

# Global Pharmacovigilance and Drug Safety Software Market 2026 by Company, Regions, Type and Application, Forecast to 2032

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## Abstracts

According to our (Global Info Research) latest study, the global Pharmacovigilance and Drug Safety Software market size was valued at US\$ 267 million in 2025 and is forecast to a readjusted size of US\$ 375 million by 2032 with a CAGR of 5.1% during review period.

Pharmacovigilance and drug safety software are digital systems designed specifically for pharmaceutical companies, regulatory agencies, and health organizations. Their core function is to collect, record, analyze, and report adverse reaction information before and after drug marketing. These tools efficiently manage safety data, ensure compliance with global regulatory requirements, aim to protect patient medication safety, and reduce the workload of pharmacovigilance specialists through automated processes.

With the increasing complexity of global clinical trials and stricter post-marketing regulations, PV software has evolved from a simple compliance tool into a core strategic asset for pharmaceutical companies. The current industry focus is on the application of next-generation AI.

The industry is driven primarily by both increasingly stringent global regulatory compliance requirements and the explosive growth of massive amounts of diverse data. Firstly, regulatory agencies in various countries (such as the FDA's updated FAERS system and the EMA's improved EudraVigilance standards) require companies to have faster reporting timeliness and greater transparency, forcing pharmaceutical companies to abandon traditional manual processing methods and seek automated software support. Secondly, the rise of biologics, orphan drugs, and cell and gene therapies

(CGTs) has brought about more complex safety characterization, and this need for long-term risk tracking has directly driven the growth of dedicated PV platforms.

This report is a detailed and comprehensive analysis for global Pharmacovigilance and Drug Safety Software market. Both quantitative and qualitative analyses are presented by company, by region & country, by Type and by Application. As the market is constantly changing, this report explores the competition, supply and demand trends, as well as key factors that contribute to its changing demands across many markets. Company profiles and product examples of selected competitors, along with market share estimates of some of the selected leaders for the year 2025, are provided.

### **Key Features:**

Global Pharmacovigilance and Drug Safety Software market size and forecasts, in consumption value (\$ Million), 2021-2032

Global Pharmacovigilance and Drug Safety Software market size and forecasts by region and country, in consumption value (\$ Million), 2021-2032

Global Pharmacovigilance and Drug Safety Software market size and forecasts, by Type and by Application, in consumption value (\$ Million), 2021-2032

Global Pharmacovigilance and Drug Safety Software market shares of main players, in revenue (\$ Million), 2021-2026

### **The Primary Objectives in This Report Are:**

To determine the size of the total market opportunity of global and key countries

To assess the growth potential for Pharmacovigilance and Drug Safety Software

To forecast future growth in each product and end-use market

To assess competitive factors affecting the marketplace

This report profiles key players in the global Pharmacovigilance and Drug Safety Software market based on the following parameters - company overview, revenue, gross margin, product portfolio, geographical presence, and key developments. Key

companies covered as a part of this study include Oracle, Veeva Systems, ArisGlobal, IQVIA, EXTEDO, RxLogix, Ennov, Uppsala Monitoring Centre, Medidata, Clarivate, etc.

This report also provides key insights about market drivers, restraints, opportunities, new product launches or approvals.

### Market segmentation

Pharmacovigilance and Drug Safety Software market is split by Type and by Application. For the period 2021-2032, the growth among segments provides accurate calculations and forecasts for Consumption Value by Type and by Application. This analysis can help you expand your business by targeting qualified niche markets.

### Market segment by Type

Adverse Event Reporting Software

Drug Safety Audits Software

Issue Tracking Software

Fully Integrated Software

### Market segment by Data Range

Pre-market

Post-market

### Market segment by Deployment Method

Cloud-Based

On-Premises

Hybrid Mode

## Market segment by Application

Pharma and Biotech Companies

Contract Research Organizations (CROs)

Business Process Outsourcing (BPO) Firms

Pharmacovigilance Service Providers

Others

## Market segment by players, this report covers

Oracle

Veeva Systems

ArisGlobal

IQVIA

EXTEDO

RxLogix

Ennov

Uppsala Monitoring Centre

Medidata

Clarivate

AB Cube

Cloudbyz

Cognizant Technology

Taimei Technology

eMedyun

Bioknow

Singedi Health

Market segment by regions, regional analysis covers

North America (United States, Canada and Mexico)

Europe (Germany, France, UK, Russia, Italy and Rest of Europe)

Asia-Pacific (China, Japan, South Korea, India, Southeast Asia and Rest of Asia-Pacific)

South America (Brazil, Rest of South America)

Middle East & Africa (Turkey, Saudi Arabia, UAE, Rest of Middle East & Africa)

**The content of the study subjects, includes a total of 13 chapters:**

Chapter 1, to describe Pharmacovigilance and Drug Safety Software product scope, market overview, market estimation caveats and base year.

Chapter 2, to profile the top players of Pharmacovigilance and Drug Safety Software, with revenue, gross margin, and global market share of Pharmacovigilance and Drug Safety Software from 2021 to 2026.

Chapter 3, the Pharmacovigilance and Drug Safety Software competitive situation, revenue, and global market share of top players are analyzed emphatically by landscape contrast.

Chapter 4 and 5, to segment the market size by Type and by Application, with consumption value and growth rate by Type, by Application, from 2021 to 2032.

Chapter 6, 7, 8, 9, and 10, to break the market size data at the country level, with revenue and market share for key countries in the world, from 2021 to 2026. and Pharmacovigilance and Drug Safety Software market forecast, by regions, by Type and by Application, with consumption value, from 2027 to 2032.

Chapter 11, market dynamics, drivers, restraints, trends, Porters Five Forces analysis.

Chapter 12, the key raw materials and key suppliers, and industry chain of Pharmacovigilance and Drug Safety Software.

Chapter 13, to describe Pharmacovigilance and Drug Safety Software research findings and conclusion.

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