

Solriamfetol - Drug Insight and Market Forecast

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Abstracts

This report can be delivered to the clients within 48 Hours

"Solriamfetol - Drug Insight and Market Forecast - 2030" report by DelveInsight outlays comprehensive insights of the product indicated for the treatment of its approved condition. A detailed picture of the Solriamfetol in Seven Major Markets, i.e., United States, EU5 (Germany, France, Italy, Spain, and the United Kingdom), and Japan, for the study period 2017–2030 is provided in this report along with a detailed description of the product. The product details covers mechanism of action, dosage and administration, route of synthesis, and pharmacological studies, including product marketed details, regulatory milestones, and other development activities. Further, it also consists of market assessments inclusive of the market forecast, SWOT analysis, and detailed analyst views. It further highlights the market competitors, late-stage emerging therapies, and patent details in the global space.

Drug Summary

Solriamfetol, sold under the brand name Sunosi, is a medication used for the treatment of excessive sleepiness associated with narcolepsy and sleep apnea. It is derived from d-phenylalanine and its chemical name is-2-amino-3-phenylpropylcarbamate hydrochloride. It is a norepinephrine–dopamine reuptake inhibitor.

SCOPE OF THE REPORT

The report provides insights into:

A comprehensive product overview including the product description, mechanism of action, dosage and administration, route of synthesis, pharmacological studies (pharmacodynamics and pharmacokinetics) and



adverse reactions.

Elaborated details on regulatory milestones and other development activities have been provided in this report.

The report also highlights the drug marketed details across the United States, Europe and Japan.

The report also covers the patents information with expiry timeline around Solriamfetol.

The report contains historical and forecasted sales for Solriamfetol till 2030.

Comprehensive coverage of the late-stage emerging therapies (Phase III) in the space with a brief snapshot of the details.

The report also features the SWOT analysis with analyst insights and key findings of Solriamfetol.

METHODOLOGY

The report is built using data and information sourced primarily from internal databases, primary and secondary research and in-house analysis by DelveInsight's team of industry experts. Information and data from the secondary sources have been obtained from various printable and nonprintable sources like search engines, news websites, global regulatory authorities websites, trade journals, white papers, magazines, books, trade associations, industry associations, industry portals and access to available databases.

Solriamfetol Analytical Perspective by DelveInsight

In-depth Solriamfetol Market Assessment

This report provides a detailed market assessment of Solriamfetol in Seven Major Markets, i.e., United States, EU5 (Germany, France, Italy, Spain, and the United Kingdom), and Japan. This segment of the report provides historical and forecasted sales data from 2017 to 2030.



Solriamfetol Clinical Assessment

The report provides the clinical trials information of Solriamfetol covering trial interventions, trial conditions, trial status, start and completion dates.

Report highlights

In the coming years, the market scenario for Solriamfetol is set to change due to the extensive research in the treatment of the indicated condition and incremental healthcare spending across the world; which would expand the size of the market to enable the drug manufacturers to penetrate more into the market.

The companies and academics are working to assess challenges and seek opportunities that could influence Solriamfetol dominance. The therapies under development are focused on novel approaches to treat/improve the disease condition.

Other approved products for the disease are giving market competition to Solriamfetol and launch of late-stage emerging therapies in the near future will significantly impact the market.

A detailed description of regulatory milestones, development activities, and some key findings provide the current market scenario of Solriamfetol.

Our in-depth analysis of the sales data of Solriamfetol from 2017 to 2030 will support the clients in the decision-making process regarding their therapeutic portfolio by identifying the overall scenario of the Solriamfetol in the market.

Key Questions

What is the prescribed dosage and strengths of Solriamfetol are available in the market?

What are the common adverse reactions or side effects of Solriamfetol?

What is the product type, route of administration and mechanism of action of



Solriamfetol?

What are the chemical specifications of Solriamfetol?

How are the clinical trials diversified on the basis of the trial status?

What is the history of Solriamfetol, and what is its future?

What are the marketed details of Solriamfetol in the seven major countries, including the United States, Europe (Germany, France, Italy, Spain, and the United Kingdom), and Japan?

