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Sommario/riassunto

Until recently, the cellular basis for sudden death, the Brugada Syndrome, has largely remained an unknown to modern arrhythmologists and cardiologists, particularly in the absence of any structural heart disease. Detailed observations of age-groups, especially the young, families and populations where sudden death frequently occurs, and improved understanding of its contributory factors and mechanisms are, however, showing the way forward. This addition to the Clinical Approaches to Tachyarrhythmias (CATA) Series, written by the investigators who discovered and probed the Brugada Syn