CLARIO.

Accelerate your study setup faster than ever before

Discover the major timesaving impact our latest tools, technology and enhancements can have in your overall study startup timelines.

Talk to a specialist

Accurate study data, captured right the first time

As a trusted eCOA provider to over **4,000 studies**, our experienced scientists, physicians and statisticians ensure your trial includes only the most suitable outcome assessments and data collection methodologies, resulting in:

- High-quality data that leads to operational efficiencies and reduced sample sizes for a shorter trial duration overall
- Regulatory-compliant, endpoint data that can be collected remotely or at study visits
- Patient-centric solutions that drive engagement and retention

Helping you meet IRB submission and FPI dates

In response to the tighter timelines and increasing complexity many trials face today, we've identified key areas to accelerate for the most impact in the study setup process.

Our Science and Operational teams can advise how your individual trial can:

Move quickly between study phases

Reduce eCOA setup times

Potentially avoid a lengthy TQT study

Access expertise across therapeutic areas, endpoint technologies, regulatory, safety and patient engagement throughout the duration of your trial. Our world-renowned Science teams:

- Are involved in every study from clinical development to close out.
- Work with sponsors and sites to oversee qualitative and quantitative endpoints so high-quality data can be collected for eligibility, safety and efficacy studies.
- Provide expert independent review, data analysis and maximum trial efficiency in support of regulatory approval for drugs, therapies and medical devices.
- Incorporate the latest developments in clinical research and regulatory compliance and contribute to the advancement of industry outcome measures.
- Provide strategies and best practices for instrument design that improve the quality of data.
- Collaborate with Clario's experienced logistics teams and local logistics hubs in the US, UK, EU and APAC to keep your trial running smoothly.

Solutions you can trust

Successful trials demand proven solutions that deliver confidence at each pivotal moment.

1,000+

compounds the number of potential medicines tested using Clario remote data collection technologies

70%

of all FDA drug approvals between 2019-2020 came from Clario-supported studies

50 years

of clinical, global regulatory and therapeutic experience

19k

clinical trials spanning more than five million patients in 120 countries

24/7

local support troubleshoots and finds answers to all your questions while reducing site and management burden

Explore how we can help your trial accelerate its study setup

Talk to a specialist