

Wilson Disease - Pipeline Review, H2 2019

https://marketpublishers.com/r/W04BE4F835EEN.html

Date: December 2019

Pages: 59

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

ID: W04BE4F835EEN

Abstracts

Wilson Disease - Pipeline Review, H2 2019

SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Wilson Disease - Pipeline Review, H2 2019, provides an overview of the Wilson Disease (Genetic Disorders) pipeline landscape.

Wilson disease is a rare autosomal recessive inherited disorder of copper metabolism that is characterized by excessive deposition of copper in the liver, brain, and other tissues. Symptoms include abdominal pain, jaundice, problems with speech, swallowing and muscle stiffness. Treatment includes chelators and Vitamin E supplements.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Wilson Disease - Pipeline Review, H2 2019, provides comprehensive information on the therapeutics under development for Wilson Disease (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 Wilson Disease (Genetic Disorders) pipeline guide also reviews of key players involved in therapeutic development for Wilson Disease and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies/Universities/Institutes, the molecules developed by Companies in Phase III, Preclinical and Discovery stages are 2, 5 and 2 respectively. Similarly, the Universities



portfolio in Preclinical stages comprises 1 molecules, respectively.

Wilson Disease (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 Wilson Disease (Genetic Disorders).

The pipeline guide reviews pipeline therapeutics for Wilson Disease (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 Wilson Disease (Genetic Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Wilson Disease (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 Wilson Disease (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 Wilson Disease (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 Wilson Disease (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.



Contents

Introduction

Global Markets Direct Report Coverage

Wilson Disease - Overview

Wilson Disease - Therapeutics Development

Pipeline Overview

Pipeline by Companies

Pipeline by Universities/Institutes

Products under Development by Companies

Products under Development by Universities/Institutes

Wilson Disease - Therapeutics Assessment

Assessment by Target

Assessment by Mechanism of Action

Assessment by Route of Administration

Assessment by Molecule Type

Wilson Disease - Companies Involved in Therapeutics Development

Alexion Pharmaceuticals Inc

Aligen Therapeutics SL

Deep Genomics Inc

DepYmed Inc

GMP-Orphan SA

Krisani Bio Sciences Pvt Ltd

LambdaGen Therapeutics

Ultragenyx Pharmaceutical Inc

Vivet Therapeutics SAS

Wilson Disease - Drug Profiles

Wilson Disease - Dormant Projects

Wilson Disease - Discontinued Products

Wilson Disease - Product Development Milestones

Appendix



List Of Tables

LIST OF TABLES

Number of Products under Development for Wilson Disease, H2 2019

Number of Products under Development by Companies, H2 2019

Number of Products under Development by Universities/Institutes, H2 2019

Products under Development by Companies, H2 2019

Products under Development by Universities/Institutes, H2 2019

Number of Products by Stage and Target, H2 2019

Number of Products by Stage and Mechanism of Action, H2 2019

Number of Products by Stage and Route of Administration, H2 2019

Number of Products by Stage and Molecule Type, H2 2019

Wilson Disease - Pipeline by Alexion Pharmaceuticals Inc, H2 2019

Wilson Disease - Pipeline by Aligen Therapeutics SL, H2 2019

Wilson Disease - Pipeline by Deep Genomics Inc, H2 2019

Wilson Disease - Pipeline by DepYmed Inc, H2 2019

Wilson Disease - Pipeline by GMP-Orphan SA, H2 2019

Wilson Disease - Pipeline by Krisani Bio Sciences Pvt Ltd, H2 2019

Wilson Disease - Pipeline by LambdaGen Therapeutics, H2 2019

Wilson Disease - Pipeline by Ultragenyx Pharmaceutical Inc, H2 2019

Wilson Disease - Pipeline by Vivet Therapeutics SAS, H2 2019

Wilson Disease - Dormant Projects, H2 2019



List Of Figures

LIST OF FIGURES

Number of Products under Development for Wilson Disease, H2 2019

Number of Products under Development by Companies, H2 2019

Number of Products by Targets, H2 2019

Number of Products by Stage and Targets, H2 2019

Number of Products by Mechanism of Actions, H2 2019

Number of Products by Stage and Mechanism of Actions, H2 2019

Number of Products by Stage and Routes of Administration, H2 2019

Number of Products by Molecule Types, H2 2019

Number of Products by Stage and Molecule Types, H2 2019



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