

Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) - Pipeline Review, H1 2018

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Abstracts

Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) - Pipeline Review, H1 2018

SUMMARY

According to the recently published report 'Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) - Pipeline Review, H1 2018'; Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) pipeline Target constitutes close to 26 molecules. Out of which approximately 25 molecules are developed by companies and remaining by the universities/institutes.

Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) - Fibroblast growth factor receptor 4 is a protein encoded by the FGFR4 gene. It plays a role in the regulation of cell proliferation, differentiation and migration, and in regulation of lipid metabolism, bile acid biosynthesis, glucose uptake, vitamin D metabolism and phosphate homeostasis. It is required for normal down-regulation of the expression of CYP7A1, the rate-limiting enzyme in bile acid synthesis, in response to FGF19. It phosphorylates PLCG1 and FRS2. Ligand binding leads to the activation of several signaling cascades.

The report 'Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) - Pipeline Review, H1 2018' outlays comprehensive information on the Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) 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 Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) targeted therapeutics development with respective active and dormant or discontinued projects. Currently, The molecules developed by companies in Pre-Registration, Phase III, Phase II, Phase I, IND/CTA Filed, Preclinical and Discovery stages are 1, 3, 6, 5, 1, 7 and 2 respectively. Similarly, the universities portfolio in Preclinical stages comprises 1 molecules, respectively. Report covers products from therapy areas Oncology, Metabolic Disorders, Central Nervous System, Gastrointestinal and Musculoskeletal Disorders which include indications Hepatocellular Carcinoma, Solid Tumor, Bile Duct Cancer (Cholangiocarcinoma), Breast Cancer, Type 2 Diabetes, Lung Adenocarcinoma, Ovarian Cancer, Bladder Cancer, Endometrial Cancer, Esophageal Cancer, Gastric Cancer, Head And Neck Cancer Squamous Cell Carcinoma, Melanoma, Metastatic Transitional (Urothelial) Tract Cancer, Non-Small Cell Lung Cancer, Obesity, Recurrent Glioblastoma Multiforme (GBM), Squamous Non-Small Cell Lung Cancer, Transitional Cell Carcinoma (Urothelial Cell Carcinoma), Achondroplasia, Adenoid Cystic Carcinoma (ACC), Adrenocortical Carcinoma (Adrenal Cortex Cancer), Constipation, Epithelial Ovarian Cancer, Ewing Sarcoma, Fallopian Tube Cancer, Glioblastoma Multiforme (GBM), Head And Neck Cancer, Hematological Tumor, High-Grade Glioma, Kidney Cancer (Renal Cell Cancer), Liver Cancer, Lung Cancer, Lymphoma, Metastatic Biliary Tract Cancer, Nasopharyngeal Cancer, Non-Alcoholic Steatohepatitis (NASH), Osteosarcoma, Peritoneal Cancer, Primary Biliary Cirrhosis, Primary Sclerosing Cholangitis, Primitive Neuroectodermal Tumor (PNET), Prostate Cancer, Refractory Acute Myeloid Leukemia, Relapsed Acute Myeloid Leukemia, Renal Cell Carcinoma, Rhabdomyosarcoma, Small-Cell Lung Cancer, Spinal Cord Injury, Thymic Carcinoma, Thyroid Cancer and Transitional Cell Cancer (Urothelial Cell 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 Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1)

The report reviews Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) 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 Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) targeted therapeutics and enlists all their major and minor projects

The report assesses Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) 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 Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) 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 Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1)

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 Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) 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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ArQule Inc

Bayer AG

Blueprint Medicines Corp

Eddingpharm Inc

Eisai Co Ltd

Eli Lilly and Co

Genosco Inc

H3 Biomedicine Inc

Hanmi Pharmaceuticals Co Ltd

Incyte Corp

Ionis Pharmaceuticals Inc

Johnson & Johnson

Merrimack Pharmaceuticals Inc

NGM Biopharmaceuticals Inc

Novartis AG

Principia Biopharma Inc

Tasly Pharmaceutical Group Co Ltd

Vichem Chemie Research Ltd

Fibroblast Growth Factor Receptor 4 (CD334 or FGFR4 or EC 2.7.10.1) - Drug Profiles**ABSK-011 - Drug Profile**

