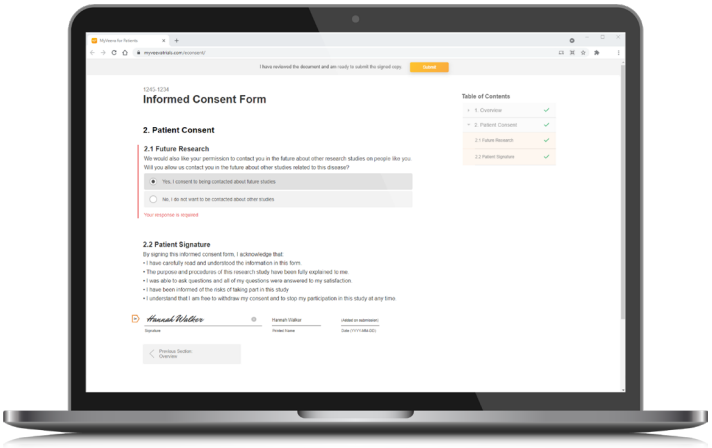
Veeva SiteVault eConsent

Overview

Veeva eConsent makes patient participation in clinical trials easier, while reducing administrative burden for study teams and ensuring compliance for sites of all sizes.

Veeva eConsent is unified within SiteVault to make the consent process efficient for study teams, and includes MyVeeva, an easy-to-use, patient-centric application to foster patient engagement.

MyVeeva for Patients offers unique flexibility for both in-person and remote consenting and empowers patients to sign and access their consent forms right from their own device.



Benefits

Better Patient Experience

Eliminate the need for patients to carry around paper and provide convenient access to study documents and site information on their own device.

Reduce Staff Burden

Eliminate printing and copying, simplify screening, and easily access documents with a single system that can be used across all studies and works seamlessly.

Faster Study Execution

Accelerate consent completion and simplify monitoring by eliminating paper processes. Enabling study teams, patients and monitors to complete consenting activities, whether onsite or remote.

Stronger Compliance

Reduce errors related to signature dates and times, stay informed with complete visibility into consent status, and automatically generate screening logs to streamline consent monitoring.

Features

Flexible Consent Options

Reach patients where they are with options for in-person or remote eConsent on any device.

Simple and Fast Consent Set Up

Transform your IRB-Approved PDFs participant-ready eConsents in minutes.

Intuitive Patient Experience

Never miss a signature with an easy-to-navigate layout for patients and a guided review to ensure all sections are reviewed prior to sign-off.

Flexible eConsent Templates

Set up your eConsent once and use it for any study participant. eConsent is built to accommodate one or more signatories with ease.

Reporting

Gain full visibility of patient consent status, date, and version gives sites and monitors the vital information needed to support compliance.

Sharing and Collaboration

Easily share and collaborate on informed consent forms between sponsors and sites.

Validated and Secure

Eliminate the burden of validation and gain peace of mind. Veeva eConsent is fully validated by Veeva and supports compliance with HIPAA and regional data privacy requirements.

Version and Audit Controls

Automate versioning and view date/time stamps for better compliance and traceability. Easily compare documents to previous versions to see what has changed.

Compliant eSignatures

Capture patients and study staff signatures in full compliance with 21 CFR Part 11.

Scan to register your site for Veeva SiteVault