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Treatment for Pulmonary Fibrosis, Volume II

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Deadline for manuscript submissions:
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Message from the Guest Editors

Dear Colleagues,

Pulmonary fibrosis is characterized by the excessive deposition of extracellular matrices and destruction of the pulmonary parenchyma. The cause or contributing factor of pulmonary fibrosis is often unknown, and some diseases, including idiopathic pulmonary fibrosis, have poor prognosis despite treatment. Recently, progressive fibrosing interstitial lung disease (PF-ILD) or progressive pulmonary fibrosis (PPF) has been advocated to this phenotype, but there are many uncertainties and problems left to be resolved in the term “PF-ILD or PPF”.

This Special Issue, “Pulmonary Fibrosis: From Pathogenesis to Therapeutics”, aims to focus on novel approaches to the pathogenesis, diagnosis and therapeutics of pulmonary fibrosis at basic to clinical levels.

Prof. Dr. Hiroshi Mukae
Dr. Noriho Sakamoto
Guest Editors

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Message from the Editor-in-Chief

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