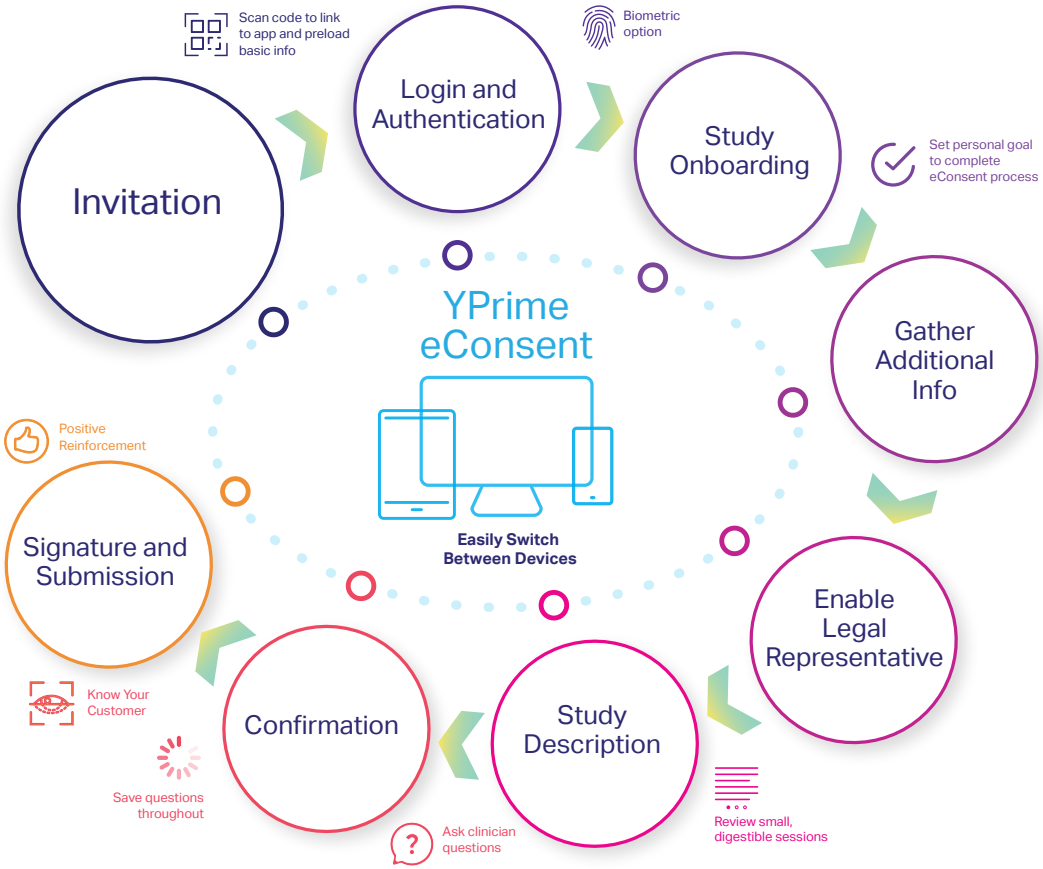
eConsent

Empower patients to make informed decisions about participating in clinical trials

Electronic informed consent (eConsent) is a powerful tool for improving patients' understanding of increasingly complex and technical clinical study designs, objectives, and requirements. Technology enables the industry to present information according to the patient's preferred method, media, and pace.

YPrime's eConsent platform and patient-centered approach ensures better comprehension of complex information, accelerates enrollment and screening, and improves patient engagement. We empower the patient throughout the eConsent process because we have taken the entire patient journey into consideration as we developed this platform.



YPrime's flexible approach means our eConsent platform can be used as a standalone service, or sponsors and sites that use other YPrime technology solutions will discover this eConsent technology integrates into their existing workflow and they will not need to learn a new workflow or system. Single sign-on convenience means sponsors and sites will not have to log into multiple systems to access eConsent and eCOA platforms.

Our Platform

- All data is encrypted in transit and at rest
- All activity is logged and time stamped with every version change stored for future compliance needs and/or FDA audits
- Platform is 21 CRF Part 11 and NIST-800-53 compliant
- Supports multimedia and can supply user-specific graphics and videos in multiple languages

Features

For Patients

- Access the eConsent form through the YPrime App on the device that's most convenient for them—mobile phone, tablet/laptop, desktop. They may begin the process on one device and continue on another.
- Enter, review, and save questions for the clinicians which can be followed up via messaging, telephone call, or in person
- Review and sign consent forms at their own pace and where they are most comfortable

For Sponsors and Sites

- Available as a standalone service
- If used with other YPrime technology solutions, seamlessly fits into the existing workflow
- Single sign-on technology eliminates multiple system logins for other platforms integrated into the app environment (eConsent and eCOA)
- Data sync eliminates need to restrict patients to specific devices
- Automatically generate all user screenshots for IRB review
- Sponsor, site, and study team receive real-time, permissions-based data
- eConsent workflow is streamlined when there are multiple members of the care team who need to sign off