

N Sulphoglucosamine Sulphohydrolase - Pipeline Review, H1 2020

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Abstracts

N Sulphoglucosamine Sulphohydrolase - Pipeline Review, H1 2020

SUMMARY

According to the recently published report 'N Sulphoglucosamine Sulphohydrolase - Pipeline Review, H1 2020'; N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) pipeline Target constitutes close to 9 molecules.

N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) - N-sulphoglucosamine sulphohydrolase is an enzyme encoded by the SGSH gene. This enzyme is involved in the lysosomal degradation of heparan sulfate. Mutations in this gene are associated with Sanfilippo syndrome A, one type of the lysosomal storage disease mucopolysaccaridosis III.

The report 'N Sulphoglucosamine Sulphohydrolase - Pipeline Review, H1 2020' outlays comprehensive information on the N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) 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 N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) targeted therapeutics development with respective active and dormant or discontinued projects. Currently, The molecules developed by companies in Phase III, Phase II, Preclinical



and Discovery stages are 1, 4, 3 and 1 respectively. Report covers products from therapy areas Genetic Disorders and Central Nervous System which include indications Mucopolysaccharidosis III (MPS III) (Sanfilippo Syndrome) and Alzheimer's Disease.

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 N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1)

The report reviews N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) 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 N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) targeted therapeutics and enlists all their major and minor projects

The report assesses N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) 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 N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or



EC 3.10.1.1) 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 N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1)

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 N Sulphoglucosamine Sulphohydrolase (Sulfoglucosamine Sulfamidase or Sulphamidase or SGSH or EC 3.10.1.1) 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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Amicus Therapeutics Inc

ArmaGen Inc

Denali Therapeutics Inc

Esteve Pharmaceuticals SA

JCR Pharmaceuticals Co Ltd

Lysogene SAS

Swedish Orphan Biovitrum AB

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Mechanism Of Action

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Featured News & Press Releases

Feb 25, 2020: Lysogene receives FDA fast track designation for LYS-SAF302 gene

therapy in MPS IIIA

Feb 12, 2020: Abeona Therapeutics announces presentation on its MPS IIIA drug candidate ABO-102 at WORLDSymposium

Dec 20, 2019: Abeona Therapeutics receives European Medicines Agency PRIME

Designation for ABO-102 Gene Therapy in MPS IIIA



Dec 18, 2019: Lysogene announces the publication of an article in the scientific journal "molecular therapy methods & clinical development" demonstrating the potential of its drug candidate LYS-SAF302

Oct 21, 2019: Abeona Therapeutics announces presentations at the 27th European Society of Gene and Cell Therapy (ESGCT) Congress

Jul 25, 2019: Abeona Therapeutics announces positive interim data from the ABO-102 phase 1/2 gene therapy clinical trial in MPS IIIA

Jul 25, 2019: Abeona Therapeutics announces positive interim data from the ABO-102 phase 1/2 gene therapy clinical trial in MPS IIIA

Jun 11, 2019: Lysogene announces first European patient treated in AAVance, Phase 2/3 clinical trial investigating LYS-SAF302, a gene therapy for the treatment of MPS IIIA (Sanfilippo Syndrome Type A)

Mar 22, 2019: Lysogene announces presentations at upcoming scientific and family conferences

Feb 14, 2019: Lysogene and Sarepta Therapeutics Announce Dosing of the First Patient in AAVance, a Phase 2/3 Clinical Trial Investigating LYS-SAF302, a Gene Therapy for the Treatment of MPS IIIA (Sanfilippo Syndrome Type A)

Jan 31, 2019: Abeona Therapeutics to present new supportive data for Novel Gene Therapies at WORLDSymposium

Dec 06, 2018: Abeona Therapeutics provides update on Sanfilippo Syndrome drug candidate ABO-102 at 2018 R&D Day

Sep 14, 2018: JCR to initiate development of a new drug candidate for sanfilippo syndrome Type A using J-Brain Cargo

Sep 05, 2018: Lysogene Announces FDA approval of IND Application to Initiate Phase 2-3 Clinical Trial in MPS IIIA

Aug 10, 2018: First patient dosed in phase 1/2 study evaluating SOBI003 for treatment of mucopolysaccharidosis type IIIA (MPS IIIA)

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COMPANIES MENTIONED

Abeona Therapeutics Inc
Amicus Therapeutics Inc
ArmaGen Inc
Denali Therapeutics Inc
Esteve Pharmaceuticals SA
JCR Pharmaceuticals Co Ltd
Lysogene SAS
Swedish Orphan Biovitrum AB



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