Clario eCOA

Helping sponsors capture the right data to prove drug efficacy
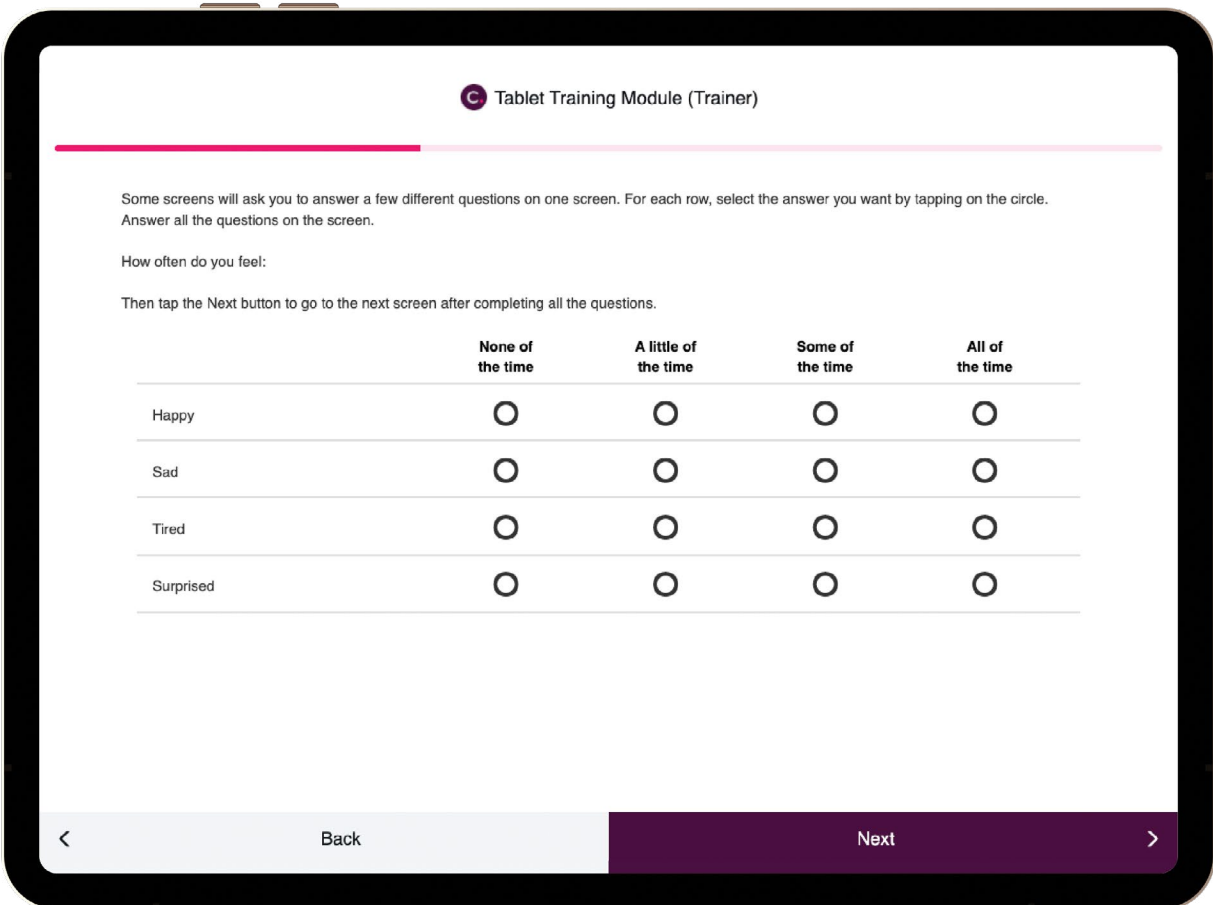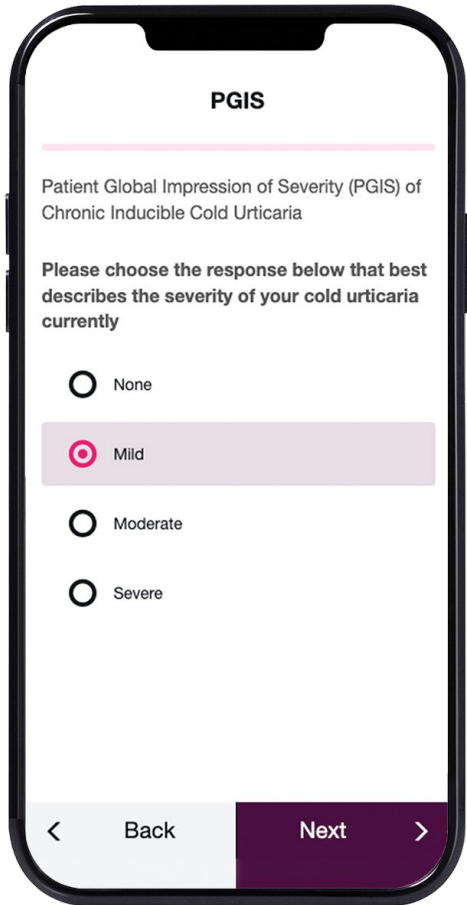
CLARIO.

