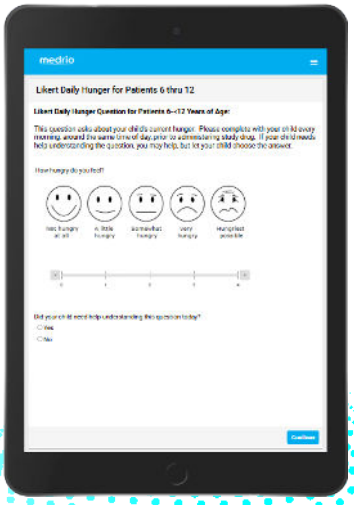
SOLUTION SHEET

Medrio eCOA/ePRO

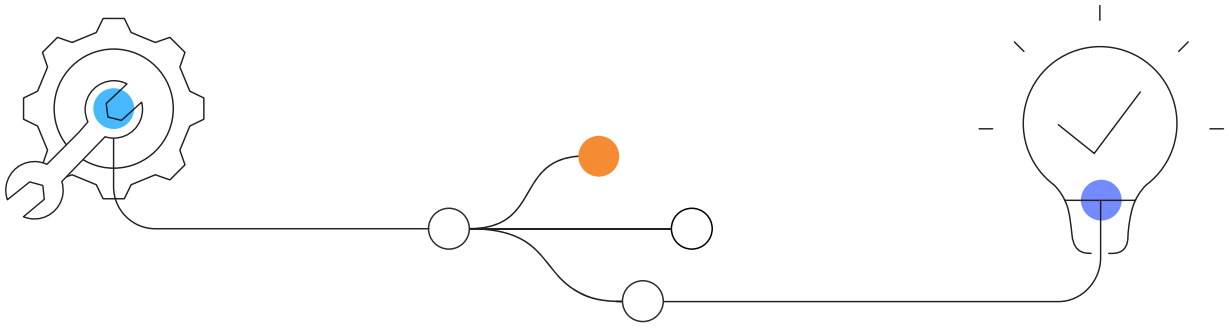
Improve data quality, engagement, and retention with a cost-effective and patient-centric solution.



With a user-friendly interface and flexible data capture options, Medrio improves participant engagement and compliance rates, so you can focus on collecting high-quality data for regulators and payers. Our web-based solution empowers participants, clinicians, and observers to conveniently and accurately report outcomes, enhancing data quality and study efficiency.

No more navigating to different software systems to view data. Medrio eCOA/ePRO automatically integrates with Medrio CDMS/EDC, allowing users to easily view all COAs in one place.

To support complex protocols, Medrio offers the flexibility of in-clinic and remote data collection, supporting both validated and home-grown surveys, as well as episodic collection. Our bring-your-own-device (BYOD) approach ensures your participants can respond to surveys anywhere at any time, with automated notifications boosting compliance, retention, and engagement.



Configurable

No custom coding or programming needed. Control your trials with real-time data visibility.

Flexible

Built-in functionality supporting remote or in-clinic workflows and quick mid-study changes.

Supportive

Unrivaled support and access to product and industry experts.

Patient-centric experiences

Make participation easy with a web-based, device-agnostic solution that centers the patient experience. An easy-to-use interface, notifications, and the MyMedrio participant portal improve participant compliance and retention while ensuring higher-quality data.

Accelerated study startup

Medrio eCOA/ePRO can be set up in as little as four weeks, accelerating the time it takes to begin capturing data. Quick, intuitive survey builds ensure you meet study timelines, ultimately reducing overall costs. Meanwhile, robust support helps you adhere to study protocols and industry standards.

Accessible data in real time

With zero delays between data collection and database population, you can view data instantly and make critical decisions in real-time. Choose Medrio to better manage participant safety and trial oversight.

Ready to learn more?

> *"A lot of people at these companies have successfully used paper in prior lives, and they focus on price not cost. But the right solution pays for itself. Don't think just about the cost of adding ePRO. As a CRO, more errors in the data may equate to a change order later to complete the work reconciling paper diaries and surveys."*
>
> – **COO,** OncoBay

CONFIGURATION

- Branching and skip logic
- Immediate notifications of participant-reported events
- Non-email usernames
- Customizable look-and-feel
- Language localization

SCHEDULING

- Activation based on consent status or date
- Customizable assessment prompts and reminders
- Adjustable preferred time zone for remote participants

CONDUCT

- Email and SMS notifications
- Automatic form save after each data entry
- MyMedrio participant portal

DATA REVIEW

- 21 CFR Part 11 compliant audit trail
- Custom reports
- Real-time data access

medrio