

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

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Abstracts

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) 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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Drugs In Development, 2022, provides an overview of the Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (Musculoskeletal Disorders) pipeline landscape.

Fibrodysplasia ossificans progressiva (FOP) is a disorder in which muscle tissue and connective tissue such as tendons and ligaments are gradually replaced by bone (ossified), forming bone outside that constrains movement. Symptoms swellings in fibrous tissue, pain, fever, muscle ossification and restricted movement of joints. Treatment includes surgery and medications to help relieve symptoms of FOP, such as pain and inflammation.

REPORT HIGHLIGHTS

Global Markets Direct's Pharmaceutical and Healthcare latest pipeline guide Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Drugs In Development, 2022, provides comprehensive information on the therapeutics under



development for Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (Musculoskeletal Disorders) pipeline guide also reviews of key players involved in therapeutic development for Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies/Universities/Institutes, the molecules developed by Companies in Pre-Registration, Phase III, Phase II, Phase I and Preclinical stages are 1, 2, 4, 1 and 2 respectively. Similarly, the Universities portfolio in Preclinical stages comprises 3 molecules, respectively.

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (Musculoskeletal Disorders).

The pipeline guide reviews pipeline therapeutics for Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (Musculoskeletal Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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 Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) (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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Involved in Therapeutics Development

AstraZeneca Plc

Clementia Pharmaceuticals Inc

Daiichi Sankyo Co Ltd

Incyte Corp

Ipsen SA

La Jolla Pharmaceutical Company

Pfizer Inc

Regeneron Pharmaceuticals Inc

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Development Milestones

Featured News & Press Releases

Oct 25, 2022: Ipsen statement on updated timeline for palovarotene FDA Advisory



Committee meeting

Jun 29, 2022: Ipsen announces U.S. FDA priority review for palovarotene new drug application in patients with fibrodysplasia ossificans progressiva following resubmission Jan 24, 2022: Health Canada approves Ipsen's Sohonos (palovarotene capsules) as the first approved treatment for Fibrodysplasia Ossificans Progressiva

Aug 13, 2021: Ipsen announces withdrawal of palovarotene NDA, confirming intention to re-submit following additional data analyses

May 28, 2021: Ipsen confirms U.S. FDA accepts new drug application for Palovarotene as the first potential treatment worldwide for fibrodysplasia ossificans progressiva (FOP) Oct 30, 2020: Regeneron provides update on the Garetosmab phase 2 LUMINA-1 trial in fibrodysplasia ossificans progressiva (FOP)

Sep 10, 2020: Ipsen to present new Insights at ASBMR for potential treatment of ultrarare disease Fibrodysplasia Ossificans Progressiva (FOP), including global phase III MOVE Trial results

Aug 25, 2020: Ipsen to present results from MOVE, the first global phase III trial in fibrodysplasia ossificans progressiva (FOP), at ASBMR 2020 annual meeting

Mar 27, 2020: Ipsen to restart dosing of palovarotene in FOP studies

Jan 27, 2020: Ipsen temporarily stops palovarotene dosing in FOP trials

Jan 10, 2020: Regeneron reports positive data from rare bone disease trials

Jan 09, 2020: Regeneron announces encouraging garetosmab phase 2 results in patients with ultra-rare debilitating bone disease

Dec 09, 2019: Ipsen initiates partial clinical hold for palovarotene IND120181 and IND135403 studies

Mar 27, 2019: Blueprint Medicines provides update on clinical progress of BLU-782

Feb 11, 2019: Clementia granted Rare Pediatric Disease Designation by FDA for Palovarotene for Fibrodysplasia Ossificans Progressiva

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