

Transforming trial execution

for faster, better decisions



**>95
studies**
using risk-based
monitoring

As much as
**25% cost
reduction**
over traditional
trial execution
approaches

A proven, data-driven approach for optimal results

Risk-based monitoring (RBM) has become a cornerstone of innovative trial execution. Supported by regulators, this approach can deliver greater efficiency while upholding data quality and patient safety.

With Quintiles, a leader in the RBM evolution, you can be confident you're taking the right approach to drive better execution throughout your trial. Our expanded RBM solution, Data-driven Trial Execution, leverages enhanced data review processes and timely insights from our breakthrough technology, Quintiles Infosario® Clinical, to provide holistic, near-real-time views of your patient, study, site and program-level data.

The result? You can proactively mitigate risk and allocate monitoring resources when and where you need them. This helps you reach smarter, faster trial decisions, cut trial timelines and experience as much as 25 percent cost reduction over traditional trial execution approaches.

Combining operational excellence with experience

Streamline trial execution across the globe with a data-driven approach informed by our experience using RBM in more than 95 studies across more than 20,000 sites and with more than 250,000 patients.

- Comprehensive risk assessment, performed upfront and then reviewed regularly to identify and mitigate scientific and operational risk
- Dynamic monitoring strategies, tailored to achieve the right balance of on-site and remote monitoring for both your entire study and individual sites
- Ongoing data surveillance, including review by medically trained staff, to help protect study integrity and patient safety through visibility into holistic trial data

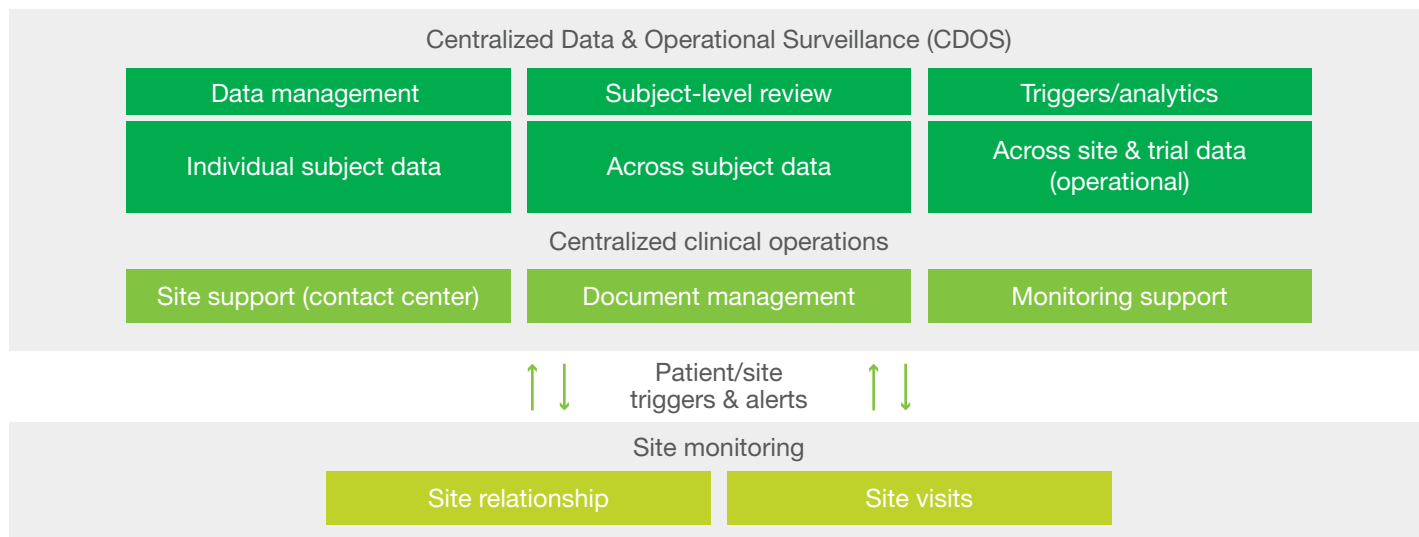
To date, this approach has resulted in customers experiencing faster query resolution, reduced data entry and source data verification backlog, and improved data quality.

See your data at a deeper level

For greater transparency into potential risks and more proactive trial management, Infosario Clinical provides access to crucial data insights. This robust technology platform integrates multiple data sources – including labs, CTMS, resource management, third-party data and EDC – to deliver timely alerts and triggers so you can continuously reassess risk and take corrective action as needed.

Integrating data for improved quality

Our centralized data surveillance and operational model



The right specialists for the right insights

How do you make analysis-ready data available faster than ever? With near-real-time knowledge from monitors and advanced analytics for ongoing, actionable insights throughout trial execution.

Using dashboards to visualize on-demand subject, site and study data, our specialized teams clean data following a strategy that supports your risk assessment goals by prioritizing the right subjects at the right time. Our medically trained reviewers can efficiently help ensure subject data are medically congruent and sound within and across subjects, helping to improve patient safety.

We then perform trend review and data analytics, comparing day-to-day safety and operational triggers across sites and countries, so our medical monitors and clinical data scientists can help optimize trial performance and support your monitoring strategy. To enhance efficiency, our centralized operational team helps resolve queries faster and supports project teams globally for improved clinical execution and site communications. Your on-site CRAs can then focus on strategic issues, such as patient recruitment.

Start off strong to speed execution

Integrating our Right Start solution with Data-driven Trial Execution can help you reduce trial delays, leading to faster time to market. When using our optimized feasibility, site selection and patient recruitment procedures, you can achieve up to 30 percent faster start-up timelines in certain therapeutic areas and as much as 45 percent faster investigator contracting timelines.

Chart a new path forward with confidence

Your risk-based monitoring strategy needs to do more than manage risk – it must also improve patient safety and quality while reducing costs. Customers are seeing these results today. Contact us to learn how Data-driven Trial Execution can deliver more for you.

**30%
faster**

start-up for certain
therapeutic areas

Up to
**50%
higher**

recruitment rates
with Partner Sites



Contact us

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