Special Issue

Cystic Fibrosis: Diagnosis and Treatment

Message from the Guest Editors

You know better than us that cystic fibrosis is an important genetic disease caused by a myriad of CFTR gene mutations and that the sweat chloride test, genetic testing, prenatal/preconception testing, and newborn screening are different ways to obtain a diagnosis. In recent years, novel drugs have dramatically changed the symptoms and, therefore, the lives of many cystic fibrosis patients who are allowed to take them. Moreover, the median age of treated patients is reducing over time thanks to new legislations. What's more, a lot of patients without F508del will hope to be allowed to take these new medications. In this Special Issue, we welcome original articles, reviews, brief reports about the diagnosis, treatment, and management of cystic fibrosis.

Guest Editors

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There has been an explosion of gene and target based research and therapeutics in the multitude of fields that compose clinical medicine. The *Journal of Clinical Medicine*'s (*JCM*) staff and editorial board are dedicated to providing cutting edge, timely, and peer-reviewed articles covering the diverse subspecialties of clinical medicine. The journal publishes concise, innovative, and exciting research articles as well as clinically significant articles and reviews that are pertinent to the myriad of disciplines within medicine. The articles published are relevant to both primary care physicians and specialists. The journal's full-texts are archived in PubMed Central and indexed in PubMed. Please consider submitting your manuscripts for publication to our journal and check us out on-line!

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