Fact Sheet

IQVIA Complete Consent

Global, compliant eConsent that improves enrollment, patient understanding, and study execution

Transform informed consent into a patient-centric, digital experience

Patient consent has historically been a slow, paper-based process. IQVIA Complete Consent transforms informed consent into a flexible, digital experience-, improving enrollment, enhancing patient understanding and strengthening compliance across global studies.

Improve understanding and consent quality across diverse patient populations

Empower patients to make informed decisions with engaging, accessible consent experience designed to support recruitment and retention across global trials.

Key capabilities

- Multimedia content improves patient understanding
- Digital consent on any device, anywhere
- Configurable to meet diverse patient and protocol needs
- Real-time review and discussion tools
- Built-in privacy and data protection controls

Proven impact

100% of patients rated Complete Consent as easy-to-use

60% reported improved understanding

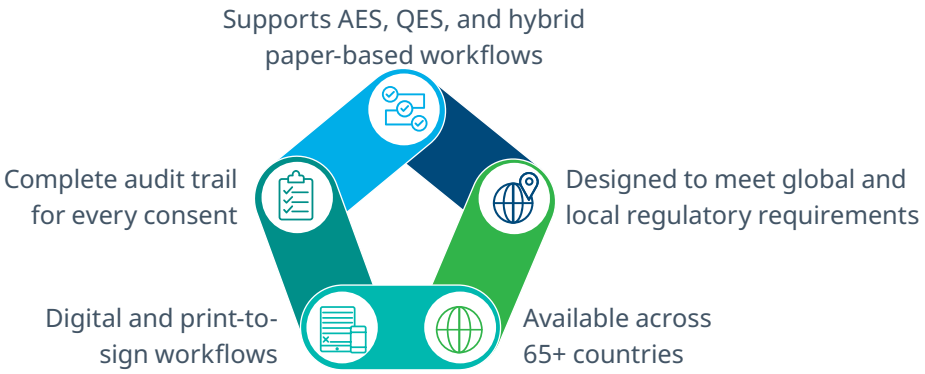
74% preferred Complete Consent over paper

57%

Reduce errors and improve compliance

Across nearly 100 IQVIA trials, Complete Consent reduced consent-related protocol deviations from 14% to 6%, helping sponsors improve compliance and reduce downstream study risk.

Flexible, globally compliant eSignature options



A versatile platform for any trial design

IQVIA Complete Consent supports any study, from simple to highly complex.

Multimedia Consent	For complex, multi-cohort trial designs, such as platform, umbrella, and basket studies, across high-complexity therapeutic areas like oncology and CNS
Interactive Consent	For mid-complexity trials (e.g. vaccine, lifestyle diseases, and simple diagnostic)
Simple PDF	For simple trials including screening, registry, observational, pilot, epidemiological, and translational studies

Deliver value across sponsors and sites

Sponsors

- Reduce study delays and costs associated with consent errors
  - » Enable consistent execution across global protocols
  - » Strengthen compliance and oversight with full audit visibility

Sites

- Streamline consent workflows and reduce administrative burden
  - » Accelerate patient onboarding
  - » Deliver a more consistent, patient-friendly experience

Integrates seamlessly with the IQVIA Patient Suite to connect consent, patient data and study workflow, improving operational efficiency and overall study outcomes.

A proven track record and unmatched scale.

400K+ Participants | **55K+** Site users | **10K+** Sites | **350+** Studies | **60+** Countries

Learn more

Explore how IQVIA Complete Consent can support your next study and improve patient engagement from the first interaction.

IQVIA TECHNOLOGIES

CONTACT US

iqvia.com/econsent

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