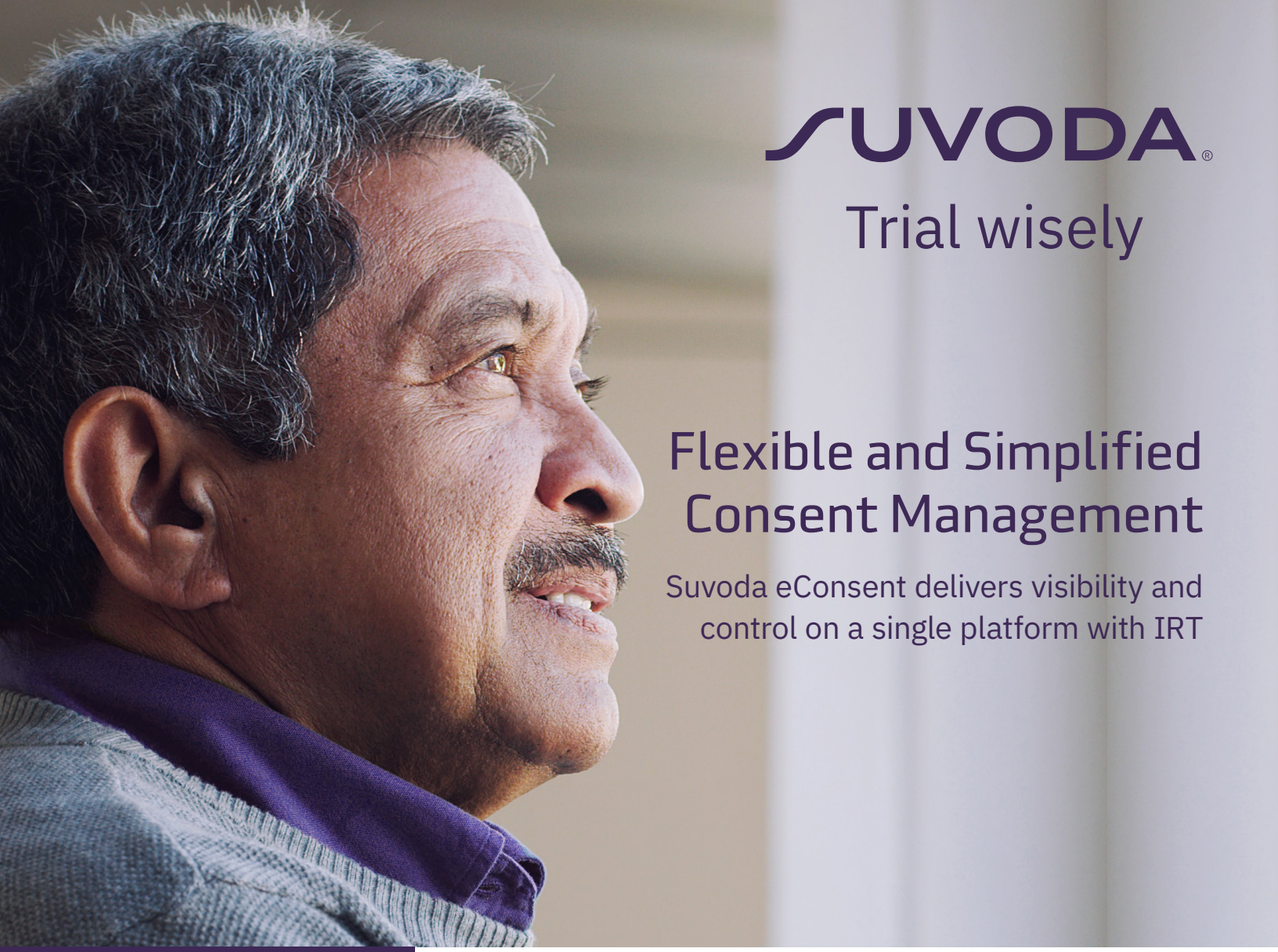


SUVODA®

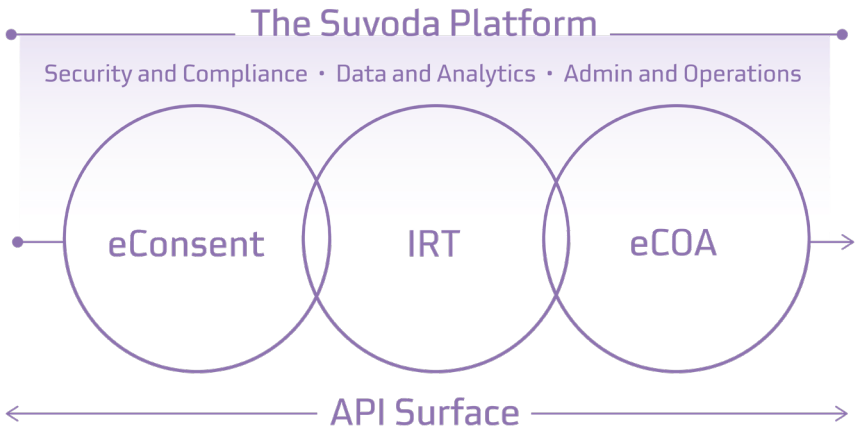
Trial wisely

Flexible and Simplified Consent Management

Suvoda eConsent delivers visibility and control on a single platform with IRT

- Enhanced user experience and reduced workflow
- Rapid design and deployment of complex trials
- Improved data workflow and reduced integrations
- Future-proofed eClinical programs

Seamlessly manage complex, mission-critical, time-sensitive moments of the patient journey in a clinical trial—be it traditional, decentralized, or hybrid—through a single, patient-centric workstream. The Suvoda Platform – a purpose-built, easy-to-use ecosystem delivering eConsent, IRT, and eCOA - enables these solutions to work harmoniously together and with other applications, to be tailored to meet the needs of each protocol, and to be upgradable to benefit from continuous enhancements to the platform. Combining the power of IRT and eConsent solutions in one easy-to-use platform allows you to track consent, screening, and randomization data in one place - gain control of the infinite variables and constant change in your complex, life-sustaining studies.



Seamless Architecture: eConsent + IRT

Suvoda's 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

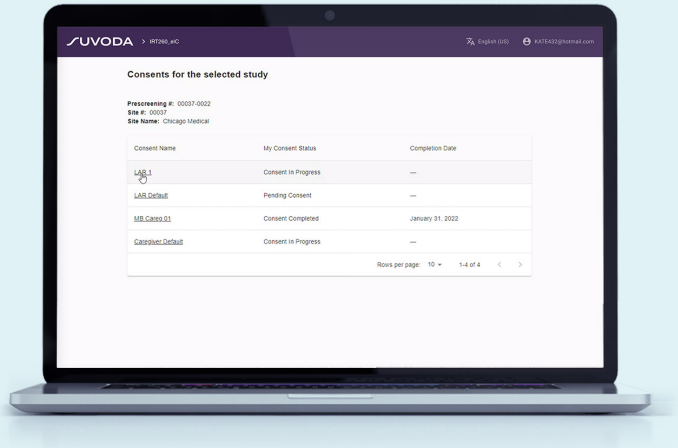
Smoothly move between IRT activities and eConsent tasks with minimal additional user training

REDUCED REGULATORY RISK

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

KEY FEATURES

+ Embedded 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



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