

Veeva eCOA

Veeva eCOA simplifies the design, management, and completion of eCOA for sponsors, sites, and patients.

With accelerated build times, automated workflows, and on-demand data access, it removes eCOA from the critical path to FPI and database lock.

The screenshot displays the Veeva eCOA interface. On the left, a 'Library' section contains a search bar and a list of surveys: 'Patient Global Assessment of Arthritis' (3 items), 'Daily Wellness Survey' (7 items), 'Weekly Injection Diary', 'Dermatology Life Quality Index', and 'Migraine Log'. The 'Daily Wellness Survey' is highlighted. On the right, a 'Rate your pain over the past 24 hours' scale is shown, featuring a row of six smiley faces (green, yellow, orange, red) and a numeric scale from 0 to 10. The number '2' is selected, corresponding to the second green smiley face. The scale is labeled 'No Pain' at 0 and 'Extreme Pain' at 10.

Library	
Survey Name	
Patient Global Assessment of Arthritis	3
Daily Wellness Survey	7
Weekly Injection Diary	
Dermatology Life Quality Index	
Migraine Log	

Rate your pain over the past 24 hours

0 1 2 3 4 5 6 7 8 9 10

No Pain Extreme Pain

Benefits

- **Build studies faster:** Centrally-controlled library of pre-validated and fully reusable eCOAs accelerates study design.
- **Streamline study management:** On-demand workflows, dashboards, and data exports increase study efficiency and control.
- **Optimize site experience:** Easily navigable workflows that automate tasks, eliminate duplicate data entry, and streamline device management.
- **Improve patient experience:** Single user-friendly app to access all study activities and make eCOA completion easier.

Features

Centrally-Controlled eCOA Library

Accelerate study design through a library of reusable and validated instruments, sourced from both Veeva and sponsor libraries.

Interactive Design Process

Engage stakeholders early with eCOA previews across all device types and available languages throughout the build process.

Automated Document Generation

Auto-generated screenshot documents for site review and IRB/ethics packs.

Self-Service Management Tools

Track study progress, make rapid mid-study changes, and export data on-demand, without extra cost or delay.

Version Control

Complete tracking and audit trails for instruments, translations, and amendments.

Patient and Site Notifications

Notify patients of ePROs requiring completion and alert sites if they are uncompleted or missed.

Guided Navigation

Indicators guide the sites through outstanding activities that need to be completed during patient visits.

Enhanced Login Options

Users can login with a username and password, PIN, or choose to activate biometric authentication (fingerprints and facial recognition).

Optimized for BYOD

Patients can use their own device including android, iOS, and web.

End-to-End Platform

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

Veeva Clinical Platform

Veeva Clinical Platform improves clinical research by providing the most complete and highest-quality solution built for the unique needs of patients, research sites, and trial sponsors. With seamless connection and data flow across all stakeholders, the Veeva Clinical Platform enables faster, more efficient trials that achieve higher data accuracy, and deliver a better experience for sites and patients.