Product Description

Mechanism Of Action

R&D Progress

BLU-554 - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

BLU-9931 - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

derazantinib - Drug Profile

Product Description

Mechanism Of Action

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Mechanism Of Action

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Mechanism Of Action

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Fusion Protein to Agonize FGFR for Type 2 Diabetes - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

Fusion Proteins to Agonize FGFR for Metabolic Disorders - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

golvatinib tartrate + lenvatinib mesylate - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

H-3B6527 - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

HM-81442 - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

INCB-62079 - Drug Profile

Product Description

Mechanism Of Action

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infigratinib - Drug Profile

Product Description

Mechanism Of Action

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Featured News & Press Releases

Apr 14, 2018: NGM Bio Announces Results From Phase 2 Study Of NGM282 In Nash Patients Demonstrating Clinically Significant Improvements In Liver Histology After 12 Weeks

Apr 13, 2018: NGM Bio Announces Late-Breaking Plenary Presentation At International Liver Conference 2018 Of Detailed Results From Phase 2 Study Of NGM282 In

Patients With Primary Sclerosing Cholangitis

Mar 23, 2018: Anticancer Agent LENVIMA (lenvatinib mesylate) Approved for Additional Indication of Unresectable Hepatocellular Carcinoma (HCC) in Japan, First Approval Worldwide for LENVIMA for HCC

Mar 15, 2018: Janssen Announces U.S. FDA Breakthrough Therapy Designation for Erdafitinib in the Treatment of Metastatic Urothelial Cancer

Mar 15, 2018: ArQule to Present Data on Derazantinib

Mar 12, 2018: Grupo Biotoscana Announces New Initiatives for LENVIMA

Mar 12, 2018: NGM Bio Announces Publication In The Lancet Of Phase 2 Study Of NGM282 In NASH Patients And Upcoming Presentations Of NGM282 NASH Histology Data And NGM282 PSC Data At The International Liver Congress 2018

Mar 08, 2018: Gastrointestinal Hormone Measurably Improved Symptoms of Non-Alcoholic Fatty Liver Disease

Feb 16, 2018: UK NICE Recommends Anticancer Agent Lenvima as Treatment for Thyroid Cancer

Feb 15, 2018: Positive Phase 3 Trial Results of Investigational Anticancer Agent Lenvatinib in Unresectable Hepatocellular Carcinoma (HCC) Published in The Lancet

Feb 10, 2018: Phase III Trial Results of Anticancer Agent Lenvatinib in Hepatocellular Carcinoma Published in the Lancet

Feb 05, 2018: NGM Reports Top-Line Results From Phase 2 Study Of NGM282 In Patients With Primary Sclerosing Cholangitis (PSC)

Jan 26, 2018: NICE Issues Technology Appraisal Guidance on Lenvatinib with Everolimus for Previously Treated Advanced RCC

Jan 25, 2018: Janssen to Present data on Erdafitinib at Prostate and Urothelial Cancers at ASCO GU 2018

Jan 22, 2018: Eisai Presents Results of Analysis of Phase III Trial of Lenvima (Lenvatinib) In Hepatocellular Carcinoma Based on Independent Imaging Review at ASCO-GI

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COMPANIES MENTIONED

Abbisko Therapeutics Co Ltd
Amgen Inc
ArQule Inc
Bayer AG
Blueprint Medicines Corp
Eddingpharm Inc
Eisai Co Ltd
Eli Lilly and Co
Genosco Inc
H3 Biomedicine Inc
Hanmi Pharmaceuticals Co Ltd
Incyte Corp
Ionis Pharmaceuticals Inc
Johnson & Johnson
Merrimack Pharmaceuticals Inc
NGM Biopharmaceuticals Inc
Novartis AG
Principia Biopharma Inc
Tasly Pharmaceutical Group Co Ltd
Vichem Chemie Research Ltd

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