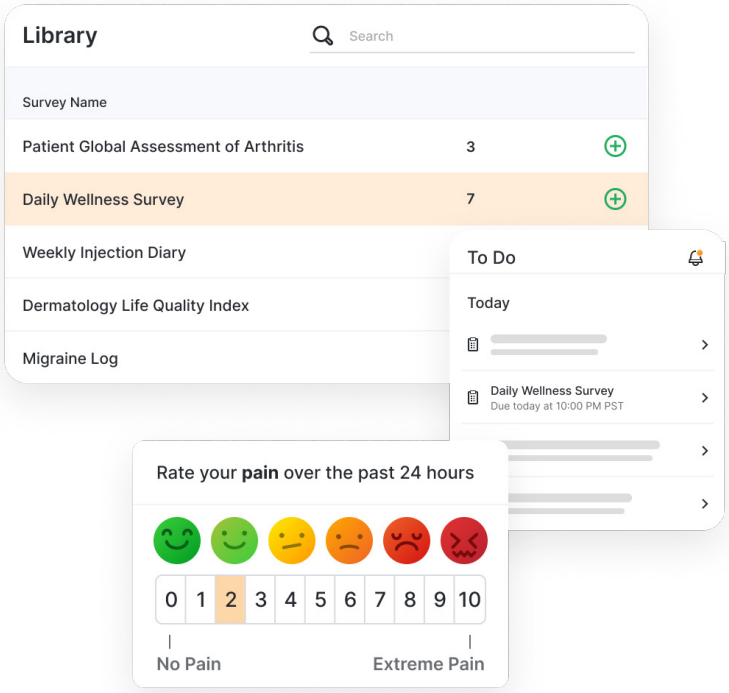
Veeva ePRO

Veeva ePRO simplifies the design, management, and completion of electronic Patient Reported Outcomes (ePRO), with seamless data flow that ensures access and transparency throughout clinical studies.

This delivers high quality patient data when you need it, where you need it through a modern platform built for the unique needs of sponsors, sites, and patients.



Benefits

- **Accelerate study design and execution.** Build, manage, and change studies faster with a modern designer and full reusability across studies.
- **Access high-quality data faster.** Ensure data is accurate, complete, and readily available from study design to close out.
- **Improve patient experience.** Make ePRO completion easier with a single user-friendly app to access all study activities.

Features

ePRO Library

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

Supports Range of Question Types

Incorporate required question options including multiple choice, numerically rated scales (NRS), and visual analogue scales (VAS).

Branching Logic

Only include relevant questions to simplify ePRO completion and enhance patient experience.

Patient and Site Notifications

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

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.

Real-time Data Access

Sites and study teams can access patient-entered data, including adherence, in real-time in the systems they work within – no more data silos.

Amendments and Version Control

Seamlessly manage amendments to the study design which are version controlled and automatically provided to sites.

End-to-end Platform

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

Validated and Secure Cloud Platform

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

myVeeva FOR PATIENTS

MyVeeva for Patients is a simple application that makes trial participation easier for patients and streamlines study execution for research sites and trial sponsors.

With capabilities for eConsent, ePRO, eClinRO, 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.