

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Pipeline Review, H1 2020

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Abstracts

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Pipeline Review, H1 2020

SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Fibrodysplasia Ossificans Progressiva - Pipeline Review, H1 2020, provides an overview of the Fibrodysplasia 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 - Pipeline Review, H1 2020, provides comprehensive information on the therapeutics under development for Fibrodysplasia 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 (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 Phase III, Phase II, Phase I and Preclinical stages are 1, 3, 4 and 4 respectively. Similarly, the Universities portfolio in Preclinical and Discovery stages comprises 2 and 1 molecules, respectively.

Fibrodysplasia 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 (Musculoskeletal Disorders).

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

The pipeline guide evaluates Fibrodysplasia 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 (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 (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 (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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Clementia Pharmaceuticals Inc

Daiichi Sankyo Co Ltd

Incyte Corp

Keros Therapeutics Inc

La Jolla Pharmaceutical Company

Pfizer Inc

Regeneron Pharmaceuticals Inc

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Drug Profiles

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R&D Progress

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Featured News & Press Releases

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

Nov 04, 2019: BioCryst launches trial to treat fibrodysplasia ossificans progressiva

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

Oct 01, 2018: Blueprint Medicines presents foundational preclinical data supporting the development of BLU-782, a highly selective ALK2 Inhibitor, for the treatment of patients with fibrodysplasia ossificans progressiva

Sep 24, 2018: Blueprint Medicines to present data on BLU-782 at 2018 ASBMR annual meeting

Jan 05, 2018: BioCryst Advancing Potential Treatment for Rare and Severely Debilitating Fibrodysplasia Ossificans Progressiva

Jan 05, 2018: BioCryst Provides Update on Drug Candidate BCX9499

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AstraZeneca Plc

BioCryst Pharmaceuticals Inc

Clementia Pharmaceuticals Inc

Daiichi Sankyo Co Ltd

Incyte Corp

Keros Therapeutics Inc

La Jolla Pharmaceutical Company

Pfizer Inc

Regeneron Pharmaceuticals Inc

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