Veeva

Regulatory, Quality and Product Claims Solutions for Specialty Chemicals

Gain agility, transparency and efficiency.

Specialty chemical companies are manufacturing ever more complex ingredients and products and substantiating stronger claims. They need better systems and processes to be able to manage critical studies, claims, regulatory and quality actions flawlessly and at speed.

Veeva provides intuitive cloud applications that digitally unify your data, documents and workflows enabling faster, more collaborative, error-free work within a single source of truth.

- Gain agility with real-time portfolio and pipeline management using dashboards and reports
- Simplify and streamline regulatory processes to accelerate business growth and compliance
- Continuously improve and harmonize quality across the enterprise and supply chain
- Strengthen claim substantiation and management to enhance portfolio value
- Enhance and protect study value with digital management tools

"Veeva Vault Platform helped us transform and speed up our regulatory processes. It streamlines product submissions and establishes a single source of truth across all stakeholders."

Dr. Landry Le Chevanton, Head of RAQM/Operations at DSM

DSM

Agility at the speed of your imagination

www.industries.veeva.com/chemical

Veeva's Solutions

Veeva's secure cloud solutions support the chemical industry and help bring innovative and safe products to market faster without compromising compliance.

CLINICAL	REGULATORY	QUALITY	CLAIMS
Veeva Clinical	Veeva Regulatory	Veeva Quality	Veeva Claims
Study Planning/Tracking CRO/Lab Management Test Reporting Study Data Management	Submissions Submissions Archive Formula Insights Registrations	Document Control Training Management QMS & Supplier Quality Risk Management	Claims Management Localization of Global Claims Challenge Response Oversight and Insights

Veeva brings agility, transparency and efficiency to:

Clinical

- Simplify the design and execution of GxP and non-GxP studies
- Collaborate seamlessly with internal and external partners
- Use templates, workflows and approvals maximize study relevance, compliance and value

Regulatory Affairs

- Global registration portfolios and pipelines
- Submission/assessment planning and workflow
- Regulatory review and approval workflow and control
- Regulatory authority communications and commitments
- Dossier and compliance document creation templates and workflow (FCN, REACH…)
- Underlying data and document management (SDS, studies, assessments, approvals…)
- Compliance document management (regulatory data sheets, SVHC declarations, certificates…)

Quality Management

- Document Control & Training Management
- Supplier Qualification & Management
- Customer Complaints
- Nonconformance Investigation & CAPA
- Internal and External Audit Management
- Risk Management
- Change Control

Claims Management

- Full detail on permissible usage by product, geography, and market
- Workflows for creation, substantiation, review and approval, with audit trails
- One click traceability from claim to substantiation and marketing assets using the claim
- Claim risk ratings with automated prompts and triggers for review workflows

"Veeva's reputation, proven cloud solutions, and commitment to meeting our industry requirements were important factors in our decision. Veeva will help us unify people, processes, and data across the organization."

Alexandre Gultzgoff, CIO of Seqens

SEQENS

Founded in 2007, Veeva NYSE (VEEV), is a global provider of cloud-based software solutions. We help quality, regulatory and commercial teams in the cosmetics, consumer goods, and chemical industries unify disconnected processes, documents and data. In doing so, we help our customers eliminate inefficiencies so they can bring innovative, high quality products to market faster without compromising compliance. Veeva is headquartered in the San Francisco Bay Area, with offices throughout North America, Europe, Asia, and Latin America.

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