Research Article

OC-STAMP Overexpression Drives Lung Alveolar Epithelial Cell Type II Senescence in Silicosis

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Cellular senescence has been considered an important driving mechanism for many chronic lung diseases. However, the specific mechanism of cellular senescence in silicosis is still unknown. In the present study, silicotic rats and osteoclast stimulatory transmembrane protein (OC-STAMP) overexpression of MLE-12 cells were used to explore the mechanism of OC-STAMP in cellular senescence in alveolar epithelial cell type II (AEC2). We found an increasing level of OC-STAMP in AEC2 of silicotic rats. Overexpression of Ocstamp in MLE-12 cells promoted epithelial-mesenchymal transition (EMT), endoplasmic reticulum (ER) stress, and cellular senescence. Myosin heavy chain 9 (MYH9) was a potential interacting protein of OC-STAMP. Knockdown of Ocstamp or Myh9 inhibited cellular senescence in MLE-12 cells transfected with pcmv6-Ocstamp. Treatment with 4-phenylbutyrate (4-PBA) to inhibit ER stress also attenuated cellular senescence in vitro or in vivo. In conclusion, OC-STAMP promotes cellular senescence in AEC2 in silicosis.

1. Introduction

Silicosis is a chronic occupational lung disease caused by long-term inhalation of free crystalline silica dust and is characterized by silicotic lesions and progressive massive fibrosis [1]. Injury, loss, and disruption of alveolar epithelial cell type II (AEC2) play a central role in pulmonary fibrosis due to its critical function in alveolar niche homeostasis through the production of pulmonary surfactant and as progenitor cells to self-renew and transdifferentiate into AEC1 [2].

Cellular senescence is now considered an important driving mechanism for chronic lung diseases, particularly chronic obstructive pulmonary disease (COPD) and idiopathic pulmonary fibrosis (IPF) [3]. Cellular senescence occurs due to replicative and stress-related senescence with activation of p53 and p16INK4a, respectively, leading to activation of p21CIP1 and cell cycle arrest [3, 4]. As silicosis is an age-related and chronic occupational lung disease, silicotic patients showed significantly shorter and telomerase gene variants compared with healthy controls in response to exposure to silica [5]. Our previous study showed increasing levels of p21, cleaved caspase-3, and phosphorylated histone H3AX (γH3AX) in rats exposed to inhaled silica over time [6]. We also found that silica, matrix stiffening, or their combination triggered DNA damage and replication stress in AEC2 [7]. Therefore, cellular senescence may be a key contributor to silicosis, but the exact mechanism is still unknown.

In our preliminary study, we found that the receptor activator of nuclear factor kappa-B ligand (RANKL) signaling pathway, a classic signaling pathway for regulating osteoclast...
2. Methods

2.1. Silicosis Model. Wistar rats (3 w old) were employed in this study. A silicosis model was induced by inhalation of 50 ± 10 μg/m³ of silica (s5631; Sigma-Aldrich, St. Louis, MO, USA; ground and then heated at 180°C for 6 h) for 32 w and inhalation of pure air as the control group. To inhibit ER stress, rats received 4-phenylbutyrate (4-PBA, P21005, Sigma-Aldrich) at 100 mg/Kg once daily from 24 w until 32 w [1]. All animal protocols were reviewed and approved by the Committee on the Ethics of North China University of Science and Technology (LX2019033), and they complied with the US National Institutes of Health Guide for the Care and Use of Laboratory Animals [14].

2.2. Cell Culture and Treatment. The MLE-12 cell line was obtained from the Chinese Academy of Sciences cell library (Shanghai, China). Cells were plated in 6 cm² dishes and transfected with pCMV6-Entry (PS100001, OriGene Technologies Inc., MD, USA) and cell lysates (2 g/L) were mixed to a volume of 400 μL with 1% protease inhibitors. Then, 30 μL of sepharose beads (FO115, Santa Cruz Biotechnology, Santa Cruz, CA, USA) and cell lysates (2 g/L) were mixed to a volume of 400 μL and incubated for 2 h at 4°C on a shaker for preclearing. The clear supernatant was incubated overnight with anti-Myh9, anti-IgG antibody, and Protein A sepharose at 4°C. The beads were collected and washed 3 times with PBS before being boiled in a 2× loading buffer at 95°C for 5 min. Western blotting was used to analyse the CoIP results.

