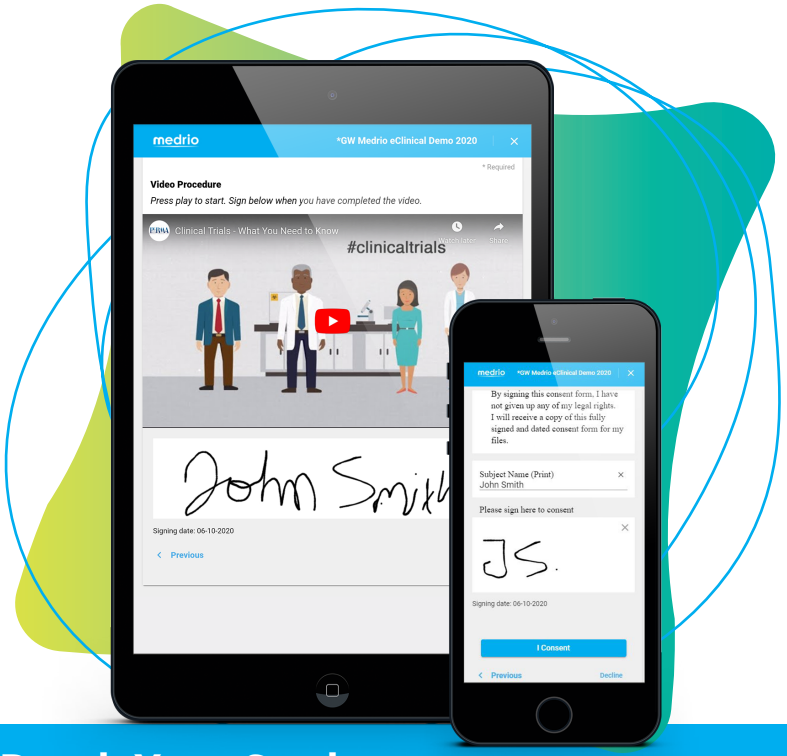
SOLUTION SHEET

Medrio eConsent

Flexible consent technology improves participant comprehension and process oversight.



The Consent Process Can Make or Break Your Study

85% fail to retain enough patients

30% of patients drop out of studies

8% of those patients drop out due to a failure to understand study requirements

Medrio's Consent solution empowers organizations to accelerate all aspects of the consent process,

from setting up and modifying forms to ensuring patient comprehension, while remaining in full regulatory compliance.

Our flexible solution supports both electronic and paper-based processes, allowing you to prioritize the patient experience and the sites' needs.

Sites may either consent participants in-clinic using tablet-based workflows or remotely using email and MyMedrio, our secure participant portal.

medrio

You've Got Options

Unlike other Consent technology solutions, our fully integrated web-based Consent and EDC platform allows you to meet your study milestones by streamlining implementation and data collection. Multimedia presentations and participant quizzes enrich patient understanding. Plus, when time is of the essence, you can benefit from quick set-up options, including uploading IRB-approved paper consent documents.

Improve Patient Experience

- Medrio's BYOD model allows patients to focus on understanding the trial requirements, not on learning how to use a new device
- Leverage in-clinic and remote options to consent patients where they're most comfortable
- Improve comprehension by incorporating videos, FAQs, and quizzes

Accelerate Setup and Changes

- Upload consent forms to a tablet app with instant site access
- Control consent version by site easily with programmer-less configuration options
- Expedite re-consenting workflows with do-it-yourself access

Improved Compliance and Oversight

- Accelerate oversight with real-time access to consent progress data
- Centralize data into a single system while supporting either electronic and/or paper-based processes
- Protect patients' PHI using encryption and granular permissions that meet or exceed global compliance regulations

Less than 5% of clinical trials use eConsent today, but we expect rapid growth in the next three years. Medrio can help you take a step away from paper.

ABOUT MEDRIO

At Medrio, we know it takes a global village to achieve a disease-free world. Since 2005, we've developed a successful ecosystem of visionary people like you who want to make the world a healthier place - our employees, customers, and partners alike. We've supported Sponsors and CROs across the life sciences spectrum, in all therapeutic areas and trial phases, to achieve critical breakthroughs and secure more than 375 regulatory approvals. Our innovative and intuitive technology solutions support your teams and sites, while reducing trial participation burden. While other eClinical vendors make these same claims, few can deliver the high-touch, white-glove customer service that makes us your most successful strategic partner and solves your most pressing challenges. To learn more, visit us at medrio.com.

 | +1 800.498.6830 | info@medrio.com | www.medrio.com