



Advancements in Diagnosis and Management of Hypertrophic Cardiomyopathy and Amyloidosis

Guest Editor:

Dr. Ahmed Abuzaid

Department of Medicine, Division
of Cardiovascular Medicine,
Jefferson University
Hospital/Christiana Care Health
System, Newark, DE 19713, USA

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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.

Dr. Ahmed Abuzaid

Guest Editor





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