### Culture and identification of EPCs

On day 1 of the culture, MNCs isolated from the murine bone marrow formed circularly, sizes of cells were almost uniform, and cells were suspended in culture media (Figure 1A). On day 4 of the culture, the cells were attached to one another and getting larger, and shapes became oval, spindle, or polygonal. The cells at this stage tended to gather to form ball-like structures (Figure 1B). On culture day 7, cells shaped into fusiform or polygon patterns and contacted one another to attempt to form capillary structures (Figure 1C).

Cell shapes at this stage displayed well in the culture medium. In addition, laser-scanning confocal microscopy illustrated that cells on culture day 7 displayed red cytoplasm when stained with Dil-acLDL (Figure 2A), green cytomembrane when combined with FITC-UEA1 (Figure 2B), and orange confocal when double-positively stained with Dil-acLDL and FITC-UEA1 (Figure 2C). The positive rate of amphophilic cells was 95.25%±3.61% on culture day 7.

#### Lung-function test

As shown in Figure 3, the maximal expiratory flow-volume curve of the CSE group (Figure 3B) and CSE + 5-Aza-CdR group (Figure 3C) showed abrupt ascents, and descending limbs showed a prolonged expiratory phase compared with that in controls (Figure 3A). \( C_{\text{dyn}} \) (mL/cmH

\[ \text{O} \]) was significantly lower in the CSE group (0.57±0.15, \( P < 0.01 \)) and CSE + 5-Aza-CdR group (0.67±0.19, \( P < 0.01 \)) than controls (1.03±0.29). \( R_{\text{aw}} \) (cmH

\[ \text{O} / \text{mL/min} \]) was significantly higher in the CSE group (2.49±0.52, \( P < 0.01 \)) and CSE + 5-Aza-CdR group (1.91±0.47, \( P < 0.01 \)) than controls (0.58±0.14). PEF (mL/second) was significantly lower in the CSE group (2.91±0.5, \( P < 0.01 \)) and CSE + 5-Aza-CdR group (3.24±0.62, \( P < 0.05 \)) than controls (4.4±0.74). The \( T/T_e \) was significantly lower in the CSE group (0.63±0.17) than controls (0.89±0.17, \( P < 0.05 \)). There was no significant difference in \( T/T_e \) between the CSE + 5-Aza-CdR group (0.7±0.15) and controls (\( P > 0.05 \)). There was no significant difference between the CSE group and CSE + 5-Aza-CdR group in terms of the parameters described (\( P > 0.05 \), Figure 4).

![Figure 1](https://www.dovepress.com/)

**Figure 1** Morphological changes in endothelial progenitor cells (EPCs) sourced from bone marrow of C57BL/6j mice during culture. **Notes:** (A) Representative microscopy of EPCs cultured with endothelial growth medium 2 in the presence of 5% fetal bovine serum on day 1. EPCs formed spherical, cell sizes were almost the same, and cells were suspended in the culture medium. (B) On day 4 of the culture, the cells became oval, spindle, or polygonal. (C) On day 7 of the culture, the cells became fusiform or polygonal. EPCs attached to one another to attempt to form capillary structures (arrows). Magnification ×100.
Histomorphological changes in lung tissue

As shown in Figure 5, lung tissue of the CSE group exhibited enlarged alveolar space, thinner alveolar septum, and destroyed alveolar wall. Lung tissue of the CSE + 5-Aza-CdR group also exhibited enlarged alveolar space, but less than the CSE group. The changes described were manifested in the MLI and DI (Figure 6). The MLI of the CSE group (67.63±9.87 μm) was significantly increased when compared with controls (29.2±4.64 μm, P<0.01). Interestingly, the MLI of the CSE + 5-Aza-CdR group (52.7±6.34 μm) was significantly smaller than the CSE group (P<0.01), though larger than controls (P<0.01). Similarly, the DI of the CSE group (42.41%±5.86%) was significantly increased when compared with controls (6.38%±1.57%, P<0.01). The DI of the CSE + 5-Aza-CdR group (33.26%±5.03%) was significantly less than the CSE group (P<0.05), though more than controls (P<0.01).

