

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

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Abstracts

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SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Becker Muscular Dystrophy - Drugs In Development, 2022, provides an overview of the Becker Muscular Dystrophy (Genetic Disorders) pipeline landscape.

Becker muscular dystrophy (BMD) is one of type of muscular dystrophy, a group of genetic, degenerative diseases primarily affecting voluntary muscles. Symptoms include pain and sensation, difficulty with running, hopping, and jumping, toe walking, breathing problems, cognitive problems and frequent falls.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Becker Muscular Dystrophy - Drugs In Development, 2022, provides comprehensive information on the therapeutics under development for Becker Muscular Dystrophy (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 Becker Muscular Dystrophy (Genetic Disorders) pipeline guide also reviews of key



players involved in therapeutic development for Becker Muscular Dystrophy and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies/Universities/Institutes, the molecules developed by Companies in Phase I, Phase I and Preclinical stages are 3, 5 and 4 respectively.

Becker Muscular Dystrophy (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 Becker Muscular Dystrophy (Genetic Disorders).

The pipeline guide reviews pipeline therapeutics for Becker Muscular Dystrophy (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 Becker Muscular Dystrophy (Genetic Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Becker Muscular Dystrophy (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 Becker Muscular Dystrophy (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 Becker Muscular Dystrophy (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 Becker Muscular Dystrophy (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

Becker Muscular Dystrophy - Overview

Becker Muscular Dystrophy - Therapeutics Development

Pipeline Overview

Pipeline by Companies

Products under Development by Companies

Becker Muscular Dystrophy - Therapeutics Assessment

Assessment by Target

Assessment by Mechanism of Action

Assessment by Route of Administration

Assessment by Molecule Type

Becker Muscular Dystrophy - Companies Involved in Therapeutics Development

ARMGO Pharma Inc

Edgewise Therapeutics Inc

Epirium Bio Inc

Italfarmaco SpA

Milo Biotechnology LLC

PhaseBio Pharmaceuticals Inc

Santhera Pharmaceuticals Holding AG

Sarcomed AB

Scholar Rock Inc

Stealth BioTherapeutics Corp

Tivorsan Pharmaceuticals Inc

Becker Muscular Dystrophy - Drug Profiles

apitegromab - Drug Profile

Product Description

Mechanism Of Action

History of Events

ARM-210 - Drug Profile

Product Description

Mechanism Of Action

History of Events

dextro epicatechin - Drug Profile

Product Description

Mechanism Of Action

History of Events



EDG-5506 - Drug Profile

Product Description

Mechanism Of Action

History of Events

elamipretide hydrochloride - Drug Profile

Product Description

Mechanism Of Action

History of Events

epicatechin - Drug Profile

Product Description

Mechanism Of Action

History of Events

Gene therapy To Activate Follistatin For Duchenne Muscular Dystrophy,

Facioscapulohumeral Muscular Dystrophy, Becker Muscular Dystrophy And Inclusion

Body Myositis - Drug Profile

Product Description

Mechanism Of Action

History of Events

givinostat - Drug Profile

Product Description

Mechanism Of Action

History of Events

nandrolone decanoate - Drug Profile

Product Description

Mechanism Of Action

PB-1023 - Drug Profile

Product Description

Mechanism Of Action

History of Events

TVN-102 - Drug Profile

Product Description

Mechanism Of Action

History of Events

vamorolone - Drug Profile

Product Description

Mechanism Of Action

History of Events

Becker Muscular Dystrophy - Dormant Projects

Becker Muscular Dystrophy - Discontinued Products



Becker Muscular Dystrophy - Product Development Milestones Featured News & Press Releases

Apr 14, 2022: Edgewise-Funded natural history trial of Becker Muscular Dystrophy (BMD) now enrolling

Mar 08, 2022: Edgewise Therapeutics to present on EDG-5506 for Becker and Duchenne Muscular Dystrophy (BMD, DMD) at the 2022 Annual MDA Clinical and Scientific Conference

Jan 05, 2022: Edgewise Therapeutics announces positive topline results from the EDG-5506 phase 1b clinical trial in adults with Becker Muscular Dystrophy (BMD)

