Letter to the Editor

The Role of Fibrocyte in the Pathogenesis of Silicosis*

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Exposure to free silica induces silicosis and myofibroblasts are regarded as primary effector cells. Fibrocytes can differentiate into myofibroblast. Therefore, the present study was designed to investigate whether fibrocytes participate in silicosis. The rat model of silicosis was established. Hematoxylin-eosin stainings and Masson stainings were used to evaluate the histopathology and collagen deposition. Flow cytometry and immunofluorescence were performed to detect the number of fibrocytes and their contribution to myofibroblasts. Results showed that fibrocytes participate in silicosis. Trend analysis of different sources of myofibroblasts during silicosis indicated that fibrocytes and lung type II epithelial cell-derived myofibroblasts play an important role in the early stage of silicosis, while resident lung fibroblast-derived myofibroblasts play a predominant role during the fibrosis formative period.

Chronic inhalation of crystalline silica (SiO₂) is associated with the development of silicosis, which is characterized by tissue remodelling, fibroproliferation, and deposition of extracellular matrix (ECM). Myofibroblasts, as key effector cells, are believed to play a major role in the pathogenesis of fibrotic diseases. The source of these cells has been intensively debated. Resident lung fibroblasts were conventionally thought to be the main source of myofibroblasts. However, recent studies have demonstrated that epithelial-mesenchymal differentiation or recruitment of fibrocytes also contribute to myofibroblasts in lung fibrosis. Little attention has been focused on fibrocytes and the contribution of different sources of myofibroblasts in silicosis. Among the possible sources of lung myofibroblasts, fibrocytes are bone marrow-derived cells that co-express CD45 and collagen type I (CD45⁺ Col I⁺)³. Fibrocytes participate in both tissue fibrosis and tissue remodelling by producing ECM and matrix metalloproteinases². Recent studies have reported that the number of fibrocytes is obviously increased in patients with idiopathic pulmonary fibrosis (IPF) and that higher numbers of them are related to early mortality³. Taken together, a growing body of evidence indicates that fibrocytes participate in lung. In our present study, we sought to evaluate the contribution of fibrocytes to silicosis. We compared lung tissue and peripheral blood from rats with silicosis and control rats for the presence and number of fibrocytes and contribution of the three sources of myofibroblasts during the process of experimental silicosis. Our findings support the notion that fibrocytes may participate in silicosis and that different sources of myofibroblasts may play roles at different stages of silicosis.

The animal experiments were approved by the Experimental Animal Ethics Committee of Zhengzhou University, and our study was conducted in accordance with the guidelines of the Chinese Association of Laboratory Animal Care. The SD rats were randomly divided into the model group and control group (n = 36 per group). The rats in the model group were administered a single intratracheal instillation of SiO₂ solution (100 mg/0.5 mL per rat) and those of control group were instilled an equal amount of 0.9% saline. All of the rats were anaesthetized with chloral hydrate and then sacrificed by cervical dislocation for further study at 1, 2, 3, 6, 9, and 12 weeks (n = 12 per week) after silica/saline instillation.

Hematoxylin-eosin (HE) and Masson stainings were used to evaluate the histopathology and collagen deposition. The extent of lung fibrosis was evaluated by the measurement of the area of

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fibroblast-like cells need to participate in pathological repair and reconstruction. At this time, bone marrow may supply more fibrocyte precursor cells to maintain a high level of fibrocytes in the peripheral blood or lung tissue. The further study confirmed CD45\(^+\) collagen I\(^+\) fibrocytes were existent in the lung tissue. CD45\(^+\) collagen I\(^+\) fibrocytes contributed to approximately 10% to 16% of the collagen I-expressing cells in the lungs of SiO\(_2\)-exposed rats which suggests that fibrocytes play a role in the ECM deposit in silicosis (Figure 2A and 2B).

