

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

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Abstracts

Encephalopathy 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 Encephalopathy - Drugs In Development, 2022, provides an overview of the Encephalopathy (Central Nervous System) pipeline landscape.

Encephalopathy is a term for any diffuse disease of the brain that alters brain function or structure. Symptoms include seizures, difficulty speaking or swallowing, trembling, involuntary twitching and muscle weakness. Causes include traumas or injuries, genetic, liver disease, hypoxic condition, Lyme disease and organ failure.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Encephalopathy - Drugs In Development, 2022, provides comprehensive information on the therapeutics under development for Encephalopathy (Central Nervous System), 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 Encephalopathy (Central Nervous System) pipeline guide also reviews of key

players involved in therapeutic development for Encephalopathy and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies/Universities/Institutes, the molecules developed by Companies in Phase II, Phase I, Preclinical and Discovery stages are 4, 5, 10 and 2 respectively. Similarly, the Universities portfolio in Preclinical stages comprises 3 molecules, respectively.

Encephalopathy (Central Nervous System) 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 Encephalopathy (Central Nervous System).

The pipeline guide reviews pipeline therapeutics for Encephalopathy (Central Nervous System) 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 Encephalopathy (Central Nervous System) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Encephalopathy (Central Nervous System)

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 Encephalopathy (Central Nervous System)

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 Encephalopathy (Central Nervous System).

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 Encephalopathy (Central Nervous System) 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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Product Description

Mechanism Of Action

ANL-302 - Drug Profile

Product Description

Mechanism Of Action

ARG-007 - Drug Profile

Product Description

Mechanism Of Action

BIO-018 - Drug Profile

Product Description

Mechanism Of Action

BM-1021 - Drug Profile

Product Description

Mechanism Of Action

cannabidiol - Drug Profile

Product Description

Mechanism Of Action

Cell Therapy for Autoimmune Disorders, Cardiovascular Disorders, CNS Disorders, Infectious Disease, Musculoskeletal Disorders and Oncology - Drug Profile

Product Description

Mechanism Of Action

CL-2020 - Drug Profile

Product Description

Mechanism Of Action

diazoxide - Drug Profile

Product Description

Mechanism Of Action

Drugs to Target SLC2A1 for Glut1 Deficiency Syndrome - Drug Profile

Product Description

Mechanism Of Action

estetrol - Drug Profile

Product Description

Mechanism Of Action

Gene Therapy to Activate SLC2A1 for GLUT1 Deficiency Syndrome - Drug Profile

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Product Description

Mechanism Of Action

MCC-950 - Drug Profile

Product Description

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melatonin - Drug Profile

Product Description

Mechanism Of Action

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Product Description

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Proteins for Hypoxic-Ischemic Brain Injury - Drug Profile

Product Description

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remestemcel-L - Drug Profile

Product Description

Mechanism Of Action

RLS-0071 - Drug Profile

Product Description

Mechanism Of Action

sovateltide - Drug Profile

Product Description

Mechanism Of Action

Stem Cell Therapy for Anemia and Hypoxic Ischemic Encephalopathy - Drug Profile

Product Description

Mechanism Of Action

Stem Cell Therapy for Encephalopathy and Periventricular Leukomalacia - Drug Profile

Product Description

Mechanism Of Action

THDG-3 - Drug Profile

Product Description

Mechanism Of Action

UDI-001 - Drug Profile

Product Description

Mechanism Of Action

Encephalopathy - Dormant Projects

Encephalopathy - Discontinued Products

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Featured News & Press Releases

Oct 04, 2022: Argenica receives \$1.38m R&D Tax Incentive cash refund for FY22

Sep 29, 2022: Further positive preclinical results for arg-007 in hypoxic-ischaemic encephalopathy

Aug 29, 2022: Pharmazz announces Indian Central Drugs Standard Control Organization (CDSCO) Clearance of IND to conduct a Phase II clinical trial of sovateltide (PMZ-1620) in hypoxic-ischemic encephalopathy in neonates

Jun 23, 2022: USA FDA grants Fast Track Designation to ReAlta Life Sciences' lead compound RLS-0071 for the treatment of Hypoxic-Ischemic Encephalopathy in neonates

Jun 02, 2022: ReAlta doses first subjects in Phase Ib severe asthma therapy trial

May 10, 2022: Germany's BfArM approves ReAlta Life Science's phase 1b inhaled LPS challenge trial to support development of RLS-0071 in severe asthma

Apr 27, 2022: ReAlta Life Sciences partners with Hope for HIE to improve neonatal brain injury outcomes and quality of life

Apr 04, 2022: ReAlta Life Sciences' RLS-0071 demonstrates excellent safety profile and confirmed target engagement in first-in-human phase 1 clinical trial

Mar 31, 2022: ReAlta gets FDA clearance for Phase II HIE therapy trial in neonates

Mar 04, 2022: ReAlta Life Sciences announces poster presentation at the American Academy of Neurology 2022 Annual Meeting

Feb 03, 2022: Pharmazz submits Investigational New Drug Application to India Central Drugs Standard Control Organization for a phase II clinical trial of sovateltide (PMZ-1620) in hypoxic-ischemic encephalopathy in neonates

Dec 09, 2021: Argenica to be granted patent protection in the United States, one of the world's largest healthcare markets

Nov 16, 2021: MPO data presented at ACR 2021

Nov 03, 2021: Argenica Therapeutics: Positive preclinical data for ARG-007 neuroprotection in hypoxicischaemic encephalopathy

May 18, 2021: ReAlta Life Sciences expands phase 1 clinical trial of RLS-0071 to include three multiple-ascending dose cohorts and an additional single ascending dose cohort

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