# Letter to the Editor

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A novel approach to pulmonary

fibrosis: Stem cell therapy

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To the Editor. Lung disease remains a pressing concern for public health, and pulmonary fibrosis

ORCID:

is a chronic and progressive manifestation of this ailment with limited treatment options.<sup>[1]</sup> However, recent research suggests stem cell therapy could provide a promising alternative for managing pulmonary fibrosis. With its unique ability to repair damaged lung tissue and modulate the immune response, stem cell therapy holds significant hope for those affected by this devastating condition.<sup>[2]</sup>

Several preclinical studies and clinical trials have examined the effectiveness of stem cell therapy for pulmonary fibrosis. These studies indicate encouraging outcomes, with many patients experiencing marked improvements in lung function and overall quality of life.[3] However, uncertainties remain regarding the optimal cell source, preferred delivery route, and appropriate dosing regimen for this treatment, which hinders the full realization of stem cell therapy's therapeutic potential.

Further research is crucial to address these uncertainties and advance stem cell therapy for pulmonary fibrosis. A key area of inves-

tigation involves determining the best cell source to treat this condition. While Mesenchymal Stem Cells are commonly used, other cell types such as Induced Pluripotent Stem Cells and Embryonic Stem Cells have shown potential in preclinical studies and merit additional exploration.<sup>[4]</sup>

Second, determining the optimal delivery method and dosing regimen for stem cell therapy in pulmonary fibrosis is essential. The method of delivery, along with the frequency and dose of stem cell administration, can influence the therapy's efficacy and safety.<sup>[5]</sup> Further studies are needed to pinpoint the optimal delivery route and dosing regimen for stem cell therapy.

Third, to further develop treatments for pulmonary fibrosis using stem cells, it is essential to investigate and understand the underlying mechanisms responsible for their regenerative and immunomodulatory properties. While stem cell therapy has showcased its potential in tissue repair, their specific effects on the immune system require deeper scrutiny.<sup>[2]</sup> Grasping these intricate mechanisms will pave the way for the development of more effective treatment methods [Fig. 1].<sup>[6]</sup>

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Figure 1: Mesenchymal Stem Cells Mechanisms (adapted from Wenzhao Cheng, 2022)<sup>[6]</sup>

TNF-α: Tumor necrosis factor alpha, TGF-β: Transforming growth factor beta, TLR-4: Toll-like receptor 4, EMT: Epithelial–mesenchymal transition, TIMPs: Tissue inhibitor of metalloprotease, MMPs: Matrix metalloproteinase, ATII: Alveolar epithelial type II cells

Fourth, it is crucial to establish the long-term safety and efficacy of stem cell therapy for pulmonary fibrosis. Currently, there is a lack of comprehensive evidence regarding the long-term impacts of this treatment, necessitating further research to confirm its safety and efficacy.<sup>[7]</sup>

In summary, while stem cell therapy offers a promising future for treating pulmonary fibrosis, additional research is imperative to address lingering questions and enhance the therapy's effectiveness. The transformative potential of stem cell therapy for pulmonary fibrosis is significant. A continued commitment to research is essential to fully harness this potential.

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