Intratracheal Transplantation of Alveolar Type II Cells Reverses Bleomycin-induced Lung Fibrosis

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Rationale: Transplantation of stem cells has been proposed as a strategy for repair of lung fibrosis. Nevertheless, many studies have yielded controversial results that currently limit the potential use of these cells as an efficient treatment. Alveolar type II cells are the progenitor cells of the pulmonary epithelium and usually proliferate after epithelial cell injury. During lung fibrosis, however, the altered regeneration process leads to uncontrolled fibroblast proliferation.

Objectives: To investigate whether intratracheal transplantation of isolated alveolar type II cells can halt and reverse the fibrotic process in an experimental model of bleomycin-induced lung fibrosis in rats.

Methods: Lung fibrosis was induced in syngeneic female Lewis rats by a single intratracheal instillation of bleomycin (2.5 U/kg). Animals were transplanted with alveolar type II cells from male animals at a dose of 2.5 × 106 cells per animal 3, 7, and 15 days after endotracheal bleomycin instillation. Animals were killed 21 days after the induction of lung fibrosis.

Measurements and Main Results: Lung fibrosis was assessed by histologic study and determination of hydroxyproline content. Engraftment of transplanted cells was measured by real-time polymerase chain reaction for the Y chromosome and by fluorescence in situ hybridization for the Y chromosome. Transplantation of alveolar type II cells into damaged lung 3, 7, or 15 days after bleomycin instillation led to reduced collagen deposition, and reduction in the severity of pulmonary fibrosis.

Conclusions: This study demonstrates the potential role of alveolar type II cell transplantation in designing future therapies for lung fibrosis.

Keywords: cell therapy; lung epithelium; lung repair

Idiopathic pulmonary fibrosis is a chronic inflammatory interstitial lung disease that has a potentially fatal prognosis, with death occurring in most patients within 5 years of diagnosis. Conventional therapy consists of glucocorticoids or immunosuppressive drugs, which are usually ineffective in preventing the progression of fibrosis (1). A number of studies showed that stem cells derived from adult tissues could differentiate into lung epithelial cells and confer a functional benefit (2–7). Therefore, the contribution of adult stem cells to the regeneration of lung tissue was considered an alternative approach to the treatment of certain lung diseases. However, despite these promising results, studies have failed to restore lung epithelium with adult stem cells (8–10).

Classically, the progenitor cells of the pulmonary epithelium are considered to be the alveolar type II cells in the alveoli.
METHODS

Animals
Syngeneic Lewis rats, weighing 170–200 g at the beginning of the experiment, were used according to European Community (Directive 86/609/EEC) and Spanish guidelines for the treatment of experimental animals.

BLM-induced Lung Fibrosis
Pulmonary fibrosis was induced in female animals by intratracheal instillation of a sublethal dose of BLM (2.5 U/kg) dissolved in 400 μl of sterile saline under halothane anesthesia (17). Control animals received the same volume of saline. During the experiment, animal body weights were recorded every 2 days.

Purification of Alveolar Type II Cells
Fresh alveolar type II cells were isolated from healthy male donor animals. The protocol for purification has been described elsewhere (18). Briefly, to isolate alveolar type II cells, the lungs were removed from each animal and lavaged six times, each time with 8 ml of saline. The lungs were digested with 0.25% trypsin (T8003; Sigma, St. Louis, MO) dissolved in saline (100 ml) and suspended in 0.9% NaCl at 37°C for 30 minutes. The trypsin was constantly topped up during the 30-minute expansion of the parenchyma.

After digestion, the lungs were chopped into 1- to 2-mm² cubes, treated with DNase dissolved in saline (7,500 U/100 ml), and filtered through nylon meshes ranging in pore size from 150 to 30 μm. The resulting cell suspension was centrifuged at 250 × g for 20 minutes. These cells were resuspended in 5 ml of DCCM-1 (Biological Industries, Kibbutz Beit Haemek, Israel) supplemented with 2% (wt/vol) L-glutamine and subjected to differential attachment on a plastic Petri dish. No adherent alveolar type II cells were collected after 2 hours and cells were counted to establish the final yield of freshly purified cells.

