

Adrenoleukodystrophy - Pipeline Review, H1 2017

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Abstracts

Adrenoleukodystrophy - Pipeline Review, H1 2017

SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Adrenoleukodystrophy - Pipeline Review, H1 2017, provides an overview of the Adrenoleukodystrophy (Genetic Disorders) pipeline landscape.

Adrenoleukodystrophy is a disorder that occurs most often in males. It mainly affects the nervous system and the adrenal glands, which are small glands located on top of each kidney. Abetalipoproteinemia is also a kind of Adrenoleukodystrophy (Adrenomyeloneuropathy/Schilder-Addison Complex) Bassen-Kornzweig syndrome is a rare disease passed down through families in which a person is unable to fully absorb dietary fats through the intestines. The disease is caused by a defect in gene that tells the body to create lipoproteins. It is sympotomized by curvature of spine, muscle weakness, decrease of vision over time. Treatment involves large doses of vitamin supplements containing fat-soluble vitamins.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Adrenoleukodystrophy - Pipeline Review, H1 2017, provides comprehensive information on the therapeutics under development for Adrenoleukodystrophy (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 Adrenoleukodystrophy (Genetic Disorders) pipeline guide also reviews of key players involved in therapeutic development for Adrenoleukodystrophy (Adrenomyeloneuropathy/Schilder-Addison Complex) 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 I, IND/CTA Filed, Preclinical and Discovery stages are 2, 1, 1, 7 and 1 respectively. Similarly, the Universities portfolio in Phase II and Phase I stages comprises 1 and 1 molecules, respectively.

Adrenoleukodystrophy (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 Adrenoleukodystrophy (Genetic Disorders).

The pipeline guide reviews pipeline therapeutics for Adrenoleukodystrophy (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 Adrenoleukodystrophy



(Genetic Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Adrenoleukodystrophy (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 Adrenoleukodystrophy (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 Adrenoleukodystrophy (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 Adrenoleukodystrophy (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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VK-0214 - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

VK-2809 - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

Adrenoleukodystrophy (Adrenomyeloneuropathy/ Schilder-Addison Complex) - Dormant Projects

Adrenoleukodystrophy (Adrenomyeloneuropathy/ Schilder-Addison Complex) - Product Development Milestones

Featured News & Press Releases

Mar 21, 2017: Minoryx Therapeutics successfully completes phase 1 clinical trial for lead candidate MIN-102

Feb 22, 2017: Minoryx Therapeutics receives Orphan Drug Designation from the US FDA for its lead candidate MIN-102

Jan 18, 2017: Minoryx Therapeutics further strengthens its Scientific Advisory Board (SAB)

Dec 14, 2016: Minoryx Therapeutics receives European Orphan Drug Designation for its lead candidate MIN-102

Dec 06, 2016: Viking Therapeutics Receives Orphan Drug Designation for VK0214 from the U.S. Food and Drug Administration for the Treatment of X-Linked Adrenoleukodystrophy

Sep 26, 2016: Viking Therapeutics Presents Positive Proof-of-Concept Data for VK0214 in In Vivo Model of X-Linked Adrenoleukodystrophy at the 86th Annual Meeting of the American Thyroid Association

Sep 14, 2016: Positive Preclinical Data With MD1003 (High Dose Pharmaceutical Grade Biotin) in X-Linked Adrenoleukodystrophy to be Presented at European Committee for Treatment and Research in Multiple Sclerosis 2016

Jul 26, 2016: Viking Therapeutics Announces Positive Top-Line Results from Proof-of-Concept Study of VK0214 in In Vivo Model of X-Linked Adrenoleukodystrophy (X-ALD) Jul 06, 2016: Minoryx Therapeutics Initiates Phase 1 clinical trial of MIN-102

Jun 14, 2016: Minoryx Therapeutics appoints Dr. Uwe Meya as Chief Medical Officer

Apr 03, 2014: Minoryx receives Orphan Drug Designation for MIN-101

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COMPANIES MENTIONED

bluebird bio Inc MedDay SA Minoryx Therapeutics sl Pfizer Inc ReceptoPharm Inc Viking Therapeutics Inc



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