

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

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Abstracts

Sarcopenia 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 Sarcopenia - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides an overview of the Sarcopenia (Musculoskeletal Disorders) pipeline landscape.

Sarcopenia is an aging-associated condition which is characterized by loss of muscle mass, strength and function. Symptoms include weakness, loss of endurance and poor balance. Causes of sarcopenia include a reduction in the nerve cells, low hormone levels and decline in body's ability to convert protein to energy. Treatment includes hormone replacement therapy (HRT), exercise and dietary supplements.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Sarcopenia - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides comprehensive information on the therapeutics under development for Sarcopenia (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 Sarcopenia (Musculoskeletal Disorders) pipeline guide also reviews of key players involved in therapeutic development for Sarcopenia 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, IND/CTA Filed, Preclinical, Discovery and Unknown stages are 4, 3, 1, 21, 9 and 2 respectively. Similarly, the Universities portfolio in Preclinical and Discovery stages comprises 2 and 1 molecules, respectively.

Sarcopenia (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 Sarcopenia (Musculoskeletal Disorders).

The pipeline guide reviews pipeline therapeutics for Sarcopenia (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 Sarcopenia (Musculoskeletal Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Sarcopenia (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 Sarcopenia (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 Sarcopenia (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 Sarcopenia (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

Feb 25, 2022: Oncocross begins clinical trial for AI-developed drug for sarcopenia

Jan 05, 2022: MyMD Pharmaceuticals announces issuance of new U.S. patent covering MYMD-1 in a method of treating sarcopenia

Dec 16, 2021: Biophytis to meet with FDA to advance Sarconeos (BIO101) development in sarcopenia from phase 2 to phase

Oct 06, 2021: Biophytis to host virtual KOL event on Sarconeos (BIO101) on its lead projects in COVID-19 and Sarcopenia

Oct 04, 2021: Biophytis announces promising full results from the SARA-INT Phase 2b Trial of Sarconeos (BIO101) in Sarcopenia at the 11th Annual International Conference on Frailty and Sarcopenia Research (ICFSR) September 29 - October 2, 2021

Sep 29, 2021: Biophytis to present full results from the SARA-INT Phase 2b trial of Sarconeos (BIO101) in Sarcopenia at the 11th Annual International Conference on Frailty and Sarcopenia Research (ICFSR) September 29 - October 2, 2021

Aug 02, 2021: Biophytis announces top line results of SARA-INT phase 2 study with Sarconeos (BIO101) in sarcopenia

Jun 30, 2021: Results of Biophytis SARA-INT phase 2 trial with Sarconeos (BIO101) in sarcopenia will be released by August 2021

Dec 16, 2020: Biophytis - Last patient completes final visit in SARA-INT, a Phase 2 clinical trial evaluating the efficacy of Sarconeos (BIO101) in the treatment of sarcopenia

Dec 14, 2020: Biophytis presents SARA-OBS study results at the annual conference of The Society on Sarcopenia, Cachexia and Wasting Disorders (SCWD)

Aug 28, 2020: Biophytis provides an update on SARA-INT, a phase 2b clinical trial evaluating the efficacy of Sarconeos (BIO101) in the treatment of sarcopenia

Mar 24, 2020: Biophytis completes recruitment of SARA-INT phase 2b study, evaluating the efficacy of Sarconeos (BIO101) in the treatment of sarcopenia

Mar 03, 2020: Biophytis to present the preliminary results of SARA-OBS and its impact on SARA-INT, the phase 2b clinical study, at the 10th International Conference on Frailty and Sarcopenia Research (ICFSR 2020) in Toulouse, France

Feb 11, 2020: Biophytis - Protocol amendment of SARA-INT, a phase 2b clinical trial of Sarconeos (BIO101) in sarcopenia, cleared by FDA and AFMPS

Dec 12, 2019: Biophytis to attend Biotech Showcase 2020 in San Francisco, USA

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