2.3. Immunohistochemistry and Immunofluorescence Staining. Immunohistochemical staining was performed using published protocols [15] with antibodies directed against OC-Stamp (1 : 100 dilution, 1501. PB1; FabGennix Inc., Frisco, TX, USA), ABCA3 (1 : 200 dilution, ab24751, Abcam, Cambridge, UK), P21 (1 : 100 dilution, ab109520, Abcam, Cambridge, UK), α-smooth muscle actin (α-SMA, 1 : 200 dilution, ab32575, Abcam, Cambridge, UK), Proliferating Cell Nuclear Antigen (PCNA, 1 : 100 dilution, GTX100539, Genetex, Irvine, CA, USA), and Phospho-PERK (p-PERK, 1 : 100 dilution, DF7576, Affinity, Cincinnati, OH, USA) at a concentration of 1 : 200. Immunofluorescence staining was performed using published protocols with antibodies directed against ABCA3/OC-Stamp, p21/α-SMA, and p-PERK at a concentration of 1 : 200.

2.4. Western Blot. Western blot was performed using published protocols [16] with antibodies directed against OC-Stamp (2051. PB1; FabGennix Inc.), collagen type I (Col I) (ab34710, Abcam), α-SMA (ab32575, Abcam), E-cadherin (ab76055, Abcam), N-cadherin (ARG23870, Arigo), p-Smad2/3 (8828 s, Cell Signaling Technology, MA, USA), Smad2/3 (5678, Cell Signaling Technology), Phospho ataxia telangiectasia and Rad3-related protein (p-ATR, DF7512, Affinity), Phospho ataxia telangiectasia mutated (p-ATM, A8410, Affinity), p-p53-S15 (AP0083, Abclonal), p21 (ab109520, Abcam), p16 (A0262, Abclonal), p-PERK (DF7576, Affinity), p-IRe1α (ab48187, Abcam), Phospho-nuclear factor-kappaB (p-NF-κB, ARG51561, Arigo), transforming growth factor-β1 (TGF-β1, ARG54629, Arigo), TGF-β receptor I (TGFβR1, A16396, Abclonal), and TGF-β receptor II (TGFβR2, ARG59501, Arigo) at a concentration of 1 : 1000.

2.5. Coimmunoprecipitation (CoIP). The interaction of OC-Stamp with Myh9 was evaluated by Co-IP. The cells were lysed in RIPA (R0020, Solarbio Life Sciences, Beijing, China) buffer containing 1% protease inhibitors. Then, 30 μL of sepharose beads (FO115, Santa Cruz Biotechnology, Santa Cruz, CA, USA) and cell lysates (2 g/L) were mixed to a volume of 400 μL and incubated for 2 h at 4°C on a shaker for pre-clearing. The clear supernatant was incubated overnight with anti-Myh9, anti-IgG antibody, and Protein A sepharose at 4°C. The beads were collected and washed 3 times with PBS before being boiled in a 2× loading buffer at 95°C for 5 min. Western blotting was used to analyse the CoIP results.

2.6. Real-Time Quantitative Polymerase Chain Reaction (PCR) Analysis. Reverse transcription (K1622, Thermo Scientific, USA orZR102, ZOMANBIO, China) was performed according to company recommendations. Amplification by real-time PCR was carried out using the 2× SYBR qPCR Mix (ZF102, ZOMANBIO, China) system [17]. The sequence details were as follows: (1) Rat Oc-stamp: Forward: TGGCTGGCCTGTTACTGAG, Reverse: GTGTGAAGTCGGAAGGCTGA; (2) Rat Gapdh: Forward: GGTGAAAGTCCGCGTGGTA; (3) Rat TGFβ receptor I: forward: TGGCTGCCCTTGGAGATGGTG, reverse: TGGCTGCCTTGGAGATGGTG. The results were calculated via the 2^-ΔΔCT method.

2.7. Statistical Analysis. Statistical analyses were performed using SPSS 20.0 software (IBM Corp., Armonk, NY, USA). Two-group comparisons were made using unpaired Student’s t-test, and multiple-group comparisons were made using one-way analysis of variance followed by Tukey’s post hoc test. Statistical significance was achieved when P < 0.05 at a 95% confidence interval.
Figure 1: An increased level of OC-STAMP in silicotic rats. (a) Positivity of ABCA3 and collagen in silicotic rats; data are presented as the mean ± SD; n = 8 per group. (b) Positivity of p21 and PCNA in silicotic rats; data are presented as the mean ± SD; n = 8 per group. (c) Coexpression of ABCA3 and OC-STAMP in silicotic rats; (d) protein and mRNA levels of OC-STAMP in silicotic rats. Data are presented as the mean ± SD; n = 3 per group.
measured by Western blotting. Data are presented as the mean ± SD.