Alveolar type II cell viability was assessed with trypan blue (Sigma), showing greater than 95% viability. Cell yield and purity was assessed with an alkaline phosphatase staining kit (Sigma). Cells were cytocentrifuged (Cytospin 3; Shandon Scientific Ltd, Grupo Taper, Madrid, Spain) and cell smears were allowed to air dry and were stained for 15 minutes at room temperature. Staining was conducted according to the manufacturer’s protocol for the use of alkaline phosphatase (18). Cells were evaluated by transmission electron microscopy. The cells were fixed in 2.5% glutaraldehyde in phosphate buffer (0.1 mol/L, pH 7.4) for at least 2 hours at 4°C. The cells were secondarily fixed in 1% osmium tetroxide and 0.8% potassium ferrocyanide for 1 hour at 4°C. After three washes with cold double-distilled water, the samples were dehydrated through an ascending concentration of acetone (30, 50, 70, 95, and 100%), and three changes of 100% acetone. They were then embedded in Spurr resin and polymerized at 60°C. The embedded blocks were sectioned with a diamond knife (Diatome, Biel, Switzerland) on a Leica Ultracut UCT (Leica Microsystems, Deerfield, IL). Ultrathin sections were placed on copper grids and stained with uranyl acetate and lead citrate before examination at the electron microscope level. More than 95% (n = 4) of purified rat cells stained positively for alkaline phosphatase.

Experimental Groups
Animals were randomly distributed into six experimental groups (n = 8 for each group):

- Control: Saline instillation
- Control plus transplantation: Saline instillation followed by alveolar type II cell transplantation 3 days after instillation
- BLM: Bleomycin instillation
- BLM plus transplantation (3 d): Bleomycin instillation plus alveolar type II cell transplantation 3 days after instillation
- BLM plus transplantation (7 d): Bleomycin instillation plus alveolar type II cell transplantation 7 days after instillation
- BLM plus transplantation (15 d): Bleomycin instillation plus alveolar type II cell transplantation 15 days after instillation

Transplantation Procedure
On Days 3, 7, and 15 after endotracheal BLM administration, female recipient animals were transplanted with alveolar type II cells endotracheally, by the transoral route under halothane anesthesia. Each animal received a dose of 2.5 × 10⁶ cells suspended in 400 μl of sterile saline. The control group received the same dose of cells 3 days after saline instillation. Animals were killed 21 days after induction of lung fibrosis.

Real-time Polymerase Chain Reaction
The engraftment of donor alveolar type II cells was evaluated by examination of the lung genomic DNA of the transplanted animals, using the real-time polymerase chain reaction (PCR) as described (19) and an iCycler iQ5 real-time PCR detection system (Bio-Rad Laboratories, Hercules, CA) using PCR Y chromosome primers 5’-CTG-GCT-CTG-CCT-CTA-CCT-AGT-C-3’ and 5’ATT-CTC-CTT-TCT-TAC-ACA-CTA-CAC-ATC-C-3’ and an iQ SYBR Green supermix kit (Bio-Rad Laboratories). Standard curves were generated by serially diluting male rat genomic DNA into female rat genomic DNA prepared from liver.

Fluorescence In Situ Hybridization for the Y Chromosome
Fluorescence in situ hybridization (FISH) assays of formalin-fixed tissue sections were performed to identify the Y chromosomes. Briefly, paraffin sections of 4 μm were deparaffinized and incubated in saline–sodium citrate (4× SSC buffer, pH 7.0) for 30 minutes, and then the slides were incubated in 1 M sodium isothiocyanate at room temperature overnight. Slices were incubated in pepsin for 5 minutes and washed twice in phosphate-buffered saline (PBS) for 5 minutes, washed once with formaldehyde-PBS–MgCl₂ for 1.5 minutes and with PBS for 1.5 minutes, then followed by serial ethanol dehydration steps (1.5 min each). Sections were denatured at 65°C for 2 minutes in preheated 70% formamide and 2× SSC buffer, pH 7.0, and subsequently “quenched” with ice-cold 70% ethanol for 1.5 minutes. Serial ethanol dehydration was done again. The rat Y chromosome probe labeled with fluorescein isothiocyanate (Star*FISH; Cambio, Cambridge, UK) was denatured at 65°C for 10 minutes and applied to the sections at 45°C. The sections were coverslipped and sealed with rubber cement for incubation overnight in a hydrated slide box at 42°C. Coverslips were carefully removed in preheated 2× SSC buffer, pH 7.0, at 45°C. Sections were preheated in 50% formamide in 2× SSC buffer for 5 minutes each at 45°C and gently washed twice in preheated 0.1× SSC buffer for 5 minutes each at 45°C. Sections were observed by confocal microscopy. Confocal images were acquired with a Leica TCS SL laser-scanning confocal spectral microscope (Leica Microsystems Heidelberg GmbH, Mannheim, Germany) with argon and helium-neon lasers attached to a Leica DMI2500 inverted microscope. Fluorescein isothiocyanate and propidium iodide emission were acquired sequentially with a triple dichroic beam splitter (TD 488/543/633 nm) and emission detection ranges of 500–553 and 592–700 nm, respectively. All images were obtained with a ×63 oil immersion objective lens (numeric aperture, 1.32) equipped with phase-contrast optics and a confocal pinhole setting of 1 Airy unit. Because of the small size of the cells an electronic zoom (×2) was necessary for stronger magnification and better image resolution. Image assembly and treatment were performed with image-processing Leica confocal software.

