

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Pipeline Review, H2 2019

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Abstracts

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Pipeline Review, H2 2019

SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Fibrodysplasia Ossificans Progressiva - Pipeline Review, H2 2019, 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, H2 2019, 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, Preclinical and Discovery stages are 1, 2, 4, 5 and 1 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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Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Therapeutics

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Involved in Therapeutics Development

AstraZeneca Plc

BioCryst Pharmaceuticals Inc

Clementia Pharmaceuticals Inc

Daiichi Sankyo Co Ltd

Ipsen SA

Keros Therapeutics Inc

La Jolla Pharmaceutical Company

Oncodesign SA

Pfizer Inc

Regeneron Pharmaceuticals Inc

Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva) - Drug Profiles

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Featured News & Press Releases

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

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

Oct 24, 2019: Oncodesign: publication of an article presenting promising results for the treatment of stone man syndrome (FOP)

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

Oct 23, 2018: Clementia announces plan to submit a new drug application for Palovarotene for the treatment of FOP based on positive phase 2 results

Oct 02, 2018: Clementia initiates phase 1 clinical trial of palovarotene eye drop formulation in healthy volunteers

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 26, 2018: Clementia announces updated phase 2 part B data on Palovarotene for FOP

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

Sep 05, 2018: Clementia to Present at ASBMR 2018 and Upcoming Investor Conferences

Aug 17, 2018: Clementia completes patient enrolment in phase 3 of MOVE Trial

May 23, 2018: Clementia Reports Positive Phase 2 Part B Data Showing Treatment with Palovarotene Significantly Reduces New Bone Growth in Patients with FOP

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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COMPANIES MENTIONED

AstraZeneca Plc

BioCryst Pharmaceuticals Inc

Clementia Pharmaceuticals Inc

Daiichi Sankyo Co Ltd

Ipsen SA

Keros Therapeutics Inc

La Jolla Pharmaceutical Company

Oncodesign SA

Pfizer Inc

Regeneron Pharmaceuticals Inc

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