
AUDREE®
Turn ideas into reality
www.audreeinfotech.com
CERTIFIED
ISO
9001:2015
COMPANY

ABOUT US

Audree Infotech is a professionally driven IT solutions provider offering a diverse range of products and services aimed at optimising customers' processes and productivity for maximising growth and efficiency. Backed by a team of experts specialised in planning, design, development, integration, and deployment of diverse IT projects, we deliver the most robust, scalable and secure solutions which inspire operational excellence in the realms of data, quality, accounting and security among others. Armed with the capabilities to deliver to the needs of diverse industries, Audree specialises in offering niche solutions to Life Sciences industry, which operate in a stringent regulated environment.

QUALITY POLICY

Audree Infotech is committed to ensure the highest degree of customer satisfaction through best professional practices in delivering software services.

We aim to be the leaders in providing the software services, using the state-of-the-art software development, information security & professional services, adopting quality management systems of national & international standards complying with customer & legal requirements.

Quality objectives are defined at relevant functions, levels & processes.

We strive for excellence through continual improvement with a team of trained, dedicated & competent professionals.

WMPS: Warehouse Management Portal System

WMPS is Used for materials creation, receipt, request, quality inspection, issuance and dispatch of product. Our product has been designed to establish a good system through a defined procedure wherein paper-work is reduced and manual Errors/Interventions are minimized

VMS: Vendor Management system

Vendor management system helps minimizing errors while dealing with multiple vendors which may cause delay in business operations due to late shipping or other issues. Vendor management system helps to get a comprehensive view of all the financial transactions. It optimizes the performance, increases administrative efficiency and increased on boarding speed.

UMS: User Management System

UMS is used for the management of individual identities, their authentication, authorization, roles and privileges within or across system and enterprise boundaries with the goal of increasing security and productivity while decreasing cost, downtime, repetitive tasks and paperless work

WMS- Work Order Management System

WMS enables you to quickly create work orders and track progress thorugh a single dashboard. WMS helps maintenance managers reduce paper-work, track and prioritize work orders, and ensure all of them are completed on time. WMS helps keep a maintenance record to eliminate recurring issues.

LMS: Learning Management System

LMS helps in real-time visibility into the performance of employee training programs with comprehensive training management. It schedules, tracks and reports on training, certifications and qualification courses for employees or workgroups.

QR Code for API

QR Code for API will help you to generate QR code for all your API export requirments. As Govt of India has mandated QR on Product label, Our QR code application will help you to run your process smoothly.

CMS : Change Management System

It is a process to ensure that changes to a product or system are introduced in a controlled and coordinated manner. Our Product has been designed to establish a good system through a defined procedure wherein paper work is reduced and Manual Intervention/Errors are minimised.

CAPA : Corrective and Preventive Action

CAPA focuses on the systematic investigation of discrepancies (failures and/or deviations) in an attempt to prevent their recurrence. Our product has been designed in line with FDA requirements emphasising on Root Cause Analysis, Investigations and an Action Plan to correct the non-conformance

LIR & AER: Laboratory Investigation Report and Apparent Error Report

LIR application is an electronic system to document every laboratory incident at one place. LIR makes it easier to track incidents and take timely actions. This easy-to use application with a simple UI makes it simpler for user to initiate, review, assess, approve and investigate incidents.

IMS: Incident Management System

This system is used to report any unplanned action or unidentified outcome which is differed from the predefined standards by further investigation through the established procedure.

OOS : Out of Specification

It is defined as a result that falls outside the predetermined specifications or established acceptance criteria set by the manufacturer and/or the laboratory. Our Product has been designed to establish a good system through a defined procedure wherein paper work is reduced and Manual Errors/Intervention are minimised

Market Complaint

In pharmaceutical industry, Market Complaint indicates customer dissatisfaction regarding quality of product, received verbally or in written format. Our Product has been designed based on type and criticality in line with FDA requirement

APQR: Annual Product Quality Review

Our APQR product is used to track the total Information of the particular product during a year/ defined schedules to improve/check the Quality of the Product by defining the Controls or action plans

DMS: Document Management System

DMS provide storage, versioning, metadata, security, as well as indexing and retrieval capabilities electronically and reduce manual paper work. Our product is capable of keeping a record of various versions created and modified by different users (history tracking).

SDMS: Scan Data Management System

Our SDMS product provides tools for scanning, indexing, searching and viewing documents.

RIMS: Regulaotry Inofrmation Management System

RIMS is used to facilitate the information related to product registrations in different Countries/Regions by regulatory affairs.

Errata

Errata helps fix common typographical errors in documents. With the use of Errata companies don't have to go over the entire document review process and manage change control ensuring data integrity.

ELMS: Electronic Log Management System

This system is used to Log usage and cleaning activities pertaining to Area and Equipment electronically. ELMS application also contains recording of product dispensing and weighing details, which will help to make error free entries for better review, monitoring, control & compliance

IMS: Intervention Management System

This system is used to replace manual recording of interventions in batch manufacturing record process, facilitates online access to perform intervention activities in processing areas, and enhances the compliance

APPLICATIONS

QUALITY APPLICATIONS

CMS : Change Management System

CAPA : Corrective and Preventive Action

LIR & AER : Laboratory Investigation Report and Apparent Error Report

IMS: Incident Management System

OOS: Out of Specification

Market Complaint

APQR: Annual Product Quality review

DMS: Document Management System

SDMS: Scan Data Management System

RIMS:Regulaotry Inofrmation Management System

Errata

MANUFACTURING

ELMS: Electronic Log Management System

IMS: Intervention Management System

PWO: Process Work Order

GWO: General Work Order

Artwork Management System

INVENTORY

WMPS: Warehouse Management Portal System

BUSINESS PROCESS AUTOMATION

VMS: Vendor Management system

SECURITY AND ADMINISTRATION

UMS: User Management System

RAMS: Remote Access Management System

TRAINING

LMS: Learning Management System

OTHER PRODUCTS

Pharmacovigilance Automated Alert System

Label Printing Management System

Analytical Research & Development

Intervention Management system

Sample Tracking Management System

QR Code with Product Label

SERVICES

CSV

We provide a cost effective framework of good practice to ensure that computerized systems are fit for intended use and compliant with applicable regulations. We adopt life cycle approach for validating the computerized system and provide technical expertise starting from Risk/Gap Assessment until Summary Report that complies with regulatory requirement.

Application Support

We have a dedicated team in place to extend support for .Net applications development and maintenance. Our team is backed by a proven track record for fast-tracking the implementation, migration and adoption of SAP solutions for clients in line with regulatory requirements. Our team is also expertise in troubleshooting IT applications process related issues and can provide recommendations for solutions in consultation with relevant

SAP Services

Our team of experts have implemented SAP for global customers, to transform their businesses. We possess talented workforce with business process expertise, capable to re-engineer the processes and to increase the operating efficiencies. We aim to offer value by re-defining the complex business processes to simplify and achieve the best results to comply with regulatory requirements.

Training Services

We conduct regular trainings and workshops on variety of topics related to cGMP, Regulations & Guidelines pertaining to Computerized System Validation. We offer excellent programs and interactive workshops to help pharmaceutical companies align with the changing regulatory requirements.

CORPORATE OFFICE

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