3.2. Overexpression of OC-STAMP Promotes EMT in MLE-12 Cells.

Increased in silicotic rats (Figure 1(d)). Activation of AEC2 showed a cellular senescence phenotype (Figure 1(c)). These results suggested that an increased level and expression of p21 but not of PCNA (Figures 1(b) and 1(c)). Although AEC2 in silicotic rats showed a "hyperplastic" phenotype, most of these cells showed expression of p21 but not of PCNA (Figures 1(b) and 1(c)). These results suggested that an increased level and activation of AEC2 showed a cellular senescence phenotype in rats exposed to silica. We also found coexpression of ABCA3 and OC-STAMP in silicotic rats (Figure 1(d)). The protein and mRNA levels of OC-STAMP were also increased in silicotic rats (Figure 1(d)).

3.3. Overexpression of OC-STAMP Promotes ER Stress and Cellular Senescence in MLE-12 Cells.

3.4. Overexpression of OC-STAMP Induces Insensitivity in Silica-Induced MLE-12 Cells.

First, the cell apoptosis in MLE-12 cells treated with or without 50 μg/mL silica was measured by flow cytometry; silica did not exhibit cytotoxicity in MLE-12 cells at a concentration of 50 μg/mL (Figure S1). As shown in Figure 5, the pCMV6-Entry and pCMV6-Octstamp cells were treated with 50 μg/mL silica. Silica promoted cellular senescence in MLE-12 cells transfected with pCMV6-Octstamp, but it did not increase the activation of cellular senescence signaling in pCMV6-Octstamp cells. Also, silica treatment did not change the EMT-related proteins in MLE-12 cells transfected with pCMV6-Octstamp.
We used 4-PBA to explore the role of ER stress in cellular senescence induced by OC-STAMP overexpression or silica, and we found that treatment with 4-PBA inhibited the activation of senescence signaling in silica-induced or OC-STAMP overexpressing MLE-12 cells (Figures 6 and 7).

3.5. OC-STAMP Interacts with MYH9 to Promote Cellular Senescence. To explore the effect of OC-STAMP in MLE-12 cells, we screened for potential OC-STAMP interacting proteins. We performed CoIP using an anti-OC-STAMP antibody followed by LC-MS/MS assay. MYH9 was identified as a potential OC-STAMP interacting protein. As shown in Figure 8, Western blot analysis of the precipitates with an OC-STAMP antibody indicated OC-STAMP CoIP with MYH9. Downregulated expression of MYH9 with siRNA also inhibited senescence signaling in Ocstamp-overexpressing MLE12 cells.

3.6. Inhibition of ER Stress Attenuates Cellular Senescence in Silicotic Rats. In the present study, the established model [1] was used to explain the effect of ER stress on cellular senescence in silicotic rats. Treatment with 4-PBA attenuated...
the activation of senescence signaling, collagen deposition, and high expression of OC-STAMP in silicotic rats, which suggested that blocking of ER stress inhibited cellular senescence in pulmonary fibrosis induced by silica (Figure 9).

4. Discussion

Hypertrophy and hyperplasia of AEC2 is one of the prominent features of silicosis and is consistently associated with
Figure 5: Overexpression of OC-STAMP induces insensitivity in silica-induced MLE-12 cells. Levels of p-ATM, p-ATR, p-p53, p21, p16, E-cadherin, N-cadherin, α-SMA in NC, and OC MLE-12 cells treated with or without silica, measured by Western blotting. Data are presented as the mean ± SD; n = 3 per group.

Figure 6: 4-PBA treatment inhibits EMT, ER stress, and cellular senescence in OC-STAMP-overexpressing MLE-12 cells. Levels of Col I, E-cadherin, N-cadherin, α-SMA, p-PERK, p-EF-2α, p-IRE-1α, p-ATM, p-ATR, p-p53, p21, and p16 in MLE-12 cells, measured by Western blotting. Data are presented as the mean ± SD; n = 3 per group.
Figure 7: 4-PBA treatment inhibits EMT, ER stress, and cellular senescence in silica-treated MLE-12 cells. Levels of Col I, E-cadherin, N-cadherin, α-SMA, p-PERK, p-EIF-2α, p-IRE-1α, p-ATM, p-ATR, p-p53, p21, and p16 in MLE-12 cells treated with silica, measured by Western blotting. Data are presented as the mean ± SD; n = 3 per group.

