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Diagnosis and Management of Cardiac Amyloidosis

Guest Editor:

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Deadline for manuscript submissions:
closed (31 December 2021)

Message from the Guest Editor

Cardiac amyloidosis has undergone major changes in recent years: first with the development of non-invasive diagnostic techniques that have led to an increase in diagnoses and, more recently, the development and improvement of treatments, particularly in the field of transthyretin amyloidosis. These developments have led the entire cardiology community to take an interest in this pathology and to set up collaborations between specialists to improve diagnosis and management. Cardiac amyloidosis leads us to reconsider cardiology from a new perspective, with a pathology that differs in all respects from other cardiomyopathies and requires dedicated knowledge and multidisciplinary management. For all these reasons, it seemed necessary to create a Special Issue on this topic. With this Special Issue, we hope to encourage submissions that discuss the current state of the art, address ongoing knowledge gaps, and focus on ongoing controversies and management related to cardiac amyloidosis.

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