

# Copiktra - Drug Insight and Market Forecast - 2030

<https://marketpublishers.com/r/CE098F66D706EN.html>

Date: February 2021

Pages: 30

Price: US\$ 2,750.00 (Single User License)

ID: CE098F66D706EN

## Abstracts

This report can be delivered to the clients within 48 Hours

### OVERVIEW

“Copiktra - 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 Copiktra 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

Copiktra is an oral inhibitor of phosphoinositide 3-kinase (PI3K), and the first approved dual inhibitor of PI3K-delta and gamma involved in the signaling that leads to malignant B-cells proliferation and cancer.

Copiktra is indicated for the treatment of adult patients with relapsed or refractory chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) after at least two prior therapies and relapsed or refractory follicular lymphoma (FL) after at least two prior systemic therapies. Copiktra is also being developed for the treatment of peripheral T-cell lymphoma (PTCL), for which it has received Fast Track status and Orphan Drug Designation and is being investigated in combination with other agents through

company and investigator-sponsored studies.

A European Marketing Authorization Application for Copiktra was submitted in November 2019 to the European Medicines Agency (EMA) seeking approval for the treatment of patients with relapsed or refractory chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) and relapsed or refractory follicular lymphoma (FL).

## **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 Copiktra.

The report contains historical and forecasted sales for Copiktra 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 Copiktra.

## **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.

## Copiktra Analytical Perspective by DelveInsight

### In-depth Copiktra Market Assessment

This report provides a detailed market assessment of Copiktra 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.

### Copiktra Clinical Assessment

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

## Report highlights

In the coming years, the market scenario for Copiktra 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 Copiktra 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 Copiktra 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 Copiktra.

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

## Key Questions

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

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

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

What are the chemical specifications of Copiktra?

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

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

What are the marketed details of Copiktra 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 Copiktra and when these patents will get expire?

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

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

What are the clinical trials are currently ongoing for Copiktra?

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

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

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

What is the historical and forecasted market scenario of Copiktra?

How is the market trend of Copiktra 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 Copiktra?

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 Copiktra
- 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 Copiktra, Description

Table 2 Copiktra, Trial Diversification

Table 3 Copiktra, Marketed Details United States

Table 4 Copiktra, Marketed Details Europe

Table 5 Copiktra, Marketed Details Japan

Table 6 Patent Details: Copiktra

Table 7 Copiktra, Clinical Trial Description, 2020

Table 8 Safety and Efficacy Results for Copiktra

Table 9 Copiktra, 7MM Market Size from 2017 to 2030 (in Million USD)

Table 10 Copiktra, US Market Size from 2017 to 2030 (in Million USD)

Table 11 Copiktra, EU Market Size from 2017 to 2030 (in Million USD)

Table 12 Copiktra, EU5 Market Size from 2017 to 2030 (in Millions USD)

Table 13 Copiktra, 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 Copiktra

Figure 2 Patent Details, Copiktra

Figure 3 Copiktra, 7MM Market Size from 2017 to 2030 (in Million USD)

Figure 4 Copiktra, US Market Size from 2017 to 2030 (in Millions USD)

Figure 5 Copiktra, EU Market Size from 2017 to 2030 (in Millions USD)

Figure 6 Copiktra, EU5 Market Size from 2017 to 2030 (in Millions USD)

Figure 7 Copiktra, Japan Market Size from 2017 to 2030 (in Millions USD)

## I would like to order

Product name: Copiktra - Drug Insight and Market Forecast - 2030

Product link: <https://marketpublishers.com/r/CE098F66D706EN.html>

Price: US\$ 2,750.00 (Single User License / Electronic Delivery)

If you want to order Corporate License or Hard Copy, please, contact our Customer Service:

[info@marketpublishers.com](mailto:info@marketpublishers.com)

## Payment

To pay by Credit Card (Visa, MasterCard, American Express, PayPal), please, click button on product page <https://marketpublishers.com/r/CE098F66D706EN.html>

To pay by Wire Transfer, please, fill in your contact details in the form below:

First name:  
Last name:  
Email:  
Company:  
Address:  
City:  
Zip code:  
Country:  
Tel:  
Fax:  
Your message:

**\*\*All fields are required**

Customer signature \_\_\_\_\_

Please, note that by ordering from marketpublishers.com you are agreeing to our Terms & Conditions at <https://marketpublishers.com/docs/terms.html>

To place an order via fax simply print this form, fill in the information below and fax the completed form to +44 20 7900 3970