

SUVODA®

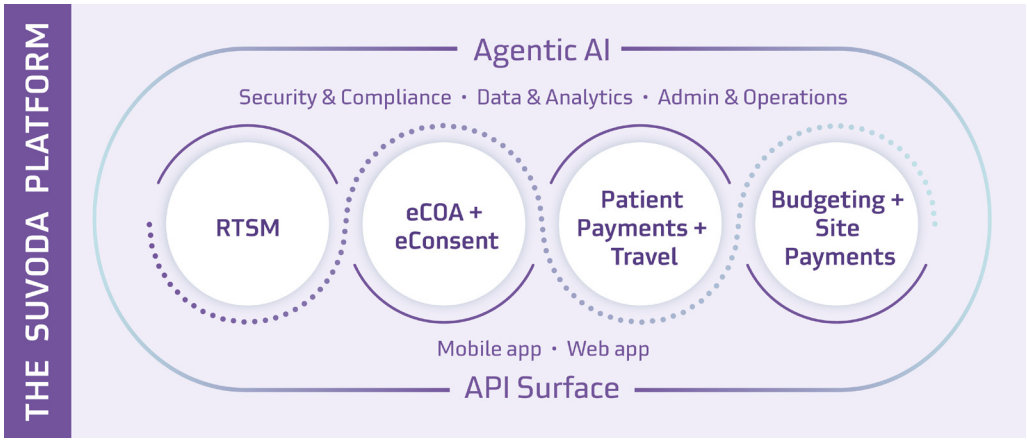
Trial wisely

Flexible and Simplified Consent Management

Suvoda eConsent on the Suvoda Platform: one app for participants and a connected experience for study teams

- A single patient app and patient-centric workstream
- Reduced integrations and a single data model
- Agentic AI built on the proven, patented Suvoda Platform
- A consultative, services team with deep expertise

Seamlessly manage mission-critical, time-sensitive moments of the patient journey with the Suvoda Platform. By delivering real-time data that informs decision-making through critical moments like randomization, endpoint collection, and patient payments, the Suvoda Platform eases the experience for sponsors, CROs, and sites. Suvoda's patented technology enables rapid and compliant design and deployment powered by agentic AI and tailored by the expert Suvoda services team that brings deep clinical trial knowledge to every trial — advising, anticipating, and partnering through the full lifespan of a study. Embrace the inherent complexities and constant change with a modern, precision-built platform that supports calm action and confident decisions to move life-changing studies forward.



Simplify consent management with the unified Suvoda Platform

Suvoda eConsent is an intuitive, flexible electronic informed consent application for clinical trials. Whether on-site or remote, eConsent allows patients, caregivers, and authorized representatives to review consent documentation, while providing consistency and compliance. Leveraging seamless integration with Suvoda RTSM/IRT, eConsent enables real-time visibility into and automated control over the patient consent process — helping to reduce regulatory risk, administrative burden, and study duration and expenses created by mid-study consent requirements.

LEARN MORE

Visit suvoda.com
to learn about our solutions

BENEFITS

A SINGLE, PATIENT APP

Patients can complete consent remotely, from home, through the Suvoda web app. With one login, they can also respond to questionnaires, access payments, manage travel, and more—all within the same app. A connected experience helps simplify study activities and support engagement throughout the trial.

VISIBILITY

Powered by the Suvoda Platform, reduce site workload through integrated reporting that provides real-time visibility into consent status across sites

CONTROL

Workflows are fully streamlined with a completely controlled process safeguarding quality and compliance

REDUCED REGULATORY RISK

Verify consent remotely and take early corrective action to reduce the risk of excluded patients and discarded data

FLEXIBILITY

Accommodate a variety of consent methods (electronic signature, wet ink upload, or print-and-sign consent), consent types (subject, caregiver, legally authorized representative), and consent configurations (one or multiple signatures in a single consent form from a participant), to meet differing requirements across trials and countries

IMPROVED COMPREHENSION

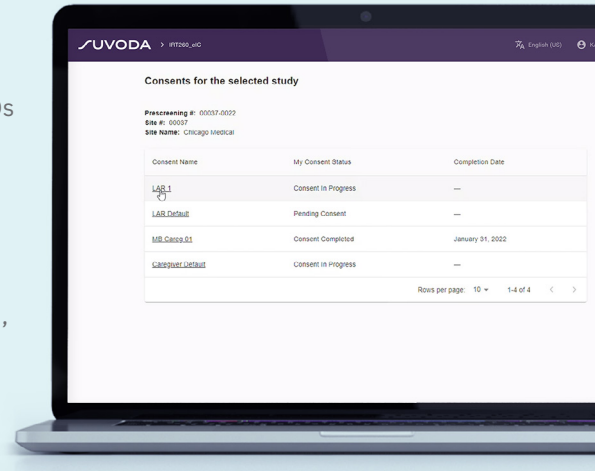
Interactive and multi-media features provide summaries, glossary, and ability to ask questions in the ICF to improve patient understanding

INTUITIVE INTERFACE

Sites can move easily between eConsent and other Suvoda Platform tasks with minimal additional user training

KEY FEATURES

+ Embedded patient-focused videos and FAQs
+ Cross-linked glossary
+ In-document discussion threads
+ Electronic signature, wet ink upload, and print & sign
+ Available on multiple devices — web-based, on-site, or patient device
+ Multilingual capability
+ How-to videos for site staff



suvoda.com | info@suvoda.com