

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.

Veeva RTSM
 Subjects Activities Shipments Kits Admin Profile Help Logout

Kits: Show status history Include kits at warehouse Make Kits Available/Unusable

For Site: 101 Filter By: Updated Since ALL

Menu	Site	Kit	Shipment	Subject ID	Activity	Status	Status Date	Requested On	Shipped On	Available On	Assigned On
	101	10643	5002	101-002	Week 4	Assigned	11 Oct 2021	30 Nov 2020	03 Dec 2020	06 Dec 2020	11 Oct 2021
	101	10578	5002	101-001	Week 4	Assigned	28 Sep 2021	30 Nov 2020	03 Dec 2020	06 Dec 2020	28 Sep 2021
	101	10667	5002	101-002	Week 8	Assigned	08 Nov 2021	30 Nov 2020	03 Dec 2020	06 Dec 2020	08 Nov 2021
	101	11064	5002	101-001	Week 8	Assigned	26 Oct 2021	30 Nov 2020	03 Dec 2020	06 Dec 2020	26 Oct 2021
	101	11399	5002	101-001	Week 2	Assigned	14 Sep 2021	30 Nov 2020	03 Dec 2020	06 Dec 2020	14 Sep 2021
	101	11427	5002			Available	06 Dec 2020	30 Nov 2020	03 Dec 2020	06 Dec 2020	

Kit Tracking

Veeva RTSM
 Subjects Activities Shipments Kits Admin IPM System Amino Profile Help Logout

Subjects: Show Cohort Show rollout report Show stratification Show next activity Add New Subject

For Country: ALL

For Site: ALL Filter By: Recorded Since ALL

Menu	Country	Site	Subject ID	Birth Date	Age at Screening	Initials	Gender	Previous Treatment	Severity	Status	Status Date	Next Activity	Target Date
	United States	101	101-001	15 Aug 1986	35	VRW	Male	Yes	Mild	Completed	23 Nov 2021	n/a	n/a
	United States	101	101-002	13 Jun 1951	70	DEP	Female	No	Mild	Week 8	08 Nov 2021	Week 10	22 Nov 2021
	United States	101	101-003	30 Apr 1972	49	NKQ	Female	No	Mild	Week 4	09 Nov 2021	Week 8	07 Dec 2021
	United States	101	101-004	11 Jan 1962	59	FIO	Male	Yes	Moderate	Week 2	15 Nov 2021	Week 4	29 Nov 2021
	United States	101	101-005	30 Jul 1954	67	BSC	Male	No	Moderate	Randomized	12 Nov 2021	Week 2	26 Nov 2021
	United States	101	101-006	27 May 2002	19	FCZ	Male	Yes	Severe	Randomized	19 Nov 2021	Week 2	03 Dec 2021
	United States	101	101-007	24 Mar 2002	19	LVA	Female			Screened	19 Nov 2021	Randomization	n/a
	United States	101	101-008	07 Oct 1937	84	FVC	Male			Screened	22 Nov 2021	Randomization	n/a
	Germany	102	102-001		40	IIS	Male	No	Mild	Completed	23 Nov 2021	n/a	n/a
	Germany	102	102-002		80	LMR	Female	Yes	Severe	Week 8	08 Nov 2021	Week 10	22 Nov 2021
	Germany	102	102-003		64	HRE	Female	Yes	Mild	Week 4	09 Nov 2021	Week 8	07 Dec 2021
	Germany	102	102-004		30	KNU	Male	Yes	Mild	Week 2	15 Nov 2021	Week 4	29 Nov 2021
	Germany	102	102-005		22	XAZ	Male	Yes	Moderate	Randomized	12 Nov 2021	Week 2	26 Nov 2021
	Germany	102	102-006		20	UGO	Male	No	Moderate	Randomized	19 Nov 2021	Week 2	03 Dec 2021
	Germany	102	102-007		86	LJL	Female			Screened	19 Nov 2021	Randomization	n/a
	Germany	102	102-008		19	DYR	Female			Screened	22 Nov 2021	Randomization	n/a

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+
sites

37,000+
system users

175+
trials

60+
countries

45+
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.