ARC360®

Your Study Submissions Copilot

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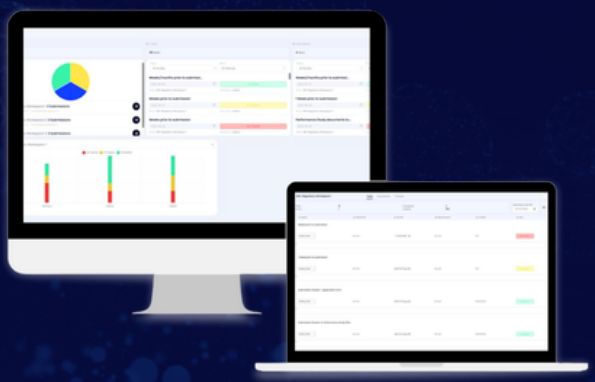

Clinical Trials Miss Timelines.

Regulatory Complexity Is Often the Hidden Cause.

- Do your regulatory teams lack bandwidth?
- Are your clinical trials consistently missing planned timelines?
- Is regulatory start up difficult?
- Is first patient often delayed due to approval or documentation issues?
- Do teams spend excessive time monitoring regulatory updates?

ARC360®

Managing Complex Requirements For Predictable Trial Approvals

- ✅ Faster study start-up
- ✅ Real Time Submission Status
- ✅ Reduced regulatory risk
- ✅ Greater confidence in delivery timelines

ARC360 protects your clinical trial investment by bringing clarity, control, and intelligence to regulatory execution across global study approvals.

Why Delays Happen

- Evolving global regulatory requirements
- Manual monitoring of regulatory updates
- Regulators' requests for information extending timelines
- High volume of complex submission documents to be drafted
- Fragmented coordination across sponsors, investigators and CROs

How ARC360 Accelerates Clinical Trials

Intelligent Regulatory Monitoring
AI detects changes. Experts stay in control.

Secure, Tailored Collaboration
The right people. The right access. The right tasks.

Real-Time Submission Status
Full visibility from kick-off to approval.

Instant Reporting & Oversight
Less admin. Improved Insights & Stakeholder Confidence.

AI-Accelerated Documentation
80+ hours reduced to seconds, reviewed by experts.

The Result

- Accelerated time to First Patient In
- Fewer unexpected regulatory delays
- Reduced manual effort
- Simplified stakeholder management & communication

PARTNER WITH ARC360®

Sign up for a demo today

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