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Lysosomal Disease Treatments: Technologies and Global Markets

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Abstracts

Report Scope:

The lysosomal storage disease treatment market has been divided by type of indication into: Gaucher's Diseases, Fabry Diseases, Pompe's Syndrome, Mucopolysaccharidosis and more. Mucopolysaccharidosis is further segmented into Hunter syndrome, Hurlers Syndrome and Moriquo syndrome. The market is also segmented by routing of administration into IV and oral. The market has been divided into hospitals, clinics and other facilities into end users by type of treatment into enzyme replacement therapy and substrate reduction therapy. The market has been segmented into North America, Europe, Asia-Pacific and RoW.

Report Includes:

37 data tables and 25 additional tables

An overview of the global market for lysosomal disease treatments

Analyses of global market trends, with data from 2017, estimates for 2018 and projections of compound annual growth rates (CAGRs) through 2023

Segmentation of the global market by cancer therapy indiscation, treatment type, route of administration, end-use, application and geographical region

Information on current regulatory environment, and a trend analysis of recent government policies and regulations for lysosomal disease treatments

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Assessment of lysosomal storage diseases by the accumulated substrate and the profitability of focusing on ultra-orphan diseases

Profiles of major players in the industry, including Astellas Pharma, Astrazeneca, Eli Lilly and Co., Merck & Co., Inc., and Novo Nordisk A/S

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Contents

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CHAPTER 5 MARKET BREAKDOWN BY DISEASE TYPE

Gauchers
Diagnosis
Treatment
Oral Substrate Reduction Therapy (SRT)
Fabry Disease
Type 1 Classic Phenotype
Pompe Disease
Causes
Treatment
Mucopolysaccharidosis (MPS)
Hurlers (MPS I)
Hunter Syndrome (MPS II)
Morquio Syndrome (MPS IV)
Others

CHAPTER 6 MARKET BREAKDOWN BY ROUTE OF ADMINISTRATION

Oral Route of Administration
Intravenous Route of Administration

CHAPTER 7 MARKET BREAKDOWN BY END USE

Hospitals
Clinics
Others

CHAPTER 8 MARKET BREAKDOWN BY TREATMENT TYPE

Enzyme Replacement Therapy Market
Substrate Reduction Therapy

CHAPTER 9 MARKET BREAKDOWN BY REGION

North America
Europe
Asia-Pacific

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List Of Tables

LIST OF TABLES

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List Of Figures

LIST OF FIGURES

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