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Cardiomyopathy: Clinical Diagnosis and Treatment: Part II

Guest Editor:

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Message from the Guest Editor

Dear Colleagues,

It is my pleasure to invite you to contribute to the Special Issue entitled "Cardiomyopathy: Clinical Diagnosis and Treatment: Part II". This is a follow-up volume to the previous one, where we published six papers. For more details, please visit:

https://www.mdpi.com/journal/jcm/special_issues/Clinical_Tako

Cardiomyopathy entails a broad group of diseases, acquired or genetic, which result in a similar phenotype. Furthermore, cardiomyopathy is a clinically heterogeneous disease with large differences depending on gender, age of onset and rate of progression, which are thought to be explained by a complex interplay of genetic susceptibility and environmental factors. Because of the wide variety of conditions that can lead to cardiomyopathy, a systematic approach is needed to facilitate the identification and management of specific cardiomyopathies. The diagnosis, management and follow-up of patients with cardiomyopathy is a multifactorial process. In this Special Issue, we would like to present the latest findings on cardiomyopathy from various viewpoints, and share this information with readers.

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