

Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist -Pipeline Intelligence, 2019

<https://marketpublishers.com/r/C92E941D342EN.html>

Date: December 2018

Pages: 60

Price: US\$ 1,000.00 (Single User License)

ID: C92E941D342EN

Abstracts

This report can be delivered to the clients within 1-2 Business Days.

CmaxInsight's, "Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist-Pipeline Intelligence, 2019", report provides comprehensive insights about pipeline drugs across this mechanism of action (MoA). A key objective of the report is to establish the understanding for all the pipeline drugs that fall under Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist.

Highlights and Scope of the Report

MoA Overview:

This section of the report provides comprehensive coverage of MoA enables the client to understand the landscape of the Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist.

Pipeline Covered:

This section mentions all the promising therapies in different phases of development including the NDA/BLA Filing, Phase III, Phase II, Phase I, Pre-Clinical and the Discovery. The section also details the products which have been dormant or discontinued during the trial stages of development.

Drug Profiles:

The pipeline guide features provide descriptive drug profiles for the pipeline products

which comprise product description, Research and development, and product development activity.

Product Description

It comprises of descriptive drug profiles for the pipeline products in terms of its mechanism of action, mode of administration, molecule type, chemical information, etc.

Research and Development

This section of the report focusing on the clinical and pre-clinical activity which provide detailed information about the efficacy, safety and tolerability of pipeline drugs. A snapshot on the clinical trial of a pipeline therapy includes information about sponsor, stage of development, trial design, enrollment number, location, study start and primary completion date, and dosage frequency and formulation of the drug.

Product Development Activity

This section of the report provides detail information about licensing and collaboration, funding & financing, designation, patent, technology, key milestones and other developmental activities.

Therapeutic Assessment

The report comprises of comparative pipeline therapeutics assessment by stage of development, therapy type, indication / therapy area, molecule type, and route of administration across this MoA.

Company Profile

Company profile includes the detail about type of company, headquarter, global presence, research focus and key financial

Methodology

The report is built using data and information sourced from proprietary databases,

primary and secondary research and in-house analysis by CmaxInsight's team of industry experts.

Secondary sources information and data has been collected from various printable and non-printable sources like search engines, News websites, Government Websites, Trade Journals, White papers, Magazines, Trade associations, Books, Industry Portals, Industry Associations and access to available databases.

Reasons to buy

Establish a comprehensive understanding of key competitor information, analysis, and insights to improve R&D strategies

Identify emerging players with potentially strong product portfolio and create effective counter-strategies to gain competitive advantage

Discover in licensing and out licensing strategies by identifying potential partners with progressing projects for Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist to enhance and expand business potential and scope

Plan mergers and acquisitions successfully by identifying major players with the most promising pipeline therapeutics in the target demographic

Our extensive in-depth analysis on therapy portfolio support the client in decision-making process regarding their therapeutic portfolio by identifying the reason behind the dormant or discontinued drugs

Develop strategic initiatives by understanding the focus areas of leading companies

Assess challenges and opportunities that influence Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist R&D

Please note: Certain sections in the report may be removed or altered based on the availability and relevance of data for the indicated mechanism of action.

Contents

1. REPORT INTRODUCTION

Executive Summary

Key Findings

2. CALCITONIN GENE-RELATED PEPTIDE RECEPTOR (CRLR) ANTAGONIST - OVERVIEW

3. PIPELINE THERAPEUTICS

An Overview of Pipeline Products for Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist

4. COMPARATIVE ANALYSIS OF PIPELINE THERAPEUTICS

5. PRODUCTS IN LAST / MID / EARLY CLINICAL STAGE DEVELOPMENT

5.1 Drug Name: Company Name

Product Description

Research and Development (Clinical and Pre-clinical activity)

Product Development Activities

Key Development Milestone

6. PRODUCTS IN PRE-CLINICAL STAGE DEVELOPMENT

6.1 Drug Name: Company Name

Product Description

Research and Development (Pre-clinical activity)

Product Development Activities

Key Development Milestone

7. PRODUCTS IN DISCOVERY STAGE DEVELOPMENT

7.1 Drug Name: Company Name

Product Description

Research and Development (In-vitro activity)

Product Development Activities

Key Development Milestone

8. THERAPEUTIC ASSESSMENT OF PIPELINE ACTIVE PRODUCTS

Pipeline Assessment by Route of Administration

Pipeline Assessment by Stage and Route of Administration

Pipeline Assessment by Molecule Type

Pipeline Assessment by Stage and Molecule Type

Pipeline Assessment by Indication / Therapy Area

9. DORMANT / DISCONTINUED PRODUCTS

9.1 Drug Name: Company Name

Product Description

Research and Development

Product Development Activities

Reason for Dormancy / Discontinuation

10. COMPANY PROFILING

Appendix

Report Methodology

Disclaimer

About CmaxInsight

Note: Certain sections of the table of contents would vary according to the availability of information

List Of Tables

LIST OF TABLES

Table 1: Total Pipeline Products for Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist

Table 2: Number of Products under Development by Companies

Table 3: Late stage Therapeutic Products

Table 4: Mid stage Therapeutic Products

Table 5: Early stage Therapeutic Products

Table 6: Pre-clinical stage Therapeutic Products

Table 7: Discovery stage Therapeutic Products

Table 8: Pipeline Analysis by Route of Administration

List Of Figures

LIST OF FIGURES

Figure 1: Total Pipeline Products for Calcitonin Gene-Related Peptide Receptor (CRLR) Antagonist

Figure 2: Late stage Therapeutic Products

Figure 3: Mid stage Therapeutic Products

Figure 4: Early stage Therapeutic Products

Figure 5: Pre-clinical stage Therapeutic Products

Figure 6: Discovery stage Therapeutic Products

Figure 7: Pipeline Analysis by Route of Administration

Figure 8: Pipeline Analysis by Stage and Route of Administration

Figure 9: Pipeline Analysis by Molecule Type

Figure 10: Pipeline Analysis by Stage and Molecule Type

Figure 11: Pipeline Analysis by Indication/ Therapy Area

Figure 12: Dormant / Discontinued Products

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