Clinical Trial Consent Solution

Transform Your Consent Process

Interlace Health's Clinical Trial Consent Solution provides the digital platform to collect and store consent of clinical trial participants and serves as a portal to access vital educational content needed to understand trial requirements.

This new way of communicating with participants and documenting engagement reduces participant drop-outs and critical study delays.

of trial participants drop out due to lack of understanding

of studies take 2x longer than planned

*Sources: NCBI, 2013. Tufts CSDD, 2012.

Achieve Your Goals

Our solution meets regulatory requirements and improves:

- Engagement with patients
- Education & understanding of participants
- Timeliness of participant onboarding
- Data quality
- Staff workflows by reducing paper burdens

Raise the bar with eConsents

- Instant form assignment
- Faster form completion
- Trial-specific content
- Enhanced education & understanding
- Virtual participant signing
- Prompt review & action
- Robust dashboard management
- Immediate archiving

Why it's Better

Interlace Health's Clinical Trial Consent solution integrates seamlessly with your EHR and clinical trial software and is customized to meet your remote or in-person consenting needs:

- Content flexibility
- Embedded educational links
- Ability to ask questions
- Remote consenting & re-consenting
- Version management
- Audit trails
- Role-based access & security
- Dashboard status management
- Meets regulatory requirements

Measure the Value

Interlace Health clients who have implemented electronic consent have experienced significant time savings.

10+ minutes saved per consent completed

2-3 days faster archiving per consent

For more information on how an electronic consenting process can benefit your organization, and a customized demo, visit **interlacehealth.com/contact**

INTERLACE HEALTH | Making healthcare *better* one process at a time.