Fact Sheet

IQVIA Patient Suite

Improve data quality and enhance trial outcomes with unified technologies, data, and delivery

Harness the power of IQVIA Patient Suite to accelerate decision-making, raise operational standards and drive better trial outcomes. Our suite of best-in-class, integrated solutions transform patient engagement and data management in clinical trials: eConsent, Interactive Response Technology (IRT), Electronic Clinical Outcome Assessment (eCOA) and Connected Devices.

IQVIA Patient Suite offers sponsors the flexibility to adopt the full Suite or integrate individual solutions within their current environments. With unmatched planning and delivery experience, our team partners with customers to configure solutions for site-based, hybrid or decentralized trials.

A proven solution to support today's complex global studies

PATIENT SUITE	ECOA	ECOA	CONNECTED DEVICES	ECONSENT	IRT
>500	**>2,500**	**>90%**	**>40**	**>350**	**>400**
Standard integrations with labs, EDC, depots, etc.	Library of pre-validated digital instruments	Average patient protocol compliance (versus 11% with paper)	Portfolio of medical-grade devices	Studies conducted across 10,000 sites	Drug supply and eClinical integrations

Focus on trial success, not operational friction

Intuitive, interoperable products and tools deliver superior study outcomes

- Best-in-breed applications support end-to-end patient-related functions
- Flexibility to interoperate with existing clinical solutions
- Broadest portfolio of medical devices, wearables and sensors
- Built-in governance ensures software management aligns with customer goals and supports compliance
- Integrate with IQVIA eSource, telemedicine, labs, home health nursing, patient concierge, and more services

Exceptional data performance and a single experienced team enhance trial quality

- Ensure the highest integrity data for regulatory submission
- Scientific, clinical, medical and operational experts engage and consult across all trial stages
- Unmatched design, delivery and ongoing support by a single, experienced team
- Combined reports surface actionable insights and study intelligence
- Integrated data infrastructure and automation enhance clinical and operational data analytics

Simplified trial operations and elevated experiences for all stakeholders

- Automated workflows eliminate duplicative data entry and save time
- Unified patient reporting across all applications in the Patient Suite
- Around-the-clock, dedicated support for sites, 24/7/365
- Streamlined project management for study conduct, protocol amendments, etc.
- Readily accommodate protocol changes and other unexpected disruptions
- Risk-based monitoring drives timely responses to emerging issues as they arise

IQVIA TECHNOLOGIES

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