Prader-Willi Syndrome (PWS) - Pipeline Review, H1 2016

Description: Prader-Willi Syndrome (PWS) - Pipeline Review, H1 2016

Summary

The report ‘Prader-Willi Syndrome (PWS) - Pipeline Review, H1 2016’, provides an overview of the Prader-Willi Syndrome (PWS) pipeline landscape.

The report provides comprehensive information on the therapeutics under development for Prader-Willi Syndrome (PWS), complete with analysis by stage of development, drug target, mechanism of action (MoA), route of administration (RoA) and molecule type. The report also covers the descriptive pharmacological action of the therapeutics, its complete research and development history and latest news and press releases. Additionally, the report provides an overview of key players involved in therapeutic development for Prader-Willi Syndrome (PWS) and features dormant and discontinued projects.

The report features investigational drugs from across globe covering over 20 therapy areas and nearly 3,000 indications. The report is built using data and information sourced from 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. Drug profiles featured in the report undergoes periodic review following a stringent set of processes to ensure that all the profiles are updated with the latest set of information. Additionally, various dynamic tracking processes ensure that the most recent developments are captured on a real time basis.

The report 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.

Note*: Certain sections in the report may be removed or altered based on the availability and relevance of data.

Scope

- The report provides a snapshot of the global therapeutic landscape of Prader-Willi Syndrome (PWS)
- The report reviews pipeline therapeutics for Prader-Willi Syndrome (PWS) by companies and universities/research institutes based on information derived from company and industry-specific sources
- The report covers pipeline products based on various stages of development ranging from pre-registration till discovery and undisclosed stages
- The report features descriptive drug profiles for the pipeline products which includes, product description, descriptive MoA, R&D brief, licensing and collaboration details & other developmental activities
- The report reviews key players involved Prader-Willi Syndrome (PWS) therapeutics and enlists all their major and minor projects
- The report assesses Prader-Willi Syndrome (PWS) therapeutics based on drug target, mechanism of action (MoA), route of administration (RoA) and molecule type
- The report summarizes all the dormant and discontinued pipeline projects
- The report reviews latest news related to pipeline therapeutics for Prader-Willi Syndrome (PWS)

Reasons to buy

- Gain strategically significant competitor information, analysis, and insights to formulate effective R&D strategies
- Identify emerging players with potentially strong product portfolio and create effective counter-strategies to gain competitive advantage
- Identify and understand important and diverse types of therapeutics under development for Prader-Willi Syndrome (PWS)
- Identify potential new clients or partners in the target demographic
- Develop strategic initiatives by understanding the focus areas of leading companies
- Plan mergers and acquisitions effectively by identifying key players and it's most promising pipeline therapeutics
- Devise corrective measures for pipeline projects by understanding Prader-Willi Syndrome (PWS) 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
- Modify the therapeutic portfolio by identifying discontinued projects and understanding the factors that drove them from pipeline

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Featured News & Press Releases

Jan 20, 2016: Zafgen's Pivotal Phase 3 Trial of Beloranib in Prader-Willi Syndrome Achieves Co-Primary Efficacy Endpoints

Jan 07, 2016: Rhythm Receives Orphan Drug Designation for Setmelanotide for the Treatment of Prader-Willi Syndrome

Dec 02, 2015: Zafgen Announces Beloranib IND Placed on Complete Clinical Hold

Dec 02, 2015: Zafgen Provides Clinical Update on Beloranib

Oct 22, 2015: Zafgen Announces Beloranib Program Update

Oct 14, 2015: Zafgen Issues Statement

Sep 25, 2015: Essentialis Presents Results From Clinical Study PC025 Evaluating DCCR in the Treatment of Prader-Willi Syndrome at Annual Meeting of the Foundation for Prader-Willi Research

Sep 24, 2015: FDA Office of Orphan Products Development Awards Rhythm $1 Million Grant to Support Phase 2 Study of Setmelanotide in Prader-Willi Syndrome

Sep 09, 2015: Essentialis Announces the Dosing of the First Prader-Willi Syndrome Patients in a Long-Term
Extension to Clinical Trial PC025

Jun 04, 2015: Rhythm Initiates Two Phase 2 Clinical Trials of Setmelanotide in Rare Genetic Disorders of Obesity Caused by MC4 Pathway Deficiencies

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