Apoptosis in lung tissue

As shown in Figure 7, numbers of apoptotic cells in alveolar septa in the CSE and CSE + 5-Aza-CdR groups were significantly increased compared with controls. Quantitatively, the AI of the CSE group (19.5%±3.16%) was significantly increased when compared with controls (2.75%±0.46%, P<0.01). Interestingly, the AI of the CSE + 5-Aza-CdR group (12.75%±1.67%) was significantly lower than the CSE group (P<0.05), though higher than controls (P<0.01, Figure 6).

Expression of p16<sup>ink4a</sup> protein in lung tissue and EPCs

As shown in Figure 8A and C, p16<sup>ink4a</sup>/β-actin in lung tissue was significantly increased in the CSE group (0.59±0.05, P<0.01) and CSE + 5-Aza-CdR group (0.46±0.03, P<0.01) compared with controls (0.32±0.02). Interestingly, p16<sup>ink4a</sup>/β-actin in lung tissue was significantly lower in the CSE + 5-Aza-CdR group than the CSE group (P<0.01).

As shown in Figure 8B and C, p16<sup>ink4a</sup>/β-actin in EPCs was significantly increased in the CSE group (0.51±0.05, P<0.01) and CSE + 5-Aza-CdR group (0.42±0.02, P<0.05) compared with controls (0.36±0.02). Interestingly, p16<sup>ink4a</sup>/β-actin in EPCs was significantly lower in the CSE + 5-Aza-CdR group than the CSE group (P<0.05).
Expression of p16<sup>ink4a</sup> mRNA in lung tissue and bone marrow-derived EPCs

As shown in Figure 9, p16<sup>ink4a</sup> mRNA in lung tissue was significantly increased in the CSE group (5.24±0.67, P < 0.01) and CSE + 5-Aza-CdR group (3.82±0.44, P < 0.01) compared with controls (1±0.12). Interestingly, p16<sup>ink4a</sup> mRNA in lung tissue was significantly lower in the CSE group than the CSE + 5-Aza-CdR group (P < 0.05). p16<sup>ink4a</sup> mRNA in EPCs was significantly increased in the CSE group (4.4±0.6, P < 0.01) and CSE + 5-Aza-CdR group (1.99±0.25, P < 0.05) compared with controls (1.01±0.13). p16<sup>ink4a</sup> mRNA in EPCs was significantly lower of CSE + 5-Aza-CdR than the CSE group (P < 0.01).

Discussion

The present study showed that the expression of p16<sup>ink4a</sup> in lung tissue and bone marrow-derived EPCs was increased in mice with CSE-induced emphysema, which suggested that CSE might induce p16<sup>ink4a</sup> expression, resulting in EPC senescence that contributes to emphysema with overexpression of p16<sup>ink4a</sup> in lung tissue of mice with emphysema. Most importantly, the present study demonstrated that the onset of replicative senescence could not be totally prevented.

Cigarette smoking is by far the most critical risk factor for emphysema and COPD. CS induces significant increases in reactive oxygen species generation. CSE contains most of the compounds inhaled by cigarette smokers, and is usually used as a surrogate for CS. CSE directly induces inflammatory cytokines and superoxide generation, resulting in increased p16<sup>ink4a</sup> expression that induces fibroblast senescence.

