

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

<https://marketpublishers.com/r/P6F4547246C1EN.html>

Date: April 2022

Pages: 107

Price: US\$ 2,000.00 (Single User License)

ID: P6F4547246C1EN

Abstracts

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SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Polycythemia Vera - Drugs In Development, 2022, provides an overview of the Polycythemia Vera (Oncology) pipeline landscape.

Polycythemia Vera (PV) is a stem cell disorder characterized as a panhyperplastic, malignant and neoplastic marrow disorder. Symptoms include itchiness, headache, dizziness, weakness and excessive sweating. The risk of polycythemia vera increases with age. Treatment includes alkylating agents.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Polycythemia Vera - Drugs In Development, 2022, provides comprehensive information on the therapeutics under development for Polycythemia Vera (Oncology), 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 Polycythemia Vera (Oncology) pipeline guide also reviews of key players involved

in therapeutic development for Polycythemia Vera 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 and Preclinical stages are 1, 2, 8, 4 and 4 respectively. Similarly, the Universities portfolio in Phase II stages comprises 1 molecules, respectively.

Polycythemia Vera (Oncology) 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 Polycythemia Vera (Oncology).

The pipeline guide reviews pipeline therapeutics for Polycythemia Vera (Oncology) 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 Polycythemia Vera (Oncology) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Polycythemia Vera (Oncology) 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 Polycythemia Vera (Oncology)

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 Polycythemia Vera (Oncology).

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 Polycythemia Vera (Oncology) 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

Mar 02, 2022: NCCN clinical practice guidelines in oncology update recommends BESREMi (ropeginterferon alfa-2b-njft) for the treatment of polycythemia vera

Feb 15, 2022: German Federal Court of Justice upholds AOP's license rights in ropeginterferon and PEC's liability for damages

Dec 14, 2021: PharmaEssentia USA selects Veeva Data Cloud to support launch of novel therapy for rare, chronic blood cancer

Dec 12, 2021: Protagonist Therapeutics presents updated phase 2 rusfertide data in Polycythemia Vera (PV) at ASH 2021 Annual Meeting

Dec 06, 2021: BESREMi (ropeginterferon alfa-2b-njft), FDA approved for treatment of polycythemia vera, available at Biologics by McKesson

Nov 18, 2021: BESREMi (ropeginterferon alfa-2b-njft) now approved for the treatment of adults with polycythemia vera

Nov 15, 2021: BESREMi (Ropeginterferon alfa-2b) approved by the US FDA

Nov 12, 2021: U.S. FDA approves BESREMi (ropeginterferon alfa-2b-njft) as the only interferon for adults with polycythemia vera

Nov 04, 2021: Protagonist Therapeutics announces updated data from phase 2 study of Rusfertide in polycythemia vera selected for oral presentations at the ASH 2021 Annual

Meeting

Oct 11, 2021: FDA lifts hold on Protagonist's rusfertide clinical studies

Sep 20, 2021: FDA puts Protagonist Therapeutics' rusfertide on clinical hold

Sep 14, 2021: PPMX-T003: Announcement on initiation of phase I clinical trial in polycythemia vera patients

Aug 10, 2021: Protagonist Therapeutics announces resolution of contract dispute with Zealand Pharma

Jun 11, 2021: Protagonist Therapeutics announces updated phase 2 data supporting long-term efficacy of Rusfertide in polycythemia vera

Jun 07, 2021: Protagonist Therapeutics to host Investor Conference Call and webcast to discuss updated Phase 2 Rusfertide results in Polycythemia Vera as presented at EHA 2021

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