



Advancements in Diagnosis and Management of Hypertrophic Cardiomyopathy and Amyloidosis

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Message from the Guest Editor

Dear Colleagues,

This Special Issue delves into the latest developments in the diagnosis and management of two significant cardiovascular disorders: hypertrophic cardiomyopathy (HCM) and amyloidosis. HCM is characterized by abnormal thickening of the heart muscle, while amyloidosis involves the accumulation of amyloid proteins in various organs, including the heart. The 100-word summary highlights the emerging diagnostic techniques, therapeutic interventions, and prognostic factors for these conditions. Key topics include genetic testing, imaging modalities, risk stratification, novel treatment options, and the impact of early detection on patient outcomes. The aim is to provide clinicians and researchers with a comprehensive understanding of HCM and amyloidosis to enhance patient care and improve long-term prognosis.

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