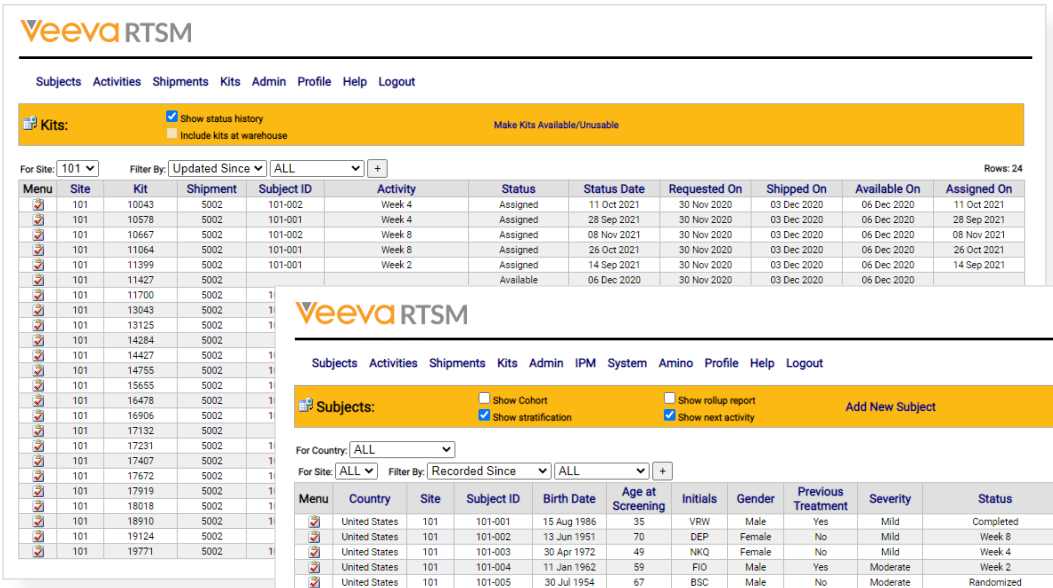
Veeva RTSM

Veeva RTSM is a robust, reliable, and user-friendly randomization and trial supply management solution designed to simplify complex processes and expedite clinical trials. It is modular and quick to configure empowering teams to implement the capabilities they need in fast startup timelines.

The extensive feature set allows Veeva RTSM to scale and support a broad spectrum of trial design needs across all therapeutic areas. Connections with Veeva clinical applications and integrations with third party solutions allow you to seamlessly incorporate Veeva RTSM into your clinical trial ecosystem.



Kit Tracking

Subject Tracking

Benefits

- **Everything you need and only what you need:** with our modular solution, you only use and pay for the functionality you need when you need it.
- **No compromises:** flexible configuration combined with appropriate customizations deliver the design you want, without traditional limitations.
- **Maximize supply & minimize costs:** minimize drug wastage and control shipment scheduling with adaptable supply management settings and expiry management.
- **Exceptional service delivery:** proactive project management and communication from Veeva RTSM experts guide you through every stage of your trial.

Randomization

Randomize patients with timeliness and accuracy.

- Multiple randomization schemas are available including: static, stratified (single or multi-variable), dynamic/minimization, forced, and adaptive.
- Re-randomize patients when needed.
- Easily configure eligibility, stratification, and other related questions to randomize subjects appropriately.
- Supports capping and can tightly control sentinel dosing.

Trial Supply Management

Optimize the supply management strategy for your trial's needs.

- Trigger site shipments according to trigger/resupply thresholds, predictive inventory control, just-in-time options, and more.
- Adapt supply settings during study execution, e.g. change resupply levels based on actual site enrollment numbers.
- Track all types of clinical trial supply materials with blinded and unblinded views, temperature management options, and connections with shipping providers.
- Supports complex patient dosing schedules including titration and dose escalation.
- Trace the entire kit lifecycle with our end-to-end drug accountability module.

A PROVEN TRACK RECORD

8,500+	37,000+	175+	60+	45+
sites	system users	trials	countries	languages

Additional Features

Tracking Subject Activities

Easily track subject activities including screening and enrolling subjects; randomization, drug assignments including titration management, completion, and follow-up; as well as unscheduled activities such as screen failure, early terminate/withdrawal, unscheduled resupply, skipped visits, and emergency unblinding.

Functionality for all trial types

Support a variety of study designs including crossover, sequencing, or extension studies, as well as multi-cohort designs and sub-study tracking.

Rapid Setup Processes

A modular and highly configurable architecture delivers faster study setup times (in as little as 3 to 4 weeks) and allows customizations to be built and deployed quickly and easily.

Easy Mid-study Updates

Reconfigure key settings during a study, such as activating a cohort or modifying limits on total subjects. Amendments requiring a system change are implemented rapidly by the Veeva RTSM services team--often in days rather than weeks.

Accessible Reporting

Filter information to generate targeted reports whenever needed. Review details on screen or export on demand to Excel and SAS.

User-friendly Interface

A simple, easy-to-use interface with role and permission-based access controls minimizes the clicks needed to complete common tasks and improves the accuracy of site data entry. It has been translated into 45+ languages to date.

Connectivity

Veeva RTSM can function seamlessly with different electronic data capture systems, reporting tools, and shipping providers. Connecting Veeva RTSM with Vault EDC and MyVeeva will optimize your data quality and trial efficiency by eliminating duplicate data entry and automating data flows.

Service Expertise

Veeva's RTSM services team has a deep understanding of RTSM best practices, system configuration, and validation, and is committed to customer success. Throughout trial execution, a project manager and 24x7x365 help desk will provide ongoing expertise and support.