

5 reasons to trust Clario with your eCOA Rescue Study

When the eCOA solution in your clinical trial is at risk, consider Clario's experience in resetting studies across therapeutic areas and indications. Here are some qualities of our **committed approach to helping studies revive and thrive**.

01 Unique rescue methodology

Clario has developed a Playbook to rapidly engage with and manage each study, including:



This approach incorporates proven best practices from prior experiences. Learn about our rigorous model to **build a tailored plan for your study**.



02 Agile team focused on your study

We assign a project manager, solutions design analyst, technical implementation manager and scientist pre-award. This 'triage team' **responds within 24 hours** to arrange a current-state assessment. And we are **prepared for the changes and fast-moving timelines** associated with challenging trials.



24 hours

03 A trusted partnership

Open and honest dialogue about every dimension of the trial turnaround.



Listen to our sponsor and CRO customers and partner with them to renew studies.



Anticipate possible obstacles and address every data, device and delivery inquiry with customer stakeholders.



04 Scientific and technical expertise



therapeutic areas



protocol design



site management



patient support

Our top priority is to deliver rich and compliant endpoint evidence to complete trials successfully. Specialists rally to support all phases of a study, the specific indication and the eCOA instruments used. For instance, a Clario data manager ensures the quality and regulatory readiness of patient endpoint data.

Learn how Clario works closely with sponsors and Clinical Research Organizations (CROs) to get clinical trials back on track at www.clario.com/ecoarescue

