Special Issue

Current Updates on Pediatric Cystic Fibrosis Care and Outcomes

Message from the Guest Editors

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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Message from the Editor-in-Chief

You are invited to contribute a research article or comprehensive review for consideration and publication in *Children* (ISSN 2227-9067). *Children* is an open access journal—research articles, reviews, and other content are published online immediately after acceptance. The scientific community and the general public have unlimited free access to the content as soon as it is published. The journal focuses on sharing clinical, epidemiological, and translational science relevant to children's health. We would be pleased to welcome you as one of our authors.

Editor-in-Chief

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