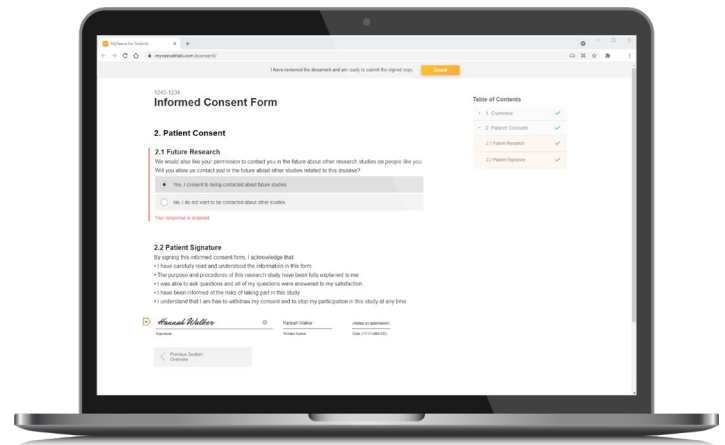# Veeva SiteVault eConsent

## Overview

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

Veeva SiteVault eConsent is unified with Veeva SiteVault eISF 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 Site 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 with Veeva SiteVault eISF.

### 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.

# Veeva eConsent

## Features

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

### 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.

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

### 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.

### Seamless Integration with Veeva SiteVault
Reduce duplicate data entry and errors with eConsent that is unified with your Veeva SiteVault eISF.

### Simple and Fast Consent Set Up
Transform your IRB-Approved PDFs participant-ready eConsents in minutes.

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

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

**Scan to register your site for Veeva SiteVault**

Learn more at sites.veeva.com