



## Postmarketing Surveillance – Japan

Fulfill all requirements for PMS studies in Japan

Viedoc PMS (postmarketing surveillance) for the Japanese market is specially developed to fulfill all country-specific requirements, including booklet data collection and Kaifu functionality. Seamless integration with Viedoc Clinic means our solution shares all the robust and flexible features of our eCRF. For data managers, project managers, and anyone involved in clinical operations, we deliver ease of use with comprehensive oversight and reporting — saving significant time and effort.

## Licensing overview

Our PMS solution for the Japanese market adds a powerful postmarketing surveillance (PMS) management and compliance tool to our intuitive EDC environment.

viedoc **EDC**  
viedoc **admin**  
viedoc **designer**  
viedoc **clinic**  
viedoc **reports**

viedoc **pms**  
PMS interface

## Features

- **Booklet data collection**  
Support booklet data collection, allowing site users to enter data and sponsor users to review data in booklet-related forms.
- **Kaifu (submit/receive) process**  
Booklet submission for review and approval process.
- **Feature-rich booklet management page**  
Intuitive UI and advanced functionality for sponsor users, including sorting and filtering data, guiding users, and identifying critical tasks.
- **Adverse events management and reporting**  
Adverse events reporting functionality through the booklet and as a standalone instance, with options to include additional data prior to regulatory submissions.
- **Query management**  
Full utilization of the query process within Viedoc Clinic, with option to send queries back to site users.
- **PMS-specific reports**  
PMS-specific dashboards and reports within the Viedoc Reports tool.
- **Progress management integration**  
Integration with external progress management systems.

## What's in it for you?

- **Streamline the PMS process**  
Collect booklet data, manage the Kaifu process, and report and review every step in a single application, ensuring compliance and audit readiness.
- **Easily follow study progress**  
PMS-specific dashboards and integration with external progress management systems provide confidence and control.
- **Ensure transparency and accountability**  
Flexible query management makes it easy to set review responsibilities between sites and sponsors.

## Want to learn more?



Get all the details – click here to watch demos, browse FAQs, and explore case studies on our website.

## About Viedoc

Viedoc delivers powerful eClinical solutions that streamline clinical trials, making data collection, management, and analysis faster and easier. Trusted in 75+ countries, powering 7,000+ studies with more than 1.6M participants, we accelerate clinical trials and bring life-changing treatments to market faster.