We partner with sponsors and CROs to ensure their trials are measuring the endpoints they need to prove drug efficacy.

eCOA solutions

Our eCOA solutions make participant data collection simple and fast and help bring drugs to market, faster.


Poor endpoint design or strategy can contribute to trial delays or even failure.

Connected devices

Connected devices make taking part in clinical studies straightforward. Clario's devices are pre-validated for use in clinical trials and integrated with the eCOA device to capture ECG, spirometry, blood glucose levels, activity levels, movement endpoints and many more.
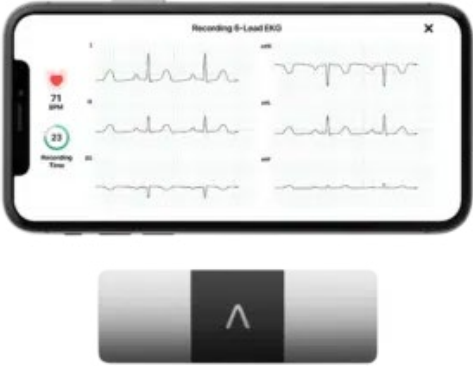
Alivecor KardiaMobile6L

iSpiro®
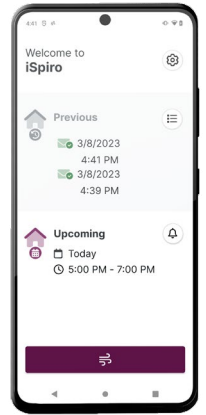
Actigraph Watch

Glucometer

Opal® Precision Motion Monitoring

01 Capture the right endpoints to prove the efficacy of your drug.

02 Quickly understand if your drug is working.

03 Differentiate your compound.

Leading endpoint technology platform for pharma R&D

Data sets, technology and domain expertise deeply embedded in customer workflows

Domain expertise

- **27,000** clinical trials supported
- **200+** therapeutic and endpoint experts
- Involvement in **60%** of all FDA approvals between 2019 and 2022

Data science

- Leading endpoint data solutions spanning medical imaging, eCOA, cardiac safety and respiratory efficacy and safety
- Generation of the richest clinical evidence

Technology solutions

- Broadest endpoint technology and service platform
- Backed by validated clinical trial management software, connected devices and artificial intelligence

CLARIO.

Medical Imaging
- >250 regulatory approvals
- >50K sites supported
- >30 years working with regulators

Respiratory
- >12K respiratory trials
- >600K participants
- >90% new respiratory approvals

eCOA
- ~4K studies supported
- >260K sites supported
- >2.5M participants supported

Cardiac Safety
- >13K cardiac safety trials
- >600 QT endpoint studies
- >580 new drug approvals

01 Scientific expertise

02 Evidence generation

03 Accelerated timelines

04 Improved workflows

05 Enhanced safety

06 Global regulatory expertise