Special Issue

Nutrition and Cystic Fibrosis in Children

Message from the Guest Editor

Cystic Fibrosis (CF) is an autosomal dominant genetic disorder that impacts approximately 105,000 persons in 94 countries throughout the world. The classic presentation of CF remains that of malnutrition, lung disease, and pancreatic insufficiency. Malnutrition was one of the early targets of therapy and focused on a high-fat, high-calorie diet based on a pivotal study in the 1970s comparing the nutritional management practices of two centers in Toronto and Boston. Since then, many treatments have been developed to decrease morbidity and mortality in CF including CFTR modulators which aid in folding and stabilizing the CFTR protein. New highly effective modulator therapy (HEMT) has shown a significant improvement in weight gain due to an improvement in whole-body inflammation, decreased resting energy expenditure, and improvements in malabsorption. The impact of HEMT along with the recognition of more than 2000 mutations with variable clinical presentations is creating new challenges for clinicians. This Special Issue aims to collect manuscripts focusing on the change in the traditional approach to nutritional therapy, unveiling the importance for personalized medicine in CF.

Guest Editor

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About the Journal

Message from the Editorial Board

Nutrients is an on-line open access journal that was first published in 2009. *Nutrients* adheres to rigorous peerreview and editorial processes and publishes only high quality manuscripts that address important issues related to the impacts of nutrients on human health. The Impact Factor of *Nutrients* has risen rapidly since its establishment and it is now ranked in the first quartile of journals publishing in the field of nutrition and dietetics research.

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