

Uterine Leiomyoma (Uterine Fibroids) Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

<https://marketpublishers.com/r/UB2788459019EN.html>

Date: March 2022

Pages: 91

Price: US\$ 2,000.00 (Single User License)

ID: UB2788459019EN

Abstracts

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SUMMARY

Global Markets Direct's latest Pharmaceutical and Healthcare disease pipeline guide Uterine Leiomyoma - Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update, provides an overview of the Uterine Leiomyoma (Non Malignant Disorders) pipeline landscape.

Uterine fibroids are benign tumors that originate in the uterus (womb). Although they are composed of the same smooth muscle fibers as the uterine wall (myometrium), they are much denser than normal myometrium. Uterine fibroids are usually round. The most common symptoms of uterine fibroids include heavy menstrual bleeding, prolonged menstrual periods seven days or more of menstrual bleeding, pelvic pressure or pain, frequent urination, difficulty emptying bladder, constipation and backache or leg pains.

REPORT HIGHLIGHTS

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

Uterine Leiomyoma (Non Malignant 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 Uterine Leiomyoma (Non Malignant Disorders).

The pipeline guide reviews pipeline therapeutics for Uterine Leiomyoma (Non Malignant 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 Uterine Leiomyoma (Non Malignant Disorders) therapeutics and enlists all their major and minor projects.

The pipeline guide evaluates Uterine Leiomyoma (Non Malignant 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 Uterine Leiomyoma (Non Malignant 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 Uterine Leiomyoma (Non Malignant 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 Uterine Leiomyoma (Non Malignant 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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Development

AbbVie Inc

Bayer AG

Context Therapeutics Inc

Daewoong Pharmaceutical Co Ltd

Endo International Plc

Eurofarma Laboratorios SA

Evestra Inc

GeneScience Pharmaceuticals Co Ltd

Immunitor Inc

Jiangsu Hengrui Medicine Co Ltd

Kissei Pharmaceutical Co Ltd

Livzon Pharmaceutical Group Co Ltd

Luye Pharma Group Ltd

Richter Gedeon Nyrt

TiumBio Co Ltd

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collagenase clostridium histolyticum - Drug Profile

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vilaprisan - Drug Profile

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Featured News & Press Releases

Feb 04, 2022: ObsEva provides update on EU marketing authorisation process for linzagolix, an oral GnRH antagonist, for the treatment of uterine fibroids

Dec 17, 2021: ObsEva Announces Positive CHMP Opinion for Linzagolix, an Oral GnRH Antagonist, for the Treatment of Uterine Fibroids

Dec 10, 2021: ObsEva hosts symposium and presents clinical data on oral GnRH antagonist Linzagolix at SEUD Congress 2021

Dec 02, 2021: ObsEva announces symposium and presentation of clinical data on oral GnRH antagonist Linzagolix at SEUD Congress 2021

Nov 22, 2021: Obseva announces U.S. FDA acceptance of new drug application for Linzagolix

Oct 20, 2021: ObsEva presents clinical data on oral GnRH antagonist Linzagolix for the treatment of uterine fibroids at ASRM 2021 Scientific Congress & Expo

Oct 19, 2021: ObsEva announces presentation of clinical data on oral GnRH antagonist linzagolix at ASRM 2021 Scientific Congress & Expo

Sep 15, 2021: ObsEva announces submission of New Drug Application to U.S. FDA for Linzagolix for the treatment of Uterine Fibroids

Jun 30, 2021: ObsEva presents clinical data on open-label pilot study of Yselty (linzagolix) for the treatment of severe adenomyosis at ESHRE Virtual 37th Annual Meeting

Jun 24, 2021: ObsEva to present data on Yselty (linzagolix) at ESHRE Virtual 37th Annual Meeting

May 20, 2021: ObsEva announces final results from the phase 3 PRIMROSE program of Yselty (linzagolix) for the treatment of uterine fibroids

May 04, 2021: ObsEva announces enrollment completion of linzagolix phase 3 EDELWEISS 3 trial for patients with moderate to severe endometriosis-associated pain

Apr 30, 2021: ObsEva presents posters at the ACOG Annual Clinical and Scientific Virtual Meeting April 30 - May 2, 2021

Apr 27, 2021: ObsEva initiates PRIMROSE 3 bone mineral density follow-up study in PRIMROSE 1 and PRIMROSE 2 trial participants

Feb 18, 2021: Ulipristal acetate 5mg (Esmya): further restrictions due to risk of serious liver injury

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