







"It's Not Software it's PaperLess GMP"



Cyclone Pharmaceuticals Pvt. Ltd. is an Indian Leading technical consultancy company with more than 15 years of quality experience in Pharma GMP & Regulatory affairs and hence can conclude on perfect development of software for Pharma industry.

Pharma industry till now is operating with traditional management of documents and there must be improvement with lack of technical & QMS knowledge many softwares failed to fulfill requirement of pharma industry.

Cyclone Pharmaceuticals Pvt. Ltd. in collaboration with PAB Internationals - A Medical softwares development company had developed key softwares for Pharma industry.

- **Repair Less GMP Software**
- **Recognition** celebrates and economic e
- ★ Digital Pharma HR
- Regulatory Affairs Software (Dossier/DMF)

OUR PRODUCTS

- **★** PaperLess GMP
- * Regualtory Dossier / DMF compiler
- ***** eCTD Conversion software
- * APQR & Trend Management
- **★ Vendor Management System**
- **★** Document Management
- 🖈 Pharma HR
- **★ Paperless QMS**
- **★ SOP Management**



WHAT WE DO

Our amazing features & services and others listed below.

Paperless GMP (Pharma Manufacturing)

Paperless GMP takes care of each step with dual authenication and checker is basic concept of the GMP and we covers the sames.



It is designed with high level of care at every point so that the chances of maual mistakes are minimized It manages all aspects of wholesale operations quickly and accurately...

eCTD Software

PAB Internationals has developed eCTD compilations and conversion Software. Cloud based software with highest security. Software is having reserved rights for admin & users.



Retail Medical Software

Pharmacy Software with Easy GST Billing & Filing. Have full control over inventory, billing & accounting, information about store items, suppliers & customers.

OPD/IPD Management

(Small Clinics)
OPD / IPD covers Appointment,
Reception, OPD ,IPD, TPA etc.
Hospitals and nursing to manage
and moniter patient data.

and

CYCLONE PHARMACEUTICALS PVT.LTI

Digital HR

A Payroll system is software designed to organize all the tasks of employee payment and the filling of employee taxes.

B2B for Pharma

Pharma Business Portal for Buyer & Sellers of Medicine and Pharmaceutical products and medical Devices.

VeriPharma Software

Indias First Exclusive Pharma Recruitment Portal with Employee Verifications System.

Regulatory Software

The hotel and restaurant manegment degree prepares students for supervisory position in hostels ,restaurants, and other hospitality-related orgnization..

Pharma ERP

Inventory Management, Material Management & Distribution Management Software for Pharmaceutical Management.



PaperLess GMP



First time in world
PaperLess GMP Digital
platform for Pharma QMS / Documentation





INTRODUCTION TO PaperLess GMP

A paperless GMP manufacturing operation sounds counter-intuitive. Almost all operations use paper forms especially pharmaceutical manufacturing.



With growing future and established documentation system CPPL decided to go one step ahead. Till now many companies have developed ERP systems for Pharmaceutical Industries but these ERP systems are identical with it's name its Pharmaceutical Entrepreneurs resources system means management information system.

CPPL Decided to change the concept and developed a GMP Software which will really take care of your all systems. It's validated GMP Software and have inbuilt quality systems.

It's Not EPR It's PaperLess GMP

You are now free from paper and you have to just implement a advanced digital pharma ERP concept we called it as Paperless GMP.

The Advanced ERP which we are providing is Paperless GMP system where from Procurement to Dispatch you will get accountability for everything along with compliance to the Current Good Manufacturing Practices.

The Paperless GMP System is Compliant to all guidelines of the world.

A unique and Novel research by Minds of Pharma and Software Technocrats.



FEATURES OF SOFTWARE



Preventive Maintenance

User Authentication

System Compliant

Regulatory Compliant

→ Validated Software

QMS Enabled

Fast Authentication

Perfect Communication

Inline with existing system

Auditors Login

SOP & Flow charts

- APQR Preparation

Equipment History

→ Validation management











GMP Software Pvt. Ltd.

PaperlessGMP Software



Powered by Cyclone Pharmaceuticals pvt. Ltd.

