

Arrhythmias Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

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Abstracts

Arrhythmias Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Arrhythmias - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides an overview of the Arrhythmias (Cardiovascular) pipeline landscape.

An arrhythmia is a problem with the rate or rhythm of the heartbeat. During an arrhythmia, the heart can beat too fast, too slow, or with an irregular rhythm. Signs and symptoms include anxiety, weakness, dizziness, and light-headedness, fainting or nearly fainting, sweating, shortness of breath and chest pain. Treatment includes antiarrhythmics and anticoagulants.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Arrhythmias - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides comprehensive information on the therapeutics under development for Arrhythmias (Cardiovascular), complete with analysis by stage of development, drug target, mechanism of action (MoA), route of administration (RoA) and molecule type. The guide covers the descriptive pharmacological action of the therapeutics, its complete research and development history and latest news and press



releases.

The Arrhythmias (Cardiovascular) pipeline guide also reviews of key players involved in therapeutic development for Arrhythmias and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies /Universities /Institutes, the molecules developed by Companies in Phase II, Phase I, Preclinical and Discovery stages are 1, 2, 10 and 7 respectively. Similarly, the Universities portfolio in Preclinical and Discovery stages comprises 5 and 4 molecules, respectively.

Arrhythmias (Cardiovascular) pipeline guide helps in identifying and tracking emerging players in the market and their portfolios, enhances decision making capabilities and helps to create effective counter strategies to gain competitive advantage. The guide is built using data and information sourced from Global Markets Direct's proprietary databases, company/university websites, clinical trial registries, conferences, SEC filings, investor presentations and featured press releases from company/university sites and industry-specific third party sources. Additionally, various dynamic tracking processes ensure that the most recent developments are captured on a real time basis.

Note: Certain content / sections in the pipeline guide may be removed or altered based on the availability and relevance of data.

SCOPE

The pipeline guide provides a snapshot of the global therapeutic landscape of Arrhythmias (Cardiovascular).

The pipeline guide reviews pipeline therapeutics for Arrhythmias (Cardiovascular) by companies and universities/research institutes based on information derived from company and industry-specific sources.

The pipeline guide covers pipeline products based on several stages of development ranging from pre-registration till discovery and undisclosed stages.

The pipeline guide features descriptive drug profiles for the pipeline products which comprise, product description, descriptive licensing and collaboration details, R&D brief, MoA & other developmental activities.

The pipeline guide reviews key companies involved in Arrhythmias (Cardiovascular) therapeutics and enlists all their major and minor projects.



The pipeline guide evaluates Arrhythmias (Cardiovascular) therapeutics based on mechanism of action (MoA), drug target, route of administration (RoA) and molecule type.

The pipeline guide encapsulates all the dormant and discontinued pipeline projects.

The pipeline guide reviews latest news related to pipeline therapeutics for Arrhythmias (Cardiovascular)

REASONS TO BUY

Procure strategically important competitor information, analysis, and insights to formulate effective R&D strategies.

Recognize emerging players with potentially strong product portfolio and create effective counter-strategies to gain competitive advantage.

Find and recognize significant and varied types of therapeutics under development for Arrhythmias (Cardiovascular).

Classify potential new clients or partners in the target demographic.

Develop tactical initiatives by understanding the focus areas of leading companies.

Plan mergers and acquisitions meritoriously by identifying key players and it's most promising pipeline therapeutics.

Formulate corrective measures for pipeline projects by understanding Arrhythmias (Cardiovascular) pipeline depth and focus of Indication therapeutics.

Develop and design in-licensing and out-licensing strategies by identifying prospective partners with the most attractive projects to enhance and expand business potential and scope.



Adjust the therapeutic portfolio by recognizing discontinued projects and understand from the know-how what drove them from pipeline.



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Aetas Pharma Co Ltd

ARMGO Pharma Inc.

Cardurion Pharmaceuticals LLC

Cynata Therapeutics Ltd

Espero BioPharma Inc

Galectin Therapeutics Inc

Gilead Sciences Inc

Jiangsu Kangyuan Pharmaceutical Co Ltd

LATITUDE Pharmaceuticals Inc.

Les Laboratoires Servier SAS

LQT Therapeutics Inc

Nissan Chemical Corp

Orion Corp

SignPath Pharma Inc

Stablix Inc

Vera Therapeutics Inc

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Gene Therapy to Activate KCNQ1 for Arrhythmias - Drug Profile

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Product Description

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M-201 - Drug Profile

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ORM-10103 - Drug Profile

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Featured News & Press Releases

Jan 06, 2022: LQTT announces award by the European Joint Program for rare diseases

for SGK1 inhibition as a novel therapeutic approach in LQTS

Oct 29, 2020: AnaBios and Orion publish scientific article describing novel cardiac

compound

Sep 27, 2017: U.S. Patent Granted for Cynata Cymerus Technology

Sep 05, 2017: Cynata to Present at the 19th Annual Rodman & Renshaw Global

Investment Conference in New York

Apr 19, 2017: Cynata Advances to Pre-IND Meeting with US FDA

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