There was evidence showing that expression of p16<sup>ink4a</sup> in aged cells may be ten times more than in young cells. Inserting p16<sup>ink4a</sup> cDNA into normal fibroblasts slowed cell growth, aggravated nonenzymatic glycosylation, increased senescence-associated β-galactosidase positivity, and shortened telomeres. On the other hand, significant delay of several senescent features was observed in fibroblasts, and the life span of fibroblasts was significantly extended by inserting antisense p16<sup>ink4a</sup> but the onset of replicative senescence could not be totally prevented. Therefore, p16<sup>ink4a</sup> could be regarded as a major dominant senescence gene. p16<sup>ink4a</sup> levels are increased in pulmonary vascular ECs in patients with COPD. A recent study showed that cord-blood EPCs in premature neonates exhibited overexpression of p16<sup>ink4a</sup>, contributing to accelerated senescence of EPCs.
Aging lungs exhibit both structural and functional alterations. The leading clinical symptom of COPD or emphysema is chronic airflow limitation, which means decreased lung function. In the present study, airflow limitation was detected in CSE-induced emphysematous mice and manifested by decreases in C_d, R_{aw}, PEF, and T/T' of the lungs. Lung tissue in emphysema mice showed enlarged alveolar space, thinner alveolar septum, and destroyed alveolar wall, manifested in increased MLI and DI. Alveolar septal cell apoptosis plays an important role in the development of emphysema. Oxidative stress also triggers apoptosis. In the present study, the AI of lung tissue, which reflects the apoptosis status of lung parenchyma, from emphysematous mice was increased compared with control mice.

In COPD, oxidative stress induced by cigarette smoking further damages the lung, leading to acquired genetic changes, including DNA methylation, due to inefficient...
DNA-repair machinery. DNA methylation is catalyzed by the DNA methyltransferase (DNMT) family and plays an important role in maintaining cell identity by affecting gene expression. 5-Aza-CdR, a DNMT inhibitor, inhibits DNMT and demethylates DNA by incorporation into DNA, leading to changes in gene reactivation. Typically, methylation in the promoter region of a gene is associated with repression. A high frequency of aberrant methylation of the gene for p16\(^{\text{ink4a}}\) has been shown in cases of non-small-cell lung cancer, heavy smokers, and advanced poorly differentiated small adenocarcinoma. Breuer et al showed that loss of p16\(^{\text{ink4a}}\) expression by promoter hypermethylation is inconsistent and occurs late in the carcinogenic process at the level of severe dysplasia. Another intriguing observation is that DNA-methylation levels within gene bodies are also dynamic in relation to gene expression. In the present study, increased p16\(^{\text{ink4a}}\) expression in CSE-induced emphysematous mice was partly suppressed by 5-Aza-CdR. Lung morphological changes and apoptosis in emphysematous mice induced by CSE were also partly reversed by 5-Aza-CdR. The molecular mechanism of active demethylation in mammalian cells is not well understood, but seems to be linked to DNA-repair machinery. We noticed that in this study, there was no statistical difference in lung function between the CSE group and CSE + 5-Aza-CdR group, despite little change in the numbers. The possible reason may lie in lung-function tests being less sensitive than morphometry.

Since methylation is reversible, it is an interesting target for intervention with specific inhibitors of DNA methylation. The antitumor effect or auxiliary-therapy effect of 5-Aza-CdR has been investigated and confirmed by many studies. It could be assumed that 5-Aza-CdR at lower concentrations might be applied in the attenuation of emphysema.

In summary, the present study indicated that p16\(^{\text{ink4a}}\) expression was increased in EPCs and lung tissue in CSE-induced emphysematous mice, and contributed to alterations in lung function, histomorphological changes, and apoptosis in emphysematous lung tissue. 5-Aza-CdR partly reversed the structural emphysematous outcomes resulted from CSE stimulation, which in turn suggested that DNA methylation may be involved in the pathogenesis of emphysema with regard to epigenetic modifications in...
terms of hypermethylation of genes and EPC senescence. DNA methyltransferase inhibitors might help potentially in clinical treatment of emphysema. Future study is expected to elucidate the exact mechanism of regulation of p16
Ink4a on EPC senescence.

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Disclosure

The authors report no conflicts of interest in this work.

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