

# Inclusion Body Myositis (IBM) Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

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## **Abstracts**

Inclusion Body Myositis (IBM) 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 Inclusion Body Myositis (IBM) - Drugs In Development, 2022, provides an overview of the Inclusion Body Myositis (IBM) (Musculoskeletal Disorders) pipeline landscape.

Inclusion Body Myositis (IBM) is one of the inflammatory myopathies that involve inflammation of the muscles or associated tissues, such as the blood vessels that supply the muscles. IBM symptoms include progressive weakness of the muscles of the wrists and fingers, the muscles of the front of the thigh, and the muscles that lift the front of the foot.

## REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Inclusion Body Myositis (IBM) - Drugs In Development, 2022, provides comprehensive information on the therapeutics under development for Inclusion Body Myositis (IBM) (Musculoskeletal 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 Inclusion Body Myositis (IBM) (Musculoskeletal Disorders) pipeline guide also reviews of key players involved in therapeutic development for Inclusion Body Myositis (IBM) 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 II, Phase I, Preclinical and Discovery stages are 2, 1, 2, 3 and 2 respectively.

Inclusion Body Myositis (IBM) (Musculoskeletal 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 Inclusion Body Myositis (IBM) (Musculoskeletal Disorders).

The pipeline guide reviews pipeline therapeutics for Inclusion Body Myositis (IBM) (Musculoskeletal 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 Inclusion Body Myositis (IBM) (Musculoskeletal Disorders) therapeutics and enlists all their major and



minor projects.

The pipeline guide evaluates Inclusion Body Myositis (IBM) (Musculoskeletal 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 Inclusion Body Myositis (IBM) (Musculoskeletal 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 Inclusion Body Myositis (IBM) (Musculoskeletal 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 Inclusion Body Myositis (IBM) (Musculoskeletal 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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Featured News & Press Releases

Nov 05, 2021: Abcuro to present late-breaking abstract at American College of Rheumatology Convergence 2021 of ABC008 inclusion body myositis phase 1 data demonstrating proof of mechanism for selective depletion of highly cytotoxic T cells Oct 06, 2021: Abcuro to present trial in progress poster on a phase 1 clinical trial of ABC008 demonstrating targeted depletion of large granular lymphocytes at the 2021 Leukemia, Lymphoma and Myeloma Congress

Jun 30, 2021: Abcuro and ImaginAb share initial results of study using novel technology for imaging T cell infiltration of skeletal muscle in patients with inclusion body myositis

Apr 01, 2021: Nobel Pharma announces notice of designation of orphan drug

Aug 22, 2017: Ultragenyx Announces Top-Line Results from Phase 3 Study of Ace-ER in GNE Myopathy

Nov 11, 2016: Ultragenyx Announces Withdrawal of Marketing Authorization Application for Aceneuramic Acid Prolonged Release (Ace-ER) in the European Union

Jul 27, 2016: Ultragenyx Announces Completion of Enrollment in Phase 3 Study of Aceneuramic Acid Extended Release (Ace-ER) in GNE Myopathy

Oct 02, 2015: Ultragenyx Announces Aceneuramic Acid Prolonged Release Marketing Authorization Application Filed and Accepted for Review by European Medicines Agency

May 28, 2015: Ultragenyx Announces First Patient Enrolled in Global Phase 3 Study of



Aceneuramic Acid (Sialic Acid) Extended Release in GNE Myopathy
Jan 12, 2015: Ultragenyx Announces Intent to File for Conditional Approval in Europe
for Sialic Acid Extended-Release Tablets in Hereditary Inclusion Body Myopathy
Oct 13, 2014: Ultragenyx Announces Interim Data From Phase 2 Extension Study of
Sialic Acid Extended-Release at International Congress of the World Muscle Society
Apr 30, 2014: Ultragenyx Announces Positive Data From Phase 2 Study of Sialic Acid
Extended-Release at Emerging Sciences Session of American Academy of Neurology
Annual Meeting

Dec 20, 2013: Ultragenyx Announces Results from Phase 2 Study of Sialic Acid Extended-Release Treatment in Hereditary Inclusion Body Myopathy Sep 26, 2013: Ultragenyx Announces Three Abstracts Accepted for Poster Presentation at 18th Annual World Muscle Society Congress

Jul 03, 2013: Ultragenyx Announces a Positive Signal in Interim Data from Phase 2 Study of UX001 in Hereditary Inclusion Body Myopathy

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