Hydroxyproline Content
Lung hydroxyproline content was measured as an indicator of collagen deposition by the method outlined by Woessner (20). Samples were homogenized and then hydrolyzed in 6 M HCl for 18 hours at 110°C. The hydrolysate was then neutralized with 2.5 M NaOH. Aliquots
(2 ml) were analyzed for hydroxyproline content after addition of 1 ml of chloramine-T, 1 ml of perchloric acid, and 1 ml of dimethylaminobenzaldehyde. Samples were read for absorbance at 550 nm in a spectrophotometer. Results are expressed as micrograms of hydroxyproline per lung.

Histology

For histologic studies, the left lung was perfused via its main bronchus with a fixative solution (10% neutral-buffered formalin) at a pressure of 25 cm H2O and immersed in the fixative for 12 ± 24 hours. Lung tissue blocks were then taken and placed in formalin, dehydrated in a graded series of ethanol, embedded in paraaffin, cut into 4-μm-thick serial sections, and stained with hematoxylin–eosin and Masson’s trichrome to identify inflammatory cells, connective tissue, and collagen deposition.

Macrophage Transplantation

An additional experiment involving macrophage transplantation was performed as a control, using an unrelated cell type to determine whether the fibrotic reversal response was specific for alveolar type II cells. Fresh macrophages were isolated from healthy male donor animals by performing bronchoalveolar lavage. Bronchoalveolar lavage fluid was obtained by washing the lungs four times with 10 ml saline aliquots through a tracheal cannula and centrifuged (300 × 2 min, 4°C). Cell viability was assessed with trypan blue and was greater than 95%. To evaluate the cell yield and purity and to characterize freshly isolated macrophages, Cytospin preparations were stained with Diff-Quik (Diagnostics Grifols SA, Barcelona, Spain) according to the manufacturer’s protocol. More than 95% (n = 4) of rat bronchoalveolar cells were macrophages (see Figure 5A).

On Day 15 after bleomycin instillation, female recipient animals were transplanted with macrophages endotracheally, by the transoral route under halothane anesthesia, at a dose of 2.5 × 10^6 cells per animal. The cells were suspended in 400 μl of sterile saline. The control group received the same dose of cells 15 days after saline instillation. Animals were killed 21 days after the induction of lung fibrosis. The animals were randomly distributed into four experimental groups (n = 8 for each group):

- **Control**: Saline instillation
- **Control plus transplanted macrophages**: Saline instillation plus macrophage transplantation 15 days after instillation
- **BLM**: Bleomycin instillation
- **BLM plus transplanted macrophages (15 d)**: Bleomycin instillation plus macrophage transplantation 15 days after instillation

Statistical Analysis

Data are expressed as means (SEM) with 95% confidence intervals. Statistical analysis was performed by analysis of variance followed by appropriate post hoc tests including the Newman-Keuls test when differences were significant (GraphPad Software, Inc., San Diego, CA). A P value less than 0.05 was considered significant.

RESULTS

BLM instillation initially causes a marked loss in body weight, which is normally recovered later. After alveolar type II cell transplantation, animals regained weight more quickly than did BLM nontransplanted animals (Figure 2A). This indicated that alveolar type II cell transplantation enhanced the health of the animals.

After lung collection, the macroscopic appearance of BLM nontransplanted lungs was fibrotic. They were large, yellowish, and had several scars when compared with saline control lungs (Figure 2B). In contrast, BLM transplanted lungs showed an improved appearance similar to that of the saline control lungs and were devoid of scars (Figure 2B). Figure 2C shows Masson’s trichrome–stained lung sections examined by stereomicroscopy to illustrate the heterogeneous topography of the fibrotic lesions (patchy areas of lung fibrosis); BLM nontransplanted lungs showed extensive areas of fibrosis compared with BLM transplanted lungs.

