Managing Veterinary Regulatory & Safety Information

RIM – Safety / PV

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Reduce errors managing your RIM data for different markets

Remove the complexity of getting the right information for the right market by letting the system guide you through what you need to provide.

Meet NVR requirements whilst maintaining corporate data

Ensure that submitted information aligns with requirements from your regulators and standard terms, without overhauling your corporate data.

Understand and control what is submitted where

Instantly view what has been approved in any given market with full traceability through global and local submissions and correspondence.

Track and manage safety compliance in a single system

Capture case data and the associated documentation in a secure, version-controlled manner with data vocabularies coming from the industry.

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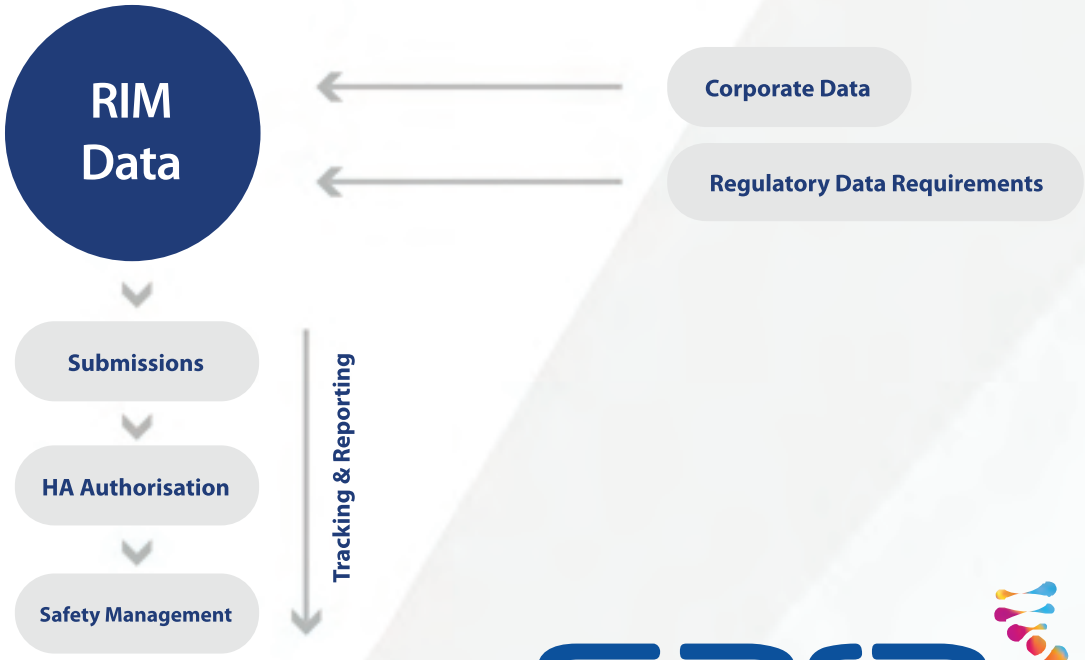
How do I understand and manage my regulatory activities?

CARA provides the optimal solution to the situation – allowing veterinary science companies to have a single UI for Regulatory, RIM, and PV that simplifies the business users' experience, with the data seamlessly connected to submission assemblies and individual documents, allowing full traceability, impact analysis and an easy Current Approved View of your products globally - all while meeting NVR requirements.

The lifecycle of veterinary products from early discovery through development and trials, submission and post-submission tracking, to updates and safety management is extremely complex.

At the heart of managing the process is a combination of the Master Data on products and the documentation associated with every step of the process. Traditionally, managing these has involved implementing multiple siloed systems with very little inter-system communication and thus data integrity.



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