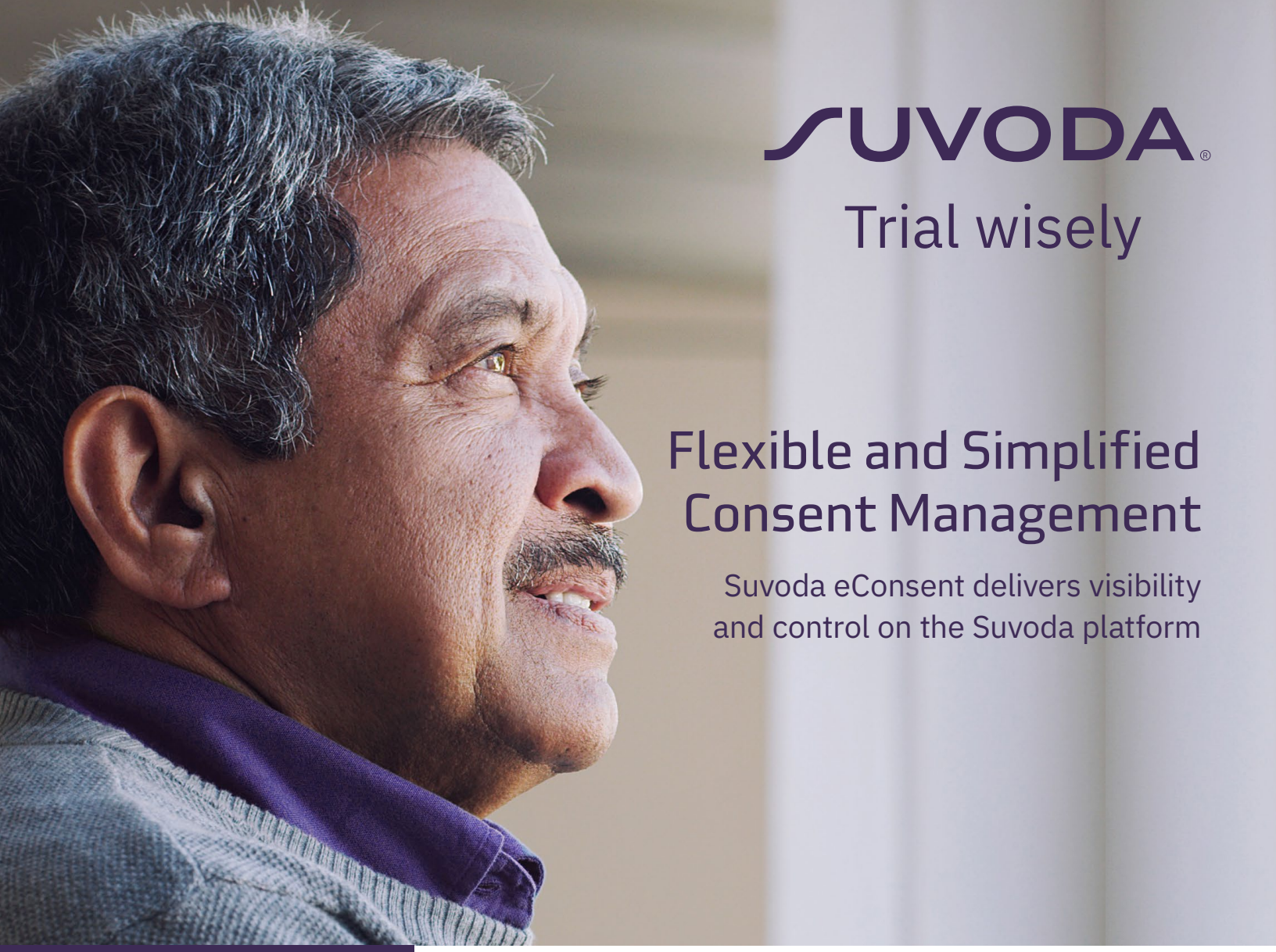


SUVODA®

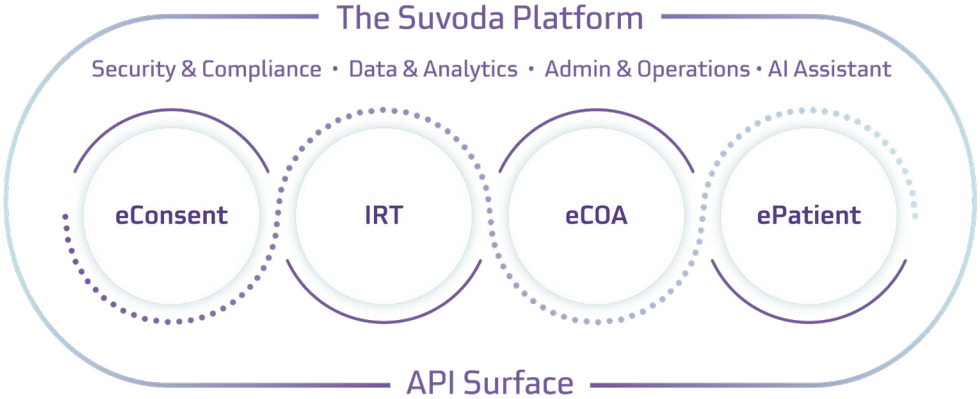
Trial wisely

Flexible and Simplified Consent Management

Suvoda eConsent delivers visibility and control on the Suvoda platform

- A single, patient-centric workstream
- Patented technology powers rapid design and deployment
- Reduced integrations and a single data model
- Future-proof eClinical programs with flexible solutions

Seamlessly manage mission-critical, time-sensitive moments of the patient journey in your clinical trial through a single, patient-centric workstream with the Suvoda Platform. Built from the ground-up, the easy-to-use ecosystem of eConsent, IRT, eCOA, and ePatient requires just one log in for sponsor teams and site professionals. And because all solutions are on the Suvoda Platform, they work harmoniously together and with other applications. Suvoda's patented technology powers rapid design and deployment, while allowing our expert services team to tailor solutions to meet the needs of each protocol. Gain control over the inherent complexities, infinite variables, and constant change in your life-sustaining studies.



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 the 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

VISIBILITY

Integrated reporting provides real-time visibility into consent status across sites within your IRT

CONTROL

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

FLEXIBILITY

Accommodate a variety of consent methods and types to meet differing requirements across trials & 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

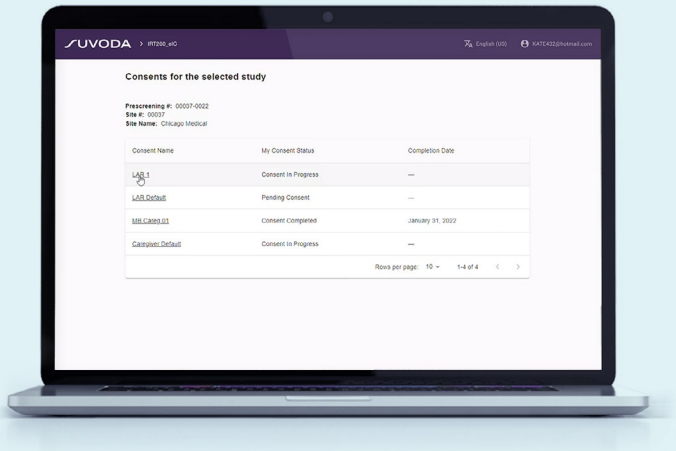
Smoothly move between IRT activities and eConsent tasks with minimal additional user training

REDUCED REGULATORY RISK

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

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



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