

Special Issue

Cystic Fibrosis in Children—Monitoring, Complications and New Therapies

Message from the Guest Editors

Cystic fibrosis remains a challenging disease, whether in terms of diagnosis and treatment points of view, or regarding its multifaceted complications. Besides lung disease, the condition that dictates the disease outcome, more other comorbidities like liver disease and diabetes arise, complicating disease management, especially in children. With the new use of modulators, the life expectancy is likely to upsurge. However, but further studies would be necessary in order to provide the best care for CF children. With increased life expectancy, more complications occur and several decisions must be taken in clinical practice. This Issue addresses the cystic fibrosis diagnosis, monitoring and management of disease's complications like lung disease exacerbations, and their relation to other non-respiratory complications like liver disease, diabetes, bone disease, endocrine dysfunctions and new treatments.

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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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