

Prader-Willi Syndrome (PWS) Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

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Abstracts

Prader-Willi Syndrome (PWS) 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 Prader-Willi Syndrome (PWS) - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides an overview of the Prader-Willi Syndrome (PWS) (Genetic Disorders) pipeline landscape.

Prader-Willi Syndrome (PWS) is a rare genetic disorder present at birth that results in a number of physical, mental and behavioral problems. PWS is due to lack of several genes on one of an individual's two chromosome 15's. In the majority of cases, there is a deletion. In remaining cases, the entire chromosome from the father is missing and there are instead two chromosome 15's from the mother (uniparental disomy). Symptoms include poor muscle tone, low levels of sex hormones and a constant feeling of hunger.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Prader-Willi Syndrome (PWS) - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides comprehensive information on the therapeutics under development for Prader-Willi Syndrome (PWS) (Genetic 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 Prader-Willi Syndrome (PWS) (Genetic Disorders) pipeline guide also reviews of key players involved in therapeutic development for Prader-Willi Syndrome (PWS) and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies /Universities /Institutes, the molecules developed by Companies in Filing rejected/Withdrawn, Phase III, Phase II, Phase I, Preclinical, Discovery and Unknown stages are 1, 2, 6, 6, 6, 3 and 1 respectively. Similarly, the Universities portfolio in Phase II, Preclinical and Discovery stages comprises 1, 1 and 1 molecules, respectively.

Prader-Willi Syndrome (PWS) (Genetic 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 Prader-Willi Syndrome (PWS) (Genetic Disorders).

The pipeline guide reviews pipeline therapeutics for Prader-Willi Syndrome (PWS) (Genetic 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 Prader-Willi Syndrome (PWS) (Genetic Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Prader-Willi Syndrome (PWS) (Genetic 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 Prader-Willi Syndrome (PWS) (Genetic 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 Prader-Willi Syndrome (PWS) (Genetic 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 Prader-Willi Syndrome (PWS) (Genetic 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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Beryl Therapeutics Inc

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Bioprojet SCR

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ConSynance Therapeutics Inc

Ferring International Center SA

GLWL Research Inc

Helsinn Group

Inversago Pharma Inc

Lipidio Pharmaceuticals Inc

Lumos Pharma Inc

Neuracle Science Co Ltd

Neuren Pharmaceuticals Ltd

Notitia Biotechnologies Co

OptiNose Inc

OT4B

Saniona AB

Soleno Therapeutics Inc



Taysha Gene Therapies Inc

Tonix Pharmaceuticals Holding Corp

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Featured News & Press Releases

Mar 09, 2022: Running for research - Prader-Willi Syndrome to fund multi-center study of DCCR in early phase Prader-Willi syndrome

Mar 03, 2022: Tonix Pharmaceuticals announces FDA Orphan-Drug Designation for TNX-2900 for the treatment of Prader-Willi Syndrome

Feb 28, 2022: Tonix Pharmaceuticals announces research agreement with the French National Institute of Health and Medical Research (Inserm) to study the mechanism of oxytocin-mediated improvement of eating behaviors in Prader-Willi Mice

Feb 03, 2022: Aardvark starts enrolment in three Phase II trials of TAS2R agonist



Jan 24, 2022: Soleno Therapeutics provides regulatory update on DCCR for the treatment of Prader-Willi syndrome

Dec 28, 2021: Saniona initiates phase 2b clinical trial of tesomet for Prader-Willi syndrome

Oct 18, 2021: Saniona completes submission of manufacturing data for Tesomet capsules to U.S. FDA

Oct 11, 2021: US patent to 2034 granted for Neuren's NNZ-2591 in autism

Sep 17, 2021: adMare Portfolio Company, Inversago Pharma initiates a phase 1 clinical trial on INV-202, a next generation peripherally-acting CB1 Blocker

Sep 08, 2021: Soleno Therapeutics announces positive data showing continued significant improvements in symptoms of PWS following one year treatment with DCCR Sep 07, 2021: Inversago Pharma initiates a phase 1 clinical trial on INV-202, a next generation peripherally-acting CB1 blocker

Sep 03, 2021: FDA grants Orphan Drug Designation for NNZ-2591 to treat Prader-Willi syndrome

Jul 26, 2021: Saniona receives U.S. FDA orphan drug designation for Tesomet in hypothalamic obesity

Jul 22, 2021: Radius Health announces plans for global Prader-Willi syndrome pivotal study

Jul 06, 2021: Soleno Therapeutics provides regulatory update on DCCR for the treatment of Prader-Willi syndrome

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