How many patents have been granted to Solriamfetol and when these patents will get expire?

What are the pros (benefits) and cons (disadvantages) of Solriamfetol?

In which countries Solriamfetol got approval and when it gets launched?

What are the clinical trials are currently ongoing for Solriamfetol?

How the safety and efficacy results determined the approval of Solriamfetol?

What are the key collaborations, mergers and acquisitions, licensing and other activities related to the Solriamfetol development?

What are the key designations that have been granted to Solriamfetol?

What is the historical and forecasted market scenario of Solriamfetol?

How is the market trend of Solriamfetol is different in the Seven Major Markets (the United States, EU5 [Germany, France, Italy, Spain, and the United Kingdom], and Japan)?

What are the other approved products available and how these are giving competition to Solriamfetol?

Which are the late-stage emerging therapies under development for the treatment of the indicated condition?







Contents

1. PRODUCT OVERVIEW

- 1.1. Indication
- 1.2. Mechanism of Action
- 1.3. Dosage and Administration
 - 1.4.1 Dosage Forms and Strengths
- 1.4. Route of Synthesis
- 1.5. Pharmacology
 - 1.4.2 Pharmacodynamics
 - 1.4.3 Pharmacokinetics
- 1.6. Adverse Reactions
- 1.7. Product Snapshot
- 1.8. Development Milestones of Solriamfetol
- 1.9. Marketed Details
 - 1.4.4 United States
 - 1.4.5 Europe
 - 1.4.6 Japan
- 1.10. Patent Details

2. SWOT ANALYSIS

2.1. Analyst Views

3. REGULATORY MILESTONES

- 3.1. Approvals
- 3.2. Research and Development
- 3.3. Clinical Trials Information
- 3.4. Safety and Efficacy
- 3.5. Product Developmental Activities

4. MARKET ASSESSMENT

- 4.1. 7MM Market Analysis
- 4.2. United States
- 4.3. Europe
- 4.4. Japan



- 4.5. Key Findings
- **5. MARKET COMPETITORS**
- **6. EMERGING THERAPIES**
- 7. APPENDIX
- 7.1. Report Purchase Options



List Of Tables

LIST OF TABLES

Table :	1	Sol	riam [.]	fetal	\Box	escription
Iabic		20	IIIaiii	ICIUI.	\boldsymbol{L}	<i>เ</i> ซอบเมนเบเ

- Table 2 Solriamfetol, Trial Diversification
- Table 3 Solriamfetol, Marketed Details United States
- Table 4 Solriamfetol, Marketed Details Europe
- Table 5 Solriamfetol, Marketed Details Japan
- Table 6 Patent Details: Solriamfetol
- Table 7 Solriamfetol, Clinical Trial Description, 2020
- Table 8 Safety and Efficacy Results for Solriamfetol
- Table 9 Solriamfetol, 7MM Market Size from 2017 to 2030 (in Million USD)
- Table 10 Solriamfetol, US Market Size from 2017 to 2030 (in Million USD)
- Table 11 Solriamfetol, EU Market Size from 2017 to 2030 (in Million USD)
- Table 12 Solriamfetol, EU5 Market Size from 2017 to 2030 (in Millions USD)
- Table 13 Solriamfetol, Japan Market Size from 2017 to 2030 (in Million USD)
- **Table 14 Market Competitors**
- Table 15 Emerging Therapies



List Of Figures

LIST OF FIGURES

- Figure 1 The Development Timeline of Solriamfetol
- Figure 2 Patent Details, Solriamfetol
- Figure 3 Solriamfetol, 7MM Market Size from 2017 to 2030 (in Million USD)
- Figure 4 Solriamfetol, US Market Size from 2017 to 2030 (in Millions USD)
- Figure 5 Solriamfetol, EU Market Size from 2017 to 2030 (in Millions USD)
- Figure 6 Solriamfetol, EU5 Market Size from 2017 to 2030 (in Millions USD)
- Figure 7 Solriamfetol, Japan Market Size from 2017 to 2030 (in Millions USD)



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