Based on animal models of fibrosis in parenchymal organs, fibrocytes accounted for approximately 5% to 20% of myofibroblasts in fibrotic organs. The highest amounts were in bleomycin-induced lung fibrosis, of which fibrocyte-derived myofibroblasts comprise 25% to 50% of the total number of myofibroblasts\(^{[6]}\). These data suggest that fibrocyte recruitment may contribute to lung fibrosis. Silicosis is a type of fibrotic disease, and the number of fibrocytes increases during the process of silicosis. Therefore, we wonder whether there are fibrocyte-derived myofibroblasts, what is their contribution, and what is the interaction of the different sources of myofibroblasts in the course of silicosis. To work out the above questions, CD45\(^+\) \(\alpha\)-SMA\(^+\) and SPC\(^+\) \(\alpha\)-SMA\(^+\) were used to identified fibrocyte-derived myofibroblasts and AT II-derived myofibroblasts, respectively. The results showed that fibrocyte-derived myofibroblasts (15%-35%, Figure 3A and 3C) and AT II-derived myofibroblasts (9% -21%, Figure 3B and 3D) were increased in the lung tissue of silicosis rats at different time points. We calculated the contribution of resident fibroblast-derived myofibroblasts as being approximately 40% to 65% according to the other two sources of myofibroblast contributions. As showed in Figure 3E, the numbers of fibrocyte-derived and AT II-derived myofibroblasts have increasing trends at the inflammatory response period (at 1, 2, and 3 weeks) and then decreasing trends at the fibrotic period (at 6, 9, and 12 weeks). However, those of resident fibroblast-derived myofibroblasts in the lungs showed opposite trends. The dynamic changes of different sources of myofibroblasts indicated that they may play roles in different time points of silicosis. Fibrocyte-derived and AT II-derived myofibroblasts may act mainly during
the inflammatory response period. Fibrocytes are well-known to be capable of inflammatory chemotaxis and secreting chemokines, including inflammatory factors and chemokines, which not only maintain the local inflammatory microenvironment but also recruit many types of cells to amplify the inflammatory response\(^7\). We assume that fibrocytes may maintain the inflammatory microenvironment through pro-inflammatory and paracrine functions in the early stage of silicosis and may initiate the later stage of the fibrotic process. Resident fibroblast-derived myofibroblasts in the lungs play roles in the

**Figure 1.** SiO\(_2\) exposure increased the number of fibrocytes in peripheral blood and lung tissue. The number of fibrocytes in peripheral blood was increased from week 1 to week 9, there were no significant difference was observed at week 1 (A and C). The number of fibrocytes in lung tissue was increased from week 1 to week 12 (B and D). Data are expressed as mean ± SD. *P < 0.05 vs. control.
pathological fibrotic process in the late stage of silicosis. The cells in the lungs released a large number of cytokines that may promote lung resident fibroblasts to differentiate into myofibroblasts, and these myofibroblasts are capable of a greater proliferation and secretion of collagens. It is remarkable that fibrocytes can lose their CD45 phenotype in vitro\(^8\). Perhaps this phenomenon exists in vivo. With an increased SiO\(_2\) exposure time, fibrocytes migrated to the lungs and lost their CD45 phenotype. According to our current calculation method, these cells are counted into resident fibroblasts. Thus, using the CD45\(^+\) α-SMA\(^+\) method to identify fibrocyte-derived myofibroblasts may underestimate its contributions and overestimate the contribution of lung resident fibroblasts.

Our study suggests that fibrocytes may participate in silicosis and play an important role in the early stage of silicosis. In addition, it is feasible to block migration of fibrocytes due to their existence in peripheral blood, so fibrocytes may represent a novel therapeutic target for developing more effective treatments against silicosis.

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No conflict of interests to declare.

Figure 2. Fibrocyte is exist in lung tissue (A) and SiO\(_2\) exposure increased the percent of fibrocyte derived collagen I secreting cells (B). The red represents CD45 and green represents Collagen I. The arrow points to Fibrocytes (yellow). Data are expressed as mean ± SD. *\(P < 0.05\) vs. control.
Figure 3. The contribution of different sources of myofibroblasts. The red represents CD45 and green represents α-SMA and the number of fibrocytes-derived myofibroblasts (yellow) was increased from week 1 to week 12 (A and C). The red represents SPC and green represents α-SMA and the number of AT II-derived myofibroblasts (yellow) was elevated from week 1 to week 12 (B and D). The trend changing analysis of different sources of myofibroblasts (E). Data are expressed as mean ± SD. *P < 0.05 vs. control.
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Figure S1. SiO₂ exposure induces lung inflammation and fibrosis. Pathological changes and morphological alteration of lung tissues were evaluated by HE staining (A, magnification ×200). Masson trichrome staining was used to determine collagen deposition in lung. Areas of fibrosis (%) were quantified with Image-Pro Plus 6.0 software. The areas of fibrosis was increased from week 2 to week 9 (B and C, magnification ×200). Data are expressed as mean ± SD. * P < 0.05, vs. control.