BLM instillation caused an increase in lung weight owing to the inflammatory and fibrotic component (17). Lung weight was significantly decreased in the transplanted groups 7 and 15 days after BLM instillation compared with the BLM nontransplanted group (Figure 2D).

Because alveolar type II cell transplant donors and recipients were sex-mismatched, engraftment levels of male donor–derived cells in the lungs of female transplant recipients were quantified by real-time PCR (19). No male DNA was detected in the control transplant animals. We found 0.18, 0.30, and 0.49% male DNA in the transplanted groups 3, 7, and 15 days after BLM, respectively (Figure 3A). These data demonstrate that the engraftment of alveolar type II cells is more effective when fibrosis has developed than in its early phases.

In addition, the successful engraftment of male donor cells in the female transplanted animals was assessed by FISH of the Y chromosome. FISH analysis was performed in paraffin-embedded sections of lung tissue, using Y chromosome paint. Y chromosome–positive FISH signals (absent in the female controls and present in the male controls) were observed in lung sections from all the transplanted groups (Figure 3B). In sections of transplanted lungs, these signals were localized mainly in areas of fibrosis.

BLM administration to rats results in an increase in collagen deposition in the lung. Therefore, the amount of hydroxyproline, a modified amino acid specifically found in collagen (20), was assessed to determine how alveolar type II transplantation could alter the clinical course of BLM-induced lung injury. BLM nontransplanted lungs showed a significant increase in the amount of lung hydroxyproline when compared with saline control rats (Figure 4A). In contrast, levels of hydroxyproline were significantly reduced in all the BLM transplanted groups. In the groups transplanted on Days 7 and 15 after BLM instillation, we observed a greater reduction in hydroxyproline levels, which did not significantly differ from levels observed in the saline control groups. These results confirm that transplantation of alveolar type II cells induces a reduction in collagen deposition and, consequently, in the fibrotic response.

To further examine the effect of alveolar type II cell transplantation in BLM-induced pulmonary fibrosis, serial lung sections were stained with hematoxylin–eosin or Masson’s trichrome and examined by light microscopy. Lung tissue sections from control rats showed no evidence of inflammation or epithelial damage. As expected, lung tissue sections from rats with BLM-induced fibrosis (nontransplanted) showed marked peribronchial and interstitial infiltration with inflammatory cells, extensive cellular thickening of interalveolar septa, interstitial edema, increased interstitial cells with a fibroblastic appearance, excessive collagen deposition, and focal cuboidal metaplasia of alveolar lining cells (Figure 4B, row a). The pattern of lesions was multifocal (i.e., patchy areas of pulmonary fibrosis) and in most cases involved the pleura. Although multifocal parenchymal lesions were still present in lungs transplanted with alveolar type II cells, the organized fibrolastic foci were smaller and considerably less frequent than were observed in nontransplanted animals. The reduction in parenchymal lesions was evidenced by large areas of undamaged tissue with normal alveolar architecture (Figure 4B, row b). Compared with the BLM nontransplanted group, BLM transplanted animals showed less edema, less collagen deposition, less septal widening, and fewer clusters of inflammatory cells.

Specificity of the alveolar type II cells to revert the fibrotic process was verified in a group of animals transplanted with macrophages (Figure 5). When macrophages were used for cell...
transplantation instead of alveolar type II cells, no beneficial effects were observed. Body weight (Figure 5B), lung weight (Figure 5C), and survival rate showed no significant differences from the BLM nontransplanted group. In addition, hydroxyproline levels remained unmodified (Figure 5D) and the fibrotic areas evaluated by light microscopy after hematoxylin–eosin and Masson’s trichrome staining revealed a similar degree of fibrosis compared with that observed in BLM nontransplanted animals.

