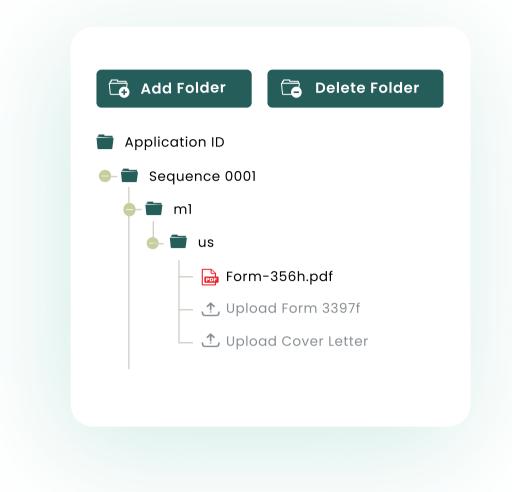


End-to-End Regulatory Affairs Software

What is **RIMS**?

RIMS, short for Regulatory Information Management System, is a centralized software platform that helps Pharmaceutical and Life Sciences organizations manage regulatory information for their products. It enables end-to-end tracking of Regulatory Activities and reduces efforts to obtain Regulatory Compliance.

The DYAZ RIMS Platform Offers



Template Creation

Pre-built Templates

Build submission plans using fully customizable inbuilt agency complaint eCTD folder structure

Smart, Reusable Content

Create variables in documents for easy and consistent reuse across submissions

Document Management

Collaborative Content Authoring

Create, edit, and share structured submission content without external documents or spreadsheets

Onedrive Integration

Open & Edit the document in Onedrive, accompanied by Bookmarking, Hyperlinking and eSignature.

Convert DOC to PDF

Render documents in the correct PDF format with auto-generated TOCs and appendices based on your content plan

	ize	Status		
19	5 МВ	Reviewed		Î
15	5 МВ	Draft	Downloo	ad
			Convert to PDF	
			Rename	e File
17–12–2023 19 ife Cycle General Ready for Pu				
'es				

eCTD Submissions

Comprehensive Lifecycle Management

Assign tasks, identify document lifecycle operations and navigate among files across all the sequences with the help of our integrated eCTD Viewer

Actions Required

App 12, Seq 0003 Overdue Revise the target date, contact RA

Lead at earlie	st
View more	01-05-2024 08:25 IST
	Document Properties:
	Name: cover-letter.pdf
	Created On: 27-12-2023 19:38 UTC
	Size: 168 MB
	Reviewer: Markus Winter
	Approver: Aaron Grant
	Lifecycle:
	Replace ~
	Link the Document:
	C Link

Submission Alerts and Tracking

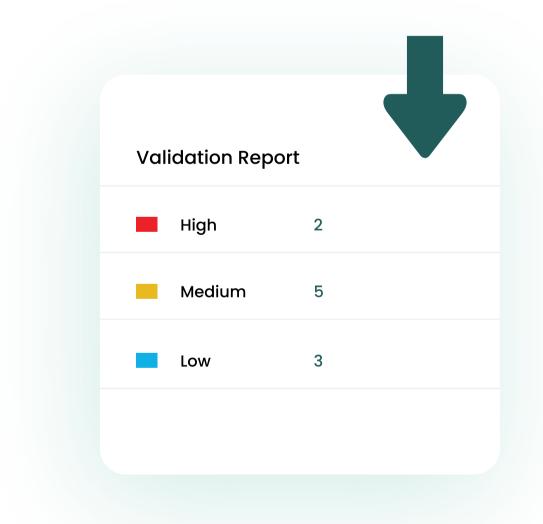
Receive automated alerts and notifications about your applications and never miss a deadline

Sequence Import & Export

Easy migration of sequences from an external platform and download submission ready sequences

Query Management

Track correspondence with health authorities and manage approval processes



Validations

Reduce Risk of Errors

Validate sequences with latest ICH Standards and Regional Guidelines, preventing technical errors and ensuring on-time approvals

Download Validation Report

Review technical errors and handle them within the system

Benefits of DYAZ RIMS

Cloud Storage



V Flexible Subscriptions

Agency Compliances



U.S. Food and Drug **Administation**





European Medicines Agency

GCC Health Council

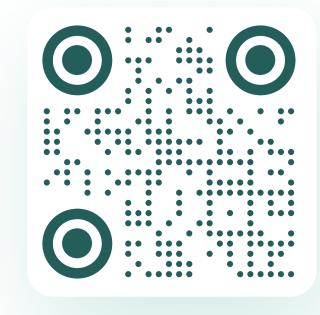
Contact Us

DYAZ INNOVATE PRIVATE LIMITED

+91 - 82000 - 84119

info@dyazinnovate.com

Hyderabad, Telangana, India - 500048



www.dyazinnovate.com