Scigeniq™ RMS

Scigeniq RMS is an all-inclusive, automated system for planning, tracking, coordinating, and documenting regulatory submissions for life sciences companies.

Scigeniq RMS provides an extensive set of capabilities to manage all aspects of the regulatory submission process, from initial planning to submission and archiving. It is a flexible, easy-to-use application that can be configured to each organization's specific processes. With Scigeniq RMS, submission activities are planned, organized, and managed to ensure compliance with requirements. In addition, the system includes complete workflows for creating, reviewing, and approving submission documents within a CSV-compliant, validated environment.

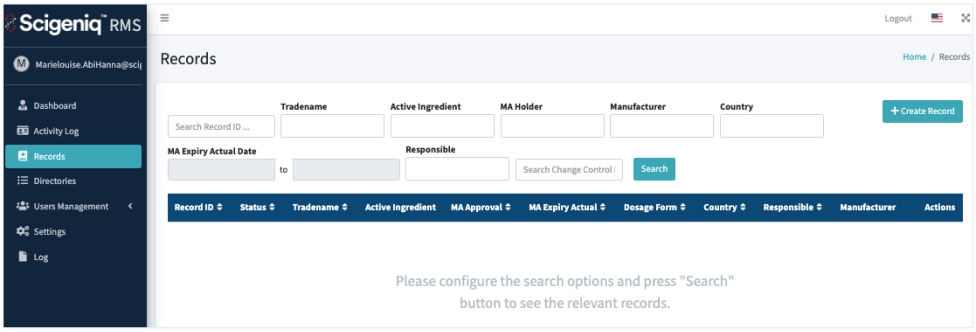
Beyond Managing the Documents

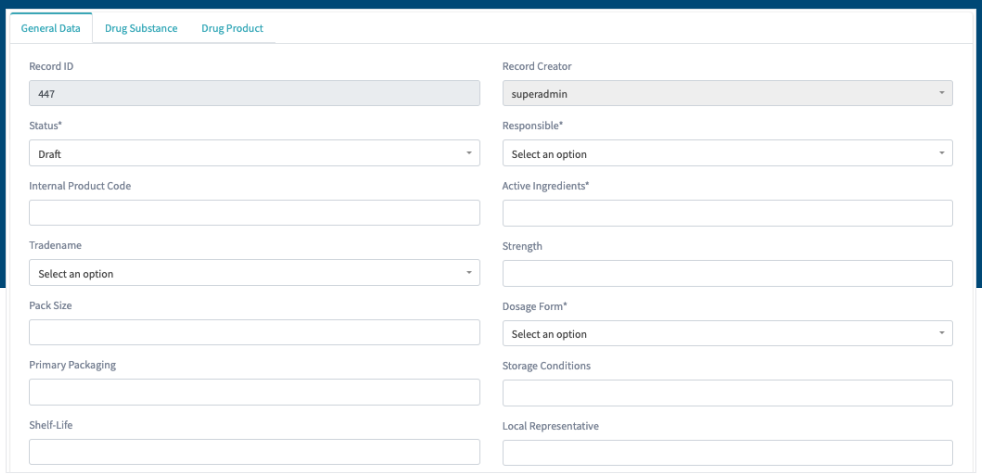
Many regulatory solutions for life sciences companies are based on traditional, legacy document management systems that focus only on the documents. Scigeniq RMS provides a complete planning and process management solution as well. The result is a complete system that tracks and coordinates all the activities involved in each submission, as well as all the documents. And Scigeniq RMS is part of the Scigeniq family of easy-to-use applications and is easily configured to align with your processes. The result is much faster implementation and ready adoption by users.

A Complete Regulatory Management System

Scigeniq RMS includes an extensive set of capabilities, including:

- Planning and Tracking Tool. Track every submission against the plan and schedule and provide notifications to everyone involved in the submission when things are due.
- International Registration Tracking. Creates records of every product's registration in every country and stores all related approvals and specifications.
- Change Control Management. Tracks Change Control approvals and activities and captures all changes (Variations, Renewals, Safety updates) in one centralized system.
- Complete Archiving Capabilities. Stores complete registration dossiers and allows easy access to all users for compliance check.





Part of the Scigeniq Family of Applications

Scigeniq RMS is part of the Scigeniq family of applications for managing quality and compliance at life sciences companies. Each application shares our commitment to delivering quality management applications that are much easier to use, much faster to deploy, and designed for mid-sized life sciences companies in the Middle East, Africa, and South America.

Our family of applications includes solutions for managing quality processes from deviation to CAPA right through change control, tracking training activities, and coordinating regulatory submissions and product life cycle management.

More Than Software

The Scigeniq team has deep, unparalleled knowledge of the pharmaceutical industry and quality processes. This expertise is embedded directly into our applications and provided to every customer throughout the implementation process.

That means that our years of expertise are available to you at every step of the process, and proven best practices are “baked into” our applications.

Scigeniq: We Deliver Technology-Based Advantages to our Customers

Scigeniq provides software and expertise to life sciences companies in the Middle East, Africa, South America, and around the world to accelerate their digital transformation. We provide expertise, guidance, and software solutions to our customers, which allows them to rapidly gain new capabilities and advantages based on proven technologies.

LEARN MORE:
Visit www.scigeniq.com
Email: info@scigeniq.com
Phone: +971 4 568 6972

Scigeniq™