How Sponsors Can Manage Risk Across Clinical Development

By pairing comprehensive protocol and site performance data throughout the clinical development process, sponsors can mitigate potential enrollment, retention, and timeline risks. Implementing these insights early on in the study design process means less reactivity to potential roadblocks, and more proactivity in planning to avoid them altogether.

Instead of:	Sponsors can:	Outcome:
Adding operational complexity with potentially unnecessary endpoints or procedures.	Create optimized designs for answering scientific questions and meeting regulatory standards.	Sponsors create a better experience for sites and participants, without sacrificing scientific soundness.
Identifying sites based on previous relationships (with your organization or via your CRO).	Leverage high-quality site performance data to identify sites likely to excel at your specific trial design.	Sponsors can feel confident building relationships with new sites — including those in new markets — to help meet recruitment goals and timelines.
Relying on sites' own enrollment estimates to forecast costs and timelines.	Look to high-quality site performance data to more accurately assess site feasibility.	Sponsors can create more accurate forecasts, and identify opportunities to improve site performance from the outset, before lagging performance impacts the trial.

ClinSphere Trial IntelX™ was built specifically for bringing together protocol data, site performance benchmarks, and outcomes history to help study teams design feasible trials, select high-performing sites, and forecast timelines with confidence.

Ready to see what your trial could look like before it starts?
Visit wcgclinical.com/knowing to learn more and schedule a demo.

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