

C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

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Abstracts

C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) Drugs in Development by Stages, Target, MoA, RoA, Molecule Type and Key Players, 2022 Update

SUMMARY

According to the recently published report 'C-C Chemokine Receptor Type 5 - Drugs In Development, 2022'; C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) pipeline Target constitutes close to 16 molecules. Out of which approximately 11 molecules are developed by companies and remaining by the universities/institutes.

C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) - C-C chemokine receptor type 5 (CCR5) also known as CD195, is a surface protein located on the plasma membrane of white blood cells and is encoded by CCR5 gene. This receptor binds and responds to a variety of chemokines (CCL3, CCL4, CCL5, CCL3L1). This protein is expressed by T cells and macrophages, and is known to be an important co-receptor for macrophage-tropic virus, including HIV, to enter host cells. It plays a role in granulocyte lineage proliferation and differentiation.

The report 'C-C Chemokine Receptor Type 5 - Drugs In Development, 2022' outlays comprehensive information on the C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) targeted therapeutics, complete with analysis



by indications, stage of development, mechanism of action (MoA), route of administration (RoA) and molecule type; that are being developed by Companies/Universities.

It also reviews key players involved in C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) targeted therapeutics development with respective active and dormant or discontinued projects. Currently, The molecules developed by companies in Filing rejected/Withdrawn, Phase II, Phase I, Preclinical and Discovery stages are 1, 4, 2, 3 and 1 respectively. Similarly, the universities portfolio in Preclinical and Discovery stages comprises 4 and 1 molecules, respectively.

Report covers products from therapy areas Infectious Disease, Central Nervous System, Oncology, Respiratory, Gastrointestinal and Toxicology which include indications Human Immunodeficiency Virus (HIV) Infections (AIDS), Coronavirus Disease 2019 (COVID-19), Multiple Sclerosis, Chronic Obstructive Pulmonary Disease (COPD), Metastatic Colorectal Cancer, Non-Alcoholic Steatohepatitis (NASH), Solid Tumor, Acute Respiratory Distress Syndrome, Alzheimer's Disease, Bladder Cancer, Bone Metastasis, Colon Carcinoma, Colorectal Cancer, Coronavirus Disease 2019 (COVID-19) Pneumonia, Diabetic Neuropathic Pain, Gastric Cancer, Glioblastoma Multiforme (GBM), Hepatocellular Carcinoma, Liver Cancer, Lung Cancer, Melanoma, Metastatic Pancreatic Cancer, Neuropathic Pain (Neuralgia), Non-Small Cell Lung Cancer, Opioid Induced Side Effects, Opium (Opioid) Addiction, Ovarian Cancer, Pancreatic Cancer, Pancreatic Ductal Adenocarcinoma, Parkinson's Disease, Post-Acute Sequelae of COVID 2019 (PASC or Long COVID), Post-Operative Pain, Testicular Cancer, Throat Cancer, Triple-Negative Breast Cancer (TNBC) and Uterine Cancer.

Note: Certain content/sections in the pipeline guide may be removed or altered based on the availability and relevance of data.

SCOPE

The report provides a snapshot of the global therapeutic landscape for C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5)

The report reviews C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) targeted therapeutics under development by companies and universities/research institutes based on



information derived from company and industry-specific sources

The report covers pipeline products based on various stages of development ranging from pre-registration till discovery and undisclosed stages

The report features descriptive drug profiles for the pipeline products which includes, product description, descriptive MoA, R&D brief, licensing and collaboration details & other developmental activities

The report reviews key players involved in C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) targeted therapeutics and enlists all their major and minor projects

The report assesses C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) targeted therapeutics based on mechanism of action (MoA), route of administration (RoA) and molecule type

The report summarizes all the dormant and discontinued pipeline projects

The report reviews latest news and deals related to C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) targeted therapeutics

REASONS TO BUY

Gain strategically significant competitor information, analysis, and insights to formulate effective R&D strategies

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

Identify and understand the targeted therapy areas and indications for C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5)Identify the use of drugs for target identification and drug repurposing

Identify potential new clients or partners in the target demographic

Develop strategic initiatives by understanding the focus areas of leading



companies

Plan mergers and acquisitions effectively by identifying key players and it's most promising pipeline therapeutics

Devise corrective measures for pipeline projects by understanding C-C Chemokine Receptor Type 5 (CHEMR13 or HIV 1 Fusion Coreceptor or CD195 or CCR5) development landscape

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



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Creative Bio-Peptides Inc

Cytodyn Inc

Forest Hills Partners Hong Kong Ltd

GSK plc

Merck & Co Inc

Novartis AG

Orion Biotechnology Canada Ltd

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Featured News & Press Releases

Jul 28, 2022: American Gene Technologies' HIV Cure Clinical Trial Enters Critical Phase: Withdrawing Participants from Antiretrovirals

Jul 19, 2022: Preclinical Data Demonstrate Potential of Creative Bio-Peptides' Multi-Chemokine Receptor Antagonist RAP-103 to Enhance Opioid Analgesia and Inhibit Opioid-derived Dependence, Withdrawal and Respiratory Depression

Jul 14, 2022: Preclinical data demonstrate potential of Creative Bio-Peptides' multichemokine receptor antagonist RAP-103 to provide opioid sparing post-surgical pain relief and treatment for chronic pain

Jul 11, 2022: CytoDyn highlights NIH Grant for HIV functional cure preclinical study of gene therapy based on leronlimab

Apr 12, 2022: CytoDyn announces publication of peer-reviewed paper, "Suppression of human and simian immunodeficiency virus replication with the CCR5-specific antibody leronlimab in two species"

Mar 31, 2022: FDA places full clinical hold on CytoDyn's Covid-19 programme Feb 24, 2022: American Gene Technologies advances closer to demonstrating a



potential cure for HIV

Feb 02, 2022: American Gene Technologies' HIV clinical trial shows blood markers of efficacy in two more patients

Jan 25, 2022: CytoDyn announces leadership transition plan to support regulatory approval and commercialization of Leronlimab

Jan 13, 2022: CytoDyn cancels webcast and live Q/A scheduled for today

Jan 10, 2022: Published paper indicates leronlimab shows activity against 4-class drug resistant HIV-1 from heavily treatment experienced ("HTE") subjects

Jan 05, 2022: Leronlimab 14-week, NASH clinical trial met primary endpoint (PDFF) and secondary endpoint (cT1) for per protocol population in 350 mg weekly dose Dec 22, 2021: FDA provides positive response to begin CytoDyn's Covid-19 antibody

trial

Dec 21, 2021: American Gene Technologies' HIV clinical trial demonstrates blood markers of efficacy

Dec 13, 2021: NASH phase 2 trial open-label portion demonstrates average 80 msec cT1 reduction in 50% of patients and reduction of nearly 50 msec in 80% of patients Appendix

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