

Dossier Managements

The Compliance Risk of Manual Document Management in Pharma

- ▶ In the Pharma and Chemical industries, regulatory compliance hinges on timely and accurate document submissions.
- ▶ Yet many organizations still rely on folders, shared drives, and email chains to manage critical documentation like ANDA files, Lab Development Reports (LDRs), and deficiency responses.



Dossier Management Systems

- ▶ Missed deadlines, version confusion, and lack of traceability in interactions with global regulatory bodies. PMSoft's **Dossier Management System** streamlines regulatory submissions with structured workflows, enhancing speed, compliance, and accountability.

With our Dossier Management Module, you can:

- Digitally organize documents for ANDA, DMF, CTD submissions and more
- Manage Lab Development Reports and regulatory attachments with traceability
- Track submission status by country, region, or regulatory body



About Us

At PMSoft, we bring decades of expertise in Project, Program, and Product Management to help organizations transform the way they work. As trusted partners to the Pharma, Chemical, IT, and AI domains, we deliver tailored solutions in Training, Consultancy, and Digitalization that empower leaders and teams to achieve measurable results.



PMSoft
Character ▶ Integrity ▶ Results

Tired of Chasing Documents When the Deadline Looms?

Streamline Dossier Submissions and Responses

Contact Us : info@pmsoft.com

25+
Years of Experience

7500+
Executive Trained

6000+
Consulting Hours

45+
Customers Served

8+
Domains Served