DISCUSSION
In this study, we demonstrate for the first time that intratracheal transplantation of alveolar type II cells can halt and reverse lung fibrosis. Alveolar epithelial damage is an important initial event in pulmonary fibrosis. Epithelial cell damage and cell death induce the formation of gaps in the epithelial basement membranes. Migration of fibroblasts through these gaps into the alveolar space leads to intraalveolar fibrosis. Normal repair of the epithelial layer occurs through the proliferation of alveolar type II cells and their subsequent differentiation to alveolar type I cells necessary for proper lung function. In lung fibrosis, however, the regenerative process is altered, with the number of alveolar type II cells decreasing markedly in areas of severe inflammation. Extensive injury and cell death follow (21, 22). In areas of severe damage, alveolar type I and type II cells die and are replaced by a large number of fibroblasts and smooth muscle cells. The injury, together with the inadequate repair of

Figure 1. (A) Cytospin preparation showing alkaline phosphatase–positive cells. Original magnification, ×200. (B) Transmission electron micrographs of alveolar type II cells showing lamellar bodies. Bar = 2 μm. (C) Detail of the lamellar bodies. Bar = 1 μm.

Figure 2. (A) Change in animal body weight over time. On Day 0, the animals received bleomycin (BLM). Arrowheads show the days of alveolar type II cell transplantation (Trp) (Days 3, 7, and 15). (B) Representative photographs of whole lungs from all the experimental groups after 21 days of BLM-induced lung fibrosis. Arrows show the scars in the BLM nontransplanted lung. BLM transplanted lungs showed an improved aspect compared with BLM nontransplanted lungs. (C) Representative photographs of lung sections from all the experimental groups. Multifocal parenchymal lesions were smaller in BLM transplanted lungs compared with BLM nontransplanted lungs. Scale bars: 1 mm. (D) Lung weight after 21 days of BLM-induced lung fibrosis in experimental groups. Alveolar type II cells transplanted 7 or 15 days after BLM instillation reduced lung weight. Data represent means ± SEM of eight animals per group (*P < 0.05 vs. control groups, #P < 0.05 vs. BLM).
epithelial cells, disturbs normal epithelial–fibroblast interactions and leads to pulmonary fibrosis.

Histologic and biochemical results show that transplantation of alveolar type II cells in rats with BLM-induced lung fibrosis can reverse the fibrogenic response. However, more studies are needed to ascertain the mechanism whereby alveolar type II cells induce recovery in the lung, as several causes could explain this effect. First, the alveolar type II cells could limit the deleterious effects of BLM by replacing already damaged alveolar epithelial type II cells, which are known to be targets of apoptotic signals induced by BLM (22). Alveolar type II cells are known to be the alveolar progenitor cells, which give rise to alveolar type I cells in response to injury (23). Thus, transplantation of alveolar type II cells may restore this pool, leading to an increased number of alveolar cells for the resolution of disrupted alveolar surfaces. In turn, this could promote the repair process. Second, alveolar type II cells have a number of physiological functions, including synthesis, secretion, and turnover of pulmonary surfactant (23). Surfactant maintains low surface tension at the air–liquid interface of the alveoli, which is necessary for proper respiratory function. Transplantation of alveolar type II cells could thus improve the maintenance of normal lung function, promoting normal epithelial repair. Third, alveolar type II cells could protect against BLM-induced fibrosis by altering the lung microenvironment at sites of engraftment, producing a number of cytokine antagonists that disrupt signal pathways leading to fibrosis (23–26). Alveolar type II cells are a rich source of chemokines, including inhibition factors for fibroblast proliferation and secretion of collagen (27). Therefore, secretion of these products from alveolar type II cells could induce degradation of the new collagen deposition, halting the development of lung fibrosis.
Figure 4. (A) Lung hydroxyproline levels after 21 days of bleomycin (BLM)-induced lung fibrosis. Alveolar type II cell transplantation reduced the lung content of hydroxyproline at all times tested. Data represent means ± SEM of eight animals per group (*P < 0.05 and **P < 0.001 vs. control groups, #P < 0.05 and $$P < 0.001 vs. BLM). (B) Representative photomicrographs of lung histopathology in all the experimental groups after 21 days of BLM-induced lung fibrosis. Row a: Lung sections were stained with hematoxylin–eosin. Alveolar type II cell transplantation ameliorated the inflammatory and pulmonary lesions. Original magnification, ×100. Row b: Masson’s trichrome of all the experimental groups. The presence of interstitial collagen (blue staining) was diminished by alveolar type II cells in all the transplantation groups. Original magnification, ×100.