LOGIN

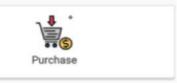


Management



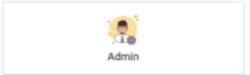




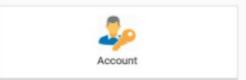


General Administration









Technical









Quality System









Dispatch



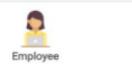














Quality Control Department

Dashboard

Testing

- Testing Allocation
- Testing

Operations

Training Needs

New Sampling

Testing

A. R. Report

Specification

■ Method of Analysis

Log Books

- Spec Report
- COA Report
- √ Equipment Usages Log
- **Equipments**



















Stability Study



Log Books



Incident



Control Sampling





How Electronic Operations improves efficiency in Human Resources



GUIDE TO
PAPERLESS HR

FEATURES OF Pharma HR

- Digital Interview , Offer letters, Appointment letters
- Attendance & Payroll
- Leave Management, Shift Management, Labour Management
- Notice & Warnings, Daily Reports
- Induction Training sessions
- Medical Checkup, Apprisal systems, Salary Slip
- Job Descriptions, Job Responsibility
- Performance Monitoring





Your Way To People Success













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REGULATORY SOFTWARE (Dossier & DMF Compilation)











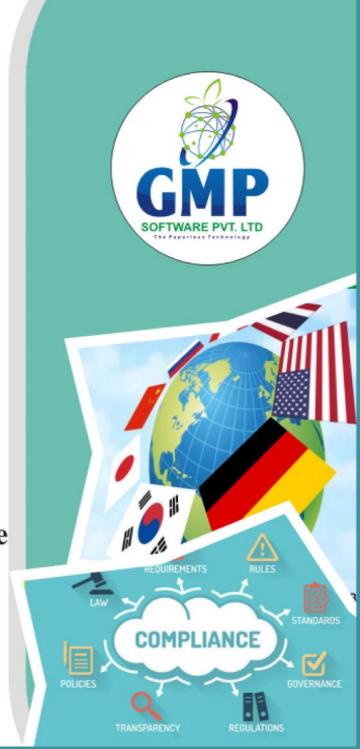
About Software

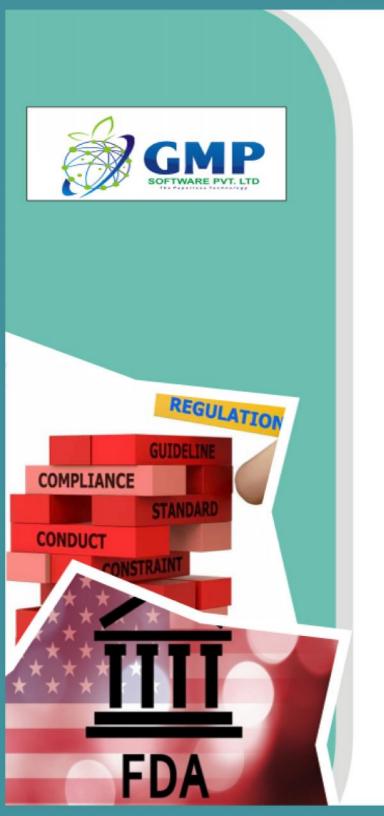
Cyclone Pharmaceuticals Pvt. Ltd. has started development of regulatory Dossier & DMF Compilation Software in 2016.

With help of CPPL Technocrats and Expert Industrial Professionals CPPL got success to in Development of Regulatory Software for Dossier and DMF Compilation.

This software is designed with high level of care at every point so that the chances of manual mistakes are minimized.

This software is 21 CFR compliance across IT subsystems





Features of Software

Department wise Access

Document managements with department user

Product wise document uploads by User

Admin and user login & rights are separate

Dossier Compilation for ACTD, CTD all regions

All regions Documents Checklist access

Automatic PDF file generation

Auto Dossier Compilation as per specific country guideline

All guidelines and registration guidance for all CIS countries, ROW market



POLICIES

REGULATIONS

STANDARDS

Contact Details

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COST . QUALITY . COMMITTMENT