

Pulmonary Embolism - Pipeline Review, H1 2020

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Abstracts

Pulmonary Embolism - Pipeline Review, H1 2020

SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Pulmonary Embolism - Pipeline Review, H1 2020, provides an overview of the Pulmonary Embolism (Cardiovascular) pipeline landscape.

Pulmonary embolism is the sudden blockage of a major blood vessel (artery) in the lung, usually by a blood clot. Symptoms include shortness of breath, chest pain, cough, leg pain or swelling, excessive sweating, rapid or irregular heartbeat and dizziness. Predisposing factors include High blood pressure and cardiovascular disease. Treatment includes anticoagulants and thrombolytics.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Pulmonary Embolism - Pipeline Review, H1 2020, provides comprehensive information on the therapeutics under development for Pulmonary Embolism (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 Pulmonary Embolism (Cardiovascular) pipeline guide also reviews of key players involved in therapeutic development for Pulmonary Embolism and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies/Universities/Institutes, the molecules developed by Companies in Pre-

Registration, Phase III, Phase II, Phase I, Preclinical and Discovery stages are 2, 1, 2, 4, 2 and 1 respectively.

Pulmonary Embolism (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 Pulmonary Embolism (Cardiovascular).

The pipeline guide reviews pipeline therapeutics for Pulmonary Embolism (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 Pulmonary Embolism (Cardiovascular) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Pulmonary Embolism (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 Pulmonary Embolism (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 Pulmonary Embolism (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 Pulmonary Embolism (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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Pulmonary Embolism - Companies Involved in Therapeutics Development

Acticor Biotech SAS

China Resources Emde Biological Pharmaceutical Co Ltd

Daiichi Sankyo Co Ltd

F. Hoffmann-La Roche Ltd

Indiana Lysis Technologies LLC

Les Laboratoires Servier SAS

Tasly Pharmaceutical Group Co Ltd

Tianjin Pharmaceuticals Group Co Ltd

Valeo Pharma Inc

Verseon Corp

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Featured News & Press Releases

Nov 14, 2019: Valeo Pharma announces filing of a New Drug Submission for Low Molecular Weight Heparin in Canada

Oct 25, 2017: IU-based drug-discovery startup aims to use nanotechnology to digest blood clots in lungs

Oct 11, 2017: IU-based drug-discovery startup wins \$10,000 in BioCrossroads New Venture Competition

Feb 23, 2017: Daiichi Sankyo Europe Receives CHMP Positive Opinion for Roteas (edoxaban)

Nov 28, 2016: Research to Be Presented at ASH Annual Meeting Shows Verseon's New Class of Anticoagulants Prevents Thrombosis While Preserving Platelet Function

Nov 09, 2016: Once-Daily Anticoagulant LIXIANA (edoxaban) Approved in Canada for Stroke Prevention in Atrial Fibrillation and for the Treatment and Prevention of Recurrent Deep-Vein Thrombosis and Pulmonary Embolism

Oct 13, 2016: Daiichi Sankyo Announces New Data to be Presented at the ISPOR EU Congress 2016 Analysing the Safety, Efficacy and Cost-effectiveness of LIXIANA (edoxaban)

Sep 01, 2016: Lixiana applies to National Health Insurance

Aug 26, 2015: UK's NICE recommends once-daily LIXIANA (edoxaban) for the treatment and prevention of recurrent deep vein thrombosis and pulmonary embolism in adults

Aug 07, 2015: NICE Recommends Once-daily Lixiana (Edoxaban) for Preventing Stroke and Systemic Embolism in People with Non-valvular Atrial Fibrillation

Jun 25, 2015: Daiichi Sankyo's Once-Daily LIXIANA (edoxaban) Approved in the EU for Stroke Prevention in Nonvalvular Atrial Fibrillation and for the Treatment and Prevention of Recurrent DVT and PE

Oct 28, 2014: U.S. FDA Cardiovascular and Renal Drugs Advisory Committee Makes Recommendation on Daiichi Sankyo's Once-Daily SAVAYSA for the Reduction in Risk of Stroke and Systemic Embolic Events in Patients with Non-Valvular Atrial Fibrillation

Dec 09, 2013: Phase 3 Data Show Daiichi Sankyo's Once-Daily Edoxaban Lowered Incidence of VTE Recurrence and Clinically Relevant Bleeding Compared to Warfarin in a Large Subgroup of Patients with Cancer

Sep 01, 2013: Daiichi Sankyo's Once-Daily Edoxaban Shows Comparable Efficacy and Superiority for the Principal Safety Endpoint Compared to Warfarin in a Phase 3 Study for the Treatment of Symptomatic VTE and Prevention of its Recurrence

Oct 25, 2012: Daiichi Sankyo Completes Enrolment In Hokusai-VTE Phase III Study Of Edoxaban For Treatment And Prevention Of Recurrence Of VTE

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Acticor Biotech SAS

China Resources Emde Biological Pharmaceutical Co Ltd

Daiichi Sankyo Co Ltd

F. Hoffmann-La Roche Ltd

Indiana Lysis Technologies LLC

Les Laboratoires Servier SAS

Tasly Pharmaceutical Group Co Ltd

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