Introduction

Atrial brillation (AF) is a common cardiac arrhythmia characterized by irregular, o en rapid heartbeats. It has been a subject of extensive research due to its association with an increased risk of stroke, heart failure, and other cardiovascular complications. While traditional risk factors for AF, such as hypertension, diabetes, and obesity, have been well-established, emerging evidence suggests that oral in ammatory diseases may also play a signi cant role in its development and progression. is comprehensive review aims to explore the connection between oral in ammatory diseases and AF, shedding light on the potential mechanisms and clinical implications of this association [1].

In addition, C-reactive protein, interleukin-6, tumor necrosis factor- , and other in ammatory factors can cause an abnormal

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cardiovascular system [5].

In ammation and atrial brillation

In ammation has long been recognized as a crucial player in the development and exacerbation of cardiovascular diseases. In the context of AF, it is believed that the systemic in ammation triggered by oral diseases can a ect the myocardium and conduction system, leading to electrical and structural remodeling. is, in turn, can promote the initiation and maintenance of AF [6].

Mechanisms of in uence

Several potential mechanisms underlie the relationship between oral in ammatory diseases and AF. ese mechanisms include:

In ammatory mediators: e release of in ammatory mediators like interleukin-6 (IL-6), C-reactive protein (CRP), and tumor necrosis factor-alpha (TNF-) from oral in ammation may promote atrial in ammation and brosis, contributing to the development of AF [7].

Endothelial dysfunction: Oral in ammatory diseases can impair endothelial function, leading to increased oxidative stress and reduced nitric oxide bioavailability [8]. is endothelial dysfunction may contribute to AF by a ecting the endothelium of the atria.

Microbiota dysbiosis: An altered oral microbiota composition may lead to the translocation of oral pathogens into the bloodstream, potentially causing systemic in ammation and impacting cardiac health [9].

Autonomic nervous system activation: Local in ammation in the oral cavity can activate the sympathetic nervous system, which is known to in uence atrial electrophysiology and increase the likelihood of AF.

Clinical Implications

Understanding the link between oral in ammatory diseases and AF has clinical implications. Dentists and cardiologists should collaborate more closely to identify patients at risk and develop integrated care plans. For patients with AF, maintaining good oral health may

be an adjunctive strategy to reduce the risk of AF recurrence and complications [10].

Conclusion

e connection between oral in ammatory diseases and atrial brillation is an emerging eld of research that holds promise for improving our understanding of the multifactorial nature of AF. While