#### 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<br>between study<br>phases | Access expertise across therapeutic areas, endpoint technologies, regulatory, safety and patient engagement throughout the duration of your trial. Our world-renowned Science teams: <ul> <li>Are involved in every study from clinical development to close out.</li> </ul>                |
|---|---|
| Reduce eCOA<br>setup times              | <ul> <li>Work with sponsors and sites to oversee qualitative and quantitative endpoints so high-quality data can be collected for eligibility, safety and efficacy studies.</li> <li>Provide expert independent review, data analysis and maximum trial efficiency in support of</li> </ul> |

#### Potentially avoid a lengthy TQT study

- regulatory approval for allogs, theraples 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 Clariosupported studies

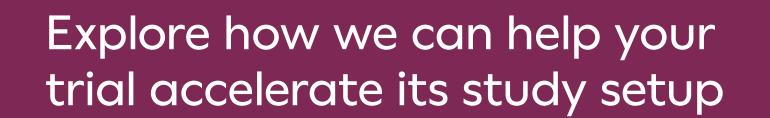
## 50 years

of clinical, global regulatory and therapeutic experience





clinical trials spanning more than five million patients in 120 countries local support troubleshoots and finds answers to all your questions while reducing site and management burden



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