Medrio CTMS

powered by Engility®

Clinical trial management built for scale

Medrio CTMS, powered by Engility®, gives sponsors and CROs a single system of record for study planning, site management, and trial monitoring. You can centralize data, strengthen governance, and maintain audit-readiness from first patient in through study close.

Regulatory-Ready Operations

Demonstrate GCP compliance with audit-ready records for any due diligence review.

Real-Time Oversight

Track enrollment, milestones, and quality metrics to identify risks before they impact your timeline.

Rapid Study Deployment

Go live in weeks, not months, with a pre-configured system for agility and ease of use.

Request a demo **medrio.com/request-demo/**

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Centralize Clinical Operations

Fragmented data kills productivity and raises risk. Medrio CTMS centralizes site, investigator, study, and monitoring information in one structured system of record. Establish a documented operational foundation that supports audit readiness and future development programs.

Scale at Any Stage

Built for emerging life sciences sponsors and CROs, Medrio CTMS scales with your portfolio. Avoid the complexity of legacy enterprise systems while gaining the features you need, including automated monitoring, visit reports, and milestone tracking.

Ensure Compliance

Remove compliance gaps and stay inspection-ready, with no manual reconciliation required. Medrio CTMS connects directly to Medrio EDC and eTMF—automatically linking operational data to study documentation.

"Our journey as a startup taught us how to spot a reliable partner. It's not about luck or magic – it's about trust, plain and simple."

VP Global Clinical Affairs

Scan the QR code or visit
medrio.com/request-demo/