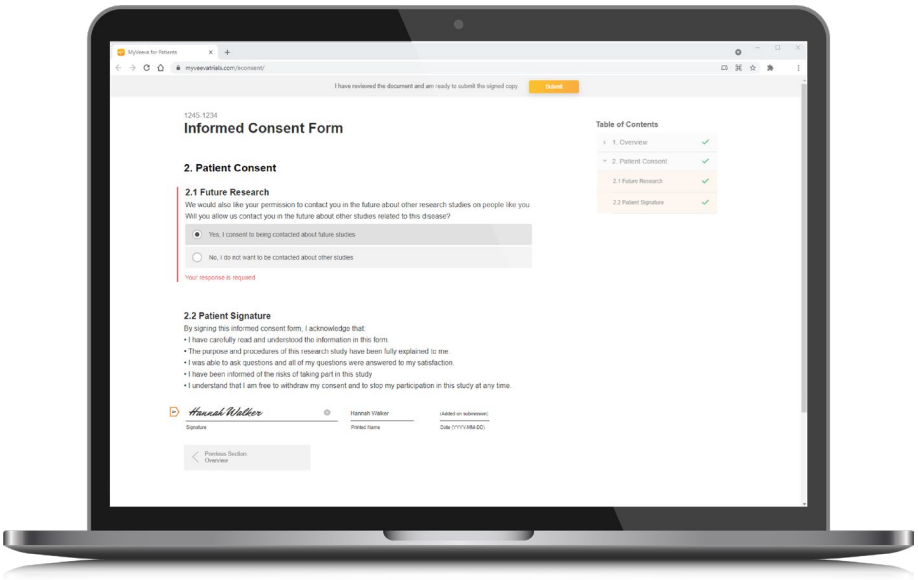
Veeva eConsent

Overview

Veeva eConsent simplifies the set-up, completion, and review of consent, reducing administrative burden and ensuring compliance for sites and study teams.

This enables study teams to make patient participation in clinical trials easier through a user-friendly application to access key information and complete consent electronically.



Benefits

Better Patient Experience
Give patients direct and convenient access to their study documents on their own device via the MyVeeva for Patients app.

Reduce Site Burden
Remove administrative burden by using one eConsent system across all studies without the need for additional systems or applications.

Faster Study Execution
Accelerate informed consent form (ICF) creation and review by eliminating burdensome manual processes.

Stronger Compliance
Stay informed with complete visibility into consent status, streamlining consent monitoring and ensuring study compliance.

Features

End-to-End eConsent Platform

Unified system seamlessly connecting sponsors, sites, and patients across the entire consent process.

Flexible Consent Options

Enable in-person or remote eConsent on any device. A signed copy of the ICF is stored in the patient's app and can be downloaded anytime.

Reduced Authoring and Review Cycles

Creation and reuse of consent templates through easy-to-use editor tools, streamlining authoring, review, and approval of consent documents.

Sharing and Collaboration

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

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.

Familiar Site Interface

Site friendly solution to manage eConsent, reducing administrative burden and training requirements.

Intuitive Patient Interface

Guide patients with an easy-to-navigate layout and ensure all sections are reviewed prior to sign off.

Interactive Content

Easily add images and videos to aid in comprehension. Add custom questions to collect additional information and enhance the consenting process.

Reporting

Full visibility of patient consent status, date, and version gives sponsor, monitor, and site staff the vital information needed to support compliance.

Validated and Secure Cloud Platform

Veeva eConsent is fully validated by Veeva and supports compliance with HIPAA and regional data privacy requirements.

myVeeva FOR PATIENTS

MyVeeva for Patients is a suite of solutions that make trial participation easier for patients and streamlines study execution for research sites and trial sponsors.

With capabilities for eConsent, ePRO, education and support, messaging, and visit management, MyVeeva for Patients simplifies the management of patient solutions through a connected clinical ecosystem that increases study team efficiency and reduces administrative burden for sites.