Figure 8: OC-STAMP interacts with MYH9. (a) Co-IP of OC-STAMP and MYH9; (b) levels of p-ATM, p-ATR, p-p53, p21, and p16 in siRNA-Myh9 treated MLE-12 cells, measured by Western blotting. Data are presented as the mean ± SD; n = 3 per group.
alveolitis, but the contribution of AEC2 in the pathogenesis of silicosis is largely unknown [18, 19]. Several studies have suggested that hypertrophic and hyperplastic AEC2 were proliferative AEC2, as well as enhanced production and secretion of phospholipids and surfactant proteins for lung injury and repair [20, 21]. In the present study, we analyzed the number of AEC2, collagen deposition, expression of p21 and PCNA in normal alveoli, alveolitis (consisting of macrophages and hypertrophic AEC2), and silicotic granulomas. As observed in previous studies, hypertrophic and hyperplastic AEC2 was mostly located in the alveolitis-affected area and showed a senescent phenotype with more collagen deposition. An in vitro study also showed activation of cellular senescence signaling in silica-treated MLE-12 cells. Most importantly, we found overexpression of OC-STAMP, one of the major factors of RANKL signaling, derived EMT, ER stress, and cellular senescence in MLE-12 cells, which showed some similar features in rats exposed to silica. OC-STAMP knockout (KO) mice showed normal skeleton, growth, and bone metabolic markers, and OC-STAMP-deficient cells isolated from bone marrow were able to differentiate into TRAP-positive osteoclasts under RANKL stimulation but could not fuse into multinucleated cells, which suggested the specific role of OC-STAMP in osteoclast multinucleation or cell fusion rather than osteoclast differentiation [9, 10]. Thus, several studies have proposed that OC-

![Figure 9: 4-PBA attenuates cellular senescence in silicotic rats. (a) Coexpression of p21 and proSPC in silicotic rats, Bar = 50 μm; (b) levels of p-PERK, p-EIF 2α, p-IRE-1α, p-ATM, p-ATR, p-p53, p21, and p16 in MLE-12 cells treated with silica, measured by Western blotting. Data are presented as the mean ± SD; n = 3 per group.]
STAMP is involved in pathogenic bone resorption rather than normal bone metabolism [11]. Furthermore, OC-STAMP induced a phenotypic switch in macrophage polarization and suppressed the M1 proinflammatory state [22]. We have been described the potential proinflammatory effect of OC-STAMP in macrophages, as a member of the RANKL signaling pathway in silicosis [8]. In the present study, we found a different mechanism of OC-STAMP in AEC2 and promoted EMT, ER-stress, and cellular senescence in MLE-12 cells, which could be blocked by Octstamp-siRNA or 4-PBA. Furthermore, ER stress has been reported to be associated with EMT and resulted in an increase in the p16 and p21 levels in lung epithelial cells in pulmonary fibrosis [23]. ER stress has been also observed in senescence induced by different stimuli and has been proposed as the consequence of senescent phenotype [24]. Combined with our previous study [1, 25], we speculated that the EMT, ER stress, and cellular senescence worked together in silicosis, at least in part, by the overexpression of OC-STAMP.

Our data showed that OC-STAMP interacted with nonmuscle myosin class II, isoform A (NM II-A, also known as MYH9), which regulated the senescent signaling pathway. MYH9 is an actin-binding molecular motor and is encoded by the Myh9 gene, which participates in many crucial cellular processes, such as adhesion, cell migration, cytokinesis and polarization, maintenance of cell shape, and signal transduction [26]. It has been reported that MYH9 localization and filament assembly can be modulated by the interaction with S100A16 during kidney injury or TGF-β stimulation to promote cytoskeleton reorganization and EMT progression in renal tubulointerstitial fibrosis [27]. TGF-β1 increased MYH9 expression, and siRNA-mediated knockdown of MYH9 remarkably repressed TGF-β1-induced lung fibroblast-to-myofibroblast differentiation [28]. Furthermore, inducible conditional knockout of Myh9 in the renal tubules of adult mice resulted in progressive kidney disease with expansion of ER tubules and activation of ER stress [29]. Our data showed that OC-STAMP interacted with MYH9, which regulated cellular senescence signaling in MLE-12 cells and may have an important role in silicosis. This study has some limitations. First, the results should be verified in clinical samples to strengthen the meaning of high expression of OC-STAMP in silicosis. Furthermore, the cross talk between ER stress and senescence in silicosis still needs to be explored to better understand age-related lung pathology and pathophysiology. Further studies are needed to consider and overcome these limitations.

In summary, we have shown that cellular senescence of AEC2 participates in silicosis formation. In the context of profibrotic insults, overexpression of OC-STAMP in MLE-12 cells exacerbates ER stress, EMT, and cellular senescence, and it may play an important role in silicosis. Blockage of ER stress protects against cellular senescence and pulmonary fibrosis in response to silica exposure.

**Data Availability**
The underlying data of the study can be obtained by contacting the authors if it is reasonable.

**Conflicts of Interest**
The authors declare that they have no competing interests.

**Authors’ Contributions**
All authors contributed to data analysis and drafting or revision of the article. All authors have agreed on the journal to which the article will be submitted, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

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**Supplementary Materials**
Figure S1: the cell apoptosis in MLE-12 cells treated with silica or not. There was no statistically significant difference in apoptosis rate between MLE-12 cells treated with or without 50 μg/mL silica. (Supplementary Materials)

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