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Current Updates on Pediatric Cystic Fibrosis Care and Outcomes

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Message from the Guest Editors

Dear Colleagues,

Tremendous progress has been achieved in the diagnosis and management of cystic fibrosis (CF) over the last few years. Highly effective CFTR modulators have become available, “adding tomorrows” to the lives of people with CF. Patients who are not eligible for modulators may benefit from the development of new CFTR function restoring therapies. Newborn screening and early diagnosis of the disease in the newborn period remains an area of intensive research. Lung infections and inflammation, CF-related diabetes, nutrition, hepatic and gastrointestinal disorders, and lung transplantation for advanced lung disease are additional exciting areas of CF research. The purpose of this Special Issue is to accommodate review articles describing the path to a cure for every individual with CF. We hope that contributors will cover as many topics as possible, reviewing the extensive body of published and ongoing research.



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Special Issue



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Editor-in-Chief

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Message from the Editor-in-Chief

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