



ARC360®

Your Study Submissions Copilot

www.arc-regulatory.com



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.

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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From Complexity to Clarity

ARC360® is now powered by AURA ai - a trusted AI platform that turns regulatory complexity into actionable intelligence.

ARC360® Powered by AURA ai combines the speed of AI with trust of human expertise.



ARC360® Powered by AURA ai Capabilities:



Regulatory Monitoring

Proactive monitoring of global regulatory changes, ensuring you stay compliant and informed.



eQMS System Integration

Embed regulatory intelligence directly into quality workflows for audit-ready, compliant processes.



Consistent & Accurate

Validated AI delivers traceable, auditable, and repeatable outputs you can trust every time.



Document Generation

80+ hours reduced to minutes -
Generate high-quality draft submission documents ready for review.



Human-in-the-Loop

AI-powered efficiency with human-in-the-loop validation for trusted, reliable results.



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