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Research progress and novel approaches in the treatment of malignant arrhythmias with potassium channel modulators**Xu Huixing***People's Hospital of Changshou Chongqing, China*

Objective: This study aims to explore the research progress in the treatment of malignant arrhythmias using potassium channel modulators. Malignant arrhythmias, including Long QT Syndrome (LQTS) and Short QT Syndrome (SQTS), pose severe cardiac conditions that may lead to sudden death. Potassium channels play a crucial role in cardiac electrophysiology, and their dysfunction is closely associated with various arrhythmias.

Methods and Materials: Based on the latest research literature and clinical trial data, this review analyzes the mechanisms of action, clinical advancements, and potential benefits of potassium channel modulators in treating arrhythmias. Special attention is given to the relationship between potassium channel dysfunction and arrhythmias, as well as the development of novel potassium channel modulators.

Results: Potassium Channel Modulators as Novel Agents: At concentrations of 1-100 μ mol/L, quinidine enhances IK1 current, hyperpolarizes resting membrane potential, and shortens action potential duration. For Long QT Syndrome, openers of *IKs* and *IKr* channels are considered promising. For instance, ML277, a selective activator of Kv7.1, enhances *IKs* current, aiding in shortening the action potential duration in induced pluripotent stem cell-derived cardiomyocytes. Polyunsaturated fatty acids (pufas) and their analogs also emerge as modulators for the Kv7.1/KCNE1 channel, shortening action potential duration and suppressing early after depolarization's (eads). The clinical effectiveness and safety of these potassium channel openers need further confirmation through clinical trials. For example, PUFA analogs effectively shorten the cardiac action potential in human-induced pluripotent stem cell-derived cardiomyocytes, but their specificity and safety concerning other cardiac ion channels require further investigation.

Conclusion: Potassium channel modulators demonstrate significant potential in the treatment of malignant arrhythmias. Dysfunction of potassium channels is a crucial contributor to arrhythmias, and potassium channel openers like ML277 and pufas effectively regulate cardiac electrophysiology, improving arrhythmia symptoms. These novel drugs show particular promise in treating Long QT Syndrome. However, their clinical application requires further research to establish their safety and efficacy. Future studies should focus on the specificity, safety, and effectiveness of these drugs in different types of arrhythmia patients. With the ongoing development of new drugs and treatment strategies, potassium channel modulators may become essential tools in treating malignant arrhythmias.

Biography

Xu Huixing, female, born on April 25, 1981, bachelor's degree, the main research direction of drug management, with many years of experience in clinical pharmacy. Dr. Xu Huixing is a leading researcher and clinician at the People's Hospital of Changshou Chongqing, China, specializing in cardiovascular medicine with a particular focus on malignant arrhythmias.