

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

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Abstracts

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SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Paroxysmal Nocturnal Hemoglobinuria – Drugs In Development, 2022, provides an overview of the Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) pipeline landscape.

Paroxysmal nocturnal hemoglobinuria is a rare disease in which red blood cells break down earlier than normal. Symptoms include abdominal pain, back pain, dark urine, easy bruising or bleeding, headache and shortness of breath. The predisposing factors include age, obesity and hormone therapy. Treatment includes surgery, chemotherapy and radiation therapy.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Paroxysmal Nocturnal Hemoglobinuria – Drugs In Development, 2022, provides comprehensive information on the therapeutics under development for Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders), 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 Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) pipeline guide also reviews of key players involved in therapeutic development for Paroxysmal Nocturnal Hemoglobinuria and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies /Universities /Institutes, the molecules developed by Companies in Phase III, Phase II, Phase I, IND/CTA Filed, Preclinical and Discovery stages are 10, 6, 6, 1, 15 and 1 respectively.

Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) 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 Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders).

The pipeline guide reviews pipeline therapeutics for Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) 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 Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) 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 Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders)

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 Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders).

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 Paroxysmal Nocturnal Hemoglobinuria (Hematological Disorders) 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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Featured News & Press Releases

Jun 12, 2022: Samsung Bioepis presents new phase 3 Study of SB12 (Eculizumab), a proposed biosimilar to Soliris, at the European Hematology Association (EHA) Congress 2022

Jun 10, 2022: CANbridge CAN106 phase 1 data presented at the European Hematology Association 2022 Congress

Jun 10, 2022: Apellis and Sobi report new data reinforcing the robust efficacy and safety profile of EMPAVELI (pegcetacoplan) for PNH at EHA 2022 Congress

May 12, 2022: Apellis announces seven abstracts in PNH to be presented at the European Hematology Association Congress

May 12, 2022: CANbridge to present CAN106 phase 1 data at the European Hematology Association 2022 Congress

Apr 08, 2022: BioCryst pauses enrollment in BCX9930 clinical trials

Mar 28, 2022: CANbridge doses first subject in Phase Ib/II PNH treatment trial in China

Feb 07, 2022: CANbridge reports positive top-line CAN106 Phase 1 data

Feb 04, 2022: Novartis Japan developing Iptacopan as priority product, initially targeting 3 indications

Jan 07, 2022: BioCryst begins patient enrollment in REDEEM-1 pivotal trial evaluating BCX9930 as oral monotherapy for patients with PNH

Dec 15, 2021: Apellis and Sobi announce EU approval of Aspaveli (pegcetacoplan) for treatment of PNH

Dec 13, 2021: Apellis and Sobi Report Empaveli (pegcetacoplan) demonstrated sustained normalization of clinical measures in a broad PNH patient population

Dec 11, 2021: Samsung Bioepis' SB12 Soliris (Eculizumab) Biosimilar Demonstrates PK, PD Bioequivalence in phase 1 study

Nov 29, 2021: BioCryst begins patient enrollment in REDEEM-2 pivotal trial evaluating BCX9930 as oral monotherapy for patients with PNH

Nov 04, 2021: Apellis to present new data reinforcing EMPAVELI (pegcetacoplan) efficacy and safety in patients with PNH at the 2021 ASH Annual Meeting

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