PRODUCT SHEET

Veeva RIM

Veeva RIM unifies regulatory systems and processes on a single cloud platform to enable end-to-end submission and registration management.

RIM applications share a common data model, which allows for regulatory business functions to run in one Vault.

Veeva Registrations plans, tracks, and reports on global health authority product registrations and associated changes.

Veeva Submissions is a content management application used to plan, author, review, and approve regulatory submissions.

Veeva Submissions Publishing produces compliant published submissions ready to send to global health authorities.

Veeva Submissions Archive provides storage, navigation, and search of submitted regulatory applications and related correspondence and questions.

PRODUCT	ANNOUNCED	STATUS	CUSTOMERS
Veeva Registrations	2015	Mature	100+
Veeva Submissions	2013	Very Mature	200+
Veeva Submissions Publishing	2017	Mature	51–100
Veeva Submissions Archive	2015	Mature	100+

PRODUCT SHEET

Veeva Registrations

Registrations allows sponsors to plan, track, and report on global product registrations along with health authority correspondence and commitments.

Events provide the ability to manage product changes, from the initial assessment of the proposed change through submission creation, health authority interactions, and final registration update. Label changes can be tracked and managed at both the global and local level. Registrations also produces compliant product data output (e.g., xEVMPD and IDMP) for EU regulations.

Dashboards and reports allow personnel to track the progression of change events and provide an understanding of product registration locale.

Announced	2015
Status	Mature
Customer type	Enterprise Pharma, Biotech, CDMO, MedTech, Consumer
Customers	100+
Platform	Veeva Vault
Integrations	Lives with Submissions, Submissions Publishing, Submissions Archive Connected with QMS

PRODUCT SHEET

Veeva Submissions

Submissions is the leading content management application used to plan, author, review, and approve regulatory documents. It provides full enterprise content management capabilities for creation, version control, approval, and real-time co-authoring for all submission-related documents. With content planning capabilities, users can build a submission outline and automatically match documents to the outline.

Clinical and non-clinical reports can be built and published using Report Level Content Plans. Dashboards and reports allow submission managers to track the status of each document in real time.

Announced	2013
Status	Very Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech, Consumer
Customers	200+
Platform	Veeva Vault
Integrations	Lives with Registrations, Submissions Publishing, Submissions Archive Connected with eTMF, PromoMats

PRODUCT SHEET

Veeva Submissions Publishing

Submissions Publishing generates electronic submissions for global health authorities. Veeva releases new templates and validation criteria to keep up with evolving regulations. Submissions Publishing leverages content plans created earlier in the lifecycle to start the publishing process as soon as individual documents are finalized.

Users can create internal and external hyperlinks to connect references in the text. They publish submissions directly to health authorities from Vault, in markets where allowed.

Dashboards and reports allow publishers to track each submission component as it progresses from authoring to completion.

Announced	2017
Status	Mature
Customer type	Enterprise Pharma, Biotech, CDMO, MedTech, Consumer
Customers	51–100
Platform	Veeva Vault
Integrations	Lives with Registrations, Submissions, Submissions Archive Requires Submissions, Submissions Archive

PRODUCT SHEET

Veeva Submissions Archive

Submissions Archive is a global, secure repository of submission published output. It functions as the authoritative source of applications submitted to health authorities.

Submissions Archive includes a viewer that supports all electronic and paper formats, with PDF link navigation provided for electronic formats. Users can view submissions and health authority correspondence alongside all previously submitted applications.

The embedded document viewer provides visibility into each document in the structure.

The Active Dossier feature displays the submission components that are currently active for any product / market combination.

Announced	2015
Status	Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech, Consumer
Customers	100+
Platform	Veeva Vault
Integrations	Lives with Registrations, Submissions, Submissions Publishing