Figure 5. (A) Cytospin preparation showing alveolar macrophages. Original magnification, ×200. (B) Change in animal body weight over time. On Day 0 the animals received BLM. Arrowheads show the day of macrophage transplantation (Day 15). (C) Lung weight after 21 days of BLM-induced lung fibrosis in the experimental groups. (D) Lung hydroxyproline levels after 21 days of BLM-induced lung fibrosis. Macrophage transplantation did not reduce the lung content of hydroxyproline. Data represent means ± SEM of eight animals per group (**P < 0.001 vs. control groups). (E) Representative photomicrographs of lung histopathology in all the macrophage experimental groups after 21 days of BLM-induced lung fibrosis. Row a: Lung sections were stained with hematoxylin and eosin. Macrophage transplantation did not reduce inflammatory and pulmonary lesions. Original magnification, ×100. Row b: Masson’s trichrome staining does not show a decrease in the presence of interstitial collagen (blue staining) in the macrophage-transplanted group. Original magnification, ×100.
The success of the method reported here could be due to the fact that we used adult progenitor cells and the intratracheal route for transplantation. This stands in contrast to earlier reports, in which stem cells derived from adult tissues, such as bone marrow–derived cells and mesenchymal stem cells, were transplanted by the intravenous route (2–6). Previous studies showed that bone marrow–derived cells transplanted intravenously engrafted in different nonhematopoietic tissues and acquired epithelial phenotypes, which suggested a new paradigm for epithelial tissue repair (2). Nevertheless, the initial enthusiasm vanished when subsequent studies either failed to reproduce these initial results (28) or concluded that engraftment events were due to transplanted cells fusing with endogenous somatic cells (28–32). One important caveat relates to the intrinsic structure of the lung, in which each alveolus comprises aseptate containing capillaries filled with circulating blood cells, which could have been misinterpreted as bone marrow–derived cells engrafted in the lung (33). In this regard, our method using the intratracheal transoral route directly provides the necessary cells for repairing the damaged lung. Because the origin of fibroblasts in pulmonary fibrosis is assumed to be intrapulmonary, the potential extrapulmonary origin of lung fibroblasts in pulmonary fibrosis merits further investigation. Evidence suggests that bone marrow–derived cells migrate to wound-healing sites and serve as sources of fibroblasts and myofibroblasts, which participate in the fibrotic process in response to the signals released during lung injury and fibrosis (1, 2, 8, 33–35). Therefore, it seems reasonable that the increase in these signals could lead to migratory recruitment of transplanted bone marrow–derived cells into the lung and could contribute to the pathogenesis of lung fibrosis. By contrast, administration of alveolar type II cells would provide the necessary cells for the repair process. This notion has been confirmed by transplantation of macrophages, which did not improve the course of lung fibrosis.

One important aspect of this study is the assignment of times for alveolar type II cell transplantation. Transplantation was performed at three different times after BLM-induced lung injury: in the inflammatory phase (3 d after BLM instillation); in the profibrotic phase (7 d after BLM instillation); and in the fibrotic phase (15 d after BLM instillation), when the disease was fully developed. This allowed us to determine the phase in which transplantation could restore the damaged lung. Surprisingly, our data indicate that alveolar type II cell transplantation can halt the fibrogenic process at any of these times. It is important to take into account that at 3 and 7 days after BLM instillation the fibroblasts are not growing yet, and therefore the administration of alveolar type II cells could in this case be preventing the fibrotic process.

For this reason, the most noteworthy result is the significant reduction of fibrosis 15 days after BLM administration, when the lungs are already fibrotic (21). Therefore, our data demonstrate that the transplantation of alveolar type II cells is able to reverse the fibrogenic process. In our opinion, this is a crucial finding given the absence of a useful treatment for this lung disease.

There are multiple therapeutic applications of this work. Because the alveolar type II cells can be obtained by in vitro differentiation of adult stem cells (36) and also by isolation from lung biopsies, it would be possible to perform transplantation in humans. Thus, adult stem cells could constitute an effective and reproducible source of alveolar type II cells, which would allow long-term treatment of lung fibrosis. Moreover, given the propensity of alveolar type II cells to engraft in damaged lung tissue and their ability to restore epithelial damage, these cells could also be an ideal target for gene therapy. Furthermore, because the intratracheal route of administration is commonly employed, this technique would be applicable to clinical practice.

In conclusion, these results show that intratracheal transplantation of alveolar type II cells could become a promising therapy for the future treatment of fibrotic lung diseases.

Conflict of Interest Statement: None of the authors has a financial relationship with a commercial entity that has an interest in the subject of this manuscript.

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References


