Fact Sheet

IQVIA Interactive Response Technology

Run faster, higher quality patient-centric studies

Accelerate study start-up and amendments, simplify site and sponsor experience, reduce trial supply costs and improve study decision making while ensuring quality standards and patient safety during your trial with IQVIA Interactive Response Technology. Built on a robust, scalable platform, our patient-centric approach to technology delivery provides exceptional value and peace of mind for sponsors and their patients in clinical trials. IQVIA IRT is part of the Digital Patient Suite of IQVIA Technologies Orchestrated Clinical Trials (OCT), an **end-to-end, patient-centric platform that provides an unparalleled data infrastructure, seamless connectivity and intuitive design to drive smarter, faster trials.**

Your challenges

Complex and lengthy site/provider workflows and interactions across multiple eClinical systems

Randomization tools that don't support complex or adaptive schemas or integrate across platforms

Site stock-outs or IP waste due to fluctuating or unpredictable enrollment or study changes

Struggles in optimizing supply management and gaining transparency

Time-consuming and costly protocol amendment and change management process

Your solution

IQVIA IRT provides randomization and trial supply management support for simple to complex trials that:

- Accelerates study start-up and protocol amendments with flexible, adaptable systems and self-service capabilities
- Simplifies site and investigator experience with streamlined workflows and reduced data entry
- Reduces supply costs through data driven insights, automation and expertise
- Improves decision making with access to real-time data and the right tools to monitor your trials
- Enhances insights and reduces administrative burden with 400+ turn-key eClinical integrations

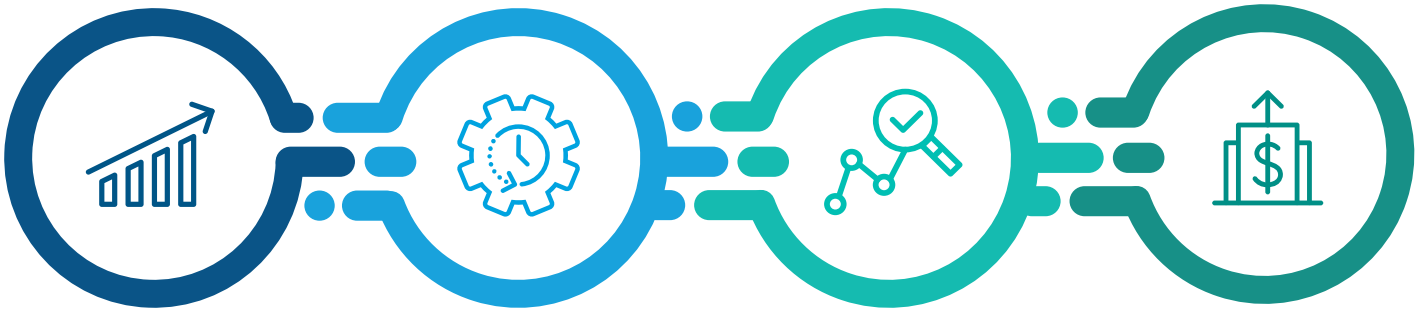
Improves site experience and supports patient outcomes

Rapid delivery of studies and protocol amendment

Intelligent supply optimization and flexible across platforms

Accelerates standard, decentrialized, hybrid or virtual trials

Delivering the right drug to the right patient at the right time has never been as easy or efficient



$2M
Overage savings[1]
Implementing optimized study specific forecasting solutions for comparator saved over $2M.

75%
Reduction in shipments[2]
Reduced overage of shipments by using automated resupply optimization tool.

$3M
Savings in shippings[3]
Saved over $3M in shipment costs in 2019 compared to 2018.

$5M
Total savings[4]
Reduced waste on comparator by 60% Reduced shipments by 20% Overall savings over $5M .

1. IQVIA internal, overall study savings for comparator 2. IQVIA internal, YOY reduction in shipments for large, multinational trial 3. IQVIA internal, YOY shipment cost savings for large, multinational trial 4. IQVIA internal, total combined sponsor savings on 2 large phase 3 trials

IQVIA IRT supports patients, sites and sponsors throughout the clinical study

For patients

- Ensures the right drug gets to the right patient at the right time
- Technology speeds drug dispensation and reconciliation on-site and remotely
- Electronic drug accountability assists monitoring of patient compliance

For sites

- Single sign-on with user-friendly interface and shared view for data and reporting with IQVIA patient solutions
- Integrated and shared workflows help sites perform faster transactions, leaving more time for patients
- Easy, turn-key integrations with 400+ trial support technologies and vendors, reduces administrative burden

For sponsors

- Self-service tools that facilitate trial activities, enhancing sponsor insights and improving communications with sites
- Rapid study design process reduces time to first-patientin and improves quality with design prototyping and templates
- Intelligent tools and technology to optimize drug forecasting and resupply
- Data analysis and insights against key study metrics

Discover how IQVIA Technologies IQVIA IRT can help optimize clinical trials through advanced technologies. Visit IQVIA.com/OCT to learn more.

IQVIA IRT is part of IQVIA Technologies Digital Patient Suite within the Orchestrated Clinical Trials (OCT) platform.

IQVIA TECHNOLOGIES

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