Dec 15, 2021: Edgewise Therapeutics to report phase 1b topline results of EDG-5506 in individuals with Becker muscular dystrophy (BMD) on January 5, 2022, and initiates ARCH follow-on open label BMD study

Oct 28, 2021: Edgewise Therapeutics announces positive topline results from the EDG-5506 phase 1 multiple ascending dose (MAD) study in healthy volunteers (HVs) and doses first Becker Muscular Dystrophy (BMD) patients

Sep 27, 2021: ReveraGen and Santhera announce FDA orphan grant funding for clinical trial with vamorolone in Becker muscular dystrophy

Aug 16, 2021: Edgewise receives U.S. FDA Fast Track Designation for EDG-5506 for the treatment of individuals with Becker muscular dystrophy (BMD)

Jun 26, 2021: Italfarmaco announces topline data from phase 2 trial with Givinostat in patients with Becker Muscular Dystrophy

May 04, 2021: Edgewise Therapeutics announces publication of data demonstrating the elevation of fast but not slow skeletal muscle fiber injury biomarkers in the circulation of patients with Becker and Duchenne Muscular Dystrophy

Feb 22, 2021: Italfarmaco provides update on ongoing clinical programs with givinostat in oral presentation at XVIII International Conference on Duchenne and Becker Muscular Dystrophy

Nov 11, 2020: Edgewise Therapeutics initiates phase 1 clinical trial of EDG-5506 in Becker muscular dystrophy

Oct 29, 2020: Edgewise Therapeutics appoints John Moore as general counsel Aug 12, 2020: Epirium Bio announces commencement of phase 1 clinical trial of EPM-01 in Becker muscular dystrophy

Apr 14, 2020: Epirium Bio receives FDA orphan-drug designation for EB 002 ((+)-Epicatechin) for the treatment of duchenne and becker muscular dystrophy Jun 30, 2015: Cardero Therapeutics Presents at 2015 Parent Project Muscular Dystrophy Connect Conference

Appendix Methodolo

Methodology

Coverage



Secondary Research
Primary Research
Expert Panel Validation
Contact Us
Disclaimer



List Of Tables

LIST OF TABLES

Number of Products under Development for Becker Muscular Dystrophy, 2022

Number of Products under Development by Companies, 2022

Products under Development by Companies, 2022

Number of Products by Stage and Target, 2022

Number of Products by Stage and Mechanism of Action, 2022

Number of Products by Stage and Route of Administration, 2022

Number of Products by Stage and Molecule Type, 2022

Becker Muscular Dystrophy - Pipeline by ARMGO Pharma Inc, 2022

Becker Muscular Dystrophy - Pipeline by Edgewise Therapeutics Inc, 2022

Becker Muscular Dystrophy - Pipeline by Epirium Bio Inc, 2022

Becker Muscular Dystrophy - Pipeline by Italfarmaco SpA, 2022

Becker Muscular Dystrophy - Pipeline by Milo Biotechnology LLC, 2022

Becker Muscular Dystrophy - Pipeline by PhaseBio Pharmaceuticals Inc, 2022

Becker Muscular Dystrophy - Pipeline by Santhera Pharmaceuticals Holding AG, 2022

Becker Muscular Dystrophy - Pipeline by Sarcomed AB, 2022

Becker Muscular Dystrophy - Pipeline by Scholar Rock Inc, 2022

Becker Muscular Dystrophy - Pipeline by Stealth BioTherapeutics Corp, 2022

Becker Muscular Dystrophy - Pipeline by Tivorsan Pharmaceuticals Inc, 2022

Becker Muscular Dystrophy - Dormant Projects, 2022

Becker Muscular Dystrophy - Discontinued Products, 2022



List Of Figures

LIST OF FIGURES

Number of Products under Development for Becker Muscular Dystrophy, 2022

Number of Products under Development by Companies, 2022

Number of Products by Top 10 Targets, 2022

Number of Products by Stage and Top 10 Targets, 2022

Number of Products by Mechanism of Actions, 2022

Number of Products by Stage and Mechanism of Actions, 2022

Number of Products by Routes of Administration, 2022

Number of Products by Stage and Routes of Administration, 2022

Number of Products by Molecule Types, 2022

Number of Products by Stage and Molecule Types, 2022



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