

Veeva RIM for Animal Health

Veeva RIM helps animal health companies streamline regulatory processes, improve visibility into regulatory business objectives, and strengthen compliance with health authority requirements. Veeva RIM supports the regulatory needs of the animal health industry with its standard animal health data model, veterinary dossier formats, and much more. There are four applications within Veeva RIM:

- ✓ **Veeva Registrations** plans, tracks, and reports on global health authority product registrations and associated changes. The flexible data model enables compliance with evolving animal health regulatory requirements.
- ✓ **Veeva Submissions** allows teams to plan, author, review, and approve regulatory submissions. It leverages animal health specific standard document classification and dossier templates for the U.S. and E.U.
- ✓ **Veeva Submissions Publishing** provides continuous publishing, hyperlinking, table of contents edits, and merging to produce animal health specific published sequences. Users can submit published sequences directly from the platform, streamlining the process in markets where it is permitted.
- ✓ **Veeva Submissions Archive** provides easy storage, navigation, and search of submitted regulatory applications and related correspondence and questions.

Regulatory trends and challenges in the animal health industry

The animal health industry is experiencing a significant shift toward digitization in regulatory affairs. Companies are focused on data collection, protection, and integrity, as they revisit regulatory information management needs.

Legacy systems are often unable to support the dynamic regulatory landscape. They provide

limited visibility, which hinders collaboration across the organization and reduces the speed and quality of decision making. With new standards like the European Veterinary Medicinal Products Regulation, regulatory teams must find new ways to drive efficiency, while remaining compliant.

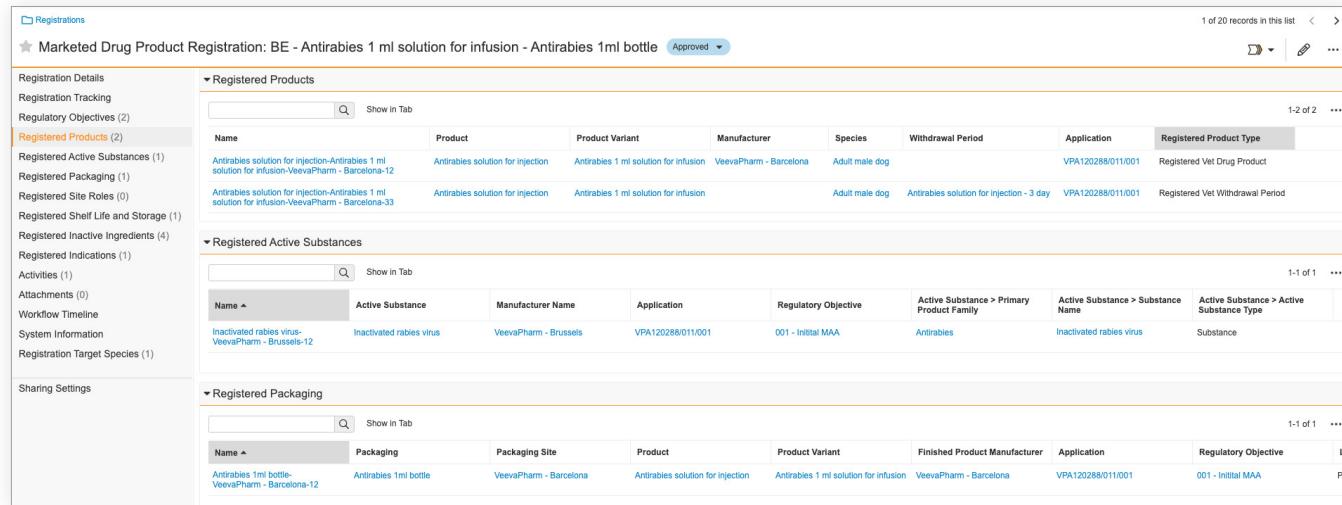


Vault has become our single source of truth.

Luis Saavedra, Head of Regulatory and Pharmacovigilance
Animalcare Group

Veeva RIM platform for animal health

Veeva RIM provides a unified, cloud-based platform that helps animal health companies align global teams, eliminate manual processes, and accelerate time to market. It is flexible, scalable, and supports compliance with industry requirements such as UPD (Union Product Database) submissions.



The screenshot displays the Veeva RIM platform's Marketed Drug Product Registration interface. The main content area shows a table of registered products, packaging, and active substances. The sidebar on the left lists various registration details and tracking options. The top right corner indicates 1 of 20 records in this list.

Name	Product	Product Variant	Manufacturer	Species	Withdrawal Period	Application	Registered Product Type
Antirabies solution for injection-Antirabies 1 ml solution for infusion-VeevaPharm - Barcelona-12	Antirabies solution for injection	Antirabies 1 ml solution for infusion	VeevaPharm - Barcelona	Adult male dog		VPA120288/011/001	Registered Vet Drug Product
Antirabies solution for injection-Antirabies 1 ml solution for infusion-VeevaPharm - Barcelona-33	Antirabies solution for injection	Antirabies 1 ml solution for infusion		Adult male dog	Antirabies solution for injection - 3 day	VPA120288/011/001	Registered Vet Withdrawal Period

Name	Active Substance	Manufacturer Name	Application	Regulatory Objective	Active Substance > Primary Product Family	Active Substance > Substance Name	Active Substance > Active Substance Type
Inactivated rabies virus-VeevaPharm - Brussels-12	Inactivated rabies virus	VeevaPharm - Brussels	VPA120288/011/001	001 - Initial MAA	Antirabies	Inactivated rabies virus	Substance

Name	Packaging	Packaging Site	Product	Product Variant	Finished Product Manufacturer	Application	Regulatory Objective	L
Antirabies 1ml bottle-VeevaPharm - Barcelona-12	Antirabies 1ml bottle	VeevaPharm - Barcelona	Antirabies solution for injection	Antirabies 1 ml solution for infusion	VeevaPharm - Barcelona	VPA120288/011/001	001 - Initial MAA	P

Dedication to customer success and innovation

Veeva takes great measures to remain current with the latest regulatory requirements and trends within animal health. The company maintains dedicated teams that are focused on regulatory intelligence and global regulation updates. Additionally, Veeva aligns closely with customers to help guarantee their success. Finally, Veeva is committed to innovation and provides new features and products three times a year to improve efficiencies in the regulatory process and ensure compliance with evolving animal health regulations.



Visit veeva.com/eu/rim/animalcare to hear how Animalcare Group improved their regulatory compliance with Veeva RIM.